U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES CENTERS FOR DISEASE CONTROL NATIONAL INSTITUTE FOR OCCUPATIONAL SAFETY AND HEALTH

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ADVISORY BOARD ON RADIATION AND WORKER HEALTH

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TBD-6001 WORK GROUP

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WEDNESDAY
JULY 7, 2010

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The Work Group met in the Zurich Room of the Cincinnati Airport Marriott, 2395 Progress Drive, Hebron, Kentucky, at 9:30 a.m., Henry Anderson, Chairman, presiding.

PRESENT:

HENRY ANDERSON, Chairman R. WILLIAM FIELD, Member MARK GRIFFON, Member*

ALSO PRESENT:

TED KATZ, Designated Federal Official ISAF AL-NABULSI, DOE*
DAVE ALLEN, DCAS
TERRIE BARRIE, ANWAG*
HANS BEHLING, SC&A*
SAM GLOVER, DCAS
STU HINNEFELD, DCAS
JOHN MAURO, SC&A
JIM NETON, DCAS
JENNY LIN, HHS
WILLIAM THURBER, SC&A

*Present via telephone

C-O-N-T-E-N-T-S

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P-R-O-C-E-E-D-I-N-G-S

(9:32 a.m.)

MR. KATZ: We will begin with roll call. Just a quick overview of the agenda and then Andy may have to more to say but we are going to both generically be addressing the TBD-6001 procedure and then as time allows, we will be specifically addressing two petitions: one for Electro Met; and the second for United Nuclear.

And then toward the end of the day we will talk about the path forward and also address, with respect to the path forward, Hooker Electrochemical, which we will not get to substantively today. But that's just to let everybody know, just sort of the general landscape.

We may run out of time anywhere in the course of this. There's a lot to cover for a day and I doubt we'll get through it all. There's a long agenda. It is available to people on the internet, on the DCAS website

1	and I think all participants have a copy.
2	So we will begin with roll call
3	and for all the agency-related individuals,
4	Board and others, contractors, please state
5	whether you have a conflict with respect to
6	this site as well, when you respond to the
7	roll call.
8	So beginning with Board Members in
9	the room, with the Chair.
10	CHAIRMAN ANDERSON: Henry
11	Anderson, Wisconsin Division of Public Health.
12	I don't have any conflicts.
13	MEMBER FIELD: Bill Field, no
14	conflict.
15	MR. KATZ: And on the line?
16	MEMBER GRIFFON: Mark Griffon, no
17	conflicts.
18	MR. KATZ: Any other Board Members
19	on the line? Okay and those are the Members of
20	the Work Group. And then NIOSH/ORAU team in
21	the room?
22	MR. HINNEFELD: Stu Hinnefeld,

1	interim director of DCAS. No conflicts at
2	these sites.
3	DR. NETON: Jim Neton, DCAS, no
4	conflicts.
5	DR. GLOVER: Sam Glover, DCAS, no
6	conflicts.
7	MR. ALLEN: Dave Allen, DCAS, no
8	conflicts.
9	MR. KATZ: And NIOSH/ORAU team on
10	the line? Okay. SC&A in the room?
11	DR. MAURO: John Mauro, SC&A, no
12	conflict.
13	MR. THURBER: Bill Thurber, SC&A,
14	no conflicts.
15	MR. KATZ: SC&A on the line?
16	DR. BEHLING: Hans Behling, no
17	conflicts.
18	MR. KATZ: Very good. Then, other
19	HHS or other agency personnel or contractors
20	to the agencies in the room?
21	MS. LIN: Jenny Lin, HHS.
22	MR. KATZ: And on the line?

1	DR. AL-NABULSI: Isaf Al-Nabulsi,
2	DOE, no conflict.
3	MR. KATZ: Welcome, Isaf.
4	DR. AL-NABULSI: Thanks.
5	MR. KATZ: Okay and then, finally,
6	members of the public on the line; there are
7	none in the room.
8	MS. BARRIE: Terrie Barrie with
9	ANWAG.
LO	MR. KATZ: Welcome, Terrie. Very
11	good. And I probably didn't even identify
L2	myself. My name is Ted Katz. I am the
L3	Designated Federal Official for the Advisory
L4	Board and we are ready to get started then.
L5	Andy, it's your agenda.
L6	CHAIRMAN ANDERSON: First,
L7	congratulations to Mark on his appointment.
L8	It's good news that you are on the
L9	line. It means there must not be some big
20	event occurring that we don't know about yet.
21	MEMBER GRIFFON: Yes, nothing that
22	you don't know about. The one you know about

			is	the	only	one	out	there.	But	thank	you
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CHAIRMAN ANDERSON: I really don't have much else. I think we ought to just get started. This, of course, Committee was broken off from the TBD-6000 because of all the different sites being addressed so the only question I would have is, I looked at the matrix and there didn't seem to be any responses yet from NIOSH. Is there? Not yet.

MR. ALLEN: There weren't any written responses.

CHAIRMAN ANDERSON: Okay. Okay. Fine. Good, I wanted to be sure I didn't miss something. So let's go on then and start up.

MR. KATZ: So, the first item is Overview of TBD-6001 from DCAS, just a brief summary and a chance for the Work Group to ask questions about it.

MR. ALLEN: Okay, I think to start with it takes kind of a brief summary of what our contractor did for us as far as TBD-6000/6001. The whole intent at the time was to

gather up some information on the sites that did very similar work and at the time it was decided to divide these into three primary categories with very different types of exposure conditions.

One being uranium metal, largely because it's a very similar type of -- you can only do so many things with uranium metals so all exposure conditions were fairly similar and the smaller sites that did that type of work were pretty abundant. And that became TBD-6000.

TBD-6001 was intended for the other chemical work, essentially with uranium, the processing of the uranium compounds like UF4, UF6, U02, et cetera, and that became TBD-6001 which was -- it's labeled as uranium-refining operations.

There was to be a third one at the time and that was for uranium-ore operations.

That was split out because of the natural-occurring decay products, with the radium,

1	thorium-230 et cetera, make for a very
2	different exposure conditions than refining
3	processed uranium.
4	CHAIRMAN ANDERSON: So, just a
5	quick question. On a product-flow basis, so
6	kind of 6000 was earlier in the process and
7	6000
8	MR. ALLEN: It was just the
9	opposite of what I introduced. It was
10	CHAIRMAN ANDERSON: Okay, so 6001
11	is the ore processing, refining, and then
12	their output went to the facilities that were
13	6000.
14	MR. ALLEN: And then the ore would
15	have been first so, pretty much exactly the
16	opposite of what I introduced.
17	CHAIRMAN ANDERSON: That was what I
18	was wondering. I thought, gee, I had looked
19	through that document, it doesn't sound right
20	to me. Thank you.
21	MR. ALLEN: But that was the
22	thought process at the time. The decision on

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the ore was that it is very different and very limited and the idea of a generic TBD was kind of dismissed partway into that process so a TBD to address that processing was never developed. That was going to be just a site-by-site type of Technical Basis Document.

