THE U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES PUBLIC HEALTH SERVICE

CENTERS FOR DISEASE CONTROL AND PREVENTION NATIONAL INSTITUTE FOR OCCUPATIONAL SAFETY AND HEALTH

convenes the

TELEPHONIC WORKING GROUP MEETING

ADVISORY BOARD ON

RADIATION AND WORKER HEALTH

ABRWH WORKING GROUP MEETING

The verbatim transcript of the

Meeting of the Advisory Board on Radiation and

Worker Health held telephonically on April 20, 2006.

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TRANSCRIPT LEGEND

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PROCEEDINGS

(11:00 a.m.)

WELCOME AND OPENING COMMENTS DR. LEWIS WADE, DFO

1	MR. GRIFFON: I think it's probably 2:00 p.m.
2	eastern time, right, Lew? I figured we could
3	do from now until 1:00 and then break for lunch
4	at 1:00.
5	DR. WADE: Okay. And then
6	MR. GRIFFON: And then pick up Rocky at 2:00
7	hopefully.
8	DR. WADE: Okay.
9	MR. GRIFFON: That's the tentative plan anyway.
10	DR. WADE: Okay.
11	MR. GRIFFON: That'll work.
12	DR. WADE: That's the plan.
13	MR. GRIFFON: All right. Okay. Thanks a lot.
14	DR. WADE: Okay. Thank you.
15	MR. GRIFFON: Bye.
16	DR. WADE: Okay. Well, I guess we have Mark
17	with us, Mike, Wanda, Ray. I think that's most
18	of what we need so maybe we can begin. This is
19	Lew Wade and I have the the pleasure of
20	serving as the designated federal official for

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the Advisory Board. And this is a meeting of the working group of that Advisory Board. working group has looked at many issues including individual dose reconstruction reviews, site profile reviews, procedures reviews. Recently flowing from the working group's efforts with regard to the site profile reviews for Y-12 and Rocky Flats the Board asked that this working group continue and look at SEC-related issues with regard to Y-12 and Rocky Flats. There have been a number of very productive calls of this working group and today we're meeting to discuss two issues, Y-12 as -- as Mark had mentioned and then followed by Rocky Flats. I would like to just take a brief moment to have the Board members identify themselves. I know Mark, Mike and Wanda are on the call. Are there any other Board members on the call?

(No response.)

DR. WADE: Okay. Just checking to see that -that we don't have a quorum. What I would like
to do is to go through our -- our conflict of
interest discussion. Let's have it relative to
Y-12 and then we will repeat that discussion.

1 Hello? 2 (Brief interruption) 3 DR. WADE: Somebody's at an airport getting 4 ready to board at Gate 43. 5 UNIDENTIFIED: Maybe I should sign off. Maybe 6 I'm too distracting. DR. WADE: I think you're right. Yeah, I guess 7 8 it would be good. 9 **UNIDENTIFIED:** Okay. 10 UNIDENTIFIED: We're not going to be able to 11 hear him. 12 DR. WADE: All right. Yeah. Okay. We're back 13 to it. We'll go through and have Board 14 members, the NIOSH team, the SC&A team identify 15 themselves on the call and any conflicts they have relative to Y-12. And then we'll go 16 17 around and let other government folks identify 18 themselves and anyone, petitioners and anyone 19 else who would like to be identified as being 20 on the call -- on the call. So I'll start. 21 I'm Lew Wade and I work for NIOSH and I have no 22 conflicts relative to Y-12. How about Board 23 members. Mark? Mike? 24 MR. GIBSON: This is Mike Gibson. I have no 25 conflicts.

1	DR. WADE: Wanda?
2	MS. MUNN: Wanda Munn. No conflicts.
3	DR. WADE: Okay. Mark, are you with us?
4	(No response)
5	DR. WADE: Okay. We'll listen for Mark. We'll
6	re-establish contact. How about the NIOSH ORAU
7	team?
8	MR. RUTHERFORD: This is LaVon Rutherford of
9	NIOSH. I have no conflicts with Y-12.
10	DR. NETON: This is Jim Neton. No conflicts.
11	DR. WADE: The ORAU team, please introduce
12	themselves.
13	MR. KENOYER: This is Judson Kenoyer, no
14	conflicts.
15	MR. TANKERSLEY: Bill Tankersley, no conflict.
16	MR. KERR: George Kerr. I have no conflicts.
17	MR. CHEW: Mel Chew. I have no conflicts.
18	MR. MCFEE: Matt McFee. No conflicts with Y-
19	12.
20	DR. WADE: Anyone else from NIOSH ORAU?
21	MR. SMITH: Yeah, this is Matthew Smith. No
22	No comments, or conflicts, rather.
23	DR. WADE: Okay.
24	MR. SUNDIN: Dave Sundin. No conflict.
25	DR. WADE: Other NIOSH ORAU?

1	(No response)
2	DR. WADE: Okay. SC&A.
3	DR. MAURO: John Mauro, SC&A. No conflicts.
4	DR. MAKHIJANI : Arjun Makhijani, SC&A. No
5	conflicts.
6	MR. GRIFFON: Hi, Lew. It's Mark Griffon
7	again.
8	DR. WADE: Okay. We're just going through a
9	conflict identification, Mark.
10	MR. GRIFFON: Okay.
11	DR. WADE: You could do yours.
12	MR. GRIFFON: Okay.
13	DR. WADE: Relative to Y-12.
14	MR. GRIFFON: Relative to Y-12 I only have a
15	conflict in changes where (inaudible) Labor
16	Council, HELC (unintelligible), is the named
17	petitioner.
18	DR. WADE: Okay. We were continuing then with
19	SC&A. Anyone else?
20	MR. BUCHANAN: Ron Buchanan. No conflicts.
21	DR. WADE: Anyone else from SC&A?
22	(No response)
23	DR. WADE: Okay. Without the need for conflict
24	identification, are there any other federal
25	employees on the line?

1 MS. HOWELL: This is Emily Howell with HHS. 2 have no conflict. 3 MR. RAFKY: Michael Rafky also with HHS. I 4 also have no conflict. 5 DR. WADE: Any petitioners or representatives for Y-12? 6 7 (No response) 8 DR. WADE: Okay. I open up to anyone else who 9 would like to identify themselves as being on 10 the call. Not necessary, but if you'd like, 11 please. 12 MS. FRANK: Laura Frank from the 13 (unintelligible). 14 DR. WADE: Welcome. 15 Thank you. I'll probably hang up MS. FRANK: 16 and then come back when you all attend to the 17 Rocky Flats. 18 DR. WADE: Okay. Thank you. 19 THE COURT REPORTER: I'm sorry. This is the 20 court reporter. Could I get your name again, 21 please? 22 Laura, L-A-U-R-A, Frank, F-R-A-N-K. MS. FRANK: 23 THE COURT REPORTER: Okay. Thank you. 24 MS. FRANK: You're welcome. 25 DR. WADE: Anyone else who would like to be

1 identified? 2 MR. LAWSON: Howard Lawson and Larry Jones, 3 Labor Council at Y-12. 4 DR. WADE: Okay. Mark, back to you. 5 MR. GRIFFON: Okay. I guess we -- you're getting ready to start the -- the agenda, Lew. 6 7 DR. WADE: Correct. 8 MR. GRIFFON: I missed a few minutes, so okay. 9 DR. WADE: We just did introductions --10 MR. GRIFFON: Yeah. 11 DR. WADE: -- and we talked about quorum issues 12 and things like that. 13 Y-12 14 MR. GRIFFON: I think the best way to proceed 15 here -- I'm almost ready to get off my cell 16 phone and onto a hard line so I apologize for 17 that. But I think the best way to proceed is 18 probably to start with what Jim had provided. 19 I think Jim included most of the outstanding 20 actions that we had in the matrix as from 21 NIOSH's standpoint anyway. And I think maybe 22 Jim can give us an overview of that and then we 23 can start into the SC&A's review report of --

of the evaluation report if that -- if that

makes sense. And if Jim -- I assume Jim is on

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1 the line? 2 DR. NETON: Yeah, I am. I'm on the line. 3 MR. GRIFFON: All right. 4 DR. NETON: I'm going to have to scramble and 5 sort of re-- recall from memory what I sent 6 out. 7 MR. GRIFFON: Okay. 8 DR. NETON: I thought we were going to go 9 through the report but --10 MR. GRIFFON: I guess -- I guess it really 11 doesn't matter which order. I thought that 12 that would be the easier thing to -- to get a handle on but --13 14 DR. NETON: I think I can do it. Just give me 15 a second here to --16 MR. GRIFFON: Okay, sure. 17 DR. NETON: The -- The issues -- The items 18 that I -- that I sent out which I think -- and 19 I think Mark is correct -- I did believe at 20 least we -- we were responsive to the closing 21 out the issues, you know, that were for you to judge whether they're sufficient to close it 22 23 out, but we sent out the remaining dose 24 reconstructions. Those were for polonium, 25 plutonium, an extremity dose as well as there's one other in there.

MR. GRIFFON: One of the exotics? Is that one of the --

DR. NETON: Nuhytrogalian (ph) 67.

MR. GRIFFON: Yeah.

Thanks. That's correct. And so DR. NETON : those -- those have been -- been put out there. We also put out a table that compared the -the databases from the CER for uranium urinalysis versus the distribution of the data that we observed in the uranium samples that were in the delta view database. remember, we determined that those uranium samples were not in the CER database and yet the issue was would those samples, if they were added to the CER database pollute the co-worker model to where it would not be an accurate depiction of what the exposures were. think the table is fairly self-explanatory in that the -- the -- the delta view data actually end up having a lower -- the distribution would end up lowering the results for the uranium urinalysis logs so therefore we don't believe there is a significant effect on our co-worker model that was developed from the CER data.

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1 There was another issue --2 MR. GRIFFON: I don't -- I don't know if we 3 have to comment on these but if SC&A, if you 4 guys have any comments on these, you know, or 5 need clarification on any of these items I 6 think it's probably appropriate to sort of discuss it. 7 8 This is John Mauro. I think that a DR. MAURO: 9 lot of the items that were covered by Jim we've 10 sort of taken the next step forward in our 11 evaluation report. Those items will -- some of 12 those items will be revisited at -- at the next tier so to speak during our discussion of our 13 14 draft evaluation report that went out yesterday 15 and that I presume most folks on the line have 16 copies of. 17 MR. GRIFFON: I think so, too. That's why I chose this order because I think, yeah, it 18 19 makes sense to -- all right. 20 DR. MAURO: Okay. 21 MR. GRIFFON: Go ahead, Jim. I'm sorry that I 22 cut in there. 23 DR. NETON: That's fine. And then -- then 24 there -- there was an item I sent out that 25 dealt with the discussion of 1951 data that

1 appeared in delta view versus what was in the 2 CER database and we put that out, about a page 3 and a half document. And I'm very certain that 4 SC&A commented on that in their review so we'll 5 get into that later. MR. GRIFFON: Yeah, we'll probably cover that 6 7 later, right. 8 DR. NETON: And then I also sent out a -- a 9 criticality -- a draft criticality -- a draft 10 report on criticality incident that occurred in 11 1958 that sections of, we believe, substantiate 12 the reasons why not all workers were monitored 13 at criticality incident and why is that not an 14 indication that, you know, the highest exposed 15 workers were monitored. That went out fairly 16 recently. I think that's -- that's -- that's 17 all the information I sent out. 18 MR. GRIFFON: Yeah, I think that does cover it. 19 DR. NETON: And all the --20 MR. GRIFFON: I think given that -- the last --21 well, most of the items as John said are going 22 to come up as we go into the review report so 23 if the -- unless there's any other questions or 24 comments or clarification by Jim I think we're 25 probably ready to go right into John's -- into

1 your report.

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DR. MAKHIJANI: Yeah, Mark, this is Arjun.

MR. GRIFFON: Yeah.

DR. MAKHIJANI: I don't think the incident list was part of the matrix -- the incident list with the exotics was part of the matrix but --MR. GRIFFON: Yeah. Jim, there was -- I -- I thought, and -- and again we didn't have to come back to and I know -- I know there's -we've -- we've done a lot of these calls so there's a lot -- a lot of work there but I thought that you had mentioned as part of the exotics dose reconstruction that there was -there was incident data that you were going to be calling on for the dose reconstructions related to the exotics. And I don't know if you -- if that is on the O-drive or if you intended, you know -- I guess that's --DR. NETON: No, we -- we can get into that maybe when -- when we get to that issue but I -- I didn't recall if the incident list was one of the closeout items in the matrix. But we -we do intend to rely on incident reports that we know are present, particularly on the delta

view system and there are over 4,000-something

images out there. Frankly we just have run out
of time to be able to catalog all those. We
just -- we know that there are -- there are a
lot of them out there and the ones that we
sampled definitely allow us to do dose
reconstructions. And that was the one intent
of the gallium example but --

MR. GRIFFON: Oh, okay. Okay.

DR. NETON: -- we didn't have time to
distribute the -- a complete compendium of all
the incidents. It would be -- it would be
quite an undertaking to do that.

MR. GRIFFON: Okay. Okay.

DR. MAURO: This is John Mauro. Along those lines while we are discussing this, Arjun and I have had an opportunity, of course, to discuss a lot of these matters before this call. With regard to the incident reports, one of our observations as we're talking about it is that the gallium report I guess represented a later time period. As an example problem or maybe a couple of example problems I think we're basically looking for kinds of information in the incident reports that are available during the earlier years for some of these exotic

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DR. NETON:

radionuclides. I guess just to provide an example that shows here's typically the kind of data that we have available to us in the reports and how we would use that data to reconstruct. Right now I guess you felt the gallium was an example that was more of a -- a later time period if I'm correct. And I guess just so that I can close the loop on the -- on this is I guess a little more reassurance that, yes, even though in the earlier years when these exotic radionuclides were handled and there were incidents, the kinds of information that are available in those numerous incidents reports by and large give you the information you need to reconstruct the inhalation doses. MR. GRIFFON: But Jim -- Jim, did you say a -a -- I -- I might have misunderstood this but are the -- are the 6,000-page images or the images that we have from the delta view database, do they include some of these incident reports that you're discussing or is it another part of the delta view database? DR. NETON: I don't know that they do, Mark. MR. GRIFFON: Okay.

That was not what we pulled the

database for at that point.

MR. GRIFFON: Right.

DR. NETON: The delta view database is searchable by certain key words and fields and when one searches the delta view database for investigation slash incidents, you end up with about 4,000 images that are -- that are resident. And that was the intent of the delta view database was to consolidate all these -- these reports and such into one -- one central data system. We just have not had the time to pull --

MR. GRIFFON: Yeah.

DR. NETON: -- all of these out and comb
through them although again we believe that
every indication that we have are that they are
there, available and we could use them. And
there's -- there's other pieces of information
that we'll be bringing to the table to
demonstrate how we can do exposures for the
Cyclotron but I don't know if we want to do
that now or wait until we get to the relevant --

MR. GRIFFON: Probably wait until we get to their report but I -- I just, yeah, just to --

I just wanted to clarify that we -- that it

wasn't in what we had so okay. So that's -
that's understandable. Okay. Anything else

John or Arjun or should we -- Should we start

into your review report?

DR. MAURO: Yeah, this is John. You know, we might as well get started.

MR. GRIFFON: Yeah.

DR. MAURO: I will make a couple of prefatory remarks before I hand the baton over to Arjun who did the heavy lifting. One of these -- the -- in our report I can't say for certain that we've captured everything that came across in the -- on the email from Jim. We were certainly attentive to the material as it came in, certainly the example problems, but I'm not quite sure whether we -- how we reflect all of the material that has come through as of the time that we -- that we sent out our report. So we may be a little bit behind the power curve in terms of capturing everything that Jim has provided. The second point I would like to make is that you may have noticed that we have not yet addressed the recycled uranium piece. There is a placeholder in our report that we

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are close to finishing up, and our intention is that after this conference call and after we sort of regroup we'll probably issue a revised version of the report to sort of catch up on those pieces of material that we have not captured, address the recycled uranium issue; and there's one more point that I feel needs to be incorporated. I think our report in general zeros in on all of the areas that we feel there are deficiencies that need to be dealt with. also feel that we probably need to incorporate some material in our report in areas where we feel the case made by NIOSH is especially strong. Right now there is -- there really is very little of that. Now, the reason I say that is I think it's important for the Board to get a sense of giving the -- the issues and the time periods of concern to -- to somewhat get a bird's eye view of in the grand scheme of things where -- where is the evaluation report strong in terms of making its case or has made its case and areas where we feel it's weak and there are some problems that need to be addressed. Right now I think our report really zeros in on the problems but doesn't help the

Board too much in terms of letting them know where we feel it's relatively strong. We're -- Our intention is to -- to issue a next draft of this report as soon as possible and -- and address many of the -- these -- these matters that I'm describing. With that as a preface I'd like to hand it over to -- to Arjun to go through the -- the major points that we -- that we have made in our -- our review of the evaluation report.

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DR. MAKHIJANI: Thank you, John. The --John and I talked this morning about some gaps and one of the -- I'd just like to preface what I'm saying about -- with a description of a couple of those gaps. We didn't review the plutonium dose reconstruction. It came in on Monday and I think I was a little too overwhelmed to review new material since it was typeset on Tuesday and Wednesday. And the other -- The other thing is that in reviewing the 147 worker data I -- I focused on table 45-B but not on table 45-A and in going back I felt that the workers at Y-12 seemed to broadly have been sorted into two large bins, low and -relatively low and relatively high as reflected

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in post-61 data. And -- And that -- that overall idea there needs to be included in that evaluation of that model. I -- I don't believe that any other bottom line comments would change but I think it will better reflect what -- what NIOSH has done. So I just -- I just wanted to give the working group a little bit of an idea of a couple of things that John and I had discussed before this call. said, the -- we -- I went through -- there was a team of people that worked on the report. Hans is unfortunately not on the call. and -- and Ron Buchanan worked on the external dose stuff. I worked with John and Joe and Kathy on various parts of this report and as we -- so let me go -- there's one finding or one comment on uranium with trace thorium where I forgot to write a conclusion paragraph in the text of the report so it didn't get pulled up into the summary. I'm sorry about that. It will be there in the final report. So start at the top. Our main finding in regard to the SEC evaluation recommendation about thorium workers was that we agreed with NIOSH that there's not enough data to reconstruct doses for workers

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who were exposed or potentially exposed to thorium or should have been monitored for thorium during the SEC petition period. And we did some research. Kathy Demers did some research on buildings where whether the buildings covered in the evaluation were -were the only ones and -- and we found evidence, documentation that there were probably other buildings where thorium was processed we think in the '50s. I want to preface -- qualify this by saying, you know, that we researched this very rapidly obviously -- but I've listed the buildings there under heading two in the summary where thorium also appears to have been processed. Whether it was always processed in the '50s there I think may remain to be determined but this is the best of our judgment.

DR. MAURO: Excuse me. This is John Mauro.

Just quickly, just to help orient, I don't know if everyone is looking at the same page but page 1 in our report at the very top says attachment one. For the purpose of this discussion it's probably convenient if you folks have not already surmised this that we

1	have prepared we have listed a number of
2	findings and and Arjun is basically going
3	down items one, two, three, four, so forth in
4	that summary of principal conclusions. So that
5	may help a little bit for
6	DR. MAKHIJANI: Thank you, John. Yeah, I'm
7	sorry. I apologize. I should have said that.
8	DR. MAURO: Yeah, just to yeah. So we'll
9	just be going through that and, of course, each
10	one of these principal findings, the main body
11	of the text gives the rationale behind it.
12	DR. MAKHIJANI: We looked at the internal and -
13	- and the CER database validation in the
14	internal and external
15	DR. NETON: Arjun?
16	DR. MAKHIJANI: Yes?
17	DR. NETON: Mark, do you think it would be
18	better if we did these one by one or if we just
19	wait until all the issues have been discussed?
20	I mean it's up to you but
21	MS. MUNN: This is Wanda. I'd prefer we did
22	them one by one, frankly.
23	DR. WADE: Is Mark on the call?
24	(No response)
25	DR. WADE: Oh, we lost Mark.

With the

1 MS. MUNN: I think he must be moving from one 2 phone to the other again. 3 DR. MAKHIJANI: It would be fine by me to -- to 4 go one by one if that's the most convenient --DR. MAURO: I think that is. It keeps the 5 story a little bit more continuous. 6 7 DR. MAKHIJANI: Okay. 8 DR. MAURO: I would also recommend one by one. 9 DR. WADE: Okay. So let's do that. Arjun has 10 gone over points one and two. Jim, do you want 11 to respond? 12 DR. NETON: Yeah. I -- I think so. issue of thorium I mean we're -- we're 13 14 certainly gratified that SC&A agreed with our 15 position that thorium could not be 16 reconstructed although we're a little perplexed 17 at the -- at the issue raised that these other buildings are involved. Even though I think 18 19 the report states something to the effect that 20 there's ample evidence or significant evidence 21 that it was processed at other buildings, the 22 only citation I could find that -- that they 23 relied on was out of this Chem-Risk report that 24 -- that says starting in the early 1950's the

Y-12 thorium began processing its weapons

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components. And then they go on to cite the buildings. We -- We take no issue with the fact that production occurred, you know, significant production operation that started in the late '50s in our opinion, or early '60s did occur in those buildings but I -- I -- I scoured the entire 490 pages of the Chem-Risk report and found no other indication as to where that information starting in the '50s came from. It's an un-cited text. They just reference it. So it doesn't seem to be a strong piece of evidence. We have relied on reports directly from Y-12 personnel. are specifically several reports that we've cited that state that the thorium operation started in the '60s. So, you know, we don't take exception to the fact that those buildings that are cited in the Chem-Risk report were where major productions occurred. literally scoured hundreds and hundreds of pages of health physics reports and frankly had a lot of trouble coming up with the buildings that we did. We're not even among ourselves sometimes convinced that those buildings had huge exposures. But -- So I -- I don't know

1 that we agree with the position that these 2 other buildings come into play. 3 MS. MUNN: This is Wanda. I question that 4 though the statement was on page 6, that there 5 was clear evidence that you had not adequately 6 explored the potential and I -- I questioned 7 what the clear evidence was because if we had 8 discussed any such evidence prior to this I 9 wasn't aware of it. 10 DR. NETON: And literally with the hundreds and 11 hundreds of pages we've gone through there is 12 not one shred of evidence to indicate that 13 thorium processing occurred in those other 14 buildings prior to 1957 --15 DR. MAKHIJANI: Well --16 **DR. NETON:** -- '58 so --17 DR. MAKHIJANI: Well, if I might just respond I 18 -- as I said the -- we -- we've given the 19 citations I think for those buildings clearly, 20 you know. The people who worked there have 21 evidence and -- and their evidence should be 22 taken into account. But I think we've cited 23 the reports. Not, as I said, not all of the 24 reports give dates that are clear. But the

Chem-Risk report was very clear. And frankly I

was very surprised. But -- But I -- I haven't read the whole Chem-Risk report but I do -- we did think that it should be evaluated since -since there were other reports as well that mentioned other buildings. It's not -- It's not really clear to us from reviewing this other than the Chem-Risk thing that -- that there were other buildings but when it is in an official report that was prepared as a result of access to all classified information and production and there was a commission I believe, was it by the Centers for Disease Control? I -- I don't remember now. that -- I -- I don't believe that that -- that should be dismissed as -- as -- as flimsy evidence or not --

DR. NETON: I'm not saying flimsy, Arjun, but you could interpret this paragraph several different ways. I mean they began thorium processing and fabrication but now there were fairly pilot operations going on. I don't think what we're citing here is inconsistent with the language in this report. We had -- we take no exception to the fact that thorium was being handled and moved about and -- and

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operated on in those periods but the -- what they say in the last sentence of this paragraph that you cited is that the majority of the thorium production scale operations. And we're saying that production scale operations did not begin until the end of 1950s. But it's not inconsistent with that. And we have cited the RCO report; it was called Atypical Radionuclide Assessment of the Y-12 National Security Complex that references the Wilcox report as well as the Hap West report, that both confirm that the -- that the production scale operations occurred in the end of the 1950s. And that's very consistent with seeing the ramp-up of the fecal sampling program, the ramp-up of the 90,000 hair samples that were taken starting in those years and everything else that we've looked at. I don't know that this is an issue that -- that we can agree with.

MR. GRIFFON: Arjun or Jim, does -- does the Chem-Risk documents cite any source documents?

DR. NETON: It makes no reference at all.

DR. MAKHIJANI: Well, I -- I, you know, I had
very little time and I kind of parceled out to

1 the various pieces. Unfortunately Kathy is not 2 on the call. I -- I did collect -- I -- I 3 researched parts of this myself but parts of 4 this part I did not so I have not actually read 5 the Chem-Risk report. And I, you know I trust Jim that there's no reference there but -- so I 6 7 -- I don't know where to go with this. I mean 8 obviously we had to cite -- we -- we were asked 9 to review the report and so we cited the 10 evidence that was available to us. You know, 11 there's -- I don't believe that we should take 12 a stand on any particular (inaudible) despite contrary information but this is the 13 14 information that was available and I thought --15 I was a little surprised as I said to see them 16 compare this to operations comparable to 17 uranium which -- which would indicate significant operations. 18 19 MR. GRIFFON: Jim -- Jim, can I ask this to --20 to try to resolve this? You mentioned several 21 documents that you had. Are some of those or 22 all of those on the O-drive or --23 DR. NETON: I believe they are. Someone at 24 ORAU can help me with this. 25 MR. GRIFFON: I mean maybe if -- it doesn't

1 have to be done on this call but --2 DR. NETON: Sure. 3 MR. GRIFFON: -- but maybe you can provide a 4 list of documents that --5 DR. NETON: We can certainly provide the source documents on the O-drive --6 7 MR. GRIFFON: Yeah. 8 DR. NETON: -- that we related that are 9 referenced in our internal dosimetry TBD. 10 quess that's where I take a little bit of 11 exception where, you know, the -- the report 12 cites ample evidence that we haven't clearly identified it but it doesn't cite the evidence 13 14 that we cited. And -- And so, you know, they 15 found one exception to -- to the rule which is 16 unreferenced so --17 DR. MAKHIJANI: But we -- Jim, we did not 18 disagree with your finding. What, you know --19 there was no need to -- there was no need to --20 to re-cite your references. And one -- one of 21 your references was not yet available to us 22 that was cited in the evaluation report. But 23 we didn't -- we didn't have -- we did look at 24 the references that you cited that were

available to us and had no disagreement with --

1 with what you said as regards to thorium processing and all that. We were just 2 3 supplementing what we found about buildings 4 that you hadn't cited. 5 DR. NETON: Right, Arjun. But the TBD which 6 you did review cites that we believe it started 7 in the early '60s and those references are 8 listed there as well, and they were not 9 reviewed at all. 10 MR. GRIFFON: Okay. I think that -- that, you 11 know, I mean maybe a follow-up we can make sure 12 that -- that either in the TBD or the -- or, 13 you know, if there's others that -- that those 14 reference are just maybe told SC&A and the 15 Board, you know, the work group what those are, 16 where they are and, you know, you might 17 consider that in this, you know. Again I think John, you're presenting -- and Arjun, you're 18 19 presenting this as a draft --20 DR. MAKHIJANI: Yeah. 21 MR. GRIFFON: -- final draft report so --DR. MAKHIJANI: Yeah, well, you know, it was --22 23 MR. GRIFFON: There's other stuff that you 24 should consider in -- in assessing this issue. 25 I think you should, you know --

DR. NETON: Right.

type of discussion.

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DR. MAURO: In fact, this goes a little bit toward -- this is John Mauro -- my prefacatory (ph) remarks in terms of capturing the bigger picture in terms of for example, disclosing the -- the arguments for when major thorium activities may have taken place. However, there is also perhaps some other information such as the Chem-Risk report which would seem to indicate that perhaps some important thorium -- in other words, try to tell the story in a way that is more inclusive as opposed to your zeroing in on those particular delta pieces of information that we've uncovered that probably need to be run to ground. So I think the report, our report, would benefit from that

MS. MUNN: This is Wanda. Again, I zeroed in on the specific language in the second paragraph on page 6 that says there is clear evidence that NIOSH has not adequately explored the potentials of thorium work. And what I'm hearing from NIOSH is that they have explored that quite extensively. So the -- the language, the way in which this question is

1 presented, raised an issue in my mind. 2 DR. MAURO: Wanda, fair enough. I hear you. 3 MR. GRIFFON: Yeah. Yeah, I think we have to, 4 you know, yeah. We -- We should look at all 5 the references that they -- that they cited or -- and if there's additional ones that are not 6 7 cited in the TBD or otherwise I think, you 8 know, that does shed light on this. I think 9 you should --10 DR. MAKHIJANI: We -- We did cite here that 11 the TBD says that processing with thorium began 12 in the '60s. I mean we -- we will go back, you 13 know, at the working group's direction, of 14 course, yeah, and -- and review the other 15 references. 16 DR. NETON: I might also add though that we did 17 reference the Chem-Risk report in the site 18 profile and clearly a weight of the evidence in 19 our mind did not include the early '50s based 20 on an evaluation of the data we had at hand. 21 MR. GRIFFON: All right. 22 DR. NETON: Yeah, Chem-Risk --23 MR. GRIFFON: And I -- Jim, I agree with your 24 point that -- that you could interpret that one 25 paragraph, that last line especially, as a

1 little bit, you know --2 DR. NETON: Right. 3 MR. GRIFFON: There's a little, you know, you 4 can interpret it either way, I suppose, you 5 know. But with your other evidence you're 6 saying, you know, you certainly don't think 7 it's inconsistent with what you found in all 8 those other documents so I think --9 DR. NETON: Right. 10 MR. GRIFFON: I think we need, you know, SC&A, 11 we need to look at those other source documents 12 and weigh the prepon-- you know, weigh the preponderance of the evidence I guess. 13 14 DR. NETON: Okay. 15 MR. GRIFFON: Can I ask one thing, Jim? 16 I know Mel Chew talked about having all this 17 sort of receipt data or ledgers or whatever 18 that showed amounts of thorium coming in, 19 amounts of all those radionuclides. 20 probably just gross receipts to the site, 21 right? It didn't talk at all about 22 distribution to any buildings or -- is that 23 true? 24 DR. NETON: Mel's on the line.

MR. CHEW: Yeah, Mike (sic). I -- I'm glad

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you brought that up because I was going to also show that's another pieces of evidence here. When we go back to the classified ledgers which are still classified it does bring in the receipts of the -- of the thorium that came into Y-12 by year and by period. Now, if you really dive down into the individual receipts there, and we didn't have -- we didn't go there exactly at the time, it also shows that in -for instance that if they move it to another materials accountability area and that certainly could be by building. And I don't want to quote that to be -- be exact. know, we could trace for instance, you know, ten kilograms or five kilograms went to this particular building, for the R&D work which makes sense. But I only took the larger number that came in for that period just to show the quantity, total quantity that was at Y-12 available here. But I said -- I want to again add to it that there is certainly evidence by many of the reports that Jim has been talking about where the processing of -- major processing for the campaign of thorium did occur. Now, thereby, I will also agree there

1 was thorium there. Remember they used some 2 thorium for the co-precipitation for the 3 Cyclotron. That -- That was there. And there 4 And there certainly was evidence that there 5 was small quantities of thorium that was used 6 for the R&D development of the processes, you know, in -- in -- in anticipation of the major 7 8 program. We saw, you know, an air sample that 9 was cited in the health physics reports that 10 talk about that particular building. And then 11 also the -- the slow ramp-up as the R&D 12 activity occur. But I would like to say that 13 in looking at item number two, those particular 14 buildings that were cited in the last sentence, 15 those activities really started even -- even 16 past the 1979 in the FCC period but in the late 17 1959 into rough 1960s and --18 MR. GRIFFON: Are you talking about --19 MR. CHEW: -- that was documented. 20 MR. GRIFFON: Are you talking about Alpha 5 and 21 Beta 4. Are those -- 9201-5 and -- and 9204-4? 22 MR. CHEW: Yes. All the ones that are listed. 23 MR. GRIFFON: Okay. 24 MR. CHEW: And then -- And we can mention them

for evidence because when the -- when the

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1 campaign really started then thousands -- many 2 air samples have showed up and you can just 3 show up -- you can actually go to show where 4 the air sampling started because that's where 5 the operation started, and those air samples 6 are by building. 7 MR. GRIFFON: Yeah. I think I probably know 8 the answer to this, Mel, but I'm going to ask 9 anyway. How difficult would it be to walk the 10 thorium data back, the ledger data back to the 11 buildings? 12 MR. CHEW: It would probably mean that we have 13 14 MR. GRIFFON: A time-consuming effort? 15 MR. CHEW: Yeah, I mean it would be going -- go 16 back to Y-12 and go back into, pull the ledgers 17 which we know are there and then try to reconstruct in how we would contract -- you 18 19 know, these are -- at that time they kept the information in -- in the ledgers, you know, 20 21 according to like numbers or something like 22 that. You would have to find the corresponding 23 -- what MBA it is. I think it could be done 24 but I think it would be time-consuming. 25 MR. GRIFFON: Okay. I assumed that.

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DR. MAURO: This is John Mauro. You're going to see a little later on one of the other items, one of our observations is that it doesn't take very much thorium airborne to contribute significantly to bone dose for example, or other organs so -- and this -- it may be related. It sounds to me that there is a continuum of operations going from I guess perhaps R&D to production where thorium is being handled in various buildings. And it sounds like we could run down, through what Mel just described, that process in terms of quantities delivered to various buildings. Now, confounding this problem is the matter that it doesn't take very much thorium airborne to be an important contributor to the dose as compared to uranium. As a result we've got ourselves what we envision as a bit of a That is, even if it's a relatively dilemma. small quantity that might have been handled, it doesn't take very much to be important. MR. CHEW: Right, John. I'd like to have a collegial discussion with you. I saw your report on the -- about the contribution attempts of one percent there, of doubling the

1 bone dose here. 2 DR. MAURO: Yeah. 3 MR. CHEW: You know, you -- you -- you clearly 4 mentioned that it was done by radioactivity and 5 I agree with that. DR. MAURO: 6 Yeah. 7 MR. CHEW: But you need to look at it from a 8 math standpoint here, okay? 9 DR. MAURO: Okay. 10 MR. CHEW: In other words, you tell me how much 11 you go back and recalculate if I had a gram of 12 uranium dust in the air how much more thorium I 13 would have to take to -- to add to that 14 contribution from a --15 DR. MAURO: You're absolutely right. 16 MR. CHEW: -- from a math standpoint. 17 DR. MAURO: And that might be the answer. 18 MR. CHEW: Yes, exactly right. Yes. If that's 19 I think it's misleading to say -- I shouldn't say that, John. Sorry. Don't take 20 21 offense at that. 22 DR. MAURO: Are you saying I'm misleading? 23 MR. CHEW: No, no. Don't take offense at that, 24 John. 25 DR. NETON: I'd like to -- I'd like to chime

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in on this if I may. Mel is absolutely right that, you know, it takes much more mass of thorium than uranium to -- to get the equivalent amount of intake. But that issue notwithstanding I think, you know, in reading SC&A's write-up on this issue, I think that they might have missed the -- the concept here in the sense that we didn't say thorium workers are covered. We said workers who were monitored or should have been monitored for thorium, that is by today's standards. we're not -- we're not -- the SEC class is not people who physically worked with thorium material. It's people who may have been in buildings that were nearby thorium and because of exactly the reason SC&A cited there could have been bleed-over of thorium into their adjacent work areas and then they would be covered as part of the class. There's a little bit of a difference there I think if you look at it from that perspective.

DR. MAKHIJANI: Yeah. This is -- This is
Arjun, and -- and, you know, this -- this -- I
-- I wrote that section so let me take
responsibility for that one at least.

DR. NETON: Okay.

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DR. MAKHIJANI: I couldn't exactly explain the other, all the details of the other one. The -I -- I did look at the work in the evaluation report and I did think of the possibility that -- that even trace amounts of thorium exposure might be covered. looked at the description of the air sampling that involved thorium in the one building I think I cited it. Only one of the 13 air samplers was described as potentially an air sampler for thorium where uranium and thorium would be mixed and so it did raise a question in my mind what -- as to what might happen to uranium workers who were breathing trace amounts of thorium and whose doses you might think that you can calculate because you had air monitoring data for alpha and uranium bioassay data in the same way that say you were -- you were trying to handle the Mallinckrodt information. And actually I didn't conclude that you could or couldn't do it. It was, in the case of uranium workers who -- whom you have bioassay and some air concentration data, I'm not clear as to whether you can or can't -

- can't calculate their doses. And the point of -- of raising that question was exactly that. Is it -- Is it -- Are you including the trace exposures in the uranium class -- in the uranium class or in the should have been potentially monitored class?

