# 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

WORKING GROUP MEETING

ADVISORY BOARD ON

RADIATION AND WORKER HEALTH

# ROCKY FLATS

The verbatim transcript of the Working

Group Meeting of the Advisory Board on Radiation and

Worker Health held in Hebron, Kentucky on January

26, 2007.

# <u>C O N T E N T S</u> January 26, 2007

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#### TRANSCRIPT LEGEND

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#### PROCEEDINGS

1 (10:00 a.m.)

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## WELCOME AND OPENING COMMENTS

## DR. LEWIS WADE, DFO

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DR. WADE: (by Telephone): Ray, are you up and running?

COURT REPORTER: Yes, sir, ready.

DR. WADE: (by Telephone): What this is is a meeting of the work group on Rocky Flats site profile and SEC petition activities that functions under the Advisory Board on Radiation and Worker Health. The group is most ably chaired by Mark Griffon, members Gibson, Presley and Munn. Also, the materials we've been providing, the principal NIOSH contact is identified as Brant Ulsh, and the principal SC&A contact identified as Joe Fitzgerald.

Just for a little bit of background, the full Board at its last meeting decided that it stated its intentions to take up the Rocky Flats' petition at its meeting in May, May 2, 3 and 4, later this year. It did that

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because of input it has had from petitioners and members of Congress and others as to the need to do a complete job in terms of the science issues that were remaining. So that's the target for the Board, taking up the petition in earnest May, 2, 3 and 4. Our intention would be to do that in Denver, Colorado.

Again, this is Lew Wade. I have the

Again, this is Lew Wade. I have the privilege of serving as the Designated Federal Official for the Advisory Board. I apologize for not being there. This is the first meeting of any type that I've not been able to be there face to face. So I'll do my introductions, and I'll be on the phone with you until noon.

And then ably Chia-Chia Chang is there, and Emily Howell is there, and they'll serve the role as Designated Federal Official as well as representing the Office of General Counsel. To them I'll say I'll have my cell with me and would be available on cell if you were to need me at any time during the course of the meeting.

I first would like to establish what

1 Board members are involved and present on the 2 call. Mark, I assume you're there. 3 heard Wanda. I've heard Robert. Is Mike 4 Gibson there as well? 5 MR. GIBSON (by Telephone): I'm via phone 6 call. 7 DR. WADE (by Telephone): Okay, are there 8 any other Board members that are either in the 9 room that I didn't identify or on the phone? 10 (no response) 11 DR. WADE (by Telephone): So my hearing is 12 that we have Mark, Wanda, Robert and Mike 13 participating. We don't have a quorum, and 14 therefore, we can proceed. 15 One quick news update, as many of you 16 know, two new Board members have been 17 identified, and it's looking very likely that 18 they would be seated at the table for our 19 February meeting. It's not completely done yet, but it's certainly looking that way. And 20 21 that's good news for all of us, particularly 22 the overworked Board members who they'll be 23 joining. 24 One quick thing on phone etiquette, if 25 you are participating by phone, please mute

when you're not talking. If you are talking use a handset. Realize that background noises of all types enter into the room and can distract the participants. And also, if you have a system where when you go on hold or go away for awhile and background music plays, please be aware of that and don't do that to the working group. It happens more frequently than we would like.

One other caution is with regard to the discussion of Privacy Act related information. This working group and those that support it have done a wonderful job of getting their documents in and cleared, and I think we have cleared documents available for anybody who would want them. But since this work group has in some of its deliberations got to look at individual records, I would just caution all of us in our discussions to be sure that we don't get into anything that would contain personal identifiers or allow an individual to be identified. I think the documents are good. Just exercise a little discipline in your verbal comments as well, realizing that protecting everyone's privacy

1	is terribly important to all of us.
2	I guess with that I would suggest that
3	we go around the table and do our
4	introductions. Please for ORAU/NIOSH team
5	members and SC&A team members, when you make
6	your introductions, please identify any
7	conflicts you have with regard to this site.
8	So I would start in the room from Mark's
9	right.
10	MS. MUNN: This is Wanda Munn. I have no
11	conflicts with Rocky Flats.
12	MR. DeMAIORI: Tony DeMaiori, petitioner.
13	DR. NETON: This is Jim Neton, NIOSH, I have
14	no conflict at Rocky Flats.
15	MR. PRESLEY: Robert Presley, Board member,
16	I have no conflicts with Rocky.
17	MR. SHARFI: Mutty Sharfi, ORAU team, no
18	conflicts Rocky Flats.
19	MS. CHANG: Chia-Chia Chang, NIOSH, no
20	conflicts.
21	MR. FITZGERALD: Joe Fitzgerald, SC&A, no
22	conflict.
23	<b>DR. MAKHIJANI:</b> Arjun Makhijani, SC&A, no
24	conflicts.
25	MS. JESSEN: Karin Jessen, ORAU team, no

1	personal conflicts.
2	MS. HOFF: Jennifer Hoff, ORAU team, no
3	personal conflicts.
4	DR. ULSH: Brant Ulsh with NIOSH, no
5	conflicts.
6	MR. LITTLE: Craig Little, ORAU team, no
7	conflicts.
8	MR. MEYER: Bob Meyer, ORAU team, no
9	conflicts.
10	MR. RICH: Bryce Rich, O-R-A-U team, have
11	two year administrative oversight conflicts.
12	MR. ELLIOTT: Hello, this is Larry Elliott,
13	no conflicts.
14	MS. HOWELL: Emily Howell, HHS, no
15	conflicts.
16	MR. GRIFFON: And Mark Griffon with the
17	Advisory Board, no conflicts.
18	And Lew, we'll go back to the phone
19	people.
20	DR. WADE (by Telephone): If there's any
21	NIOSH/ORAU team or SC&A team members on the
22	line, please identify yourself.
23	MR. SMITH (by Telephone): Yeah, this is
24	Matthew Smith, ORAU team, no conflicts.
25	MS. BRACKETT (by Telephone): Liz Brackett,

1	O-R-A-U team, no conflicts.
2	DR. WADE (by Telephone): SC&A?
3	DR. MAURO (by Telephone): John Mauro, SC&A,
4	no conflicts.
5	MR. BUCHANAN (by Telephone): This is Ron
6	Buchanan, SC&A, no conflicts.
7	MS. LOPEZ (by Telephone): Teresa Lopez,
8	ORAU team, no conflicts.
9	DR. WADE (by Telephone): Any other team
10	members, SC&A, ORAU/NIOSH?
11	MR. LaBONE (by Telephone): I'm sorry, this
12	is Tom LaBone, O-R-A-U team, no conflicts.
13	MS. DeMERS (by Telephone): This is Kathy
14	Robertson-DeMers, SC&A, no conflicts.
15	MS. THOMAS (by Telephone): This is Elyse
16	Thomas, O-R-A-U team, no personal conflicts.
17	DR. WADE (by Telephone): Anyone else?
18	MR. GIBSON (by Telephone): Lew, this is
19	Mike down at the complex.
20	DR. WADE (by Telephone): Welcome.
21	Any other federal employees who are on
22	the line by virtue of their federal
23	employment?
24	MS. BOLLER (by Telephone): This is Carolyn
25	Boller with Congressman Mark Udall's office.

1	DR. WADE (by Telephone): Welcome.
2	MS. ALBERG (by Telephone): And Jeanette
3	Alberg with Senator Allard's office.
4	DR. WADE (by Telephone): Welcome.
5	MS. ESCOBAR: And this is Felicia Escobar
6	with Senator Ken Salazar's office.
7	DR. WADE (by Telephone): Welcome all.
8	MS. HOMOKI-TITUS (by Telephone): Liz
9	Homoki-Titus with Health and Human Services,
10	and I have no conflicts.
11	DR. WADE (by Telephone): Jeff?
12	MR. KOTSCH (by Telephone): Jeff Kotsch with
13	Department of Labor.
14	MR. BROEHM (by Telephone): Jason Broehm,
15	CDC Washington Office, no conflicts.
16	DR. WADE (by Telephone): Anyone else who
17	would like to be identified, petitioners,
18	representatives, workers at the site? Anyone
19	else who would like to be identified for the
20	record?
21	MS. BARRIE: Terry Barrie with ANWAG.
22	DR. WADE (by Telephone): Thank you for
23	being with us, Terry.
24	MS. BARRIE: Kay Barker will be on the line
25	later.

DR. WADE (by Telephone): She's more than welcome.

Okay, Mark, I think it's all yours.

MR. GRIFFON: Before we start I think Emily wanted to take the floor for a second.

MS. HOWELL: I just wanted to kind of reiterate what Lew was saying a minute ago about Privacy Act protected information. I think that we work really hard to try and review all of the documents for these meetings, but this working group has in its quest to really delve into information has gone to the level of looking at individual files.

And I think that there's some older versions of the matrix that are out there floating around that may include some protected information. As members of the working group or our contractors you're able to have those.

But please just be aware that you may have a version if it was, if you printed it or received it prior to yesterday afternoon then the version you have probably does have protected information in it, and I think that

Brant has some redacted matrices if you would like a new copy. So please just be aware of that if you go to speak off of the matrix.

If there's a name in it, and it's not someone who's sitting around this table as an OCAS, ORAU or SC&A employee, you probably should not be saying that name. And if you ever have any questions, you're more than, I'm more than willing to take any questions at any time.

MR. GRIFFON: I think we've been, I mean I think we can work with that with the work group here today. The only person I have, I think the prior matrices have been made public in the Board meetings so they're out there, not just internally circulated within the work group.

MS. HOWELL: Right, and like I said, this is kind of a continuing area of concern, and we're trying to work on this, but there may be other matrices that include information that we would have liked redacted that are still out in the public so we're just trying to get better from here on out.

DR. MAKHIJANI: Could I ask a question?

MS. HOWELL: And if you do have copies of the old matrices that have the Privacy Act protected information in them or if you're not sure, if you could discard them and do so in a careful manner, that would be helpful.

#### Arjun?

DR. MAKHIJANI: Yeah, could I ask a question about that? Now there is one individual I know who NIOSH asked for Privacy Act release, and he did sign an appropriate form --

MS. HOWELL: Okay, I would prefer you talk about that off line with me. Without having it in front of me I can't be sure, and any time we get into these discussions around the table, we could be making the problem worse. So we can talk about that during the break.

DR. MAKHIJANI: All right.

MR. GRIFFON: I don't think this in any way affects what we have to discuss here in this meeting, but we should all understand the rules so let's just, we can work from the redacted version of the matrix today.

Actually, I have an abbreviated agenda which we've been using for most of the last two or three work group meetings. And then

I'd like to use that same sort of format, cover the main action items of interest and then go back through this redacted matrix toward the end of this meeting and make sure we've captured everything and sort of update ourselves.

The agenda I'd propose, and partly this was to frontload the items I thought were the most pressing and also to allow some SC&A personnel to be online when they need to be and not necessarily for the whole day and the same goes for some ORAU people maybe.

I'd like to start off with number one, data completeness; number two, the '69 data question; number three is the coworker models, and that's mainly as they pertain to this question of completeness. Number four is other radionuclides, and I think we're down to thorium as our other radionuclide outstanding action here. Number five is log book analysis; six is the safety report analysis. Seven is the Super S item; eight is neutron issues, and nine is D and D worker questions.

So if that's okay with everyone, I'll think we'll stick to that. I tried to, like I

said, frontload some of the more, items I thought were going to take a little longer and were maybe more pressing. Any comments on that; otherwise we'll proceed.

#### COMPLETENESS OF DATA

All right, let's start off with the data completeness. I think where we stand on this is that SC&A provided a report, and did we get the whole report or were there still pieces that were Privacy Act review or at this point we have the whole section of the report?

MR. FITZGERALD: Yeah, the whole section of the report has gone through Privacy Act clearance and was distributed on January 10<sup>th</sup> to the working group. So it's been through all the review.

MR. GRIFFON: Can I ask NIOSH is that distributed to petitioners and other interested parties? It went through Privacy Act review. I thought we were going to try to get those components out.

Lew, do you know if that was made available to the petitioners and other interested --

MS. BARRIE: Mark, this is Terry Barrie and

1	no, I have not received any reports that have
2	been cleared by the Privacy Act.
3	DR. WADE (by Telephone): I guess I would
4	ask, Larry, if you would make a note to see
5	that that's distributed.
6	MR. ELLIOTT: Yes, I will.
7	MR. GRIFFON: I think we agreed that this
8	was okay, right? These draft sections of the
9	
10	DR. WADE (by Telephone): Yes, once they've
11	been cleared, yes.
12	MR. GRIFFON: So I think there's a couple
13	MR. ELLIOTT: I just don't know that I've
14	had approval that they've cleared. Well,
15	we'll check into it.
16	MR. GRIFFON: Once we're sure they've been
17	Privacy Act reviewed and cleared, we'll get
18	those distributed as soon as possible, right?
19	MR. ELLIOTT: Yes, as soon as I hear that
20	they're approved for distribution, we'll put
21	them on the website and we'll share it with
22	the petitioners and interested external
23	parties.
24	MR. GRIFFON: I didn't know that it had to
25	be on the website, but

DR. MAKHIJANI: Yeah, what didn't happen is we sent the materials to CDC to Emily, and we got instructions about what to take out that was covered by the Privacy Act, and we did that. And then put together a version that was completely de-identified as per the instructions. And that's what --

MR. FITZGERALD: That's what on January 10<sup>th</sup>, at least for this particular section, we transmitted to NIOSH and the work group.

That's as far as it went. It went to the work group as well as NIOSH, and with a notation and a cover e-mail that we would assume that NIOSH would handle outside distribution.

MR. GRIFFON: Okay, that's fine. Larry, you can check on making sure that it's been finally approved for distribution. I think the way, part of the reasoning here is that these are draft sections, final draft sections of SC&A's overall report, and we kind of, the notion was to keep the petitioners and other interested parties informed along the way instead of dropping one big report at the end of --

MR. ELLIOTT: I understand. I share the

interest in making that happen. I can't do it until I hear I'm okay to do it.

MR. GRIFFON: With that maybe we'll just start, Joe, if I can, I'll give you the floor to give us an overview of what you've put in this report.

MR. FITZGERALD: On this particular issue we caught in shorthand the completeness issue but again, we're going back to an old issue in a sense because when we reviewed the Rocky Flats' site profile, we did spend quite a bit of time focusing on concerns, certainly concerns expressed by the former workers that there were unexplained zeros, perhaps gaps in records, notations, such as no data available.

So we certainly raised those questions in the site profile and certainly the petitioners have raised similar issues in the petition. So certainly going into this evaluation we wanted to focus on that from a number of different vantage points. And we spent a great deal of time on the data reliability side looking at whether or not we could corroborate that question by virtue of the log book reviews that we've done, and

we'll talk about that later, evaluating quite a few safety concerns that were available and also looking at certain, the affidavits and certain data integrity issues.

On this particular issue though we were also concerned about looking at the database as a whole. Looking at whether or not one could look at the database and ascertain how complete in fact it is. We looked at HIS-20, which is the electronic database, first. And in our comparative reviews we certainly began picking up some issues, some discrepancies.

It took some time, I think, to get into a level of understanding as to what we were looking at. But over the course, I think, of middle and last, late last year we certainly established that we had a number of issues regarding the completeness of the electronic database, HIS-20.

Certainly, at that stage we turned to the, what we would call, the claimant file, the raw data itself that the HIS-20 is based on, certainly wanting to know how complete that is in fact because that would be, in

fact, the original data that would be the baseline for doing dose estimations.

We did a sampling analysis initially which was basically just doing a limited number of samples, in this case a dozen, just to get some sense of what we were looking at. That sort of reconnaissance survey demonstrated a number of gaps that we were, I think, taken aback by. We didn't expect to see in that kind of a limited sample the kinds of gaps that we did see.

That led to our going to this working group, and certainly looking for their guidance which they suggested that we expand that sampling to include what ended up being 40 more cases from the original raw data file. So this is a total now of 52 cases that we were sampling out of the original claimant file. And that sampling plan was designed to be, given the sample size, as representative as we could be.

And so in any case the analysis that we presented right after New Year's is in essence our report of what that sampling of the 52 cases showed us. And quite frankly,

and I'm going to turn to Arjun in a second to walk through the details, the bottom line, it corroborated, I think, our initial concerns on that smaller sample that there are, in fact, are significant gaps in the claimant file upon which your coworker models and other estimations would be based.

whether or not this database is going to be sufficiently complete to do coworker modeling and to do the necessarily gap filling that one has to do when you have unmonitored workers. So with sort of a prelude and background I think what we want to do is really walk this work group and this group carefully through this report. I think this report's a very, very critical report on this whole question of completeness. And I think coupled with what we did on data reliability, I think does tackle what I think is the cornerstone of this review.

MR. GRIFFON: Excuse me. If you're on the line, could you mute your phone? We're hearing some people talking so you might want to mute your phones, please.

DR. MAKHIJANI: Thanks, Joe. Basically the frame work of the review was provided by the Board's criteria on the pedigree of the data and the methodology and internal consistency, and also the Board's approval of our interim SEC review procedures which require us to review completeness of data.

As Joe said, we reviewed two different kinds of samples from the data. One was a set of 32 developed over time which were random samples divided into two periods, the early period when there wasn't universal monitoring which goes up to '63, and then from '64 when the ID badge was an integrated badge with the dosimeter and the identification in a single badge.

Now we understood later on that subcontractor workers were not covered by the universal monitoring policy, but the jump in the percentage of monitored people in '64, percentage of people wearing badges, that indicates that the percentage of people badged were in the 90 percents except for 1969 through about 1990. So this analysis goes up to 1992.

Nineteen ninety-two is a little bit anomalous here because that production stopped early in that year, so they went through the transition year to go into a decommissioning mode which was formally declared in 1993. So I'll walk you through our findings.

We divided this into external dose and internal dose findings, and we also just as a caveat on what kind of completeness review this is. This wasn't a detailed completeness review where the whole record was examined for every gap. This was a broad screen review where we said the data had a gap, when there was no record for that year at all. If there was one zero reported, for instance, we did not count that as a gap.

So it doesn't mean that the record is complete or there's a record for every badge cycle. So it's a very broad screen that allowed us to go relatively efficiently to determine whether the gaps were large or not. And the use of the broad screen means, especially for external dose, that this is a minimal estimate of the completeness of the data.

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So in the first table where we did the external dose analysis we found that for the early period we defined two kinds of data gaps. One was the proportion of workers who had a gap for one year or more. And then we defined a cumulative gap which is if you add up all the years for all 32 workers that were in each period and how many percent of those years were not monitored cumulatively.

So just to run you through the numbers for external dose in the '51 through '63 period. There were 14 workers in the sample and 29 percent of them had gaps of one year or more. And the cumulative gap was 21 percent, that is, of all, we multiplied the 14 workers by the 13 years. Twenty-one percent of those years were blanks.

Then '64 to '92 there were 30 workers in the sample, and the percentage of workers with gaps of one year or more were 33 percent. Now there's a statistical artifact in that by the way we defined the period because of the 1992 transition year. Of the ten workers who had a gap of one year or more, four workers were in that, had a gap only for 1992. So if

you exclude 1992, and we included it because that was the instruction of the working group and that's how it was decided here, but if you exclude 1992, the gap goes down to about 20 percent.

And the cumulative gap for 30 workers multiplied by the number of years, '64 to '92, 29 years, is ten percent. So gap is smaller for the second period which is what we would expect since there was this quasi-universal badging.

Now in these numbers we did not include 1969 which is separate. Nineteen sixty-nine had partial or full gaps and so we have to, we just analyzed that separately, and it's a separate action item on the working group. And so these numbers don't reflect 1969. And it's also important to remember that these are gaps in the record, not zeros. So zero entries are counted as positive indications of recorded data.

Then we looked at internal dose and the same data for the same set of workers, and for the '51 to '63 period number of workers with a gap of one year or more was about the

same, 29 percent. Now for internal dose we defined the gap when there was no measurement of any kind, either urinalysis or fecal samples or an in vivo count. So if there was one count of any of these types we counted as a year that had a measurement, and it was not a gap. So no internal dose measurement for the full year of any kind counts as a gap. So again, it's a very broad screening.

The cumulative gap for the early period was 12 percent of all the workers, 14 workers, multiplied by 13 years. Twelve percent of those years were unmonitored cumulatively.

One surprising finding was in the second period we found that 73 percent of the workers had a gap of one year or more, and the cumulative gap was 33 percent. So this, of course, raises the question of how the gaps are going to be filled. The reason to look at the highly exposed workers was to examine the question of whether the gaps can be filled and what questions it raises for coworker models.

The highly exposed workers, quote/unquote, were defined by Rocky Flats

review, retrospective review that was undertaken in the 1990s. They had a set of, I don't remember how many workers they reviewed, but they had a set of workers that they reviewed for cumulative exposure where the internal and external was added up. And then they classified workers from one to four into exposure categories.

And category four was the most exposed cumulative dose. Category three was the less below that, and category one had the least cumulative exposure. Now this doesn't separate internal and external dose, and it doesn't separate periods. So if somebody who had a high cumulative may not have been in the highest exposed category at some period.

But it was an approximate guide to examining whether there were data available for filling in the gaps. It's not definitive but indicative. What we found was, we again did separation of external and internal dose. We did not find any full year gaps in the internal dose for the highly exposed workers. We had ten cases from category four and ten cases from category three. And all of them

1 had at least one measurement.

Now it doesn't say it's complete in all respects but there was in this broad screening no gap in the internal dose measurements. Now we don't include a screen as to whether every relevant radionuclide was monitored and so on. But every one of these 20 workers had at least one measurement during each year.

For external dose we did find some gaps -- there's no table here so I'm trying to find the number. Excuse me. But the most significant gap we found was in group three workers, that is, in the workers just below the highest exposed workers in the '51 to '59 period. So it turned out in the early period there were external dose gaps.

And if you looked at a part of that early period, '51 to '59, the gaps were particularly large. Sixty-two percent of the cumulative employment years in group three workers were missing -- I shouldn't say missing -- had blanks in their external dose records which indicate that maybe they were not badged in that period.

Now this was a period of partial badging of workers. In the earliest period there were the fewest workers badged and that proportion of badging went up as the '50s went on. And then in the '60s we did not find external dose gaps.

We looked at the job descriptions of the workers, the job cards of the workers who had these external dose gaps and the group four workers in order to determine whether there was any pattern in their job assignments for why they may have been badged or not badged.

Most of the full year gaps were associated with work in Production Plant B where depleted uranium and enriched uranium was processed. So as NIOSH has said that the uranium areas were the areas that were thought to have low exposures. So there was a systematic decision, at least the sampling indicates that there was a systematic decision that workers in plutonium areas would be badged and workers in uranium areas were. I don't know if they were all done that, but the gaps are focused in those areas.

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 2
 3

There were some gaps in Building 81, the QC Lab, something called the Pipe Shop.

There were no full year gaps for people who worked in Production Plant C which were the plutonium workers. So that's important to say. So there's a clear division in how these gaps emerged.

Now we looked at documents indicating exposure potential in the uranium areas and in a particular uranium area, the foundry operations, the foundry operations result — let me go back. Uranium as a homogeneous material builds up its first decay products to equilibrium relatively rapidly, thorium—234 and protactinium—234. And its external contact dose, shallow dose and deep dose, are relatively well—known to be maximum two millirad for hour for deep dose and about 240 for shallow dose.

But if, in foundry operations, there's a separation of the decay products, and the shallow beta dose rates can be pretty high, this is documented in an evaluation of the uranium areas. The depleted uranium castings where in the early years there were shallow

dose rates as high as 2,000 to 3,000 millirads per hour, so in order of magnitude bigger
than the homogeneous uranium metal contact
dose that were documented.

These were thought to be very high or extremely high and the problem persisted though apparently in lesser degree through 1982 when the report was written. I think either this report or the 1969 report actually contains the extended quotation from the year 1982 report documenting that.

Now we have external dose data for the plutonium areas, and the question arises plutonium also itself does not have, plutonium-239, does not have a strong shallow dose component or deep dose component. There is americium-241 in the plutonium areas that provides the main shallow dose component, I mean deep dose component.

And the question arises from this analysis is how do the plutonium -- well, there are two questions. How do the external doses in the plutonium areas, especially in the 1950s, correspond to the external doses in the uranium areas especially for shallow dose

and would they be bounding? And that's a question we could not answer from our analysis.

And then secondly, from among the monitored workers in uranium, was there monitoring from the areas that had the higher exposure potential so you could possibly construct a coworker model? And that also we could not tell from this analysis. So that's the main, other than 1969, which is separately covered, that's the main issue which emerged from the completeness evaluation.

There are two other issues, one I've already mentioned in regard to subcontractors. We did not find any evidence of any systematic avoidance, that subcontractors were sent into radiological areas and without badges, and then they were exposed in some nefarious way. We did not find any evidence that there was a systemic violation of the policy that would have resulted in unbadged subcontractor workers going into radiological areas.

But we thought that some verification of enforcement would be desirable since there were unbadged subcontractor workers on site

apparently throughout the period. And one possible verification that might settle the question which is at the present time sort of an expert statement which is somewhat in opposition to the statement that has been made by the petitioners is that the blanks in the records correspond to a systemic problem. For this particular problem one verification procedure might relate to the clean up of the fires after 1965 or 1969 to see if there were any subcontractor workers that might have been sent into radiological areas but had no badge records.

The other issue that has arisen in the context of working groups discussions was that, on the completeness investigations was that of one particular individual who had, who was not a subcontractor worker, who was a prime contractor employee who had an eleven-year exposure data gap from 1963 to 1973 inclusive, and only one of those years was a year of when universal badging was not in place.

Now this worker is thought not to have a high exposure potential, but ten of those

1 years do relate to a universal badging period. 2 And the question arises, especially when we 3 discuss the 1969 gaps. Maybe you want to 4 defer that question to 1969. What was the 5 policy of badging for prime contractor 6 workers? Or were badges issued and not read 7 from 1964 onward in some way and what happened 8 to those records. So that's kind of an open 9 question that also arose from this 10 investigation. 11 Sorry it took so long to present, but 12 it was pretty complex. 13 MR. GRIFFON: That's okay. I just have one 14 follow-up on the production workers. You 15 said, I heard the external group three workers 16 you found some gaps in the '50s. I think you 17 said no gaps from '64 through '92 for the 18 production workers? 19 DR. MAKHIJANI: Yeah. From, I believe if I 20 remember all the details correctly, from '61 21 onward. 22 MR. GRIFFON: So it was mainly that '52 23 through '59. 24 DR. MAKHIJANI: Yes, '51, yeah. 25 MR. GRIFFON: And I know that at this point

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I guess we don't need to cover '69 yet, but I know at this point NIOSH has this report, and they're reviewing it and going through it. I know that, and talking to Brant I know you're not in a position to give a full response. guess the only question I had was this, you know, just in terms of when we do get your response to this I still have this question of what are we measuring against? And I don't know if you made any headway in resolving this question of, remember the '69 memo where this policy apparently was put in place for not reading some of the badges from people that were wearing badges. But we never really understood if it was one year, if it continued, and I know you said you were going to look into that. I don't know if you've got any update on that, but I think it would be nice for us to know. The better we understand the policies over time, the better we can understand the results here, you know, if they, if there's anything to these results or if they are well explained by policies that were in place at the time.

DR. ULSH: Well, you raise a good point

there, Mark. We've got to keep in mind the overall purpose of this analysis. I think SC&A's report gets us partway there in terms of how many years -- well, no, what time periods do you see where individual workers did not, there's no monitoring results in their record.

But the other part of the question is would you expect there to be records there. And I don't think that we have that answer yet. That's what we are doing. We have, we've had some discussions with SC&A on this. They've interviewed some of the site experts, and what we've suggested was -- and Arjun talked about this -- was that you have to look at the worker's job exposure history card to determine whether or not you would expect them to be monitored. And that's what we're doing in our analysis now.

I can tell you that we are seeing a lot of instances where, for the periods where there is no monitoring, that's exactly what we would expect given the worker's job duties at the time. I'm not in a position to tell you that that is the case in everyone because

we're not done yet. But we are seeing, again, just like in the first 12, that the workers' job duties do explain at least some of these periods when there are no records for them.

And, of course, as you mentioned, there are a couple of questions that this goes to. One is data integrity. I mean, do you have people who should have been monitored according to the badging policies in place at the time, and there's no records, in which case you might conclude that those records are missing. The other question which is number three on your agenda is what effect might that have on the coworker data. And I'll save that discussion for maybe that topic.

Now in terms of the '51 to '59 period, so you've got to keep this separate from the '69 issue as Arjun mentioned, you know, that's handled separately. But '51 to '59 I think it's fair to say, Arjun, that we found, I think your report said that you found gaps in the Plant B workers, Building 81 more so than others, not exclusively, but there were more there.

DR. MAKHIJANI: Yeah, I think you can say

that there were not gaps in Plant B workers for full years, and then the gaps that were there were concentrated in the Plant B area.