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TBD-6000, 6001 are both somewhat generic type of documents. The intent is to look at the exposure conditions for a type of work with the type of material and then to apply that to other work sites that did that same type of task with the same materials.

The idea all along was to have this generic and then have an appendix for every site that it would apply The appendix would analyze and document the exposure estimate based on data from particular site and use the generic TBD-6001 and 6000 to fill in the blanks where there was no data. That was the concept when we put all that together.

And about the only other thing is

that SC&A was tasked with reviewing both 6000 and 6001 some time ago. As you know there was a previous Work Group to deal with both has now been split up, obviously and in that Work Group we never really addressed 6001 issues because at the time there was no sites they were interested in that actually used the defaults out of 6001.

As I said, the appendices would use defaults where there was lack of data, but the sites we were looking at, there was no lack of data, there were no defaults used at that point in time.

So even though they were assigned as an appendix to TBD-6001, there was no data, no generic data that was actually pulled into that appendix. And I think that is all the background I have for you. I think that brings everybody up to speed.

DR. MAURO: As a preface, though, also I would like to add one thing, is this, the use of TBD-6000 and 6001 is actually, does

go to the heart of the surrogate data issue and I know we all are aware that this is a very important issue to the Board and to the public, claimants.

So we are really in the trenches on what I consider to be probably the biggest surrogate data issue because as we process this, and we less, or dismiss certain issues, there's always the overarching question, well wait a minute, plausibility, applicability, all the criteria that have been developed by the Board, now officially -- I believe the criteria are official -- is sort of like an overarching issue that we always have to sort of keep in mind as we move through this, whether we are doing it for TBD-6000 or any of the sites, such as United Nuclear or Electro Met.

DR. NETON: I guess I would just point out, though, that these are generic documents and really the test at the end of the day comes to how it is applied to the

1	specific site. The data that are collected
2	and assembled in TBD-6000 and 6001 are not in
3	and of themselves bad or wrong, one needs to
4	make the judgment when you apply that to a
5	particular site: is it appropriate or not.
6	So I think there's a real
7	distinction there. These are certainly the
8	data but until it really gets applied, you
9	cannot judge it against
10	DR. MAURO: Agree with you.
11	CHAIRMAN ANDERSON: It's
12	applicability. It looked to me like it was
13	mostly kind of a library of available data.
14	DR. NETON: And that's what I was
15	going to point out. I was going to get to
16	this maybe a little bit later, but you almost
17	really have to judge it against the individual
18	site because we have three DRs out there now,
19	these little pieces
20	DR. MAURO: Again, we're
21	overarching now, United Nuclear and Electro

Met: I don't know the degree to which they do

1	draw upon 6001 so since this is on our table,
2	I know Hooker does, I know we're getting to
3	Hooker later, but I'm not sure whether I
4	know whether or not so we may not have,
5	I don't know if either of them do. Electro
6	Met? No. I am trying to think of if there's
7	any place where they did that and I don't
8	recall.
9	DR. NETON: I don't know what they
10	are doing in this Working Group.
11	MR. ALLEN: It's Appendix C.
12	DR. MAURO: Yes, way things are
13	grouped, there are I think five
14	CHAIRMAN ANDERSON: But there's
15	considerable data.
16	DR. MAURO: For the particular
17	sites we will be dealing with starting today,
18	Electro Met and United Nuclear, I believe that
19	they didn't use too much if any data from
20	DR. NETON: See, and that's what I
21	was getting at it because if there's a lot of

data, it's --

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1	DR. MAURO: You are right. Good.
2	CHAIRMAN ANDERSON: So, we can
3	knock it all off today.
4	DR. MAURO: We can just knock
5	everything off.
6	CHAIRMAN ANDERSON: Okay, so
7	
8	MR. KATZ: What is SC&A's
9	DR. MAURO: Oh, we're on.
10	CHAIRMAN ANDERSON: You're on.
11	Well, of the, this is basic TBD-6001.
12	DR. MAURO: Just by introduction,
13	Bill Thurber and I for several years, have
14	been really been the heart of doing all of the
15	AWE work, which means TBD-6000, 6001, all of
16	these appendices, all of the AWE work somehow
17	fell with us and a few others and really Bill
18	has been doing the heavy lifting on a lot of
19	this and I'd like to turn it over to Bill, who
20	is, I guess we will start with the TBD-6001
21	matrix and take it away.
22	MR. THURBER: Just a further

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comment on how this intertwines with, how TBD-6001 is intertwined with the appendices, we originally prepared our review about two and a half years ago and it has been since that time that we have looked at some of these sitethe specific appendices or SEC petitions related thereto and in the course of that, a number of problems have surfaced as to how you can actually use TBD-6001 appropriately and where the numbers come from that initially apparent when we did review and we will get into a little bit of that as we go along.

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With regard to TBD-6001, all of the -- virtually all of the internal exposure data come from a journal article published by two guys named Christifano and Harris in 1960 and these gentlemen worked for the AEC Health and Safety Laboratory and in the course of their work between 1948 and 1956 they had accumulated some 20,000 samples, air samples, from seven different AEC locations:

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Mallinckrodt, Harshaw, Electro Met et cetera.

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And so it was on the basis of this Christifano and Harris journal article that the air concentration data and the attendant exposures in TBD-6001 were derived.

Obviously, that's а very rich resource. Some of the problems with it -there are some problems with it, though, for example there are no supporting references so, as with a journal article, you don't really understand all the details of how they averaged their numbers and where all the data came from and whether any of the data was relatable to specific sites; you can't discern any of that from the Christifano and Harris paper. But that's where the internal exposure data from TBD-6001 came.

In terms of external exposure, where the exposure was related to drums of uranium, uranium oxide, UF4, whatever, the drum exposure data in part came from modeling studies using MicroShield and workers standing

different distances from the source, and also in the case of pitchblende, it was actually based on measurements made at Mallinckrodt of workers standing various distances from the source.

In terms of the rest of the operations that are involved in the uranium-refining process, the digestion and nitric acid, the solvent extraction, et cetera, et cetera, the data on external exposure for those operations came from Mallinckrodt, where all of these operations were conducted.