DR. NETON: I think we're getting into an issue that the Department of Labor is going to address for us at the Board meeting, which is how do they determine or define who is a member of the proposed class, in particular in light of the fact that the definition says was monitored or should have been monitored. That -- That's not under our purview. You know, we define the class as, you know, what we can and they -- they make the determination. And whether or not they take in, you know, account for trace potentials or not I think we need to hear -- hear them out.

MR. GRIFFON: Yeah. I -- I -- I think you're

-- I think we do need to hear them out, Jim.

I think you're right. I -- I mean I -- I've

been wondering about this issue myself that,

you know, my understanding was that it's up to

the Department of Labor to identi-- you know,

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you define the class and then the Department of Labor identifies claimants who meet the definition of the class. And now, you know, to me this -- the only concern I have is that does the Department of Labor have enough information to actually -- to understand the definition of the class and how the claimants fit into that class, you know, to -- I guess it's a different scenario. You know, Larry, in the last call, brought up the idea of -- of Paducah but really it's -- it was, you know, monitored or should have been monitored for the whole plant site and they might exclude like administrative assistants or something like that and send them for dose reconstruction but I think it's a little -- little harder for the Department of Labor to discern who, within these large, you know, production buildings might have been near or nearby a thorium process when they don't even know where these things took place.

MR. ELLIOTT: Well, let's just --

MR. GRIFFON: I don't know if we're giving them enough -- enough information to do the job.

And then how do they deal with it, you know.

MR. ELLIOTT: Yeah, this is Larry Elliott. Let

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me again make a comment here. We -- We, as part of the process in developing the evaluation report, when we arrive at a recommended definition for the class we vet that with DOL and there's a discussion about does it -- is it suitable and does it give them all that they need and do they have all -- all that they need to determine eligibility of the claim for inclusion in that class. certainly had done this on Y-12 in this particular case. Also, I would remark again that this is not new to the Department of They are -- Pete Turcic will be at the Labor. Board meeting next week to provide you with a presentation and examples on how they go about doing this. It's not only just for -- they don't determine just eligibility for a given class but they determine eligibility of a In fact, if you look at like Chapman Valve and Building 55, if you look at the Iowa Army Ammunition Plant and line one, when you get into those kinds of covered facility designations, those have to be clearly and carefully handled, and DOL has developed their experience in that regard.

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MR. GRIFFON: You're right, Larry. We need to hear their presentation, so you're right.

DR. MAURO: But I think we're in a very interesting grey area that in defining the class effectively what we're saying is while the class of thorium workers, and identifying the buildings, but the implication that the other buildings are, you know, limited to uranium workers and therefore, we can do the dose reconstruction. I think the key to parsing the two and -- and bringing this issue to ground goes to what Mel has just described. I think -- I'm thinking about, you know, how do you -- how do you get to grips with making sure that the -- that the buildings we say we can do the dose reconstructions for are in fact buildings we can do the dose reconstructions for. We need to go to somehow getting a handle on, as Mel mentioned, how much material en masse may have been transported to those buildings at a given point in time. And --And this becomes very much a technical health physics kind of question. Is that enough material to create -- in terms of mass now, to create a situation where you could have

1 picocuries per cubic meter, that could 2 contribute significantly to the inhalation 3 dose. I mean this becomes -- I'm trying to 4 find a way to make sure that the boundary can 5 be found. And I think the -- the key to that 6 boundary lies with the information that Mel 7 just described. 8 MS. MUNN: There's also the question of what 9 form the thorium was in at the time. Later in 10 SC&A's recent report here there's a long list 11 of precisely what activities and therefore we -12 - we know what form thorium was in in the '60s. 13 But in these early days when I believe I heard 14 expert comment from individuals who knew the 15 site well that all thorium use in these early 16 years that we're looking at for the SEC 17 petition revolved around its use as 18 precipitation in the Calutrons. Was that not 19 correct? 20 MR. CHEW: No. 21 MS. MUNN: Okay. 22 MR. GRIFFON: Not all -- Not all of it. 23 DR. NETON: In the very early years --24 MR. GRIFFON: Oh, very early years. Okay. 25 MS MUNN: Right. Right. And -- And that's

1 what we're looking at here. 2 MR. GRIFFON: But not all during the SEC 3 period. 4 No. In the later years, in the '56 DR. NETON: 5 time frame in particular there is evidence of people working with thorium. 6 7 MS. MUNN: It was starting to ramp up. 8 DR. NETON: In the research building, right. 9 MS. MUNN: Right. But -- But early on we, 10 perception and perhaps it's my lack of 11 understanding of the Calutron process but my 12 perception was that that would have been a wet 13 process? Yes? No? 14 DR. NETON: It was a co-precipitation process; 15 that's correct. 16 MS. MUNN: All right. So -- So extreme 17 concern over airborne would seem to be 18 questionable. 19 MR. GRIFFON: But see, and I don't -- I don't 20 necessarily disagree with you, Wanda, here. 21 The question I have more is could -- defining 22 that potential, you know. It seems to me that 23 -- that, you know, exposed or could have been 24 exposed; well, now it's in DOL's court and they 25 have to determine, you know, geez, what kind of

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processes were in these buildings, what kind of -- who is making that determination as to a -- a real, significant potential for exposure.

DR. NETON: I really think, though, we need to hear the Department of Labor out.

MR. GRIFFON: Yeah, I agree, Jim.

DR. NETON: Especially in all the areas of how
they --

MR. GRIFFON: No, no, no. I agree and Larry -- Larry's right on that point so --

DR. MAKHIJANI: Yeah. This -- Okay, this is Arjun, just to say why I wrote that part is the -- the evaluation report distinguishes between uranium workers or those who were exposed to uranium and those who should be monitored for thorium. And the point I was raising is the dose reconstructibility for those who worked with uranium and may unknowingly to them or to the people who were involved at that time in monitoring. In that building where they had 13 monitors they only defined one as a thoriumuranium mixed area. So unknown to them -- so these workers -- there's a group of workers that would be defined as uranium workers which would fall within the purview of NIOSH's

1 assertion that you can calculate dose. 2 course agreed there's quite a lot of uranium 3 bioassay data. And that's the group of workers 4 that I raised the question about and -- and it 5 may be possible or not possible to calculate their doses. I -- I don't have a judgment 6 7 about that. 8 DR. NETON: Arjun, again the definition is not 9 uranium or thorium worker. 10 DR. MAKHIJANI: I agree. 11 DR. NETON: I mean, so, you know, you can't 12 presume what we're going to do here. 13 DR. MAKHIJANI: Okay. 14 MR. GRIFFON: We have to wait on this. Yeah, 15 we -- I think, I mean we're discussing one and 16 two, right? We sort of went on to seven a 17 little bit I think but -- or not seven but section seven. 18 19 DR. MAKHIJANI: Section seven. 20 MR. GRIFFON: Yeah. But anyway, is there 21 anything else on one and two that we can 22 resolve now? I mean I think one thing as a 23 follow-up, Jim, it would be good to make sure we have all the references if -- and you can 24

just say if they're as -- as cited in the TBD

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1 and -- and maybe just to expedite things if you 2 can kind of point us in the right direction 3 where they are in the O-drive that would be, 4 you know, helpful. And then SC&A should 5 consider them in the final draft of this section on the -- the other buildings, the ones 6 7 particularly cited in Chem-Risk doc. 8 Will do. DR. MAKHIJANI: 9 MR. GRIFFON: And then is there anything else 10 on one and two? I'm looking at the time, too, 11 at 12:00 o'clock here. I'd like to get through 12 most of this before lunch, take -- taking lunch 13 at 1:00 again I think. Is there any more on 14 that -- those two sections or any --15 DR. NETON: Not from our end, no. 16 MR. GRIFFON: Okay. 17 DR. NETON: Okay. MR. GRIFFON: 18 And the big thing I think we're 19 going to have to wait for is DOL's, you know --20 we need to hear what DOL has to say on that so 21 okay. 22 DR. NETON: I guess -- I guess I do have one 23 more thing just -- just for completeness is 24 there was an issue raised about the ponds and 25 the exposure out there and we have to track

this down but I -- I've got to believe that I haven't been able to definitively define this this morning but those ponds were -- were being dredged after the SEC period. It makes no sense that they would be dredging ponds for thorium when they had such limited use and there was huge concentrations of thorium that they were finding in the bottoms. You know, while the material was being discharged in the pond we don't feel there's any credible exposure scenario to the workers.

MR. GRIFFON: Okay.

MR. CHEW: Jim and Mark. This is Mel. I'd like to just make one more comment to John Mauro. John?

DR. MAURO: Yeah.

MR. CHEW: I think -- and I appreciate -- I appreciate your expertise and I did a backup (unintelligible) calculation here. It would take about a hundred grams of thorium to -- in addition to one gram of uranium to equal the amount of radioactivity that would be present and so -- so please look at it from a math standpoint to make -- to come to your conclusion, okay?

1	DR. MAURO: Yeah.
2	DR. MAKHIJANI: Did you use enriched uranium or
3	natural uranium or DU?
4	MR. CHEW: I think I used nata probably just
5	the what the concentration in natural
6	uranium at that particular time. And this is
7	just a rough calculation here.
8	DR. MAKHIJANI: It would be about a factor of
9	six or seven if you take the half-lives. When
10	you throw in thorium 228 it's about a factor of
11	five, not a factor of a hundred.
12	MR. GRIFFON: Anyway, you can you can
13	consider that in your final draft, right?
14	DR. MAKHIJANI: Sure.
15	MR. GRIFFON: On a math basis, yeah.
16	DR. MAKHIJANI: Sure.
17	MR. GRIFFON: All right. Thanks, Mel. Go
18	ahead, Arjun. You're going to go on to number
19	three?
20	DR. MAKHIJANI: Number three.
21	MR. GRIFFON: Yeah.
22	DR. MAKHIJANI: We thought NIOSH had done a lot
23	of work on the internal dose verification of
24	the CER, of the verification validation of
25	the CER database on the internal dose point of

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view from 1952 onwards. There had been a suggestion in the working group meetings that there were some raw data from the SEC period to which it could be compared and I don't know what happened, what was the status of that. There was some raw data comparison from -- from the 1970s. I'm just looking at my summary if I remember correctly, and I think that there is a lot more confidence in -- in the -- in the database from 1952 onward but we thought there were still some gaps. 1950 and '51 served different issues in the sense that there's -there's not been an effort that we saw for validation in those two years and we had a concern about those two years particularly because in the external database there were a lot of problems. Didn't find a parallel problem of zeros for the record in -- in the -in the internal dose database but did think that specific -- specific verification of -- of those two years to some extent or some -- some part, some piece of -- modest piece of that should -- should be done.

DR. NETON: This is Jim. I'm a little confused
because -- not confused -- What SC&A is now

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asking for us to validate '50 and '51 when in fact we have not been able to find any raw data to my recollection in the -- in the SEC period. You know, we had to rely on secondary, you know, analyses of -- of looking at -- at data outside the period. I think we need to keep in mind a couple things here. One is that at the outset we determined that the CER database or we -- it was our belief and we were provided some at least secondary evidence to the fact that the CER database was accepted by the Department of Energy as being the data of record for exposures of workers. And in that sample a lot of work went into making sure the data accurately represented what, you know, what the samples, you know, measured. that sense, you know, we believe that we've got -- we're a little bit above the bar here because it has been validated to a certain extent. But at least I feel we were not able to establish, you know, show the pure documentation but at some point one needs to -to accept it as it is for these dose reconstructions. We tried to validate it against various pieces of information, the

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delta view data, punch cards and that sort of thing. And in fact in some cases as SC&A points out we were successful in demonstrating that the data are reasonable. However, there are discrepancies. I would point out that the discrepancies that we've observed both in the internal and the external areas have consistently provided data that would -- that would bias the results low, in my opinion anyway, especially if you're -- if you're using them for developing co-worker data. In other words, the data in the '51 time period for external with a significant portion of zero results, you know, that sort of thing. delta view database that had uranium had lower results than what the averages that were for the CER database. So given that, we believe the data that are -- are present in the -- in the CER database are reasonable to use for dose reconstructions and reasonable to use for coworker development. We see no reason, and SC&A asserts, that the data in '50 and '51 are invalid in the CER database. I don't think anyone has come to that conclusion.

DR. MAKHIJANI: I don't believe we said that

1 about the internal dose. We did say that about 2 the external dose and I -- and I thought that 3 you agreed with us that there was some kind of 4 problem that you couldn't identify. But that -5 - that's a separate -- the term invalid was not applied I believe either in the fine print or 6 7 in the summary in regard to the internal dose. 8 DR. NETON: Well, I think there are statements 9 made though, Arjun, that says that we could not 10 use them for dose reconstructions for --11 DR. MAKHIJANI: Well --12 DR. NETON: -- or by inference because of 13 issues with the external you -- you have 14 equated that to issues with the internal. 15 DR. MAKHIJANI: Well --16 DR. NETON: That's what it says. 17 DR. MAKHIJANI: Well, yeah, we did feel that the 1950 and '51 -- I mean if you take -- if 18 19 you take the statement that the DOE 20 certification of this as the dose -- as the 21 database of record at face value, then you have 22 to take that statement in its entirety both for 23 internal and external and it is very clear that 24 for 1950 and 1951 the -- the CER database is 25 wrong because it contains all zeros contrary to

1 the information in the raw data --2 DR. NETON: Well you have --3 DR. MAKHIJANI: -- for external dose. Please. The -- The -- It also contains 4 5 information that at least to us felt that when shallow and penetrating dose did not seem to 6 7 make scientific sense in that neutron seemed to 8 be included in shallow dose but not in 9 penetrating dose. So because you're trusting 10 the DOE statement in regard to the whole 11 database, not for internal or external, I -- I 12 -- I think that some verification for -- for 13 the years 1950 and '51 is needed, especially 14 because as discussed in another section, the 15 types of work done in three buildings in those 16 years were different and were terminated in 17 1951. So you need the data from those years to 18 reconstruct for dose -- for those workers. 19 DR. NETON: Are you talking about the internal 20 exposures? 21 DR. MAKHIJANI: Internal and -- and external. 22 DR. NETON: Well, let's -- let's --23 DR. MAKHIJANI: Unless --24 DR. NETON: I think George wanted to say 25 something.

1 MR. KERR: Yeah, I -- I want to say something 2 because there's a misstatement up here in the 3 front as well as back on page 11. And the fact 4 is that in the early years the beta doses were 5 more concern than the gamma doses. 6 DR. MAKHIJANI: Okay. 7 MR. KERR: And if you look back at '50 and '51 8 there are beta dose data that are not zeros. 9 There are significant beta dose exposures in 10 '50 and '51 among employees. In '50 there is 11 one gamma dose in -- or '50 there's one person 12 that has a recorded gamma dose that's not zero. 13 In '51 there are -- there are no recorded. But 14 keep in mind there is beta dose data in the CER 15 database. 16 DR. MAKHIJANI: I don't believe -- I believe 17 that gamma and beta in the CER database are all zeroing. 18 19 MR. KERR: No --20 DR. MAKHIJANI: Maybe I'm --21 MR. KERR: -- no, no. That's wrong. That's 22 wrong on page 11. 23 DR. MAKHIJANI: Well --24 MR. GRIFFON: Okay. Can I -- Can I ask one 25 thing? Can we go back to number three and --

1 and focus on the internal just for one second 2 and then we'll do more on -- we'll come back to 3 the external. 4 MR. KERR: Okay. 5 MR. GRIFFON: I'm sorry. I just -- Jim, can you tell me just -- just as a summary specific 6 7 items that you did? I mean I'm trying to think 8 of -- of the various items that you did to 9 check the reliability. We've got the letter, 10 of course, that's your -- that's your 11 overriding thing here. But then you have the 12 HP reports percentile data mainly. 13 DR. NETON: Right. 14 MR. GRIFFON: And then you have if I'm not mistaken 8 -- 8 or so or 8 or 20 -- I don't 15 16 know if --17 DR. NETON: There were 20 -- I think there were 18 20 workers who we found that had reference to 19 bioassay results in the health physics report 20 and they were cross-walked to the database in -21 22 MR. GRIFFON: Twenty individuals. 23 DR. NETON: -- virtual 100 percent agreement 24 with the exception of one bioassay. 25 MR. GRIFFON: Right. Twenty individuals so

1 from the HP report again. 2 DR. NETON: Correct. 3 MR. GRIFFON: And then you have the -- the --4 the --5 The punch cards. DR. NETON: 6 MR. GRIFFON: -- so urine cards, right? 7 DR. NETON: Right, the punch cards which were 8 in a later time period where the samples 9 matched up. We weren't able to reconstruct the 10 bioassay results very well because we didn't 11 have all the background. 12 MR. GRIFFON: Right. 13 DR. NETON: Now --MR. GRIFFON: Now, can -- can you tell me 14 15 'cause I -- I remember bringing up this 16 question and I -- I don't think it was a 17 follow-up action but you were going to -- or --18 or there was a question as to whether you had -19 - no, you didn't have punch cards from the --20 from the time period in question, right? 21 DR. NETON: That's correct. 22 MR. GRIFFON: Okay. So that was --23 **DR. NETON:** So we -- we really were not able to 24 establish any -- any direct validation or 25 reliability check of -- of the data in the SEC

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period. But -- But getting back to the 1950 and '51 era, you know, I think there's a misunderstanding -- we'll get into this later in one of the questions -- about how NIOSH has modeled the internal exposures in 1949 and '50. We have no bioassay data in that period. what we did is we didn't assume that the bioassay would have been excreted to the same level as 1951 and '52. We took the excretion in 1952 and said, what could these workers have possibly inhaled in '49, '50 and early '51 and still be excreting what they are today in 1952. That's a very different analysis. In other words, we used the workers as long term integrators of their exposure in the earlier years. And we believe that sufficiently brackets the exposures in those areas and actually does a fairly nice job at it. So we did not assume that they were excreting the same amount in their urine. We used them as actual predictors to back calculate what the maximal exposures could have been from a chronic exposure scenario.

DR. MAURO: Jim, that's -- this is John.
That's very helpful.

1 MR. GRIFFON: That's a good clarification, yes. 2 DR. MAURO: (Unintelligible) strategy. 3 DR. NETON: Yeah, I felt --4 DR. MAURO: Perhaps I should have known that 5 but I didn't. DR. NETON: This will answer a couple questions 6 7 I think where SC&A was -- was -- had some 8 serious issues with those time periods. 9 DR. MAURO: So in effect what you're saying --10 what you're effectively saying is what you're 11 seeing in the urine of workers when you do have 12 the bioassay data -- I'm looking at your table 3 now, for example. In table 3 you have --13 14 well, I'm looking at table 3 in our report on 15 page 15. What I'm hearing you saying is for 16 urinalysis we have 166 employees measured and 17 you're seeing certain concentrations. 18 assumption is being made that what you're 19 observing there in those workers is the result of chronic intake, as an integrated intake that 20 21 the workers experienced prior to that date. 22 MS. MUNN: Is it my phone or is John fading 23 away? 24 MR. GRIFFON: Prior to that date maybe all the 25 way back to 1950 is what you're saying, right,

1	Jim? Depending on the workers' circumstance I
2	guess. Hello?
3	DR. NETON: Prior to that date and all the way
4	back to 1948.
5	MR. GRIFFON: Oh, '48. Yeah, yeah.
6	DR. NETON: Yeah, we're saying
7	MR. GRIFFON: Right.
8	DR. NETON: We're saying
9	MR. GRIFFON: Right.
10	DR. NETON: what could these workers have
11	inhaled on a chronic basis and be excreting
12	what we're measuring in that time frame in the
13	early '50s.
14	MR. GRIFFON: Okay.
15	DR. NETON: And so that that we believe
16	MR. GRIFFON: That
17	DR. NETON: provides a bounding analysis of
18	what the exposures were in those years.
19	MR. GRIFFON: That wasn't clear to me so that's
20	helpful, yeah.
21	DR. MAURO: Excuse me.
22	MR. GRIFFON: It should have been but it
23	wasn't.
24	DR. MAURO: Yes, that's that's very helpful.
25	DR. NETON: I have to admit that the TIB I

1 think it's in there but, you know, it's those 2 dosimeters sometimes use shortcut language and 3 it's not obvious I don't think. 4 MR. GRIFFON: Okay. Well, that's helpful. And 5 Jim, can you tell me one other clarifying point here? 6 7 DR. NETON: Sure. 8 MR. GRIFFON: And without having to look it up? 9 In your evaluation report the HP reports that 10 you looked at the percentiles for, was it --11 was it multiple years? Was it one year? 12 13 DR. NETON: I -- I think it was only for one 14 year. Bill Tankersley did that analysis. 15 Bill, could you --16 MR. GRIFFON: It was like '53, wasn't it? 17 MR. TANKERSLEY: Yes, it was for one year, and Mark, it was for 1952 for all 26 weeks I think, 18 19 the latter part of '52. 20 DR. NETON: So if it was only one year I mean I 21 -- I fully admit that we've had limited success in -- in demonstrating the reliability of the 22 23 data, you know, particularly in the SEC period. 24 But again I went back and looked at our -- our 25 discussion, Mark, that we had back in November

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of last year about this exact issue and in the -- in re-reading the transcripts of that meeting it was clear to me that we were concerned more with -- with reliability when there were issues raised particularly by petitioners about, you know, certain activities that may have occurred. And secondly, if these were secondary databases such as CEDR data which were -- were summary data obtained from epidemiologic studies. And so here we have what we think is about as close as we're going to get to a -- a -- a very good quality database. And the fact is, and I've raised this issue back in November, that for 50 years later it's very difficult for us to obtain raw data to validate all these individual points. And the working group and the Board are going to have to decide what level of -- of proof they're -- they're comfortable with.

MR. GRIFFON: Well, I -- I also think, and I'll -- I'll offer this up as -- as a -- maybe a bit more to support the reliability case, that there's other HP reports that have the same percentile data and I think I've done back -- and I admit back of the envelope sort of

1 calculations on -- on those other periods and I 2 think they would bolster your argument so --3 But I -- But I think just to present one in 4 the evaluation, you know, at least -- at least 5 you might have that in your -- in your hip pocket to -- to better defend. And it would 6 7 also, you know, say that because we're, you 8 know -- I think that is probably one of the 9 most powerful arguments because that's --10 that's the summary data for that whole half a 11 year. I think it's about half a year on most 12 of the reports. 13 DR. NETON: Right. 14 MR. GRIFFON: And it -- It virtually agrees, 15 you know, pretty dead on with the numbers in 16 the database. 17 MR. TANKERSLEY: Excuse me. 18 MR. GRIFFON: But just to present one half year 19 of it, I think, you know, makes a less powerful 20 argument. 21 MR. TANKERSLEY: This is Bill Tankersley. 22 Mark, I was just about to add, and I appreciate 23 your comment there. It sounded like an 24 inference a moment ago was that this was the

only analysis that -- that we found to match.

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That -- That's not the case. It's the only one we tried, and the reason why is because it takes quite a bit of work to extract the percentiles from their graphs and then to calculate the percentiles, you know, by week for these things among all of the other things that, you know, the team is doing.

MR. GRIFFON: Yeah. No, I see --

MR. TANKERSLEY: All other -- Not in every one of the reports, but there are other of those graphs that could be done. I'm not in a position to say what the match would be. It sounds like you've done the matching.

MR. GRIFFON: Well, and again, I -- I did a quick and dirty but I didn't have to put it in the report either so -- so I understand you'd have to be a little more precise and it takes a little more time, yeah.

DR. NETON: Yeah, I hear what --

MR. GRIFFON: But I think it would bolster your argument and that's the reason I bring it up is that what's before the Board is an evaluation report with one, you know, where that was done through one half a year. And it suggests to, you know, all my colleagues on the Board and

1 the public that, you know, that's the piece of 2 evidence you had so I don't know. I think that 3 might be worth pursuing if it wasn't going to 4 be a tremendous amount of person hours, you 5 know. DR. NETON: Appreciate that, Mark, and we'll --6 7 we'll take that to heart and do the best we can 8 prior to the Board meeting. 9 DR. MAKHIJANI: Mark, this is Arjun. 10 MR. GRIFFON: Yeah. 11 DR. MAKHIJANI: Guide -- Guide me here a 12 little bit. And guide the SC&A team. We took our cue from the Board's decision on criteria 13 14 for -- for approaching SEC evaluations in 15 preparing our review. But that's the one --16 that's the one Board approved document that we 17 have. We don't have approved procedures but we 18 do have that. 19 MR. GRIFFON: Yeah, I think that's appropriate. 20 I think we agreed to that. DR. MAKHIJANI: And -- And data validation so 21 22 it's -- so data validation and -- and 23 representativeness -- those are separate issues 24 -- are very prominent and central in that 25 document and -- and are kind of limited to what

you can show. And I think -- And I think I -- I don't disagree with Jim in that a lot of effort has been made and I think of -- to the -- to the extent that the validation has been done from '52 onward there appear to be matches and so on. But we did, if you take your cue from the Board's document then you do have to -- then you do, in our review, do have to reflect that the validation was partial. If you don't want us to do that, of course, then -- then that -- that -- that we will -- it will be at your pleasure.

MR. GRIFFON: No, I -- I think those are our guidelines and -- and that's what I'm saying, you know, NIOSH has -- has -- has pulled a lot of different information. This is my -- my point of view anyway. NIOSH has pulled a lot of information. Came up short in some cases as Jim just said but -- but, you know, they have a fairly strong case, you know, for the internal section especially, and I think they put that forward. I think that you, Arjun -- I think SC&A appropriately should say, you know, that this is what it was. Is it, you know, and -- and you know, maybe to be careful with

1 adjectives but describe it as -- as what it is, 2 as what you per--, you know --3 DR. MAKHIJANI: Yeah. 4 MR. GRIFFON: -- perceive it to be. And, you 5 know, that it clearly wasn't, you know -- there -- there wasn't data, you know. There just 6 7 wasn't raw data available for every time period 8 for every, you know -- So I think present it 9 as is and then the Board has to weigh the 10 evidence I guess. You know, okay, it is 11 partial but there are powerful arguments made 12 here, you know. So I think we have to weigh that evidence so -- but I -- I don't think you 13 14 addressed, you know, from our policy document I 15 think you approached it correctly. Other 16 people may disagree with me. I don't know. 17 (No response) MR. GRIFFON: 18 I guess not. 19 DR. MAKHIJANI: Thank you. 20 MS. MUNN: This is Wanda. We have to at some 21 juncture come to grips with the issue revolving 22 around the original wording of our charter 23 which is more or less the definition of how 24 much is enough. There's no question we're

never going to have perfect information. Since

we're not going to have perfect information the issue is how much information can be considered relative to the overall issue so that we can define an acceptable limit. We're not going to be able to define acceptable limits in each case. I don't believe that's possible. So we're back to the same question, how much is enough? And you're right.

MR. GRIFFON: Right.

MS. MUNN: I believe this is a question that the Board has to face every time we have an SEC and this one is probably more difficult than some other decisions the Board must make.

MR. GRIFFON: Yeah, you're right, Wanda. And - And, yeah, I think we can -- we're probably
only going to be able to take the policies so
far but then -- then there -- there are going
to be sort of site-specific things that have to
weigh into that definition of how much is
enough. But yeah, you're -- I don't disagree
with that at all. So can we move on to number
four? Have we -- Arjun or Jim?

DR. MAKHIJANI: Sure.

MR. GRIFFON: I think we touched on this a little. I'm sorry to cut you off, George. I -

1 - I just was trying to keep going item by item.

MR. KERR: That -- that's really -- that's fine. I just wanted to clarify the fact that there was some dose -- beta dose in -- in '50 and '51.

DR. MAKHIJANI: Yeah, we -- We looked at the external dose, the database and the internal one. I at least -- I -- I at least did not find any non-zero entries, and there may be one. I can't say that I looked at every single one but I did not find any non-zero entries in -- in the gamma or beta entries in the CER database.

MR. KERR: Well, I --

DR. MAKHIJANI: There are non-zero entries in several ones, all -- all of which happened to be for 1951 so I don't know about 1950 in the delta view database that some of which I put in a table. There are also non-zero beta doses in the delta view database which -- which I did not compile but I just mentioned them -- mentioned them in the text. And -- And so there -- and I -- and I believe in the -- in the communication that NIOSH sent us this week

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NIOSH did acknowledge that there are these zeros and -- and had some kind of preliminary idea of where they might be coming from although they said the origin of these zeros is unknown, and that maybe that maybe they're due to some computer glitch. That -particular thing did -- did -- we discussed it and that -- that raises the bigger question, because that was a little bit of a surprise I have to say in that the -- the later years' validation seemed -- seemed to work from '52 onward to the extent for the various things. There are some differences and as NIOSH has pointed out, most of those differences appear to be claimant favorable. I think I cited that on page 13 or someplace in -- in the details. But -- But this question of why those zeros were there in '50 or '51 we -- we didn't have any -- any idea where they came from but now NIOSH said they might be due to a software problem and that does raise a question of what -- where else that software problem might show up and what the DOE did to -- to -- to ensure that -- that these problems were not occurring in a widespread way in the -- in the database.

1 To the extent that the evaluation was done for 2 '53 mostly it -- it did appear to be okay. 3 DR. NETON: The software problem was related to 4 delta view database though, not --5 MR. KERR: I don't think it was -- I also got printouts from the Y-12 database and -- and 6 7 knowing that the Y-12 database is what CER has, 8 I asked Y-12 to look for me back in the early 9 years. And if you look in both of them there 10 clearly is beta dose data for '50 and '51 in 11 both the printouts from the CER and the Y-12 12 database. MR. GRIFFON: Well, when you said, George, for 13 14 15 MR. TANKERSLEY: -- Tankersley -- and George is 16 absolutely right. There are positive data from 17 1948, 1949, 1950, 1951 and onward. And --18 DR. MAKHIJANI: Bill, in the CER database? 19 MR. TANKERSLEY: (Inaudible) have not looked at 20 the correct fields. 21 MR. GRIFFON: Bill or George, I'm just -- I'm 22 just doing this right now and -- and I want a 23 clarification. 24 DR. MAKHIJANI: I'm going to go off, too, 25 because maybe --

1	MR. GRIFFON: You're looking at
2	DR. MAKHIJANI: (inaudible) and I looked at
3	the wrong one.
4	MR. GRIFFON: Well, you're looking at at the
5	S-millirem field?
6	MR. TANKERSLEY: The skin and the penetrating,
7	that's exactly right.
8	MR. GRIFFON: Okay. Because I have '50,
9	there's no penetrating. There is skin but
10	there's no beta beta gamma fields is all
11	zeros.
12	MR. TANKERSLEY: That's correct as George said.
13	MR. GRIFFON: Okay.
14	DR. MAKHIJANI: No, I have believe what I
15	said is that all of the beta gamma fields are
16	zero. That's what is in our report. And among
16 17	zero. That's what is in our report. And among the other two fields, the S-millirem and P-
17	the other two fields, the S-millirem and P-
17 18	the other two fields, the S-millirem and P-millirem I did not observe any non-zeros in the
17 18 19	the other two fields, the S-millirem and P-millirem I did not observe any non-zeros in the P-millirem but I did observe some in the S-
17 18 19 20	the other two fields, the S-millirem and P-millirem I did not observe any non-zeros in the P-millirem but I did observe some in the S-millirem.
17 18 19 20 21	the other two fields, the S-millirem and P-millirem I did not observe any non-zeros in the P-millirem but I did observe some in the S-millirem. MR. GRIFFON: That's correct.
17 18 19 20 21 22	the other two fields, the S-millirem and P-millirem I did not observe any non-zeros in the P-millirem but I did observe some in the S-millirem. MR. GRIFFON: That's correct. DR. MAKHIJANI: Since the gamma and beta are

1 millirem button. No non-zero readings in the 2 P-millirem. That's sort of the substance of 3 the comment there. 4 MR. GRIFFON: Maybe we just need a clar -- can 5 -- George or Bill, can you clarify that? 6 MR. TANKERSLEY: Through the years people 7 reported the -- the doses in those two sets of 8 fields differently and I do not know why that 9 And to understand the data in that -- in -10 - in that set, which again is the Y-12 set; 11 everyone continues to refer to it as the CER 12 database. 13 MR. GRIFFON: Right. 14 MR. TANKERSLEY: It's simply a copy, of course. 15 You have to -- You have to get into it deeper 16 than -- than perhaps some have. But there are 17 definitely positive values in -- in 1950 and 1951 and then, of course, I'm assuming everyone 18 19 is pretty comfortable with the 11,000-plus 20 records, you know, in '48 and '49, PIC data and 21 -- and film badge data. The -- The -- The records in the '50 and '51 22 23 are not from the neutron data. 24 MR. GRIFFON: So -- So it's sort of unknown 25 why the beta fields would be zero and the S-

1 millirem would have positive value. 2 MR. KERR: Well, I guess what you've got to do 3 is -- is for some of those years you also got 4 to go look at the -- sometimes it was the --5 the penetrating and then -- in the skin. 6 you can go to those and you can clearly 7 separate those doses out. Now, you know, 8 that's where in the early years, you know, I 9 guess the -- as a matter of fact what I do have 10 from Y-12 is slightly different than what I got 11 from CER. But from Y-12 for each of the years 12 starting back in 1950 up through I think 2003 or '04 gives me penetrating, they give me the 13 14 skin and they give me the neutron. And from 15 those three -- those items I can go back 16 through and separate out such things as -- as 17 the gammas and the betas and the neutrons. DR. MAKHIJANI: I -- I have this database open 18 19 before me. 20 MR. GRIFFON: Yeah, me, too. 21

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MR. GRIFFON: Yeah, me, too.

DR. MAKHIJANI: And the file, table Y-12, PBL
Y-12, External 1950 to 1957. Every -- Every
single entry in the beta and gamma dose -well, there's one I believe in the gamma, not
in the beta that I just found that is non-zero.

1	MR. KERR: Okay. What about your skin and your
2	penetrating?
3	DR. MAKHIJANI: Well, the the
4	MR. GRIFFON: Nothing for penetrating.
5	DR. MAKHIJANI: There are a number of entries
6	as I said in the report in the skin that are
7	non-zero but no entries in the penetrating that
8	are non-zero. All zeros. And if all of the
9	entries in beta and gamma are zero then one
10	must presume that the only remaining source of
11	dose would be neutron that would appear in the
12	other two fields.
13	MR. KERR: I think the problem early on then is
14	the way that the doses were recorded.
15	DR. NETON: Yeah, Arjun, I think that, you know
16	I think you're
17	MR. KERR: That's the problem right there
18	DR. NETON: interpreting those fields
19	MR. KERR: is the way they were recorded.
20	They just recorded some as skin and some as
21	penetrating in the earlier years.
22	DR. NETON: Right. Rather than fill in the
23	beta gamma fields independently
24	MR. KERR: Yeah.
25	DR. NETON: they just report skin and deep

1 which is a fairly common notation for doing 2 dosimetry. 3 MR. KERR: And it's fairly common at a lot of 4 sites just for getting your doses that way. 5 DR. MAKHIJANI: Shouldn't you have a non-zero 6 badge reading to enter something in the other 7 two fields? 8 MR. KERR: Well, no, it was originally how it 9 was --10 DR. MAKHIJANI: (Inaudible) was a zero. 11 MR. KERR: It was originally how it was 12 recorded probably on the cards that went into 13 the database. 14 Right. In other words --DR. NETON: 15 It was recorded as skin unless they 16 put it in the skin column. If it -- If it was 17 recorded in gamma beta they subbed them to get 18 the skin dose. 19 DR. MAKHIJANI: We -- We were asked to 20 evaluate what we saw in the CER database and 21 whether it was validated or not. The -- We --22 We did find non-zero beta and gamma entries in 23 the -- in the beta and gamma column. 24 explicit columns in the delta view database and 25 the record numbers for that are cited in the

1 report. All of the corresponding values for --2 for those times in the -- the database are zero 3 and -- and so -- and -- and NIOSH then did send 4 us a document saying that the database does not 5 -- the CER database for those years does not 6 appear to be correct and the origin of these 7 zeros is unknown. 8 MR. GRIFFON: Yeah. I -- There's two issues 9 going on here, too, Arjun, right? The delta 10 view compared to the database --11 MR. KERR: Yeah. 12 MR. GRIFFON: -- versus just the database 13 itself? 14 MR. KERR: Yeah. Yeah, I agree that their --15 their data in -- in the delta view that does 16 not appear to be in the Y-12 database but I'm 17 saying that the reason you're seeing zero in 18 some of those columns were the things -- the 19 way things were recorded back in the early 20 years. 21 MR. GRIFFON: So in -- in '53 it changed, 22 George, is what --23 MR. KERR: Well --24 MR. GRIFFON: I mean obviously. I'm looking at 25 the database and in '53 you have beta -- I got

1 one example here. Beta is 188; gamma 4901, S-2 millirem is 5089 which is the sum of those two. 3 MR. KERR: Right. 4 MR. GRIFFON: And then P-millirem is 180. 5 MR. KERR: And I think in some of the earlier 6 years they may have already summed them and had 7 no way to split them back out so, you know, 8 they may have just put them in as skin dose. 9 MR. GRIFFON: And then in this particular case 10 P-millirem is 188 which it probably should be 11 4901 but -- but that's another issue I quess. 12 DR. NETON: Yeah, see, I -- I think what's 13 clear is that there's the -- the CER database 14 had to accommodate all ways of reporting so 15 there are fields there that may not have been 16 used in the early years which is what George is 17 trying to say. MR. GRIFFON: Right, right, right. No, I -- I 18 19 gather that, Jim. Now, here's another 20 question. When you did your models did you do 21 the -- which fields did you use? Did you use 22 certain ones throughout or did you --23 MR. KERR: Oh, we -- We -- We used the beta 24 gammas fields when we did our models. 25 MR. GRIFFON: Okay.