DR. ULSH: Right, and this was an issue that when you interviewed Roger that Roger brought up, that people in Plant B, Building 881, were not monitored in the '50s up until I believe it's the fourth quarter of 1960. So we are aware of that. And the badging policy that was in place at the time, I believe, was that people who were expected to get less than ten percent of the tolerance weren't required to be badged. And so that was why those people, that was the thinking about why those people wouldn't have been badged.

Now you have raised the concern in your report, but what about people in the foundry? I mean, they could get some pretty high doses, especially if they're coming into contact with the castings or the sculls from the foundry. I would point out to you, however, that the foundry is not in Plant B. The foundry is in Plant A.

DR. MAKHIJANI: Building 444.

DR. ULSH: Four forty-four.

1 DR. MAKHIJANI: That's why I said the gaps 2 are, the way I characterized the gaps is that 3 the gaps are not in Production Area C. They 4 are in other areas. Right, we caught a lot of 5 gaps in Plant B. DR. ULSH: Now Area C, I think, is the 6 7 plutonium area, right? 8 DR. MAKHIJANI: That's right. 9 DR. ULSH: So you've got, in the early years 10 you've got A, B, C and D. Plant A is Building 11 That's where the foundry was. 12 where you might see some of these high doses, 13 like you said in your report. Plant B where 14 you're seeing a lot more of the gaps is 15 Building 881. It's not the foundry. 16 MR. GRIFFON: Is it your understanding, or 17 Roger's understanding that those foundry 18 workers should have been -- you know, based on 19 the ten percent? DR. ULSH: We're looking at that right now. 20 21 I think, the indications that we have so far 22 are that their doses were low, but, Mark, that 23 may not be my final word on this. We're 24 looking at it right now. 25 MR. PRESLEY: Brant, it's Bob Presley,

1 question. Those early years in that building. 2 Those production years are those years that 3 Rocky was going in there and fabbing the 4 buildings and fabbing the process equipment 5 and things like that. 6 DR. ULSH: Bob, I can tell you Building 881 7 was built in 1952, I believe, maybe '53, 8 something like that. I can't tell you when it 9 actually came on line and became hot. 10 MR. PRESLEY: Then that may be one reason 11 you've got gaps in those things. 12 DR. ULSH: Could be, I'm not in a position to say that that's it. 13 14 MR. PRESLEY: But I mean that's a 15 possibility that those people were not 16 production workers, but they were construction 17 workers, and that building wasn't hot, thus, 18 they weren't monitored. And that was a 19 practice that we did from day one throughout 20 the complex. 21 DR. ULSH: That is true, Bob, and that's why 22 we're looking at the job exposure histories to 23 come at it from that question. So I mean, to 24 get at would you expect there to be monitoring 25 here and we don't have any. That would say to

you that there might be a data integrity issue here. There should be monitoring results. People should have been monitored, and they're not. We're not in a position to offer an opinion on that just yet for all of the workers and say that some of them --

MR. GRIFFON: Again, to me it's, I think it's much more useful to see as best you can, and I know you found some of this pieces of this, but the policies of the time, you know, the written policies, indication that here's how we made this decision. Here's who was badged.

I mean, I think you need to look at both, but you can look at these jobs, and you probably have a lot that are clear-cut one way; a lot that are clear-cut the other way. And then you've got some of these murky ones. But if you have a policy then we have a brighter line to test, so as best we can find some of those things, that'd be helpful.

DR. ULSH: I agree with you, Mark, and after our last working -- well, after one of our previous working group meetings when we discussed the first 12, and there was some

confusion about what the badging policy was,

Jim Langsted put together a write-up on the

badging policies. The date on the copy I have

here is December 6<sup>th</sup>. I don't know exactly

what date that was sent out, but that has been

sent out.

MR. GRIFFON: I'm sure we got that, yeah.

DR. ULSH: So you're right, that's the other piece of the puzzle in answering that question. But I did want to point out, I mean, going to the second question, what effect might it have on coworker models. You would obviously be concerned about the areas, the operations in the uranium areas where there was a potential for high exposure and the foundry may be one of those places.

But again, the foundry is in Plant A, not Plant B. Plant B is where you're seeing a lot of the gaps. So I just wanted to point that out. We still have to go through the individual cases.

Arjun, I think there at the end -okay, well, one other thing I wanted to talk
about was the group three workers you
mentioned, I think, that there were some gaps

in the group three workers in the '50s. And again, the group three and four workers were the ones that were identified in the Medical Recall Program as having had the highest exposures onsite as Arjun mentioned.

But the other point that you have to consider is when did they become, when did they achieve that status? When did they become highly exposed individuals? So you shouldn't assume just taking one of these group three or group four people if you see a year when they were not monitored in 1953 and say, gosh, this is a highly exposed worker. Why wasn't this guy monitored? He might not have had a big uptake, a big intake incident or a big external exposure incident until much later. I mean, that's a possibility.

MR. GRIFFON: And a lot of them moved from uranium areas to the plutonium area.

DR. ULSH: Yes, that is true. So again, you have to look at the job exposure history even for those people as well, and we are doing that.

Now in terms of the '69 fire I believe that you brought up a point about were subs

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possibly going into the building.

DR. MAKHIJANI: No, no, just to clarify what I said. In regard to subcontractors, we didn't find evidence of systemic violation. What we suggest, and so that would appear to corroborate to some extent the site expert statement that has been made, but there's no documentation of enforcement of that policy because obviously it leaves some room for That you had subcontractor workers; problems. you had a problem onsite; you had a fire; you had an incident, and you had a need for workers and people were sent in without badges. Because there are statements in the petition about those kinds of problems, not directly about subcontractors. And so the suggestion was that some kind of verification of enforcement be made. But if subcontractor workers can be identified who were sent in, for instance, to clean up after the fire that they were actually badged. We haven't made any statement about whether they were or not or whether there are gaps in that area or not.

MR. GRIFFON: I was just going to ask, and I know you sent out that paper Jim Langsted

1 wrote it on the policies -- maybe refresh my 2 memory. By the time I found it on my hard 3 drive it'll probably be one o'clock. So does 4 it speak to this question of when did this policy of reading, you know, everybody had the 5 security badge with the TLD or film in it, and 6 7 they didn't read them all and in '69 that 8 started. 9 DR. ULSH: You're talking in '69. Let me 10 give you --MR. GRIFFON: Well, it started in '69. 11 12 saw that in the memo, right? But we're not 13 clear when it, how long that policy was --14 DR. ULSH: That is correct, Mark. 15 MR. GRIFFON: Well, how long was that policy 16 in effect or do you know? Okay, that's the 17 question. DR. ULSH: I know. 18 This is the question 19 that we've been hoping to come up with an 20 answer for for awhile. 21 The '69, the policy not to read badges 22 of workers who were on quarterly badge 23 exchange cycles in non-plutonium areas, the 24 first indication in the documentation that 25 we've seen of that was in a monthly progress

report, I believe, in April of '69. And that's in actually SC&A's report on that. It pulls out the quote. That was the first indication of when it started. We still have not found any indication of when that policy might have been rescinded.

Now, one thing that we are checking on, you've also got to remember that the justification for this that was given in a letter -- I don't want to say the name, but -- was that the reason they wanted to do that is because these people -- let's define cohort people -- had low exposure potentials, right? And it was a lot of effort involved in reading badges, using densitometers to read the film badges, and it may not be the best use of your resources, something like, it's not a direct quote.

In late '70 or maybe it was '71, the site switched to TLDs, and so that same argument may not hold in terms of being a lot of effort that could be better spent elsewhere. Now, we are trying to look and see whether those people that would have been covered by this policy, the non-plutonium

1 areas on quarterly exchange cycles, whether 2 their TLDs were read. We're checking on that. 3 I don't have an answer for you there, but 4 that's a possibility. I don't know. 5 MR. GRIFFON: And then the other question, 6 and I don't know if, I've got to admit I 7 forget, I did read it, believe it or not, but 8 I forget what it said. This question I 9 remember in one of the meetings it came up 10 with not only was there this new or this memo 11 that we found about this policy, but there was 12 also another explanation that I thought was 13 given which was that some personnel or some 14 subcontractor -- this gets into the 15 subcontractor question -- some subcontractors 16 did not require any monitoring? Was that 17 brought up as a policy after '64 or any time 18 period? I'm trying to remember. I thought 19 that was brought up as a possible explanation. 20 MS. MUNN: We have this one quote that was 21 in a (inaudible), subcontractor (inaudible) 22 not have been issued a dosimeter. 23 MR. GRIFFON: What are you reading from? 24 Which document is this? 25 MS. MUNN: This is the Rocky Flats Badging

1 (inaudible). 2 MR. GRIFFON: So it's SC&A's report? 3 MS. MUNN: No, I think it's (inaudible). 4 MR. GRIFFON: Can you read that, Wanda? 5 MS. MUNN: Yeah. On the badging report that 6 was sent out in December that the statement 7 was, however, subcontractor personnel with low 8 exposure potential may not have been issued a 9 dosimeter, and then it quotes paragraph 10 6.15.1, conditions where general Health 11 physics surveillance was not required: (a) in 12 areas where penetrating radiation levels are 13 not likely to exceed an average of 0.2 14 millirem per hour, comma, outside contractor 15 personnel may be utilized. Film badges will 16 not be provided under these conditions unless 17 on advice of Health physics, badging is 18 desired for assessing a possible criticality 19 exposure, end of quote. Health physics guide 20 for Rocky Flats division compiled by EA 21 (unintelligible), first issued July 1961, 22 reviewed January 1967, re-issued January 1970. 23 Section renumbered 5.1.7. 24 MR. GRIFFON: Well, I guess that's the 25 question. If some of these people that we

1	reviewed for completeness fall into that
2	subcontractor category, then they may not have
3	been required according to this policy to
4	DR. ULSH: That's what it appears to
5	indicate.
6	MR. GRIFFON: And we don't necessarily know,
7	or Arjun, you don't know whether these
8	individuals were subcontractors or primary
9	contractors based on the records you were
10	looking at, do you?
11	DR. MAKHIJANI: No, we didn't go to that
12	depth.
13	MR. GRIFFON: So that's where we're headed.
14	I knew that statement came up before.
15	DR. MAKHIJANI: Well, actually in the job
16	cards that I reviewed, I didn't see, I mean,
17	they were pretty much production workers in
18	category three and category four so far as I
19	remember, and there may have been exceptions.
20	MR. GRIFFON: The broader, the other
21	DR. MAKHIJANI: The other sample? No, we
22	didn't look at that.
23	MR. GRIFFON: So that's what we're engaged
24	in right now.
25	DR. ULSH: Okay, Mark, so that's where we

are with the question of when did this nonbadge policy end. We are pursuing that. I don't have an answer. I know it's still an

issue of concern.

Now in terms, Arjun, I understand your concern about the subs, that it would be nice to have some verification that this policy was actually enforced. And I can't speak to the general issue, but the specific example that you used, the 1969 fire, I can tell you that we spent a lot of time on that '69 fire and some of the interviews I believe that Mel conducted and maybe even some documentation.

I can't remember exactly, but right after the fire, the immediate aftermath of the fire, only professional staff were sent in.

There were very little number of them. I don't want to give a number because I don't remember exactly what it was. And they were badged. It wasn't subs that went in immediately after the fire to recover the plutonium because there was a concern about criticality. So that's the immediate aftermath.

Now, after the plutonium that was

involved in the fire was removed from the building then there were a lot of cleanup activities involved. Of course, this is another thing, and we see a couple of times here in SC&A's reports, in the aftermath of the fire, once the plutonium has been removed from the building, and there's no production operations going on, of course, the external exposure potential goes down. You can't assume that the same exposure potential exists as there was in full production. In fact, it's exactly the opposite.

DR. MAKHIJANI: There shouldn't be a misunderstanding about what the suggestion is. The suggestion is in fact that subcontractors were at high exposure potential or not when they went into radiological areas. The suggestion was simply a response to the statement that was made, I believe in November, that subcontractor workers were badged when they were sent into radiological areas which would fit.

If they were sent for cleanup, that would in my definition at least be a radiological area. And they may not have been

at risk of external exposure potential, but
they may have been at risk of internal
exposure potential. If they were not badged
and sent into radiological areas, then there
would be a kind of systemic, if that were a
problem, then there would be some
corroboration of what the petitioners have
been saying.

And so the concern that is being raised is, yes, there's been a statement by NIOSH, and yes, we did not find any evidence that that statement is incorrect. But it stands so far undocumented and uncorroborated, and it would be useful to have a corroboration not of high or low exposure potential, but whether there were subcontractors who were sent into cleanup areas whose records indicate that they were not issued badges.

That I think is an important item because it corresponds directly to the concerns that the petitioners have raised.

And I think we have tried to address those systematically and this one should be also.

Some verification is desirable. That's the statement that we've made.

DR. ULSH: Okay, Mark, I think that's all I've got on that at the moment.

DR. MAKHIJANI: Just a comment about the badging policies, I mean, we didn't raise any, I mean, the part of the reason to look at the job cards was to verify the statements that NIOSH has made. And I think broadly we did verify it. We did only a preliminary look at these job cards. It was rather rapid, and it did result in a confirmation that the non-badging and the gaps were in the non-plutonium area. So actually, we did confirm what has been said many times by NIOSH. And so I think that much has been validated.

The question about dose reconstruction that it raises is not confined to what the badging policy was and whether it was regarded as reasonable or done in good faith at the time. And we haven't said that it wasn't, and I think in fact what we found that it was. It was done in good faith. People who were thought to be at not high exposure potential were not badged.

In retrospect though NIOSH can't use the policies that were in effect then to do

dose reconstruction. They have to do dose reconstruction according to today's standards, and that's the question that I think is raised by the gaps in '51 to '59 is are the gaps in the areas that had high exposure potential where there were systemic lack of badging and how are those gaps to be filled by today's dose reconstruction standards.

And we're not saying that in this particular case some policy at the time was violated. We didn't find that. On the contrary we found that NIOSH statements were generally correct.

MR. GRIFFON: And I think we can't go much further with that until we see your findings -

DR. MAKHIJANI: Exactly.

MR. GRIFFON: -- and you do your final analysis on that yet.

MR. ELLIOTT: Let me just make sure I understand what I hear Arjun saying here that in essence, and this goes back to the verification comment he made, suggestion he made to us. In essence you validated what we have made as our stated position, and you're

1 suggesting that there's a way here to further 2 verify and confirm that validation. 3 DR. MAKHIJANI: No, no, you're mixing up two 4 separate issues. One issue was sort of a 5 subsidiary issue for a small proportion of 6 workers who were subcontractors, whether they 7 were sent into radiological areas without 8 badges. And that was just a verification. We 9 found no evidence of that. So that was a 10 verification suggestion for that. 11 MR. ELLIOTT: That's exactly what I was 12 referring to. 13 DR. MAKHIJANI: Yeah, but that's completely 14 separate from the data gaps for the 1950s --MR. ELLIOTT: I understand. 15 DR. MAKHIJANI: -- which were a result of 16 17 policy of who was badged and who was not 18 badged, and we didn't find any evidence that, 19 you know, there was some systematic effort in 20 the '50s to hide doses of highly exposed 21 people so therefore, they were not badged. We 22 found some confirmation that people in 23 plutonium areas were badged, although as I say 24 it's a broad screen what the gap is. And that 25 people in non-plutonium areas were not badged,

1 not all of them, but the non-badging was 2 focused in the non-plutonium areas. 3 MR. ELLIOTT: I think I'm saying the same 4 thing. 5 Okay, okay. MR. GRIFFON: 6 DR. MAKHIJANI: But that latter problem is a 7 coworker model problem, not a verification 8 problem. 9 MR. GRIFFON: I guess with respect to the 10 first question, the subcontractor question, I 11 think is there an action there is my question. 12 Is there an action on NIOSH's behalf? We've 13 got a statement by, concerns various other 14 petitioners, and you're saying we've had a 15 response that it was the policy at the time to 16 badge them if they were in the RAD areas, but 17 we have nothing to really support that, and 18 you know, I think the action would be give us 19 some data that supports that. I'm not sure 20 what exactly that would be, but is that sort 21 of what you're raising here? MR. ELLIOTT: What I'm hearing is their 22 23 understanding is no different than ours. 24 They're suggesting a way to verify and confirm 25 that we're both on the same --

1 MR. GRIFFON: Right, we haven't seen --2 MR. ELLIOTT: -- it goes to helping out the 3 petitioners understand what we've all done 4 here, and how we're reacting to their --5 MR. GRIFFON: Arjun's saying we haven't found any problems there, but we haven't 6 7 either seen the verification of that either, 8 right? 9 DR. ULSH: Well, what I'm wondering, Arjun, 10 is if -- and I really don't know this -- in 11 the '52 that we're looking at are there subs 12 included in there? 13 MR. GRIFFON: That's what I asked. 14 DR. MAKHIJANI: I did not look at the job 15 cards for 32 random sample workers so I don't 16 know. I do not believe that from my 17 preliminary review there were subs in the category three and category four workers. 18 19 MR. ELLIOTT: I'm simply saying what I'm 20 taking away from this is that you see things 21 the same as we stated, and you're providing us 22 a constructive suggestion on how to verify 23 what we both have seen. 24 DR. MAKHIJANI: Yes, in terms of the reasons 25 for the gaps and in terms of the overall

completeness picture apart from how you're going to do dose reconstruction, yes, we have no differences, and we didn't identify that any statement that NIOSH made was incorrect. We're just suggesting one, there is one outstanding question that relates more to the '69 but goes not only forward from '69 but back from'69 is what happened in the case of the types of individuals that have, that were primary contractor employees who have blanks before '69. And that's a separate --

MR. GRIFFON: We'll hold that.

DR. MAKHIJANI: Yeah, we'll just hold that.
That's the one exception.

MR. GRIFFON: So that's a question that is a potential action item. It would help, I think you're right, Larry, it would help support that position and verify that to a further extent.

MR. FITZGERALD: In the context of this step
I think more to validate the earlier limited
sampling that, in fact, these gaps were there,
were real and a larger sample helped answer
that question, but I think that's as far as it
went, just to, in fact, substantiate that

these gaps are, in fact, real. Now as far as the reasoning and interpretation, I think all that needs to follow.

MR. GRIFFON: And it may be if you look through these 32 job histories, and you find some subcontractors, and it shows that they were, in fact, monitored, that might answer the question. That might give us some evidence to support that position.

Now you may look at those 32 and find that none of them were subs, you know? But I don't know if you can easily identify within the claimant files, it may not be so easy to pull out who was a sub without looking -- is that something easy within the database that you can look up individuals and --

DR. ULSH: It seems to me that it would be fairly easy, Mark, to look at the job exposure history cards if they exist for subs.

DR. NETON: I believe they do.

DR. ULSH: Something to check. I mean, this is a way to go in and pick out a case by hand and say there's a card here. Is he a sub?

Well, nothing here, next guy. But in terms of like doing a sort on NOCTS or something to

1 pull out subs, I don't think we have a way to 2 easily do that. 3 MR. ELLIOTT: Unless they reported their 4 employment as with a specific contractor who 5 was at the site. 6 DR. NETON: I think we need to check with 7 Mel. He's done a lot of work on this with 8 Rocky Flats was one of the example sites 9 that we used to determine whether 10 subcontractors and Rocky Flats main workers 11 had differences in external dose profiles. 12 fact, that analysis found no difference, and 13 they went to a large extent to pull out as 14 many records as possible to identify 15 subcontractors. I think we ought to look at 16 that. 17 There were building trades workers, 18 subcontractors, but I think we ought to look 19 at that because a lot of work has been done in 20 that area. 21 MR. ELLIOTT: That's true. That didn't 22 occur to me. 23 MR. GRIFFON: That may be useful. that might be an action item, but I would also 24 25 say -- I don't know how to say this. I mean I

think we need to control the scope of that, and I think you need to control the scope. It would be good to have some evidence toward that, but I don't think that I would expect you to pull hundreds, you know, if you find a few subcontractor files or randomly find four or five or whatever that seem to support this position, that would be pretty good evidence toward the, you know.

DR. ULSH: Going forward, Mark, I just want to make sure that I know what the work group is expecting. Are we going to finish up our analysis of the 52 and let you know whether or not subs are included in there and then get a feeling from the Board whether or not that will suffice or is there an action item at this time that you want us to go pick out some subs and --

MR. GRIFFON: Well, I would say pending your analysis of the 32 if there's no subs in there, I think you should go and find some subs. And I'll let you define some, but I would think keep it a small sample.

MR. ELLIOTT: And look at this TIB-52.

MR. GRIFFON: Yeah, and maybe the TIB-52

1 already has done some of that. 2 MR. ELLIOTT: I would ask that you put that 3 second, and then if that doesn't produce 4 enough, then you go find some subs to look at. 5 MR. GRIFFON: If the work is done already 6 obviously don't spend additional resources, 7 right. But I'm not, well, I don't know. I 8 don't know how the subcontractors work, but if 9 buildings trades are in TIB-52, and these were 10 subcontract maintenance people, it may not, I 11 don't know if it's apples and apples but you 12 can find that out. 13 DR. NETON: But we need to at least look at 14 that so we don't duplicate our --15 MR. GRIFFON: Certainly as efficiently as 16 you can. If you've got the work done already, 17 use that, yeah. 18 MS. MUNN: And the methods Mel used might be 19 useful, too. 20 MR. GRIFFON: Yeah, I guess that's what I 21 would say. 22 And I think we have the '69 data gap 23 coming up, but I've got a request for a short 24 break, maybe ten minute break. We've got some 25 microphone issues, and we'll take a comfort

break. So ten minutes, I think we're going to keep the line open if that's okay with everyone. And we'll be off the record now.

(Whereupon, a break was taken at 11:14 a.m. and the meeting resumed at 11:29 a.m.)

MR. GRIFFON: I think we're ready to reconvene here, and I think, unless there's anything else on that first item, I think item two --

DR. MAURO (by Telephone): Excuse me, Mark, this is John Mauro. Before we leave the subject I just was, Arjun had mentioned during his presentation it sounds like one of the issues is that a group of workers where we're finding gaps, and I think everyone agrees that there are gaps. Are these workers either in the uranium areas or foundry areas where they were not monitored for externals.

But I also heard Arjun say that one of the concerns there is that under certain conditions the contact dose is on the order of two-to-three rem per hour when you have this thorium and protactinium sort of surface, and also that there might be, in other words not the standard 240 millirem per hour for contact

and two millirem per hour penetrating. I think that two millirem per hour is at one foot.

I'd like to hear a little bit about has NIOSH been looking at that particular dose reconstruction challenge. In other words how to deal with workers who may have a gap, not monitored, and may have been involved in handling or working with foundry operations. Is that something that they feel they've got a good handle on and it's tractable? Or do they think there are certain aspects to that aspect of dose reconstruction that they're still struggling with?

MR. GRIFFON: I think what we heard is it's a little premature for that because they're still looking at the Plant B versus foundry question, but I don't know. Brant might have something to add onto that. I think it's a discussion better saved for the next meeting. If we find that some of the foundry workers were not monitored then that might be a more relevant topic of discussion. I don't know. Brant, do you have anything?

DR. ULSH: That's exactly right, Mark.

That's why I made that distinction,

John, between Plant B and the foundry. The

gaps, as I understand it, that Arjun saw were

more for Plant B workers. I don't want to say

only for Plant B workers, but more for Plant

B. But that would not include the foundry

workers because they were in Plant A.

Now the remaining question is were those foundry workers in Plant A monitored or not. And that's what we're going to try to establish as we go through the 52 cases and see whether or not we have some input on that.

DR. MAURO (by Telephone): Okay, thank you.

DR. MAKHIJANI: Just from my recollection of the review of the job cards I don't remember a lot of workers in that sample from Plant A.

So I don't know whether we might have a small numbers problem in for the 1950s because we weren't initially screening for that, to sample that particular population. And so we might have a small numbers problem there that you'll need to do a more definitive look at that.

DR. ULSH: Right, that's the first step.

The first step as you said, Mark, is to look

at these 52. But if we don't find the answers there, we agree, Arjun, that it would certainly be an issue if there are people who were not monitored and had high exposure potential. We just have to look at that and see whether or not those foundry workers were, in fact, monitored.

DR. NETON: I think this is primarily a skin dose issue that we're talking about here which is a beta-type exposure. So you need to be looking at things like extremity exposures and shallow dose to the skin primarily because these are not two R per hour deep penetrating doses for the most part.

DR. MAURO (by Telephone): Jim, John again.

One of the calculations we have not done is to see, we are aware of the contact dose issue.

We weren't quite sure whether or not the two

MR per hour at one foot would also be affected by the fact that you have more of the protactinium close to the surface and whether or not that would change anything. I don't have a sense of that.

DR. NETON: You're going to have a little more (unintelligible) coming off of there, but

1 I think it's primarily a beta dose issue is 2 what these contact doses are. But if that's 3 the case, then it really becomes a skin dose 4 issue which is then not an SEC cancer although 5 it doesn't mean it couldn't be --6 MR. GRIFFON: Although we're looking at it 7 in the construct of completeness, so if these 8 people weren't monitored. And if they're 9 monitored for skin, they would also be 10 monitored for deep. You should see their 11 records, right? So that's really the issue 12 there, but I understand your point. 13 DR. MAKHIJANI: You also brought some deep gamma issues in relation to the daughter 14 15 products. 16 MR. GRIFFON: Yeah. 17 DR. MAKHIJANI: Potential. 18 DR. NETON: But it's just not in the order 19 of two rem. MR. GRIFFON: No, no, certainly not. 20 21 DR. NETON: Most of that is beta. MR. GRIFFON: Yeah, we agree on that. 22 23 All right, I think this discussion is better 24 served for when we have NIOSH's response and 25 have more data in front of us.

## 1969 DATA GAP

Let's move on. The second item, the '69 data gap, and I think this overlaps quite a bit with the first topic. But obviously, we've been looking at the specific problem for awhile of questions about blanks in data in the '69 time period. So I'll throw that to Joe or Arjun. I'm not sure who's --

MR. FITZGERALD: Yeah, I want to take a second just to sort of summarize how we got here because it does have a kind of convoluted, somewhat long history. I think some history would be a little dim at this point.

But during the course -- and I'm paraphrasing from our review -- during the course of SC&A's review a number of former workers from Rocky Flats were interviewed, and a number of them expressed concerns about the '69 fire. And Kathy Robertson-DeMers, in fact, spent a lot of time looking at the HIS-20 database and some of the records and found blanks and certainly zeros reported for a number of these workers.

And from that we compared notes with

work that Ron Buchanan was doing looking at external dosimetry pretty much records in terms of dose distribution, and in parallel, I think he established for '69 and '70 that there were a higher preponderance of zeros recorded for that time period.

And that was something that we shared with NIOSH, and I think it was sometime in the summer. And NIOSH went forward to basically look at the historic record to see if there's any explanation for why we would see these zeros, in some places blanks, particularly in '69-'70 timeframe.

NIOSH came back, I think it was sometime in the fall with at least two or three explanations, hypotheses, as to what may have occurred. And one was certainly that it was an administrative decision. I think we touched on that already that there was, in fact, a policy that was put in place. And then there was maybe a computer programming error, but I think clearly we've established that the policy, in fact, was the reason that, a large reason why we were seeing these zeros.

And the evaluation that we, in fact,

performed was to go back and look at the claimant files for the individuals that were, in fact, identified for this period. And there were 136 claimants that NIOSH identified originally of which 35 had no external dosimetry data at all for 1969. That's kind of a starting point for the review that Arjun and Kathy Robertson-DeMers did because I'm going to just let you catch up on the review.

And I don't think this has actually cleared Privacy Act review yet, so this is something that with that one step we can certainly, NIOSH provided.

DR. MAKHIJANI: Yeah, we've examined the data relating to claimants, but I just want to put a caveat about that in relation to all workers. We had this discussion in relation to the Nevada Test Site that really the goal in an SEC is to examine the data for the class of workers who are covered by the petition, not the class of claimants.

And there's no systematic procedure as yet for relating claimants to the whole group of workers statistically. That said, we all agree I think that it's very important to look

at the data that we have accessible which is the data of the claimants. And that is fairly plentiful so it's important to look at it.

As Joe said we looked at the HIS-20 database and found a number of zeros in 1969-1970 went up and that was the cause of the whole investigation. At first it might have been hypothesized that it may be associated with the fire, but NIOSH said that it was not associated with the fire. There may be other explanations, and I think we're in broad agreement that it was not primarily associated with the fire.

We looked at the various explanations. Just to make it short we have some questions about whether the computer error is relevant to this particular issue or not. But there was an error in the computer records, and these gaps might be related to that. We do agree that prior to the fire there was a policy put in place not to read badges of workers, or certain badges of workers in non-plutonium areas and that that the gaps appeared to be, at least were largely related to this policy.

Now when this policy was, now we know the policy started before the fire in 1969 and then went on after that for some time, probably into 1970. We don't know how far beyond 1970 it might extend. And as I said earlier in the completeness presentation, there's also some indication that it may precede 1969.

Now whether primary contractor workers were not being issued badges in the '64-'69 period and so there are gaps or they were not, they were issued badges that were not read is not so clear. NIOSH had said the former, but we don't have any documentation about that or any policy statement that's comparable to this 1969 report where a policy was adopted.