Now a lot of the operations were conducted at other places, as well, but the data that was built into TBD-6001 was based on Mallinckrodt data.

With those additional background comments, let me get into our findings and I think, hopefully people can read them. Our first finding was that -- and this speaks to the Christifano and Harris data -- as we said, while there is a lot of it presented there,

it's	diffic	cult t	to ur	ndersta	nd th	ne pe	digree	of
the	data	and	we	tried	to	vali	.date	the
Chris	stifano	and	Harr	is dat	a, if	you	will,	by
compa	ring	it wi	lth 1	Mallino	ckrod	t da	ta, wl	nich
presu	umably	was a	a suk	oset of	it	but	which	was
well	docume	nted	in th	ne Mall	inckr	odt r	reports	3.

And in doing that, we found that - and there are a number of examples in our
review and I won't go into them -- but we
found that in a number of cases, it looked
like the Mallinckrodt data yielded higher
exposures than the numbers in TBD-6001.

So that is the basis for finding one. Finding two, again --

MR. KATZ: I'm sorry, Bill, could I just make a suggestion, if we are going to - it's probably easier to go finding by finding and have the others have a chance to be prepared on any of these to provide their input on them.

MR. ALLEN: Are you done with finding one then, or --?

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MR. THURBER: Yes.

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MR. ALLEN: of the big One there's some flaws in 6000 and 6001, I mean I'll admit that freely, and the biggest one in my book is what I was mentioning earlier was, the intent was to separate out ore-processing pre-processed uranium refining Mallinckrodt did ore-processing. They processed radium-bearing ores as well as other uranium compounds.

TBD-6000, the flaw I mentioned was that it was definitely not clear in there that this was not to include ore. In fact it mentioned pitchblende and some other things that it really should not have. It has not been used for any kind of ore-processing but it's certainly not clear in TBD-6000 that it is not supposed to be used for that.

DR. NETON: Six thousand one.

MR. ALLEN: Six thousand one, sorry. A lot of the findings in SC&A's report, since TBD-6001 is not clear on that, they

reference Mallinckrodt data that actually is higher due to the ore, the radium-bearing ores et cetera, that should not be applied with 6001.

And obviously there is a -- we would have to revise 6001 to make that clear, obviously, and I think that's going to end up being the --

CHAIRMAN ANDERSON: So is it possible in the Christifano to sort that out?

Or are we sorting it out -- you can do it in Mallinckrodt, but --

MR. ALLEN: It's possible to sort out the tasks and eliminate some of those tasks that could be either and in worst case use ones that are ore in Christifano as a bounding but I don't think we have to do that.

MR. THURBER: Christifano and Harris as a category, do consider ore digestion, which is, after the sampling, is the first step, assuming you have obtained concentrates from somewhere else or you have

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1	obtained pitchblende as a feeding material.
2	CHAIRMAN ANDERSON: The raw data is
3	not available from Christifano
4	MR. THURBER: For ore
5	CHAIRMAN ANDERSON: Twenty-thousand
6	samples or something?
7	MR. THURBER: Yes, yes it is. It is
8	available and they point out, they discuss to
9	some extent in their article the differences
10	in processing pitchblende or concentrates, not
11	only in the ore digestion step, but also in
12	the solvent extraction step, in terms of
13	external exposure. We have to be careful,
14	sometimes we are talking about one and some
15	times the other. But there is, they do
16	consider
17	DR. NETON: Was Christifano and
18	Harris not really a journal publication that
19	was based on a lot of AEC
20	MR. ALLEN: It was Health and
21	Safety Lab
22	DR. NETON: Health and Safety

1	Laboratory reports and to my knowledge it was
2	a fairly thick report that contained a lot
3	more background information than what it is in
4	the journal article.
5	MR. ALLEN: That's what I was going
6	to say, just to clarify what Bill said
7	CHAIRMAN ANDERSON: The article is
8	sort of a summary. I mean, you have got ranges
9	and stuff but it really doesn't help you, so -
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11	MR. ALLEN: We have a lot of Health
L2	and Safety Lab data and I think that's what
L3	Bill was referring to; just to be clear, the
L4	article itself does not contain all the
L5	individual samples.
L6	DR. MAURO: There's 20,000
L7	measurements.
L8	CHAIRMAN ANDERSON: No, I mean
L9	DR. MAURO: In my perspective,
20	external is a lot more manageable issue,
21	simply because, beside measurements, you could
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always resort to modeling, as you know --

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1	CHAIRMAN ANDERSON: Good, we can't
2	do that.
3	DR. MAURO: I know, modeling is the
4	but we are just talking about physics
5	modeling.
6	CHAIRMAN ANDERSON: Yes, I know, I
7	know.
8	DR. MAURO: Which are pretty I
9	mean, you are running MCNP, you run a point
10	kernel, you do it by hand. You really cannot
11	be too wrong, if you know the source.
12	CHAIRMAN ANDERSON: The bounding is
13	pretty easy.
14	DR. MAURO: The internal always is
15	the one that you trip over.
16	MR. THURBER: Okay, are we ready to
17	move on to finding two?
18	CHAIRMAN ANDERSON: So for finding
19	one, are you going to revise something, I
20	mean, is this something that as a Committee we
21	can say, okay, it was identified, it's sort of
22	been addressed, we are aware of it? I am not

sure every time you find -- the effort to, when you write it, as opposed to it's a working document we know about it, is this one that we need to do anything, do we need to discuss anymore or is it basically --

MR. ALLEN: Well, I think the problem I am having is a lot of the findings that SC&A put together, you know, they look for some examples and examples they have found in a lot of cases are this ore stuff, and that's the fault of TBD-6001 not making that clear.

CHAIRMAN ANDERSON: Okay.

MR. ALLEN: But, that means you throw a paragraph in TBD-6001 and the findings don't seem to be valid anymore but they were examples.

CHAIRMAN ANDERSON: So just to keep our matrix alive, it may be worth putting in there, if you agree that's what the issue is, then, if others, these findings, really are reflecting the same issue, then if, in fact,

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the ore is taken out is a -- or the processing or however we want to state that -- is the issue, let's identify that with the finding and then we can kind of lay that out and get a paragraph or we just remember every time.

MR. KATZ: I think you wanted a finding-by-finding response, even if it may apply to more, I think you want to go finding by finding. And so, I mean --

CHAIRMAN ANDERSON: I mean --

MR. KATZ: -- suggested a response to the first finding, which still needs I think to be confirmed that everyone agrees if that's the resolution and then you document that and in the future you look for it to be resolved by actually changing the document.