1 DR. NETON: But that was only after a certain 2 year. We didn't use any of the --3 MR. GRIFFON: That's right. 4 DR. NETON: '51 data for the model. 5 MR. KERR: Yeah. 6 MR. GRIFFON: Okay. Okay. DR. NETON: See, that's the other point here is 7 8 that the co-worker model is not based on these 9 data at all. The only relevance of this issue I think is if we received -- if we have a 10 11 claimant who has monitoring data in '50 and '51 12 then -- and then maybe Arjun has a point. But 13 I think there's a strong argument to be made 14 why there are zeros in the beta gamma field in 15 the early years based on changes in reporting 16 practices when the database covers all years. 17 MR. GRIFFON: But that -- that's just -- that's 18 just speculation, Jim. 19 DR. NETON: Well, I --20 MR. GRIFFON: You don't have evidence of that. 21 You're just saying that it could have happened. 22 DR. NETON: I don't, but it certainly makes 23 sense to me. 24 MR. GRIFFON: Yeah, I don't disagree. It's an 25 argument. But I don't think you have -- run

1 that data --2 DR. NETON: I think it's just as speculative, 3 Mark, to say that -- that zeros there imply 4 that the beta -- the skin and deep dose are 5 invalid. DR. MAKHIJANI: Well, this -- This is --6 7 MR. GRIFFON: No, I'm not trying to imply that 8 -- I think part of the issue for me was '50/'51 9 is that you have S-millirem data and you have 10 no penetrating data at all and no gamma or 11 beta. 12 DR. NETON: Well, I think that's not 13 inconsistent with low level beta exposures --14 DR. MAKHIJANI: It's --15 DR. NETON: Or below the detection limit of the 16 badge. I mean hopefully they would --17 MR. GRIFFON: And they just weren't recorded in 18 the beta field is your argument? 19 DR. NETON: Yeah, sure. 20 MR. GRIFFON: You know, that's a possibility. 21 MR. KERR: Yeah, the problem is is back in the 22 early days they were changing badges every 23 week. And you can measure beta sometimes, I 24 mean if your LD -- your lower limit of 25 detection is -- is 30 you could probably

1 measure betas but on your gamma dose it may 2 show up as zero. 3 DR. NETON: Right. But you --4 MR. KERR: And -- And I mean, you know, the 5 beta exposures were really what was concern in 6 the early days. And with the -- with the 7 people in -- that working with in -- in the 8 foundries in natural and depleted uranium. 9 I'm not surprised that you see all these zeros 10 for gammas. 11 DR. MAKHIJANI: This is Arjun. 12 DR. NETON: It's not about the -- the ten to 13 one --14 DR. MAKHIJANI: I believe there are two separate issues here. The delta view documents 15 16 from 1951 that I've seen clearly are from that 17 period so they should reflect the way in which 18 doses were recorded in that period. 19 They have four fields in the delta view database. They have beta, they have gamma, 20 21 they have neutron and they have extremity dose 22 if I remember correctly. And the -- there are 23 -- there is a corresponding column for beta and 24 gamma in the CER database. And when you

compare those two things the -- the fields with

1 the identical headings, the entries do not 2 match. I believe that what -- the 3 interpretation of what's in the SM and PM -- P-4 millirem and S-millirem is a different issue. 5 The -- The -- That's how, you know, how you use the dose information for dose 6 7 reconstruction. The -- The point of that 8 particular section is are the data -- is this 9 database good for the years '50 and '51? 10 the observation is that for those years the 11 beta and gamma fields do not match the delta view database and therefore they do not match 12 13 the raw data records that are available so they 14 have to be declared to be invalid. I do not 15 see how these beta and gamma entries can be 16 considered reasonable or appropriate or correct 17 in any way. 18 DR. NETON: Okay. 19 DR. MAKHIJANI: I fail to see that. 20 DR. NETON: If we grab that argument, Arjun, 21 and I'm not saying I'm willing to do that, but 22 if we did what's the practical significance of 23 this? 24 MR. GRIFFON: It's the reliability of the

overall database I think.

1 DR. MAKHIJANI: Yeah. The practical 2 significance --3 DR. NETON: Well, no, no, no. 4 DR. MAKHIJANI: There's no explanation for it. 5 DR. NETON: No, you're saying that 1950 and '51 6 are invalid and that's your position. say that '53 appears to be okay. 7 8 DR. MAKHIJANI: Right. 9 DR. NETON: And so what we're saying is if --10 if the practical significance is that -- that '50 and '51 are invalid we have a co-worker 11 12 model which we're going to discuss yet that --13 that fills in those values so what -- I don't 14 know what the practical significance of the 15 argument is anyway. DR. MAKHIJANI: Well, Jim, until -- until we 16 17 got your note about -- which -- which assessed 18 why these zeros might have been there I -- I 19 don't know that I could have -- have given you 20 a more nuanced answer to that question but 21 since there is the issue of whether there was a 22 software glitch in how these zeros occurred it 23 -- it definitely raises in my mind at least the 24 question of what else did this software do and

is the '53 validation that you did, which --

1 which appropriately was all right, does -- do 2 you need to do some more checking or not? 3 it was a software glitch what -- what's the 4 investigation of the software or what is the 5 other explanation for this problem? There's got to be an explanation for -- for why zeros 6 7 were entered when the raw data from the time 8 clearly had non-zeros in these same fields. 9 MR. TANKERSLEY: This is Bill Tankersley. You 10 need to discount the comment about a software 11 problem producing those zeros. That person 12 simply misspoke when he put that into the 13 report. As I explained probably a month or two 14 ago, there are database managers. I'm talking about a program that will insist in a numeric 15 16 field putting in zero instead of nulls and the 17 new programs won't insist on that. But there's 18 not a software error that put in zeros when 19 there should have been, you know, positive 20 numbers. So any discussion about, you know, 21 that is -- is not useful at this time. 22 Okay. Well, here -- Here --MR. GRIFFON: 23 Here's, Jim, just to -- to -- from my 24 perspective, here's what I'm looking at with 25 this item. Is -- is the weight of the overall

1 evidence for demonstrating the reliability of 2 the -- the Y-12 or as we're calling it CER 3 database? And, you know, the way I look at it 4 right now is you have several cases -- several 5 people from the delta view in '53 that you backtracked and -- and found doses to be in 6 7 agreement -- in pretty strong agreement but 8 then you have all this in '51 that's in 9 disagreement so -- and then that's all we have. 10 And, you know, that's my concern is that we're 11 -- we're -- I think we're a little thinner on 12 our --13 **DR. NETON:** Okay. 14 When Wanda asks how much is MR. GRIFFON: 15 enough, you know, I think -- I feel like our 16 arguments are a little thinner on this -- the 17 external database than they are for the 18 internal database. 19 DR. NETON: Well, right. We couldn't -- We 20 couldn't go back and find the original data but 21 22 MR. GRIFFON: I'm not saying you didn't make 23 all kinds of effort, you know. I'm just --24 DR. NETON: Right. But again, you know, we 25 have -- we're not relying on anything in the

early years for reconstructing doses for workers. I mean we have gone, you know, George Kerr has demonstrated pretty conclusively that the data that we have in those years do not fit any good distribution and so we're not using them to -- to reconstruct doses. Now, when we get into the '56 time frame, I don't know. I guess we're going to -- you're going to -- the argument is that if '50 and '51 don't match and '53 did then we need to go back and look at more years after '53. I mean is that what we're hearing? And then if we can't what's the ultimate answer? I don't know.

MR. GRIFFON: Well, yeah. Yeah, I'm just saying that -- that the SEC -- I know you're not using that earlier -- that early data but it is all part of the database so -- and we've heard explanations of why this might have occurred; you might be right. But, you know, and so far we have sort of two, yeah, two pieces to -- to answer this question of reliability of the -- of that '50 to '57 database. Now, you know, later -- I mean you can't -- and we've talked about this before, Jim. You can't sort of have it both ways with

this. I mean in the other case you -- you pulled some data from the '70s to demonstrate the -- the '50 to '57 period of the urinalysis database is good, you know, so --

DR. NETON: Well, but Mark, we've looked at '53 and we've looked at the '70s now. I guess I'm hearing the intervening years need to be checked. I mean that's what I'm hearing. I don't know what else we can do.

MR. GRIFFON: I'm just making observations about where we're -- where we're at right now. I'm not saying whether we have to or not.

MR. KERR: The only importance of that data back -- that we had back in '48, '49, '50, '51 period is if we take our co-worker model that we have and -- and apply it. We're -- We're making conservative estimates of what the doses were back in those days because, yeah, we're way above the doses people received. And, you know, that's the only reason I think they're important is it's a basis of comparison for what we're predicting doses to be. And everything I see we're very considerate and very claimant favorable. That's the importance of the data back in early --

1	DR. NETON: I think we understand that, George.
2	But what Mark is saying is are the data that
3	we've used for the co-worker model even valid
4	now? And I'm not sure that '51 and '51
5	mismatch after we've done a '53 comparison and
6	a `70s comparison is enough to invalidate
7	MR. GRIFFON: But you didn't do a `70s
8	comparison, did you, for external?
9	DR. NETON: I thought that's what we just said
10	we did.
11	MR. GRIFFON: I I said for the internal you
12	brought in some data from the '70s.
13	DR. NETON: For the internal, yeah.
14	MR. GRIFFON: The urine punch cards.
15	DR. NETON: Yeah.
16	MR. GRIFFON: So right now you have one one
17	data point, '50s, you know, one one set of
18	results which which I I, you know, it's
19	good. It's encouraging that they match. But
20	I'm, you know, I'm just I'm just throwing
21	out there, Jim. I'm not saying you have to go
22	back and do more. I'm just saying that, you
23	know, is how much is enough?
24	DR. NETON: Yeah. I I agree. And I don't
25	know if there's much more we can do.

1 MR. GRIFFON: Right. 2 DR. NETON: And that's the problem. 3 MR. GRIFFON: And then I think you -- you use 4 that and you present to the Board just sort of 5 the same arguments that you've used along with 6 what George said that that, you know, those 7 early periods the co-worker model is going to you believe, you know, be very conservative 8 9 anyway, yeah. So all those -- all those 10 bolster your arguments sort of. 11 DR. NETON: Yeah, I think that -- that's pretty 12 much our position at this point. 13 DR. MAKHIJANI: Could I ask George a clarifying 14 question, please? If -- If the co-worker 15 model is to be judged to be claimant favorable for '48 and '49 for internal dose where we have 16 17 no data and for external dose for '50 and '51 18 where all the entries are zero, any non-zero 19 entry would appear to be claimant favorable. 20 Well, you see -- Okay. MR. KERR: What I'm saying --21 22 DR. MAKHIJANI: How do you make a judgment --23 how do you make a judgment about claimant 24 favorability when the -- when the database 25 itself doesn't appear to contain material

contents?

MR. KERR: Okay. We do have '48 and '49 data. We do not -- and here we come back to your argument. We -- If you go back to the '50/'51 data you do not have entries as true for the gammas and betas separately. But you do have penetrating and you do have the skin dose. And my contention is you can derive or you can get estimates of what these people had from those two. In the case of -- of part of it was beta. It was penetrating. It was gamma. You can subtract and get some idea of what the beta doses were people were receiving. And you can compare with those.

DR. MAKHIJANI: But the fact is that all of the penetrating dose entries are zero.

MR. KERR: That's okay. But we -- we can still get beta doses out of there. We're -- We're -- We're developing a beta dose model, too.

And you still have the '48/'49 data. And as a result of it, even in the delta view, you say those are zero. We still have the delta view to go to to compare with doses that are recorded in there with the co-worker model.

And -- And even doing that they look very

1 conservative. 2 MS. MUNN: Here's Wanda. It appears that one 3 could make a very good case of having verified 4 the data for an immediately subsequent year, in 5 the CER database. And (inaudible) year in the CER database (inaudible) the type of recording 6 7 that you see in '50 and '51 clearly was 8 overcome in 1953 and therefore the 9 extrapolations that are made from subsequent 10 data (inaudible) in the obvious absence of 11 unusual events (inaudible) in that '50/'51 12 period. Do we have unusual events recorded in 13 that period? I wasn't aware of any if we did. MR. GRIFFON: And Wanda, can I ask, are you on 14 15 a speaker phone? 16 MS. MUNN: Yes, I am right now. 17 MR. GRIFFON: Because I hear every fourth word 18 or so. You're cutting in and out on me. 19 don't know if that's happening to everybody but 20 21 MS. MUNN: It must be happening to everybody. 22 One never can trust a speaker phone. 23 MR. GRIFFON: Sorry. 24 MS. MUNN: So did what I say come through 25 enough to make any sense?

MR. GRIFFON: Yeah. Yeah, I -- I think so. I mean I, you know -- if -- I guess it comes down to, you know, it would be more concerning to me if the -- the '50/'51 issue and not matching was in the middle of the time period, you know, not on the front end I suppose. I don't know but, you know, I come back to you have some, you know, some data in '53 that are supporting the argument of reliability and -- and I suppose this letter that says the DOE accepted this as the database of record, correct? I mean that was for both external and -- and internal, correct, Jim?

DR. NETON: Right. I believe so.

MR. GRIFFON: So, you know, it comes down to the -- the weight of the evidence.

MS. MUNN: We know from our own experience and from information that we have from individuals who were in those positions at that time that the particular period we're talking about, the '48, '49, '50, '51 period was a period of enormous change not only in plant process but in administrative process and in health physics process as well. We have some data prior to that confusing time and significant data

following that time. If we've been given two very valid points of comparison following that time that agree, then the question becomes very simply is that reliable enough for the Board. It's reliable enough for me.

When we have times that are -- are confusing for everyone and have differing methods of -- of computation, differing methods of calculation, differing methods of recording then we must either say as one argument has gone, that we can't use any of that data; or we must say those problems were worked out and all data from there on is reliable. That essentially in my view is the question we're going to have to put before the Board.

MR. GRIFFON: That -- That -- Yeah, that's the question and it's just, you know, be -- being convinced of those arguments she just made. That's -- That's the important part and I think the stronger the arguments can be made, the -- the better, you know, so I mean -- so look at this, you know. It seems like what has been mentioned for '50 and '51 are -- are likely explanations, you know, but I don't know that I've seen documents indicating that, you

know. So -- So there's good explanations, you know, possible good explanations. I don't know that we've seen that as, you know, any health physics report saying or any -- and I don't know that there would be any report saying that that kind of thing happened, you know, and this is why.

MS. MUNN: No, but it may be helpful to put that rationale very crisply in print and even if it's just a letter report to provide for the Board because what we're -- the agony we're going through here in the working group is not going to be --

MR. GRIFFON: Right.

MS. MUNN: -- manageable in the Board setting.

MR. GRIFFON: Right. I agree. I agree. I mean, yeah. And I think we -- I think what I'd like to do from the working group is summarize where we're at on different items and I'm not sure how much I'm willing to connect the dots, you know. But we'll lay out the -- the facts as they've been presented to us and the arguments that -- that have been presented to us. And then I think, you know, we present that to the Board and it's, you know, so -- so

1 that we don't have to go, you know -- obviously 2 we don't want to go through all the details at 3 the Board level. I -- I agree, Wanda. 4 we'll -- we'll -- we'll have to work on that. 5 MS. MUNN: My personal feeling is that such report from us is going to be crucial in the 6 7 discussions in Denver. 8 MR. GRIFFON: Yeah, so --9 MS. MUNN: (Inaudible). 10 MR. GRIFFON: Yeah, we're going to have a long 11 weekend. 12 MS. MUNN: -- the language needs to be right 13 and very clear and very factual. 14 MR. GRIFFON: Yes. Okay. And factual, I 15 agree. Okay. I don't know that we can -- can 16 we do any more on this topic? I don't know. I 17 missed -- Arjun, one thing I might want 18 clarification on from George is just in looking 19 at this database if -- if I'm looking at P-20 millirem in the later years when there's 21 actually recorded numbers --22 DR. MAKHIJANI: Yeah. 23 MR. GRIFFON: -- that should in most instances 24 be equal to the gamma or gamma plus neutron or 25 is there a more sophisticated algorithm?

1 DR. MAKHIJANI: As we understood it the P --2 the P-millirem dose column should include the 3 gamma plus the neutron dose, yes. 4 MR. KERR: That's right. 5 MR. GRIFFON: Okay. MR. KERR: And then the -- the -- where they 6 have millirem or the skin dose it should be the 7 8 gamma plus the neutron plus the beta. 9 MR. GRIFFON: Right. 10 DR. MAKHIJANI: That -- That's exactly how we 11 interpreted it and wrote it up. 12 Okay. Okay. Anyway, yeah, and MR. GRIFFON: 13 just glancing at a few of those I just spotted 14 some that were -- but I'm -- I can't do this 15 and talk on the phone but I think there is some 16 interesting ones that the gamma and -- and 17 penetrating don't seem to line up but I'm --18 and there's no neutron dose on those ones that 19 I'm talking about so -- but -- and that's in Anyway, that -- that's sort of why I was 20 21 wondering which -- which columns were actually 22 being used in the co-worker model --23 MR. KERR: Yeah. 24 MR. GRIFFON: -- out of -- out of those data 25 and is it the -- which columns are being used?

1	Which Which parameters?
2	MR. KERR: We used We've used the gamma and
3	the beta.
4	MR. GRIFFON: Gamma and beta? Okay.
5	DR. MAKHIJANI: Then you apply the neutron to
6	photon ratio, right?
7	MR. KERR: No.
8	DR. MAKHIJANI: That's what was in the sample
9	dose reconstructions anyway.
10	DR. NETON: That was for a person who was
11	potentially exposed to neutron but not
12	monitored.
13	DR. MAKHIJANI: Right. Right. I mean in your
14	co-worker model.
15	DR. NETON: Well, no. The co-worker model is
16	for is for gamma and is for beta.
17	DR. MAKHIJANI: Right. Right. For somebody
18	who is not monitored for neutrons you use a
19	neutron to photon ratio.
20	DR. NETON: We have done that in the example;
21	that's correct.
22	DR. MAKHIJANI: Right.
23	MR. GRIFFON: Okay. Should we move on to five?
24	I don't think we're going to get through all
25	eleven of these before

1	DR. NETON: I think, Mark, some of these
2	MR. GRIFFON: Yeah.
3	DR. NETON: next couple we've talked about
4	in relation to internal dose reconstruction and
5	and the co-worker model that used the 1952
6	bioassay data to back-calculate the maximum
7	intake that could have occurred based on, you
8	know, what we're observing in '52.
9	MR. GRIFFON: That's five and five and six,
10	right?
11	DR. NETON: I think five and six
12	MR. GRIFFON: Yeah.
13	DR. NETON: are related to that issue. And
14	in fact in number six I think SC&A said that
15	example five does not address the issue of
16	unmonitored worker. There is a clear co-worker
17	model dose intake applied there. I'm not sure
18	where they they got that idea.
19	DR. MAKHIJANI: I'll I'll go back and check
20	that; maybe if it's my mistake it will be
21	corrected.
22	DR. NETON: Yeah. I mean I think that the
23	confusing part of number five where it says the
24	worker was monitored and it only implied that
25	he was monitored for a certain period prior to

1 '50. Of course he could not have been 2 monitored and we applied the co-worker intakes 3 so they're there. 4 DR. MAKHIJANI: Okay. 5 DR. NETON: Okay. DR. MAKHIJANI: Well, you said the worker was 6 7 monitored and you assumed zero -- zero bioassay 8 results. 9 DR. NETON: Well, right. But see it was a 10 little misleading. He was monitored after 1950 11 12 DR. MAKHIJANI: All right. 13 DR. NETON: But there is no monitoring data 14 prior to '50 so we --MR. GRIFFON: -- there was just confusion. 15 DR. MAKHIJANI: Okay. So if -- If there was a 16 17 misunderstanding that arose from how the thing 18 was written up I guess. 19 DR. NETON: I believe so. DR. MAKHIJANI: Okay. All right. I'll go back 20 21 to that. But -- But the only point was I 22 think here that we haven't discussed in 23 relation to five and six is that it's the piece 24 of -- of the operations at Y-12 that's 25 indicated in the site profile terminated in '51

if I remember correctly that was called a recycle and salvage, etcetera, where they were reconditioning pieces of -- of -- of the -- of the site for -- for new operation. And then those operations were terminated at that time and never redone. I -- I have not seen anything, any calculations that show that the available data for from '52 onward would bound the internal doses for those particular workers so there's a question -- there's an explicit question about the salvage and recycle operations in those three buildings that are named, 9206, 9207 and 9211.

DR. NETON: Right. But -- But we discussed this a little earlier. We took the urine data from the workers in '52 who would have been working in those time frames and assumed that they had chronic intakes all the way through those periods and -- and did a bounding analysis using what was being excreted in their urine in 1952.

MR. CHEW: Jim, this is Mel. Arjun, I think -you know, I don't -- I fell into the same -- a
little bit of the same trap that -- well, I was
claryifying (inaudible) in submitting the

1 report. But people would talk about recycled 2 uranium and recycled uranium there are --3 looking at the details there are two different 4 things as you probably, well, you well know. 5 DR. NETON: Yes. They -- They basically out of the 6 MR. CHEW: machine shops they tried to save every piece of 7 8 uranium they had and they recycled it and they 9 called it recycled uranium. And then in 1952, 10 even late '52 was the first entry of what you 11 and we have been talking about as RU with the 12 contaminants of the neptunium and plutonium and technetium in here and I -- I just want to make 13 14 sure that we -- we often fall into the same 15 trap here that I did earlier on, too. 16 DR. MAKHIJANI: No, no. I -- I didn't. 17 didn't misunderstand that. 18 MR. CHEW: Okay. 19 DR. MAKHIJANI: I am not -- I am not calling 20 recycled -- in fact I didn't even think about 21 it until you mentioned it. 22 MR. CHEW: There's recycled and there's 23 recycled. 24 MR. GRIFFON: Right. Right. Right. 25 DR. MAKHIJANI: No, no. This -- I'm not

raising a recycled uranium trace contaminants issue.

MR. CHEW: Okay. And so therefore if it's recycled uranium in the earlier days, then the bioassay for uranium was certainly bound and I was making sure that you were not talking about the contaminant, okay?

DR. MAKHIJANI: Yeah. No, I'm talking about
the specific jobs that occurred in those years
--

MR. CHEW: Uh-huh.

DR. MAKHIJANI: -- that stopped, you know, in the conditioning of the facilities and cleaning up the places and so on. There was a kind of a decommissioning and recommissioning operation as I understand that went on. And -- And I -- I have not seen where the workers were involved in those specific jobs which seemed -- which seemed to involve different exposure conditions than the production workers. It seems to me that -- that job-specific analysis is necessary to show that -- that you've covered those workers with your co-worker analysis. And that's the thrust of the comment here. It isn't that -- I didn't mean that the co-worker

model would not bound these doses. It's just that for those workers do we have the information say from '50 or '51 for those job types to demonstrate that you've got them covered in your co-worker model.

DR. NETON: But I think if the issue is if they bounded it then the answer is we have.

DR. MAKHIJANI: Yes. No, I -- I didn't see that -- that any -- any -- any demonstration for those groups of workers. Perhaps it's there and I missed it but -- but I -- I'm not aware that such a thing has been done. But as -- maybe -- maybe it's just my -- my not having seen the right document.

DR. NETON: What we're saying though, Arjun, is that of all the workers that were there in '51 and '52, they're leaving urine samples and -- and these are the workers, these are the production-type workers, the workers who would have been working with the uranium. And we've taken those workers and -- and -- and looked at their urine samples and said if they were working in '48 and '49, how much could they have breathed then and -- and still be excreting what we're measuring in '51 and '52.

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But we -- We're trying to bound it based on using the workers as their own sort of standard.

DR. MAURO: Jim, this is John. I -- I understand where you're going and I think I see the subtlety of the -- the issue that's now on the table. Again going to table 3 on page 15. Let me see if I can articulate this. have here is you've got this urinalysis data for 1950 and '51 for 166 and 367 employees. That urine data -- Now -- Now, we also could look over to the second column. We see there basically is the same number of employees, '48, '49, '50. And of course, it increased in '51. But what I'm hearing you saying is we -- the -the 166 employees that were monitored, that the activity you're looking at in the (unintelligible) is the -- is the result of an integrated exposure that they -- that those workers experienced while they were working in 1948 and '49 and -- and I completely understand and agree of taking that tack. And it would certainly be a very good surrogate for the fact that the workers in '48 and '49 weren't monitored. If you're looking at that 166 and

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you go back in time and say that this is what they took in in order to get the -- whatever reading you're getting for the 166. The -guess the distinction now -- to get to the point where I think that there might be a distinction is -- is there a -- of the 2,248 workers that were working in '49, what I'm hearing is there might be a -- a subgroup of those workers that were performing activities that were substantively unique, whatever they -- the -- what I hear, recycle of the scrap or other operations that were substantively unique. And in effect you were saying that okay, that -- that's fine because we caught them in the 166 people that we did monitor in 1950. So I think what I'm hearing is that you've got it covered. It really then becomes a matter of, all right, you've got these 166 monitored employees and you -- and you have a worker that worked in 1949 and you're going to want to reconstruct what he might have inhaled. Now, if you were to take the high end of the distribution for the 166 you certainly would be placing an upper bound, perhaps an overly conservative upper bound. Or you could take

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the full distribution which you would argue would be a claimant neutral approach. What I'm hearing is that if there was a fundamental -if there were some activities going on in '49 and '48 that were not going on in 1950 among said subgroup, and though -- and there's reason to believe that that subgroup had activities that created a greater potential for them to be exposed, the implication would be that when you go to the 1950 data, the 166 people that were monitored, you would probably have to use the high end of that distribution to make sure you captured that subgroup. Alternatively if you could demonstrate there was nothing about the activities that were going on in '49 and '48 that were substantially different than -- than we're going on 1950 -- then I can see you using the full distribution. So I -- I guess I --I'm working my way through this as we're working the problem. I think you've got a tractable situation. I'm just not quite sure if, you know, do we have a situation in '49 and '48 where the activities were substantially different? What I'm hearing from Arjun is that there was such activities but I'm not quite

sure whether those activities created the circumstance which had a substantially high potential for exposure than let's say the other activities that were going on and that continued into 1950 and '51.

DR. NETON: I think -- Mark, go ahead.

MR. GRIFFON: I was just going to say, just to flip that around, do you have any reason to believe, Arjun, that these operations -- I mean you picked these out particularly because you thought that these may not be bounded by the approach or --

DR. MAKHIJANI: Well -- Well, I picked them out particularly for two reasons. One -- One is that since the co-worker model starts in 1952 the going back into the era where work that was being done that was different than these three buildings, I felt that the validity of that co-worker model should be applied to the job types in these three buildings because there was different types of work. And the second reason is, yes, you know, the decommissioning and recommissioning operation involved substantially contaminated equipment. They were dealing with scrap and recycling

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uranium and scrap recovery operations are often -- have often been pretty dirty, at least in those early periods. They -- They involved -involved kind of difficult work. If you go to Ames in 1945 for instance, you know, you -- you -- you have pretty highly exposed workers. there's no judgment here that the -- the -- the data from 1952 wouldn't bound the earlier doses but the kinds of job types were different and were of the type where significant exposures were certainly possible. I -- I think that demonstration has to be made and that's the point of the comment, not that the doses can't be reconstructed or -- or that this is an SEC issue. But it has to be ruled out as an SEC issue or by the construction of a specific demonstration.

MR. GRIFFON: Jim, do you know of any air sampling data during that time period that you might be able to use to make your argument to say that, you know, we -- we're applying two years of chronic or three years of chronic exposure up to when we have a urine sample, and here's the dose we would have received in air sampling, limited air sampling that we have in

1 these buildings suggests that, you know, we're 2 over-estimating if nothing, you know --3 DR. NETON: Right. 4 MR. GRIFFON: I mean is --5 DR. NETON: I'm not aware right now --6 MR. GRIFFON: That might be a way to --7 DR. NETON: But what I -- what I'm concerned 8 about here is --9 MR. GRIFFON : Yeah. 10 DR. NETON: -- why do we believe -- do we believe that all of a sudden in 1951 or '52 11 12 this is an entirely different work force that's 13 monitored? I mean that would have to be the 14 case for this to be invalid. DR. MAKHIJANI: No, that's not the argument. 15 16 DR. MAURO: I don't think we're saying that. 17 We're saying within the work force which were 18 the number of people were about the same 19 throughout those years. 20 DR. MAKHIJANI: Yeah. No, they doubled in --21 they went up. 22 DR. MAURO: There was a subset. 23 DR. NETON: But what my -- But my point is, 24 though, that if -- if that subset is included 25 in this analysis --

1 MR. GRIFFON: Then it's appropriate. 2 DR. NETON: Then it's appropriate and what John 3 said is true. It's -- It's a decision whether it's the 50th or the 95th percentile I mean but 4 5 6 MR. GRIFFON: Right. 7 DR. NETON: But if -- If this subset is 8 covered in this monitoring then these people 9 are their own long-term integrators of their 10 own exposure in 1949 and '50 or '48 and '49. 11 mean that's the whole concept here and I'm not 12 sure Arjun was quite grasping that. 13 DR. MAKHIJANI: Maybe not. MR. GRIFFON: Additionally I got to say --14 15 additionally I didn't --16 DR. MAKHIJANI: (Unintelligible) you know that 17 the recycle workers were there in the later 18 years and were monitored and therefore you know 19 what their exposures were and that you iden -- -20 - I -- I haven't seen the recycle workers 21 identified as a subset in the later years for -22 - for checking whether their exposures were 23 comparable to or less than production workers. 24 DR. NETON: Now, my point is -- is if these are 25 the same workers or similar groups of workers

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that were working in '48 and '49 -- I don't think they laid everybody off in '49 and hired new uranium --

DR. MAURO: Yeah, Jim, in a way I -- I see
exactly where you're going.

MR. GRIFFON: Right.

DR. MAURO: In the extreme, in the limit, and we're going to write this story as to what's the worst possible thing that can happen. Okay. Out of these 2,500 workers that were working there in 1948 there's this large group of them that were doing decommissioning work, that were getting these very large exposures and held large -- large -- large amounts of material, much larger than anything anyone experience, let's say from 1950 onward, and they all left in 1949 and we never caught them. And we never caught -- and so therefore their -- the urinalysis data that we picked up in '50 -- I would -- I for one will argue that that is a scenario that certainly would defeat your -your methodology. But I think it's really hard-pressed to postulate if such a thing occurred. So I guess I'm coming down where you are.

1 MR. GRIFFON: I would tend to agree with that 2 and --3 DR. NETON: So -- Okay. 4 DR. MAKHIJANI: I -- I believe I -- I've 5 stated what my issue was. 6 DR. NETON: Yeah. 7 DR. MAKHIJANI: And it's up to the Board, of 8 course, to go where it should. 9 MR. GRIFFON: Okay. And it's also -- I think 10 we've done five and six. What I'd -- what I'd 11 suggest right now is can we break for lunch and 12 then we'll pick up on seven and hopefully --13 because Rocky people are going to be on the 14 line at 2:00 p.m. or thereabouts. 15 DR. WADE: We can work some of them. 16 MR. GRIFFON: Well, yeah. Hopefully we can 17 complete Y-12 fairly quickly and not --18 DR. WADE: Right. 19 MR. GRIFFON: -- you know, and then get to 20 Rocky. Is that -- Is that okay with everyone? 21 DR. WADE: Okay. So back at 2:00 ready to 22 work. 23 **MR. GRIFFON**: 2:00 p.m. 24 DR. NETON: Okay. Great. 25 MR. GRIFFON: All right. Thank you. Bye.