There was a rather more unfortunate aspect to the 1969 policy. So we did corroborate that the non-reading, which was about 1,000 badges in every cycle were not being read, so quite a large number of badges. We found that to issue badges and not read them is not a sound policy, but the problem was made much worse by discarding, the policy of discarding the badges after a few weeks.

So there's now no way to go back and verify whose badges were actually not read because apparently the badges were discarded.

We looked at the various external dose databases to see who this affected and how it affected them and what was actually entered into the record. There were four, the four different databases where external dose data are entered that are available in the claimant files, there's the occupational dose report, which is the handwritten summary dosimetry in the Health Physics file.

There's a dosimetry history by individual which is a computer printout generated prior to the HIS-20 database creation. There's the Health Physics External Radiation Exposure Report, which is a quarterly summary report. And there's the HIS-20 computerized database which NIOSH is using for its coworker model.

And so we looked at all four of these to examine the evolution of how these zeros got entered into the HIS-20 database. We looked at 19 individual workers who are all in the HIS-20 database. And there was one case

of a worker who's not in the HIS-20 database just missing as part of the group of workers that are not in that database whose case was also important because there was a gap in 1969 in the data processing, the original log books.

We looked at NIOSH's explanation that where in the original badge processing logs there's a zero across the place where the dose is entered with arrows going down that --

MR. GRIFFON: Excuse me, everyone on the phone. We've got some interference coming through from someone's phone. I don't know if someone's on a cell phone or changed through speaker. Whatever you're doing now, do what you were doing before.

DR. MAKHIJANI: Where was I? Yes, the zeros that were in the film processing logs that went down with an arrow at the dose, but there was no densitometer reading, seemed to indicate the times, at least in the year in 1970 when the badges were not read. And in those records the Health Physics External Radiation Exposure Report generally shows gaps or there's nothing. But the dosimetry history

by individual in the HIS-20 database actually replace the blanks by zeros.

So as a result now the HIS-20 database has got two kinds of zeros that are mixed up. There are zeros that were entered because a result, a badge was read and was less than the detectable limit, and there are zeros because a badge was not read that was actually issued. And because the badge has been thrown away, this raises some pretty serious questions about data integrity.

We didn't find that there was an intent to fabricate data, to actually write zeros when there were high readings or something like, but to write zeros where there was no actual reading of a badge is a problem in data integrity. And this database, therefore, where there are zeros is fundamentally flawed in that respect.

So unlike the earlier discussions we had about gaps, there is this extra problem in the HIS-20 database where --

MR. GRIFFON: Arjun, you might have said it already, but how many individuals did you find this flaw?

DR. MAKHIJANI: Well, we looked at 19 individuals who had complete blanks, we looked at 19 individuals whose badges we determined could not be read, and I think we had that same understanding of whose badges were not being read, as NIOSH.

MR. GRIFFON: Excuse me, we're still getting bad interference from someone's phone, so I don't know if people, if you've done something differently maybe hang up and dial in again or try a different phone. We'd appreciate it, very loud static coming through. And it's stopped now.

DR. MAKHIJANI: So maybe I should have said something about how we pick the 19 claimants to look at. Nineteen sixty-nine is peculiar in that we actually looked at workers with partially complete, partially incomplete and partially complete gaps. So there are workers who had gaps for part of 1969. There are workers who had gaps, that is, badges not read for all of the year.

So we thought to focus on those workers who had gaps for the whole year because that way we could determine whether

the zeros that are in the HIS-20 database actually correspond to gaps or not because otherwise it'd be rather more difficult. And so we focused on 19 individuals who, among the 1969 workers who did not have any external dose records in their original badge processing files, whose badges were not read, and examined what was there in the other databases where their doses were recorded.

And as I said the result was that in the dosimetry history by individual computer printout, out of these 19, there were zeros for ten individuals instead of the blanks or gaps, and there was no dosimetry history by individual file for nine individuals. And for all 19 cases the gaps or blanks have been replaced by zeros in the HIS-20 database.

So that was the result of that analysis. And you have the case-by-case analysis in the full report which is covered by the Privacy Act, with you. So there could be a number of, so the conclusion is that unread badges were being entered as zeros in the HIS-20 database. And to some extent also in the dosimetry history by individual

computer printouts. So the problem seems to have originated somewhat prior to the HIS-20 database but was carried over and aggravated by the HIS-20 database compilation.

There are a lot of reasons for concern about this conclusion. We looked at the same workers' records for 1968 and 1971 to see perhaps whether the zeros might correspond to a situation where they had no exposures in the earlier years but their badges were actually read. And we found that there was an increase in the number of zeros from '68 to '69, that there were workers who had non-zero doses recorded in the earlier years immediately preceding who suddenly had zeros in the HIS-20 database and whose badges were not read.

So it's not clear that the badges of workers, that the workers whose badges were not being read had all uniformly low exposure potential. We looked at the case of one worker who was not in the HIS-20 database but whose case seemed to be quite important even though there's no record in the HIS-20 database of his file, and to bring up his sample.

1 MR. GRIFFON: Careful not to reference too 2 specifically. 3 DR. MAKHIJANI: Yes, I will not. 4 MS. HOMOKI-TITUS: You might want to tell 5 them somebody who --6 DR. MAKHIJANI: Well, maybe, maybe not. 7 Maybe not. 8 MR. GRIFFON: Hold off on that. 9 DR. MAKHIJANI: Yes, just hold off. I need 10 to find the, yes, here. 11 This person is listed as a nonplutonium worker so he would fit into the 12 general description of workers who were 13 14 covered by the non-reading of badges, but 15 actually worked in a lot of different areas. 16 And so there's the question of whether the 17 workers who are described and officially 18 assigned to non-plutonium areas were more 19 frequently also working in plutonium areas. 20 Also, we found quite significant, more 21 than the ten percent limit doses recorded in 22 every year for four years prior to the, prior 23 to 1969. And then the badges were not read in 24 1969. So this is a sort of a particular case and NIOSH has the data, claimant numbers and

1 all of that information on this particular 2 person. So that example actually raises a lot 3 of different questions about job assignments 4 as they are in the, as they actually happened 5 compared to the policy of whose badge was not being read in 1969. 6 7 MS. MUNN: Arjun, what, do you have the 8 figures post-1969 for the same individuals? 9 DR. MAKHIJANI: We do. 10 MS. MUNN: And do you see better comparisons 11 between the post-'69 and the pre-'69 figures 12 such that you can, so that you are convinced 13 that there was not likely a change in their 14 work history during that particular period? 15 DR. MAKHIJANI: We looked at people whose 16 work histories did not change over certain 17 periods of time prior to and after 1969, and 18 maybe Kathy can elaborate on what I'm saying. 19 And part of the reason for the concern is both 20 immediately prior and immediately after some 21 of the people whose badges are not being read 22 do show non-zero doses. 23 MS. MUNN: The static is back. 24 MR. GRIFFON: Yeah, can I ask everyone on 25 the phone, I know this is kind of a hassle,

1	but if everyone could just hang up and redial
2	back in. And we'll pause for a minute and let
3	
4	UNIDENTIFIED (by Telephone): I think
5	someone out there may have a Blackberry, and
6	they need to disable that. That's what the
7	static is often associated with a Blackberry
8	trying to pick up mail from the server.
9	That's what it sounds like to me.
10	DR. WADE (by Telephone): I'm going to leave
11	you now, too, Mark. I wish you well.
12	MR. GRIFFON: Lew, thanks.
13	MS. DeMERS (by Telephone): Wanda, this is
14	Kathy. Arjun is right. We selected
15	(unintelligible) table can you guys hear
16	me?
17	MS. MUNN: Yes, we can.
18	MR. GRIFFON: Yeah, we can hear you.
19	DR. MAKHIJANI: Be careful, all right,
20	Kathy?
21	MS. DeMERS (by Telephone): so that they
22	had the same job title before 1969 and then to
23	1970, and they were also assigned to the same
24	area.
25	MS. MUNN: Yeah, okay, I just, I understand

that consistency. It's just it's difficult for me to not expect that there would have been some change in the way operations were conducted for awhile after that major fire. I would just assume that a lot of people, even though they went back to their former jobs afterwards, would somehow have been in some other part of the site doing some other kinds of work for a short period surrounding that massive incident. I don't know how we could show that, but you've answered my question, thank you.

MS. DeMERS (by Telephone): This is Kathy.

Remember that we're talking about uranium

workers here. And the only shift they would

have was to the plutonium area. We're

actually going from an area that's supposed to

be lower exposure to higher exposure.

MS. MUNN: Except that the plutonium had all been taken out, and so it's a different ballgame. But I understand and my question has been answered. Thank you.

MS. DeMERS (by Telephone): One thing I would like NIOSH to provide to us is some sort of actual indication that the doses dropped in

the 777 service area after the fire because we're talking about that, but we haven't seen anything like a survey report particularly on the external dosimetry or anything like that that did occur.

MR. GRIFFON: Okay, we'll go back to that, but let Arjun finish up his report.

DR. MAKHIJANI: Let me just finish up the conclusions about 1969 data integrity and where we wound up.

So there are four major bullets of issues that arise in relation to data integrity in the 1969 investigation. Not reading badges, not consonant with sound practice and throwing away the badge after a few weeks converted a problem of unsound practice to a problem of data integrity because it cannot now be verified as to what the doses were and whether the assumption of lower exposures was valid.

We tried to examine that in other ways, and we would have definitely had some questions about whether that assumption was generally valid for the people whose badges were not read. There are indications that it

was not.

Non-reading of badges appears to have been done with the intention of minimizing work, but there's some indication that this was, at least for some workers, an erroneous belief and notably for shallow dose. The entry of zeros in regard to the entry of zeros in some data records when badges were not read is a data integrity question about data recording practices for as long as that practice went on.

And we have some question whether it was initiated as a de facto practice without being declared before 1969. And it was there definitely in '69, continued into 1970, and when exactly it was stopped is not clear as we've discussed before. And we have some examples of workers whose badges were not read whose formal assignment did not include plutonium areas who did work in plutonium areas.

And there's also the case of one worker who was involved with the fire who apparently, whose badge was not read. So there's that problem that also arose in our

1 survey. 2 MR. GRIFFON: The one with the fire, it was 3 a response capacity? 4 DR. MAKHIJANI: Yes, in a response capacity. 5 MS. DeMERS (by Telephone): Now, he was a 6 fireman who fought the fire. 7 DR. MAKHIJANI: Yeah, that's what I said. 8 So that's why I said in a response capacity to 9 the fire. 10 MR. GRIFFON: Apparently, I see Brant maybe 11 not agreeing with that. 12 DR. ULSH: Well, I don't agree or disagree. 13 What I need to check though, I think the 14 report says fire suppression. And we do know 15 that there were people, and I don't know if 16 this applies to this individual or not, there 17 were people who were involved in fire watch, 18 fire suppression, who never went in the 19 building at all. And I don't know if this is 20 one of those people or not. What we have 21 heard from the people who were involved in the 22 immediate aftermath of the fire was that only 23 the professional staff went in, and they, 24 after the fire, and they were monitored. So I 25 don't know. We've got to look a little closer

on this one.

2 3

MS. DeMERS (by Telephone): Do you want me to respond to that?

4 5 MR. GRIFFON: I don't know if it's worth it

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that we've left up to NIOSH.

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at this point. There's just a question there I think. Let's leave it at that for now. DR. MAKHIJANI: And as before in the completeness investigation, we haven't kind of, we felt that our instruction from the

working group was to examine the issue to see

if there was one, and that the detailed work of resolving it was NIOSH's work and not our work. So we haven't tried to track all these issues down to the last, and so it may well be that there's a good explanation or not.

The conclusions in regard to the overall situation with the 1969 data is that overall the gaps don't appear to be related to the fire although there's some fire-associated questions but largely to this policy of not reading badges. And there is questions of data integrity and also about how the coworker models are going to be developed with uranium workers which we also raised earlier, the

complication with 1969 compared to the 1950s just to be clear.

In the 1950s there was a policy of not badging a certain group of workers, and they were not badged, and there was no data integrity problem there. Here there were badges that were issued -- maybe I can just talk very loudly, overcome the static.

Here there were badges that were issued that were not read and were subsequently thrown away so this is a much different issue. And that to some extent at least corroborates the petitioners that there were zeros entered into some dose records when the badges were not being read. That's an issue that's been raised in the petition, and while in the earlier period we did not find that, it was simply people who were not badged. Here there is a more substantial issue of data integrity.

MR. GRIFFON: Can I ask, Arjun, did you, I know you were focused on the '69 period, did you find this practice in other time periods beyond, I think you said '70 there might have been some, too --

1 DR. MAKHIJANI: It goes into 1970, but we 2 did not actually go into individual dose 3 records --4 MR. GRIFFON: -- prior to or after that? 5 DR. MAKHIJANI: Well, there's the one case 6 that we have of a person with 11 years of 7 blank records from '63 to '74 that came up in 8 the data completeness question in the very 9 first set that we examined. Actually, I think 10 it was the very first one that I looked at 11 which caused me to sit up. And --12 MR. GRIFFON: And in that instance there 13 were blanks all through the hard copy file, 14 but the database had zeros for that --15 DR. MAKHIJANI: Yes. 16 MR. GRIFFON: -- those 11 years? Or do you 17 know? 18 DR. MAKHIJANI: Well, I don't know that the 19 database has zeros because I've not tracked 20 the individual's HIS-20 file. I know that 21 this was a worker that was supposed to be 22 badged because they were a prime contractor 23 worker. NIOSH said in a working group meeting 24 that they were not badged, but we have no 25 documentation of that.

1 MR. GRIFFON: Well, we have no policy sort 2 of thing to see, right, right. 3 DR. MAKHIJANI: And so they may have been 4 issued a badge, and there may be no readings. 5 Or they were not issued a badge. But in both 6 cases it kind of raises a pretty big question of was there really ever a universal badging 7 8 policy in place for prime contractor workers 9 and who was not badged, if NIOSH's statement 10 is to be taken at face value? Or if not, was 11 this policy initiated more formally in 1969 of 12 not reading large numbers of badges to save 13 work, actually de facto introduced at some 14 earlier time and then formalized in '69, and 15 then stopped at some time maybe when TLDs were introduced. It's unclear. There are a lot of 16 17 questions there. We can make some guesses, 18 but --19 MR. GRIFFON: Yeah, let's not make guesses. 20 Brant, I'm sure you pretty much --21 That's it, Arjun, for a summary? 22 DR. MAKHIJANI: (no audible response) 23 MR. GRIFFON: Go ahead, Brant. 24 DR. ULSH: We certainly agree that there are 25 more zeros in '69, and it looks like from the

figures that SC&A has put in their report, possibly into '70 as well. As Arjun mentioned when this issue was first brought to our attention, we considered a number of hypotheses, the first of which, I mean the most obvious of which would be some connection with the fire. And we eventually determined that it was not related to that.

MS. MUNN: Static again.

DR. ULSH: But I want to be clear. The population that this issue affects, this policy of not reading the badges of people who were in the non-plutonium areas, there's another criteria in there who were on quarterly badge exchange cycles. Now these were people who were judged by the health physics staff to be at low exposure potential. And we can discuss whether or not they could make that determination adequately.

But we are not talking about all people who are in non-plutonium areas, only those who are on quarterly. There were also people who were on monthly exchange cycles, more frequent than quarterly. So the issue that we have here is for the people who are on

quarterly, and that's the cohort that was judged to be a low exposure potential.

Now we also have discussed in the past that in terms of 1970 there was a strike that occurred in the summer of 1970, lasted for about 70 days. And also you have the '69 fire, of course, which disrupted production in the highest exposure areas of the site. It took the people onsite who, in general, got the highest exposures and put them in a non-production role in some cases. So you would absolutely expect to see higher frequencies of zeros. That's not surprising.

Now in terms of this policy, you know, it's certainly not the way I would have done things, but that's not really the question.

The question is does it prevent us from doing dose reconstructions with sufficient accuracy.

Now, I don't want to commit to a work around on this until we finish our analysis, but one thing that I would like to put on the table for the working group and SC&A to consider is that given that we have a cohort of people who were judged by the health physicists at least to be at low exposure

1 potential, what would be, I mean, if we decide 2 that those people, that cohort of people, the 3 quarterlies, could not be used because we 4 can't differentiate real zeros from really 5 unmonitored doses that were artificially 6 converted to zeros. 7 If we decide at the end, and I'm not 8 saying we are, but if we decide at the end 9 that that population of data is corrupt, and 10 we can't use it, we can't rely on it, what would be the effect of not using that data? 11 12 Of taking the quarterly badge exchange cycles 13 and not considering them? 14 Well, if you think about --15 MS. MUNN: Static again. 16 DR. ULSH: I hope you're talking about the 17 reception and not me. 18 MS. MUNN: No, but that is a Blackberry. 19 do wish you'd take your PDA and place it 20 elsewhere. 21 DR. ULSH: So if you look at the 22 distribution of coworker, the distribution of 23 the population of doses that you have, and you 24 take out the people who are on quarterly, what 25 are you doing? Well, in general, and I'm not

saying in every single case but in general, speaking from a population standpoint, what you're doing is you're lopping off the lower end of your coworker distribution so that when I pick a 95 percentile value or a 50 percentile value or whatever it is that we pick, the value that I pick without the data would be higher than the value I would pick with that data in there.

So in effect at the end of the day, if we say this data's no good, we take it out. That's a claimant favorable thing to do in terms of assigning coworker doses to unmonitored people. And that's what we're talking about here. These people whose badges were not read, in fact, they're unmonitored. That's what they are. They wore badges, but those badges weren't read. So it's the same as if they weren't wearing them.

And if you look at the coworker data that, the values of coworker data that we are proposing, that we have in our TIBs for 1969, it is far, far, far higher than you see for people, you know, the lowest exposure people on the site. So I'm not committing to that

1 approach. I'm just saying that when you think 2 about this in terms of what is its relevance 3 in terms of an SEC consideration, that is 4 something that we should think about. 5 Now one of the conclusions that SC&A 6 has in their report here I have to express 7 some caution about. It says while the non-8 reading of badges may have been done with the 9 attempt to minimize work -- well, let me just 10 read it in total. 11 "While the non-reading of badges may 12 have been done with the intent of minimizing 13 work related to reading badges of workers 14 judged to have low exposure potential, the 15 facts relating to at least some non-plutonium 16 work indicate that this was an erroneous 17 belief, notably for shallow dose." 18 Now am I correct in assuming that that 19 refers to the uranium foundry workers or does 20 that refer to something else? 21 MS. DeMERS (by Telephone): No, it's not 22 related to them. 23 DR. MAKHIJANI: Right, Kathy. Go ahead. 24 MS. DeMERS (by Telephone): When I went and 25 investigated gaps, I did a spread from various

1 buildings and just for example, there's a --2 MS. MUNN: Static, there's static out there. 3 MS. DeMERS (by Telephone): It's probably 4 coming from my phone. 5 MS. MUNN: Well, I hope not. We need you. 6 MS. DeMERS (by Telephone): Just to give you 7 an example, there's an administrative building 8 9 MR. GRIFFON: A little louder if you could, 10 Kathy. 11 MS. DeMERS (by Telephone): There's an 12 administrative building, 111, and normally 13 they're not routinely in the field, but they 14 do periodically go into the field. And in one 15 case on our table we have an industrial 16 photographer. Well, he happened to have 17 photographed all those wonderful pictures of 18 the fire, and subsequent pictures of the 19 cleanup. So there's an example where, no, 20 they're not coming from the foundry, they're 21 not coming from the immediate uranium area, 22 they're just a professional that had the 23 opportunity to intermittently go out into the 24 field.

Some of the people we looked at were

1 machinists. 2 MR. GRIFFON: Machinists. 3 MS. DeMERS (by Telephone): Some of them 4 were equipment operators. We have an 5 electrician. 6 Okay, so it's broader than the MR. GRIFFON: 7 uranium and foundry workers is what you're 8 saying. 9 MS. DeMERS (by Telephone): Right. 10 DR. MAKHIJANI: Mark, the data are in your 11 write-up in Table X-4, and it does show a 12 sampling of different buildings, different job 13 types. They are largely from Building 444, 14 but there's also the 881 and 883 and 111, 331. 15 And you see quite large beta doses, relatively 16 speaking, in at least two, three, four cases. 17 And these are all cases in the last, where 18 these beta doses are recorded in the last 19 quarter of '68. And in all these cases if you look at the first quarter 1969, the badges 20 21 weren't being read. 22 DR. ULSH: Okay, now I understand what 23 you're saying. Okay, we'll take a closer look 24 at the Table X-4. Arjun, I'm looking at the 25 de-identified version. I think you guys also

1 sent us one where it does identify these individuals, right? 2 3 DR. MAKHIJANI: Yes. 4 Kathy, did we send all the claimant numbers -- I do not remember now -- to NIOSH? 5 MS. DeMERS (by Telephone): If you don't 6 7 have them, I can pull those up. 8 MR. GRIFFON: We'll make sure it happens if 9 you don't. 10 DR. MAKHIJANI: I don't know if we actually 11 in this case sent you a version with claimant numbers. At least I don't remember seeing one 12 13 with claimant numbers. 14 MR. GRIFFON: And this report from what I 15 understand is still under Privacy Review. that correct? 16 17 DR. ULSH: I think, Joe, is that what you --MR. FITZGERALD: Yeah, that's my 18 19 understanding. The copy I have has a notation 20 that it's subject to Privacy review, and I think we sent it to you and Brant back in the 21 10<sup>th</sup> of January. I think that's the only one 22 23 that hasn't come back. Now there's been so 24 much going back and forth that I think we just 25 need to validate that that's where it stands.

1 But I believe that's where it is. 2 MR. GRIFFON: I just want to, again, and 3 this goes for all the, those four items that 4 we'll discuss the rest of the day, that we'll 5 commit to getting these to the petitioners' 6 interested parties as they're cleared for 7 Privacy concerns. 8 DR. ULSH: So like the other sections of 9 SC&A's report, we are still reviewing this. 10 And this is one where in contrast to some of the other sections, I think SC&A sees a 11 12 potential at least for SEC implications. 13 don't see any disagreement. 14 DR. MAKHIJANI: Well, this is, of course, up 15 to the Board. We did raise the question about 16 dose reconstructability given the state of the 17 data for this group of workers. But there's 18 also that to some extent we raised the 19 question also in regard to some workers in the 20 1950s that it's not a separate question that 21 the dose reconstructability for 1950s is 22 settled for the uranium workers. 23 DR. ULSH: Well, where I'm headed with this 24 is that this is one where since there's at 25 least some belief that there might be SEC

implications, that this is one that we're going to focus on more than some of the other ones like, you know, we're going to get into log books and whatnot. This would be a higher priority for us.

MR. GRIFFON: It's certainly a high priority I think. All of the data included in this section I think is a high priority and with '69 maybe being the highest of that.

DR. ULSH: So yeah, we're going to be preparing a detailed response. We'll look at each of the individual cases here. And like I said, the point that I just want to make is that we are not talking about all 1969 data. We are talking about quarterly data for non-plutonium areas.

DR. MAKHIJANI: This is correct that I should have mentioned that we're talking about quarterly data so that I omitted that fact.

And I also omitted something else which is that there were some badges that were contaminated after the fire. And both during worker interviews and in some documentation it's indicated that some badges at least were thrown away. Not again, for nefarious

1	purpose, but simply because you don't want to
2	contaminate the reading equipment and so on.
3	But some badges of people involved in the fire
4	would have been thrown away. So some of the
5	zeros may be related, or gaps, may be related
6	to that.
7	MR. GRIFFON: Did you do follow up to see
8	because the suggestion I saw in some of the
9	log books was that badges were destroyed for
10	those contamination reasons, right?
11	DR. MAKHIJANI: Right.
12	MR. GRIFFON: But I thought I saw, at least
13	one of the log books suggested that health
14	physics assigned or assigned a dose to the
15	individual
16	DR. MAKHIJANI: Kathy would have looked at
17	that.
18	MR. GRIFFON: and I don't know if you
19	cross-walked any of those.
20	DR. MAKHIJANI: Kathy, did you?
21	MS. DeMERS (by Telephone): I took all of
22	the discussion of badged individuals and
23	included them in the log book review.
24	Unfortunately, only two of them had claimant
25	files. And those two, if I remember, did not

1 have the dosimetry investigation in their 2 file. 3 MR. GRIFFON: Did they have anything in 4 their record, in their recorded dose for that 5 quarter or year? That may be a little inconclusive there because they may have had 6 7 other periods where they had monitoring and 8 the whole year would add something. 9 MS. DeMERS (by Telephone): I don't think we 10 were able to define the particular badge 11 period. 12 MR. GRIFFON: Okay, maybe we'll catch that in the log book stuff, too, but I just 13 14 wondered if any of that had been tracked 15 through. 16 DR. MAKHIJANI: Kathy would have looked at 17 this. I certainly did not. 18 DR. ULSH: Well, there were a couple of 19 points there that I can maybe speak to now. 20 All of our indications are that when people 21 went in immediately after the fire to assess 22 the situation and to remove the plutonium, 23 that those people were fully suited and double 24 badged. So I mean, it is certainly possible, 25 and I think it was even probable, well, I

don't think there's any question that on occasion badges became contaminated and were disposed of. And those are noted in Kittinger's logs and the notation said that that was witnessed by other people.

Now in terms of the lack of investigation reports, I don't know. I'd look at the agenda again. We may get to that some other place on the agenda, but as we've discussed in the past, you would expect to find the abbreviated or extended dosimetry, sorry, dose reconstruction investigation I think is the title of it, in the later periods. You know, we provided those procedures. In the earlier periods you would not necessarily expect to see the same kind of documentation.

MS. MUNN: Worse than static.

DR. ULSH: That's not to say though that investigations weren't done, just that they didn't have a formal mechanism like they did in the '90s, I don't know what the time period was, where they had a standard form that they wrote up the investigation on. We have seen from various documents that I believe Kathy

might have supplied, although don't hold me to that, there were some individuals where, just for example, the, I think it was the neutron badges were higher than they expected.

And it clearly indicates that that situation was investigated and that doses were assigned. So I mean there are situations that indicate that investigations were done. It's just the documentation is not the same as it was in the '90s.

MR. GRIFFON: I guess that was my point is that the log books seem to suggest that they did attempt to assign some dose. And if you cross-walk that and see there was a blank then that might be an issue. But if there's some dose recorded --

DR. ULSH: Right.

MR. GRIFFON: -- then, you know, it might be within their procedures.

DR. ULSH: Well, we interviewed a couple of people on this, one of whom was, okay, I pause, but I think that's okay, one of whom was Bob Bistline who was the site expert that SC&A employs, and he strongly disagreed with the suggestion that they would have just

1 assigned zero in a situation. They always, 2 okay, I don't want to say always again, but it 3 was the policy to investigate situations where 4 badges were suspect or had to be discarded. 5 So I mean, we've heard that from a number of -6 7 MR. GRIFFON: Well, I'm just simply asking 8 the simple question, these log books actually 9 have specific badge numbers, and if you can 10 cross-walk them that may actually strengthen 11 that. 12 **DR. ULSH:** Do I hear an action item? MR. GRIFFON: No, but I think we might pick 13 14 that up in the log book analysis. I think 15 it's already in that so that discussion --16 Kathy, unless you want to, some of 17 this I think gets into our log book 18 discussions. 19 MS. DeMERS (by Telephone): Actually, in all 20 the entries we pulled out of the log books I 21 think there was one that indicated that there 22 was an investigation in the field. 23 MR. GRIFFON: Okay, but again, we don't know 24 if, the documentation process in the early 25 years versus the later years, I mean, I think

we can discuss this further. I think I'd like to save it for the log book because I know we have it in there.

I mean, I guess for the '69 question really, you're looking into this, the only thing I would say is that we keep saying '69 and I guess the thing that jumps out at me in this discussion is the blanks and having zeros entered in the HIS-20. And it brings up this whole question of the integrity of the HIS-20 database period.

And I think to the extent we can, and this might involve SC&A, I'm not sure who should have this action, but I know out of those other cases it seems to me that it wasn't only necessarily a '69 issue. Or you had that one example of the first twelve cases where you had 11 years, but you didn't then look and see if that carried through to the database if the blanks in that individual's records carried through to zeros in the database.

And I think that, you know, I have a question in my mind is was this just in '69 where you had some blanks inadvertently or for

1	whatever reason entered as zeros in the
2	database or is that in other time periods as
3	well. And maybe all of this can be resolved
4	like you said. I hear your option about
5	potentially you could just drop this quarterly
6	and that may
7	DR. ULSH: And this ties back again to that
8	question you asked earlier, Mark, about when
9	did that non-read policy end, right?
10	MR. GRIFFON: Right, partially, yeah.
11	DR. ULSH: If that question could be
12	answered it might put a
13	MR. GRIFFON: Shed some light on all of
14	this, yeah.
15	MS. DeMERS (by Telephone): I have a
16	suggestion.
17	MR. GRIFFON: But it also that one example
18	is prior to '69, too, which is a little
19	DR. ULSH: This is the individual with 11
20	years, the administrative assistant?
21	MR. GRIFFON: Yeah.
22	MS. MUNN: Kathy's trying to say something.
23	MR. GRIFFON: Go ahead, Kathy.
24	MS. DeMERS (by Telephone): I have a
25	suggestion on when the policy may have ended,

and you can check this out. The uranium areas were among the last people to be assigned the TLDs towards the end of '70, and that policy may have ended with the assignment of TLDs to different groups.