MR. THURBER: I would, I have no disagreement at all with what David says but I would point out that the comment is broader than just very front end of the process. The validation work, if you will, that we did using the Mallinckrodt data against the

Christifano and Harris data indicated that for other operations like denitration and oxide reduction and recasting, the Mallinckrodt data were higher.

So the comment is broader than just the ore end, there's no problem with the ore end relationship but it's a broader comment.

DR. MAURO: By way of process, the matrix usually is our score card.

CHAIRMAN ANDERSON: Yes.

DR. MAURO: And one of the things I should you should perhaps, you may want to decide, I know like Mark, when he runs his Work Group, likes to prepare the matrix himself or in other cases, you know, we work with NIOSH to prepare the next version of this. In other words, for example NIOSH may provide a response, NIOSH makes a response, or tracking what is going on, trying to find a way to see, this discussion we just had, it's certainly captured in the transcript, but

there's always a degree to which, do we want
to try to capture it, the essence of this
discussion, in the matrix and how we are going
to do that.
I think this is something that the

I think this is something that the work with, would like, I would like to go forward.

CHAIRMAN ANDERSON: Yes, I don't know, Bill, how you feel. Mark, you have been doing it longer than most of us so any thoughts on how we should do this?

MEMBER GRIFFON: Well, I mean, I think, I don't necessarily, I don't think it matters who prepares the matrix so much as -- but I think it would be worth, you know, carrying it through and seeing a written response from NIOSH before you close it out, you know.

CHAIRMAN ANDERSON: Okay.

MEMBER GRIFFON: Because sometimes we have caught ourselves with that, that we think we closed it out in discussions but we,

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you know, never formally closed, and then we have to reopen it and so I think it's always better to, for me, it would be nice to see the written-out response to make sure I am in agreement and then have the Work Group Members vote it through and, you know, then we are done with it.

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CHAIRMAN ANDERSON: Okay, good. So that's the process, I guess, well.

MR. KATZ: I think for the time being I think, if SC&A, they've started this matrix, DCAS can send them a written response documenting -- and they can keep it up to date at this point. Moving down the road, I mean, there's this work underway to do this all online as part of this sort of database effort that's being used for some other Work Groups least for t.he Subcommittee or at Procedures. Eventually that will be expanded and that could be done online.

But anyway, for the meantime, it seems like it's good for SC&A to keep up the

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matrix and to receive input from DCAS for their responses.

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DR. MAURO: On this particular item, there are some issues where let's say it's very specific. I'll give you an example, we may have some of these, where we say gee, I think you guys made a mistake. Your number is off by a factor of five, and you know, can we explain why, and then I say, Dave, I say, yes, you are right, this is not reasonable.

And then at that point, it becomes very simple. NIOSH usually responds with the, we agree that there is an error. We will correct it. And what typically is done, during other procedures and other Work Groups, that item is not really closed. What it is, is it's put in this place that we call in abeyance, which means that okay, all we technically, this is the approach and it will be closed when the next issue or revision comes out where it's fixed, and then it's closed.

I mean this is really a choice you folks have, or we could close it right there and say, listen, it's technically agreed upon that the solution is yes, at some time in the future, we will be revising this particular procedure and we agree that this factor of five needs to be fixed.

This is purely, you know, whether you want to close it on that basis and not to back to it, or say no, let's leave it open and we call it, in abeyance, which means where we all agree but until it's actually fixed in the document itself, we are not going to close it out. Again, this is a choice that each one of you makes for themselves.

MEMBER FIELD: Myself, I would prefer the second.

DR. MAURO: Okay then and that would be what we would do on any others.

CHAIRMAN ANDERSON: Well, as time goes by, then you tend to forget and if it doesn't get --

1	DR. MAURO: That's exactly the
2	reason for it.
3	CHAIRMAN ANDERSON: On the agenda,
4	I mean, one or two, you can keep in your mind,
5	but as it piles up across all the
6	DR. MAURO: But, now, where I was
7	leading, though, is, in this particular
8	finding
9	CHAIRMAN ANDERSON: This will stay
10	open forever.
11	DR. MAURO: This is an important
12	finding. If it was simply that the work that
13	Bill has done in showing the examples where
14	the generic numbers really are not bounding
15	because when we look at Mallinckrodt and as
16	Bill pointed out it's more than just the ore
17	issue. I think the ore issue is going to solve
18	a lot of it.
19	But I think that where I'm
20	heading is I think we need a White Paper for
21	this issue and maybe others, namely, it's not
22	a simple story. There is enough richness to

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this issue where we have, okay, here's the ore and perhaps in your White Paper you could explain, you know, this, the concern that SC&A raised about the examples, for example in our work, where the, where we feel that the 6001 underestimated for some people, because of the ore issue and that would solve that.

But as Bill pointed out, things get a little bit more complicated when you leave the ore issue and you actually get into the process issue which is part of 6001, appropriately, where we also have some problems.

And I think that there's going to need to be some work done initially, let's say by NIOSH, to say, okay, let's take a look at this and see if perhaps the distributions that are currently in 6001 need to be broader, or the median has to be shifted based on a closer look at some of the data that I guess originally Battelle compiled and it was 20,000 measurements.

And bear in mind this is I believe all air sampling measurements and that is always an issue too. When you depend on air sampling measurements, you have to be a little careful and we all ran into this before, whether it was breathing zone, general air, so in other words, where I am getting at is that I think that item one is a simple statement but embedded in it is a richness and I think a White Paper would be appropriate. This is my recommendation to deal with this.

MR. ALLEN: Well, I would agree with you except that I tried to sort through the actual write-up and knowing what I knew about the ore and you guys didn't realize that at the time, and I tried to narrow it down to what was significant and honestly, in the examples, like I said, they were examples, and I realize that, for finding one, I couldn't find anything.

In fact there's a statement in the review that says plant four and six data are

1	significantly higher for ore one operation,
2	parentheses, ore digestion and about the same
3	as Christifano and Harris and all others.
4	DR. MAURO: Well, you know, if it's
5	that simple, I mean, I am not disagreeing with
6	you
7	MR. ALLEN: No, I am not saying
8	that it's that simple but I am saying I am
9	kind of stuck, I think in all honesty it's
10	kind of in your ballpark to take this new
11	information that I have given out today and
12	kind of re-look at the issues.
13	DR. MAURO: Bill, right now, do you
14	feel that there are steps in the process that
15	are appropriately part of 6001 where we still
16	think there are some problems?
17	MR. THURBER: Well, what I
18	mentioned, if you go to section 8.1 of our
19	review, we showed that, for the steps I
20	mentioned before, for digestion, denitration,
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oxide reduction and recasting, that the data

from Mallinckrodt were higher than those from

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TBD-6000 by factors of two to five.