1 MS. MUNN: Rocky, be back at 2:00. 2 (Whereupon, a recess was taken from 1:05 p.m. 3 to 2:05 p.m.) 4 DR. WADE: I think there were some Y-12 issues 5 open. I think some of our friends from Rocky 6 Flats are on the line but we need to do what we 7 need to do. 8 I think what I'd ask is if MR. GRIFFON: Yeah. 9 we can just try to conclude Y-12 and then move 10 into Rocky understanding that the folks from 11 Rocky are on with us. We didn't quite finish 12 this morning. We're going to try to wrap up. And I just -- just to -- I just want to go back 13 14 to five and six for one second, Jim and John 15 and Arjun. 16 MS. MUNN: Are Jim and John and Arjun on yet? 17 MR. GRIFFON: Oh, are they on? 18 DR. MAURO: John Mauro, I'm here. 19 DR. NETON: Yeah, NIOSH is here. 20 MR. GRIFFON: Okay. And SC&A is on? 21 DR. MAKHIJANI: Yes. 22 MR. GRIFFON: Yeah? Okay. For five and six, I 23 just wondered if -- the only question I had 24 there was I had mentioned whether NIOSH had any 25 data that could sort of, you know, such as air

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sampling data that could demonstrate for these particular I guess D&D salvage, whatever -whatever kind of workers they were, that this co-worker approach is bounding. And I -- I guess, you know, that may, you know, once and for all sort of put this -- this concern to bed. I mean I guess the -- the real question that's still out there, it seems as though if -- if those workers were in that monitoring pool then -- then the co-worker approach described by Jim may well be bounding. But if there was other data, you know, if this was followed up to -- to at least look at -- at the concerns as to whether they were monitored later, in the later years, you know, or a set of those people that did that kind of work were actually monitored. You know, it seems reasonable to believe that they might have been. And -- And a second follow-up might be, you know, is there any like summary air data in any of the HP reports that might say here's, you know, average levels and if we compare intake based on the co-worker approach versus air sampling data, you know, the co-worker model seems very claimant favorable or whatever. I --

1 that it would at least strengthen that case if 2 NIOSH could demonstrate that. 3 MR. TANKERSLEY: Hey, Jim. This is Bill 4 Tankersley. 5 DR. NETON: Yeah. 6 MR. TANKERSLEY: We certainly can identify 7 those people easily enough if you choose to go 8 that direction. 9 DR. NETON: Yeah, thanks, Bill. I think that -10 - that would be one approach to go back and 11 show that, you know, they didn't fire everybody 12 in 1950 and hire a new work of -- group of 13 uranium workers or something to that extent. 14 don't know about air monitoring data, Mark. 15 think in '48 and '49 it's going to be pretty --16 pretty small and then -- then you always get 17 into the issues of representativeness and 18 because it's BZ versus GA and for us to put 19 that --20 MR. GRIFFON: Yeah. 21 DR. NETON: It sometimes causes -- raises more 22 questions than it answers. 23 MR. GRIFFON: Right. 24 DR. NETON: But, you know, and I just recognize 25 that if -- if -- it's going to be Friday here

1 pretty soon and I'm flying to Denver on Monday. 2 MR. GRIFFON: Yeah, I know. 3 DR. NETON: And I don't know what we can 4 realistically expect by then but we will do the 5 best we can. We hear what you're saying and 6 all those are great strategies to try to -- to 7 bolster our position and we'll do what we can. 8 MR. GRIFFON: Okay. Okay. I just --9 DR. NETON: Yeah. 10 MR. GRIFFON: All right. Let's move on to 11 seven then I think, Arjun. If you can present 12 13 DR. MAKHIJANI: Yeah. Seven -- Seven is 14 partly the same issue as -- as five and six for 15 external dose in that except for the co-worker 16 model you've got 56 to 65 doses where the work 17 was completely different than these 18 decommissioning workers. And again I -- I'm 19 not sure what -- and then for 1950 and '51 you've got all the beta and gamma entries being 20 21 zero in the database. So at least I -- I 22 couldn't see where one would find a piece of 23 information to validate that co-worker model. 24 I'm not saying that it isn't valid or bounding

but that it hasn't been demonstrated to be

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bounding. I did take a look also at the number of records available and then looked at the fine print in -- in the NIOSH documentation and it seemed to me that while the -- the table in -- in the ER, table 6-2, says there are 11,000and-odd records, the number of -- there are there are -- the records that are counted are -- are four records actually and the -- the PIC records and the film badge records are all counted separately even though the film badge records are not regarded as reliable up to 1950. And then the film badge records are kind of questionable. Most of them are either zeros or limit of detection and it's not clear that there was -- NIOSH itself says, you know, that they were 30 millirem or zero entered and it seems both were used as the equivalents of limit of -- below limit of detection. And then the film badge data are not to be used because they were unreliable. So one's left with ionization chamber data and it seemed to me that the non-zero record -- I didn't do an actual count. I -- I -- I did a kind of a little bit of a sampling as to how many nonzero records there may be and -- and it seemed

1 like there were only about 1,000 or 1,500 2 records or one -- one per worker per -- per ten 3 weeks. And -- And that seemed a pretty slim 4 basis on which to compare the co-worker model, 5 especially for this group of workers so that's 6 -- it's sort of -- it's a little bit more 7 involved than the -- than the internal dose 8 question because there's no monitoring at all 9 for '48 and '49 on internal dose. 10 MR. KERR: I'd like to speak to that because I 11 think you're taking the fact that the film 12 badge data for '48 and '49 was unreliable. 13 You've taken that out of context. That's not 14 what -- That's not what the TIB says. It was 15 thought at one time it was unreliable --16 DR. MAKHIJANI: I quoted --17 MR. KERR: but we went back --18 DR. MAKHIJANI: I quoted --19 MR. KERR: to look at that data --20 I quoted the TIB actually. DR. MAKHIJANI: 21 MR. KERR: Now, but you took it out of context 22 is what you did because earlier it was thought 23 that that was unreliable. We went back and 24 showed that there was good agreement between 25 the PICs and the film badge data.

1 DR. MAKHIJANI: So are you saying you're using 2 the PIC at POC? 3 MR. KERR: No, we're not using it but --4 DR. MAKHIJANI: (Unintelligible) data were to 5 be used. The reason we went back and looked 6 MR. KERR: 7 at that data was so if we could see our co-8 worker model of predicting doses back in '48 9 and '49 was truly claimant favorable. And if 10 you go back and look at the '48 data and you 11 look at the PIC data and you look at the film 12 badge data and you compare with what we predict 13 back in '48/'49, our -- our estimates of dose 14 for the workers back in those days on the -- on 15 the co-worker model that we're using are 16 extremely claimant favorable. 17 DR. MAKHIJANI: Now, why did --Tankersley is on the phone. I guess maybe he 18 19 can -- he can explain his 1987 paper and -- and 20 whether I took it out of context. 21 quoted it saying -- I'm trying to find the 22 quote here. It's in the report somewhere. 23 MR. KERR: It's -- It's in the discussion 24 section. 25 DR. MAKHIJANI: Yeah. And where he said that

1 the earlier data were regarded as unreliable 2 and I -- I --3 MR. TANKERSLEY: (Inaudible) 4 DR. MAKHIJANI: Sorry, I can't hear. 5 MR. GRIFFON: We can't hear. UNIDENTIFIED: Can't hear. 6 7 MR. TANKERSLEY: (Inaudible) 8 DR. NETON: Bill Tankersley, are you on the 9 phone? 10 MR. TANKERSLEY: Yes, I am. I'm not quite sure 11 what paper he's referring to. 12 MR. GRIFFON: We've got a lot of interference 13 all of a sudden. 14 DR. MAKHIJANI: Well, it's table 6-2 in the 15 evaluation report. And --16 **UNIDENTIFIED:** -- that interference --17 DR. WADE: (Inaudible) I don't know what it 18 That's better. 19 DR. MAKHIJANI: Let me -- Let me see here. 20 Okay. Table 6-2 in the evaluation report for 21 '48 and '49 says that 3,599 records for 162 22 monitored employees in '48 and 7,893 for 49 23 monitored employees in -- in 1949. So I could not match up the 49 monitored. It seemed there 24 25 were more monitored employees than the 49 but I

1 couldn't resolve the differences. And then I 2 found the issues described in section 5.2 of 3 the SC&A reports above those records including 4 the statement from you as to the -- well, I 5 won't characterize it so you can -- about -about the quality of the film badge data prior 6 7 to 1950, referring to a 1987 paper by you. 8 MR. TANKERSLEY: Well, actually I don't 9 remember -- I don't remember writing that. 10 We've never questioned --11 DR. MAKHIJANI: Well, in -- in 0-TIB-47 on page 12 13 it says that the film badge readings prior to 1950 were "considered questionable because 13 14 of frequently changed procedures and a 15 perceived general lack of monitoring quality 16 control during this period". And I'll -- I'll 17 just open --18 MR. KERR: Bill? 19 DR. MAKHIJANI: Open the TIB because it sites -20 21 MR. KERR: Bill? 22 DR. MAKHIJANI If I remember correctly it sites 23 a 1987 paper by you. MR. KERR: No, it's an '82. It's '82 and it's 24 25 a memorandum to Shirley Fry (ph).

1 DR. MAKHIJANI: Let -- Let -- Let me 2 go to the TIB and so I can verify my memory 3 here. Okay, 47, page 13 -- page 13 -- yes. 4 Pre-1982, you're right, George. But it is 5 Tankersley, 1982. MR. KERR: Right. Okay. But now, read the 6 7 next to the last sentence in that same 8 paragraph, the 1948, 1949. 9 DR. MAKHIJANI: Yes. 10 MR. KERR: Read that sentence. 11 DR. MAKHIJANI: Yeah. (Reading) Personnel --12 '48/'49 personnel dosimetry study that Y-12 13 demonstrated that film badges provided a 14 reliable and convenient method for monitoring 15 shallow doses both in low energy photons and 16 penetrating whole-body doses from gamma rays. 17 So what -- what was the 1982 paper about? 18 MR. KERR: It was because the data had never 19 been looked and detailed before. It was just 20 thought or perceived that it wasn't very 21 reliable and -- because of frequently changed 22 procedures and -- and a general lack of 23 monitoring quality control and it was a 24 perception in that data up until this study. 25 MR. TANKERSLEY: Let me -- Let me add this,

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too, please. Keep in mind that was 1982, more than 20 years ago. That may very well -- I mean I don't have the paper in front of me -that may very well have been before we even had the original data. I assure you that we -- we did get those original data. I held the cards in my hand. I looked at them again a week ago or something. And we ultimately got the original data; I don't mean photocopies of it, the original double-sided cards and so forth. And I don't -- I don't think any of us now question the -- I mean obviously there are shortcomings in any -- any monitoring data but none of us questioned the credibility of those data, neither the film badge nor the -- the PIC data. I don't know exactly -- I'd have to look at that paper and -- and also think about it in light of it being a 1982 memo to -- to Dr. Fry. DR. MAKHIJANI: So I -- I must confess I'm confused because normally your practice is to use film badge data as the data of record. MR. KERR: No, in the early days the PIC data was used as a -- as the -- as the measurement of record. And that's true at both Oak Ridge National Laboratory, that's true of Hanford,

1 and that's true of Y-12. In the early days the 2 PICs were considered the -- the dose of record. 3 DR. MAKHIJANI: And what was the limit of 4 detection on the PICs? 5 MR. KERR: It depends on how -- the model you chose and -- and typically there were 200 6 7 millirem per day, 2 to 300 millirem per day. 8 DR. MAKHIJANI: Okay. But I saw entries as low 9 as five millirem. 10 MR. KERR: Well, you could read them down to 11 that if the scale on them, depending on what 12 scale you used and what sensitivity you used, 13 you could read them down to probably five. 14 wrote a paper, there's a paper on -- on the Oak 15 Ridge website where we went back and looked at the PIC data and the badge data and ORNL in the 16 17 early days and we used the PIC data to compare 18 with -- with the -- with the film badge data. 19 You can see what kind of comparisons you get 20 when you do the two. 21 MR. GRIFFON: George, can you explain to me, 22 and I understand you said the limit of 23 detection was 2 to 300 millirem per day but you 24 could read them down to five? 25 MR. KERR: Yeah. Typically the scale, on them

1 you could read some of them, say if they were 2 200 millirem per day, the scale was such you 3 could probably read down to five, ten -- five 4 or ten millirem. 5 MR. TANKERSLEY: Keep in mind that one of the reasons why the -- the PIC chambers have such a 6 7 poor reputation is because, you know, the 8 readings can be thrown off by dropping the --9 the badge, things like that. That's the reason 10 why they typically wore them in pairs. 11 remember correctly on that set of data, the 12 '48/'49 data, both of the PIC chamber readings 13 are on there. 14 MR. KERR: Right. 15 MR. TANKERSLEY: And then, you know, when they 16 have good agreement that's the reason why 17 they're still used today because they have good 18 agreement; it's generally accepted that it's a 19 reading. DR. MAKHIJANI: I -- I only saw one PIC entry 20 21 in the database. 22 MR. GRIFFON: It usually had a slash, didn't 23 it, Arjun? 24 DR. MAKHIJANI: Yes. 25 MR. GRIFFON: That's the two readings I think.

1 DR. MAKHIJANI: I -- I don't recall that. 2 Yeah. 3 MR. KERR: Yes, and -- and then also, Mark, 4 sometimes if they do not put both readings on 5 there I -- I know that sometimes it's on there; I've seen it. But they also have a field there 6 7 called TSR which is the total significant 8 reading and that I think typically means that 9 they have, you know, put the two together and 10 averaged them or whatever. I can't quite 11 remember what that looks like. 12 MR. CHEW: George, this is Mel. Just have 13 clarification for Mark, made a comment about. 14 It's not 200 millirem per day (unintelligible). 15 And yet, the chamber can read from zero to 200 16 millirem --17 MR. KERR: It's zero to 200 millirem but 18 typically they --19 MR. GRIFFON: Okay. 20 MR. KERR: -- they wore it (inaudible) each 21 day. 22 MR. GRIFFON: Well, that sounds more like it. 23 Okay. 24 MR. KERR: They -- They wore them each day. 25 MR. GRIFFON: I was confused, but the

1 terminology was throwing me off there. Okay. 2 MR. CHEW: I just wanted to make sure you --3 you got that, Mark. 4 MR. GRIFFON: Yeah. Thank you. 5 MR. CHEW: Good. You're welcome. DR. MAKHIJANI: But if they're wearing them 6 7 every day the number -- the number record will 8 indicate that. 9 MR. KERR: Well, at Oak Ridge, you know, they -10 - people wore PICs every day to work in 11 radiation zones. I, you know, wore -- they 12 were wearing them up into -- they still wear 13 them. And when I went to work at ORNL in the 14 '60s and '70s we wore -- I wore a set of pocket 15 ionizations chambers every day. 16 DR. MAKHIJANI: But I guess -- I guess the 17 question --18 MR. KERR: And those were not -- those were not 19 now part of the official records. 20 DR. MAKHIJANI: No, but then were they read and 21 recorded every day or --22 MR. KERR: Yes. Yes, because we got weekly, 23 monthly and quarterly printouts of the -- of 24 the PIC totals. And when they exceeded 500 25 millirems we pulled the workers' badges and had

them developed, if they were over 500 millirems we restricted them from going back in a radiation field for the rest of the quarter, because we limited their yearly doses to two rem.

MR. TANKERSLEY: I believe if you'll look at the -- those data, well, it could -- looking at electronic data, the cards actually have a -- a field, a block for each day. And I think one side of the card -- help me remember, George -- I think it covers two weeks at a time or --

MR. KERR: Right.

MR. TANKERSLEY: -- or something like that.

MR. KERR: Yeah.

MR. TANKERSLEY: And so you -- they add -- they do have the individual daily readings across the card and then at the end there's -- there's about six fields, film badge, open window, shielded and maybe one other. Then -- Then they have the -- the PIC chamber that's sum of the week and then (inaudible) significant reading. You'd really have to see the original, you know, card to see. Heck, no, we certainly didn't put in all of that. We put in the -- you know, the -- the added data, the

1 summary data at the right side of the card. 2 DR. MAKHIJANI: These are summed like for a 3 week or two? 4 MR. TANKERSLEY: I think -- I think a week. 5 I'd have to --6 MR. KERR: Yeah, because the -- the film badge 7 data was for a week. DR. MAKHIJANI: Yeah, that -- That puzzled me. 8 9 MR. KERR: Okay. 10 DR. WADE: We have to move on. 11 MR. GRIFFON: Yeah, let's -- Let's go. Arjun, 12 where do we stand on this issue then? 13 DR. MAKHIJANI: I don't know. I guess --14 MR. GRIFFON: Yeah. 15 DR. MAKHIJANI: -- if Hans might -- you know, 16 I'm not the internal -- external dose person 17 here and I guess it'll be up to the rest of the 18 team to figure out and tell me what to write 19 here 'cause as I said I -- I -- I just have 20 coordinated a lot of this and -- and --21 MR. GRIFFON: Right. 22 DR. MAKHIJANI: -- maybe Hans and John can tell 23 me where to go on it. MR. GRIFFON: Well, at least I mean I think we 24 25 have a better understanding, too.

1 DR. MAKHIJANI: Yes, right. 2 MR. GRIFFON: And we -- We just did receive 3 this database so it's hard to --4 DR. MAKHIJANI: Yeah. Yeah, no question I think I -- I understand the -- the -- the 5 numbers better. 6 MR. GRIFFON: Okay. 7 DR. MAKHIJANI: We'll just have to go back and 8 9 see what we can do. 10 MR. GRIFFON: Yeah. 11 DR. NETON: I'd like to --12 MR. GRIFFON: And take this discussion into 13 account for the final draft. Go ahead. I'm 14 sorry. 15 That's okay. I just want to point DR. NETON: 16 out we need to look at what kind of work was 17 going on at '48/'49 versus when there was 18 really uranium there. I mean '48 and '49 as we 19 talked about was cleanup of residual uranium in 20 the Calutron. 21 MR. KERR: No, I think they were starting to 22 already mill depleted uranium back in '48 and 23 ′49. 24 DR. NETON: Okay. Okay. 25 MR. KERR: Because one thing they did was they

1	were making shields for sources out of depleted
2	uranium.
3	DR. NETON: There is a source term available
4	for external. That's what I was trying to get
5	at.
6	MR. KERR: Uh-huh. Okay. Okay.
7	DR. NETON: I've got one more question and then
8	we can move on. George, you mentioned that it
9	the the co-worker model over-predicts
10	what we would estimate based on the '48/'49
11	data. That stands for about how much?
12	MR. KERR: Jim, I'd have to go back and look at
13	it. I I I can't
14	DR. NETON: My sense was that this was
15	MR. KERR: It's extremely conservative, let me
16	say that. How much does it over-predict doses
17	to people, back in those days, I can't give you
18	a figure off the top of my head.
19	DR. NETON: I bet this is well above the 95 th
20	percentile.
21	MR. GRIFFON: When When you did that
22	comparison, George, did you compare against
23	these PID readings in the in the database we
24	had, this '48/'49 database?
25	MR. KERR: Yes, sir. Yes, sir, I did.

1 MR. GRIFFON: So that was the basis for --2 MR. KERR: Yeah. 3 MR. GRIFFON: All right. We might have to, you 4 know -- SC&A, we might need a little more to 5 look at that and reconsider this issue. 6 MR. KERR: And you could see what would predict 7 -- back -- if you'll -- that last handout that 8 I gave out on the -- on the gamma and -- and 9 beta regression. You can go back to there's 10 five dose reconstructions at the end of that 11 report. And go back to the one where the 12 scaling factor was one and you can take those 13 doses off yourself and compare what's in that 14 report. 15 MR. GRIFFON: All right. Let's -- Because the 16 Rocky folks are on, too, let's move on to 17 number eight. I think we got a good sense of 18 what was in there so... 19 DR. MAKHIJANI: Yes, okay. I guess this is --20 this is the big item. 21 MR. GRIFFON: Yeah. 22 DR. MAKHIJANI: The -- We looked at, you know, 23 there's a -- there's a lot of stuff in the 24 evaluation report and as I said, a little bit 25 of disclaimer in the beginning, focused on

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table 45-B and didn't -- not enough on 45-A so let me say here that it seems -- it seems that there was a broad kind of sort in -- that was fairly successful in the early period of putting people into these two bins in the various departments. And -- And the comparison -- the -- the reason we focused on the 45-B is if you -- that's where the high exposed workers are supposed to be, more than 30 millirem average dose from 61 to 65 and by department. And Harry Hariminsky (ph), the statistician on our team, took a look at that data and did some correlations between the -the -- those departments that had relatively high doses from the -- that one table. Did they have what -- what they correlated were the relatively high doses from the earlier period of monitoring. And there was a correlation but it was weak. And then there was a question of who was monitored in the earlier period and was there a correlation between the percentage of monitored people in the earlier period with those who were shown to have -- those departments that had the higher doses when everybody was monitored? And --

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And the assumption underlying the analysis is when everybody was monitored the average doses are -- are somewhat representative of exposure potential because as they go up you expect the distribution to shift to the right. And that was also a pretty weak correlation and Harry concluded that the pre -- that the pre-1961 workers moni-- who were monitored didn't belong in the same distribution as the -- as those who were identified as having the highest doses in table 45-B from the '61 to '65 period when everybody was monitored. And so -- so it -- it seems that putting -- putting all of those -- the data for all of those workers into a single co-worker distribution doesn't -doesn't seem appropriate. When we looked at -at -- at the data it seemed that the supervisors -- you know, Hans had quoted, and I hope that Hans is on the line, so, Hans, a lot of the technical work is yours and correct me if I'm -- if I'm wrong. But it seemed like the -- the supervisors were -- had some idea of who was at high risk and then they were badging people according to that. And they made some good judgments and then badged nearly everybody

or the majority in those departments. And then some of the judgments were shown to be off in a later period. And that's the problem with the lack of correlation. And so while they had the intent of catching people with high exposure potential, the lack of or weak correlations indicate they didn't always succeed. And so we think that while it seems possible to make a co-worker model that would be claimant favorable with the available data, that that hasn't been demonstrated with the existing model.

MR. KERR: Well, we have because you go back and look at those five dose reconstructions we did. You -- Keep in mind that we scale these. We've got a -- a way to scale. If you are going to assign 95 percentile to workers you're going to have five workers out of 100 that have doses higher than that 95 percent you're going to assign if you're basing it on actual distributions. Okay. We scale up based on the workers monitoring between 1961 and 1965. We are less apt to miss those high exposure people than you are with a co-worker model.

DR. MAKHIJANI: Anyway, I mean that -- that --

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that was our conclusion is that -- that the way the model is put together by -- by -- by using the data from these two periods is -- is -- is not appropriate.

MR. KERR: Let Bill address that because we picked out workers that had the most monitoring data over a ten-year period and used them. the only thing we were trying to do was to get a time trend in the data. And I don't think there's any question that the time trend shows that the gamma doses got smaller over time because of one, the fact that -- that -- that the -- the rate guides were reduced and -- and the fact that more and more workers were monitored with time which meant that you were constantly bringing in some more lowly exposed workers so there is a time trend in the data. And that's the only thing we were trying to do was that group, one group that went from '56 to '65 was to look at a time trend. And then that model is fit to where you have actual monitoring data and I cannot believe that if we picked out monitored workers and you apply that without scaling that you're going under-predict for unmonitored workers.

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DR. MAKHIJANI: Well --

MR. KERR: And if they do have monitoring data we scale the doses upward.

DR. MAKHIJANI: Well, time trends are not -not so clear, at least as I saw them because both for the gamma and beta doses in the 1950s, well, for the gamma first in the early '50s the number of zeros went up from the early '50s some 10 or 20 percent to 80 or 90 percent and then it went down to 10 percent. And for the beta doses the trends -- trends were reversed. So -- But it seemed to indicate that -- that people were honestly trying to find who was at risk but there was some -- some -- some experimentation or some -- some trial and error involved in what was happening there.

MR. KERR: There's three problems with the data before 1960 -- before 1956. That is you had small monitored worker population. You had frequent exchange of the badges. And you had a lot of assigned dose. And those things really mean that -- that for a lot of -- if you're trying to go back and use the actual data for that period that you're going to see you can't fit it to a model. There's no way you can

1 develop a co-worker model from going back from 2 the actual data. I mean you get -- you get 3 some things that are ridiculous. You get 4 extremely -- the values scatter a lot. You get 5 extremely in some cases small uncertainties in the data because where you have a lot of 6 7 assigned dose to people their -- their high 8 doses are all coming in in a single band, a 9 small band. And it doesn't make sense to do it 10 that way and I -- I'm telling the way we 11 constructed that model made sure it was 12 claimant favorable. 13 DR. BEHLING: Arjun, this is Hans. I'm on the 14 line and I am not sure if this is the right 15 time to bring up an issue that I had discussed 16 with you, and that is the issue of quarterly doses prior to 1958 --17 DR. MAKHIJANI: Yes. 18 19 DR. BEHLING: -- defending full term exposure 20 monitoring and -- and I think we might want to 21 talk about that. 22 DR. MAKHIJANI: No, go ahead. I mean you 23 developed the issue. 24 DR. BEHLING: Yeah. The issue is one of the

following. Obviously prior to 1958 people

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monitored on a weekly basis meaning that if there is a quarterly dose record prior to '58 there is the potential that a person may be part of that database having had a guarterly dose when in fact he was monitored for as few as one week out of 13 or all 13 weeks. when I looked at the -- we don't have the original data but I did a spot check and I will give you an example. For the -- For the 25th week of 1958 which -- which the date after the criticality accident at Y-12 -- there is an inhouse memo that identifies the names and -- and badge numbers of all people who were monitored. And it turns out to be for that week, the 25th week of 1958 there were 378 -- that would be 378 people who were monitored that week. Yet when you go to, for instance, table 4-4 in the evaluation, in the appendix 1 of the SEC evaluation and you look at the third quarter you identify a total of 689 persons who were monitored in that quarter. And of course, there's now a -- almost a factor of two discrepancy which leads me to believe that you may have entered into the database people who were monitored in any given quarter who were

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not monitored for the full 13 weeks during which the dosimeters were being handed out and read meaning that a person with potentially as few as one weeks of exposure will be part of that database and the database the way it is currently constructed which assume in that whatever quarterly badges -- quarterly dose records are available, that that person was monitored for each and every 13 weeks. after 1961 when the cycle was extended to quarterly cycles, that does not affect when you deal with monthly, and worse yet with weekly, just because you have a record for an individual does not necessarily mean that that individual was monitored for the full duration of that particular quarter. And so what I'm saying is that just based on that one single spot check -- check involving the 25th week of 1958 where you only had 378 people monitored, that is almost a factor of two lower than the total number of people monitored for the counted quarter, the third counted quarter of 1958.

MR. KERR: Okay. Those ones that you picked out of the table, those were coming out of this

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TIB-47 that we talked about earlier. were not -- those were just estimates that said how many people were being monitored during that period. And it just took the number of records that were turned in and divided by 13 weeks per quarter to get an estimate. that's -- that's clearly explained in that report. So you shouldn't be comparing that with the other more -- what do I want to say -fundamental thing of going in and identifying workers. But you've got to consider the way that the quarterly doses were -- were obtained. And the quarterly doses, and I -- I -- I hate to quote on this right now but I have a couple memos here of how quarterly doses were done. And they took each of the individual positive records they had and summed them up for that individual. And then they tried to correct that quarterly total for missed dose. And the way they did that was they took the number of film badges each and divide that by the number of positive records. So if that person had, say, was issued 13 film badges for the whole quarter and then they come back in and said okay, he was -- had 10 positive records, we'll

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up his dose by the ratio of 13 over 10. you know, this is one of the reasons why I think we sometimes had trouble going back into the database. And the reason for coming up with quarterly doses and yearly doses was I forget what year it was, you know, they started saying, well, you got to have -- you got to keep the dose under a certain limit depending on age. And when they did those quarterly doses they did in fact try to account for any missing dose or quarters or weeks in which they did not have a record for that -- that worker. And he could have been on vacation. He could have been off sick. He could have been transferred to another job temporarily or something. But I'm saying that they've -they've tried to adjust those for missing dose. MR. TANKERSLEY: This is Bill Tankersley and that procedure is well documented. written by C. M. West if I'm not mistaken and I know that document is on the O-drive.

MR. KERR: Yeah, it sure is.

DR. BEHLING: Could you make that available because as I said, right now I have not had any reason to come to that conclusion that for

1 instance a person who is part of that quarterly 2 record --3 MR. KERR: As a matter of fact that may be in 4 that gamma report. I'd have to look and see. 5 DR. BEHLING: Okay. Could you -- Could you identify that document? 6 7 MR. KERR: And -- And I'm sure like Bill says, 8 I'm almost positive that that -- that is on the 9 O-drive. 10 MR. TANKERSLEY: Yes, I know it is. It's been 11 sent up there, you know, months or years ago. 12 MR. KERR: When this question came up before. 13 DR. WADE: Well, can you let Hans know where it 14 is then and --15 MR. KERR: Yeah, okay. I'll get the record 16 number. 17 MR. GRIFFON: Okay. The next thing that I had, a 18 DR. BEHLING: 19 person who could have been monitored for 20 (inaudible) that he would be part of that 21 database in -- in that -- that would be 22 necessary to adjust. That -- That's the 23 central question that I have. 24 MR. KERR: Well, it would have been adjusted; 25 if he -- if he was missing some weeks it

probably would have been adjusted upward to try to account for any missing dose that he might have had due to a damaged film badge, due to a zero reading, due to the fact it wasn't turned in, it was lost. And those when they didn't have the full 13 weeks there was an adjustment made.

DR. MAKHIJANI: Or -- Or people were taken off
monitoring, too. I mean --

MR. KERR: Well, that's true, too.

DR. MAKHIJANI: -- of the examination that you did of, I don't know, 15/20 workers or 30 workers I guess, there were examples of seven workers who were found to have low doses and then were taken off monitoring. So those -- I don't know if they are partial quarters or full quarters but -- but there certainly seem to be people who went on monitoring and off monitoring.

MR. KERR: They took the transferred workers from one to the other on a -- on a quarterly basis or semi-yearly basis or yearly basis.

They didn't -- They didn't take people off just in the middle of the year unless they, you know, were terminated, the people quit or

looked at like every quarter.

whatever.

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DR. BEHLING: I guess to -- to finalize this issue and get on with other issues, but I do still have a problem in trying to reconcile the number of 689 that is in table 4-4 as defined as I guess in -- defined as the 378 people who were in fact identified by name and -- and badge number who were monitored in the 25th week of -- of 1958. To me I can certainly understand a minor discrepancy where maybe ten people, maybe somebody left -- left employment, etcetera, would come in or leave the -- the -the -- the -- the database and -- and essentially not be part of the full number for that count a quarter. But I can't see a factor of two being -- being something that you can reconcile with the explanation such as retirement or -- or --

Typically those -- those rolls were

MR. KERR: Well, I'm just saying that -- that
- that those others were just a very crude

estimate by dividing the number of records by

assuming 13 and saying, well, that's how many
- that's possibly how many that -- that's the

minimum number of people who were -- who were

monitored. And I even think you -- you -- it is possible that the values that -- and this is quoting from your report on page 15 -- it is possible that the values in ORAU O-TIB-47 are incorrect because they were deduced from the number of records assuming there would be about one record per worker per week. And that's essentially how those values were determined.

DR. MAKHIJANI: Yeah, but --

MR. KERR: Where the other went in and looked at -- at the number of workers that were involved in detail.

DR. MAKHIJANI: Yes, but --

MR. KERR: If there's a factor two difference, so be it. I, you know, that's just -- that's just the way the two tables were differently constructed.

DR. MAKHIJANI: But -- But George, the example that Hans is giving has a factor of two difference in the other direction. He had the example from the number of workers who were monitored in that week being a factor of two less than the ones that were calculated by dividing by 13. And what you're arguing is that the -- the number of 600-and-odd should be

1 a minimum number so --2 MR. KERR: Well, that could be -- that could --DR. MAKHIJANI: -- should be larger so --3 4 MR. KERR: That could be someplace --5 DR. MAKHIJANI: -- direction. 6 MR. KERR: You know, that could be a place 7 where they adjusted a number of workers. I 8 don't know. You know, we just had to go back 9 and look at it. I have no idea why there's 10 that difference. 11 DR. MAKHIJANI: So if it's actually --12 MR. KERR: It's just there. DR. MAKHIJANI: The direction that's the 13 14 troubling part. 15 MR. KERR: It's just there and that may be a 16 place where they did adjust workers back in the 17 early days by, you know, in -- in some interim 18 period. 19 DR. BEHLING: I guess I don't know what 20 footnote 12 in table 4-4 says. Footnote 12 21 which represents the N value and the footnote 22 says N therefore is the total number of 23 quarterly doses which to me suggests that you 24 monitored a total of 689 people in that 25 calendar quarter.

1 MR. KERR: All I can say is those tables were 2 constructed differently and I don't know 3 whether that reflects the way the tables were constructed or reflects a difference in the 4 5 data that -- that is -- was used to make them. The only way we could tell what -- what's 6 7 happened there is to go back and look. 8 MR. TANKERSLEY: This is Bill. I'm a little 9 bit confused here but I heard Hans say a moment 10 ago if those were the number of quarterly doses 11 would equal the number of people. 12 wouldn't be true typically and then we verified 13 this a number of times against the health 14 physics report; it would be one-fourth of the 15 number of people. 16 DR. BEHLING: I don't understand that 17 relationship. 18 MR. TANKERSLEY: Well, because they're 19 monitored -- the -- the results are recorded 20 per quarter. 21 DR. BEHLING: If you had -- Let's assume that the number of people that they monitored in the 22 25th week of 1958 were in fact a stable 23 24 population of people. They were monitored 13 25 weeks each. You would expect in table 4-4 for

quarter number (unintelligible) to have 378 as the value of N and that's what I'm contesting or questioning.

MR. TANKERSLEY: Again, I -- I don't quite follow you there but you'd expect to have about four times that number of records, one -- of one for each quarter for each person.

DR. BEHLING: No, no, these are quarterly dose values that I'm citing to you in table 4-4 in appendix 1. I'm referring to page 25, bottom of page 25. It has 1968, 2-3, 3rd quarter, and the number of records, quarterly records are 689. And yet when I as a single spot check checked the number of people badged for the 25th week there were only 378 which is approximately a factor of two lower. And as I said, I cannot reconcile that big difference realizing that perhaps maybe certain people came into the system or left the system so that the number of 378 would be potentially perhaps greater by a factor of 10 people or 20 people but not by a factor of two.

MR. KERR: The only thing I can say is we'll just have to look at the tables and see why there's a discrepancy between them. I don't

1 really know. 2 MR. GRIFFON: I would -- I would also suggest 3 that, you know, maybe prior to the Board 4 meeting, Jim, you know, you -- maybe you should 5 review this -- the statistical approach offered by SC&A and, you know, if you have a rebuttal 6 7 to that or -- or, you know, because I think we 8 still have a difference of opinion. And of the 9 last question, I think --10 DR. NETON: Well, I think Mark, we can do that 11 but --12 MR. GRIFFON: Yeah. 13 DR. NETON: -- this has been on the table for 14 two months and we just got a 20-page report for 15 statistical analysis yesterday. It's going to be hard to do that. 16 17 DR. MAURO: Jim, this is John Mauro. 18 would ask is there are two -- there are figures 19 1, 2 and 3 in -- in the appendix to this 20 report. 21 MR. GRIFFON: Right. 22 DR. MAURO: This statistical workup, there's 23 three figures. One of the figures, figure 3, 24 based on the analysis, actually supports your 25 position that there was a concerted effort to

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monitor more people who were in the departments that had the greater potential for exposure which argues that it wasn't a pure cohort sampling situation. It was a concerted effort to monitor more of the people in those departments that were expected to have the highest exposures so -- so figure 3 in this attachment provides some evidence, speaks for itself, that -- that -- there was that tendency going on. What -- However, figures 1 and 2 provide information that -- that says that there is -- it's very hard for you to say something about a given department. That is, a department that may have experienced high exposures post-1961 may not have experienced high exposures pre-1961. There was almost no relationship between the two. And -- And that figure, figure 1 and figure 2 is troubling to It's almost as if they were -- the relationship between post- and pre-exposures do not follow any predictable patterns by department or within department. To try to bring this to closure, if you wouldn't mind, just take a look at that figure 1 and figure 2 on page 30 of our report and maybe we could

talk a little bit about that. And -- And it would be fine with me that we could even talk about it, you know, tomorrow or -- or Monday because it does tell us a story that -- that raises questions whether the extrapolation approach that you folks have adopted can really work. I think if those questions could be answered maybe we can put -- put this thing to bed.

MS. MUNN: Hans, I have one question. Did you run a similar spot check on any other week?

Did you do only that one week?

DR. BEHLING: Well, that's the only data I could find. I guess it would like be nice if we could look at multiple time frames but it turns out that apparently in the aftermath of the Y-12 criticality accident I guess there was some concern about who did we monitor and what are their exposures and how close did they come to meeting regulatory or admin limits, etcetera, etcetera. So it turned out that that was just perhaps useful interoffice memos that allowed me to look at that but if there's any other data out there, Wanda, I don't have it. And so it was just a -- just a snapshot in

1	time, allowed me to look at those individual
2	numbers and then compare it to table 4-4 in the
3	appendix 1 of the SEC evaluation report.
4	MS. MUNN: Right. I just was trying to make
5	the point for myself that a single instance
6	where we have these puzzling numbers doesn't
7	necessarily cause me to jump to the conclusion
8	that virtually all of the numbers might suffer
9	from that same defect.
10	DR. BEHLING: No, well
11	MR. GRIFFON: That's interesting, too, Wanda,
12	because let's remember the reverse.
13	MS. MUNN: Yeah, exactly. Exactly.
14	MR. GRIFFON: You know, so
15	MS. MUNN: And it's but but I
16	MR. GRIFFON: Yeah.
17	MS. MUNN: I'm trying to identify
18	MR. GRIFFON: I agree.
19	MS. MUNN: whether that was the only week
20	that anyone even looked at.
21	MR. GRIFFON: Yeah. I think everybody's
22	limited on the amount of raw records we can
23	find to
24	MS. MUNN: I understand.
25	MR. GRIFFON: do comparisons, yeah.

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DR. NETON: I guess -- I guess I want to get back to the original point that we had reached I thought several months ago. See, I've looked at these graphs and I have not had time to digest this 20-page analysis, I'll be honest with you, because it came in at noon yesterday. But the point is I think if -- if it's true, what you're saying is true, that there is -that the highest workers were not monitored, then we have a sampling of the workers. why is that an SEC issue at that point if -- if then it's a matter of picking the appropriate metric to -- to use for reconstructing unmonitored workers, that is, the 95th percentile or the 50th percentile. What is the What is the -- Am I missing the issue here?