MR. GRIFFON: Yeah, I think Brant kind of suggested that earlier, so if we can, you know, have that may, but then that raises the question of if individuals have blanks beyond that, why? So it's the same I've raised as before. That if the policy were short term it may explain this spike in the number of zeros for that one year, but it doesn't explain the gaps or these blanks in other years beyond that time period.

DR. ULSH: What I would suggest to you,
Mark, is that we get back to you after we have
looked at those TLD worksheets or results, TLD
results, and see if we see the same pattern.
Then we can go forward.

DR. MAKHIJANI: Mark, I think maybe two action items and probably for NIOSH, but two action items that would help in clarifying these issues at least. If these same individuals that we've looked at could be

tracked into the TLD period and before 1969, that would be very useful to see what kind of records they have and whether there was a non-reading policy.

And in regard to that one case with 11 years of blanks, if we could look to see if that person is in the HIS-20 database and what might be entered there. And similarly for the cases that have gaps in the '50s where there were no badges issued whether who was there in the HIS-20 database and what's in the HIS-20 database. That'd be obviously important because of its use for the coworker model.

So I think that would clarify at least some of the issues that are on the table in regard to what's in that database and the status of these individuals and maybe when these policies began. I'm not sure that it will, but it may.

MR. GRIFFON: And I think this will be, probably you can roll this into your response to these, the first two items anyway, so that makes sense.

MS. JESSEN: Three items. Weren't there three?

1	MR. GRIFFON: Were there three?
2	DR. MAKHIJANI: Maybe there were.
3	MR. GRIFFON: Two and I thought you rolled
4	into a third. Should we restate those? Just
5	for clarity restate those, Arjun. Track the
6	individuals found in your '69 review, right,
7	back in the prior years and future years?
8	DR. MAKHIJANI: And future especially if
9	there's a clear suggestion that their TLDs
10	were being read then we will have a pretty
11	good answer about the ending date for this
12	policy.
13	And then there's the one case of the
14	administrative assistant
15	MR. GRIFFON: Yeah, 11 year case, yeah.
16	DR. MAKHIJANI: and then there's the gaps
17	from the data completeness, so those are
18	three, three things.
19	MR. GRIFFON: The last one is the gaps from
20	the
21	DR. MAKHIJANI: Gaps from the data
22	completeness, whether they're zeros in the
23	HIS-20.
24	MR. GRIFFON: From your '52 case review,
25	right?

1	DR. MAKHIJANI: Yeah, from the case reviews
2	in the first paper that we discussed.
3	MR. GRIFFON: Anything else on '69 because I
4	think we're ready to
5	DR. MAKHIJANI: I think that's it.
6	MR. GRIFFON: break for lunch.
7	Anything else, Brant?
8	Okay, I think we'll break for lunch
9	and come back at 1:30, and we're off the
10	record now.
11	(Whereupon, a lunch break was taken at 12:27
12	p.m. and the meeting resumed at 1:35 p.m.)
13	MR. GRIFFON: We're reconvening the work
14	group meeting, the Rocky Flats work group
15	meeting, and I think we're on to item three.
16	Now, I did have a follow-up question
17	on the first two items we covered, the data
18	completeness item, specifically, I know that
19	we talked about the policies around the
20	external monitoring program. And I think it
21	might be worth, at least for me I would like
22	to see if you can incorporate into your
23	response something about the internal
24	monitoring requirements and how they pertain
25	to the gaps or non-gaps in the SC&A report.

For instance, if you have someone that has annual plutonium samples, but the policy said for that building they should have been on quarterly or something, you know. I don't think that they got into that detail, but I think that might be, you know, if we're looking to see if policies are consistent with what we're finding in the data, I think that might be useful.

I'm still getting that echo. I was hoping that would go away.

DR. MAKHIJANI: And, Mark, as I said in my introduction to the completeness thing, we used a very broad screen and besides the time periods and the intra-year data, within the year results of the radionuclide question. So there, you know, this is just one aspect of completeness that we did.

MR. GRIFFON: And I should also say that I don't know if this is a complete new subaction on this item or anything because it may, I'm not, I was going to go back and I didn't get a chance to go back to the internal dose site profile so you may have laid a lot.

I think you have some of that

1 information in there on who, what type of 2 monitoring was done, what time periods and 3 things like that. So some of that may be 4 there already. I'm not looking to recreate, 5 you know, if you already have it there, it's 6 Just maybe bring it to our attention. fine. 7 MR. BUCHANAN (by Telephone): Mark, this is 8 Ron Buchanan. Can you hear me? 9 MR. GRIFFON: Yes, go ahead, Ron. 10 MR. BUCHANAN (by Telephone): One thing that 11 I have suggested, and I don't think that 12 anyone's ever done is on the '69-'70 that if 13 we go back and look at the internal dose data 14 and see if it has the same spike and number of 15 zeros, it might shed a little light on why 16 these occurred in external doses. 17 MR. GRIFFON: Yeah, I don't know that we 18 looked at that. I'm not, I don't think that's 19 been raised before. 20 MR. BUCHANAN (by Telephone): I don't know. 21 It just might shed some light on why there was 22 or wasn't a reason for the increase in the 23 zeros and see if the same thing happened in 24 the internal doses. 25 MR. GRIFFON: How would that shed some

1 light? 2 MR. BUCHANAN (by Telephone): Well, if we'd 3 seen the same thing. If we didn't see that, 4 well then we would see that these people were 5 being monitored and there's something wrong 6 with the external dose. But if there was the 7 same scenario in the internal dose then that 8 would probably shed a different light on why 9 this was being done. 10 MR. GRIFFON: It may I guess I 11 would say for these 20 or so cases that were 12 brought up for the '69, is it 20 individuals? 13 DR. MAKHIJANI: The 19 individuals who had 14 the zeros in the HIS-20 database, and then there's the one individual that's not in the 15 16 HIS-20 database. So there are 20 in all. 17 MR. GRIFFON: See, I'm not sure it 18 definitively answers, addresses any questions 19 because they, I think Brant was suggesting 20 earlier that it was possible that they were on 21 some sort of internal program but not required to be badged or didn't read their badges. I 22 23 don't know. 24 DR. ULSH: Yeah, I mean, keeping in mind 25 that I think we've agreed, SC&A and NIOSH have

agreed that it's at least consistent with this badge non-reading policy to see this spike in zeros. But that only applies to film badges. That doesn't tell us anything at all about what they did with internal. So I don't know if you would expect to see a spike in zeros or not. I don't know.

MR. PRESLEY: You'd almost have to have the same number. You'd have to be looking at the same people, wouldn't you?

MR. BUCHANAN (by Telephone): What I was thinking is if you had an overall increase in the internal dose not being monitored during '69 and '70, that might lend itself to say, okay, we didn't have people in radiation areas so we weren't badging them where we had a lot of zeros. Internal monitoring remained a constant from '68, '69, '70, '71 and such, then we'd say, well, these people were being monitored internally so that we know they were probably working in radiation areas is what I was thinking.

It might help us sort out whether
these people were not badged but had about the
same jobs or whether they had different jobs

1 during this period of '69 and '70. And 2 internal dose would show a difference during 3 those two years also if they had a change in 4 function. 5 MR. GRIFFON: I think I would leave that as 6 a suggestion for SC&A to consider in how they 7 answer the question of the '69, you know. I'm 8 not sure it's going to be definitive though, 9 that's my --10 DR. NETON: This is Jim Neton. I think if 11 they took people off external monitoring at 12 the uranium facilities is my understanding because the dose potential was low for 13 14 external. But as we learned at Y-12, the 15 internal potential was really the hazard from 16 working with uranium. And so it would be 17 totally consistent to have a routine 18 monitoring for internal continual all along 19 and drop the external, and wouldn't really 20 tell you anything. 21 MR. GRIFFON: At least it could be, so it's 22 not definitive. 23 DR. NETON: Wouldn't cut it one way or the 24 other I don't think. 25 MR. GRIFFON: I think it might be

inconclusive but I'll leave that to, you know, in your response if you want to consider that 3 suggestion. I don't think it, at least in my view, I don't think it's an action right now.

> MS. DeMERS (by Telephone): This is Kathy. In my X-2 Table that I have in the 1969 report, I did give you a yes or no answer as to whether they were monitored May 11th through the end of '69 both in vitro and in vivo.

MR. GRIFFON: Okay, so maybe consider that table that SC&A's provided. Thank you, Kathy.

## COWORKER MODELS

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Okay, can we move to item three, realizing that we're in the afternoon crunch as usual. I do want to stick to closing by 5:00, really. I know a lot of people have planes to catch, and it is a Friday.

So we've got coworker models here, and I think, and I'm not sure how far we can go on this, but it comes up in, the way I bring it up here is in the context of the data completeness related to the two prior items and Brant alluded to this a little bit with the quarterly badging question and the possibility of looking into that and maybe

modifying the coworker model, not committing to it as you said, but the possibility.

But the question of why it was here on the agenda was basically to say, you know, and I don't think we're really here yet because we don't have, you haven't had a chance to fully investigate all of SC&A's report. But if there were gaps, as it was raised before in the meetings I think Jim brought this up, too, that if, in fact, we do find these blanks or gaps and do the coworker models or can the coworker models adequately account for that or be used to reconstruct doses for those gaps. And I think we were thinking both internal and external so I just put coworker models here.

The other reason I bring this up here is this question of pedigree that I've brought up through numerous meetings of the discussion of the coworker models and the HIS-20 versus the CER and then we have this question, and I'm not sure we're ready to discuss this because you're still reviewing these other reports, but the question of if you have blanks that were turned into zeros it raises some questions, at least in my mind, of how

good is this database. How reliable is this database for use as coworker models.

Now we have analyzed this in several different ways. You've looked at CER versus HIS-20 and basically on the internal side you gave us a report that concluded that the intakes would be very similar, equivalent, I guess, over all years. So that that was sort of a bottom line question was even if there's differences, the bottom line is we get similar intakes.

We also have the Donna Cragle
analysis, and I'm laying these things out so
that as we go forward we address these. That
report, that Donna Cragle analysis, that
looked into the question of CER versus HIS-20,
and to me, this raised a problem in my mind
when you look down the years and there's a
table in that report that says the data over
each year and how, and I originally in the
matrix defined it as large discrepancies. I
think there are some pretty large
discrepancies in a number of data points just
looking at the number of data points.

And so the question is if you're

missing two or three thousand data points, but at the end of the day the intakes are the same, I think we need to look at that closer I guess is my challenge going forward as we consider the data completeness if we realize the coworker models are going to be into play more significantly for the Rocky claimants, I think we need to consider those pieces again and make sure that we're comfortable with them. You know, SC&A but also ultimately the Board I guess, that we're comfortable that they're going to be adequate.

And I think there's a lot of ins and outs on this. I'm not sure. I'm trying to remember the original analysis, but the, comparing the CER and the HIS-20 when it concluded that the intakes were the same, I think we want to see what exactly, you know, maybe look closer at that and see what exactly that --

MR. FITZGERALD: And we haven't done that yet.

MR. GRIFFON: Right, right, I don't think that SC&A dug into that.

MR. FITZGERALD: No.

MR. GRIFFON: And it's important to again emphasize the history of this that we kind of didn't look into the coworker model so strongly because we were basically under the impression that very few claimants were going to require coworker models to reconstruct their dose. So I think this is kind of queued up depending on the completeness analysis, and also depending on, I mean, I have kind of a follow-up question for NIOSH.

I think that since these TIBs have been published, the coworker TIBs, I think you've indicated that there were probably more claimants that might require at least partial use of these coworker models. And I don't know if you have any sense of, because when we first talked it was a very small fraction that we thought, and now I think that might have changed or shifted or whatever.

Do you have any sense now of, you know, because if it's very few out of 1,100 or how many ever claimants, then I think we have to weigh this differently than if it's a lot, right?

DR. ULSH: I can't give you exact numbers,

Mark. But what I can tell you is that, you know, and we talked, I don't know, middle of last year at some point. I think at that time we only had just a couple, maybe two or something like that cases on hold because they needed coworker data to finish them.

Now subsequent, and what I said at that time was that, at least I think I said it at that time, was that the need for coworker data at Rocky Flats is not zero, but it's minimal compared to what you see at other sites based on what we've seen so far. Now we were only 700 out of a thousand cases.

But subsequent to that the TIBs were published, the coworker TIBs, and what we have indicated is that after the publication of those TIBs, it's not necessarily that the claims require the coworker data, it's just that that might be the easiest way to do it.

Now there might be other approaches available, but since the coworker TIB is available, and it's easy to use, we use it.

So I can pretty much say with confidence that the amount that those TIBs have been, the coworker data has been used has

1 gone up since publication of the TIBs. 2 still don't think that it's a major usage 3 compared, when you compare like to other 4 sites, but I don't have the numbers for that. 5 I can get those kind of numbers for you. 6 MS. MUNN: Mark, I'm not sure that I 7 understand exactly what you're asking here 8 with respect to, I had been under the 9 impression that once we had the new procedures 10 put together that we were pretty close to some 11 sort of usable data having compared these two 12 databases and knowing where there were any 13 discrepancies that we could move over from 14 there. But am I understanding that you're 15 asking SC&A to do something specific now with 16 this data comparison? And if so, I guess I need some clarification here. 17 18 MR. GRIFFON: I'm asking SC&A, I don't 19 think, I think we had that deliverable with 20 the analysis that concluded that the intakes 21 from CER and HIS-20 were essentially the same. 22 MR. FITZGERALD: That was Joyce's work on 23 the model cases that were --24 MR. GRIFFON: No, no, no, that was --25 DR. MAKHIJANI: That was NIOSH.

MR. GRIFFON: That was Lockamy's, and I'm asking for them to, I don't think that you've ever reviewed that specifically.

## MR. FITZGERALD: No.

MR. GRIFFON: And I think we kind of let that go, and we've just, it's been brought up again, and I just want to make sure that we all are in agreement with that conclusion that, in fact, you know, and also, it's not so much, I think we've been through the coworker models, and I think SC&A's pretty comfortable with the model itself.

Now the question is the data that went into the model, so the construct they're comfortable with, I think, it's the data, whether the data is, you know, this goes back to this data completeness, data reliability question. This question that Arjun raised about blanks being put in as zeros. That certainly skews your data. And then on the other hand the use of the coworker model. How is it being used, implemented?

That gets back to things that Jim brings up like are you, the 95<sup>th</sup> can bound, you know, but I don't know that, so if they're

using the 95<sup>th</sup> in most cases then we'll probably, you know, we have one level of comfort. If you start using the median, then you're worried about these zeros probably a little more. I mean, there's a bunch of subquestions there, I guess.

MR. FITZGERALD: Where they're coming at it from, I think you described it pretty well, coming at it from three different directions. Certainly looking at the concept itself, how the model's constructed. We had quite a bit of discussions on OTIB-38, internal for example. I think we've satisfied that issue. We're pretty far along on OTIB-58 with what Ron's been doing with ORAU.

The other aspect is we're looking at the completeness issue which is kind of catch me up to the coworker review because again, we have to look at that first before we can judge that. Certainly, we talked about that.

The third one we talk about in terms of application. Now you've got this model, you know, how's it going to be applied. And we've actually got into that a fair amount. I think we had discussions on an issue-specific

call that dealt with OTIB-38 and how it could be applied, question at  $95^{\rm th}$  percentile, issues like that.

But we're also looking at questions such as, but on the application side I think we have started looking into that, had discussions on how the models would be applied. And certainly one issue since we now have, had OTIB-49, high fired, you know, how would OTIB-38 apply, certainly, in those cases as well. So we'll look at the application. Those are the three facets.

MS. MUNN: So for my benefit I guess what I'm trying to do is get my intellectual arms, as short as they are, around how large this issue is because I have some concern about this size, the magnitude, of the task we're asking SC&A to perform here at this point in our deliberations. I just --

MR. GRIFFON: Well, right now I think it's to review a report that was on the table a long time ago, but they didn't, but we sort of didn't say, it wasn't a pressing action at that time.

MS. MUNN: So you don't see this as a big

1	thing?
2	MR. GRIFFON: No.
3	MR. FITZGERALD: Well, it's at the point now
4	actually, we've been working at this for
5	awhile on these other fronts, but we haven't
6	looked at that specific question, and we
7	haven't quite finished up with some of these
8	others. But really, we've looked at the
9	construct. We've looked at the data
10	completeness. So we're pretty far along.
11	This doesn't represent a new avenue per se.
12	MS. MUNN: So we really wouldn't be
13	expecting this to be a long-term problem for
14	you?
15	MR. FITZGERALD: No.
16	MS. MUNN: It's an issue that we can hear
17	back on fairly soon?
18	MR. FITZGERALD: I would think so, yes.
19	This isn't the first time we've looked at HIS-
20	20 and CER, but we did it in the context of
21	what work that Joyce has been doing. So we
22	need to look at it a little broader than that.
23	MR. GRIFFON: I guess for me, you know,
24	another troubling piece of this for me is that
25	if you, I mean, every time I seem to look at

1 this I find discrepancies between these two 2 databases, and not one or two or 20 or 30 data 3 points. It's extensive. And to me you start 4 to wonder. I wonder about the validity of 5 either one. And we're looking at it saying, 6 okay, we're getting a similar result at the 7 end of the day so don't worry about it, you 8 know? It makes me a little uneasy of being 9 struck down. 10 MS. MUNN: I guess the basic question still 11 hasn't changed. The question is still how 12 good is good enough. Because certainly we're 13 not ever going to get absolute correlation 14 between any two databases anywhere as long as this --15 16 MR. GRIFFON: This is a half full and half 17 empty issue. 18 MS. MUNN: Yes, it is. It is. 19 MR. GRIFFON: How bad is too bad. 20 the other side of it. And I mean, I just, 21 again, I've looked at some of this. Some of 22 this comes up in the log book review. 23 looked into some exposure IDs that come out of 24 the log book analysis, and when you track them 25 back, I almost found like every time I was

finding exposure ID in the 100,000 series that didn't exist in HIS-20.

So this is very odd. It seems like trends like this keep popping up, and it's not one or two individuals. I did a little minianalysis looking at how many exposure IDs, NIOSH provided us with a spreadsheet with exposure IDs for individuals so that we can link that to the ACCESS HIS-20 database.

And if you compare, you know, I just saw this, and this is, again, no scientific sampling method, but I just sort of saw this trend of individuals I was finding in the log books with six-digit IDs starting one-zero-zero-zero-zero. And when I looked, they just weren't in HIS-20. So then I said, well, how many individuals are in the exposure ID Excel sheet versus how many are in the ACCESS database. And it seems like there's about 15 percent as many in HIS-20 as were in the exposure IDs.

Now maybe there's a good reason for that. Maybe those people were not in the plutonium areas or whatever. It certainly wasn't by year as far as I can tell. But

again, it's just this, you know, adding up these pieces when you have these kinds of discrepancies in the data it just makes me a little uneasy. And that's why I'm saying at least one sub-test for me I think is for us, we need to go back to that old report that we kind of went past in a work group meeting and

DR. NETON: Well, there's two things, Mark, one is you're talking about comparing the CEDR and the HIS-20 for comparability of film sets to internal intakes.

## MR. GRIFFON: Yeah.

DR. NETON: But then I think, I know Brant and I had talked, and he had done an analysis where similar to what was requested for us to do at Y-12. He went back and looked at the environmental reports from the health physics monthly reports. He compared the numbers, and I haven't looked at it for awhile, but I thought it was a pretty good comparison so at least the numbers that were being reported in the monthly reports were matching up with what was in HIS-20. And I think we better take a look at that and see --

1 MR. GRIFFON: Yes, and he just told me 2 about, I think I didn't get that e-mail for 3 whatever reason, but --4 DR. NETON: There were some discrepancies, 5 but I think it would be useful to look at. DR. ULSH: What I found was that --6 7 MR. GRIFFON: That would be useful to look 8 at though because that's what helped us in Y-9 12 was looking at that summary sort of --10 DR. NETON: We did that in Y-12 and it 11 seemed to satisfy people a little bit. Let's 12 go down that path. 13 DR. ULSH: Jim, I haven't looked at it 14 probably since the last time you looked at it 15 but just going from memory, just to summarize 16 what was there, it started, the progress 17 reports that I had covered the timeframe 1952 18 up to 1971, I believe. Again, all of this is 19 approximate because I'm going from memory. 20 There was one year in that period 21 where I didn't have the progress reports, so I 22 didn't do an analysis for that year. 23 was another year where I had the monthly 24 progress reports for January through November 25 so I, there was another year where I had the

1 progress reports for January through November. 2 I was missing December so I made an 3 extrapolation there. With those caveats I 4 think what I saw was that the average 5 difference between a number of bioassay points 6 in CEDR versus the monthly progress reports 7 was about five percent. They were within five 8 percent of each other. 9 MR. GRIFFON: Of CEDR versus the monthly 10 progress or HIS-20? 11 DR. ULSH: CEDR versus the monthly progress. 12 No, CEDR versus the monthly progress reports because recall --13 14 MR. GRIFFON: Because you were using CEDR, 15 right, for your --16 DR. ULSH: Well, yes, that's one reason. 17 But the other reason is that we know that in 18 HIS-20 some individuals' data is not in HIS-20 19 because they terminated employment prior to 20 That issue is there. So what I saw with 21 the CEDR, you know, I don't want to make too 22 much of this comparison. I mean, you can't go 23 beyond what the bounds of the data are, but 24 over that time period there's very good 25 agreement, at least I think there's very good

agreement, within five percent on average, between CER and the progress reports. Now that's an average over the years, but I will send that report out to you again and hopefully that can go partway towards --

MR. GRIFFON: And the entire work group I guess.

Did you get the e-mail?

DR. ULSH: I don't remember when I sent it.

It wasn't recently. It was at least a couple of weeks or maybe more.

MR. GRIFFON: At any rate that's a new piece of information. So that's helpful. I mean, any prong we can come at to answer this question I think is helpful. I think we're down to a weight of the evidence. We've got some discrepancies, but how bad are they or how small are they, and can they be overcome by the fact that we have some assurance that the, you know, it won't affect the intake estimates or things like that, the bottom line sort of thing. So if SC&A can look at both this report that Brant's discussing and that prior report by Lockamy, the earlier report, I think that would be helpful.

1	MR. FITZGERALD: Do you recall if the
2	Lockamy report had the spreadsheets or the
3	background stuff associated with it or not?
4	But that might be something.
5	DR. ULSH: I'm not sure what you mean in
6	terms of
7	MR. FITZGERALD: The Lockamy report itself.
8	DR. ULSH: Yeah, I know that, but what
9	MR. FITZGERALD: In terms of the back, the
10	supporting data.
11	MR. GRIFFON: Did he have his analysis in
12	there or was it just a summary of what he
13	MR. FITZGERALD: I recall the analysis
14	itself.
15	DR. ULSH: I'll have to go on the O drive
16	and look because there's a folder here for
17	that.
18	DR. LIPSZTEIN (by Telephone): Let me just
19	say something?
20	MR. GRIFFON: Yeah, Joyce, go ahead.
21	DR. LIPSZTEIN (by Telephone): I think,
22	well, first of all two things. One, we are
23	going to the 95th percentile, and I think we
24	favor any of the possible problems,
25	uncertainties that we have (unintelligible).

1 And other thing is that this is a model for 2 like for example, there was (unintelligible). 3 It's just one intake rate from '52 to '61, and 4 of course, there's a big difference between 5 one year and another year, like for example 6 from the first year to the second year there 7 are three times difference from the intake. 8 The intake is what's calculated in those 9 They have a lot of difference from one 10 year to the other. 11 MR. GRIFFON: But Joyce, I think this 12 comparison document that we're talking about, 13 I think he looked at year by year. I think 14 there's enough --15 DR. LIPSZTEIN (by Telephone): This is year 16 by year, but when they come to the model, it's 17 a model, so it's just one intake for ten 18 years. Then there's another intake that was 19 calculated for seven years. So --20 MR. GRIFFON: But I thought the application 21 was that --22 DR. LIPSZTEIN (by Telephone): What I mean 23 is that some uncertainties on the database wouldn't make too much difference in a model 24 25 like that, but I think it's acceptable when we

1	go to the 95 <sup>th</sup> percentile.
2	MR. GRIFFON: When you're at the 95 <sup>th</sup> , right.
3	DR. LIPSZTEIN (by Telephone): Because, you
4	know, it's not exact anyway. You know, it's a
5	lot of estimation. Imagine just one intake
6	for ten years.
7	MR. GRIFFON: Yeah, I think, I know what
8	you're saying, Joyce. I think we should at
9	least look at that prior piece though and look
10	at the
11	DR. LIPSZTEIN (by Telephone): Okay, okay.
12	MR. GRIFFON: And I think this Health and
13	Safety Report thing might be very helpful,
14	too. Did that have internal and external or
15	just
16	DR. ULSH: No, just bioassay.
17	MR. GRIFFON: Just bioassay, okay, just
18	curious.
19	MS. MUNN: But there's such a wealth of data
20	here that it's hard to imagine in light of no
21	evidence of programmatic error that we can't
22	resolve this.
23	MR. GRIFFON: That's why we're following
24	this up because we've got the, at least
25	potential that some blanks were zeroed out in

1 the database. So that's evidence of programmatic problems. I'm not saying 2 3 intentional or otherwise. I'm just saying, 4 you know. MS. MUNN: Yeah, but if you're constructing 5 6 a dose then, well, that's --7 MR. GRIFFON: I'm not saying it's 8 insurmountable either. I'm not, but I think 9 we're, all I'm saying if we can be looking at 10 these things in parallel, SC&A can look at 11 that while NIOSH is finishing their data 12 completeness response, then at the next 13 meeting we'll be ready to maybe discuss and 14 the application of those coworker, you know, 15 whether they're okay and if SC&A has any input 16 on how they should be applied, you know, that 17 sort of thing. 18 DR. MAKHIJANI: I have a question on this 19 one number, the 95 percentile for all years. 20 What happens if a worker worked for just one 21 year? What do you do with that in terms of a 22 claimant favorable number? And if he happened 23 to have worked in the years where he had --24 DR. NETON: He would get the intake for that 25 one year, but if you're asking what Joyce is

1	alluding to which is these differences, I
2	don't know.
3	DR. MAKHIJANI: Would you use the number for
4	that year or would you use the same coworker
5	model number that you're using for one number
6	for ten years?
7	MR. GRIFFON: This is sort of an application
8	question?
9	DR. NETON: Mutty Sharfi might know.
10	MR. SHARFI: Then you'd likely go back to
11	the raw coworker bioassay data and then
12	construct its own coworker intake rate based
13	off the coworker bioassay data. You wouldn't
14	use those broad long-term periods.
15	MR. GRIFFON: The one year that they were
16	employed, right?
17	MR. SHARFI: You would look at the more
18	refined periods.
19	MR. GRIFFON: Seems to make sense.
20	MR. SHARFI: You can refine the coworker
21	intake numbers to fit the specific case
22	scenario.
23	DR. MAKHIJANI: Oh, great, okay.
24	MR. SHARFI: You're not locked into those
25	numbers.
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DR. LIPSZTEIN (by Telephone): That's not
what's said on OTIB-028. Actually, it states
that but I can't
MR. GRIFFON: Well, wait, I think we're
getting into the application question. We can
talk about this more. I don't think right now
is the place, Joyce.
DR. LIPSZTEIN (by Telephone): Okay.
MR. GRIFFON: I think the sense I had was
that the application is you use that broader
average and then you have sort of an if-then
tree. And then you might go back to the
annual data if you need to, right.
MR. SHARFI: That's why the tables are
provided in the appendix of the OTIB is that
if you need a more specific for the case you
can.
MR. GRIFFON: Then you go annual, right.
MR. FITZGERALD: Say you had a number of
questions that sort of get into how would you
apply it because I think that's a
MR. GRIFFON: Application question, yeah.
So I mean, I think we're there on the
actions we need on the coworker stuff and that
may be very helpful, the latest report.

The

1 MR. FITZGERALD: And I think these are 2 pretty narrowly defined, and I think we can 3 again get that. 4 OTHER RADIONUCLIDES - THORIUM 5 MR. GRIFFON: I think we're on to item four, thorium, other radionuclides, thorium, I think 6 7 is what we're down to. And I know that there 8 was a technical call in between meetings, and 9 I apologize for not having that in my 10 Blackberry and missing it. But anyway --11 DR. MAKHIJANI: Do you have the minutes from 12 that? 13 MR. GRIFFON: Yes, the minutes were sent 14 around to, I think, you got those as well, Wanda? 15 16 MS. MUNN: Yes. 17 MR. GRIFFON: So anyway you can summarize 18 where, Arjun or Brant? I don't know who wants 19 to start. 20 DR. ULSH: As Mark mentioned we did have a 21 conference call about a week and a half, two 22 weeks ago, something like that, on thorium 23 issues. Just to bring you up to date with 24 where we are, you know, we've been looking at

the thorium issue for months now I think.