DR. MAURO: So that's what I am saying, I think that it's all in, all I am saying is I think the ball is in your court and with those examples, I guess if you could convince yourself that they are fine, great and then usually the next step is that a White Paper shows up on SC&A's desk or the Work Group and we will take a look and say, oh, okay, it looks like they have fixed the problem.

But right now I think our position is we believe it extends beyond just the ore issue and as Bill just --

CHAIRMAN ANDERSON: Well, we need some, I think we need some written, you know, I think we need a written response and then --

DR. MAURO: And then --

CHAIRMAN ANDERSON: And you know, rather than just talk further here, I think that, you know, and if you think a White Paper is appropriate or not or how -- or if your

points are, compare an actual document which is two and a half years old, it may be that needs to be updated as well so let's get it in writing and then we can go back and forth.

MR. ALLEN: I will go through the issue, review SC&A did and, you know --

CHAIRMAN ANDERSON: Yes. Great.

MR. ALLEN: -- start parsing out that this is an ore, you know, this is ore, this is ore, and try to find what is not associated with it and respond in a White Paper. If I find nothing, I will at least fire off an email to everybody.

MR. KATZ: Well, I mean a White Paper is just a generic term, you know what I mean, a memorandum, whatever, it doesn't --

CHAIRMAN ANDERSON: I mean that would be, but I am not sure we need that amount of effort but let's see what you find and if you think it's all ore issues that make a significant difference, then clearly you need to look at it and try to sort out what

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1	others remain.
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issues, is there any further -
DR MAIRO: And right now we think

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DR. MAURO: And right now we think there are.

CHAIRMAN ANDERSON: You think there are and NIOSH hasn't been through it enough I guess to really be able to --

DR. MAURO: It's good, by the way, as we go through each finding -- who has the action? And when we are done with the meeting, usually what happens is I put together what I believe to be action items and I will send them off to NIOSH, say do you agree and then this becomes our sort of score -- okay, and everyone knows who has the action.

MR. KATZ: SC&A would list their action items and DCAS would list their action items and then you would have the whole pool there.

CHAIRMAN ANDERSON: Okay. Finding two.

1	MR. THURBER: There's only 37
2	findings.
3	CHAIRMAN ANDERSON: Well, some seem
4	to be somewhat related to each other. Maybe we
5	can
6	MR. THURBER: In part, finding two
7	is related to finding one because some of the
8	points we raised relative to finding two deals
9	with ore, deal with radium removal from the
10	ore and things like that. They don't all but
11	again some of them deal with that.
12	We quoted in section 4 a number of
13	other areas like hydrofluorination where we
14	felt that the TBD-6001 numbers may be
15	understating things.
16	DR. MAURO: Were there also some
17	steps that were in the process as you know it?
18	MR. THURBER: Yes.
19	DR. MAURO: Is that part of two or
20	is that?
21	MR. THURBER: Yes, no, it is, it
22	may come up later too but let's touch on

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it. The authors of TBD-6001 said we haven't done solvent extraction yet. We are going to do it. And so, that still remains open.

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DR. MAURO: Another, we have all these findings, and we have a lot of findings, but we are going to find that -- and it looks like they are all separate but they are not separate. They cluster nicely very often.

And here's a perfect example, for example, I think if you're going to be working on a White Paper, try to -- and if they're all linked, if there's coupling, that makes sense because then there's a story here. For example the story is perfect, you are going to start talking about the steps in the process.

Let's say there are 10 steps to the refining process, the first three might be ore-related, but then the rest may be, here's step where it's addressed what problem because we think it's not a bounding number that you're using or you captured the because have upper end we

examples.

But then there's another aspect to it. There are steps in the process that are not even explicitly addressed in TBD-6001 or perhaps Christifano and Harris.

So this is all, these are all related. This all has to do with the steps that comprise the process so I don't think we should hold ourselves hostage to the findings the way they are. If you find a way of putting a White Paper together that tells a story that knocks off three, four, five findings in one shot, that's great.

CHAIRMAN ANDERSON: And the other thing, as we go through these, that would be, I mean the real guts of what we need to do are the three sites that are in the appendices so, as you say, with 37 here or however many, some of these, like, what we just talked about, if that directly impacts the United Nuclear, Hooker or those, then that, I would say that pushes that them up to be addressed earlier

than these other --

DR. MAURO: In terms of priority of issues.

CHAIRMAN ANDERSON: Priority of issues so, you know I think there's a lot of work, I mean, we have already identified a couple of things where there's already work for somebody. So you know, if we don't need to do it quite as rapidly, let's identify that now because I think I would like to at least in a meeting or two, if we can, get through the sites that we need to address, unless they are directly impacted by some of these.

Because then if we are, then we need to knock these off before we apply that to the other sites.

DR. MAURO: I think we are going to find, I mean you bring up an interesting point in principle, if the real thrust here is let's deal with United Nuclear and Electro Met and the two of those don't even depend on TBD-6001. You know we might be, perhaps we

1	shouldn't have started here.
2	CHAIRMAN ANDERSON: No, I think we
3	needed to start here. As we go through a few,
4	say "and this relates to"
5	DR. MAURO: It's Hooker, that's
6	going to be the one that is going to get
7	hammered.
8	CHAIRMAN ANDERSON: Well, let's
9	identify that, even though today we are
10	talking about these two.
11	MR. THURBER: Well, we are going to
12	show you an example of how an appendix is
13	intertwined with the numbers in here to give
14	you a feeling as to how those things meld
15	together or attempt to meld together.
16	CHAIRMAN ANDERSON: Okay.
17	MR. THURBER: Finding three?
18	MR. KATZ: So finding two, is sort
19	of wrapped up with finding one, is that what
20	it is?
21	MR. ALLEN: Same kind of thing like
22	John said, I'll try to see if I can intertwine

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responses into one White Paper then I'll do that.

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THURBER: Good. Finding three MR. deals with an external exposure question and namely it's how you estimate the external dose to a worker who is standing on a contaminated surface and this is also included in TBD-6000. At the time we prepared our review, we didn't think it had been appropriately addressed. Since then, David has written a White Paper which was presented to the TBD-6000 Work Group and as indicated in the third column there, we satisfied that this issue had been were addressed.