DR. MAKHIJANI: If -- If you look at figure 2 in which the percentage of monitored workers in the '56 to '60 period is correlated against the dose -- average doses in the '61 to '65 when there was universal monitoring, the correlation is -- is very weak. And so what -- what that says is that actually some of the departments that were at high risk were monitored at high

1 percentage times and some of them were 2 monitored a low percentage of the time. And so 3 actually what were the actual -- to establish 4 that you know the actual exposure conditions in 5 the high risk departments in -- in the -- in the '56 to '60 period seems -- at -- at this 6 7 stage that job hasn't been done. DR. NETON: Well, my point, Arjun, is if we 8 assign the 95th percentile of all the monitored 9 10 workers -- you know, we're not -- you know, the 11 only way this would not work I don't think is 12 if they preferentially monitored people who 13 weren't exposed. 14 DR. MAKHIJANI: Well, I think that that's 15 clearly not true. DR. NETON: Well, then, okay. If that --16 17 Given that's the case then I don't know why a 95th percentile co-worker model would not work. 18 19 DR. MAKHIJANI: Is that the -- Is that the one 20 we have? 21 DR. NETON: No. We -- We're -- The argument 22 or the discussion that we've been having is 23 were the highest exposed workers monitored; and 24 our position was if they were then we can assign the 50th percentile to the unmonitored 25

1 workers. 2 DR. MAKHIJANI: Okay. 3 DR. NETON: That's the issue. And you -- you 4 were arguing, and I need to look at your 5 analysis, that that may not be true. 6 DR. MAKHIJANI: Right. 7 DR. NETON: So now we have a sampling of the work force. And given that as a sampling then 8 I would agree if that's true that the 50th 9 10 percentile might not be appropriate and something like the 95th percentile might be --11 12 might be a better estimate. But why that would 13 be an invalid model then I'm not sure. 14 DR. MAKHIJANI: No, we haven't said that. 15 DR. NETON: Right. 16 DR. MAKHIJANI: In fact -- In fact, what --17 what is in the report, it -- it makes no 18 judgment about whether this is an SEC issue or 19 not. 20 That's what I'm trying to --DR. NETON: 21 DR. MAKHIJANI: It makes no judgment about --22 DR. NETON: Yeah. 23 DR. MAKHIJANI: Well, Jim, you're -- the amount 24 on the table is what it represents. 25 DR. NETON: That's what I'm trying to get at,

Arjun, is we have a very limited amount of time here to deal with issues --

DR. MAKHIJANI : Yeah.

DR. NETON: And -- And if this is not an SEC issue then I would prefer not to spend my entire weekend analyzing it.

DR. MAKHIJANI: This is -- This is -- I guess I -- I will defer to Hans on this. As I said, this is -- I'm -- you know, this is a piece I'm coordinating. Ron and Hans have looked at this. It -- It's your judgment call, Hans, not mine.

DR. BEHLING: Yeah, I would say, and I will agree with Jim, it's possibly not an SEC issue. In fact, I was just reading the recent draft for co-workers at Rocky Flats and where you give the option of using a 95th percentile value for unmonitored workers to -- who should have been monitored, and that to me is a very nice and claimant favorable approach that is clearly claimant favorable for the Rocky Flats dose reconstruction projects. There the co-worker data is divided into 50th percentile value for people who are possibly only exposed part of their work period as opposed to the

95th percent value for people who were 1 2 routinely or should have been routinely 3 monitored. And I would concur if we were to default to a 95th percent value that would 4 5 settle most of the questions and concerns. MR. GRIFFON: Hans? Hans, just can I offer 6 7 maybe what SC&A needs to do in -- in -- in 8 finalizing this report or a final draft of it 9 is -- is to make that sort of statement or 10 something, you know, if you're comfortable with 11 it, of course -- make that sort of statement 12 within the body of the report. And then, you know, then it's out there that, you know, you 13 14 feel that based on your analysis a 95th 15 percentile model may be more appropriate 16 because X, Y and Z as you presented but that it 17 -- it would preclude -- it wouldn't necessarily 18 be an SEC issues. 19 MR. KERR: And I -- And I would like to really 20 see the -- a solid basis for the --21 MR. GRIFFON: Right. 22 MR. KERR: -- for the argument. 23 MR. GRIFFON: Right. 24 DR. NETON: Yeah, George. I don't think 25 anybody's arguing that, you know, we would

1	adopt it if they so explained.
2	MR. KERR: No, I understand that.
3	MR. GRIFFON: Right, right, right.
4	DR. NETON: But, you know, I I'm just
5	trying to move things along, you know.
6	MR. GRIFFON: I agree, Jim. I was going to say
7	the same thing before you went into that is
8	DR. NETON: Sorry I pre-empted you.
9	MR. GRIFFON: And we've We've We've
10	said this before actually that this has been or
11	the borderline of SEC site profile for awhile
12	so I think maybe you can make a statement to
13	that effect in your report, SC&A.
14	MR. BUCHANAN: Yes, this is Ron Buchanan and I
15	think that it's been our position is that this
16	would not be an SEC issue if you modified the -
17	- the final. It isn't so much the missing data
18	as how it's being used.
19	DR. WADE: Okay. We need to move on. We
20	really do.
21	MR. GRIFFON: Yeah, yeah, yeah. I'm
22	saying I think the next three we can wrap up
23	fairly quickly actually but maybe I'm wrong.
24	Let's go on to number 9.
25	DR. NETON : Yeah. Can I just get a little

1 clarification that, you know, for number 8 SC&A 2 may -- may modify their -- their -- their 3 documents so that we don't have to provide 4 these analyses at this point or is that -- I 5 mean I want to make clear what we're going to 6 provide. I mean we -- we're certainly going 7 to -- we're certainly going to become familiar, 8 you know, with the entire --9 MR. GRIFFON: It sounds to me -- I mean Hans 10 and Ron weighed in there for SC&A. It sounds 11 to me like that's right, Jim. 12 **DR. NETON:** Okay. 13 MR. GRIFFON: That you don't need any more 14 analyses I mean --15 Well, we will eventually but --DR. NETON: 16 MR. GRIFFON: Although, yeah. For site profile 17 concerns. DR. NETON: Yeah, okay. Very good. All right. 18 19 Number 9 gets into the polonium 208 issue and 20 actually 9 and number 11 are somewhat related 21 because they're both Cyclotron issues. 22 MR. GRIFFON: That's, yeah. 23 DR. NETON: And so I'll try to cover it 24 somewhat in the same way. I think there's a little bit of confusion as to what we meant to

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do with the examples for the Cyclotron, that is the gallium and the polonium. Given our position, and I think SC&A understood that pretty clearly in their review, that -- that for the Cyclotron these are -- these tend to be episodic exposures over a period of time that were -- were followed up and tracked to ground and monitored, and we have a lot of indications we believe from the documents that we have in hand that that's true. I'd emphasize that by doing a gallium intake assessment for -admittedly the only one we could get our hands on quickly to get the analysis done admittedly is outside the 1957 period by three years, but it spoke to the issue of -- of not only were these things tracked to ground and -- and they do follow-ups on -- on incidents when there were target ruptures but also the -- the -- the relative magnitude of the deltas involved with these so-called exotic radionuclides that have very typically fairly short half-lives in the body and are fission products that -- not alpha emitters. They're more beta gamma emitters. That was the intent of those examples that we provided. We -- We believe and we -- we still

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have not provided to you but we believe we have sort of a five-prong approach (unintelligible) constructing these incidents. Through the HP reports that we have -- and there are some gaps in those reports because a few of them are still classified. Our folks have looked through them and they believe that they support our case that there is Cyclotron information in there that we can use to support these dose reconstructions. There are interoffice correspondences that we -- we have available, division reports and individual claimant files. We've looked through a number of individual claimant files looking at the CATIs that were done and out of the entire population right now we can only identify 11 or so individuals who indicate that they were involved in -- in Calutron/Cyclotron operations and -- and had -maybe had some reference to incident. working through those files now to identify the bioassay data, etcetera. But I want to point out that this is not a huge population of workers. This is a Cyclotron operation that -that is involved. Some technical people, some maintenance folks and those types, but our

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estimation is that -- that the affected population is somewhere in the vicinity of maybe 40 individuals because this was a unique isolated operation. Now, the Cyclotron targets were for the most part cladded. That is, you know, they were contained in cladding, exposed, pulled out and as Mel Chew nicely described it, had pictures, when the radiation was done and those targets were processed over at ORNL. the few cases there were ruptures though, again we feel that we can track these bioassay follow-ups and incident reports through either the DOE submittals for the claimants or in the investigation reports that we talked about in delta view. The polonium period is slightly more problematic in the sense that in 1951 and '52 polonium exposures were -- were non-clad. They couldn't get enough energy into these targets with the cladding in place so they were essentially bare targets that did dispense -disperse some fairly significant levels of airborne alpha activity, although if you look in the 1951 and '52 health physics reports there are indications where there are air sample results. I think there's probably about

100 individual air sample results indicating they recognized the problem, they were controlling for it, they restricted access, all those sort of things. So I think between the incident reports, some of the air monitoring data we have and the nature that these were episodic, you know, discrete events, we -- we feel fairly confident that we can go back and reconstruct exposures to these workers.

DR. MAKHIJANI: This is -- This is Arjun. The
-- I -- I actually want to separate the
polonium from the -- from the gallium example
because even though they're in the same area
because --

DR. NETON: Right.

DR. MAKHIJANI: -- we found different issues with them. I think Jim -- Jim covered some of them. There is -- There -- There is a set of samples from 1953 that does appear to relate to an incident for polonium in 1953 and those seem -- I think most -- almost all but two of the samples relate to that incident best I could tell. I don't have a description of the incident, just from the dates or how the sampling was done.

1 DR. NETON: Right. 2 DR. MAKHIJANI: Is that right, Jim? 3 DR. NETON: Yeah. 4 DR. MAKHIJANI: Okay. The -- The -- But it -5 It seems to me that we don't know the years 6 of production of polonium well because --7 UNIDENTIFIED: Yes -- Yes, we do. 8 DR. NETON: Yeah, I think we do. DR. MAKHIJANI: Okay. Because I found -- I 9 10 found that the appendix 2 compilation was --11 was not -- didn't have anything for '51 and '53 12 even though there was an accident in '53. And 13 so what -- what I -- what I mean to say is that 14 I didn't -- I didn't see that the compilation 15 was complete and so I don't know whether you 16 have a complete set of data about that. 17 DR. NETON: We -- We actually have, Arjun --18 I'm sorry I -- I usurped your introduction 19 there. 20 DR. MAKHIJANI: Oh, no. No problem. 21 DR. NETON: You saw my zeal to get --22 DR. MAKHIJANI: No, that's -- shortness of 23 time. 24 DR. NETON: We have a production of polonium 25 208 report from Oak Ridge National Laboratory.

1 It was the final report on termination of 2 project, ORAU -- ORNL 1392, that goes in -- in 3 -- in a lot of detail as to how much production 4 there was by month --5 DR. MAKHIJANI: Okay. DR. NETON: -- from the initiation of the 6 7 polonium runs in 1951 through closure in August 8 1952. 9 DR. MAKHIJANI: Okay. If it's unclassified it 10 would be useful to see it. 11 DR. NETON: Yeah, we can put that on -- on the 12 O-drive for you. MR. RUTHERFORD: Sorry. I'm sorry. This is 13 14 LaVon Rutherford. In fact that is already on 15 the O-drive under Cyclotron and Calutron --16 MR. GRIFFON: Oh, it is? 17 MR. RUTHERFORD: -- of the A-B (inaudible). It's already there. 18 19 MR. GRIFFON: Okay. 20 DR. NETON: So we do know production and again, 21 we have some of these air sample data. The 22 1953 data we -- we analyzed show that 23 (inaudible) reconstruction for polonium 24 (inaudible). 25 DR. WADE: Jim, you're cutting in and out.

1 MR. GRIFFON: Yeah, yeah. 2 DR. NETON: We could do dose reconstructions 3 for -- for polonium 208. There was some 4 concern about that given bioassay data. And --5 And we've used to -- to demonstrate proof of 6 principle that we can actually do that if in 7 these incident reports we run across a polonium 8 208. 9 MR. CHEW: Jim, this is Mel. Yeah, we -- We 10 also were aware of there was an incident with 11 polonium 210 from a polonium drilling neutron 12 source that was -- was different from the 13 polonium 208 and that could be the bioassay 14 result because they just mentioned it was 15 polonium. DR. MAKHIJANI: Oh, I see. Yes, that's right. 16 17 That was a question, too, because he had three 18 different isotopes of polonium going on --19 DR. NETON: Yeah. 20 DR. MAKHIJANI: -- binary. And the -- And the 21 data actually only mentioned the element of the 22 isotope. 23 DR. NETON: That's correct. 24 DR. MAKHIJANI: Okay. Okay. 25 DR. NETON: So anyway --

1 DR. MAKHIJANI: Yeah. 2 DR. NETON: I guess that's about all I can say 3 on our position right now. We -- We wish we 4 had all these investigation reports out there 5 for you to look at but we just don't. 6 DR. MAKHIJANI: Yeah. No, I mean, Jim, I -- I 7 just wrote up what I saw. That's all. 8 DR. NETON: Sure. 9 MR. GRIFFON: Jim, I was going to -- just going 10 to ask. You mentioned this five-prong 11 approach. 12 DR. NETON: Uh-huh. 13 MR. GRIFFON: And I guess in the spirit of --14 of sort of proof of principle the -- the better 15 you can lay that out the --16 DR. NETON: Yeah. 17 MR. GRIFFON: -- you know, before the Board the 18 better, you know, it will be in the situation 19 that --20 DR. NETON: I understand. 21 MR. GRIFFON: Yeah. 22 DR. NETON: It's just --23 MR. GRIFFON: I know. 24 DR. NETON: It's all coming out in time. 25 MR. GRIFFON: In your situation, too, I know.

1 DR. NETON: Yeah, because I'm not --2 MR. GRIFFON: We've been here before. 3 DR. NETON: I'm not making apologies. 4 MR. GRIFFON: Yeah. 5 I'm just trying to be realistic. DR. NETON: DR. WADE: So what do we have left now in terms 6 7 of -- of --8 MR. GRIFFON: Wait. Maybe we should just pick 9 up on the gallium there. Arjun, were you --10 DR. MAKHIJANI: There's a plutonium and a 11 gallium, Mark. And I think I haven't examined 12 the plutonium dose reconstruction, nor I think has anybody else on our team because it does 13 14 seem put up pretty recently. And -- But the 15 plutonium data as we say here is more copious 16 and it is from the period and there's --17 there's one year that seems to possibly be missing but it could possibly be filled in by -18 19 - by co-worker data. It doesn't seem to have 20 the same kind of issues as we picked up from 21 polonium. Does the gallium --MR. GRIFFON: Is it obvious -- let me stop on 22 23 the plutonium. Is it obvious who would --24 would -- would be exposed to plutonium in those 25 years?

1 DR. MAKHIJANI: Well, NIOSH has said based on 2 limited information that there were only 3 limited production parts there for a limited 4 time that were solid and did not pose a 5 potential for internal exposure. And so we've just re-quoted that and cannot make a judgment 6 7 about it so for -- for the moment that's where 8 it stands. And haven't come across any 9 evidence to the contrary to NIOSH's position 10 certainly. 11 MR. GRIFFON: There are a large number of bioassay samples from '52 to '56 it says. 12 13 were they doing bioassay if there was no 14 potential threat? 15 No, there were -- I think that DR. NETON: 16 these, and Mel Chew can correct me if I'm 17 wrong, but this was the plutonium separations 18 in the Calutrons. 19 MR. CHEW: Right. That's correct. Uh-huh. 20 DR. NETON: All right. And -- And so, you 21 know, it's clear in 1951 that they were thinking about it. It's even mentioned in the 22 23 health physics reports they mention that we 24 need to think about getting ready for 1952 25 production of plutonium. And so there was a

1 fair amount of separation going on in those 2 years and -- and that's why we have these 3 bioassay samples. I think I would just like to 4 comment on one of SC&A's comments that, you know, we don't have a co-worker model. 5 6 example that we provided went through and --7 and -- and as a bounding analysis we proposed 8 to use, and we identified the 95th percentile 9 of all of the monitoring data we have. And as 10 -- as a bounding analysis we would propose to 11 use that in a -- as a -- as an intake, chronic 12 intake scenario for plutonium. So we think we 13 -- we have a handle on the upper limit of 14 exposures based on the I think there are 600 or 700 plutonium samples in the -- in this period. 15 16 DR. MAKHIJANI: That's correct. 17 DR. NETON: Which is not inconsistent necessarily with the number of workers that may 18 19 have been working at the operation. 20 DR. MAKHIJANI: No, no. That's correct. 21 I agree there -- there -- there are 22 that number. Joyce, are you still on the line? 23 MR. GRIFFON: Back to my question. 24 DR. LIPSZTEIN: Yes, I'm still on the line. 25 DR. MAKHIJANI: Can we -- Can -- Will you

1 have the time to look at that? 2 DR. LIPSZTEIN: Yes. 3 MR. GRIFFON: Let me -- Yeah, let me ask this, 4 too, Jim. Back to my question on how do you 5 know who was working in the -- in this area? 6 Is it obvious by department or --7 DR. NETON: Well, this would be --8 MR. CHEW: Mark, let me try to answer that 9 question. The primary work during that 10 particular periods was using the Calutron to 11 separate some of the plutonium isotopes for the 12 research to look at cross-section work for the 13 different isotopes of plutonium. That's why 14 the pockets were there. And so I would say 15 it'll limit it to the people who were basically 16 either cleaning out the -- the Cyclotron 17 pockets and potentially the (unintelligible) 18 and recovering the specific isotopes that were 19 being separated at the Calutrons for the 20 plutonium here. So I think -- I think the --21 the class -- I mean the number of people and 22 the category of people can really be well 23 defined. 24 MR. GRIFFON: Yeah. No, that -- that all makes

sense to me, Mel. The question I'm asking is

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1 retrospectively, you know, do these people fall 2 out from department number from -- from their 3 own questionnaire? Do they self-identify that 4 they were working in Calutrons in that time 5 period? Do they, you know -- how do you -- how 6 do you place people in -- in time in that area? 7 MR. TANKERSLEY: Mark, again, I think this is 8 one of the best places where use of the work 9 history database can identify those people 10 really pretty accurately. 11 DR. NETON: Yeah, I think --12 MR. GRIFFON: Because there was a small number 13 and they were well controlled, right, or 14 whatever. 15 Right. And these department DR. NETON: 16 numbers are fairly small. 17 MR. TANKERSLEY: Job titles, departments, job 18 codes and year, you know. You can track the 19 people, you know, by every job they had, every 20 department they had. 21 MR. GRIFFON: Okay. Okay. 22 MR. TANKERSLEY: That would be pretty -- pretty 23 straightforward. DR. NETON: But we would certainly start with 24 25 the CATI and if there was any indication in the

1 CATI report that they worked with this material 2 it would -- it would certainly get us going 3 down that path. 4 DR. MAKHIJANI: And what happens with the 5 survivors? Well, that's another issue. 6 DR. NETON: 7 percent of our cases are survivors. 8 remember that these plutonium values were, we 9 believe, and this is what ORAU or Y-12 folks 10 have told us, is that these samples, if they 11 were taken there should -- should be showing up 12 in their urine samples because remember, they 13 go through the delta view database and look for 14 people who have those samples and provide them 15 with the records. So anyone who would monitor 16 for plutonium were -- we believe that these are 17 going to come across and that's what we've been 18 told in -- in our -- in the DOE submittals. 19 MR. GIBSON: This is Mike. Jim, have these --20 some of these individual cases involving the 21 Cyclotron and the Calutron, are they pended or 22 have they started to have reconstruction done 23 and -- and been adjudicated? 24 DR. NETON: Well, that's a good question, Mike. 25 We have not universally pended

1 Calutron/Cyclotron operators but I will say 2 that using the efficiency process, there's a 3 number of methods in case those could go out, 4 you know, ones that certainly would qualify, 5 you know, over 50 percent. And I don't think that -- I'm not -- I'd have to go back and 6 7 check to see where -- where any of it may have 8 been Calutron operators went out, if they were 9 -- it seemed to be less than 50 percent. I 10 don't know that any have. 11 MR. GIBSON: Okay. That would be interesting 12 to find out. 13 MR. GRIFFON: Good question, yeah. 14 DR. NETON: It's a good question. I think among the -- the cases that we've done we -- we 15 16 can take a look at that and provide some 17 information. 18 MR. GIBSON: Okay. 19 MR. GRIFFON: Okay. I think we can look at 10 20 -- or 11 just for a second, Arjun. 21 DR. MAKHIJANI: Yeah. 22 MR. GRIFFON: Then I'll try to probably take a 23 break and go to Rocky. 24 DR. MAKHIJANI: Okay. Well, 11 is -- is -- is 25 simpler. There, you know, the -- the gallium

1 internal dose was considered. I guess you were 2 only considering gallium and not trying to 3 illustrate all radionuclides to which this 4 person was exposed. 5 DR. NETON: That's correct. We were just 6 trying to show, you know, we can do these dose reconstructions using ICRP model given that the 7 8 incidents will track to bed. 9 DR. MAKHIJANI: Okay. And -- And so --10 the -- The big question is what -- how to 11 establish the relevance of a 1968 incident 12 through what went on in the SEC period, and that, there's no discussion of that. And how 13 14 do you -- how do you bound the doses or show 15 their maximum plausible for the period in 16 question? 17 DR. NETON: Right. And -- And again, we 18 believe, you know, this five-prong approach 19 that I mentioned --20 DR. MAKHIJANI: Right. 21 DR. NETON: -- that we just have not found one 22 in the SEC period yet that -- that we can -- we 23 can show you. 24 DR. MAKHIJANI: Okay. 25 DR. NETON: But the data that we have in hand

1 leads us to believe that these are -- are what 2 was John Mauro's --3 MR. GRIFFON: Jim -- Jim, is there any way 4 short of -- I was just wondering if there's any 5 sort of interim product to provide with regard to these incidents like if you had a printout 6 7 of -- of what came up on your search. I don't 8 know if that's --9 DR. NETON: Yeah. 10 MR. GRIFFON: -- any faster, that would give us 11 an indication of how much insufficient data you 12 had, how, you know, and what radionuclides were 13 covered or something. 14 DR. NETON: Yeah. 15 MR. GRIFFON: It may not be that easy but I 16 don't know. 17 DR. NETON: Yeah, I can assure you, Mark, we're 18 working towards that end --19 MR. GRIFFON: Okay. Okay. 20 DR. NETON: -- as fast as we can and --21 MR. GRIFFON: Yeah. 22 DR. NETON: -- you know, we're not -- we're not 23 sitting on our hands here but I -- it's a good 24 comment and I think if we can make this picture 25 clearer for the Board and working group we're

1 going to try. 2 MR. GRIFFON: I -- I know that, Jim. 3 DR. NETON: Yeah. 4 MR. GRIFFON: I know you're not sitting on your 5 hands. 6 DR. NETON: I know. 7 MR. GRIFFON: Okay. 8 MS. MUNN: I don't think any of us thinks 9 you're sitting on your hands. 10 DR. NETON: What I meant to say though is this 11 is an issue that, you know, as of this morning 12 we were conferencing and working to try to --13 to see, you know, the maximum amount of -- of 14 light we can shed on this the better. We know 15 that. 16 MR. GRIFFON: Okay. Okay. The -- The only 17 other question I have on the gallium was this 18 example -- I haven't even looked at the example 19 but the -- only discusses internal dose; is 20 And -that true? 21 DR. NETON: Correct. 22 MR. GRIFFON: And are there any reasons to 23 believe that you'd need any sort of other 24 method for estimating external dose in the

Cyclotron or would they all be badged and --

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1 DR. NETON: We -- We don't think so. 2 very clear that Cyclotron workers were badged. 3 We've got some -- some control procedures that 4 speak to that, you know, this -- of any place 5 at Y-12 --MR. GRIFFON: 6 Yeah. 7 DR. NETON: -- this would have been the highest 8 potential exposure. 9 MR. GRIFFON: Okay. 10 DR. NETON: In fact, we had toyed with the idea 11 of using the badge results to impute the internal doses but it didn't work out as you 12 13 can imagine. 14 Okay. All right. Because I --MR. GRIFFON: 15 Anything else on that, Arjun? I guess we --16 DR. MAKHIJANI: No. No, I think that's it. 17 MR. GRIFFON: Okay. All right. I think --18 mean what -- what -- I think SC&A has some --19 some, you know -- you're going to provide us with a final draft on this so I guess right 20 21 before the meeting. 22 DR. MAKHIJANI: Yeah. 23 MR. GRIFFON: I wouldn't expect it any sooner, 24 you know. 25 DR. MAKHIJANI: Yeah.

1 MR. GRIFFON: There's only a few days left 2 here. 3 DR. MAKHIJANI: Just -- Just so I -- I 4 understand, Mark, though, it'll be the recycled 5 uranium section. MR. GRIFFON: Oh, yeah. Yeah. 6 7 DR. MAKHIJANI: There'll be some comments on 8 the 147 worker question including comments on -9 - on 95 percentiles and -- and maybe table 4-10 Yeah, and --5A. 11 MR. GRIFFON: And then possibly some other 12 fine-tuning of -- of language that -- from the 13 discussions today, right? 14 DR. MAKHIJANI: Okay. 15 MR. GRIFFON: The other --16 DR. MAURO: Yeah, Mark, this is John. 17 MR. GRIFFON: Yeah. DR. MAURO: For each one of these 11 items I --18 19 I took a lot of notes about the response that 20 was given and I think that we're in the 21 position where we can re-craft this report in a 22 way that would communicate that we posed this 23 issue; here is the response and -- and the 24 degree to which we consider to be the issue to 25 be resolved based on the information that we've

25

been given or -- or we may be in a place where we haven't yet had an opportunity to run it down. But I guess it'll effectively be as complete as we possibly can make it and bring down with us -- perhaps we can discuss it at the sub-committee meeting on Tuesday morning.

MR. GRIFFON: That -- That sounds like a plan. And -- And I would -- I would offer that what I'm going to try to do over this weekend and maybe with the work group's help to the extent I can get it, is to sort of do a -- a summary report. And this, a real over, you know, more over-arching, not as much -- not meant to have the kind of detail that we have in these other reports. But a summary report of where we are on the -- on the SEC evaluation. And it might -- it might, you know, to some -- I'm not sure how, if it's going to be a strong recommendation to the Board, but it's going to be, you know, I guess the work group's impressions of different areas of concern with regard to the SEC and then that'll -- that'll be hopefully, you know, be useful in our Board deliberations.

DR. WADE: And remember, Mark -- this is Lew --

1	that the Board will take up the Y-12 SEC
2	petition on Wednesday so we do have Monday
3	night, Tuesday night, you know.
4	MR. GRIFFON: Oh, yeah, we've got plenty
5	DR. WADE: Yeah, plenty of time.
6	MR. GRIFFON: plenty of time.
7	DR. WADE: The work group ought to get together
8	and look at the work product.
9	MR. GRIFFON: That's right. Well, I mean I,
10	you know, I would what I would offer is I
11	would try to draft something and and email
12	it as soon as possible and then maybe when we
13	get out there we can meet at night
14	DR. WADE: Right.
15	MR. GRIFFON: as a work group separately and
16	and, you know, fine tune language or
17	whatever.
18	DR. WADE: Okay. Just let me know your
19	pleasure and we'll make the arrangements for
20	the meeting.
21	MR. GRIFFON: Okay.
22	MS. MUNN: Are you going to make an effort to
23	tie your comments to the original matrix or
24	not? Well, that's a question that we can
25	develop later.

1 MR. GRIFFON: Yeah. I haven't thought that 2 part through. 3 MS. MUNN: That's not pertinent right now. 4 Just a thought. 5 MR. GRIFFON: Yeah. MR. GIBSON: This is Mike. Would it -- would 6 7 it helpful -- could -- I mean would it be 8 possible if perhaps NIOSH could have a -- a 9 little presentation ready for the Board, the 10 whole Board, about the status of the 11 Cyclotron/Calutron worker cases, the numbers 12 and the status for dose reconstruction so that 13 they would have a better overview and not just 14 try to take stuff from our matrix and then our 15 recommendations? 16 DR. WADE: Well, I think NIOSH could take those 17 comments to -- to heart as it prepares its 18 comments for the Board and do what it can do. 19 MR. GRIFFON: Yeah. And I certainly think anything -- I think Jim's, you know, you've --20 21 you've got the message that anything that 22 you've gleaned from this call today that you 23 think would strengthen your position I think, 24 you know, might not be in your evaluation 25 report but in your presentation you could

1 certainly --2 DR. NETON: Yeah. I'm a little bit sensitive 3 though in -- in terms of, you know, breaking 4 new information, you know. We -- We -- We 5 try to fix things and, you know --MR. GRIFFON: Well, yeah. 6 7 DR. NETON: -- before that but, you know --8 MR. GRIFFON: Maybe just if -- if they can be 9 presented as clarifications rather than --10 DR. NETON: Yeah. 11 MR. GRIFFON: -- modifications, you know. 12 DR. NETON: Right. I think everything we have 13 here right now is clarifications on these 14 issues. 15 MR. GRIFFON: Right. That's the way --DR. NETON: I'd like to do that. In some sense 16 17 I see sort of a -- sort of a different 18 framework for this presentation as compared to 19 other SEC petitions because, you know, we have 20 the SC&A report and I think -- I think the 21 Board -- full Board would probably want to hear 22 our -- our position on these issues, you know, 23 independent of the working group and --24 DR. WADE: All right. This is Lew. Just very, 25 very briefly, Y-12 will come up a number of

1 times. The first time it'll come up it'll be 2 the -- the sub-committee dealing with the 3 matrix as it related to the site profile so all of the issues can be talked about then. Later 4 5 that first day then I'll ask John Mauro to make a presentation of SC&A's work with regard to 6 7 the SEC review for Y-12, and there would be an 8 opportunity there then for you to do what 9 you're talking about, Jim, if need be to put some issues on the table. All of that 10 11 channeling into a Wednesday formal presentation 12 of the evaluation report hearing from the petitioners, the working group making its 13 report and the Board deliberating. So I think 14 15 when -- when SC&A presents its report would be 16 an opportunity, Jim, for you to put some things 17 on the table outside of the formal SEC 18 evaluation report. 19 DR. NETON: Okay. Sounds good. 20 DR. NETON: Yeah. 21 MS. MUNN: And excuse me, Lew. You said sub-22 committee. Did you mean working group? 23 DR. WADE: Well, I think when the sub-committee 24 meets on Tuesday morning I would expect that

the working group would talk to them about the

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1 matrices on the different site profiles so Y-12 2 will be talked about there in the more general 3 sense of the broad work that was done and, you 4 know, what remains to be done will get more 5 focused then on the SEC issues later that day and the next day. 6 7 MS. MUNN: All right. Just wanted --8 DR. WADE: Okay. 9 MS. MUNN: -- to get clarified. 10 MR. SMITH: This is Matthew Smith with the ORAU 11 team. Before you move off of Y-12 I want to 12 take 30 seconds just to let everyone know that 13 when we do apply the external co-worker data that we've been talking about, we do apply it 14 15 into IREP as a lognormal distribution so we're not just considering 50th percentile value 16 17 only. We're also applying a GSD value, a geometric standard deviation value that takes 18 into account the 95th percentile dose as well. 19 20 And that's just a point of procedure I wanted 21 everybody to know. 22 MR. GRIFFON: Okay. 23 DR. MAKHIJANI: Can you say that again? 24 didn't quite get that. 25 MR. SMITH: When we -- When we apply the

1 external co-worker data for Y-12, the data set 2 that's been under discussion all morning, when 3 we take that dose information and put it into 4 IREP, we apply it in the lognormal distribution. We do not just put in the 50^{th} 5 percentile value as a constant. We let IREP 6 know that the 50th percentile value is a 7 geometric mean of a lognormal distribution. 8 9 DR. MAKHIJANI: Right. 10 MR. SMITH: And then we also define a geometric 11 standard deviation and in doing that, that takes into account what the 95th percentile 12 13 value is. 14 DR. MAKHIJANI: But you're not using a fixed 95th percentile value? 15 16 MR. SMITH: No, we're not. 17 DR. NETON: That's right. And --18 MR. GRIFFON: That's a good question. 19 DR. NETON: -- before was like Bethlehem Steel 20 for example. And I appreciate Matt's comment. 21 That's very true. I'm not sure that gets us 22 past this other issue, though, of, you know, if 23 the workers weren't monitored properly then one 24 needs to think about the 95th. 25 MR. GRIFFON: Right. Good to clarify that.

1	DR. MAURO: Yeah, it was good clarification.
2	We weren't thinking in those terms.
3	DR. NETON: Yeah, it is. The GSDs are fairly
4	large. I think they're around 3.7 or something
5	like that for those distributions.
6	MR. SMITH: They're They're usually above
7	3, That's correct.
8	MR. GRIFFON: Okay.
9	DR. WADE: Okay. So let's close the chapter on
10	Y-12.
11	MR. GRIFFON: Yeah.
12	DR. WADE: And open it on Rocky Flats.
13	MR. GRIFFON: Well, all I would say is can we
14	take a five-minute because I know people from
15	Rocky are on the line. Can we take a five-
16	minute break to get our documents in order and
17	
18	DR. WADE: As you wish. And then when we come
19	back we'll do some introductions and make sure
20	we get the conflict of interest statements
21	done.
22	MR. GRIFFON: Yes.
23	DR. WADE: Then we can begin our discussions.
24	MR. GRIFFON: Okay. All right. Five minutes.
25	DR. WADE: Five minutes.

1 MR. GRIFFON: Bye. 2 (Whereupon, a brief recess was held.) 3 ROCKY FLATS 4 DR. WADE: Those are the principals. This is 5 Lew Wade. I'll keep the introductions very short. I think everyone knows the working 6 group, what the working group is about. We're 7 8 now going to look at issues related to the 9 Rocky Flats SEC petition. I would like members 10 of the NIOSH ORAU team to identify themselves 11 and state their conflicts or absence of, and 12 then the same with the -- the SC&A team. 13 are no conflicts with regard to Rocky Flats for 14 the Board members involved. Brant, for ORAU 15 NIOSH? 16 DR. ULSH: Sure. This is Brant Ulsh with NIOSH 17 and I have no conflicts at Rocky. 18 That might be it, Lew. It's awfully lonely 19 here. 20 DR. WADE: Okay. 21 MS. JESSEN: This is Karin Jessen from ORAU and 22 at this time I have no conflicts. 23 MR. ROBINSON: This is Al Robinson of the NIOSH 24 team. No conflicts.

MR. FALK: And this is Roger Falk. I am part

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1	of the ORAU ORAU. And yes, I have
2	conflicts with Rocky Flats.
3	MR. LANGSTED: This is Jim Langsted with the
4	ORAU team. I have conflicts at Rocky Flats.
5	MR. KENOYER: Judson Kenoyer with the ORAU
6	team. No conflicts with Rocky Flats.
7	MR. SHARFI: Mutty Sharfi with the ORAU team.
8	No conflicts with Rocky Flats.
9	MR. WOLFE: This is Craig Wolfe with the ORAU
10	team. No conflicts with Rocky Flats.
11	MR. MCFEE: This is Matt McFee with the ORAU
12	team. I have no conflicts.
13	MR. STEMPFLEY: This is Dan Stempfley with the
14	ORAU team. No conflicts.
15	MR. MEYER: This is Bob Meyer with the ORAU
16	team. No conflicts.
17	MR. SMITH: This is Matt Smith, ORAU team. No
18	conflicts.
19	DR. WADE: Okay. SC&A?
20	DR. MAURO: John Mauro, SC&A. No conflicts.
21	MR. FITZGERALD: This is Joe Fitzgerald. No
22	conflicts.
23	DR. MAKHIJANI: This is Arjun Makhijani. No
24	conflicts.
25	DR. BEHLING: Hans Behling. No conflicts.