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latest document that we've put out was

December 27<sup>th</sup>, and after that SC&A issued some

comments and evaluation of that report. That

was in early January along with the others I

think. The topic of the conference call,

there were two main topics.

The first was the question of magnesium alloy which contained up to three percent thorium according to the workers who were involved. There's a long story involved with this and it pulls in another site, the Dow Madison site in Illinois. I was going to say Wisconsin, but it's Illinois.

And there's an active SEC petition at that site, and there were some interviews conducted not by NIOSH but by some of the petitioners involved in that site with Dow Madison workers. And there was a question brought up about, well, first of all it was whether or not thorium was shipped from the Dow Madison site to Rocky Flats and/or vice versa. We later clarified that we were talking really about magnesium-thorium alloy.

And the Dow Madison worker who talked about this said it was up to three percent.

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And there's still some questions I think, at least there were at the time of our conference call, about to what extent this might have occurred.

We presented at the conference call, I think we sent out the interview notes after the conference call, but we talked to four or five, I don't remember which, Rocky Flats workers, and they did not recall large uses of magnesium-thorium alloy at Rocky Flats.

They did recall one specific use of the magnesium alloy, and it wasn't clear whether it was magnesium thorium or not. And that was in the pennates (ph) in the conveyor line in Building 776, I believe, the one where the Mother's Day fire occurred. That was it. That was the only big use of magnesium alloy that they were aware of.

And I should qualify that. It's not big, but we're talking about maybe a couple of hundred pounds according to one worker's estimate. That was really it. I mean, we didn't see, the workers didn't recall any program, large scale program to use magnesium alloy at Rocky.

Let me see, I'm trying to think of all the other developments that have occurred. We have since learned, we've received some documentation that tells us the source of the thorium that came into Dow Madison. It was from Canada, and that was with pellets.

Ingots came from England I think is what the workers said, and we see nothing to dispute that. So it doesn't look like the source of the thorium going into Dow Madison was Rocky Flats.

But that doesn't speak to the question of whether or not the destination was Rocky Flats. Again, the four workers that we've talked to, and these are folks who were involved with the Operations Board at Rocky Flats. I might have the names of those committees wrong, and I don't know, maybe you can correct me on that.

But the folks that were in charge of all shipments of radioactive materials that came into the site and then also the Operations Committee, they didn't have any recollection of large scale use of magnesium-thorium alloy at Rocky. And in terms of the

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pennates that they did know about, they did state specifically that those were prepared by an outside vendor and delivered to Rocky Flats.

So that was one issue. I think that's pretty much where we left it. We were going to get the notes to SC&A, and we've done that. And that's pretty much where we left that issue on the conference call.

Now, Arjun, do you want to talk about

that before I move on to the other topic? DR. MAKHIJANI: Yeah, I have the same document that you have, I think, regarding the thorium pellet supply from Canada. I just wanted to note that in the amount of thorium we had at Dow Madison for this alloy program was quite large, 80 tons up to 1960. So that would mean we got about 2,400 tons of alloy, three percent thorium. So they were fabricating making the alloy on a very large scale at Dow Madison. Of course, we don't know where it went, but as I understand, I was talking to Tony at lunch about whether they did fabrication of things, you know, like trays and other parts at Rocky Flats. And I

do understand that ad hoc or custom parts were fabricated at Rocky Flats.

Correct me if I'm wrong, Tony, or misinterpreting our conversation.

MR. DEMAIORI: No, absolutely, we did a lot of fabrications at Rocky Flats. We built our own part carts with the lead shelvings. We used stainless. We used all kinds of different materials. We had our own fabrication baths for the trucks to ship the nuclear components themselves. I mean, we armed them. We put the bulletproof glass in them. We did all that. We had the hobby shop where they did some secret fabrications in there.

This issue of thorium, as a rad tech I can tell you that we had thorium in almost every toolbox on the plant site, that's in the way of welding rods. So welders commonly use thorium rods for the different welds that they were making. You know, pretty much we did our own thing, but yeah, we did a lot of fabrication out at Rocky Flats.

Now when I was there, and that was late `70s, 1979 and on, it was mostly

stainless steel because it was very acid resistant. But we definitely built our own carts and did that sort of thing. We built a lot of things that they used at Rocky Flats. And so, you know, I don't know on this magnesium-thorium alloy, but I can find out. I know a lot of machinists that would do the fabrication, and you know, I can definitely check into that.

The people who did the shipping and receiving, as far as shipping and receiving went, we got a lot of things into Rocky Flats that we didn't expect to get. Talk about americium (unintelligible) all the salts from the United Kingdom were extremely hot. So nothing is perfect, but I can definitely tell you that we did a lot of fabrication out at Rocky Flats. That's all of our research and development, 779, on the hot side, and then we would take 887 on the cold side. They did all the beryllium operations, all the R&D for that. We fabricated everything.

MR. ELLIOTT: But the numbers that Arjun quoted on thorium production at Dow Madison, we know, I think it's a common understanding

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here that Dow Madison had other work, large thorium commercial component that they performed. And much of that was dedicated to Department of Defense effort. So I'm only concerned here about what we can say about the production for the AEC portion and where that went. I don't want to get the two confused.

DR. MAKHIJANI: No, I don't want to get them confused, and it is correct that the documents that you sent to Senator Salazar's office, which I have here, they do indicate that magnesium-thorium alloy was I guess in large measure, in some measure at least, being considered for like aircraft parts and things like that. And the other document we have about thorium-magnesium alloy, the NUREG-1414 also said the same thing, that that was --1770, sorry -- so I would agree with that. It's just the extent, my only question, reason for raising that is the extent of use at Rocky Flats is, well, based on the recollection of talking to a few people. And there's not a definitive trail of what happened when it started, when it stopped, how much, and most importantly, whether it was, there was any

fabrication work there. We understand from 1 2 these claimants that there was no, they were 3 not fabricated on site. 4 DR. NETON: We don't even know that material 5 was shipped to Rocky Flats from Dow Madison. 6 That's not a foregone conclusion. 7 that's speculation. 8 MR. ELLIOTT: We have no indication in the 9 documentation that we have obtained from DOE 10 thus far there was any transfer of thorium-11 magnesium alloy from Dow Madison and Rocky 12 Flats. We haven't seen that. 13 DR. NETON: I'm just wondering if we're 14 chasing down something that might not have 15 happened. I mean, I think we just need to 16 establish the material was actually shipped 17 there before we start --18 MR. ELLIOTT: We don't doubt that there was 19 fabrication at Rocky Flats, but I don't 20 believe we have any indication that says the 21 processes at Rocky Flats required fabrication 22 of thorium-based metal parts or et cetera. 23 we're still interested in understanding how 24 thorium may have been introduced in the site. 25 But to date we have no indication in the

1	documentation we have that there's a large
2	thorium component here.
3	DR. NETON: Thorium alloy, component
4	magnesium-thorium alloy.
5	DR. MAKHIJANI: Yeah, there obviously was
6	some thorium alloy at, in the form of welding
7	rods now
8	MR. DEMAIORI: Oh, absolutely.
9	DR. MAKHIJANI: in the form of magnesium-
10	thorium alloy. There was some presence at
11	least according to interviews that you've
12	done.
13	DR. ULSH: There was magnesium alloy
14	present. We don't know if it was magnesium
15	thorium.
16	DR. NETON: And that was manufactured off
17	site.
18	DR. MAKHIJANI: I didn't understand that
19	from our conference call. I thought that your
20	reference to magnesium alloy was magnesium-
21	thorium alloy, from our conference call.
22	DR. ULSH: Here's the minutes. All* also
23	stated that magnesium alloy was used in the
24	Building 776 line. The pennates were made of
25	magnesium alloy. Now we don't know whether

that was magnesium-thorium alloy. It was magnesium alloy. We also know from the chemrisk report, task one, that went through an extensive list of like 300 pages, I think, of all the chemicals and all of the materials on site.

They talked about magnesium in terms of magnesium salts and small quantity of magnesium alloy. Now it's not clear whether it's magnesium thorium, just magnesium alloy, 0.5 kilograms in the chem-risk report. And that was at that time. So this was not identified in the chem-risk report as a material of concern.

And then we have the testimony from the four workers, which this actually came up I think at the last working group meeting.

And what we committed to do or what we were asked to do was to go, Mel suggested that he would go talk to these folks, and that's what he did. He told us that they had no recollection of, certainly not extensive use of magnesium-thorium alloy or magnesium alloy at all.

DR. MAKHIJANI: I'm concerned in reviewing

the material unless the working group desires otherwise, I mean, we've said what needs to be said in terms of raising the questions about the source term. And some new questions have been introduced and a new source term, you know, there were two items in the December report.

And subsequent to that the 80 ton ingots and the magnesium-thorium alloy, and we know the 80 ton, 80 kilogram ingots, sorry about that. So I don't know where, I mean it's the pleasure of the working group whether there's anything further to be done.

MR. GRIFFON: I think we've left it with two questions kind of. One was on the source term side, and the other was this question of empirical models. Have empirical models be developed. So on the source term side, I don't know, the question on the ingots raised my, I mean I guess what NIOSH concluded was that the latest report that gives more information on those ingots, the amounts were consistent with the earlier mass balance review. Is that fair?

DR. ULSH: Yes.

MR. GRIFFON: I'm trying to remember myself.

And then on the other side the assembly, the mock-up assembly, sort of operation, it seemed to be that those would have been smaller uses certainly, and probably less exposure generating tasks. I'm not sure.

DR. ULSH: We're still talking about that.

I don't want to put words in SC&A's mouth, but
--

MR. GRIFFON: Well, just on the source side though, smaller amounts I guess.

DR. ULSH: Yes. As I mentioned in this topic-specific conference call, there were two major topics. One was magnesium-thorium alloy, and number two was the concerns that SC&A's expressed about our empirical approach to estimating possible bounding thorium intakes. That was topic number two. Before we move on to topic number two, I guess I'd like to get your feel on like what Arjun said, what comes next with the magnesium alloy? Are you okay with where we are? Do you want to see some more specific actions? Or what's the pleasure of the working group I guess.

MS. MUNN: Certainly, this working group

member thinks we've beaten the magnesium alloy thing to death, and I see, I don't know where we could go from there. It appears to me that we've researched whether we did, in fact, have magnesium alloys, and we've identified four individuals who had total plant oversight at one time or other and were part of the Operations Board, all of whom would have known of any extensive use of magnesium alloys. They all had the same general response. Unless there is some significant issue with respect to magnesium—thorium alloy specifically, I can't see that we can take magnesium much further. It seems clear to me that it's not a major issue.

MR. GRIFFON: Magnesium-thorium alloy would be the question of concern. But, yeah, I can't, you know, short of additional information, which I don't see on the horizon, I don't think there's much we can do with that. And I think all the evidence, I mean, from what I can see so far and this is just my personal opinion right now from what I've seen, is that it doesn't, you know, it looks like if any was shipped we can't find any

1 evidence that there was any significant 2 fabrication or with that material at Rocky. 3 So from an exposure standpoint I don't 4 think it's worth chasing, so to speak, unless 5 other clear evidence is brought before us. 6 But I don't think there's any further action. 7 Now, Arjun, what's your sense of that? 8 It's sort of up to the work group. 9 DR. MAKHIJANI: Yes, it's entirely up to 10 We raised the question. NIOSH followed 11 up and did the expert interviews, and I think 12 13 MR. GRIFFON: So that's my feeling on the 14 magnesium-thorium alloy is that I don't know 15 that we're going to, you know, we've got some 16 statements. We've tried to check them the 17 best you could. You've come back and we don't 18 have any indication of, certainly no 19 indication of fabrication at the site and very 20 little indication of any use, and limited if 21 any use, you know. So I think we've chased 22 that about as far as we can at this point. 23 MS. MUNN: Pretty well documented. 24 MR. GRIFFON: The only other thing is maybe 25 if Tony does get back to us from talking with

some of the former workers that have knowledge of this then that's new information before us. So I think we have to address that. But short of new information I think we've chased it as far as we can at this point.

Now were you going to, I mean, on the other source term side before we get into the techniques for bounding, I guess on the other source term side I was looking at some of the Health and Safety reports as were probably some of the ones you were looking at for the database completeness question. In some of those reports they talked about developing a thorium urinalysis program.

And they did talk about some limited air sampling data. You might have seen some of these same reports. So it made me wonder if any of these reports, because you've gone through certainly more than I have of these, if any of these reports shed further light on a more extensive operation with the ingots, with the rolling, with that kind of thorium work.

DR. ULSH: They do shed some light on that,
Mark. It was in the, yeah, I think it was

those same progress reports that you mentioned that you do see occasional mention in 1960 that they were attempting to develop methods for thorium urinalysis. And that was in advance of that ingot project in 1960 that was covered in that report.

In fact, in our December 27<sup>th</sup> report we talked about all of the thorium bioassays. It was easy because there weren't many. They were developing those things for the ingot operation. There were a couple of individuals who I think one of them wound up, they concluded that he had worked in a thorium refinery prior to Rocky Flats, and that's why they saw, you know, I think they saw some thorium in his urine.

MR. GRIFFON: Yeah, I remember that exact distinction, yeah.

DR. ULSH: So that was some of the urinalysis results, maybe two, three, I don't know. There was another individual that came up with a funny peak in his, or an unexpected peak in his whole body count. Bob Bistline wrote a report on that one, and it turned out that it was not related to thorium.

And then there were a few confirmatory bioassays taken related to the ingot, thorium ingot project. And that was pretty much it. That's all we've seen with regard to thorium bioassay. So I don't think that those would speak to a wider thorium use program than what we've already identified.

MR. GRIFFON: And those reports where they discuss the urinalysis program, those timeframes are consistent with what you saw -- DR. ULSH: Yes.

MR. GRIFFON: -- as the thorium project in other documentation.

DR. ULSH: The thorium project, the main document that we had describing the ingot project was authored by a guy named Callabra (ph). He wrote an extensive report on that operation, and it gave exact times of when it happened.

And we also found notations on this project in the Kittinger logs of the time. So we know when that ingot project happened. It happened on eight working days spread out over the latter part of 1960s. And this development work that you're talking about

1 where they talk about the progress reports, 2 that was earlier in 1960. They were trying to 3 develop that in advance of that project. 4 MR. GRIFFON: And where did this ingot 5 processing happen? What building was that in? 6 Do you recall? 7 DR. ULSH: Primarily in Building 881, but I 8 think they did some acid, processes that 9 involved acid. I think etching maybe in 331 10 maybe, one of the 300 buildings, 331, 334. 11 That was also laid out in the Callabra report, 12 but primarily in Building 881. 13 And we also saw, that was consistent 14 with what we saw in the chem-risk test three 15 report. It talked about the uses of thorium 16 and where it occurred. And that was 17 consistent with what was in that report as 18 well. 19 MR. GRIFFON: And then the assembly, the 20 mock-up assemblies? 21 DR. ULSH: Okay, now, that's a second 22 category of operations. So in terms of the 23 quantity of thorium that was used at Rocky, 24 that was primarily in that ingot operation. 25 They had three ingots of the dimensions, they

1 were 12 inches by 12 inches by three inches. 2 So we're talking, physically anyway, they were 3 fairly small ingots, and there were three of 4 them. That was about 240 kilograms, and 5 that's a majority of the thorium inventory 6 that we saw in ledgers at Rocky Flats. 7 Now in terms of the number of 8 operations, there were occasions when Rocky 9 Flats would receive these finished parts from 10 Oak Ridge to use in their models. 11 MR. GRIFFON: That was (unintelligible). 12 DR. ULSH: So that's a separate operation 13 there. 14 And then you had smaller uses of --15 MR. GRIFFON: We don't know the extent --16 I'm sorry to interrupt -- we don't know the 17 extent of, I mean, that one memo suggested 18 that those uses were sort of below what would 19 have been recorded on the mass balance. Although Mel suggested, or Bryce, I forget who 20 21 had suggested, that if the building exceeded a 22 certain amount, it would have been rolled into 23 that mass balance summary. Is that accurate? 24 Am I off-base on that? 25 DR. ULSH: Well, you might be talking about

1 yet another, there were, I think, four uses of 2 thorium that we've identified. One was ingot. 3 Two was the use in weapons mock-ups. 4 was the use in laboratory standards. 5 MR. GRIFFON: Now I think I'm talking about 6 two, the mock-ups. 7 DR. ULSH: The mock-ups. 8 MR. GRIFFON: Where they said that they 9 would have been, I think the one paragraph in 10 that memo it was the history of thorium use, I 11 think, at Rocky Flats, in that memo. 12 DR. ULSH: Okay, the Bob Bistline report. 13 MR. GRIFFON: Yeah. And it indicated in 14 that one paragraph that currently I think it 15 said seven kilograms onsite, but it might have 16 been a 0.7. It was a blurry copy that I was 17 looking at. 18 DR. ULSH: Yeah, I think that was 19 cumulative, totally. 20 MR. GRIFFON: But anyway, it suggested that 21 earlier period would have had more but still 22 each individual use was below what would be 23 reported on these forms. I don't recall the 24 exact language. I'd have to go back to the 25 reference, but --

**DR. ULSH:** I don't either. I don't have that in front of me.

MR. GRIFFON: My question was how extensive was that operation. Are we talking about a lot of small uses that when you aggregate them, and how long did it go on sort of that -

DR. ULSH: Well, in order to answer that definitively, in other words, we could get some indirect evidence from the MDA ledgers in terms of we could look at when the inventory changed and get some indication of when they might have received a part from Oak Ridge, and when they might have sent them back. Then we can, if it's the working group's pleasure, then we can do that.

In fact that's one of the concerns I think that Arjun's report talked about was that we have reported inventory numbers and not through-put numbers. Now what -- I did not actually do this. Mel and Bryce did and Mark Rolfes. What they did was they reported the highest inventory for the year. In order to get, and what Mark told me, he looked at these ledgers, was that there were very few

times when the inventory actually changed.

But we can get those numbers if --

MR. GRIFFON: I don't know that we necessarily need them. I thought maybe from your interviews you can give us the scope of -

DR. ULSH: It was not, all the interviews that we have conducted have indicated, number one, that this was a very unusual occurrence. I don't even want to hazard a number, but it wasn't like an everyday occurrence. It was special order work, and it was not extensive.

Now we have talked to, regarding these uses in weapons mock-ups, we've talked to four R&D machinists just recently, and we're about to send out these interview notes to everyone because SC&A has expressed continuing concerns about how we're going to bound the dose of this particular use of thorium. So we went back and talked to these machinists.

And I think we need to be very clear here that we don't confuse the machining operations that occurred when the only indication that we have that these parts from Rocky Flats were machined was one interview

1 that we conducted with a site expert, and he 2 said the parts might have been lightly 3 trimmed. 4 MR. GRIFFON: Parts from Rocky Flats or from 5 Oak Ridge? 6 DR. ULSH: From Y-12. 7 MR. GRIFFON: From Y-12, you said from Rocky 8 Flats. 9 DR. ULSH: Oh, I'm sorry. 10 MR. GRIFFON: From Y-12, I'm just trying to 11 stick with it here. 12 DR. ULSH: Yes, they received the parts at 13 Rocky Flats from Y-12. And that individual 14 was not a machinist. He was very 15 knowledgeable at the site. He was one of the 16 people that we talked to about maybe using 17 thorium alloys. We had a general picture of 18 where things were at the site. 19 But we have talked to four R&D 20 machinists. These guys worked in Building 991 21 where this operation would have occurred, and 22 none of them could recall ever actually 23 machining these parts. I mean, the problem is 24 I can't say to you with 100 percent certainty,

well, they never ground off a high spot.

25

1 I can't say that because I don't know 2 that, but that's the only indication that we 3 have that they did anything other than take 4 them out of the box and bring them 5 (unintelligible) so far. And I think it's fair to say that NIOSH and SC&A have not yet 6 7 reached closure on how to bound doses for that 8 particular operation. Fair enough? 9 DR. MAKHIJANI: Yes. 10 MR. PRESLEY: Are those interviews for those 11 machinists going to be seen? 12 DR. ULSH: I hadn't planned on it, Bob. 13 can show them to you. I've got a hard copy 14 here. They don't go into detail, I mean, it 15 was pretty much, it was very focused. And it 16 asked do you ever recall machining these 17 parts, something like that, one question. 18 MR. PRESLEY: Okay. 19 DR. ULSH: I'll talk to you afterwards about 20 that before we send it out. If there's any 21 concern at all, we don't want to cross any 22 lines. 23 MR. DEMAIORI: What year was that --24 DR. ULSH: The ingot operation or the parts? 25 MR. DEMAIORI: Parts for the machining.

DR. ULSH: I can't really tell you, Tony. I don't know exactly without going back to get a more detailed look at the MDA ledgers. It was early on in Rocky Flats history, but I don't know the exact years.

MR. PRESLEY: Late '60s.

MR. GRIFFON: Nineteen-sixties?

MR. PRESLEY: Mid-to-late '60s.

MR. DEMAIORI: Because most of the machining R&D was 887. So I was just wondering. They may have not built 887, you know, in the late '60s. But when I was there, the bulk of the R&D machining was 887.

DR. ULSH: Yeah, this is certainly in the earlier years, in the '60s, maybe in the '50s. I don't know exactly when. In order to answer that question we would need to go back and get a look at the, closer look at the MDA ledgers.

There were two other thorium activities, thorium strikes, and what that involved was some special order work that Rocky Flats conducted with uranium-233. And that uranium-233 had trace contaminants in the beginning 50 ppm and later on down to seven ppm of U-232 which daughter products of which

1 are thorium-228. 2 And so in order to work with this 3 uranium-233, they had to remove those thorium 4 daughters. And this occurred on -- well, I'm 5 looking at the chem-risk report right now. 6 Twice during the '64 to '69 time period, but 7 there were some other ones later. Again, not 8 a common occurrence but a handful of 9 occurrences. 10 And unless you look at this you may 11 not know, but there is a very great external 12 exposure hazard with this kind of an 13 operation. We have seen notations about this 14 in the Kittinger logs and describes when it 15 occurred, and that they did cover it with 16 health physics support. It was a small 17 operation in terms of the number of people 18 involved. 19 MR. GRIFFON: Do you have a sense of the 20 people, how --21 DR. ULSH: Pardon me? 22 MR. GRIFFON: How many people? 23 DR. ULSH: A dozen at most and that's 24 probably an overestimate.

But they removed the thorium

25

daughters, the thorium-228, and the daughters from the uranium-233 before they processed the U-233. Now for those operations we have proposed a NUREG-1400 approach. But again, SC&A's expressed some concern about that approach, and we are currently considering their concerns about that.

And finally, we have these various miscellaneous, very small uses of thorium like in laboratory standards, that kind of thing.

I think that covers the four categories, right, Arjun?

## DR. MAKHIJANI: Yes.

MR. FITZGERALD: The only clarification I would add, I think the way we had left it because the concern was the semi-empirical approach I think from our standpoint didn't demonstrate conservatism necessarily, and I think the response was to consider a, perhaps a bounding analysis for the three activities other than the ingots that were, you just talked about. I think that's the way we left it there at the end, and I think there was some agreement that you would look at that.

Is that still the case?

1 DR. ULSH: That is the case, Joe. 2 the way I recall it, too. But I think it was 3 not SC&A's position that they couldn't be 4 bound. It's just that you were not yet 5 convinced with any bounding analysis that we 6 had yet presented. 7 MR. FITZGERALD: Yeah, I think that was it. 8 MR. GRIFFON: So that's an outstanding 9 action. 10 DR. ULSH: That is an outstanding action. 11 MR. GRIFFON: And what about, you said other 12 than the ingot operation? MR. FITZGERALD: Yeah, I think we were 13 14 focusing, I think the conclusion of the review 15 that we presented was that we were okay with 16 the ingot operation in terms of the monitoring 17 that was done, but the other three where 18 NUREG-1400 would be essential, those were the 19 three that --20 MR. GRIFFON: What is the proposal? 21 probably told me this before, but what's the 22 method for the ingots for dose reconstruction? 23 If they were involved in that operation, how 24 do you reconstruct their dose? You have some 25 urinalysis --

DR. ULSH: We did have, okay, there was some urinalysis, more confirmatory-type urinalysis to show that intakes didn't occur. There was, basically, our approach was to show that there was no significant intake potential for that job.

MR. GRIFFON: Well, that's what I wanted to, the Health and Safety reports that we're both looking at apparently, what I read is they do have references to air sampling. And it wasn't clear, obviously, we run across this again, it wasn't clear. I think one was over -- well, I don't want to quote numbers, but it was high, but it said an operational sample so it's not clear if it was in the rolling area or it wasn't probably in a breathing zone. Then they gave an average of for people in the area I think it said, a reported number said 20 or 30 percent of the mpl on average for the --

DR. ULSH: If we're thinking of the same air sample, and I think we are, they did have the location listed on the air sampling card where the air samples were taken. One was taken like three feet from the ingot. Another was

taken by --

MR. GRIFFON: Okay, I didn't see the air sampling cards themselves, so maybe I --

DR. ULSH: Well, our report considered that and we had the detailed analysis of the job in terms of it was almost an hour-by-hour blow of the whole project. So we calculated, well, we felt at least, the maximum credible intakes.

I don't know. Arjun?

DR. MAKHIJANI: Well, we didn't evaluate the question of whether there were maximum credible intakes or not. We looked at the documents and saw that there were air monitoring data, and there were some high results, did not evaluate whether they were breathing, you know, they were area samples if I remember correctly.

We haven't looked at whether the model provides a conservative dose estimate, we just did note that there are air monitoring data and some bioassay sample data of which, and the bioassay sample data probably used to bound the dose with a minimum --

DR. NETON: That's pretty high because --

DR. MAKHIJANI: Just from the point of view

of an SEC issue, and I just want to say what we did in saying the dose reconstruction is feasible, but I think we know that that we haven't actually examined the details of --

I know that there was a calculation in your December 27<sup>th</sup> report, but we haven't critically evaluated that calculation to sign off that we would agree that that's the appropriate method to use because there are bioassay sample data that I think would be used to bound the dose. And I think Jim Neton is agreeing with that.

DR. NETON: Certainly, yeah.

DR. MAKHIJANI: Now whether the issue of what should be done, and what would be appropriate would be a separate task. And certainly we could do that, but we haven't done it.

MR. GRIFFON: I guess we get to the question of do we have sufficient data that, is it plausible, the bounding dose. And I think we kind of stopped there, but we do have to at least make that determination.

DR. MAKHIJANI: Well, I think with the bioassay data being all below the minimum

1 detectable limit, you could, in principle, use 2 the minimum detectable limit for bioassay 3 dose. I don't know, Joyce is not on the line 4 so maybe --5 MR. GRIFFON: And you have a lot of air sampling. I mean, I've got to admit --6 7 DR. ULSH: Not a lot. MR. GRIFFON: -- not a lot. I shouldn't say 8 9 a lot, but you have some Health and Safety 10 reports indicating air sampling. 11 MS. MUNN: But there wasn't a lot of 12 activity going on. There wouldn't be any 13 reason for a lot of --14 DR. MAKHIJANI: Well, they did have an 15 incident. Now in the third rolling they had a 16 failure and the high, if I recall correctly 17 now, I did a quick look at these documents 18 some time back. If I recall correctly, some 19 of the high air concentrations were associated with that incident. And so you'd have to, in 20 21 order to do an assessment, you'd actually have 22 to identify those air samples, where the 23 workers were and what you would do -- now 24 they're area air samples so we will get into 25 the question of how the area air samples are

to be related to what the workers were actually breathing, which is an argument that we've had in other contexts. But it was only one out of three if I recall in the other two cases the air concentrations were considerably lower.

DR. ULSH: They were canned. If you look at the details of this operation, they took these thorium ingots, and they were trying to form them into shapes. And in order to do the operations on these ingots, they canned them.

Now Arjun is correct that on that third ingot, there was a can failure. They started to see fractures in the joints of the can so they stopped. They're done.

And then they had to remove that ingot from the can, and that's where you saw the air sampling, when they were taking the ingot out of the can. And Arjun's right. I mean you're both right. They did see detectable activity in those. They did.

DR. MAKHIJANI: Well, there was detectable activity in the air samples, in earlier air samples as well, wasn't there?