There was a minor comment in our response there, if you will, that there was -- it would be helpful if there was some explanation as to why seven days was used in TBD-6000 while the year, I'm sorry, there's a typo there. It should say, "while a deposition period of one year was used in TBD-6001."

But the basic concept of using a

terminal settling velocity over a period of time to come up with a surface contamination level has been addressed.

DR. MAURO: Yes, this was a longstanding concern that I raised quite a while ago. The fundamental approach -- I think there is a point where you conceptually understand these things, because you can see, this thread runs through so many sites.

The fundamental approach that is taken in TBD-6000 and 6001 is that, if you are concerned about the accumulation of uranium on surfaces, one way you could -- and you don't have good measurements of what that is -- one way you could estimate it is, is well, if you know the dust loading in milligrams per cubic meter, you can assume that that dust is falling out of the air, settling out of the air at 0.00075 meters per second and that's a good deposition rate, the particles that are of this nature, like five micron particles.

I originally said, gee, you know,

but that's not how surfaces get contaminated. They don't get contaminated by just this respirable dust that is falling on surfaces. They get contaminated because of these big flakes of junk coming off machining operations.

Well, it turns out I was wrong, you know, this was an intuitive thing, I said I don't believe it. But David took the data from a very good study that measured the amount, the rate that stuff is falling out of the air as 0.00075 and showed that that's conservative.

So we accept that that deposition rate -- so if you know the dust loading, you can, from the dust loading alone, figure out the rate at which milligrams per second per meter squared, the rate at which it's coming down can be estimated, very reliably, with the approach that they are using.

Now, the only issue we have is that, well, you have to assume some time

1	period, how long is that going to go now. And
2	I think our question is, seven days we know
3	from the Adley report, that equilibrium
4	this is coming down, right? It sure is coming
5	down but at the same time it's leaving and we
6	know that, in the Adley report, it was longer
7	than seven days, where we believed things
8	became sort of stabilized, where the
9	accumulation sort of stayed the same, the rate
10	of deposition equaled the rate of removal, and
11	I think seven days is too short.
12	So we still have an issue here,
13	but it's a narrow one.
14	MR. ALLEN: So you're actually, I
15	mean, from that White Paper, the time frame,
16	depending on what parameters you ratio et
17	cetera, it was anywhere from 5.8 to 27 days.
18	DR. MAURO: That was that short,
19	27?
20	MR. ALLEN: TBD-6001 uses 365 days.
21	The seven days was in TBD-6000.

DR. MAURO: Oh, okay.

1	MR. ALLEN: Why there's a
2	difference, you are right
3	MR. THURBER: It would seem to me
4	that it would be good for everybody at some
5	time, if there, if TBD-6000 and TBD-6001 are
6	set side by side and differing assumptions are
7	reconciled. But if
8	DR. MAURO: This is tractable by
9	the way, very often, I mean this is really
10	coming to some agreement that, you know, what
11	assumptions are we there is some assumption
12	that makes the most sense.
13	CHAIRMAN ANDERSON: So is the
14	assumption then, if you use seven days, that
15	there is seven days' worth of accumulation on
16	the floor?
17	MR. THURBER: Right. And it stays
18	there.
19	MR. ALLEN: And it stays that way
20	with no removal.
21	CHAIRMAN ANDERSON: And you are
22	calculating the dose not from the particles as

they are dropping, you'll have dust in the air 1 2 that is emitting external --3 MR. THURBER: Yes. 4 CHAIRMAN ANDERSON: Ι mean the 5 amount of --6 ALLEN: The route of exposure 7 that is accounted for but this particular just the 8 issue is amount of contamination. 9 10 DR. MAURO: External exposure. Good way to think about it, again, overarching, you 11 12 are going to run across this every time. When 13 you are in this kind of working environment, there's submersion, there's the 14 external 15 exposure you experience because you are in a 16 cloud. CHAIRMAN ANDERSON: Yes. 17 That dose is 18 DR. MAURO: always 19 very, very, very small and easy to calculate 20 and it never contributes. The other thing is the stuff that accumulates on the surfaces. 21

That is -- and you are standing there, so you

are getting both beta and gamma radiation from it. That's a little bit more significant and that becomes important, not only during operations, while the work is going on, but it becomes the residual radioactivity that people are exposed to after operations start, stop, and so that becomes a very important issue.

And right now our position, again this is a recurring theme, is that we believe the approach they are using is good, can be applied for external exposure to residual radioactivity on surfaces, not only during operations but also during the residual period as a way to place a bounding estimate.

That third, and that's an important contributor, but by far the biggest contributor to external dose is standing next to a 55-gallon drum of yellowcake, of ore or a slab of uranium or uranium rods. That's the big driver.

And on that aspect, we have long since, and this goes back to TBD-6000, we have

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1	long since resolved that. In other words, we
2	are all in agreement that there are
3	measurements and there are models that confirm
4	the measurements and so I mean, I think that
5	that aspect, that third piece of the external
6	exposure contribution is by and large
7	resolved.
8	The amount of the surface is close
9	to being resolved if you could agree how long

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allow this stuff you are going to accumulate. Is it one year? Is it 28 days? But I think once that's resolved that problem goes away.

And the other one, the external exposure from the cloud, we agree with the way we do that and not only that it doesn't matter anyway because it never contributes.

Yes, well CHAIRMAN ANDERSON: that's, that's --

DR. MAURO: I'm sorry --

John, this MEMBER GRIFFON: is Mark.

1		DR.	MAURO:	Yes
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MEMBER GRIFFON: I just thought it might be useful to -- the, you are correct, we discussed this in TBD-6000, I guess. But you might want to give a little background, especially for Bill and Henry on the Adley report and I think NIOSH developed a White Paper off of that, right, to sort of support their position on the deposition?

But my other point here is that it is, as I understood it anyway, it all hinges on this one study, this Adley report that was done in Hanford, is that correct?

DR. MAURO: Yes.

CHAIRMAN ANDERSON: So it, was it a laboratory story?

MEMBER GRIFFON: So I think it's worthwhile for them to get familiar with that report, at least to -- if you can give me an overview that would be great.

DR. NETON: It was a little more than that.