1	MR. BUCHANAN: Ron Buchanan. No conflicts.
2	DR. WADE: Okay. Board members on the call,
3	please identify yourselves.
4	MR. GRIFFON: Mark Griffon.
5	MS. MUNN: Wanda Munn.
6	MR. GIBSON: Mike Gibson.
7	DR. WADE: Anyone else?
8	(No response)
9	DR. WADE: Okay. So we do not have a quorum
10	and we can conclude can conduct our
11	business. Mark?
12	MR. GRIFFON: Okay.
13	DR. WADE: Well, we should have petitioners
14	identify themselves.
15	MR. GRIFFON: Yeah.
16	MR. DEMAIORI: Tony DeMaiori, USW.
17	DR. WADE: Thank you Tony, and thank you for
18	your patience.
19	MR. GRIFFON: Okay. I think the best way to
20	proceed on this is probably going to be we
21	we had some matrix responses from Brant Ulsh
22	from NIOSH and we also have a a a summary
23	report that that SC&A agreed to provide
24	regarding the data integrity issues that arose
25	in the latter part of our matrix, many of them

1 out of the SEC petition items. So let's see. 2 I -- I think, and I'm -- I'm -- I'm hesitating 3 a little because I just now opened the report 4 that Brant forwarded so -- but I -- I -- I 5 imagine it might make sense to go through your 6 responses first to the matrix items and then -and then bring in SC&A's report and discuss 7 8 that. Is that -- Is that okay or does it make 9 sense to reverse that order. I'm -- I'm open 10 either way. 11 DR. ULSH: That works for me, Mark. Whatever -12 13 MR. GRIFFON: Okay. 14 DR. ULSH: -- you'd like to do. 15 MR. GRIFFON: We'll start with your report, 16 Brant. 17 Okay. I only focused on --DR. ULSH: 18 MR. GRIFFON: Does everyone have this report 19 first of all? Did the petitioners get this? 20 DR. ULSH: I don't know. I sent it out to SC&A 21 and to the working group members. 22 MR. GRIFFON: Maybe you can just tell the title 23 and stuff just to see if people have it. 24 DR. ULSH: Okay. I think it's called 12 April 25 Working Group Comment Responses. And that's on

1 the O-drive. I -- Again I don't know who --2 MR. GRIFFON: Yeah, 12 April Matrix Item 3 Responses. Did -- Tony, did you -- you have 4 access to this or --5 MR. DEMAIORI: I'm checking right now. believe I do. 6 7 MR. GRIFFON: Okay. 8 DR. MAURO: This is John Mauro. Did that deal 9 with a full range of issues or solely the data 10 reliability? 11 DR. ULSH: No, John. This was just -- actually 12 it's even more narrow than that. This is just 13 the outstanding action items that NIOSH had on 14 Mark's latest matrix that was sent out I 15 believe the day after our last meeting. 16 MR. GRIFFON: Right. 17 DR. MAURO: Mark, would it be of any benefit 18 to, in a broad way, to set the table so to 19 speak of the -- the range of issues and -- and 20 where we're going to sort of narrow it down and 21 focus in on within the con-- the overall 22 context of the petition at this point just for 23 orientation? 24 MR. GRIFFON: Sure. You know, I -- I know that 25 you didn't have time to do a review report at

this point, John, so I felt like that might be premature. But if you, you know, if you want to generally give a broad overview of where -- DR. MAURO: I guess it goes back to, yeah, there was an issues matrix for Rocky -- MR. GRIFFON: Right.

DR. MAURO: -- that -- that covered the full territory. And if you think that it's inappropriate or it's premature, to try to just sort of set the table but we certainly could just zero right in on the data reliability issues and get to work on those. That -- That's fine.

MR. GRIFFON: I think, yeah. I think that's probably best. I mean most -- most everybody has been on these calls before so there -- they know the matrix. They know the general items that we have on the matrix and I think we -- let's hone in on the work to be done understanding that we, you know, we -- you didn't do a review of -- of NIOSH's evaluation report yet. So let's just -- just hammer through this work I think and see where we're at if that's okay.

DR. ULSH: Okay. Mark, would you like me to

proceed?

MR. GRIFFON: Yeah, go ahead, Brant.

DR. ULSH: The -- The first action item that was still open for NIOSH related to comment number 9, action item number 6, and that's on page 1 of my handout. I don't know if you also have access to Mark's matrix. Maybe you do and that's on page 4 of 13. Now, this issue dealt with the Case 16 shift, and we discussed that at the last working group meeting. Jim Langsted gave a verbal response and I think Mark -- I think it was you who requested that we provide that in writing.

MR. GRIFFON: Yeah.

DR. ULSH: And that's what you see here in this response. The -- The bottom line is pretty much the last paragraph of that response, and that is that the Rocky Flats dosimeter algorithm does not utilize one chip specifically for the K-16 spectrum and it does not use a correction factor specific for that photon energy. So we don't believe that this is a -- an -- an issue with SEC implications but that -- that's -- that's our response on that one. I don't know if we want to discuss

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1 that or -- further or --2 MR. GRIFFON: Any -- Any comments? A lot of 3 us are receiving this real time so I don't know 4 if -- if SC&A has any comments on it. I really 5 think we just wanted a written documentation on that one. 6 7 DR. ULSH: Yes. So that's about it. That --8 That basically counts as a -- a written summary 9 of what we said at the last meeting. 10 MR. GRIFFON: Right. 11 DR. ULSH: So unless SC&A or anybody else has 12 any comments I can move on to the next one. DR. WADE: Go ahead. 13 14 DR. ULSH: Okay. 15 MR. GRIFFON: I wouldn't -- I wouldn't -- Let 16 -- Let me just clarify. I wouldn't assume 17 just because we don't comment that -- that 18 we're -- that these items are closed at this 19 point because --DR. ULSH: No, certainly not. 20 21 MR. GRIFFON: -- given that we just received 22 these so --23 DR. ULSH: Certainly not, yeah. I -- I 24 realize that we're operating them pretty close 25 to real time.

MR. GRIFFON: Right.

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DR. ULSH: In between when I write it and when you read it is pretty short a time.

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MR. GRIFFON: Exactly.

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DR. ULSH: Okay. The next one is also comment

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number 9 and it's action item number 7. And this deals with the nature and extent of the

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criminal investigations and/or security

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investigations that were mentioned by the

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petitioner in some of our previous work group

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MR. GRIFFON: Right.

meetings.

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DR. ULSH: Just to bring you up to speed we

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sent a letter -- I -- I sent a letter to Tony

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on, let me see, I believe it was March 16th and

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he responded. And there's a -- a copy of his

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response letter there on page 2 of my handout.

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And basically Tony recommended in that letter

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that we talk to Lisa Bretsler (ph) who is a

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person that works in records for -- I believe

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for DOE and we did in fact talk to her. At the

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last Board meeting I reported -- or at the last

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working group meeting, sorry -- I reported that

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she had also directed us to Jackie Baridini

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(ph) who is with the Kaiser Hill legal

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department. And basically what -- what -- what we found out in talking to those two individuals, Tony had suggested that we look for all abnormal radiation dose records that have resulted in a criminal and/or internal investigation at the Rocky Flats site for the last 50 years. And we ran that by Ms. Bretsler and she indicated to us that that -- that was going to be a pretty tough request to fulfill because it was so general. So we were looking -- she suggested and we kind of agreed that what we really needed were some specific examples. So to that end I had a phone conversation with Tony I believe it was Monday of this week and that was very helpful. was able to provide four examples that he thought were relevant to this issue and gave us enough specifics that we could go after some more information on this. And so I'd like to walk through those four and tell you where we are with them. I would caution, well, I guess everyone that some of this information deals with Privacy Act protected information and so we have to be very careful about how we talk about it. And I'm -- I'm not trying to be

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evasive or anything. I'm just trying to protect, you know, Privacy Act material. the -- the first example that Tony gave us was an individual who upon termination from the site, and this is pretty recently, gave a urine sample and also had a whole-body count. whole-body count came back negative and the urine sample came back high for plutonium and this initiated an investigation. Well, we basically accessed this person's file and we found the investigation report. Specifically what happened was Kaiser Hill convened a team of outside experts -- well, I'm going to clarify that. A team of experts that included most noted internal dosimetry authorities and also people who were familiar with Rocky --Rocky Flats operations to investigate this incident. We were able to locate the report that that expert team issued. I did place that report in the O-drive, the Rocky Flats folder that is, you know, there's a chain there, rather than email. But I talked to Mark over the lunch break and he was still not able to access that so --

MR. GRIFFON: It's still not there.

1 DR. ULSH: Still not there? 2 MR. GRIFFON: I'm on the O-drive now so --3 DR. ULSH: Yeah. I don't know. That's -- I'm 4 not sure what the issue is there but we will 5 try to get that report to you as -- if that would be of interest. I have, for the benefit 6 7 of the working group, reproduced the executive 8 summary of that report and I'd like to just 9 walk you through parts of that. That is shown 10 -- the executive summary is shown on pages 4 11 and 5 here. And what this expert panel 12 concluded I've summarized here on page 3. 13 14 I should update. It's there now. MR. GRIFFON: 15 DR. ULSH: Okay. This is real time. 16 MR. GRIFFON: 17 DR. ULSH: Yes, it certainly is. So refer to 18 the email that I sent out to you giving the 19 location of these files if you'd like to look 20 at it in its entirety. But the main 21 conclusions are listed on page 3 of the handout 22 here and here's what they say. They considered 23 several possible intake scenarios from this 24 incident and they found them to be implausible. 25 They considered inhalation, ingestion, wound,

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and they found that -- they concluded that these were not plausible intake scenarios. The other thing that sends up flags I think was the isotopic composition of the plutonium that was found in the urine sample. It didn't seem to match material that was present at Rocky Flats. And I believe -- keep in mind I just got this report about a day ago. I believe that the issue was that it was almost pure plutonium 239 which is not what you'd expect to see from the material at Rocky Flats. And also please keep in mind that I am speaking for NIOSH. trying to make any value judgments on -- on any of this. I'm just reporting what this expert investigation concluded so what the team, the expert team considered was the likelihood of external contamination of the sample prior to it entering the Kaiser Hill chain of custody. They also considered, due to the isotopic composition, almost pure plutonium 239, that this was consistent with a (unintelligible) source that could have been easily removed from the site. And the team concluded that deliberate contamination of the urine and fecal samples from an (unintelligible) source was

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plausible and could be accomplished with little risk to the person doing the tampering. Now, they didn't go into any detail beyond that as far as I can see in terms of hypothesizing when such tampering might have occurred. However, they did conclude -- the expert panel concluded that Kaiser Hill has implemented a very effective program, and I'm quoting now -- "a very effective program for determining the cause of the anomalous high urine bioassay results. The team felt that Kaiser Hill had been very thorough and complete in their approach to this unexpected occurrence." I -- I recognize that some individuals might take exception to the conclusions of this expert investigation. All I'm doing is presenting what this investigation concluded. They did not conclude that there was fraud on the part of the dosimetry staff at Rocky Flats and really if you want more details on -- on that particular incident I would refer you to the full report which apparently as of about three minutes ago is now available. That was the first example. The second example that Tony provided was one that we had actually

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already considered. I believe it was in the last working group meeting although they are all kind of blurring together for me. This was the one where the individual had submitted an affidavit as part of the SEC petition. copy of that affidavit is again presented on page 7 of my handout. And the main allegation, the main issue that was raised in this petition was that the worker stated that an entire year's dose record is missing from a time when he worked in a radiation area with dose rates ranging up to eight I guess Renkin per hour and this was during the 1982/1983 time frame. page 8 of my handout you'll find the dosimetry results for this individual. And again this is a recap because we've already discussed this in a previous meeting. And what you see here is that in fact in 1982 there are quarterly results for three of the four quarters and the monthly result that falls during the one quarter where there's not a quarterly result. And then in the next year, in 1983, there are quarterly results for all four quarters. And in addition there's another monthly result. the dosimetry for this particular individual

1	does not seem to support the claim that his
2	entire year's dose record is missing. And
3	that's really all I can say about that one.
4	The next example was an individual, a specific
5	individual that that Tony was able to
6	MR. GRIFFON: Which one did you just cover the
7	figure 4 that you were looking at?
8	DR. ULSH: Oh, sorry. Let me see. It is
9	figure
10	UNIDENTIFIED: Yes, it was.
11	MR. GRIFFON: I'm intentionally slowing you
12	down, too, so I can scan through the documents
13	as you're talking. I'm sorry.
14	DR. ULSH: Yeah, I apologize. Maybe I am going
15	too fast.
16	MR. GRIFFON: Yeah.
17	DR. ULSH: Figures 3 and 4 are the ones that
18	are relevant here, Mark. Figure 3 is the
19	affidavit that was provided in the SEC
20	petition.
21	MR. GRIFFON: Right.
22	DR. ULSH: And then figure 4 is the dosimetry
23	relevant to that particular individual for the
24	time frame that he cited.
25	MR. GRIFFON: Now, this is no different than

1 what you provided last time? 2 DR. ULSH: Exactly right. 3 MR. GRIFFON: Right, right. 4 DR. ULSH: It's just that this is one of the 5 examples that Tony mentioned in our conversation on Monday. 6 7 MR. GRIFFON: Okay. 8 DR. ULSH: So I -- I just presented it for 9 completeness. 10 MR. GRIFFON: Okay. 11 DR. ULSH: The next example -- and please feel 12 free to jump in if, you know, you want to 13 discuss any of these further. Example three 14 was an individual who Tony named for me, and we 15 were able to look at the dosimetry results for this particular individual. The -- The issue 16 17 here was blackened neutron badges and this 18 would be an issue during the era of MTA films. 19 And for this particular individual he began 20 work at the very end of the NTA film era in 21 1969. And the concern about blackened neutron 22 badges, I did a little digging on this and what 23 I found is in the neutron dose reconstruction 24 project protocol there's a phenomenon described

on page 16 of that document about gamma

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fogging. And what that involves is when a neutron badge is exposed to a high gamma field, and we're talking about 500 to 1,000 millirem, it can start to cause fogging on the film that progressively makes it more and more difficult as the doses get higher to read the film for neutron results. So I took a look at the -the -- the -- the gamma results for this individual and it doesn't seem like that would be the issue here because the highest -- the highest NTA film badge result that occurred for this individual during the period of 1969 was about 430 millirem. And so it doesn't appear that gamma fogging would have been an issue. And I should mention that there's no indication in this person's file that, in other words, a film where blackening was a problem. also during that period you might not expect to see such a notation.

MR. GRIFFON: I don't understand; maybe you can explain to me why -- why seeing 430 made you feel that there wasn't a problem for the one badge where I think he only --

DR. ULSH: No, what --

MR. GRIFFON: Did this individual say that

1 happened once or -- or multiple times? 2 DR. ULSH: It wasn't clear. What I'm saying 3 is, Mark, if gamma fogging becomes an issue 4 starting at approximately 500 millirem. 5 can still read the badge at around 500 but as 6 you progress up to about 1,000 millirem it becomes progressively more difficult to read 7 8 the badge. And since the highest result that I 9 saw during this film badge era for this individual, 1969 -- because remember in 1970 10 11 they began to switch over to TLDs. 12 MR. GRIFFON: Right. 13 DR. ULSH: So we're only talking about one year 14 here and the highest individual badge read 15 gamma dose that this individual had was about 16 430 millirem. All the rest of them were lower. 17 So I wouldn't really expect to see gamma 18 fogging on any of these particular badges. 19 That's the only point I was trying to make 20 there. 21 MS. MUNN: Brant? 22 DR. ULSH: Yeah? 23 MS. MUNN: Do you have a typo on this third 24 line? 25 DR. ULSH: Entirely possible.

1 MS. MUNN: Three? Shouldn't that have one more 2 zero? 3 DR. ULSH: Yes, it should. 4 MS. MUNN: Just checking. DR. ULSH: Thanks for the catch. 5 MS. MUNN: You bet. 6 7 DR. ULSH: That probably will not be the last 8 typo. Yes, that should be 1,000 beq. 9 doesn't appear that gamma fogging would explain 10 -- I mean if in fact there was --11 MR. GRIFFON: I guess what -- what -- I just 12 don't understand the rationale of that 13 argument. If -- I mean if -- if the 14 individual believed those doses as recorded 15 then there wouldn't be any issue at all. So I mean I don't -- I don't know that this sort of 16 17 demonstrates that he couldn't have one quarter 18 where he -- he was into some other area or 19 whatever and got higher exposures and that's 20 where the badge fogged. And -- And -- And 21 he's -- I mean here -- I don't know what the 22 claim specifically is here but are they 23 claiming that, you know, that it wasn't -- that 24 whatever dose was assigned was not accurate

because he had this badge fogging problem or --

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1 DR. ULSH: I don't know. I was just looking at 2 3 MR. GRIFFON: I'm just trying to understand, 4 too. 5 DR. ULSH: No, I understand. What I was trying 6 to do, Mark, is consider -- let's assume for 7 the -- for a minute that this individual did 8 have NTA films that were blackened. 9 trying to come up with and consider all 10 possible explanations for a blackened film 11 badge. And the first possible explanation that 12 I considered was gamma fogging. MR. GRIFFON: 13 Okay. 14 DR. ULSH: Now, his gamma results don't appear 15 to be consistent with gamma fogging. Again, if 16 you assume that the gamma results are --17 represent reality. 18 MR. GRIFFON: Okay. All right. 19 DR. ULSH: The second -- Really I didn't see 20 anything else in his file. I mean there was no 21 specific mention of -- of film blackening. But 22 however, we do know that it is possible that 23 NTA films can be blackened and there are a couple of situations that can lead to that. 24 25 One of them is that if NTA films are exposed to

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high temperatures and some pretty moderate humidities you can get thermal blackening. And I have provided some references there from peer review journal articles. That's at the top of page 9. That is one possibility if in fact there were some blackened films. Now, like I said, I wasn't able to locate any but let's just assume that, you know, that that was the case. And certainly it -- it happened at Rocky Flats that some people did have blackened film badge -- film badges. And another possibility is that -- is light contamination. As you -as you may or may not know, these NTA films were in light-proof packets and those packets could be damaged, could be ruptured. And just like any other photographic film, if it is exposed to light that could blacken a film badge. So I mean it certainly is possible that, you know, we would have film blackening. I didn't see any indication of it in this individual's file but certainly it occurred at Rocky Flats. But that was about as far as I could go with this one in the time frame that we have available. That's what I know on that individual. The last example that Tony

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provided to me was an individual and the petitioner, oh, some time ago after one of our earlier working group meetings, submitted a list of -- of about I think it was about 12 or 13 questions that resulted from the discussion that they heard and participated in during the working group meeting. And as it turns out one of those questions is relevant to this particular situation. And you'll see that question reproduced on the bottom of page 9 and I'd like to just read it to you. It says (reading) how are you addressing the fact that when a person received an abnormal or unexpectedly high dose and an individual -- oh, I'm sorry -- an internal investigation could not identify the source, the person received a zero for a dose? I know this to be true because it happened to me when I was pregnant in the 1999/2000 time frame. My dosimeter showed a high reading for ionizing radiation and an investigation was con-- was conducted and the reviewers could not find the source so they decided not to follow conduct of operations which said you have to trust your indicators, in this case, my dosimeter, and

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decided to enter a zero for my exposure. sure there are hundreds of examples like this so now my dose record is inaccurate and there is obviously no way to reconstruct it accurately since they failed to do so at the time. Now, in response to that question, I think this was back in March when this question was submitted to us. We provided the -- a response but you'll see at the bottom of page 9 and the top of page 10. There's a little bit of confusion here with regard to conduct of operation. What that refers to is that in order to ensure that workers are not overexposed when they're in the field, when they're in the presence of potentially hazardous environment, if you get an indication on instruments such as chirpers or Geiger-Mueller counters or anything like that, that you're in a high dose field, conduct of operations tells you that you should not question that result at the time; you should remove yourself from that environment and then an investigation can be conducted to determine whether or not the instrument was malfunctioning or whether you were actually in

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a hazardous environment. So that's to protect the worker, just to say don't question the instrument when you're standing in the potentially hazardous envir-- environment. Get out. That conduct of operations guidance doesn't necessarily apply to film badges and I think that was a little bit of a con-confusion because of course the worker is now, you know, out of the environment and we can -they can conduct an investigation. And that's exactly what they did actually. We were able to, since Monday when -- when Tony gave me this one, I was -- he gave me enough specifics that I was able to pull the records. Actually the ORAU team was able to pull the records for this particular individual, and what you'll see I combed -- well, we combed through the entire record and we did find an extended external dose reconstruction for approximately the right time frame and you'll see that on pages 11, 12 and 13. And here is what -- here is the conclusion from that investigation, and I'll just read you that. That's on page 10, summarized in the text, and it's also in the actual report which is on page 13. It says

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that this individual -- again I'm not going to use actual names -- but this individual became separated from her dosimeter while in the building 371 RBA thereby necessitating this extended dose reconstruction. She forgot to remove it from her anti-C (ph) clothing while doffing at the room 3408 step-off pad. individual was on a tour with two listed coworkers and was separated from her dosimeter for approximately 30 minutes. The individual is being assigned the zero dose listed on page 1 for the time that she was without her dosimeter. This dose is equal to the dose received by the listed co-workers who were with her on the entire tour. So what they concluded was during the brief time that the individual is not wearing her dosimeter but she was with the other people on the tour they took a look at the doses received by those other individuals and concluded that the dose to be assigned was less than the limit of detection So we didn't see any evidence that or zero. this investigation was in error. You know, I -- I suppose that a person could take issue with it but it wasn't clear to us that this was a

clear-cut indication of fraud because an 1 2 investigation was conducted and placed in the 3 individual's file. 4 MR. DEMAIORI: This is Tony DeMaiori with the 5 steel workers. I'm intimately familiar with 6 this case. And the individual was in fact on 7 tour, was giving a tour in building 371, a 8 communications person. And when the dose was 9 discovered it was almost six months later when 10 they were questioned and the investigation 11 occurred. And they were simply told that they were going to model after their co-workers who 12 13 worked in communications and received no dose. 14 And that's how the zero was going to be 15 applied. That's even though this individual 16 routinely toured the production areas and gave 17 tours. So this -- what you have is nowhere 18 near what the individual was told; not even 19 close. 20 It does sound like if that's what DR. ULSH: 21 the individual was told, it does sound like 22 there was some miscommunication going on 23 certainly. 24 MR. DEMAIORI: Hugely so.

DR. ULSH: What I have here though is -- is the

1 report, the extended dose reconstruction report 2 that's in the file. So I mean I -- that's I 3 think is what was done and that was the reason 4 for it. 5 MR. DEMAIORI: Yeah, I think they used what you guys call the worker model and they 6 7 reconstructed the dose to the other folks in 8 communications who never entered RA's. 9 DR. ULSH: Well, now, that's actually not what 10 the report at least says. It says that the 11 assigned dose -- hold on. Let me pull it up 12 here. This dose is equal to the dose received 13 by the listed co-workers who were with her on 14 the entire tour. 15 MR. DEMAIORI: And that was a zero. 16 DR. ULSH: Right. So I think that --17 that rather than the communi--18 MR. GRIFFON: So I guess the question there is 19 that if Tony's presenting this, you know, if 20 I'm understanding Tony, this person was the 21 tour guide --22 MR. DEMAIORI: Yes. 23 MR. GRIFFON: -- and would have done several of 24 these tours and got assigned a dose based on 25 two people that were taking a tour on a given

1 day. So maybe a whole quarter's worth of 2 information was zeroed. I don't know. 3 -- I guess that's the question, you know. 4 Maybe -- Maybe it was appropriate to use the -5 - to assign a co-worker exposure but were those representative co-workers? I know they were 6 7 only in the area for one tour and this 8 individual was in there giving tours all the 9 time. 10 DR. ULSH: Well, let's see. 11 MR. GRIFFON: That's not clear, I mean --12 DR. ULSH: I'm trying to track down the date of 13 the incident. Let me see if I can find that. 14 MR. WOLFE: Brant, I have it in front of me. 15 DR. ULSH: Okay. It was -- well, now I say that. Go 16 MR. WOLFE: ahead. May 2^{nd} , '01 was the date of the -- the 17 18 -- there was a radiological improvement report 19 that was part of the investigation report and 20 the event happened on that date, May -- May 2^{nd} , '01. 21 22 DR. ULSH: Okay. 23 MR. WOLFE: Part of the report, it said she --24 she was separated from her badge for 30 25 minutes.

1 DR. ULSH: Yeah. 2 MR. WOLFE: And when they found the badge in 3 the -- still attached to her anti-contamination 4 clothing in the laundry bag and surveyed it, 5 and it was uncontaminated. And I see --(inaudible) -- was contaminated. 6 7 DR. ULSH: But Craig, I also see on page 11 of 8 my handout there's a section, section 2, 9 dosimeter, and it says -- that section gives 10 the -- the needle date and the issue date, the 11 assign date, the return date. Those are all May 2^{nd} , 2001. 12 MR. WOLFE: Yeah. 13 DR. ULSH: So that seems to indicate that the 14 15 dosimeter was retrieved on the day this 16 incident happened and was read that day. 17 MR. WOLFE: Yes. So I -- I don't think that would 18 DR. ULSH: 19 represent the entire quarter. 20 MR. DEMAIORI: And when did that guarter end? 21 DR. ULSH: Oh, well, I don't know. The date is May 2nd so let me see. 22 23 MR. WOLFE: It would have been the end of June 24 most likely. 25 DR. ULSH: Yeah, but they pulled this -- pulled

this badge on May 2nd. At least that's what it 1 2 appears to indicate. 3 MR. WOLFE: Because the co-workers who are --4 who were -- who were used that their -- their date for their badge was May 2nd through May 5 9th, '01. 6 7 DR. ULSH: Yeah, that's listed on page 12 at 8 the bottom. 9 MR. DEMAIORI: And who were the co-workers? 10 The people on tour or --11 DR. ULSH: Yeah. 12 MR. DEMAIORI: -- the other communications folks that never entered the work area? 13 14 DR. ULSH: According to the report anyway on 15 page 13 you see this individual is being 16 assigned a zero dose listed on page 1 for the 17 time she was without her dosimeter, the dosage 18 equal to the dose received by the listed co-19 workers who were with her on the entire tour. 20 MR. DEMAIORI: Okay. 21 DR. ULSH: So it is the individuals who are 22 with her on the tour. 23 MR. DEMAIORI: I tell you what. I'll have the 24 individual affidavit, the -- the entire 25 incident to you because it's not the same.

1 DR. ULSH: Okay. 2 MR. DEMAIORI: I know it's --3 DR. ULSH: Tony, are you saying she was 4 separated from her badge for six months? 5 MR. DEMAIORI: No. No, no, not at all. 6 way it was described to me in detail is as 7 media relations manager of Rocky Flats part of 8 their duties was to give tours in production 9 areas, something the other communication folks 10 never did. And that they gave a tour and then 11 six months later she was informed that there 12 was an abnormality reading in her badge and 13 they wanted to know where she was. And she 14 told them she couldn't tell them; she didn't 15 know, that was six months ago. So they 16 assigned her a zero. Now, this is what I was 17 told. 18 DR. ULSH: Okay. Tony, I -- I agree with you. 19 If that's actually what occurred, I mean if it 20 was a situation where this individual was 21 assigned doses based on other people in the 22 department that weren't even on the tour or 23 giving tours that would certainly be a concern. 24 MR. DEMAIORI: Right. This may not even be the

same incident.

This doesn't even sound like --

1 it -- it remotely sounds like the same --2 DR. ULSH: Yeah. 3 MR. DEMAIORI: -- because the person was 4 pregnant at the time and they didn't waive 5 their right to go in the area. So it sounds remotely the same. 6 7 DR. ULSH: I do have --8 MR. DEMAIORI: But there's huge discrepancies 9 in the reporting in that. 10 DR. ULSH: Okay. Like I say, if you can -- I 11 mean if there's other information that would 12 indicate that we've got the wrong 13 interpretation here we would certainly --14 MR. DEMAIORI: Well, without you giving me a 15 name over the phone I couldn't tell you it's 16 the same incident even. The name of the individual? 17 DR. ULSH: 18 MR. DEMAIORI: Right. Give me their initials. 19 Give me something so that I can --20 MR. GRIFFON: Maybe offline you can do that. 21 DR. ULSH: Yeah, yeah. I'll -- I'll tell you 22 what, Tony. I'll get with you offline so that 23 we can talk about Privacy Act material or --24 MR. DEMAIORI: Okay. Because this is, you 25 know, what your reports are aren't even close

1 to what the individual had reported to me. 2 DR. ULSH: Okay. 3 MR. DEMAIORI: And this supports what we're 4 saying, that, you know, when doses aren't 5 believed they're given out as zero. 6 MR. GRIFFON: Okay. Brant, you should follow 7 up with Tony on that offline and --8 DR. ULSH: Okay. 9 MR. GRIFFON: -- you know. 10 DR. ULSH: Sure. Okay. That's the only 11 information I have on the four individual cases 12 that Tony provided. We also invited Tony to 13 provide, you know, if he can think of any others where you can give us some details so we 14 15 can run them down just like we have with this 16 one -- these four, that would be great. 17 invited him to do that by email and you're certainly welcome to do that. 18 19 Brant, this is John Mauro. DR. MAURO: 20 DR. ULSH: Yes, John. 21 DR. MAURO: On the first example, the 22 individual that had the high reading that might 23 have been -- there's going to be some follow-up 24 investigation, was there additional urinalysis 25 taken subsequent to see if in fact the person

1 had body burden or was in fact an after-the-2 fact contamination of his sample as you -- as 3 you described? 4 DR. ULSH: I -- John, I would be speculating 5 on -- on that because I got this report yesterday late in the day so I haven't had a 6 7 chanced to read through the details to 8 determine the exact sequence of events. 9 are --10 MR. DEMAIORI: John, I can give you that 11 information. I'm intimately familiar with the 12 investigation. 13 DR. MAURO: Okay. 14 MR. DEMAIORI: The individual had a high 15 bioassay urine sample as pure plutonium. 16 the individual was sent to Los Alamos National 17 Labs and they were poked and prodded and 18 absolutely nothing in their body, not in their 19 urine samples; not in their lungs. They were 20 brought back to Rocky Flats, given another 21 urine sample kit. It returned high plutonium. 22 DR. MAURO: And -- And there -- a continuing 23 follow-up related to that? 24 MR. DEMAIORI: I don't know. You know, Rocky 25 Flats is very sensitive on a happy closure.

DR. MAURO:

Uh-huh.

MR. DEMAIORI: And so I think everybody decided that the chain of custody was the real problem and that there was no way anybody could prove how the samples got the plutonium and so there was no follow-up after that. The recommendation was not to assign dose. They decided that the chain of custody, you couldn't prove anything one way or another because the chain of custody was weak and that's what the report will tell you.

DR. ULSH: Actually I'm looking at the -- the - well, at least the executive summary of the
report and the report -- and again, I'm just
quoting from the report. I'm not issuing a
value judgment from NIOSH. All I'm saying is
that the report concluded that Kaiser Hill
implemented a very effective program for deter- for determining the cause of the anomalous
high urine bioassay result. And the team felt
that Kaiser Hill had been very thorough and
complete in their approach. However if you
look on page 5 of my handout the team does
recommend additional analyses and actions and
that's on page 5; obtained three additional

urine and three additional fecal samples, and said that those samples were collected on September 23rd -- well, it gives you the dates there. They performed a radiological survey of the individual's home. They sent (unintelligible) to the analytical lab and to Los Alamos where they did thermal ionization mass spectroscopy. And they -- based on the first three recommend-- recommendations they recommended the team reconvene. So those are the follow-up actions that are at least listed in the executive summary. Again -- Again Tony, I haven't had a chance to really look at the bulk of the report and that is available on the O-drive.

DR. MAURO: Yeah. All I'm saying is that it sounds like Tony indicated that those results did come back and they came back negative.

MR. DEMAIORI: Yeah, everything from Las Alamos came back negative and then the -- the final urine sample came back positive again. That's when the team came to the conclusion that the sample itself was injected with the plutonium and not the individual. And, you know, to give you a point, the suspicion was the RAD sources.

We had the (unintelligible) plated RAD sources that were uncontrolled, literally hundreds of them.

DR. ULSH: All right. So that's -- That's what we have so far in the more specific examples. If there's -- Is there any further discussion on this one?

(No response)

DR. ULSH: Okay. Mark, would you like me to move on?

MR. GRIFFON: Yep.

DR. ULSH: All right. Comment number 9, action item number 8. And that's on page 14 of my handout. NIOSH ORAU to demonstrate the reliability of bioassay and external database data for the compensation program. And just to refresh your memory on what we've talked about in previous meetings. In terms of co-worker data I think that's one issue that we need to talk about. And I would remind you that the need for co-worker data at Rocky Flats is far less than what you might expect based on other sites. This is getting to be old information. It was, you know, a few weeks ago that I got this information. But to my knowledge we only

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have two identified cases that require external co-worker data and I don't think we're aware of any that require internal co-worker data at this point. But keep in mind there are 300 -approximately 300 cases left to do at Rocky Flats out of the 1,100 or so that we've received. Okay. So what we did -- what we've already done, the call, is we talked about the external co-worker data. The remaining question was for internal data. And remember that what we proposed to do is use the CEDR database to use -- to generate internal coworker data distribution. And previously we had compared CEDR to HIS-20 and we found at least what I would characterize as pretty good agreement. The remaining thread here I think this action is referring to was then going from HIS-20 back to some of the earlier records like the bioassay cards and the other database printouts that are contained in individual files. And we have made some (unintelligible). We took about 300-plus -- 306 worker samples from about 38 separate individuals and we compared what we see in HIS-20 with those earlier data sources, the Health Sciences data

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system and also in the earlier time period when they were using bioassay cards. And what we found is that for instances where there was data we found very good agreement. About 97.1 percent of the samples from the earlier data sources agreed with HIS-20 so we felt pretty good about that. In the remaining three percent where there was an imperfect match the data found on the bioassay cards, I think that was about seven of the samples, seven of the individual results, and six of those seven we found that the value in HIS-20 was larger than the card data. And then we also found that for 22 of the entries that there was an indication that the worker was not involved in the bioassay sample program because there was nothing in HIS-20 for them and there was nothing on the earlier bioassay cards so that's actually in agreement. Now, as you might expect there were some discrepancies. were about 41 individual results that we saw bioassay card data but we didn't see -- we have not yet located anything in HIS-20. We have some theories about why that might be but we're still running those down. But the point I

1 think that you have to consider, the next --2 the next obvious question would be, well, gee, 3 what does that do to your co-worker data? 4 Well, first of all, keep in mind that we're 5 using CEDR data which is the most complete data set for the early years -- early years we 6 7 think. And also, 40 of the 41 results that we 8 didn't find were below detection so -- and the 9 remaining one was just slightly above the limit 10 of detection. So what we concluded here is 11 that this doesn't appear to indicate that 12 there's a systematic censoring of high data. So I -- I think we still have pretty good 13 14 confidence in the co-worker data should we ever 15 have to use the co-worker data. 16 MR. GRIFFON: Brant? 17 DR. ULSH: Yes. 18 MR. GRIFFON: This -- I'm getting a little 19 deja vu here when I ask this question so excuse 20 me if I've already asked this question. 21 DR. ULSH: Sure. Go ahead. 22 MR. GRIFFON: But you're -- you're -- you're 23 presenting this as co-worker data. 24 DR. ULSH: Yes.

MR. GRIFFON: And I -- I think the real

1 question for me has always been the data 2 reliability more so. And -- And to what 3 extent do the claimants have raw data within 4 their file or is it often a printout of like 5 HIS-20 or CEDR data? And I don't know that answer. That's -- I might have asked it 6 7 before, too. 8 DR. ULSH: Well, I can give you -- I can speak 9 in generalities and maybe I'll let some of the 10 other site experts speak in more specific. 11 MR. GRIFFON: Because that's where it would be 12 more important is if a lot of the individual 13 claimants that you say have data, they don't 14 need co-worker data. 15 DR. ULSH: Yeah. 16 MR. GRIFFON: If it's only printouts from the 17 database then -- then you're back to the same, 18 you know. 19 DR. ULSH: Well, I think, Mark, and again I'm 20 going to rely heavily on the site experts here 21 but in the early years before the computer era 22 the bioassay cards were the dose -- the dose of 23 record. And I don't know exactly what years. 24 Roger or Craig, can you give me the years when 25 bioassay cards were the dose of record?

1	MR. FALK: Yes, the cards were the means to
2	the means to record the bioassay data through
3	1969.
4	DR. ULSH: Okay. Then after that, Roger, came
5	a database. Which one? Health Science?
6	MR. FALK: That was the Health Sciences
7	database.
8	DR. ULSH: Okay. And was that the official
9	dose of record then?
10	MR. FALK: Yes. Also for the people who were
11	active at that time all of the card data was
12	actually manually transposed into the Health
13	Sciences database.
14	DR. ULSH: Okay. After the Health Sciences
15	database then came I don't know what. Then
16	came what?
17	MR. FALK: Then we started to have the HIS-20.
18	DR. ULSH: Okay. And that year that was in
19	the late '90s or maybe 2000, HIS-20; is that
20	right?
21	MR. FALK: That was in the '90s.
22	` DR. ULSH: Okay.
23	MR. FALK: I don't know I don't know the
24	exact date of that.
25	MR. DEMAIORI: Late `90s.