DR. ULSH: There were smaller air --

1 DR. MAKHIJANI: I'll have to go back --2 MR. GRIFFON: I mean, detectable activity, I 3 mean, I'm looking at, and I haven't seen that. 4 I know I probably haven't read through all 5 your details because you said you did a blow-6 by-blow. 7 DR. ULSH: Yes. 8 MR. GRIFFON: I probably skimmed through 9 I haven't looked at it thoroughly. But 10 the one Health and Safety report says while rolling coated thorium, they got a reading of 11 12 1332 percent of the mpl. Now I don't know if 13 that sample was 1332 percent. So that's a 14 high air sample. They go on to say routine 15 samples, and I'm not sure how they define 16 routine samples, but I would think maybe 17 something that they would assume the workers -18 19 DR. ULSH: Are you sure it was 1332, Mark? 20 I thought it was 132. 21 MR. GRIFFON: Well, I have pretty shaky 22 writing, but I thought it was 1332. Anyway, 23 routinely, to on to say routine, the average 24 samples were two percent of the mpl and the 25 highest was like 10.1 percent. So obviously,

1 that might be something more related to personnel exposures. I don't know. But it 2 3 certainly raises a question in my mind of was 4 this data being considered in any way to 5 establish your bound. It sounds like you're 6 saying that they had minimal overall potential 7 for doses, right, in this operation. 8 DR. ULSH: Well, maybe I should just stop 9 and say that I think we are in agreement that 10 dose could be bound. In terms of how high 11 that might be, that might be more of what John 12 would call a tractable issue that we could delve into. But I mean, I think with the 13 14 bioassay samples and the air samples that you 15 have, you can come up with a methodology to 16 bound that should you ever come up with a 17 claimant that was involved. Now we also have 18 the names of the people that were involved in 19 this operation. That was listed in the log 20 books. 21 MR. GRIFFON: Okay, so you have that much 22 detail, yeah. 23 DR. NETON: How many bioassay samples were 24 there? 25 MS. MUNN: You had 18 or something like

1	that?
2	DR. ULSH: Something like that.
3	DR. NETON: Large number for a small
4	operation like that.
5	DR. ULSH: I think the way it was, Jim, was
6	in the progress reports I found indications of
7	like maybe 18-ish bioassay samples. And I
8	did, and then that one log book that we looked
9	at had a couple of them in there, so on the
10	order of ten to 20.
11	DR. NETON: It's a reasonable number, not
12	one or two.
13	DR. MAKHIJANI: Were there sorry.
14	MS. MUNN: No, go ahead.
15	DR. MAKHIJANI: Were there bioassay samples
16	after the incident? Do you remember?
17	MR. MEYER: Eight individuals involved in
18	the operation, urinalysis was requested, four
19	individuals identified by name in the Health
20	Physics log book were involved in that
21	specific operation and no detectable urine
22	activity was observed and maximum intakes
23	using air activity data was not of
24	consequence.
25	DR. MAKHIJANI: I mean, that's why when I

looked at, Brant, actually I'm looking at my e-mail, and Brant actually had e-mailed all the three references, the Kittinger logs and the Callabra report. And I'm looking at them, and that's probably why we signed off on this. I'm trying to reconstruct what we did.

And so I think if you take the bioassay, my bottom line on this as an SEC issue would be that if you take the bioassay data, and like Joyce could comment on it if she has a comment, that if you take a bounding dose approach, and you have bioassay data, quite a number of samples for a small number of workers including after the incident, then you should be able to bound the dose. And so it then becomes a question of what's reasonable if you have a claimant rather than whether it's an SEC issue. It doesn't seem to me to be an SEC issue.

MS. MUNN: It doesn't seem to be a major issue at all if you're reading the report as I read it. It's very straightforward. The fabrication of the thorium metal parts from three 80 kilogram ingots, it gives you eight hours of cold rolling on June 3<sup>rd</sup>. It gives

1 you the dates, 4.62 dpm per liter square, 2 approximately 30 hours of other. I mean, it's 3 very clear in the final statement here you're 4 using data from both the general report and 5 the specific data from the one can opening. 6 Excellent agreement is demonstrated as one 7 Becquerel from the single can/uncanning task 8 and three Becquerels intake from 30 hours of 9 work with all three of the ingots. 10 calculated doses using different software 11 programs also shows consistency of resultant 12 doses less than 100 millirem to any organ. 13 that's --14 MR. GRIFFON: So you've been using the MDAs of the time. The intakes were that small? 15 16 DR. MAKHIJANI: Well, no, --17 MR. GRIFFON: The calculated intakes were 18 that small? That surprises me a lot. 19 DR. MAKHIJANI: This is how you would have 20 done the calculation back then. I think that 21 the reason for going on about thorium is if you read the documents back then they weren't 22 that concerned about thorium as a radioactive 23 24 material in the '50s. 25 MS. MUNN: We know that.

DR. MAKHIJANI: And so we understand that for how they viewed things then, but how we view things today obviously is quite different. And so that's the reason for raising this as an issue is if the dose reconstruction is done with today's science, then we have to take the approach of today's science. And then you wouldn't calculate a hundred millirem dose with today's science.

MS. MUNN: But the real issue is do you have data that was taken at the time that can be used in today's world. And if I'm reading this correctly, we have adequate data, we have specific data from then.

DR. MAKHIJANI: Oh yes, Ms. Munn, that's exactly why I said that especially the air samples are relatively few and they would raise questions about whether you could adequately bound the dose. And we would be in the arena of how many air samples, and where was the person versus where was the sample, and how long the sample was.

But because there are bioassay data, I think it's not a question of whether the dose can be bounded. Now how you would calculate

the dose I think --

MS. MUNN: Is something else.

DR. MAKHIJANI: -- and whether their approach we would sign off on is a different issue. I mean, we can address that if it's your pleasure that we should do it, but my own feeling is that in this context this is not --

MR. GRIFFON: No, no, I was actually refamiliarizing. I didn't remember how many urinalysis samples and all that. The other question with these kind of situations always for me is if you know the individuals involved, you have identifiers and everything.

And that sort of answers another question that I have is are you going to apply this to anybody who ever went in the building? Well, probably not if you know it was only limited to ten or 12 or whatever number of people you have specific names. You don't have to broadly apply it. So I think that answers another question I had in my mind.

But let me just, I just want to break from our agenda for a second. Tony has to leave, but I want to offer him the floor if you had anything to say that the work group

hasn't seen you in a few meetings.

MR. DEMAIORI: And I'd like to apologize for my early departure. I'm actually here on a job for CH (inaudible) Hill, and I wasn't notified of this till after I'd scheduled my flight. In fact, I haven't been getting notified for about six months of some of the working group meetings. So that was a problem that's been corrected.

However, I'd like to thank everybody for the hard work that you're doing on behalf of all the people that worked at Rocky Flats.

I know they really appreciate it. I know it's a thankless job, and I know that everybody here has worked very, very hard.

I would like to remind everybody that there's two different ways that this world actually works. There's the procedure that tells you this is how you do what when. And then there's what actually goes on in the workplace. And sometimes they're not necessarily verbatim the same.

To give you an example I'm a rad tech, and I don't think we ever passed a nod (inaudible) personnel. So I think we always

failed in turbo frisking, I think is the term. Supposedly you frisk at one-to-two inches a second. On a person it's about two minutes a person. Historically, we have two people on the step-off pad. We're moving three to five hundred people in a 45 minute period. So statistically that's impossible.

Not that we didn't try to do our jobs and do them very well, you know, but sometimes production methods, not the ideal conditions. So people modify and adapt and I would just hope that everybody would take that into consideration when they do go over this material. And that applies to all aspects. I don't care if it's machining or operations or dose reconstruction.

So I just would hope that everybody would take that into their final consideration. And once again thank you very much. We really appreciate it. And I know that unless people get exactly what they want you probably won't be thanked. However, you have to know that your hard work is definitely appreciated at the end of the day so thank you very much.

1	MR. GRIFFON: Thanks, Tony. We'll make
2	sure, we'll also make sure you get all these
3	pieces of the reports e-mailed to you and keep
4	you in the loop better.
5	MR. DEMAIORI: Yeah, I'd appreciate it.
6	MR. GRIFFON: I apologize for the
7	miscommunication.
8	MR. DEMAIORI: You know, I guess they
9	changed e-mail systems and somehow we didn't
10	get on it. However, we're there, so it's been
11	corrected.
12	MR. GRIFFON: Okay, thank you.
13	Let's take a quick break then. I
14	think people want a comfort break, and it's a
15	good time. Let's keep it at ten minutes
16	though because I do want to get out of here by
17	5:00. I'm still
18	(Whereupon, a break was taken at 3:00 p.m.
19	and the meeting resumed at 3:13 p.m.)
20	MR. GRIFFON: We're going to reconvene, and
21	I'm not sure we wrapped up the, let's just
22	finalize thorium here for a second, and then
23	we'll move on in the agenda.
24	I'm trying to understand the question
25	of source term. I'm not sure that, unless

1	more evidence is presented to the work group
2	that we can take this source term question
3	much further. The question on the bounding
4	analysis, where do we stand with the excuse
5	me, somebody's on not muted there.
6	DR. ULSH: Hey, Jim Langsted, that's you.
7	Jim, push your mute button.
8	MR. GRIFFON: He can't hear us.
9	Jim. I don't know how to, he's not
10	hearing us.
11	For the meantime those on the phone,
12	I'm just going to kind of talk over. We hear
13	another voice here, but just try to talk over
14	him.
15	So the source term question is, I
16	don't think we can go much further with it as
17	far as actions unless we get more data that
18	says there's additional source term there.
19	DR. ULSH: When you say the source term
20	question, Mark, are you talking about
21	magnesium-thorium alloy?
22	MR. GRIFFON: No, I'm talking about any
23	thorium, any thorium use, period, across the
24	board. Did you have a comment on that?
25	DR. MAKHIJANI: The thing that Tony brought
	1

1 up about the welding rods? 2 MR. GRIFFON: Yeah, I'm not, I don't know 3 the extent. I think that was pretty common at 4 a lot of DOE sites. DR. ULSH: At a lot of sites period. 5 MS. MUNN: Yeah, it can't be very large. 6 7 hesitate to use the word significant, but it 8 can't be, it cannot be a huge contributor. 9 DR. MAKHIJANI: This is discussed in NUREG-10 1717, and the range of intakes is pretty 11 varied from small to at least not 12 insignificant if you look at NUREG-1717. while it would have probably been considered 13 14 as not important at the time that it was done, 15 no doubt. Again, it's the same question. 16 Until Tony told me at lunch I wasn't, it was 17 completely unanticipated. I didn't even think 18 to ask. But in the context of just asking him 19 about magnesium-thorium alloy, he brought it, 20 well, you were there. And since NUREG --21 MR. GRIFFON: It's going to be a complex-22 wide issue, if we --23 DR. MAKHIJANI: This would be a complex-wide 24 issue. So I just want to raise that. It's 25 not necessarily a Rocky Flats-specific issue,

1 but since NUREG-1717 does cover it, we did 2 write about that and cite the specific numbers 3 in our report. 4 DR. NETON: Is this something we should add 5 to our complex-wide tracking list? It's certainly not just a Rocky Flats --6 7 MR. GRIFFON: Right, it's not just a Rocky 8 Flats, many other sites, yeah. 9 MS. MUNN: My thought would be we ought to 10 do that and put it to bed quickly because 11 there'll be a limited number of people who are 12 involved with it. It will affect primarily individuals who did that type of work, and I 13 14 wouldn't be surprised if they aren't fairly 15 easy to identify. 16 MR. MEYER: And there are a number of good 17 peer review papers that estimate the dose from 18 that. 19 DR. NETON: So there's no doubt we could do 20 something with that. We just had heretofore 21 not considered that as an exposure. I'm not 22 sure all welding rods are thorium either. 23 MS. MUNN: No, I don't think so. 24 MR. GRIFFON: I guess the question would 25 also be, for Rocky Flats for now the question

1 is could we, would there be a plausible method 2 to bound, and maybe this 1717 would answer 3 that. Not that I'm saying that you would have 4 to do that, but if we think it's a significant 5 source or maybe a significant source. MR. ELLIOTT: Well, I think you've also got 6 to look at it in conjunction with some of the 7 8 other TIBs we've got out there like 9 construction trades TIBs, and does the 10 construction trade TIB adequately account for 11 this kind of an exposure. I don't know. 12 DR. NETON: Thorium rolling rods is not 13 something that we really addressed at this 14 point. 15 MR. ELLIOTT: I understand that, but the 16 design, the exposure design in the 17 construction TIB gives a very conservative, 18 generous estimation of dose. 19 DR. NETON: But it's the separate nuclide 20 source term that we haven't really, we'd have 21 to look at it, and it's a good point. My 22 feeling is that we could certainly bound it 23 somehow. I mean, you know how much is in a 24 welding rod, and you know the estimation of 25 dose, and that's where --

1 MR. GRIFFON: And I think for the most part 2 you would expect a local exposure environment 3 to the welder, right? 4 MS. MUNN: Right. 5 MR. GRIFFON: I don't think we're talking broad exposures to others so you could 6 7 probably do something where you knew job 8 titles and do --9 DR. MAKHIJANI: I would tend to agree with 10 I haven't studied this issue. I looked Jim. 11 at NUREG-1717 for magnesium-thorium alloy to 12 see what was there. And actually I believe 13 you brought up the question of thorium welding 14 rods when we were first discussing --15 MR. GRIFFON: Mel did. 16 DR. MAKHIJANI: Okay, NIOSH brought it up 17 anyway, and because I wasn't aware of it, and 18 I did note that there was this German study 19 which I cited that has been discussed in 20 NUREG-1717. So I think that this has been a 21 studied issue. So I don't know what the 22 universe of data is. 23 MR. GRIFFON: I think we should probably 24 keep it on our radar screen in terms of 25 thorium use, but my gut feeling there, and we

should look at this further, is that if you have all these job cards, a lot of job card data. And I'm not saying the DR would necessarily go down this path, but we have to answer the question of could you estimate a plausible upper bound.

And if you know welder and if you know something about source term as Jim said, I think you can probably estimate an upper, but that's maybe for you to consider. But I think we should keep it on the radar, but I don't --

DR. ULSH: I think maybe the question, Mark, is where is the appropriate context to consider this issue. I mean, since we've already said that in, this is not a Rocky Flats specific. It's all over. So it might be one of those things that we handle wherever we're handling overarching issues. I don't know.

DR. NETON: There's overarching issues. The overarching dose reconstruction issue list is what we're maintaining now.

MR. GRIFFON: I think it rolls into a complex-wide question, but just in terms of being able to answer the issue at hand, do we

1 have any SEC concerns here, I think we want to 2 be able to answer that for Rocky now. 3 might evolve into --4 DR. ULSH: So do I hear an action item then? 5 Would it be to consider intakes from --MR. GRIFFON: Yeah, I think we should --6 7 DR. NETON: I think we have to maybe sketch 8 out a position of maybe not in extreme detail, 9 but a position that would convince people that 10 this is a tractable problem. Put it to bed 11 but not necessarily come down to the actual 12 dose assignments but, and then table the ultimate model for the overarching issues 13 14 list. And that's to make sure we don't lose 15 track of it because that's going to apply to 16 many other sites. 17 MR. GRIFFON: I think you've probably got the pieces, too. You've got this NUREG, and 18 19 you've got enough information about the source 20 term that you can --21 DR. NETON: Feel comfortable with that so 22 they're like three, four percent thorium, a 23 similar amount. And let's just vaporize it in 24 the presence of the Board. 25 MR. GRIFFON: We'll add that on to the

1 2 3 4 5 6 7 8 9 10 assume they're all less than MDA. 11 12 13 14 15 in that limited operation. 16 17 18 right? 19 20 21 22 23 on with the other three, I --24 MR. GRIFFON: Yeah, then we're on to the 25

thorium source term question and NIOSH will give us some kind of response on it.

DR. MAKHIJANI: That takes care of it.

MR. GRIFFON: Yeah, now on the other, as far as the other, we sort of got sidetracked with rolling and that was probably my fault, but so I think we're pretty comfortable, when Tony was leaving, I think we were saying you've got some urine data and at worst case you could

You've got something to plausibly upper bound it. I knew there was limited thorium urinalysis data. I didn't understand that it covered most all those people involved If SC&A's comfortable with that, I think we, it seems like there's a reasonable way to bound it,

MR. FITZGERALD: I think again through the call we've had and this discussion, I think we're pretty comfortable. And I think assuming there's an upper bound analysis going

other three, and you're going to, NIOSH, still

1	is going to provide some sort of empirical
2	analysis on those other three.
3	DR. ULSH: Yeah, I think the ball's in our
4	court. We took a shot at it and SC&A
5	MR. GRIFFON: On the technical call, right.
6	DR. ULSH: Right, SC&A didn't necessarily
7	agree with our approach. So we have agreed to
8	take another look at
9	DR. NETON: I think it's a matter of
10	parameter selection. What parameters were
11	selected.
12	MR. GRIFFON: You're really looking at sort
13	of a bounding analysis now, right? Is that
14	what I understand? I wasn't on the call, but
15	
16	DR. NETON: We haven't given up on the fact
17	that NUREG-1400 isn't applicable, although we
18	agree that there are parameter selection
19	issues there that would maybe not make it
20	quite as conservative as we maybe heretofore
21	thought. We need to look at it a little
22	closer.
23	MR. GRIFFON: So you've got those actions?
24	DR. ULSH: Yes.
25	MR. GRIFFON: And the only additional thing

1 is the welding rods, and that question. 2 All right, moving passed thorium at 3 3:26, plenty of time. This is where we speed up. I have next --4 5 Do you want to put these two items 6 together, Joe? 7 MR. FITZGERALD: Yes, I think so. 8 LOG BOOK ANALYSIS AND DATA INTEGRITY AND SAFETY CONCERNS 9 MR. GRIFFON: Log book and data integrity. 10 MR. FITZGERALD: Right. 11 MR. GRIFFON: So log book analysis and 12 before I had safety reports, but Joe says that 13 there wasn't really outstanding actions on 14 that. I don't know if we really need an 15 update. 16 MR. FITZGERALD: The safety concerns piece, 17 and certainly, Kathy's on the phone as well. 18 We certainly finished first and distributed it 19 to the work group back in November. 20 Certainly, I think maybe the easiest thing is 21 for Kathy to just sort of capsule all three, the data integrity examples, log book review, 22 23 and maybe just provide a short overview of not 24 the specific details but the sort of bottom 25 line conclusions in the SEC context and just

1	sort of walk the work group through. Would
2	that help?
3	MR. GRIFFON: That would be helpful, yes, so
4	we're going to log book analysis, data
5	integrity and safety concerns.
6	MR. FITZGERALD: Yeah, they overlap, and I
7	think it would probably be helpful not to
8	separate them out even though that's how we
9	issued them. This way it could be digestable.
10	MR. GRIFFON: Kathy, if you can give us a
11	fairly quick summary of each of those items,
12	then maybe we can get into some more detail on
13	the log book question.
14	MS. DeMERS (by Telephone): I will try. If
15	you remember in the petitions, a lot of people
16	brought up a concern that that something was
17	wrong with the badge
18	MR. GRIFFON: Hold on, Kathy, you might have
19	to speak a little louder. Are you on a
20	speaker phone?
21	MS. DeMERS (by Telephone): No, no.
22	MR. GRIFFON: Ray's having a little trouble
23	hearing you so go again.
24	MS. DeMERS (by Telephone): If you remember
25	from the petition, several people brought up

concerns about receiving years when they were working in high dose rate areas, and we were trying to get to the bottom of this in our review with the safety concerns and the log book review. The (inaudible) of (inaudible) NIOSH most of those from the list of probably 4,000 safety concerns. And NIOSH looked at them and evaluated them for their relevance to the SEC petition. And SC&A was asked to evaluate their evaluation.

And in general, there was good agreement, but there were some areas where there was not agreement. For example, there were some disagreements regarding how they handled external dosimetry investigations. There was some disagreement in relation to some of the assertions regarding individuals involved with the concern that said basically I got a zero. I don't believe this happened in X,Y area; the dose rate was X, Y, Z.

This kind of spills into the data integrity example. The basic answer was that the areas were closest at the maximum dose rate but that that was not necessarily the dose rate that the individuals were receiving

from where they were standing. Now we looked at a couple of these situations and that explanation doesn't really pass with us.

There's got to be more to this because it's been brought up in the safety concerns. It's been brought up multiple times in the comments in the data integrity section. We haven't really found I would say conclusive evidence of a systemic problem, but it sure does come up frequently in different formats.

And with respect to assigning zero when people were in high dose rate areas, is it that there should be some further investigation into that? And one of the contentions was that the people who were communicating those dose rates in the petition were not knowledgeable of the dose rate levels and how they're measured. So in fact, most of these people were radiological control technicians so they did have the knowledge to interpret the readings. In fact, they were responsible for recording them.

And with respect to dosimetry investigations basically what we have is the word of the RADCON staff that they occurred.

We have a few, actually, I think only one log book entry that said he went out in the field and tried to figure out the dose for (unintelligible) because the badge was damaged. So either these investigations didn't happen in the field or they weren't as significant to be recorded in the log books or someone else did them if we assume that the RADCON personnel are correct. Basically, we have no paper evidence that these occurred prior to the mid-'80s. So that's kind of where we stand on that.

One of the things that I noted in the data integrity analysis was that you really had to go in and understand the entire comments, and I ended up re-describing quite a number of comments as a result. And I felt that NIOSH wasn't really answering the question or the comment that was brought up by the person providing the affidavit or the comments. Those were really the two major issues where there was some disagreement amongst the safety concerns and data integrity examples.

Another thing that I noticed with

respect to the data integrity example is that they missed about seven or eight comments which were quite important as they went through and captured comments from the Advisory Board meeting in Denver. And I went ahead and added those, and NIOSH, of course, had not had a chance to respond to them so we're not quite sure where we stand on those seven or eight comments.

MR. GRIFFON: I think we did say that for the safety concerns report, I think we pointed out at one of the previous meetings that there was some individual items that there was disagreement, but at least as a work group we told NIOSH don't, we don't want you to, we don't expect you to further investigate these individual cases. So that may be, that may have been our decision.

For the data integrity, I get these things confused sometimes, these three or four kind of do overlap a little bit in my mind. I can't remember if there were specific items that we have asked for follow up on or --

MR. FITZGERALD: Well, this was the specific 73 page, very detailed compilation that NIOSH

1 put together. 2 MS. DeMERS (by Telephone): This came out or 3 this came out in mid-January, very recently. 4 MR. FITZGERALD: August, yeah, right, 5 August. 6 MS. DeMERS (by Telephone): A lot of this 7 concern is raised with respect to the safety 8 concerns are also raised in the data integrity 9 example response. And it's almost more 10 beneficial to address them in terms of the 11 data integrity examples. 12 MR. FITZGERALD: Yeah, Kathy, this is Joe. 13 I think it's fair to say though for both of 14 those documents that we do have, in fact, 15 interpretive differences on the specific cases 16 within the safety concerns as well as in the 17 data integrity examples. And I think you 18 added that there were maybe seven or eight 19 examples that may have came out at the Denver 20 meeting that were not necessarily in the data 21 integrity compilation that we've added for 22 comment that we have since provided on January 3<sup>rd</sup> to NIOSH for review. But I think the 23 24 overall conclusion again with those 25 differences that we weren't able to

1 conclusively demonstrate a pattern or systemic 2 problem, any evidence of fraud necessarily in 3 the broader sense. Is that fair? 4 MS. DeMERS (by Telephone): We were not able 5 to conclusively identify fraudulent data 6 entries with respect to those two reviews. 7 However, you need to be aware that there are a 8 lot of examples, and I guess you will have to 9 think about what the threshold is for 10 determining if that, if you're going to accept 11 NIOSH's explanation in regard to that. 12 MR. GRIFFON: Hold on, Kathy, Brant wants to 13 reply. 14 DR. ULSH: But at the end of the day, as Joe 15 said, I think you discovered no evidence of 16 fraudulent data entry. I mean, it's SC&A's 17 position that you have discovered no evidence 18 that would support that. Is that correct? 19 MS. DeMERS (by Telephone): I'm not 20 disagreeing with that. I would emphasize we 21 don't have any conclusive evidence. 22 MR. FITZGERALD: We can't find it. 23 DR. ULSH: Well, okay, I guess the other 24 part of that question then is how hard have we 25 looked. And I would put on the table that we

put out a 73-page document and considered every concern expressed at the Board meeting with the exception possibly of seven that we might have missed. We've looked at 5,000 safety concerns. We looked at -- what? -- 60 log books? And if you don't find something after all of that, can't you draw some kind of a conclusion from that?

MR. ELLIOTT: I think we've surpassed the threshold here. In a world of limited resources I understand and appreciate the need to respond to these individual allegations and assertions that were made in these affidavits. But I think we've more than enough addressed this issue.

MR. GRIFFON: That's why I didn't ask for any follow up from NIOSH on these items because I think at this point it's one of those questions, like with Y-12. I mean, we've got a bunch of prongs, and we're going to look at the weight of this evidence --

MR. FITZGERALD: And I think you have --

MR. GRIFFON: Everything here points to at the worst I think inconclusive is sort of where SC&A is weighing in. At the best it is

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MR. FITZGERALD: And you have the reports, and in a sense we were very careful I thought about how to lay this out. And I think our conclusions are not inconsistent with what you've said that we've looked hard, that we've looked far, and we did find a number of specific cases where we were concerned. But in the final analysis there wasn't any evidence of fraud or a systemic pattern that we would be concerned about and that's based on what we did look at.

MR. GRIFFON: How about for the log book analysis, the last piece?

MR. FITZGERALD: Well, that was the most recent, and I realize that was because of the Privacy Act review that got over kind of late. That got over the 19<sup>th</sup>, but again, Kathy, do you want to summarize pretty much, that has a long history. Maybe you can briefly summarize where that came from way back when.

MS. DeMERS (by Telephone): Back in March I started work with the Rocky Flats records folks to pull back some records, primarily dosimetry and procedures, processing logs,

that type of material. And then I went and visited Rocky Flats and talked to the petitioners on the 27<sup>th</sup> or 29<sup>th</sup> of March. And in the course of talking to the petitioners they indicated that the log books may disagree with the dosimetry records.

And so I went back to the records folks, and I said can you search for RTC log books, contamination control log books, et cetera, and provide me with the results. And of course, they didn't have time to pull that from the Records Center in one day. So they weren't available for review when I was there.

The plan was to take some of these affidavits and see if we could locate the person in the log books and follow through the dosimetry processing logs and make sure that they didn't have a damaged dosimeter or some sort of situation where a zero would be, situations that would require a dosimetry investigation. Then we were to go to the actual dosimetry records and do a comparison back to the original number in the log book.

Well, what happened when we went through several log books was that we couldn't

find these particular people. So we decided to evaluate people with dose information that were in the log books. But let me back up here a little bit.

The original intention with the second request to Rocky Flats for the log books was to have them scanned and then I was going to go through and walk through this process. In April the working group decided to turn this over to NIOSH. And we went through several iterations of trying to explain, you know, what my original plan was and how they might implement it. But in the meantime there were these log books sitting there.

And when it was turned over to NIOSH, they were to go through those log books, and there were boxes of them. And at least say this will be useful; this will not. And if there were log books that were useful, they were to scan them and provide a copy also to SC&A and the working group.

Kind of on a separate path, ITT was to make a very, very specific request of Rocky
Flats in the process of my own review being centered around identifying the main

supervisors over the buildings that may have had log books. They centered around the 1969 fire. So those '69 field log books were not intended to cover a large range; whereas, the initial log books that I had asked the records people to pull, were. As a matter of fact they ranged from 1957 through 1996.

In July NIOSH came back with a review of the famous Kittinger log which spans from '58 or '56 through '68. And they extracted quite a number of individuals from the log book and compared the doses in the log book back to the health physics file. A lot of these were external dosimetry results.

Unfortunately, they were for a cycle for some of them rather than a quarter when the health physics file really had a quarter.

So we determined whether they were an exact match or consistent, and we had no problems with the exact matches or the consistent matches. Then they came out with - - and Brant, you're going to correct me if I'm wrong -- with about 94 percent agreement.

DR. ULSH: Yeah, that's about right, Kathy.

MS. DeMERS (by Telephone): There were

1 others that they didn't have files for, and 2 they had to go back to Rocky Flats and request 3 them. And at that point, that review, there 4 was a discussion about the fact that, well, 5 that probably knocked people out of that log 6 book. But that log book doesn't represent the 7 entire span of time, and it also doesn't 8 represent the uranium area or the non-9 plutonium area. 10 DR. ULSH: Wait a minute, Kathy. Are we 11 talking about the Kittinger log book or are we 12 talking about --13 MS. DeMERS (by Telephone): Yes, the 14 Kittinger log book. 15 DR. ULSH: Oh, okay, the Kittinger log book, 16 yeah, I reviewed half of the data --17 MR. GRIFFON: That first one. 18 DR. ULSH: Yeah, that first one. I don't 19 remember the number, but I found a pretty high 20 21 MR. GRIFFON: Agreement. 22 DR. ULSH: -- level of agreement, yeah. The 23 94 percent number though is for our overall log book analysis. We found overall we found 24 25 94 percent agreement.