1	DR. MAURO: It's a great report.
2	MR. ALLEN: It was a work area at
3	Hanford, there were various tasks and I'm
4	trying to remember.
5	DR. MAURO: It did everything. It
6	is a very large operation
7	MR. ALLEN: In a small area.
8	DR. MAURO: Hanford metalwork
9	facility, something remelt and just
10	about every type of uranium processing
11	activity is there and an immense amount of
12	measurements were made, every aspect, every
13	step in the process was measured and data were
14	gathered including setting plates out to
15	measure the rate at which uranium accumulates
16	from falling out.
17	And that was done over an extended
18	period of time. I forget how many plates were
19	set out over different seasons, to see if
20	winter is different than the summer. It is
21	quite a report, quite frankly. When I read

that I said, this is sort of like the Holy

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Grail. With this document you have the data indeed to determine whether or not reports like TBD-6000 and 6001 hold up and have the data behind it that says, yes, these are good number. So we rely very heavily on that, that and certainly Christifano and Harris and there one other one. There are several source documents important that are underpinning of everything we have talking about, not only on TBD-6000 and 6001 but also on many of the appendices that we have been talking about.

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CHAIRMAN ANDERSON: Okay.

DR. MAURO: I know I am talking a lot but there is one more thing that is important, and this is this, from the external point of view I made it a little too simple. The idea that you have to get a slab of uranium. Now we know that if you have a slab of pure uranium, it's two mR per hour. That's it. That's what you get, it's a penetrating dose. And if you hold your hand against it,

it's about 200 mR per hour, penetrating and non-penetrating dose.

Now, but there's one exception. And we thought that this is more for TBD-6000. It is when you have freshly cast uranium ingots, you do get a build-up of some of the shorter-lived progeny of the uranium, namely thorium-234, 234 or 238, 234, on the outside of the ingot, which creates a field that is 10 to 20 times higher than the numbers I have just told you, very unusual metallurgical phenomenon.

Bill's a metallurgist and to this day you say you are not quite sure why that happens, but it happens. It doesn't always happen but it happens often. So that's a little nuance to external exposure that's been a fly in the ointment but I think we have got to the point in TBD-6000 where that issue has been resolved.

I am trying to paint the broader picture. We sort of left TBD-6001 but when I

1	was talking about external I wanted to make
2	sure that everybody had a sense that a lot has
3	been accomplished in coming to grips with a
4	lot of these issues.
5	CHAIRMAN ANDERSON: So what we have
6	really to resolve here is seven days versus a
7	year.
8	DR. MAURO: That, for this issue,
9	yes.
10	DR. GLOVER: One thing I would like
11	to mention. Many TBD-6000 sites are mom and
12	pop shop, small, short, they may only have
13	four days of rolling in their history, and so
14	I don't how many TBD-6001 sites have a similar
15	short-term operations. There may be some very
16	different reasons why you would perhaps have
17	some of these well, different, they only
18	roll, they only have four days of operation,
19	that's it. Why do I use a year?
20	DR. MAURO: And that will be good.
21	Right?

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NETON: You have a year of

1	operation, typically you are going to have air
2	samples taken.
3	CHAIRMAN ANDERSON: Yes, I hope so.
4	MR. THURBER: There's no question
5	that the assumption used in TBD-6001 was
6	conservative.
7	CHAIRMAN ANDERSON: Yes.
8	MR. THURBER: The 365 days.
9	CHAIRMAN ANDERSON: Yes, okay.
10	MEMBER FIELD: Let me ask a
11	question about the contaminated dust. Is
12	contaminated dust always the same or there's
13	room for the score to vary depending on the
14	source?
15	MR. ALLEN: This TBD-6000, 6001
16	are for uranium operations.
17	DR. MAURO: And we are talking
18	about basically a five micron, AMAD, uranium
19	oxide. It could be type M, it could be type S
20	and that's basically
21	CHAIRMAN ANDERSON: So, it's not
22	total dust we are really

1	DR. MAURO: It's not total well
2	the reason, certainly, there are probably
3	other dusts but these operations generate
4	dust.
5	And the measurements could be in
6	milligrams per cubic meter and we assume that,
7	you know, all the measurements and then
8	CHAIRMAN ANDERSON: And the
9	assumption is it's all
10	DR. MAURO: Yes. Well, they take a
11	sample and they do a fluorometric analysis or
12	they will do a gross alpha analysis so you
13	know it's uranium.
14	DR. NETON: There's one slight
15	added twist that after 1952 it could be
16	recycled uranium as it went through some
17	transuranic contamination to a small extent.
18	MEMBER FIELD: And the
19	concentration of the uranium in the dust is
20	not a factor?
21	DR. MAURO: It's all uranium oxide.
22	DR. NETON: When you say uranium

1	dust, you mean all uranium.
2	MEMBER FIELD: All uranium.
3	MR. THURBER: Or it's reduced to
4	that in the analyses, one way or the other.
5	DR. MAURO: And it's not an
6	unreasonable assumption because these places
7	were
8	DR. NETON: Literally there were
9	visible clouds of uranium in the air. You get
10	to 30 milligrams per cubic meter you can see
11	it pretty easily.
12	CHAIRMAN ANDERSON: Yes. Just as
13	the uranium does it stick to the floor at
14	all? I mean is it like
15	DR. NETON: Become fixed?
16	CHAIRMAN ANDERSON: Yes, so that
17	you, you know, it's like if you have lead, you
18	would end up with a slick floor from the
19	MR. ALLEN: It's not as malleable
20	as lead.
21	CHAIRMAN ANDERSON: No, I know
22	that. But the issue is when you say it

accumulates over a year, if they were cleaning on a regular basis, it still would have some residual that would be, you know, in the cement floor.

MR. ALLEN: Right, and that's been the rub all along kind of, it's -- I mean, you get cracks in the floor, some of this stuff is very fine powders and it can get down in there and not come up. The assumptions in the TBDs are it's all essentially loose and available for resuspension and that was, when we looked at Adley, that allowed us to get the rates and the actual contamination versus airborne concentrations et cetera and get a better handle on that.

DR. MAURO: Bear in mind, we are talking about AWE facilities that operated in the late '40s, early '50s, maybe up to the late '50s, where these are very dirty operations and this stuff accumulated on the floor, they didn't clean it up every day. And when you talk to workers, what was it like,

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you could see it walking, you would kick it and it would come up. It is not like -- you could have a significant amount, but you don't see it, you know, you could see it, this is it was re-suspendable. like -- and It gets crunched on and even if it was large а particle that fell, it gets crunched up and it becomes certainly re-suspendable.

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Now later on, when you have decontamination, they will say, you didn't remove that and do some clean-up, well then there's always an issue, okay, how much of it is re-suspendable now, and you eventually will encounter some of that when we get to some of these sites.

CHAIRMAN ANDERSON: Moving right along, so you guys are going to resolve one year versus --

MR. THURBER: It's not relevant to this. It's just a matter of, it's more relevant to TBD-6000's in a sense, to explain why you picked a short period.