1 DR. ULSH: Okay. So -- So Mark, the -- the 2 point that I'm making is there were different -3 - if you go over the years of operation of the 4 plant there were different systems for keeping 5 track of the dose of record. MR. GRIFFON: Well, when you say the -- the 6 7 Health Sciences database that's -- I -- I think 8 that's the first time I've heard that one but -9 10 DR. ULSH: Okay. 11 MR. GRIFFON: -- but if, you know, that -- and 12 you say that's -- that was the dose of record, this gets back to the same discussions we've 13 14 had with the Y-12, you know. That -- That --15 Maybe it's -- and I -- I don't, you know, I --16 I would -- would say you're -- you're probably 17 presenting it accurately but, you know, we went 18 through that with the Y-12 database that there 19 was I guess a letter from Y-12 and they sort of went through a process with DOE to accept the 20 21 database as the dose of record. Is there 22 anything like this in Rocky or --23 DR. ULSH: I don't know. I'm going to defer to 24 the experts. 25 MR. GRIFFON: Because otherwise I think you're

1 -- you're -- you know, the same question 2 applies. How do we, you know -- you haven't 3 chall-- you haven't checked that against the 4 raw records or -- or you did just do some of 5 that I guess in the --DR. ULSH: Well, we did respond. We certainly 6 did for the years when the bioassay cards were 7 8 -- were the dose of record. We did that. 9 MR. GRIFFON: Okay. 10 DR. ULSH: Now, the question I think would be 11 then if during the years when the HSDS, Health 12 Sciences Data System I think, was the original 13 dose of record, I'm not sure, you know, what --14 what kind of a validation you might be looking for here. 15 MR. GRIFFON: Well, I'm -- I'm -- it's the 16 17 first I heard of it so I'm just laying it out 18 there. I'm not sure either. 19 DR. ULSH: Sure. 20 MR. GRIFFON: But it seems to be that covers 21 '69 through '90-something, right or --22 DR. ULSH: Yeah, I think so. 23 MR. GRIFFON: -- thereabouts. 24 DR. ULSH: I think that's accurate. Yeah. And 25 then later HIS-20.

1 MR. GRIFFON: Right. 2 DR. ULSH: So --3 MR. GRIFFON: So you didn't really have any raw 4 records to compare against for those years from 5 '69 on but you did the earlier period? DR. ULSH: Yes. 6 7 MR. GRIFFON: What you presented here is from 8 the earlier period? 9 DR. ULSH: Well, when you say raw records, we 10 didn't have any handwritten records. 11 MR. GRIFFON: Right. 12 DR. ULSH: Yes. 13 MR. GRIFFON: Okay. 14 DR. ULSH: So the summary of -- of what we've 15 done is at the bottom of page 14. And we did 16 find pretty substantial agreement between the 17 bioassay cards, the HSDS database and the HIS-18 20 database. It is worth pointing out that 19 when we actually do dose reconstructions 20 however, we utilize all three sources of data 21 and that's to maximize completeness. Say for 22 instance there's nothing in HIS-20 but we have 23 earlier results on bioassay cards or maybe the 24 HSDS. We will certainly use those earlier

records. We'll supplement what we get from

1 HIS-20.2 MR. GRIFFON: And then this might also be 3 review but I think someone present -- I forget 4 who presented the HIS-20 CEDR comparison. 5 DR. ULSH: Yes. 6 MR. GRIFFON: And I was just, you know, looking 7 at some of that -- not that I had a lot of time 8 to look at it. 9 DR. ULSH: Yeah. 10 MR. GRIFFON: But HIS-20, there were -- there 11 were tables somewhere developed breaking this 12 down --13 DR. ULSH: Right. 14 MR. GRIFFON: -- HIS-20 '53 to '57, and CEDR 15 '53 to '57, and looking at that I remember 16 something in a discussion of the discrepancy in 17 the total number of samples was possibly due to 18 a lot of extra zeros which shouldn't have 19 actually been -- been put in the CEDR data is -20 - is what I recall. Maybe Roger indicated 21 that. But I -- I see, for instance, this time 22 period I have 10,158 samples in HIS-20 for that 23 time period. Of them I -- I -- that was 10,158 24 zeroes out of a total of 12,041 total data

points which was 84 percent. And then if I

1 look at the same time period for CEDR it was 2 16,412 zeros out of 18,888 -- 886 total data 3 points which was like 87 percent zeros. I just 4 wonder, you know, the difference in raw records 5 there is about 6,800 and you might have 6 answered this already but I -- just maybe to clarify that, why was --7 8 DR. ULSH: If I did --9 MR. GRIFFON: -- what would have caused that 10 difference there? 11 DR. ULSH: If I did answer it, Mark, I don't 12 remember so --13 MR. GRIFFON: I think Roger discussed it or 14 someone else. I know we discussed it on the 15 last call but --16 DR. ULSH: It might have been Joe Locktemy 17 (ph). I'm not sure. Roger, do you recall anything? 18 19 MR. FALK: Well, I was only commenting on the 20 lung count data --21 DR. ULSH: Oh, right. MR. FALK: -- about with regard to zeros but 22 23 I'm also thinking that the HIS-20 did not 24 capture the urine data for the workers who --25 for the workers who had retired from the Rocky

Flats site or -- or were terminated from the Rocky Flats site prior to 1977 and were not part of the benefits program. And so therefore the -- and therefore I would expect that the CEDR database would -- would contain more -- more -- more of the urine results than the CEDR database for those early years.

MR. GRIFFON: Okay. And I think -- and I have to look back on that analysis, too, but I think the general conclusion that he was making was that the -- the co-worker models would not have differed that much using either one of these approaches. Or there were some small differences but --

DR. ULSH: I think that's accurate, Mark.

MR. GRIFFON: Yeah.

DR. ULSH: I think that is what he was indicating. And -- And keep in mind that if what Roger says is -- is the explanation for the difference in the -- the number of records we do have -- for that earlier time period we do have the original dose of record which up to '69 would have been the cards and then from '70 up through -- up through '77 we would have the Health Sciences Data System printout.

1	MR. GRIFFON: Okay.
2	MS. MUNN: Mark, I lost you.
3	MR. GRIFFON: I lost myself once.
4	MS. MUNN: When you when you started giving
5	figures from the I was looking at the tables
6	from the database
7	MR. GRIFFON: Well, these these
8	MS. MUNN: assessment and follow-up
9	evaluation. Were you looking at something
10	else?
11	MR. GRIFFON: I have to the numbers I got
12	were from the on the O-drive within the co-
13	worker folder.
14	MS. MUNN: Oh, all right. Fine.
15	MR. GRIFFON: And there was a breakout of HIS-
16	20 versus CEDR, so yeah.
17	MS. MUNN: Fine. All right. All right. So I
18	no wonder I didn't have the numbers.
19	MR. GRIFFON: Yeah, right.
20	DR. ULSH: One of them was called a comparison.
21	MR. GRIFFON: Yeah.
22	DR. ULSH: I'm not going to get the titles
23	right but one of them was comparison and the
24	other one was follow-up
25	MS. MUNN: Follow-up.

1 DR. ULSH: -- comparison or something --2 MS. MUNN: Correct. 3 DR. ULSH: -- I think. Are those the two 4 documents you're talking about, Mark? 5 MR. GRIFFON: No, no, no. 6 MS. MUNN: Those are the two I was looking at -7 8 DR. ULSH: Oh. 9 MR. GRIFFON: No, I actually --10 MS. MUNN: -- and they are not the ones that --11 that Mark was looking at. 12 MR. GRIFFON: This is on the O-drive because I -- I -- we haven't received any of the HIS-20 13 14 or CEDR databases so I thought, well, maybe 15 they weren't put in the AB folder so I looked 16 in the co-worker data and to be honest with 17 you, I'm not sure exactly what sub-folder they 18 were in within the co-worker data but there was 19 -- there was actually -- I think someone broke 20 out the full database into -- into year span, 21 '53 to '57, '57 to '61, something like that. 22 MS. MUNN: Yes. 23 DR. ULSH: Now, that sounds a lot, Mark, like 24 what was in those two documents I mentioned but 25 I --

1 MS. MUNN: Yeah. 2 MR. GRIFFON: It might have been what you used 3 to create those doc-- yeah, those documents. 4 MS. MUNN: Yeah. 5 DR. ULSH: It might have been that. MS. MUNN: Very possible. 6 7 MR. GRIFFON: Because these were access 8 databases that I was looking at. 9 DR. ULSH: All right. 10 MS. MUNN: It was just --11 MR. GRIFFON: Yeah. 12 MS. MUNN: -- I was confused --13 MR. GRIFFON: Yeah. 14 MS. MUNN: -- because I couldn't find where you 15 were getting your numbers but --16 MR. GRIFFON: Well, the other -- the other 17 thing I no -- I noticed in there, and this just 18 might come into play in the -- in the models, 19 and like you said, that -- that may not be such 20 an issue because co-worker models are probably 21 not going to be used much, but in the CEDR 22 database, in the particular one I was looking 23 at anyway, for 19-- the last 20 or so values 24 all were -- all were in excess of -- let me --

let me present this correctly. HIS-20 had like

1 20 or so values that were greater than 935 and 2 I think we're talking DPM here. And when I 3 looked at it closely they all fell 4 approximately on the same -- I think all on the 5 same day or thereabouts, 6/15/57. In CEDR all those values were truncated off, and I wondered 6 7 if that was because they were related to some 8 incident and not thought to be applicable to a 9 general co-worker model or what -- what the 10 rationale was for that. And I -- I think that 11 -- that -- that's just a question on the co-12 worker models period, you know. Do you --13 DR. ULSH: Yeah. 14 MR. GRIFFON: Would -- Would that tend to be 15 an approach if you had incident data, would you 16 -- that was clearly from one incident 17 involving, you know, specific people, would you 18 tend to truncate that off your general co-19 worker models? So I don't -- yeah. If you 20 want to answer or don't have an answer --21 DR. ULSH: I -- I -- I don't really have an 22 answer to that right now, Mark. If you could 23 maybe provide the specifics in an -- in an 24 email to me I'll --25 MR. GRIFFON: Okay.

1	DR. ULSH: try to find an answer for you.
2	MR. GRIFFON: I'll type that up for you, yeah.
3	But it's 6/15/57 were the samples.
4	DR. ULSH: Okay.
5	MR. GRIFFON: And HIS-20 had high values and
6	CEDR has nothing.
7	DR. ULSH: Okay.
8	MR. SHARFI: Mark, this is Mutty. Those
9	Those can be chelated samples so in a co-worker
10	study you might not want to include those.
11	MR. GRIFFON: You know, could yeah. There -
12	_
13	MR. SHARFI: I'm thinking by chelation you
14	would not want to include them in your co-
15	worker study.
16	MR. GRIFFON: Right. And when I saw them all
17	in the same day it may just be that and I've
18	seen
19	MR. SHARFI: Depending on how high they were
20	they're they could they're they're
21	more like to be chelated.
22	MR. GRIFFON: And I think I've seen notations
23	in some of your co-worker spreadsheets where
24	there's a note in red at the bottom that says,
25	you know, this and this data point were dropped

-- found to be involved in a incident and not deemed applicable to co-worker model. So I don't know if that, you know -- there could be good rationale for this. I was just trying to understand it and whether that, you know, if that's a general approach. I thought maybe if, you know, a follow-up on that one, Bill, so maybe in general is that done for the co-worker models or for -- for Rocky for their source model.

DR. ULSH: Okay. Yeah. Mark, like I said, if
-- if you can mail that off to me I'll --

MR. GRIFFON: Sure.

DR. ULSH: -- I'll get you an answer or I'll do
my best to give you an answer.

MR. GRIFFON: Yeah.

DR. ULSH: Okay. So let me think about where we are here. I think we're on page 15 which is comment 12 from the matrix, and this deals with the no data available issue. And in previous discussions what NIOSH has said is that no data available could indicate two situ-- at least two situations that we can think of. One is a missed badge exchange. You know, a worker was on vacation or sick or, you know, maybe just

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forgot to exchange his badge in which case he would continue wearing the badge for an extra cycle. The second possibility is that the badge was turned in and there was a problem with the badge. And what I mean is there might be an investigation, there might, you know, such as during the TLD era; maybe a crystal was missing or something that would have required some additional processing so that the results from that badge weren't available at the time the report was issued. And what we've discovered as we've gone along with this issue is that the fact the place where these no data available entries appeared, but in the reports that were issued to the supervisors, they were computer printouts sent to the supervisors showing the dosimetry results for the people that -- that reported to that supervisor. And we actually over the past week, we have located some of those supervisor reports at the records center. We have pulled those and we are currently in the process of trying to go back and chase down some individual incidents -sorry, individual instances where no data available was on the supervisor reports and

1 compare those to maybe the dosimetry records 2 for the individuals. That is not complete yet. 3 We just got those records yesterday, but we 4 have made some progress. And that's really 5 about as much of an update as I can give you on that at the moment. 6 7 MR. GIBSON: This is Mike Gibson. If I could 8 step back for just a second, I had a -- I'd 9 like clarification from Roger Falk I believe it 10 was who was talking. How many people received 11 chelation at Rocky Flats? Do you have a record 12 of that, database of that, any documentation? 13 MR. GRIFFON: I think Mutty brought -- brought 14 up that possibility, didn't you? 15 MR. SHARFI: Yeah. 16 MR. GRIFFON: Yeah. 17 MR. SHARFI: Usually when you're assessing a case it's very clear in the record, both --18 19 both a part of the incident report and part of 20 their medical report about exact dates on when 21 they -- when they were receiving chelation, what -- how much, what type of chelating agent 22 23 was prescribed. Every -- Every case that I've 24 dealt with that had a chelation scenario had all that information inside their DOE file but

1 I don't have any good idea of the -- the total 2 number of chelating scenarios that they had at 3 the facility. 4 MR. GIBSON: I guess I can open up the question 5 up generally then. Does anyone with experience from Rocky out there know that -- basically 6 7 that number or is there any kind of --8 DR. ULSH: Roger, are you out there? 9 MR. FALK: Yes, I am out there. I'm trying to 10 -- I'm trying to draw that up. It is something 11 over 100 but probably less than 140 but I don't 12 have the specific number right -- right at 13 hand. 14 MR. GIBSON: That could be -- that could be dug 15 up out of some sort of data file if needed? 16 MR. FALK: Well, I'm not sure if it's really 17 pertinent but I think that the basic -- that 18 the basic statement is that if a worker was 19 actually chelated it would be in the claimant's 20 file that were captured by the -- by the 21 project so that the -- so that the dose 22 reconstructor would have that available. 23 MR. GIBSON: Well, whether it's pertinent or 24 not, that -- as a member of the Board I just 25 asked the question, is that available?

1 MR. FALK: I'm not sure. 2 MR. GIBSON: Okay. 3 MR. GRIFFON: You know, and this is a little bit of an aside here but I -- I'm also -- just 4 5 wanted to mention and I think maybe something 6 that might be important in the super-S model. 7 I think today the cases used for the super-S 8 TIB, were they chelation cases or were they 9 not? 10 DR. ULSH: Oh. 11 MR. GRIFFON: Do you recall that? I mean I 12 think you only -- at the end of the day you 13 used two cases, right, for your --14 MR. FALK: Six of the Rocky Flats cases were 15 the chelation cases and three were not. 16 MR. GIBSON: This is Mike Gibson again. 17 it be pertinent to a co-worker model? 18 MR. FALK: It would be pertinent to actually 19 exclude the urine samples that were actually 20 perturbed by the chelation and those were 21 generally coded as a code one in the Health 22 Sciences Database data. 23 MR. GIBSON: (Unintelligible) separated out by 24 a some kind of asterisk or notation. 25 MR. GRIFFON: Yeah, and that -- and that --

1 that could be one explanation of those ones 2 being separated out that I mentioned but --DR. ULSH: Well, it could be. 3 4 MR. GRIFFON: I think it's worth following up 5 on. DR. ULSH: Sure, Sure. And -- and Mike, I 6 7 think the answer to your question is yeah, it 8 sure would be relevant to make sure that those 9 chelation sample results don't make it into the 10 co-worker model. 11 MR. GIBSON: Right. 12 DR. ULSH: Yeah, that would certainly be something you'd want to do. 13 MR. FALK: But now, I would also like to point 14 15 out that if they did get into the co-worker 16 model it would be claimant favorable because it would tend to elevate the data set. 17 18 DR. ULSH: Well, and if there were only between 19 100 and 140 I'm not sure how much of an impact 20 it might have. The values of course would be 21 pretty high but you wouldn't expect all of 22 those to fall in the same year. 23 MR. SHARFI: Actually the people chelated were 24 using daily samples so they would have a 25 sizeable number of samples.

1	DR. ULSH: Oh, okay. All right.
2	MR. GRIFFON: Right. Thanks for that.
3	MS. MUNN: I can't imagine those would be used
4	for co-worker
5	DR. ULSH: No, I
6	MS. MUNN: co-worker data under any
7	circumstances. We have such a few number of
8	claims that are likely to be a part of co-
9	worker data.
10	DR. ULSH: Right. We haven't identified any
11	for internal that I know of.
12	MR. LANGSTED: This is Jim Langsted and I
13	specifically recall Joe Lochemy talking last
14	time about the fact that he did take that data
15	out of the co-worker data set.
16	MR. GRIFFON: Okay. Like I said, that may well
17	be the explanation for what I saw so that, you
18	know, that and I I don't remember Joe
19	saying that but he sure could have and so
20	DR. ULSH: Well, like you said, Mark, it could
21	be. But if you send us the statistics
22	MR. GRIFFON: Yeah.
23	DR. ULSH: we'll follow up on it.
24	MR. GRIFFON: Okay. Worth following up on,
25	yeah. All right.

1 DR. ULSH: Mike, did you have anything else or 2 do you -- should I move on or --3 MR. GIBSON: Yeah, go ahead. 4 DR. ULSH: Okay. All right. Let's see. Ι 5 think we were on comment 15 which is coincidentally on page 15 of my handout. 6 And Mark, I don't know. I may be confused. 7 I -- I 8 think that we addressed this issue on comment 9 9, action item 7. This was the follow-up with 10 the -- the petitioner on the -- on the 11 particular example. 12 MR. GRIFFON: Yeah. If you recall -- if you 13 recall it said I moved the comments from 9 to -14 15 DR. ULSH: Oh, okay. MR. GRIFFON: -- their individual comments so 16 17 they're the same one, yeah. DR. ULSH: All right. So we've already covered 18 19 that? 20 Yes. MR. GRIFFON: 21 DR. ULSH: Okay. Then comment 18 is the next 22 one, and this has to do with workers who 23 frequently did not wear badges in production 24 area and did not report non-use of the badge. 25 So this -- this is that I left my badge in my

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locker issue. And we had talked last time about some statistical analyses or -- or, you know, just looking at some of the data at least from the post-'77 years that we might be able to look at to get a feeling for this issue.

And we have done some of that. We have actually located I think, Jim Langsted, was the number 121 work reporters?

MR. LANGSTED: 239.

DR. ULSH: 239? Wow. Okay. So what we -what we've done is we started assembling graphs that you see on page 17 that indicate -- it's a cumulative dose graph. And what you might be wanting to focus in on on these graphs is instances where you might see a concave down shape. So as you go from left to right on the graph if you saw a flattening of the curve, that would be consistent with two situations at least that we know of. One would be the worker was approaching a dose limit of some kind and was removed from radiation work. The second situation that it would be consistent with is the worker continued in his job but his badge was removed from that environment. In other words, maybe left in his locker. I'm still not

1 clear on how we're going to separate that out 2 if we see it. Here are a couple of 3 representative graphs, about six of them on 4 page 17 where we did not see the kind of 5 flattening that we're talking about. And also 6 it's worth noting that --7 MR. GRIFFON: Each of these graphs represents 8 one individual or what -- what do these --9 DR. ULSH: I think it's one individual; is that 10 right, Jim? 11 MR. LANGSTED: Each -- the -- each one is a 12 different individual. 13 DR. ULSH: Right. 14 MR. GRIFFON: Yeah. 15 MR. LANGSTED: And these individuals were 16 picked because first of all they were exchanged 17 -- badges were being exchanged on a -- a semi-18 monthly basis which means they were identified 19 at the time as the highest potential dose 20 people and needed to be controlled periodi -- or 21 more periodically than others. And also we 22 selected from those the ones that had the 23 highest total dose for the year thinking that 24 those would be the individuals that would be

most likely to need dose control where you

1 might see that. 2 MR. GRIFFON: All right. 3 MR. FITZGERALD: This is Joe Fitzgerald. 4 facilities do these graphs or these curves 5 represent? 6 MR. LANGSTED: Don't have that data specifically, Joe. We just randomly picked 7 8 individuals. In fact the ID of the individual 9 was protected from me. 10 MR. FITZGERALD: Okay. 11 MR. LANGSTED: But my guess is there they'd be 12 plutonium production individuals. 13 DR. MAURO: This is John Mauro. So out of the 14 237 cases you looked at, how many of those had 15 this flattening someplace in the -- in the --16 in the cumulative distribution or the 17 distribution we're looking at? 18 DR. ULSH: Well, we're -- we're actually still 19 looking at -- at the data, John. We just got these data over the past week. I think in the 20 21 graphs that Jim sent me I saw one where there 22 might be some flattening but we're -- again 23 we're still looking at -- through the rest of 24 these. 25 DR. MAURO: I think the intent was not so much

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to say whether we could make a distinction of whether it was deliberate or leaving in the locker room versus a person who's taken off the job because of exposure. But just to see how often that occurred --

DR. ULSH: Sure.

DR. MAURO: -- that would be 237. You see it three times, I would say, well, that's not very often. Out of the 237 if you see it 237 times, then we have something that I guess we have to pay a little more attention to.

DR. ULSH: Right. And --

MR. FITZGERALD: Right.

DR. ULSH: -- keep in mind, John, I mean I'm just speculating here because again our analysis isn't complete yet. But as the years went on, as you got into the '90s say, and the dose limits, you know, got progressively lower over the history of the plant, and what you might expect to see is that as the limits got lower people may be approaching the limits more perhaps. And so you might expect to see more flattening. I don't know if that's what -what -- whether it'll turn out that we see.

We're just going to have to --

1 DR. MAURO: Yeah. 2 DR. ULSH: -- finish the analysis but --3 MR. GRIFFON: Brant, the reason you picked '77 4 was because you had monthly data or -- or --5 DR. ULSH: Jim, you had a pretty good 6 explanation for that and I'm not sure I'd do it 7 justice. Can you? 8 MR. LANGSTED: Okay. Yeah, Mark, in 1977 is 9 when the HIS-20 database first started 10 recording or -- or kept the exchange by 11 exchange data. 12 MR. GRIFFON: Right. 13 MR. LANGSTED: So if I was going to -- to do 14 that I -- I've got to be using more than just 15 quarterly totals. 16 MR. GRIFFON: That's what I thought. Okay. 17 MR. LANGSTED: That's why I started that. And 18 then I -- I thought probably 1989 is -- is 19 about the last time you want to look at this 20 because at that point production shut down at 21 the plant --22 MR. GRIFFON: Right. 23 MR. LANGSTED: -- and dose became less of an 24 issue. 25 MR. DEMAIORI: This is Tony DeMaiori. Prior to

1 shipment of the plutonium the last dose went 2 back up in the 2000s and we were doing a PUSPS 3 operations. 4 MR. LANGSTED: That's very true. 5 DR. ULSH: Okay. 6 So that's an explanation of what MR. GRIFFON: 7 -- Okay. So that's ongoing, Brant, is what 8 you're saying? 9 DR. ULSH: Exactly. Exactly. I just wanted to 10 update you on our progress so far. We had some 11 progress. 12 MR. GRIFFON: Okay. 13 DR. ULSH: Oh, also before we leave this issue, 14 as I was trolling through some of these 15 dosimetry files I did find an interesting 16 letter and that is on page 18 of my handout. 17 And what this is, it's a letter to a worker 18 notifying the worker that he's going to be 19 placed on radiation exposure restriction. 20 this is the first situation that we described 21 where a worker might be approaching a limit of 22 some sort and so that he's going to be pulled 23 out of the radiation environment. And the 24 interesting thing that you see here is that --

is that last paragraph that it says you will

1 remain in this classification, that -- that is, 2 say, you know, pulled out, restricted from 3 radiation work. You will remain in this classification 'til the end of this calendar 4 5 year. Your rate of pay will remain the same as 6 it is now. At the end of the year you will be 7 returned to Chem-Op Building 77-1. Now, I 8 don't want to make too much of this because I 9 don't know how generally it applies across the 10 years or across, you know, the plant for that 11 particular year, 1979. It does indicate though 12 that there might be less of an incentive for a 13 worker for financial reasons to engage in this 14 kind of manipulation of his dosimetry. 15 However, it should also be pointed out that the 16 petitioner mentioned that one reason a worker 17 might want to do this is to remain eligible for 18 overtime work and this letter certainly does 19 not speak to that situation. 20 MR. GRIFFON: Right. 21 DR. ULSH: But it's just one more piece of 22 evidence to add to the weight of evidence 23 approach that we're building here. 24 MR. DEMAIORI: I -- I guess -- this is Tony 25 DeMaiori. I'd like to speak on that. Wе

1 actually negotiated that into our collective 2 bargaining agreement. 3 DR. ULSH: Right. 4 MR. DEMAIORI: Which would speak just the 5 opposite. 6 MR. GRIFFON: Exactly, yeah. 7 MR. DEMAIORI: We wouldn't have wasted our time 8 if in fact people weren't suffering financial 9 loss. That's -- I'd also like to point out 10 that, you know, your rate of pay, your base 11 rate of pay, that's not premiums. That --12 Like when we re-entered the beryllium areas we paid time and a half for papper (ph) pay. 13 14 when we removed the beryllium hazard the papper 15 pay was removed. And we had two different 16 instances, one in 707 and one in building 444 17 where the people took their lapel samplers and 18 swept the floor, trying to restore the papper 19 pay. So, you know, for -- for this case you're 20 building that there was no disincentive that's 21 totally incorrect. 22 DR. ULSH: No, no, I'm not -- I'm not -- in 23 fact I was trying to be very clear that I'm not 24 saying that there was no disincentive because 25 this letter certainly does not speak to those

1 situations like overtime or the premiums that 2 you mentioned. It doesn't speak to that at 3 all. And that's the point I was trying to 4 make. 5 MR. DEMAIORI: Yeah. No, what that letter 6 speaks to is the language in the collective 7 bargaining agreement. 8 DR. ULSH: Right. And you see that in the 9 first paragraph of the letter, the article 4, 10 section 6 of the company union agreement. 11 MR. DEMAIORI: Absolutely. 12 DR. ULSH: Right. That's -- as you said, Tony, 13 that would be the basis for this no penalty in 14 the base rate of pay. So I -- I -- I don't 15 claim that this letter makes that issue go away 16 at all. 17 MR. DEMAIORI: Okay. 18 DR. ULSH: It's one piece of information to add 19 to what we've got. Should I move on or --20 MR. GRIFFON: Yeah. DR. ULSH: -- does anyone have any questions on 21 22 or discussion on that issue? 23 MR. GRIFFON: I think go ahead through. 24 DR. ULSH: Okay. That takes us to the last 25 page, page 19, comment 22, there was an action

1 This goes back to the instances of no 2 data available in situations of high exposure. 3 Again we -- we've located some of these 4 supervisor reports that we're trying to run to 5 ground now and I think the other concern that 6 was raised in -- in this particular situation 7 was the blackening of film and I think we've 8 already covered that under another comment, 9 too. I can go through it again if anyone would 10 like but if you're satisfied with that for now, 11 I can just leave it. 12 MR. GRIFFON: I think we're okay with that. 13 DR. ULSH: Okay. Well, then we're on to the 14 last item, comment 26. And this is the action 15 item was that we would provide co-worker 16 methodology to the Board and to SC&A. At the 17 risk of speaking without sitting in front of my 18 computer to see what's actually available out 19 there I -- I did see the co-worker data in the 20 location I've listed at the bottom of page 19. 21 I sure hope that all of that is out there now 22 for you guys to -- to review at your 23 convenience. 24 MR. GRIFFON: Question on that.

DR. ULSH: Yeah.

1	MR. GRIFFON: I mean as I'm pulling it open
2	again, did you put the Excel analysis files
3	with that, too, in the
4	DR. ULSH: Yeah, I think so, Mark.
5	MR. GRIFFON: It should be I know it's
6	somewhere else on there, too, but
7	DR. ULSH: I'm thinking it's in the co-worker
8	data folder and then there were some sub-
9	folders. Oh, boy, I'm trying to go from memory
10	here. I know that there's a folder for
11	americium and for plutonium and for uranium.
12	MR. GRIFFON: So all those all those folders
13	are there? Okay.
14	DR. ULSH: I think if you open those folders
15	there's a whole long list of spreadsheets in
16	there.
17	MR. GRIFFON: Okay.
18	DR. ULSH: But again, I'm trying to go from
19	memory so
20	MR. GRIFFON: All right.
21	MS. MUNN: Do you have the number of that of
22	those TIBs?
23	DR. ULSH: Yes, that is O-TIB 38 and O-TIB 58
24	although I can never keep it straight which is
25	external and which is internal.

1	MS. MUNN: That's okay.
2	MR. BUCHANAN: External is 58.
3	DR. ULSH: Okay. Thank you, Ron.
4	MS. MUNN: Thanks.
5	MR. GRIFFON: And at this point I'm not sure,
6	you know, we can really discuss 38 or 58 or any
7	of this extensively because I think we've
8	most of us have just been focused on Y-12 last
9	week so
10	MS. MUNN: Yeah.
11	MR. GRIFFON: Yeah.
12	DR. ULSH: Well, that takes you to the end of
13	my status update.
14	MR. GRIFFON: One other question on on the
15	data provided I'm just looking in the co-
16	worker folder.
17	DR. ULSH: Yeah.
18	MR. GRIFFON: I see the HIS-20 database from
19	(unintelligible); is that the one?
20	DR. ULSH: Yeah, that's I don't know if
21	that's internal or external, Mark.
22	MR. GRIFFON: Anyway, I see that but is there
23	also a a CEDR one or is
24	DR. ULSH: I don't think we provided the CEDR.
25	I don't know. I'd have to look again.

1 MR. GRIFFON: There wasn't -- you indicated you 2 had the CEDR in Access format, not in CEDR 3 format. 4 DR. ULSH: There was an issue about CEDR data. 5 We -- we have to -- according to the agreement, 6 to use CEDR data you have to only provide this 7 to an authorized CEDR user. And so I think 8 there at least was an issue about whether or 9 not we were free to do that. 10 MR. GRIFFON: Oh, okay. 11 DR. ULSH: However, if you are an authorized 12 CEDR user I think, again I'm going from memory 13 here, I think in the evaluation report, the 14 data sufficiency section, I listed the names of 15 the files from CEDR that we used. And this was 16 for I want to say the internal. And if you're 17 a CEDR user you could actually look at those 18 files in CEDR. 19 MR. GRIFFON: Yeah, yeah. I am a --20 DR. ULSH: I -- I know that's -- I know that's 21 not the most convenient but --22 MR. GRIFFON: It's not the best format to go in 23 CEDR either. I mean --24 DR. ULSH: Yeah. 25 MR. GRIFFON: -- (unintelligible) was better

1 but anyway. 2 DR. ULSH: I mean if you'd like us to pursue 3 that, Mark, we can investigate it further but -4 5 MR. GRIFFON: Well, I'm not sure how much we --6 we need it. I mean, yeah, I guess we'll push 7 through that when we get to the co-worker 8 models more but --9 DR. ULSH: Okay. 10 MR. GRIFFON: I mean really it was -- it was 11 for the purposes of comparison of the two. 12 DR. ULSH: Yeah. 13 MR. GRIFFON: And you provided that analysis 14 but just to have the raw materials there would 15 have been helpful. 16 DR. ULSH: Sure. I understand what you're 17 saying. 18 MR. GRIFFON: I mean I'll leave it at that for 19 now. 20 DR. ULSH: Okay. 21 MR. GRIFFON: Okay. I think -- is there 22 anything else on -- on Brant -- I mean this is 23 really a status report on these actions, many 24 of which you've completed but some are 25 outstanding and we've got those notes. And I

1 think the last thing we'll do -- it is getting 2 late. 3 DR. ULSH: Yeah. 4 The last thing we should do here MR. GRIFFON: 5 is get a presentation from -- from John --6 from SC&A or maybe it's Joe. I'm not sure who's presenting on the -- on their report. 7 8 And -- And, you know, the same probably 9 applies here. I'm not sure how much we can 10 discuss it because most people just received it 11 but at least have a little initial discussion 12 on it. 13 MR. FITZGERALD: Yeah, Mark. I -- I -- I have 14 Kathy here. We're in Los Alamos. 15 MR. GRIFFON: Oh, okay. 16 MR. FITZGERALD: What we can do is just 17 clarify. I think the last work group meeting 18 it was pointed out that because of the way a 19 lot of these issues were combined we -- we did 20 take the issues from the petition, combine them 21 with some from the site profile, came up with the 17 at Dr. Ziemer's request. And then we 22 23 tried to clarify where things stood and also

about the same time as we had the last work

group meeting, as we indicated, Kathy was, in

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fact, out at Rocky Flats talking to petitioners and beginning a process of trying to identify some additional documentation for the purposes of corroborating really, additional corroboration of -- of some of the issues that were -- were identified. I think it was the sense of the work group at the last meeting that it was kind of confusing tracking all these various issues. Some of them were overlapping and some of them had certainly different origins. Some of them were in fact in the process of being closed because they were recognized as not being SEC issues. so there was a lot of things in motion. we wanted to do for purposes of this discussion and the -- the SEC discussion of data integrity or data reliability was to sort of simplify it somewhat, and this is the purpose of the April 20th document which was to clarify both the major issues and the -- in a -- in a somewhat (unintelligible) the basis and reasoning behind our seeing these as sort of the key issues that need to be addressed in providing a pathway, which I think was the important suggestion that came out of the discussion last time.

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pathway to come up with sufficient corroboration that I think, you know, all would be satisfied that, you know, there was a -- a reasonable pursuit of whatever documentation could be obtained. And obviously this is a work in progress. We're still getting documentation in. I think it was the recommendation of the work group, however, that we clearly identify that which NIOSH was in a better position perhaps to pursue and -- and in the same breath maybe reserve some of the things that we were in the process of doing that we would like to complete. And that was the intent of this document was to clarify the basis but also identify a path forward that -that we would -- we could continue doing but also offer up as simply a suggestion for work group discussion of documents that could be obtained and what those documents may tell us that would be of usefulness in this process. So that's the backdrop. And I don't know how you want to go through this. Certainly I --MR. GRIFFON: Well, yeah. I think it's worth stepping through. I mean if you can summarize, Joe, but stepping through section by section

1 and then coming out -- each -- at the close of 2 each section I think you have some recommended 3 actions or -- for NIOSH and for SC&A, correct? 4 MR. FITZGERALD: That's correct. 5 MR. GRIFFON: So maybe just if you could step through in a concise --6 MR. FITZGERALD: Probably with Kathy since 7 8 she's right beside me here. And certainly we 9 start with data access as a backdrop. 10 MS. ROBERTSON-DEMERS: There are two -- two 11 sets of data that have not been reviewed. 12 (Unintelligible) dosimetry log sheets, that 13 type of information that I copied when I was 14 there that has not been shipped to me yet. The 15 other are these outstanding records. 16 DR. WADE: Kathy, we're not hearing you. 17 don't know if -- if you're on a speaker phone 18 but we're not hearing you. 19 MS. ROBERTSON-DEMERS: Is this better? 20 DR. WADE: Yes. 21 MR. GRIFFON: Yeah. 22 MS. ROBERTSON-DEMERS: Okay, there's two 23 outstanding sets of records. One set is -- is 24 the set that I copied while I was there and 25 that they were supposed to ship to me which I

haven't received yet. And that's from the boxes I did review. And then there were the records that I requested that were not pulled back from the Denver Federal Center while I was there. And I kind of summarized in table 1 the documents that I originally was looking for and which ones I -- I walked away with. And the -- the box of -- there's probably about 1,000 sheets of paper. There's just kind of a mishmash of all -- all sorts of things, everything from tritium to TLD log sheets to external dosimetry technical documents, that type of stuff.