1 MS. DeMERS (by Telephone): That's pretty 2 good agreement with the Kittinger logs to go 3 back and look at consistent results and exact 4 results. 5 However, 19 of these people from the 6 Kittinger log were picked up in the second 7 review which was to cover from 1969 through 8 the '90s. 9 MR. GRIFFON: Right, this is the final 10 report that you issued to us. 11 MS. DeMERS (by Telephone): Right. 12 MR. GRIFFON: The log book review, yeah. MS. DeMERS (by Telephone): I kind of 13 14 segregated them into two separate reviews. 15 But there were a couple that got dropped in 16 the transition, but that was primarily because 17 we decided enough was enough with that log 18 book. 19 There was a second review that was 20 done, and like I said it was supposed to cover 21 1969 through the 1990s. In July, NIOSH was 22 asked to come up with a sampling plan for how 23 they were going to sample further log books, 24 and these log books were to cover various 25 processes onsite including plutonium and non-

1 plutonium areas. 2 And what they came back with was not a 3 sampling plan but a review of a subset of 4 those 50, and that's an approximate number, 5 log books that I had retrieved and subsequent 6 from the original log book request. And as I 7 said these were more specific to particular 8 time periods, and they were targeted at the 9 RADCON field supervisors. 10 MR. GRIFFON: Yes, so now we're on the final 11 report from NIOSH. And what time period did 12 that cover, Kathy? 13 MS. DeMERS (by Telephone): That covers from 14 1957 through 1971, and that included 20 15 urinalysis log books and 16 field log books. 16 As a matter of chance there were log books 17 from the earlier years from the uranium area, 18 and there were log books for select years for 19 the plutonium area. And that was primarily 20 because the supervisors that I had requested 21 were from all over the plant. Is that, I'm thinking 20 22 MR. GRIFFON: 23 urinalysis logs, is that accurate? 24 DR. ULSH: I don't know how many urinalysis 25 logs there were.

MR. GRIFFON: Anyway, we can do that later.

Go ahead, Kathy, sorry.

MS. DeMERS (by Telephone): Well, again, various items were extracted from the field log books, urinalysis results, whether a person was involved in an incident and was sent to the in vivo counter. I think there was in one count and some external dosimetry results. With respect to the log books, they actually select a claimant from within the log books and made a comparison of that urinalysis log book to the health physics file, and in general, we had compared the results to the health physics file, which I call the hard copy record here, 94 percent of the bioassay results agreed or were consistent.

In the case of in vivo counts, and a lot of times they will just reference that they sent somebody to the in vivo counter so we were virtually just looking for evidence that they had an in vivo count. Eighty-six percent of the entries polled agreed with what was being stated in the log books. There were four individuals where we didn't have files available, health physics files available for.

And the agreement with the dosimetry results listed in the log books versus the health physics records were fairly good.

One log book that they selected individuals from was called, I'm going to refer to it as the 1966 through 1969 Special Analysis Log Book. And this had bioassay results for radionuclides other than uranium and plutonium. And they polled 24 people I believe out of that log book, and again, compared the log book results back to the health physics file. We did not have the means available to do this comparison so we were not able to verify their results. But in their write-up it indicated that they did not get agreement in eight cases out of the 24.

MR. GRIFFON: That's for the special radionuclide log?

## MS. DeMERS (by Telephone): Yes.

We also did a comparison of the log book results to the HIS-20 external database and the urinalysis database, and this is where we started to discover some issues.

We had 68 percent agreement between the bioassay results and the log versus the

1 health physics. Let me back up. First is the 2 data available in HIS-20, and the remainder of 3 those people were not showing up in HIS-20. 4 So I went and I scanned the people who were 5 not showing up in the HIS-20 to determine 6 whether they had zero doses or positive doses. 7 And there was a mixture so we're not just 8 talking about people who didn't have positive 9 urinalysis data in this case. 10 MR. GRIFFON: This could be important back 11 to our discussion on coworker models 12 obviously, that last point. 13 DR. ULSH: (Unintelligible). 14 MR. GRIFFON: Yeah, and then there could be 15 explanations if they retired before '77 or 16 whenever. We know, we know the explanations. 17 Go ahead, Kathy. Was that it for --18 MS. DeMERS (by Telephone): There are two 19 other things. 20 The problem was not as bad with the 21 HIS-20 data, between the HIS-20 database and 22 the external data, but it was still there. 23 And the final concern that we had was 24 going back to the original log book request 25 that I made of Rocky Flats of the RCT logs,

1 the contamination logs, et cetera, which got 2 turned over to NIOSH, and there's no 3 indication by NIOSH that whether those log 4 books were useful or not for the SEC petition 5 review. And they have indicated that they 6 reviewed 450 boxes of documents; however, 7 we're looking at 59 log books here and there's 8 a delta there that SC&A doesn't know anything 9 about. 10 We don't know if those log books are 11 useful or whether (unintelligible) is not 12 having any individual data in it. We don't know the contents of these 450 boxes, and 13 14 there's a concern that they concentrated on 15 these 60 log books that are on the O drive 16 which were not representative of the '69 17 through '99 timeframe. 18 MR. GRIFFON: 'Ninety-two. 19 MS. DeMERS (by Telephone): I think Arjun 20 said through --21 DR. MAKHIJANI: Two. 22 MR. GRIFFON: 'Ninety-two. 23 DR. MAKHIJANI: Early '92. 24 MS. DeMERS (by Telephone): So we were left 25 wondering, okay, what about these other time

periods. Did you find any log books in here? What about the representativeness of what you did choose to review? And this kind of goes back to the sampling plan and the fact that the sampling plan was not produced but instead this review was produced.

MR. GRIFFON: Kathy, I guess I just want to, I mean, from my read on this two things jumped out to me to follow up on anyway. One was this question of, you know, we did, there was an action item to do a sampling plan. I understand. You did a final report, but the real question, at the end of the day we just want to make sure we cover all time periods of concern and all potential, all operations of concern. So if we, I think it does look like - is it from '71, you know, '72 through '92 seem to be not represented very much in this review.

DR. ULSH: And the reason that they're not represented, Mark, is that log books that contained the kind of data that we were looking for did not exist after '71 or '72 and they went to an electronic --

MR. GRIFFON: Yeah, we know there weren't

1	any urine logs, and I don't know if there are
2	any other logs that
3	DR. ULSH: The same with external, now if
4	there are other logs
5	MR. GRIFFON: No, not just, yeah, okay, I
6	mean, the other field logs that would have had
7	some
8	DR. ULSH: There are other logs, the RCT and
9	contamination log books that exist into the
10	later time periods, but as we discussed at
11	previous working group meetings, those logs
12	are not the kind of logs that contain the
13	useful information that we needed for cross
14	walking. We talked about that, that there
15	wasn't a lot of information in those logs.
16	MR. GRIFFON: We talked about the foremen
17	logs not being very useful. I don't remember
18	talking about the RADCON or DECON logs. Maybe
19	we did.
20	DR. ULSH: I think we did, didn't we?
21	MR. GRIFFON: But I thought the foremen logs
22	were the ones that we were
23	MR. FITZGERALD: I think it was clear that
24	the foremen logs were not useful in any sense,
25	but I think it was

1	MR. GRIFFON: I mean, I haven't looked at
2	these things so if you reviewed and
3	DR. ULSH: Well, I'm trying to think back
4	from months ago, but I think I presented an
5	example of not just a foremen log but I
6	thought a contamination control log book. I
7	don't know about RCT. I can't remember. It's
8	too far back, but there just wasn't, that they
9	were very, very data poor. I didn't, I don't
10	think I saw any data in there.
11	MR. MEYER: Oh, we looked at RCTs.
12	DR. ULSH: We looked at RCT logs?
13	MR. MEYER: Yes.
14	DR. ULSH: Okay.
15	MR. MEYER: Microfilm.
16	DR. ULSH: Yeah.
17	MR. GRIFFON: With microfilm, too, you said?
18	DR. ULSH: There are some things here that I
19	do need to respond to.
20	MR. GRIFFON: So just let me, so that's one
21	question, and I think you've responded to it.
22	The only other thing that jumped out at me
23	from this review is the tie back into the HIS-
24	20 database. And I think there are some
25	possible explanations for some of these

1 things, but I wanted to ask did we, I thought 2 at some point early on in this process you all 3 did an analysis of hard copy records versus 4 HIS-20 and came out with very strong 5 agreement. Was that with log books or what 6 were you comparing that, that point? I think 7 it was RAD files with HIS-20. 8 DR. ULSH: Wasn't it TLD worksheets? 9 MR. MEYER: TLD worksheets and then I think 10 11 MR. GRIFFON: So it was external or was it 12 both? 13 MR. MEYER: Yeah, external. 14 I presented those DR. ULSH: External. 15 results at the April Board meeting when we 16 gave our ER presentation. I don't remember 17 the exact numbers. But, yeah, I'm sure it was 18 TLD worksheets versus HIS-20. And then I 19 can't remember about internal. 20 MR. MEYER: We did some internal, too. 21 MR. GRIFFON: And TLD worksheets would have been beyond the, I forget what time period you 22 23 covered. We'll have to resurrect that 24 document, but it could have been started in 25 the early '70s, right?

1	DR. ULSH: Yes.
2	MR. GRIFFON: So it would have been through
3	the `70s, yeah.
4	DR. ULSH: I don't know. I can't remember
5	the time period that the ones we looked at
6	covered. I don't remember.
7	MR. GRIFFON: I mean I think we might want
8	to reflect on that piece because that may help
9	us, you know, like I've said again and again,
10	we've got several prongs, no one item is going
11	to cover this perfectly.
12	Did you write that, Craig or was that
13	
14	MR. LITTLE: Pardon?
15	MR. ELLIOTT: Was that yours or, you wrote
16	that, right?
17	MR. LITTLE: Yeah.
18	MR. GRIFFON: I can't remember when it was
19	provided. I think it goes back to
20	MR. ELLIOTT: Or when you issued it? It was
21	before the April meeting.
22	MR. GRIFFON: Yeah, I think it was in March
23	at the Boston meeting that that first came
23	
24	out. Anyway, we might want to reflect back on

1	Go ahead, Brant. That was my two.
2	UNIDENTIFIED (by Telephone):
3	(Unintelligible) some of that information is
4	captured in OTIB-58.
5	MR. GRIFFON: OTIB-58 has that reference?
6	UNIDENTIFIED (by Telephone): Yes.
7	MR. GRIFFON: Okay, we'll look in OTIB-58
8	for that reference.
9	UNIDENTIFIED (by Telephone): It's Section
10	Six.
11	MR. GRIFFON: Section Six, thank you.
12	MS. DeMERS (by Telephone): Let me clarify
13	something. In the log book analysis we did
14	not go from HIS-20 to the health physics file.
15	We went from the log book to HIS-20.
16	MR. GRIFFON: I understand, Kathy. I'm just
17	saying that there was another type of
18	analysis. I was getting a little history
19	lesson. I couldn't remember if they looked at
20	log books or if they looked at TLD worksheets
21	as it was.
22	DR. ULSH: Okay, well, I'll make this brief.
23	It might have been Kathy's original plan to
24	look at the log books as they surrounded the
25	affidavits, but we were never tasked by the

working group with focusing on, in any way, on the particular individual instances listed in the affidavits. In fact, we were tasked to do a random sample which is exactly what we did. This analysis was never meant to be exhaustive. It was never meant to look at every log book that existed. What we were to do was to come up with a random sample.

Now in terms of why we didn't look at the log books past 1971 as I've already stated, the types of log books that were most useful, the urinalysis log books primarily, and also the ones that contained a lot of external dosimetry data did not extend past 1971, '72, right around the time when they started going to electronic recording. So those log books simply don't exist.

Now there are, as I said, RCT contamination control various kinds of log books, but those are of the type that we determined and discussed with the working group that they were not, they didn't contain the kind of data that we needed to cross-walk to draw a conclusion.

Now Kathy mentioned the 60 log books

that are on the O drive, and she mentioned that we reviewed a subset of them. I don't believe that is accurate. I believe that we reviewed every log book that has been posted. In terms of those 60 log books, I mean it is presumed that those 60 log books were requested by SC&A because they related to a specific concern in the affidavit. So to that extent I think that our analysis is even more targeted. I mean if what Kathy's saying is true that those log books supported the particular concerns that she had --

MR. GRIFFON: Let's step back and remember a meeting where we talked about the overall goal of this. There is specific references in the transcripts where we said a lot is derived from some of the affidavits and comments, but we're really, our goal here is the SEC, the entire population covered by the SEC. So we want to do a plan that would cover all the decades, '50s, '60s, '70s, '80s, up to '92. I think we cut it off because the D&D was kind of a separate thing, and different types of operations, relevant operations.

DR. ULSH: Right, uranium and plutonium.

MR. GRIFFON: Yeah, I don't particularly remember. I'm not doubting that you said this, but I don't particularly remember. I remember the foremen's logs being dismissed because we had several examples of those. But the other stuff I guess I don't remember that as well. I don't doubt that you mentioned it in a meeting. We've had quite a few meetings.

MR. MEYER: Well, an example I guess of the RCT log book is the first notice of the '69 fire was in an RCT log book, and we did talk about that. So that was one of them that --

MR. GRIFFON: Yeah.

MS. DeMERS (by Telephone): I want to clarify something here. The original plan was to target people who issued affidavits, but we found that we could not find those specific people in the log books. So we expanded it to people with data in the log books.

MR. GRIFFON: Okay. But anyway, I'm just saying we also expanded it to cover the SEC petition as a whole. So I mean I guess the question would be, you know, it sounds like you reviewed, you know, you're telling the work group that you've reviewed other of these

1 other RCT or DECON log books, and you've made 2 determinations that there's just nothing there 3 worth, nothing to cross-check basically. 4 DR. ULSH: We looked at the log books in 5 their hard copy form to determine whether it was worthwhile to scan them because as we said 6 7 8 MR. GRIFFON: Well, that was my next 9 question, were they available in scan so we --10 DR. ULSH: There were 450 boxes of log 11 books, and this is the most resource-intensive 12 part of the whole operation is scanning these 13 things. So we looked at them before they were 14 scanned, before they would have gone to 15 scanning to see if it was worthwhile to do 16 that. And we determined that it really 17 wasn't. So everything that was scanned was 18 posted on the O drive. There are no more log 19 books scanned that we are sitting on or hiding 20 or anything like that. You have everything 21 that we had scanned. 22 MS. DeMERS (by Telephone): We understand 23 that. MR. ELLIOTT: And we don't see the utility 24 25 in going forward and looking, scanning the

other log books because they don't bring anything to bear.

## DR. ULSH: Exactly.

Now in terms of HIS-20 versus the health physics files, we focused on the health physics files because remember that the original concern was is the data that's being used for dose, individual dose reconstructions reliable or are there some problems here that would indicate fraud or whatever the concerns were. So we thought it was most relevant to go to the RAD files themselves even though that was, you know, a bit more work.

But there was another motivation for doing that, and that is the known limitations at this point. We talked about the in vivo issues with HIS-20, and we've also talked about what Mark mentioned that individuals who terminated employment prior to 1976 or seven, something like that, their data was not initially loaded into HIS-20. Some of it was later restored if they were part of the medical recall program but certainly not all.

And therefore, we figured that it would be most informative for those reasons to

look at the data in the hard copy RAD file.

That's what's used in individual dose

reconstructions. That's why we didn't --

MR. GRIFFON: That's fine. This is why we go down this path all the time, but that's what's used in many of the individual dose reconstructions. I mean you have coworker models for a reason. You're going to use them eventually.

DR. ULSH: Yes, yes, we did.

MR. GRIFFON: And there was a third reason.

It's because the work group tasked you to look at the individual RAD files. We didn't ask you to look at the database stuff. We wanted you to look at the RAD files. That's correct. I don't disagree with that.

DR. ULSH: And you know I've got to go back to, I mean if you've never taken part in any of the lead up to this meeting, you would think from listening to the summary here that there were a lot of big important issues that SC&A did conclude at the end they didn't find anything, that indicated systematic problems. And I go back to something that John Mauro said when I presented the first results from

the Kittinger log book. I reviewed half the data points in it and John said something like, you know, if it walks like a duck, and it quacks like a duck, you know, there's really no reason to go look at the other half of this log book.

And I would extend that argument here. I mean, there are some concerns, I guess, being expressed that we didn't look at a representative sample. My question is, I mean, if these log books included at least, the log books that Kathy identified that correspond to places where the petitioner was saying that there were issues, and we've looked at these, and we haven't found, we found 94 percent agreement, what are the odds that the other log books that exist contain some problem that we haven't come across. That's analogous to the argument that John Mauro made after the Kittinger log.

MR. GIBSON (by Telephone): Brant, this is Mike. Can I just cut in for a minute? You know, I take a little bit of exception to the way you just characterized what you just said, if it walks like a duck, quacks like a duck,

1	it must be a duck.
2	DR. ULSH: I didn't say that. John Mauro
3	said it.
4	DR. MAURO (by Telephone): No, I didn't
5	quite say that. I said this house is clean.
6	DR. ULSH: Oh, yeah, something about rotten
7	in Denmark or
8	MR. GIBSON (by Telephone): I just want to
9	say that, you know, having lived in the DOE
10	world if it walks like DOE, and it talks like
11	DOE, you can't be assured of nothing.
12	DR. ULSH: Mike, my point was if we've
13	looked in a large sample of log books, and we
14	haven't discovered an issue, what is the
15	likelihood that we're going to find an
16	undiscovered problem if we look at more
17	similar log books. That was my point. That's
18	all I'm saying. I'm not saying anything about
19	DOE's credibility or anything like that.
20	MS. DeMERS (by Telephone): This is Kathy.
21	Can I make two requests?
22	MR. GRIFFON: Sure.
23	MS. DeMERS (by Telephone): One would be
24	that we get the names associated with those
25	pulled from the Special Analysis log books so

that we can include that in our review. 1 2 the other issue is can NIOSH document what 3 happened in their review of these log books 4 that were not scanned. 5 MR. GRIFFON: I think you're documenting to 6 an extent on the record here. I think Brant's 7 documenting it to an extent. I mean, I don't 8 know if you can describe that better how you 9 got those 450 boxes I think is the question. 10 You looked at how many and you determined that 11 they weren't very useful for in terms of, I 12 mean, I think that --13 DR. ULSH: I can't say off the top of my 14 If that's something you want, we can do head. 15 that. 16 MR. GRIFFON: Yeah, just a description of I 17 think is what --18 MR. FITZGERALD: Well, the presumption, too, 19 then is the balance of those log books were in 20 fact perhaps bridged these time periods so we 21 talked about they were the basis for the sampling plan so that's the other kind of 22 23 implication although I'm not sure that's, 24 whether you confirmed that or not. Because we 25 don't know what's in those boxes.

1 that's what she's saying. 2 DR. ULSH: Let me clarify. I misspoke earlier when I said that we reviewed 450 3 4 boxes. We pulled 450 boxes, and we looked 5 through those for log books, and it wasn't 450 6 boxes full of log books. I don't want to 7 imply. 8 MR. GRIFFON: So you looked through log 9 books, and you sampled from, but you can give 10 us a little description of that field 11 activity. And the first item I think you can 12 provide us with those names for the Special 13 Analysis log book. Were they in the report? 14 I don't know if --15 That's a very, I'm familiar with DR. ULSH: 16 that log book. I've looked at it a lot. 17 There are, I think, maybe 40 individuals in 18 that log book. In terms of identifying the 19 exact individuals that were in there, we'll 20 find out. 21 MS. DeMERS (by Telephone): What I'm 22 referring to is the individuals you did the 23 comparison for. 24 DR. ULSH: Okay, yeah, I'll try to track 25 that down. But I will point out though that

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that log book contains results, urinalysis results for what we've categorized as other radionuclides. I'm thinking of, you know, curium, neptunium for which SC&A's already agreed that there's not an issue with those radionuclides.

MS. DeMERS (by Telephone): Well, this is a matter of data completeness and not a matter of whether people were monitored.

DR. ULSH: I'll get the names.

I'm not sure, you know, this MR. GRIFFON: question of the, this crops up in many of the pieces we've done. It seems to me after '69 there's nothing to look at to verify. That's a concern I have, and you know, there's no log books. There's no, I'm curious to maybe understand this, and I know it was brought up at a previous meeting. I think Gene Potter might have brought this up. That after '71 or whatever year it was. I'm not sure of the exact year. You went to this electronic entry with the data. There was no urine logs, or they just live entered. But I don't know that, I mean, did they have, what types of computers were in place at that point to do

1 live entry or was it card entry or was it, I'm 2 surprised there's no hard copy record at all. 3 MS. MUNN: It would have had to be card 4 entry. It was Sperry Rand. That's why I was 5 there. 6 MR. LITTLE: I don't think it's quite 7 accurate to say there were no log books after 8 1971 because there were log books clear up 9 into the D&D era for certain things. 10 MR. GRIFFON: No urine logs it was saying. 11 MR. LITTLE: Well, not that I'm aware of. 12 MR. GRIFFON: Right, no urine logs. 13 MR. LITTLE: And we looked at log books of 14 various titles. We looked through reams, 15 literally, microfilm, boxes and boxes of 16 microfilm. 17 MR. GRIFFON: I mean, those urinalysis logs 18 were very helpful because you had obviously 19 quantitative data to compare against with 20 individuals, but then it stopped in '69 or '70 21 or whatever. And the question is beyond that 22 what happened? Apparently there's no hard 23 copy record to compare against after that. 24 That's your experience. I'm not saying, if 25 it's not there, it's not there.

1 MS. MUNN: Speaking from memory not at this 2 site but what was going on in the computer 3 world then, it had to be, you had card decks 4 and mag tape. That's what you had. 5 MR. GRIFFON: It wasn't laptops. 6 MS. MUNN: Yeah, no laptops and certainly no 7 Blackberrys. It was, but direct entry would 8 have probably been card stock. But also from 9 my memory one didn't maintain cards for a 10 long, long time. After you had, after you 11 transferred them to mag tape, you usually 12 discarded them. MR. LITTLE: Yeah, the (inaudible) becomes 13 14 the record at that point. 15 MR. MEYER: The mag tape might still exist 16 at the archives. 17 MS. MUNN: Might. 18 MR. MEYER: We'd have to re-invent the tape 19 drive. 20 MS. MUNN: Yeah, and finding the platform is 21 something else again. If you can find the 22 tape, you can't find the platform. 23 MR. GRIFFON: Well, some of the sites 24 certainly had the card data, saved, archived, 25 but I guess we just didn't have it here.

1	DR. ULSH: I don't know, Mark, I don't know
2	that we specifically looked for card
3	MR. GRIFFON: You haven't found it I would
4	say.
5	DR. ULSH: We don't have it in our
6	possession.
7	MR. MEYER: We haven't encountered them. We
8	haven't looked at, I mean, my experience at
9	Oak Ridge was that at that point it would have
10	been line-printed output and the cards by now
11	certainly would have been discarded. It would
12	be just a massive
13	MR. GRIFFON: So I don't know, you have this
14	log book report. You haven't given us a
15	response to this or did you?
16	DR. ULSH: No, we haven't given a response
17	mainly because the conclusion at the end was
18	that they didn't see any SEC issues.
19	MR. FITZGERALD: Remember that the charter
20	for the work group was to review what NIOSH
21	had evaluated. We, in fact, did so and based
22	on that scope I think our conclusion wasn't,
23	we didn't see any systemic issues. I think
24	the only asterisk we're adding is that we have
25	some reservations about the record as we've

discussed, and we certainly didn't have any information on the way the boxes were used. I think some of this information is now forthcoming. So that's kind of where we are.

MR. GRIFFON: Which I don't know, you know, that's what I'm looking for is another prong that sort of fills that gap of the '70s and '80s, you know. I mean, it seems like, at least in my experience of what I've seen on post, I'm limited to what I can view obviously, but the log books look like they only go into the early, I mean the Health and Safety reports also look like they, unless you have later ones that go through the '70s into the '80s.

DR. ULSH: I haven't located any beyond '71.

MR. GRIFFON: And my experience is that on most of the sites the utility of those safety reports decreases as time goes on. There's more data in the early ones and less data in the later ones. There's more language in the later ones but less data. It's just the way they, yeah, maybe for legal reasons, whatever. But this is not inconsistent with what we ran across at Y-12 actually that we sort of in the

1 '70s we started having difficulty with 2 anything to compare. 3 MR. FITZGERALD: Yeah, we can't conclude 4 anything beyond the scope that we looked at. 5 But what we did look at certainly we didn't 6 see a systemic. 7 MR. GRIFFON: I guess at this point the only 8 outstanding action then would be for this 9 description of your sampling or review of those 450 boxes, and --10 11 DR. ULSH: Are you looking for anything 12 further on data integrity and safety concerns 13 or are we --14 MR. GRIFFON: I don't think so at this 15 point. 16 MR. FITZGERALD: Not unless the work group 17 directs because otherwise we've been through 18 the review of what you dealt with the 5,000 19 and I, you know, that's the scope, and we've 20 looked at it, and we've applied conclusions. 21 MR. GRIFFON: Yeah, I don't think we want 22 any more action there. Taking a deep breath 23 at that point. But the only, you know, other 24 than the scope I think, and you may disagree 25 with me, Brant, on this, but this question of

1 the comparison of the logs versus the database 2 still raises my concern as it applies to the 3 coworker stuff that we're going to run into. 4 And so we've got several prongs that are 5 pointing to this data completeness and 6 coworker models. 7 DR. ULSH: I hear your concern, Mark, but 8 what I'm wondering is I haven't looked at that 9 particular section of SC&A's report. 10 wondering if there are individual identifiers 11 where we can see where the HIS-20 data, where Kathy couldn't find it. If we can --12 Do we have that, Kathy? I mean, you 13 14 know what you sent over so I mean do we have 15 that kind of information in what you sent over 16 already or is that something we need to 17 request in addition? 18 MS. DeMERS (by Telephone): I will send you 19 a key. I don't know if it's available in what 20 you got or not. 21 DR. ULSH: So I mean we could take a look, 22 Mark, and see whether or not --23 MR. FITZGERALD: I suspect it is because of 24 Price Anderson. I'm not sure what ended up, 25 you know, in terms of identifiers. I doubt

1	much went over with identifiers, so we'd have
2	to get the key
3	MR. GRIFFON: The Privacy Act.
4	MR. FITZGERALD: provide a key that would
5	cross walk.
6	DR. ULSH: That would be helpful because
7	we'll take a look at that.
8	MR. GRIFFON: And again, this number or one
9	of the concluding statements here says 32
10	percent were absent in the HIS-20 database.
11	Now you may want to also look at CER since
12	that's the, your bioassay coworker models is
13	based on CER so you may want to look at HIS-20
14	and CER, right
15	DR. ULSH: Okay.
16	MR. GRIFFON: or primarily CER? I mean,
17	I'm not, I'm more asking than telling. I
18	think that's what your coworker is based on,
19	CER, right?
20	DR. ULSH: The internal coworker data is
21	based on CER data.
22	DR. NETON: It's got to be identified.
23	DR. ULSH: Oh, that's right, it is, isn't
24	it?
25	MR. GRIFFON: It has to be identified.

That's the problem. That's why I always have been doing mine in HIS-20, and the only time I can compare against CER is when I have these high values in certain time periods and they stand out. But these were not, if I understand SC&A's report correctly, these 32 percent absent from the HIS-20 were not only zeros or thinking that they were high values as well or mid-range values or whatever, it wasn't just simply zeros. It was kind of all over the place.

DR. ULSH: Keeping in mind that we only covered up to 1970, '71, whatever. I have to look and see whether or not these were people who you wouldn't expect to be there because they weren't injured. When their employment terminated, I guess, is what's going to be the --

MR. GRIFFON: Yeah. Well, that whole thing is troubling to me, too, that people were taken out and then some were put back in, and we don't know who and what.

DR. ULSH: Well, they weren't, no, actually, we do know who it was. I mean, they weren't taken out. They were never initially loaded.

1 The people who, I mean, this is described in 2 the evaluation report. The people who 3 terminated prior to that year, whatever it 4 was, '77, something like that, their data 5 didn't go into HIS-20 at all initially. Now 6 later a subset of those were loaded back in if 7 they were in that highest exposed workers that 8 were part of the medical monitoring program. 9 We know that. 10 MR. GRIFFON: The highest exposed workers as 11 defined by that later program, right? 12 DR. ULSH: Right, that medical monitoring 13 program. 14 MR. GRIFFON: That was a semi-voluntary 15 program though as well, right? 16 DR. ULSH: Well, I suspect. I mean, they 17 didn't compel people to come in. 18 MR. GRIFFON: Right, it was follow up. 19 Okay, I think that, I think we need to, those reports we mentioned previously need to be re-20 21 examined and that very short Donna Cragle 22 report has some interesting reading in it to 23 compare. But let's stay on schedule here, 24 4:30. 25 Is there any more on log book analysis

1	specifically?
2	DR. ULSH: No, I don't think so.
3	MS. DeMERS (by Telephone): No, not really.
4	SUPER S (TIB 49)
5	MR. GRIFFON: Moving on to Super-S. Joyce,
6	are you still with us?
7	DR. LIPSZTEIN (by Telephone): Yes.
8	MR. GRIFFON: Super-S, where do we stand,
9	Joe and maybe Joyce can
10	MR. FITZGERALD: Just generally, obviously,
11	we presented back in June at the Board meeting
12	in D.C. the review on the conceptual approach
13	in OTIB-49, and I think certainly the
14	conclusion at that point in time was that we
15	thought that was scientifically valid. We
16	have since proceeded to look at the
17	direction of the work group look at the
18	case model, the model cases that were
19	included. Was it 25? Twenty-five model cases
20	and Joyce has been evaluating those, and I
21	think - am I right, Joyce, the only issue
22	there was this lung adjustment factor? Are
23	you making headway?
24	MR. GRIFFON: Which now you have.
25	MR. FITZGERALD: Which I think we got

1 recently, but is that, have you concluded that 2 that's what you need to finish? 3 DR. LIPSZTEIN (by Telephone): Yeah, what we 4 got to see just the adjustment factor, but we 5 still didn't get everything because we got the adjustment factors that were used until '95, 6 7 and from '95 on after '95 we didn't get what 8 were the adjustment factors. 9 MR. GRIFFON: Well, maybe you can follow up 10 with Roger or Brant. 11 DR. NETON: I thought we sent everything, 12 but --13 MR. FITZGERALD: It sounds like there's, 14 what, seven or eight years, just the tail end 15 that's missing. 16 MR. GRIFFON: But I guess the outstanding 17 action is for you to complete this review. 18 MR. FITZGERALD: Complete the review --19 MR. GRIFFON: This was the idea to see if 20 the model cases selected were inclusive of the 21 other 25. MR. FITZGERALD: Joyce, we're going to 22 23 pursue this with NIOSH, but I assume though 24 that with the correction factor that you have, 25 you can, in fact, pursue a large number of

1	these?
2	DR. LIPSZTEIN (by Telephone): Yes, yes.
3	MR. FITZGERALD: Okay, so really just to
4	make sure that we have for those individuals
5	with data beyond '95 you need that additional
6	factor.
7	DR. LIPSZTEIN (by Telephone): Yes.
8	MR. FITZGERALD: Okay, so you're in the
9	process of going through those calculations
10	now?
11	DR. LIPSZTEIN (by Telephone): Yeah, and
12	another point that is not only relevant to
13	these 49, the high fired, if there was, if
14	there is an adjustment factor, they have to be
15	used on all claimants' lung results. And they
16	haven't been from few claimants that were
17	involved in the '65 fire.
18	DR. NETON: I don't know that they were
19	using
20	MR. GRIFFON: I don't think they relied on
21	that data.
22	DR. NETON: the lung analyses for dose
23	reconstruction, Joyce.
24	DR. LIPSZTEIN (by Telephone): I'm sorry?
25	DR. NETON: We talked about that. We

1	weren't really planning on using the lung
2	measurements
3	MR. GRIFFON: For dose reconstruction.
4	DR. NETON: the urinalysis was the
5	primary method. Is that right?
6	MR. SHARFI: Are you talking about the chest
7	sample?
8	DR. NETON: Yeah.
9	MR. SHARFI: There's been a little. I would
10	not limit not using the chest sample.
11	DR. NETON: Well, I didn't say not use it,
12	but I didn't know that we were
13	MR. GRIFFON: I thought it was discussed
14	that the primary in this case for Rocky you
15	were primarily looking at the urine, and you
16	might use the lung to bound.
17	DR. LIPSZTEIN (by Telephone): All
18	claimants? No. If you have someone that with
19	a lung cancer, and you have the lung counting,
20	why would you use urine?
21	MR. SHARFI: Yeah, I agree. I would not say
22	we would not.
23	DR. NETON: Well, I'm thinking about the
24	Super-S issue here though. I mean, I don't
25	know. I need to think about this. At one

1 point we had determined that we weren't going 2 to rely on the --3 MR. SHARFI: On for like Hanford it's more 4 likely to use the urinalysis because it's more 5 sensitive, but and much lower MDA, but the 6 Rocky Flats more often chest count that is more viable. 7 8 DR. NETON: Yeah, but see that's americium 9 there, and if you're talking about plutonium, 10 you can't see the broad side of a barn with a 11 lung counter with plutonium. So I doubt that 12 we would have hardly anyone with a measurable 13 PU-239 burn. I mean, the detection limit's 14 somewhere around a couple hundred nanocuries, 15 easily. 16 MR. SHARFI: No, we've seen measurable 239 17 in americiums in some of that Super-S 18 materials so --19 DR. NETON: Well, they'd have to be pretty 20 large lung burdens. 21 MR. SHARFI: Yes. Yes, these are big 22 intakes. But the americium will be bounded in 23 the matrix so it will exhibit the same type of 24 biokinetics to the lung that you're going to 25 talk about plutonium too.

DR. NETON: Well, I don't think that's what we're saying for the Super-S model. We can look at that. We'll take that information and

DR. LIPSZTEIN (by Telephone): And another thing that was raised actually by Bob Anigstein is that some people they were exposed to fire at Rocky Flats, and that these were not reported. And there was some fires in the glove box that were put out by the workers, resulted in this fire department being notified.

And he said that sometimes those people they were monitored, but it was not recognized that they were exposed to high-fired oxides. So when you calculate that dose, if the (unintelligible) is below the detection limit because it was high fired and we wouldn't catch up in the beginning, and you don't recognize it as high fire so you might be unfair to those workers. And I don't really know how you would address this kind of exposure.

DR. NETON: I thought our default assumption here, unless we can determine otherwise, was

1	going to be high fired for virtually everyone
2	that we couldn't determine.
3	DR. ULSH: Well, for lung cases.
4	DR. NETON: For lung cases, right.
5	DR. LIPSZTEIN (by Telephone): Yeah, and for
6	systemic
7	MR. SHARFI: It'd be one of the solubility
8	choices that you would have to consider in all
9	cases if you could not rule out that scenario
10	by means of
11	DR. NETON: You pick the most claimant
12	favorable in any analysis.
13	MR. SHARFI: Yeah, regardless of organ, so
14	it wouldn't be limited to the lung. It'd be
15	all
16	DR. LIPSZTEIN (by Telephone): But would you
17	use high fired even if you did not know it was
18	a high
19	DR. NETON: Yeah, if it's claimant
20	favorable.
21	DR. LIPSZTEIN (by Telephone): Okay, then
22	that's fair.
23	DR. NETON: Because we recognize it. I
24	think we determined it's not just fires where
25	high fired exists anymore. I mean, there are

mechanisms to have high-fired plutonium without a fire.

DR. LIPSZTEIN (by Telephone): Yes, that's (unintelligible) be exposed to high-fired oxides without being exactly exposed to a fire. And the all these results would be below the detection limit and people would not have been (unintelligible) because either the (unintelligible) or they weren't found -- well, for any reason they didn't come up for the (unintelligible) but even though if you treat them not as high fired then it would be unfair, but --

DR. NETON: Right, we agree. I mean, we would use high fired.

MR. GRIFFON: Well, the other case, and this came up actually from, I'm not sure it's still related to Super-S, but it is sort of related to this inadvertent sort of unknown exposure question. And the reason it came up for me was reviewing these Health and Safety reports, and then I found that a lot of them were logging the high urine data points, and many of them were associated with wounds.

And prior to this, so I tracked this

back and found this Chapman document, a health physicist in the early years there, that was tracking some of this. And he raised a question, at least for me, that until the time of his paper, which I think was in the early to mid-'60s, '63, '64, that they had no good method of wound monitoring.

They came up with a method to monitor wounds, and he said prior to writing this paper he felt like that the biggest hazard for plutonium doses was inhalation. But he's convinced now that it's wounds. And I actually thought that prior to this that wounds would have been the exception rather than the rule. But apparently these were fairly common to get punctures in the glove box work and stuff.

So my question would be if you had an acute exposure from an injection basically instead of an inhalation, is the chronic model going to be bounding? I know we've looked at acute inhalation spikes, and we've convinced ourselves always that the chronic is bounding. But in that situation --

MR. SHARFI: The dose associated with the

1 wound incident is usually much less because 2 you get so much straight excretion to all you 3 see in high peak in urine so quickly. 4 MR. GRIFFON: But is that true for all 5 organs is my question. I mean, certainly --6 MR. SHARFI: Obviously, the lung is not 7 impacted because it's straight to blood. It's 8 going to bypass the --9 DR. NETON: The organ dose is directly 10 proportional to the --11 MR. SHARFI: To the excretion. 12 DR. NETON: -- the excretion, right? So you have a handle on that. 13 14 MR. SHARFI: Correct. 15 MR. GRIFFON: But if you have non-detect 16 urine then the coworker chronic assumption 17 should still be bounding? 18 DR. NETON: I think so. I think --19 MR. SHARFI: Usually inhalations almost 20 always a more claim favorable assumption over 21 wound incident because of the direct injection 22 would cause a quick spike versus the long-term 23 buildup of the, you have a much longer exposure scenario with a, inhalation takes 24 25 time to get into the blood system or the --

1	DR. NETON: I don't want to say too much
2	without
3	MR. GRIFFON: Yeah, I looked at this quickly
4	and that's why I'm asking because I've got to
5	look at it further, too, but I wanted to raise
6	it here so you could also consider that as
7	another, and this is really on the
8	implementation of the internal coworker.
9	MR. SHARFI: And there is a wound TIB that
10	covers how to assess wounds.
11	MR. GRIFFON: Oh, I didn't know that.
12	DR. LIPSZTEIN (by Telephone): Yeah.
13	DR. NETON: The NCRP model.
14	MR. SHARFI: We have a, ORAU has a, there's
15	an OTIB, I can't think of the number right
16	now, but there's one that does cover wound
17	intakes.
18	DR. LIPSZTEIN (by Telephone): Yes, and
19	that's to cover when you have the cancer of
20	the lymph nodes.
21	MR. GIBSON (by Telephone): This is Mike.
22	Did I miss on the wound monitoring, did they
23	actually use a wound, or did they use a
24	bioassay sample after the fact of a wound?
25	MR. GRIFFON: Well, I think they had a field

1 technique to do both, yeah, both is the 2 answer. 3 MR. SHARFI: Yeah, there are reports inside 4 the claimants' files that had the wound 5 counting data and then they also did a lot of 6 full bioassay. 7 MR. GRIFFON: But his report kind of says, 8 you know, he wonders how many of these went 9 undetected, and that sort of raised my 10 question of whether, my question is not 11 necessarily a wound TIB because the assumption 12 is wounds happen. But we're looking at the case where we didn't know it was a wound. 13 14 MR. SHARFI: If you want to look up the same 15 bioassay result in an intake from an 16 inhalation versus an intake versus a wound 17 incident, I don't know if the scenario where 18 the inhalation does not give you a bigger dose 19 than --20 MR. GRIFFON: Okay, I got to run --21 MR. SHARFI: You can use that wound TIB to 22 run your scenarios and if you want to test 23 that. 24 MR. GRIFFON: Yeah, okay. I just wanted to 25 raise it while we're here so check it.

DR. LIPSZTEIN (by Telephone): If you assume there was an acute intake and knew there was a wound, and you take the bioassay data, and you go back and everything is okay. When there is no indication that there was a wound, so but -

MR. GRIFFON: That's why I questioned that.

DR. LIPSZTEIN (by Telephone): -- all the, let's say all results were below the detectable limits, and then acute inhalation intake and that will be very unfair to some because if you (unintelligible) it's okay in a short time period, but if you take it for a long time period then it's (unintelligible).

MR. GRIFFON: Okay, maybe I can ask Joyce if you can look at that as well as Jim or Brant, whoever on your team wants to look at it.

of the problems is that the wound model which is being done by NCRP, it's almost ready, but it's not ready yet. So you have the wound OTIB which is good for now, but it doesn't cover everything that you have. If someone has a cancer of the lymphatic system, it's not well covered by this wound OTIB. I think in a

short time NCRP is going to publish a model for a wound and will solve everything.

MR. GRIFFON: So let's, it's on our radar, and we'll have SC&A and NIOSH look at that, but let's move on to our last items. I know we'll be done by 5:00. I know we will.

## NEUTRON ITEMS

Neutron issues, I'm going to need your help, Joe to kind of frame it, and maybe Ron can weigh in.

MR. FITZGERALD: Yeah, this has been a longstanding discussion we've had. And I think there's no one specific fundamental problem with the information that has been provided in terms of how neutron doses are being estimated, just that there were a number of pieces of data and information that we wanted clarified. And I think Brant itemized these very well in terms of five action items that I think the work group's very familiar with. And frankly, at this stage, we've had Ron Buchanan working directly with ORAU to see if we can actually put this to bed because essentially it's information that we need some clarifications, interpretations, corrections

and it's sort of a grab bag of different items which, I think at this point, we believe isn't moving in an SEC direction, but we wanted to make sure of that fact. And that's what we're really looking to, from this discussion, to understand if that's the case. So Ron, with that introduction, how does it look?

MR. GRIFFON: And knowing that we all want to head for the airport.

MR. BUCHANAN (by Telephone): Yeah, we've been going over this neutron issue and the main thing was two areas I wanted to look at is the proposed procedures, recommended procedures by NIOSH, claimant favorable and workable. They produce reasonable results for the doses assigned. And so that involved reviewing all the OTIBs and procedures and a lot of intertwined documents. And at this point, of course, and then we came along and we had OTIB-58 recently released a week or two ago which was a revision of the earlier one.

My review to this shows that OTIB-58 has made an effort to do a reasonable coworker dose model, and coupled with the NDERP report and OTIBs-50 and -52 and some others. And my

take on it at this point is that I'm trying to clarify where some of the data came from that's entered in the tables.

I don't necessarily have a problem with the procedures, and I don't necessarily have a problem with the data. I just have not yet been to all 14, and I'm working with Matt and some of the others and Brant has forwarded some of the answers. I got those yesterday or the day before, and then I just opened the door and received the CD with six megabytes of neutron gamma data on it about an hour ago, and I haven't had time to digest that.

And so I know everybody's wanting to close this meeting so what I, my bottom line is that at this point, that SC&A, the review of the recommended procedures indicates that there are no outstanding SEC issues at this point with our understanding of the recommended procedures. Now that hinges on the fact that the data put into these procedures is reliable and adequate. And I did not address all that issue as we spent most of the day on that.

At this time I'm looking at the

1 procedures do they, are they claimant 2 favorable, produce reasonable results, and at 3 this time I do not see SEC issues with the 4 recommended procedures and the data I've 5 reviewed today. Now we haven't completed that 6 data accuracy and availability issue 7 completely, but that's where I stand on this 8 point. 9 MR. FITZGERALD: Which reflects what we 10 talked about with the coworker model with 11 different prongs we're looking at. So he's 12 been on one prong we've been dealing with. MR. GRIFFON: We're looking at the data 13 14 issue on another prong which we spent a lot of 15 time on. 16 MR. FITZGERALD: So it sounds like really 17 The loose end is to look at the data 18 behind the new tables and the new OTIB 19 revision? 20 MR. BUCHANAN (by Telephone): Well, actually 21 there's the tables, yes, there's two 22 additional tables in the OTIBs that I'm 23 looking at and also the original tables which 24 you didn't have the data for before. So the 25 four tables I'm looking at, plus there's three

1 additional tables for the construction workers 2 which I have not touched on that aspect 3 because I'm still trying to sort out the 4 regular workers, and I'll get to that after I 5 understand the first tables. 6 And so I have read the response of 7 NIOSH on my questions, 14 questions I think, 8 and I'm reviewing those. And I'm going to 9 send another couple clarification questions to 10 Matt here in the near future and also review 11 the CD and make a decision whether I 12 understand where all the data's coming from, 13 if it looks like it provides favorable 14 results. 15 MR. GRIFFON: And I would also request that 16 if need be, you continue some of those 17 technical calls on the neutron issue 18 specifically. I mean, if you need a follow-up 19 call --20 MR. FITZGERALD: Yeah, we're seeing how this 21 goes and --MR. GRIFFON: -- in the interim, right. 22 MR. FITZGERALD: -- if it turns out that the 23 24 disk and the additional information is 25 sufficient, I think then we'll just wrap it

up.

### D AND D (TIB 0014)

MR. GRIFFON: Last but not least, the D&D worker question.

MR. FITZGERALD: Yeah, I think we had some pretty productive calls on that. Certainly the information provided by Gene Potter in terms of looking at the top tier dose distribution versus the, I guess two sets really, all the subcontractors as well that were identified as the D&D subcontractors, demonstrated the dose distributions were equivalent and could be in fact enveloped by a common coworker model. I thought we were pretty satisfied and thought that kind of addressed the issue we were concerned about most.

Really what's left on that issue is the OTIB-14 extension of OTIB-38 which is this whole issue of what's the coworker in it. If you buy into again that distribution, which we do, then how are you going to apply OTIB-38 to the D&D era. That's what OTIB-14 is directed at. And there we spent some time looking at that, in particular --

1 Joyce, are you still on? 2 DR. LIPSZTEIN (by Telephone): Yes, I am. 3 MR. FITZGERALD: Okay, well, the only 4 question that we come up with that is 5 certainly workers were reviewed from the 6 standpoint of fecal analysis, and some were 7 reviewed on the basis of in vivo counts, and 8 the OTIB isn't very clear on how those would 9 be accommodated within the model. And I think 10 we just got, what, a few days ago, some 11 additional information that Gene put together 12 or somebody put together on that issue. 13 Joyce, have you had a chance to -- I 14 know it's only been a day or two -- had a 15 chance to look at that? 16 DR. LIPSZTEIN (by Telephone): Yes, I looked 17 at it, and now I'm going to analyze it. 18 basically said that most D&D workers were 19 after '95, and so we don't have to worry too 20 much about the required fecal samples, fecal 21 samples from before '95. So we just have to 22 verify that, and I think now it looks like if 23 it's okay. 24 MR. FITZGERALD: Yeah, this is a case where 25 we looked at the concept, went to the

completeness of data, and now we're looking at the application so in a way I think this will kind of sew up OTIB-38 and the D&D issue at the same time.

#### MATRIX UPDATE

MR. GRIFFON: Now we come to the matrix. I want to walk through every item -- just kidding. I did want to bring up one specific thing on the matrix, item -- we did edit this matrix so I would ask maybe the work group and all others to read through these final actions. And I left, I usually try to track actions so there should be a little chronology in most of these, and Brant reviewed this as well as Joe and Emily apparently so we have this redacted version.

But I wanted to point out on item 18, this is one place where I put a note that it seems like one of these that we might have just kind of forgotten a little bit, or it was inconclusive and I didn't know if we needed to take this any further.

Brant, maybe you can give us some background on this one.

DR. ULSH: Yeah, Joe and I talked about

this, Mark, because I noticed that while the phrase that caught my eye was closure not achieved. And this goes back to what we call detailing analysis, and it was related to the concern that was expressed by the petitioners that sometimes workers would leave their badges in their locker and wouldn't wear them at the radiation area.

And so it was suggested, and we did some preliminary analyses to look at whether or not you see evidence of tailing off of dose that would indicate that the rate at which a worker accumulates dose or at least that's reflected on his badge, tails off. Now we did some preliminary analyses and presented that to SC&A, and it was our conclusion that we didn't really see evidence of that.

And I think SC&A concurred with that; however, they also said that they noted that the sample size was small. And it was. The reason we didn't keep going with that -- I think we discussed it at a working group meeting -- was that it wasn't clear to us that that would really give us a definitive answer to the question.

MR. GRIFFON: I do remember that.

DR. ULSH: Because it wouldn't tell you whether or not the worker was actually pulled out of the area or whether his badge was only pulled out of the area. So we decided, I think as a group, to pursue this on some of these other prongs in terms of the safety concerns and the data integrity.

MR. FITZGERALD: Yeah, I think that's kind of where we left it although I agree that was kind of, you know, that prong was left, and we went, pursued it elsewhere. So it was sort of a loose end, but I don't disagree with that conclusion. We went as far as we thought was useful to go through, and this was maybe a less perfect way of reaching that conclusion. We don't disagree with the analysis at the time.

Ron, do you want to add anything?

MR. BUCHANAN (by Telephone): No, not

particularly. We agreed with what was stated

by NIOSH. It's just that we didn't feel that

you could take those few cases and extrapolate

into all the years to all the workers.

MR. GRIFFON: So my question sort of is do we in our other prongs get at this question.

I guess we attempted to. Did we?

MR. FITZGERALD: We attempted to. I mean, certainly in the work that Kathy did trying to run these individual situations where there were some allegations about workers not wearing badges and what have you. I think we looked at that certainly on the completeness side. It's kind of a, you know, how many prongs do you need in order to put something like that to bed? I don't know. That's kind of a judgment call.

MR. GRIFFON: And there seem to be ways to possibly get at it. I don't know if we've explored all of them, but if there's people that are, you know, if you scan through the annual summary data, and you see people near the limits, if we have their files already pulled, you know, you might look at if there was any indication, I mean, I see log book indications that people were pulled from areas. It certainly looked like restrictions were applied especially on the internal side. I saw notices of that.

1	DR. ULSH: Yeah, and sometimes you'll see in
2	the worker RAD files memos that say that there
3	are work restrictions. I don't, it's nothing
4	I've looked for systematically, but I have
5	seen examples of that.
6	MR. GRIFFON: So I don't know, this was
7	derived from an individual allegation? Is
8	that correct?
9	MR. FITZGERALD: Yeah, it was actually
10	affidavit issues as well as some of the worker
11	input, but
12	MR. GRIFFON: You mean did they give us
13	specifics of individuals where it happened?
14	MR. FITZGERALD: No.
15	MR. GRIFFON: No, they just kind of said
16	they thought this practice
17	MR. FITZGERALD: Right.
18	MR. GRIFFON: And we haven't seen any
19	MR. FITZGERALD: Not in terms of running it
20	down from safety concerns and data integrity.
21	DR. ULSH: And this is on the overarching
22	issues list. This particular issue.
23	MR. GRIFFON: Right.
24	MS. MUNN: Well, and the other side of that
25	coin actually is how many workers would have

1 been expected to be included in a group that 2 would have done this. 3 MR. GRIFFON: Right, how much will that 4 affect the overall class. 5 MS. MUNN: Exactly, exactly. 6 In fact, Wanda, that prods my DR. ULSH: 7 memory. I did talk about this issue in 8 evaluation for a presentation in April. 9 MS. MUNN: I think you did, uh-huh. I think 10 we did go over this. 11 MR. GRIFFON: And if we had heard, I don't 12 know that we heard many, many people saying, 13 yeah, this happened all the time. I don't 14 think that's the --MR. FITZGERALD: We didn't find evidence of 15 16 a systemic issue. 17 MR. GRIFFON: So I don't know there's any 18 further action we can do here other than that 19 in the matrix I'll note that we tried to pick 20 this up and cover this in the individual RAD 21 file reviews and data completeness and other 22 prongs as we're describing. 23 MS. MUNN: At the time I read this, I saw 24 the further discussion by the work group. My 25 instant reaction was, why? I thought we had

1 talked about it already. 2 MR. GRIFFON: That was the only, I think we 3 were in agreement on the matrix otherwise. 4 DR. MAKHIJANI: The main issue that's come 5 up in this kind of regard was, you know, the 6 going up the zeros in '69 and so on, and we 7 know there was another explanation for that. 8 There was a policy for not reading badges and 9 so on. And we tracked that in a different 10 direction. It did not have the same 11 explanation as that item 18. 12 But I also think Wanda's MR. GRIFFON: 13 statement's important in this regard that 14 we've got to consider how much the class could 15 have been affected if this happened a little, 16 a few times or whatever. MR. FITZGERALD: Well, I think you had to do 17 18 a reconnaissance review just to see whether it 19 had any corroborating evidence. We didn't 20 find any, but certainly we wanted to take it 21 seriously and pursued it but didn't find 22 anything. 23 MR. GRIFFON: I think that's it on our 24 agenda. I would ask if anyone on the phone 25 has any final thoughts. I know it's getting

1 kind of late in the day, but especially the 2 interested parties from the congressional 3 staff. Any comments? 4 UNIDENTIFIED: Could someone keep us posted 5 when that report comes out that was referenced 6 in the last working group call? 7 MR. GRIFFON: We will. I think we committed 8 already to any reports that, from here on out, 9 that once they're through Privacy Act review 10 and approved for distribution, we'll get those 11 to you ASAP so NIOSH has that out there on 12 their agenda, and we'll do that for sure. 13 UNIDENTIFIED: Thank you very much. 14 MR. GRIFFON: Thank you for staying with us 15 all day. 16 DR. MAKHIJANI: Mark, is there a scheduling 17 question about wrapping things up that we --18 MR. GRIFFON: Yeah, I guess, I was looking 19 at the calendar, and we now have until May, 20 but if we walk that backwards, I think we 21 really want to get something to the 22 petitioners and the interested parties by the 23 end of March, first of April. That gives 24 everybody 30 days or so. I think the 25 meeting's in early May if I have my dates

1	right, yeah. So given that I don't know. It
2	seems to me that any responses from NIOSH
3	would be helpful if you had them by, what, the
4	end of February?
5	MR. FITZGERALD: Yes.
6	MR. GRIFFON: Do you think that's, I think
7	data completeness is probably the big
8	DR. ULSH: Let's start with that, the '69
9	section and the data completeness section.
10	MR. GRIFFON: Let's try for that. I mean,
11	we're all working toward the same goal here,
12	same end date. And we also have to keep in
13	mind that any final product from anyone has to
14	go through Emily's review. So that's going to
15	maybe slow us down a little bit, but I know
16	they'll turn around things quickly.
17	MR. FITZGERALD: Well, we're hoping to get
18	as many sections cleared beforehand.
19	MS. MUNN: So are we going to talk about
20	this, are we going to set another meeting
21	date?
22	MR. GRIFFON: Well, we have the Board
23	meeting coming up. I thought maybe we'll, if
24	you want to look at calendars now
25	MS. MUNN: No, I was thinking about, this is

1 not going to be, what we've been talking about 2 here today is certainly not going to be ready 3 at the Board meeting. 4 MR. GRIFFON: Right. 5 MS. MUNN: And so --6 MR. GRIFFON: We're going to have to have 7 something beyond that, but I was going to wait 8 and kind of check in with you guys in 9 Cincinnati at the Board meeting and then say, 10 okay, let's look at our calendars to see what makes sense between February 7<sup>th</sup> and the next, 11 12 our next milestone, the end of February I 13 guess. **DR. ULSH:** Probably not February 7<sup>th</sup>. 14 15 MR. GRIFFON: Probably like the third week 16 in February we'll meet. 17 MS. MUNN: Maybe we could do the first week of March. Wouldn't that give us --18 19 MR. FITZGERALD: That'd be a little tight. 20 See, I'm a little concerned that we have to 21 both accommodate any final changes and go 22 through final tech editing which we didn't do 23 on this one, which we're finding all kinds of 24 glitches. And then certainly make it 25 available to NIOSH for final Privacy Act

1 review so that can then go to the outside 2 world. And the logistics I think we need most 3 of March for is my guess by the time it's all 4 done. 5 MR. GRIFFON: Yeah, I think the third week 6 in February, although I'm not really ready to 7 pick a date. I apologize because I know 8 there's the Savannah River classified meeting 9 that I have to be at somewhere in that third 10 week in February. I think it's the 20<sup>th</sup> and 21st, but we haven't pinned it down. But I'm 11 12 thinking the third week in February. That way 13 if there's any -- does that make sense? 14 MR. FITZGERALD: Yeah, I think that --15 MR. GRIFFON: The final changes by any final 16 things by NIOSH then everything, you know, 17 we're shooting for that end of February, early 18 March. And that'll give you guys a month to 19 try and get something out for full 20 distribution to everyone, right? 21 MR. FITZGERALD: We're kind of looking at 22 maybe a couple weeks to get it to NIOSH so 23 they have a couple weeks to then make 24 distribution. So I think, really, we're only 25 talking a couple weeks from our side,

1 hopefully. 2 MR. GRIFFON: Well, let's look at the third 3 week in February tentatively, maybe toward the 4 5 DR. MAKHIJANI: Maybe the end of February? DR. ULSH: The third week of February for 6 7 another working group meeting? 8 MR. GRIFFON: Yeah. 9 DR. ULSH: End of February for whatever 10 responses we're going to provide. Is that 11 what you're --12 MR. GRIFFON: Yeah, yeah. DR. MAKHIJANI: Well, if there's going to be 13 14 a working group meeting, presumably since most 15 of the action items, I think, are in NIOSH's 16 court at this time, maybe we should have 17 something from NIOSH so we can actually 18 discuss it. 19 MR. GRIFFON: Well, that was the, yeah --20 DR. MAKHIJANI: Well, I don't know. 21 MR. GRIFFON: -- I mean, I don't know if 22 it's worth meeting if, you know, until we have 23 these products so that's the question. Maybe 24 we should just do it the last week in February 25 and then you have your sort of almost final,

1	near final product.
2	MS. MUNN: We're still planning for the
3	first week of March.
4	MR. GRIFFON: I would also say as you
5	Brant, you've been doing this very well as you
6	complete things certainly get them around to
7	us as you can.
8	DR. ULSH: Will do.
9	MR. GRIFFON: That's it, right? Anything
10	else?
11	(no response)
12	MR. GRIFFON: Meeting adjourned.
13	(Whereupon, the working group meeting
14	concluded at 5:03 p.m.)
15	
16	
17	

#### 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 January 26, 2007; 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 26th day of February, 2007.

\_\_\_\_\_

STEVEN RAY GREEN, CCR

CERTIFIED MERIT COURT REPORTER

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