DR. ULSH: Kathy, this is -- this is Brant.
With regard to the first set of data that
you're talking about, the ones that you've
copied but they've not yet been shipped to you.
MS. ROBERTSON-DEMERS: Uh-huh.

DR. ULSH: Is there anything that you can think of that NIOSH can do to maybe assist in that process? I mean are -- do you know what the issue is as to why they haven't been shipped?

MS. ROBERTSON-DEMERS: I do not know. The person that -- my contact has not gotten back to me for about a week.

1 DR. ULSH: Oh, okay. 2 MS. ROBERTSON-DEMERS: And probably what the 3 best action is is for me to try again. 4 DR. ULSH: Okay. 5 MS. ROBERTSON-DEMERS: And then if they don't turn around and provide it then for NIOSH to 6 7 step in and say, hey, we want those records 8 sent. 9 DR. ULSH: Yeah, I mean we've got people out 10 there who have a, you know, fairly good 11 relationship with records, you know, the DOE 12 records personnel. And, you know, again, I --13 it's hard for me to say without knowing what 14 the issue -- what the holdup is but, you know, 15 I mean if -- if it's just a matter of one of 16 our people driving down to the records center 17 and saying hey, you've got some boxes on hold 18 for Kathy DeMers, we'll take 'em and get 'em to 19 her, I mean we can do that. 20 MS. ROBERTSON-DEMERS: Let me catch up with 21 Andrea. 22 DR. ULSH: Oh, is this Andrea Wilson? 23 MS. ROBERTSON-DEMERS: Uh-huh. 24 DR. ULSH: Oh, okay. Okay. Yeah, that's --25 that's one of our contacts actually.

1 MS. ROBERTSON-DEMERS: Yeah, and I've been out 2 of town --3 DR. ULSH: Okay. 4 MS. ROBERTSON-DEMERS: -- this week so --5 DR. ULSH: All right. MS. ROBERTSON-DEMERS: At least let me get home 6 7 and make sure that they --8 DR. ULSH: Yeah, they may have been shipped, 9 right, so --10 MS. ROBERTSON-DEMERS: -- don't show up. 11 DR. ULSH: Okay. 12 MS. ROBERTSON-DEMERS: Now, there's a -- a later table in here that -- they're more into 13 14 table 1, table 4 and what I did was I scrunched -- it doesn't look like it but I scrunched the 15 16 type of records that would be helpful if they 17 were pulled. These -- these are really those records that I wanted to see but didn't get to 18 19 see. And those years correspond to individual situations in the SEC petition so I tried to 20 21 overlap. This person said he worked on this 22 job in this area for 1982, 1983 so I tried to 23 pull the logbook from that area for 1982/'83. 24 And I guess the -- the important thing about

those logbooks is that I'm told that there's

1 personnel dose information in them and that 2 that dose information doesn't correspond to the 3 dosimetry record. So that's what I was trying 4 to -- to check on. Now, you'll see that I said 5 select years on some of these. I realized 6 that's a lot of logbooks but those are the --7 those are the years that cover particular 8 people and the intention was to just pick a 9 couple of them through that period for that 10 building and compare it back to that person's 11 dosimetry record. 12 DR. ULSH: I would -- I would say if you're 13 doing that, too, it might be worthwhile if possible to make sure we have a good coverage 14 15 of the years, you know, the decades I should --16 I should say. Like let's not pick them all 17 from the '80s or all from the '90s or --18 MS. ROBERTSON-DEMERS: Right. 19 MR. GRIFFON: -- you know, yeah. 20 MS. ROBERTSON-DEMERS: Well, you'll see there's 21 quite a variety. 22 MR. GRIFFON: Yeah, there's a range I see in 23 your table. 24 DR. ULSH: Just from -- to get a point of 25 clarification, Kathy, I'm looking at table 4 --

1	MS. ROBERTSON-DEMERS: Okay.
2	DR. ULSH: the ones where you see select
3	years between '63 and '95.
4	MS. ROBERTSON-DEMERS: Uh-huh.
5	DR. ULSH: When you see that are are you
6	indicting that there are specific years that
7	you're looking for or rather that you're
8	interested in a random sampling of of those
9	years?
10	MS. ROBERTSON-DEMERS: That's what I was
11	talking about.
12	DR. ULSH: Yeah.
13	MS. ROBERTSON-DEMERS: A random sampling. I
14	don't expect
15	DR. ULSH: Oh, I see. Okay.
16	MS. ROBERTSON-DEMERS: this whole every
17	logbook for that building from '63 to '95.
18	DR. ULSH: Okay. Right.
19	MS. ROBERTSON-DEMERS: You know, I'm just
20	pull five or something. Those years are
21	associated with a particular person being in
22	that building over that time period because
23	they didn't specify a particular year.
24	DR. ULSH: Okay. I think I see now. Okay.
25	MS. ROBERTSON-DEMERS: Okay. I tried to under

1	the RFP-SEC petition matrix I tried to kind
2	of come down to
3	MR. GRIFFON: What page are you on, Kathy?
4	MS. ROBERTSON-DEMERS: I'm on page 4.
5	MR. GRIFFON: Okay. (Unintelligible) backup.
6	MS. ROBERTSON-DEMERS: And I already see an
7	error in this list. It's the bulleted area.
8	MR. GRIFFON: Okay.
9	MS. ROBERTSON-DEMERS: I tried to kind of boil
10	it down to to the issues, the core issues.
11	And one of those, the other radionuclides we
12	dropped in the back but apparently we didn't
13	drop from this list.
14	MR. GRIFFON: Which bullet item is that?
15	MS. ROBERTSON-DEMERS: That's the last one.
16	MR. GRIFFON: Last item, okay.
17	MS. ROBERTSON-DEMERS: And I tried to tie it to
18	one of the matrix issues just to give you
19	MR. GRIFFON: Okay.
20	MS. ROBERTSON-DEMERS: a reference back to
21	that to what particular matrix issue
22	brought this particular situation up but
23	MR. GRIFFON: That bullet is dropped is what
24	you're saying?
25	MS. ROBERTSON-DEMERS: Yeah. But really for

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several of these I would follow the same process in trying to evaluate it, going from the dosimetry record to the processing logs to the logbook like the no data available, false entries, zeros where they were -- where they expected to have high dose, those types of things. The same process will be used.

MR. GRIFFON: Okay.

MS. ROBERTSON-DEMERS: With respect to the external dosimetry investigations, really what -- what that's about is how did they do it, especially before the time period of the procedures that NIOSH has cited, and how did they document it or how were they told to document it and did they indeed document that. And I provided a table from actually a TLD problem logbook and each of these entries have had an issue which -- which I listed out. I'll try and give you the table number here, and that's on page 7. I would have expected them to say something in -- in the worker's file about it or have some policy on how to deal with that situation. And those ID numbers should -- should allow you to track back to an individual. Basically what we want to see is

1 whether they actually indeed did do -- did have 2 a process in place to assign doses when there 3 was a problem with the -- with the film badge 4 or it was lost or there was an overexposure 5 like when the film was black. And was that process formally documented. In other words, 6 did they do an extended external dose 7 8 reconstruction and go out and talk to co-9 workers and find out where this person was 10 working and that type of thing. 11 MR. GRIFFON: Kathy, can we -- I -- I guess if 12 we could step through section by section now --13 you've kind of given an overview there. 14 Section 1 you have suggested NIOSH follow-ups 15 and SC&A follow-ups. I think --16 MS. ROBERTSON-DEMERS: Okay. Let's go back --17 MR. GRIFFON: -- it might be worthwhile to be 18 clear what we expect, you know. 19 MS. ROBERTSON-DEMERS: Okay. Well, let's go to 20 page 5. 21 MR. GRIFFON: Okay. 22 MS. ROBERTSON-DEMERS: And this has to do with 23 the safety concerns. 24 MR. GRIFFON: Right. 25 MS. ROBERTSON-DEMERS: Ones that -- that have

1 been filed -- filed. What I did is I got a 2 list of the safety concerns and really it just 3 had brief descriptions. And I picked out those 4 that were relevant to dosimetry. And hopefully 5 NIOSH has gotten 71-4 but that -- but I guess you'll get -- NIOSH will have to tell me if 6 7 they've gotten that one. 8 DR. ULSH: Kathy, who did you send -- I don't 9 think I've got -- got it. Who did you send it 10 to over at NIOSH? 11 MS. ROBERTSON-DEMERS: Probably it would have 12 come through formal general. John, are you 13 still there? 14 DR. MAURO: Yes, I am. Okay. I quess I don't have it. Or if I do I don't know I have it. 15 16 MS. ROBERTSON-DEMERS: Okay. 17 DR. MAURO: You thought it was forwarded to me? 18 MS. ROBERTSON-DEMERS: Yeah. Probably a CD I 19 sent. 20 DR. MAURO: You sent it recently? 21 MS. ROBERTSON-DEMERS: Yeah. A CD. 22 DR. MAURO: Oh, okay. I did receive a set of 23 CDs from Judy. Are you referring to CDs that 24 went first to Judy and then to me? 25 MS. ROBERTSON-DEMERS: Right.

1	DR. MAURO: I have it. Yes, I do. I have
2	those CDs.
3	MR. GRIFFON: So you have to SC&A can still
4	work on providing that to NIOSH, Brant, yeah.
5	DR. MAURO: Okay. So okay.
6	MS. ROBERTSON-DEMERS: Yeah, it's really just a
7	single sheet of paper.
8	MR. GRIFFON: Okay.
9	DR. MAURO: Okay. So but I received two
10	I two separate days I received two sets of
11	CDs. Now, just let me know what you'd like me
12	to do with those because I distributed them
13	internally to SC&A folks but I did not forward
14	anything on to NIOSH.
15	MS. ROBERTSON-DEMERS: Okay. Well, that safety
16	document should be on the Rocky Flats CD.
17	DR. MAURO: Okay. So you would like me to send
18	
19	MS. ROBERTSON-DEMERS: Forward that.
20	DR. MAURO: Okay. The Rocky CD, and I'll send
21	that to whom?
22	MR. GRIFFON: TO NIOSH.
23	DR. ULSH: Yeah, if you could send it to me,
24	John Brant Brant Ulsh, that would be
25	good.

1 DR. MAURO: Okay. I'll take care of that. 2 It'll go out tomorrow. 3 DR. ULSH: Thanks, John. 4 MS. ROBERTSON-DEMERS: Okay. And then there's 5 several other safety concerns and some of them kind of track very well with concerns that were 6 7 in the petition and I would just recommend that 8 those safety concerns be pulled. And some of 9 the files, well, there's -- there's -- there's 10 a company response to each safety concern so 11 the concern is listed and the company response 12 is listed. 13 MR. GRIFFON: Okay. So we have a limited set 14 of safety concern reports here, and the recommendation is for NIOSH to pull these and 15 16 evaluate 'em, right? 17 MS. ROBERTSON-DEMERS: Right. MR. GRIFFON: And SC&A is also going to 18 19 evaluate a couple that you already have, three 20 that you --21 MS. ROBERTSON-DEMERS: Well, we have 71-4 --22 MR. GRIFFON: -- that are coming under --23 MS. ROBERTSON-DEMERS: -- but we'd like to see 24 probably 87-206 and 92-036. 25 MR. GRIFFON: Okay. So you can do a few in

1 parallel is what you're suggesting? 2 MS. ROBERTSON-DEMERS: Right. 3 MR. GRIFFON: Okay. 4 DR. ULSH: On -- on that list of safety 5 concerns, Kathy, were -- were these documents that you requested and -- and DOE was not able 6 7 to provide them? 8 MS. ROBERTSON-DEMERS: The -- I originally 9 requested 71-4 when I was at Rocky Flats 10 because I just discovered them when I was 11 there. 12 DR. ULSH: Oh, I see. 13 MS. ROBERTSON-DEMERS: And the remainder have 14 not been requested. 15 DR. ULSH: Okay. Not yet requested, okay. 16 Okay. I guess my thoughts are that I mean we 17 can certainly try to get them. It will take 18 some time I think. Pretty much everyone agrees 19 that I mean that certainly can't be 20 accomplished before the Board meeting. We can 21 try. Our -- our experience is that the 22 classified records are fairly well organized. 23 The unclassified records not so much. I quess 24 if -- if the group decides that we want to 25 pursue these documents or any other documents

1 in this -- in this -- the SC&A's report then 2 the next step for us would be to talk to our --3 our contacts, the records people, and find out, 4 you know, what kind of a time frame we're 5 looking at on getting these. I mean I'm not sure that we'll be able to do any better than -6 7 - than Kathy did but --8 Right, right. MR. GRIFFON: 9 DR. ULSH: -- we can try. 10 MR. GRIFFON: Yeah. And maybe you can, you 11 know, we can just -- we just have to keep on 12 top of this and you can give us an update on -on how you -- you -- I mean I think it's 13 14 important especially since at least a few of 15 them -- SC&A is arguing that they directly tie 16 back to some of our matrix items and they're --17 they're -- these are issues that were raised by the petitioner. 18 19 DR. ULSH: Right. 20 So I think to that extent I think MR. GRIFFON: 21 it would be good at least to attempt. And, you know, then if it's taking -- I mean we'll --22 23 we'll just try to keep on top of it and --24 DR. ULSH: Okay.

MR. GRIFFON: Yeah. If we're not getting

1 anywhere then we -- we at some point have to 2 pull the plug. We understand that. 3 DR. ULSH: Mark, a point of procedure. Are you 4 capturing these in a -- and --5 MR. GRIFFON: Yeah. DR. ULSH: And it will be coming out in a 6 7 matrix? 8 MR. GRIFFON: Yeah. 9 DR. ULSH: Okay. Good. 10 MS. ROBERTSON-DEMERS: Okay. The -- the next 11 section is the external dosimetry procedures. 12 And this somewhat goes back to the lost chip 13 issue that's in the petition but it -- it's 14 really broader and covers all sorts of -- of 15 issues. And again it's getting back to 16 verifying that they actually did do a valid 17 external dosimetry investigation when there was 18 a problem with the badge. One of our concerns 19 was that the extended external dose reviews 20 procedures were from the '80 -- from the '90s. 21 And what we'd like to -- to see is the --MR. GRIFFON: And the one that Brant mentioned 22 23 on the last call is from '83, right? 24 MS. ROBERTSON-DEMERS: Well, that -- that --

that's an actual processing procedure.

1	MR. GRIFFON: Oh, okay.
2	MS. ROBERTSON-DEMERS: Not a dosimetry
3	DR. ULSH: Yeah, that's correct. I think
4	that's the Lincoln Penox (ph) document?
5	MS. ROBERTSON-DEMERS: Right.
6	DR. ULSH: Yeah, I think I think you've
7	accurately described it. That's a dosimetry
8	processing procedure.
9	MR. GRIFFON: Oh, okay.
10	DR. ULSH: Yeah, I don't know.
11	MR. GRIFFON: I just did that on the last call
12	I think. I thought that was similar to to
13	the other investigation procedures but
14	DR. ULSH: There are some overlap but not
15	complete overlap.
16	MR. GRIFFON: Okay.
17	DR. ULSH: And I I don't know. I mean
18	without looking we're certainly not aware of
19	earlier procedures. We haven't been able to
20	locate any but but I don't know. I mean
21	MR. GRIFFON: Okay.
22	MS. ROBERTSON-DEMERS: In this case I was able
23	to in table 3 give you examples from the
24	'85/'86 logbook because that's what I had
25	access to at the time. But certainly you need

1	to do snapshots in time including the era
2	before 1983 and just kind of work your way
3	backwards.
4	DR. ULSH: Now, what I guess I'm just trying
5	to clarify what we're going to do on these. So
6	we've got some examples here on table 3 where
7	there were some problems with I guess in this
8	time frame it would have been the crystals in -
9	- in the TLD badges.
10	MS. ROBERTSON-DEMERS: Or Or the badge was
11	contaminated or
12	DR. ULSH: Right. Right.
13	MS. ROBERTSON-DEMERS: There were reader
14	errors.
15	DR. ULSH: Right. Problems with the TLD of
16	various types. And what I guess what are
17	we looking for to further inform us about
18	these? Are we looking for
19	MS. ROBERTSON-DEMERS: Let's just Let's
20	just walk through one.
21	DR. ULSH: Okay.
22	MS. ROBERTSON-DEMERS: Let's say I was looking
23	at 514479.
24	DR. ULSH: 514479 Okay, I see it.
25	MS. ROBERTSON-DEMERS: Okay. The first thing I

1	would do is compare it back to that person's
2	actual dosimetry file.
3	DR. ULSH: Okay. That's easily enough
4	easily enough accomplished I think.
5	MS. ROBERTSON-DEMERS: And see if there's any
6	indication of this in how they investigated
7	that and how they ultimately assigned the dose.
8	DR. ULSH: I I can Jim, can you Jim
9	Langsted, can you I know that in the later
10	period, certainly in the '90s sometime forward
11	they put extended extended and abbreviated
12	dosimetry investigation reports in the file.
13	How far back in time does that go? Do you have
14	a feel for that?
15	MR. LANGSTED: Probably mid-90's.
16	DR. ULSH: (Inaudible) in the logbook that we
17	see here in table 3?
18	MR. LANGSTED: My guess is no there would not.
19	MS. ROBERTSON-DEMERS: Then the question
20	becomes how did they assign the dose for that
21	particular situation?
22	DR. ULSH: Yeah, I mean I don't know that we're
23	going to get any more information than what you
24	might see in the logbook. I mean I I'm
25	trying to go from memory from the example pages

1 that you provided at the last working group 2 meeting and there was a justification for 3 change. I think there was also maybe a column 4 that showed the dose that was assigned but I 5 could be mistaken in that. 6 MS. ROBERTSON-DEMERS: And the question is how 7 did they determine that dose? 8 MR. LANGSTED: Well, I mean, again --9 MS. ROBERTSON-DEMERS: Because I guess the 10 contention by the petition is well, the badge 11 was blacked out and I got a zero. Well, they 12 had to have a reason for assigning zero. 13 MR. GRIFFON: Well, I think the -- the other --14 I mean I'm not -- I'm hearing what -- what 15 Brant's saying is that they likely wouldn't 16 have anything in the file to show how they 17 treated these. Or is that what you're saying, 18 Brant? I mean --19 DR. ULSH: Well, yeah. In terms of a separate 20 document over and beyond what you might see in 21 the logbook. I mean again, I'm --22 MR. GRIFFON: And they probably wouldn't have 23 any field in the -- in the database with like a 24 flag indicating, you know, bad crystal or 25 whatever. Not in the earlier time period

1

probably.

DR. ULSH: I don't know. Jim or Roger, do you have some insights on that? Oh, okay. Okay. Hold on a minute. I've just found the samples that I guess Kathy provided in the last set of comments. And what I'm looking at are the -- a few pages from a logbook and they do show, let's see -- I see the -- I see the ac-- I see the activity date; I see the gamma and the penetrating. There's a column for that where it has at least for some of them there's numbers there. Same for neutrons and then there's penetrating skin and beta they put some numbers. And then there's a justification column. And I assume that that talks about why those doses were assigned. I mean it doesn't -- I'll grant you it doesn't go into much detail but I guess the point I'm trying to make is I don't know that we could expect to find much more than what's in the logbooks. Jim or Roger, if I'm off-base here, please jump in and correct me but --MR. LANGSTED: I believe you're correct.

23

24

MS. ROBERTSON-DEMERS: Well, I guess the

25

contention by the petitioners is that when you

1 get into these issues, zeros are being 2 recorded. Now, obviously there are doses other 3 than zero in the logbook. 4 DR. ULSH: Yeah, but there are some zeros, too. 5 MS. ROBERTSON-DEMERS: But --6 MR. LANGSTED: Well, now, one thing that would 7 be possible --8 MS. ROBERTSON-DEMERS: -- It comes down to the 9 question where did these people work and does 10 that make sense? 11 DR. ULSH: Okay. 12 MS. ROBERTSON-DEMERS: Another -- Another 13 thing is if you can find these earlier 14 investigation reports it might give you some indication of if there is a record out there. 15 Maybe it's in the field. 16 17 DR. ULSH: Okay. Here's what I would propose 18 maybe. And Kathy, you've given us some ID 19 numbers here. Maybe we can chase those back to 20 individuals. We can -- if we can then we can 21 certainly look at what's been assigned in the 22 dosimetry file. We could also maybe for a 23 limited number go to the DOE records people and 24 pull the dosimetry files for those people, and 25 we could tell you what's in there. I don't

1	know what we'll find. It sounds like
2	MR. GRIFFON: It may be it may be
3	inconclusive, let's put it that way.
4	DR. ULSH: Yeah, exactly.
5	MR. GRIFFON: Right.
6	DR. ULSH: Yeah. But we won't know until we
7	look at the dosimetry files.
8	MR. GRIFFON: Right. Right.
9	DR. ULSH: So I mean that's certainly something
10	we can do I would think.
11	MS. ROBERTSON-DEMERS: And And
12	DR. ULSH: Okay.
13	MS. ROBERTSON-DEMERS: that's kind of what I
14	wanted you to do but I really think that you
15	need to pull the processing log for the time
16	period prior to 1983 and and look at the
17	frequency of of the loss of crystals that
18	are that's talked about in the petition.
19	MR. GRIFFON: So you're talking about in the
20	earlier time period with the
21	MS. ROBERTSON-DEMERS: The The earlier TLD.
22	MR. GRIFFON: The Harshaw badges. The Harshaw
23	TLD's. Is that the
24	MS. ROBERTSON-DEMERS: And how that was
25	handled.

21

22

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DR. ULSH: Again, I -- without seeing the logbooks I'm getting on dangerous ground because I don't want to speculate too much about what we're going to find but if -- if -to the extent that the logbooks give a complete So for instance I'll say for a record. particular quarter in, I don't know, 1983, if we can find the logbooks for -- that would re-represent all of the badge reads for that particular quarter then I guess you could go through and count how many have problems and how many don't. I'm not sure how big an effort that's going to be because I mean there are -certainly there's thousands of employees at the site. I guess what I'm saying is we could get

MS. ROBERTSON-DEMERS: Just we -- we just want a general -- general feel.

DR. ULSH: Yeah, but I -- I guess what I'm saying is without seeing the logbooks I'm not sure how they're going to be listed in terms of are all the problems going to be listed on -- in one logbook on a couple of pages or is it going to be sprinkled throughout? If it's sprinkled throughout then we might be able to

1 take a representative sampling but if it's --2 all the problems are listed, you know, in one 3 place, like they saved all those for last or 4 something, then in order to get a feel for what 5 proportion of -- of the logbooks represent or what proportion of the measurements had 6 7 problems, you know, how frequently badges were 8 lost or were -- crystals were lost or whatever 9 then we'd have to look at the entirety of the 10 logbooks for that quarter. I -- I just don't 11 know without looking at the logbooks. 12 MR. GRIFFON: You really need to get the 13 logbooks. 14 DR. ULSH: Yeah. 15 MR. GRIFFON: Yeah. 16 MS. ROBERTSON-DEMERS: The --The --The 17 dosimeter processing logs that I recommended in 18 table 4 give you some years for the Harshaw TLD 19 so you can use it as a dual purpose. 20 MR. GRIFFON: Maybe that's the action at this 21 point is to look at those logbooks and evaluate 22 the possibility of -- of following up, Brant. 23 DR. ULSH: Yeah, I think that's --24 MR. GRIFFON: I think that's as far as you --25 you really can take it. I mean you don't want

1	to
2	DR. ULSH: Exactly.
3	MR. GRIFFON: Yeah. Okay.
4	MS. ROBERTSON-DEMERS: Okay. Now, with respect
5	to the field logbook like the RCT logbook and
6	the shift supervisor logbook, what I've been
7	told is that there is documented dose
8	information in those logbooks for people and
9	that that does not match the dosimetry record.
10	Neither does the survey do that.
11	DR. ULSH: Okay, Kathy. I'm a little unclear
12	on this. Were these logbooks that were taken
13	while the jobs were actually were were
14	recorded while the jobs were actually being
15	performed?
16	MS. ROBERTSON-DEMERS: Tony, are you still on
17	the phone?
18	MR. DEMAIORI: Yep.
19	MS. ROBERTSON-DEMERS: Well, my understanding
20	was yes; am I correct?
21	MR. DEMAIORI: Yeah, daily logs. The CC logs
22	was negotiated in the collective bargaining
23	agreement.
24	DR. ULSH: Okay. So I guess it's it's not
25	clear to me how those could contain dosimetry

1	results because the dosimetry badges wouldn't
2	have been processed yet unless you're talking
3	about
4	MS. ROBERTSON-DEMERS: Well
5	DR. ULSH: I don't know.
6	MR. GRIFFON: Unless there were secondary
7	MR. DEMAIORI: CC logs you would have all your
8	high RAD areas as they were discovered,
9	contamination incidents, contaminated
10	individuals. All that would be in the CC logs.
11	That's in the dosimetry logs for like the
12	EPD's, those came out of the RWP offices.
13	Those logs would have all the EPD information
14	that you can cross-reference to your actual
15	TLD's. Those also were daily logs by the job.
16	MS. ROBERTSON-DEMERS: In other words, does the
17	field data show indication that this person
18	should have gotten more than zero or do they
19	corroborate each other?
20	DR. ULSH: Okay. So you're saying then, Kathy,
21	that it's not dosimetry data that's in these
22	logs?
23	MS. ROBERTSON-DEMERS: I, you know I was
24	just told as dose for people.
25	DR. ULSH: But vou didn't get a a a

1	feeling for how that was measured?
2	MS. ROBERTSON-DEMERS: No. But there were
3	there was special dosimetry assigned by job and
4	
5	MR. GRIFFON: Well, maybe it is secondary
6	dosimetry or maybe it's exposure rate measures
7	and sta You know, I don't know without
8	seeing I guess.
9	MR. DEMAIORI: Well, a bit of everything you
10	just said.
11	MR. GRIFFON: Yeah.
12	MS. MUNN: Would Would Wouldn't
13	contamination control logbooks be specifically
14	the area surveyed? Wouldn't that be what they
15	contain?
16	MR. DEMAIORI: Contam the contamination
17	control log logbook would be all your ab
18	abnormalities.
19	MS. MUNN: Yeah, area area of readings,
20	right?
21	MR. DEMAIORI: Oh, yeah, that's, you know, if
22	there was a high dose area that was discovered
23	during a routine survey that would be reported
24	in the CC logbook.
25	MS. MUNN: Yeah. Yeah. and RBP would survey

1 that and record it, right? 2 MR. DEMAIORI: Absolutely if there's 3 contamination incident; if, you know, you lost 4 a room that would be recorded there. It would 5 say how many people were involved. MS. MUNN: Yeah. 6 7 MR. GRIFFON: So it would have some -- some 8 more than just the survey data maybe. 9 DR. ULSH: I guess, Mark, maybe we're at the 10 same follow-up item. We can --11 MR. GRIFFON: I think so. You got to pull some 12 logs at least to see what kind of information -13 14 MS. ROBERTSON-DEMERS: Yeah. 15 DR. ULSH: Well, we can evaluate the 16 plausibility of doing that. 17 MR. GRIFFON: Right. 18 DR. ULSH: And we can try to get 'em. 19 MR. GRIFFON: Yeah. 20 MS. ROBERTSON-DEMERS: And what -- what I was 21 trying to do there is this is what I was told 22 they were called, okay? 23 MR. GRIFFON: Right. 24 MS. ROBERTSON-DEMERS: Now, through time they 25 were probably called something else but this is

Do

1 the type of record that you're looking for. 2 MR. GRIFFON: Okay. 3 MS. ROBERTSON-DEMERS: I just had the 4 opportunity to look at a -- a logbook for a 5 similar area at LANL, and what I noticed is 6 that when they went in to do a job, a 7 particular job, the individuals involved in 8 that job were listed so there was some linkage 9 to names. And Tony, I would assume that yours 10 are similar? 11 MR. DEMAIORI: Yeah, we -- we had a bunch of 12 different logbooks on the floor, not just a 13 contamination control logbook. During 14 processing days we had the processing logbooks 15 and most of those will be classified. Also the 16 shift manager kept logbooks. If it was 17 (unintelligible) breathing air job we kept 18 separate logbooks with dosimeter readings, 19 (unintelligible), that sort of thing in those. 20 And our RWP desk did all the issuing of the 21 dosimeters, DPD's, the pencil dosimeters, 22 whatever. They kept a day-to-day log of all 23 those records of penetrating. 24 MR. GRIFFON: Kathy, do you have -- the last

item says SC&A to conduct inter-comparison.

25

1	you have any logbooks currently, any of these
2	logbooks currently?
3	MS. ROBERTSON-DEMERS: No.
4	MR. GRIFFON: You haven't received any of these
5	yet? Okay.
6	MS. ROBERTSON-DEMERS: No.
7	MR. GRIFFON: But your your notion here is
8	to have NIOSH do an inter-comparison or or
9	for SC&A to do it or for parallel? What
10	What What's I'm unclear on that I
11	guess.
12	MS. ROBERTSON-DEMERS: Well, the first thing I
13	need to do and I didn't put it in this document
14	for obvious reasons is to provide NIOSH with
15	the names that go along with these logbooks.
16	MR. GRIFFON: Right.
17	MS. ROBERTSON-DEMERS: I I think that we
18	kind of wanted to do it independently and
19	compare results.
20	MR. GRIFFON: Okay. I'm just I'm trying to
21	think through the logistics of how that would
22	work. I mean would you both take the same a
23	copy of the same logbook and and go back?
24	How How do you envision that working?
25	MS. ROBERTSON-DEMERS: We We could very

1 well do that. 2 MR. GRIFFON: Okay. 3 MS. ROBERTSON-DEMERS: And I would really be 4 happy to --5 MR. GRIFFON: I think -- I think, Brant, we're on that, too, is -- is the first step is to see 6 7 if we can find these logs or -- or logs that 8 generally fit this title or these types of 9 titles and, you know, maybe bring them back to 10 the work group or subcommittee or wherever 11 we're at and -- and talk about the plausibility 12 of doing such a, you know --13 DR. ULSH: I mean I think -- I think where 14 we're going to be, Mark, is if -- I mean this 15 is common. We've -- We've been here with 16 other SEC petitions. It's a question that I 17 think I heard Wanda say this morning on -- on 18 Y-12, it's a question of how much is enough. 19 And I -- I really don't have an answer for 20 that. 21 MR. GRIFFON: Well, and I --22 DR. ULSH: Certainly the things that we're 23 talking about here are going to take some time 24 and we're willing to do that, given enough 25 time.

1 MR. GRIFFON: No, I think the other -- the 2 other factor here, and this is why I said one -3 - another reason I said this this morning was 4 how much is enough may vary from petition to 5 petition. I mean the -- the petitioner in this 6 case made -- made, you know, several specific 7 allegations within the petition so I think to 8 the extent we can we need to follow up on the, 9 you know --10 DR. ULSH: Right. 11 MR. GRIFFON: So it's just -- more than just a 12 general review but also address their specific 13 allegations. 14 DR. ULSH: No, I understand. 15 MR. GRIFFON: Yeah. 16 DR. ULSH: And -- And really I'm not trying to 17 I know, I know. 18 MR. GRIFFON: 19 DR. ULSH: -- influence what the Board decides. 20 I mean that's really not my place to do that 21 but we -- I guess I also have to make it clear 22 that NIOSH is under a statutory obligation to 23 issue the evaluation report, and to do that we 24 had to operate on the data that we had on the 25 table at the time. And certainly we recognize

and three months from now there will still be more records out there and, you know, I mean we have to -- we're put in a position -- NIOSH is put in a position where we have to issue the evaluation report and make a recommendation.

And I -- I'm still comfortable, you know, with the report that we've issued. But again, I mean if the Board decides that you would like us to take more time and, you know, delay things a bit we'll certainly do that. We'll do whatever we can to support it.

DR. WADE: The rule -- This is Lew. The rule allows for that. I mean NIOSH will present the evaluation report and then the Board can, you know, ask for additional information or, you know -- you know, let its desires be known. But we'll deal with that next week. I mean --MR. GRIFFON: Yeah, yeah, I think -- exactly. MS. ROBERTSON-DEMERS: Well, what I can try to help you do is to get to the right logbooks because there was a -- there's -- during my trip there was an interchange between the records people and myself on what might be the right logbooks. And we could have them pull a

1 couple and copy sample pages and see if that's 2 really the logbook we're looking for. 3 MR. GRIFFON: So that's something you can work 4 with Brant offline on. 5 MS. ROBERTSON-DEMERS: MR. GRIFFON: That -- That'd be great. 6 7 DR. ULSH: Yeah, anything that you can do to 8 narrow the search, that would -- that would 9 only help things. 10 MR. GRIFFON: Right. And I would say, John, 11 you know, this -- this -- this rule always 12 applies in between meetings that, you know, if you guys need to have offline conversations to 13 14 expedite this process, you know, as long, you 15 know -- if it's noteworthy I guess keep minutes 16 but, you know, I think that's fine and -- and 17 encourage that at this point. 18 DR. WADE: Right. Common sense. 19 MR. GRIFFON: Yes. Is there anything else on 20 the -- on this report, Kathy or Joe? 21 MS. ROBERTSON-DEMERS: No. That's --22 pretty much it. 23 MR. GRIFFON: Okay. 24 DR. WADE: Okay. Well --25 MR. GRIFFON: I think -- I think where --

1 where we stand, Lew, is, you know, we've got an 2 update on the matrix. I will update the 3 matrix, Brant, and to include these things as 4 well. 5 DR. ULSH: Yeah, I heard you typing, Mark, so -6 MR. GRIFFON: Yeah. And -- And it may not be 7 8 as quick as the last turnaround but I'll try. 9 But the other thing is I think we need to at 10 least give a status report at the Advisory 11 Board meeting and maybe a plan forward because 12 we also need SC&A to -- to review the 13 evaluation report. But I think, you know, with two days left before the meeting, two working 14 15 days or whatever, I know we're all going to be 16 working on the weekend but, you know, the focus 17 on most folks is going to be Y-12 to -- to finalize that -- I -- I would definitely 18 19 prioritize that for SC&A if I had -- I mean 20 not that I'm the -- the task -- prioritizing 21 your work but I think that -- that seems to be 22 a priority at this point. 23 MS. MUNN: I think that's appropriate. 24 MR. GRIFFON: Yeah. Try -- Let's try to -- to 25 -- to fine tune that one and -- and -- and then

1	we'll we'll give a status report on Rocky
2	and go forward and the Board can advise on what
3	direction we need to go with, you know, with
4	Rocky.
5	DR. WADE: Sounds like a plan.
6	MR. GRIFFON: All right.
7	DR. WADE: Well, you you're all to be
8	complimented and
9	MR. GRIFFON: Another long day.
10	DR. WADE: A long day but a productive day.
11	Mark, I'll I'll give you a call
12	MR. GRIFFON: (Unintelligible) if other work
13	groups go shorter than (inaudible)
14	DR. WADE: We'll talk I'll give you a call,
15	Mark, tomorrow and we can talk specifically
16	about next week and the organization but thank
17	you, and thank everyone who participated.
18	Thank the petitioner, Tony, we appreciate your
19	forbearance. And, you know, we'll be seeing
20	you all in beautiful Colorado next Tuesday.
21	MR. GRIFFON: Thanks, everyone, for the hard
22	work, too. I know it's these are crunching
23	weeks.
24	DR. WADE: Thank you all.
25	MR. GRIFFON: Take care.

(Whereupon, the working group meeting was adjourned at 5:35 p.m.)

CERTIFICATE OF COURT REPORTER

STATE OF GEORGIA COUNTY OF FULTON

I, Steven Ray Green, Certified Merit Court Reporter, do hereby certify that I reported the above and foregoing on the day of April 20, 2006; and it is a true and accurate transcript of the testimony captioned herein.

I further certify that I am neither kin nor counsel to any of the parties herein, nor have any interest in the cause named herein.

WITNESS my hand and official seal this the 12th day of June, 2006.

STEVEN RAY GREEN, CCR

CERTIFIED MERIT COURT REPORTER

CERTIFICATE NUMBER: A-2102