1	CHAIRMAN ANDERSON: Yes.
2	MR. THURBER: But 365 days for this
3	group of sites is certainly conservative.
4	MR. ALLEN: In any case it's a
5	different document altogether, I mean, I
6	honestly think, and I agree that the
7	documents, the assumptions made should be
8	consistent, you know, throughout the
9	documents.
10	MR. THURBER: Yes.
11	MR. ALLEN: But in this particular
12	case, I think we can close this particular
13	finding out, for TBD-6001.
14	MR. THURBER: As far as TBD-6000, I
15	am satisfied.
16	DR. MAURO: The only point I wanted
17	to make is, your starting point, though, is
18	airborne dust loading has been, given that the
19	airborne dust loading is a good number for a
20	particular category of operation, that
21	approach works. But we don't necessarily agree

loading

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was

1	selected for a particular operation on the
2	TBD-6000 over the distribution that was
3	selected is necessarily the appropriate
4	distribution.
5	So, you see what I am, that aspect
6	of it
7	CHAIRMAN ANDERSON: The cascade,
8	the front end of the cascade
9	DR. MAURO: Exactly
10	CHAIRMAN ANDERSON: might be
11	problematic, Okay.
12	MR. KATZ: So, is this finding
13	closed?
14	DR. MAURO: I would say yes.
15	MR. KATZ: This is for this Work
16	Group.
17	MR. THURBER: Yes. For this Work
18	Group, this finding is closed, because what we
19	are really agreeing on is that the use of a
20	terminal settling value of 7.5 times 10 to the
21	minus four meters per second is a good number.
22	Now, the rest of the pieces that

1	go to get to an exposure, a dose, as John
2	says, some of those are still open issues.
3	MR. KATZ: But that's separate. MR.
4	THURBER: Right, that separate.
5	MR. ALLEN: Clear as mud?
6	CHAIRMAN ANDERSON: No, no, I mean,
7	well in the initial response, NIOSH talked
8	about seven days but in 6001, it specifically
9	uses one year.
10	MR. THURBER: And that's fine, it
11	might increase
12	CHAIRMAN ANDERSON: I mean this
13	probably started when 6000 and 6001 were
14	together, so as far as closing it out, you
15	need to be very clear it's 6001.
16	MR. KATZ: Right, this Work Group
17	can't close 6000 issues.
18	CHAIRMAN ANDERSON: Do we need a
19	vote on that?
20	MR. KATZ: Just as long as it's
21	clear for all Members that it's closed.
22	CHAIRMAN ANDERSON: Anybody object

1	to	 Mark?
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MR. KATZ: Mark? Mark are you still with us? Are you on mute? We may have lost him.

DR. GLOVER: He's trying to find a copy -- we are going to send him a matrix copy, apparently he doesn't have one so we are going to try to get him.

CHAIRMAN ANDERSON: Oh, okay.

DR. GLOVER: We are taking care of it.

CHAIRMAN ANDERSON: Okay.

MR. THURBER: Finding four?

CHAIRMAN ANDERSON: Finding four.

MR. THURBER: In TBD-6001 there are two, comprehensive summary tables relating to external exposures and how external exposures should be calculated for various job descriptions, denitration operator, solvent extraction operator, whatever. But, and within those categories, the document provides how to calculate the external dose for operators, for

laborers, for supervisors and for clerical personnel. It provides data on various durations of work weeks because in the early days, a lot of the operations were probably on a six-day work week.

So these are quite comprehensive tables. The fundamental problem we have had with them is that the transparency and the traceability of the data that is in them is not apparent from TBD-6001 and when we get done with this, I would like to give you an example of that.

So if we could, I would like to pass on from this until we get through the rest of the list and then come back to this with some specific examples of the transparency/traceability issues related to external exposure.

Oh, the tables also cover the different external exposure modes that we are talking about: direct contact; submersion in a cloud; standing on a contaminated surface and

1	it provides exposures for all of those
2	different environments.
3	CHAIRMAN ANDERSON: And the data
4	source for that is?
5	MR. THURBER: The data source for
6	the external exposure is, a lot of it comes
7	from Mallinckrodt where there were comparable
8	operations, or it comes from, in the case of
9	exposure to 55-gallon drums of whatever, it
10	comes from MicroShield calculations, or from
11	actual measurements of 55-gallon drums
12	containing pitchblende ores, that impacts on
13	the ore thing a little bit, but so that's
14	where the data, virtually all the data in
15	these tables come from.
16	MR. ALLEN: Yes, plus the type of
17	calculation we were already talking about, the
18	air submersion, the surface contamination
19	MR. THURBER: Yes, that's for the
20	direct
21	CHAIRMAN ANDERSON: The source-
22	terms is more than the calculation

MR. THURBER: Yes, that's for the direct exposure, the standing-on-a-cloud comes from the calculations we have discussed, the submersion comes from the kind of calculations we have discussed.

CHAIRMAN ANDERSON: Okay.

MR. THURBER: So, I would like to pass --

DR. MAURO: I know, but when you talk about Mallinckrodt data, you are talking about, we have a group of workers that we know were doing a particular operation, they were wearing a film badge and you can see what their exposures are, so that is one way you can get at what kind of exposures.

Because those exposures reflect not only the radiation field the person is in, but also how long they worked and how close they worked. Because you know, when you run the model, what it tells you is, here's your millirem or microR per hour, at this distance, at this distance, and you

know those numbers are good.

The real question is, well, for real workers, how long were they standing -- how long were they in contact? And that's where the Mallinckrodt experience helps you get an insight of what assumptions should be made.

So it's like, the field is one thing, but those are based on physics calculations and it's straightforward. The big question is, how long do we assume a person is, and at what distance, from this -- what source, the size of the source, and the type of source?

So, and I think, correct me if I'm wrong, it's both of those that is the Mallinckrodt experience and the models that are married and that's how these tables were produced, the 7-3, I believe, it's for any given worker, I would say it's like you said, a denitration operation or whatever it is, the exposure distribution that's in there for,

1	let's say a supervisor, a matrix table, that
2	has embedded in it some assumption regarding
3	duration of exposure, like hours per year?
4	MR. THURBER: Now, I believe that
5	those are based on actual measurements.
6	DR. MAURO: Okay, so, all right,
7	all right.
8	CHAIRMAN ANDERSON: So this is
9	really a surrogate exposure?
10	MR. THURBER: Yes.
11	CHAIRMAN ANDERSON: Because you
12	are, I mean, mostly your table is generated
13	based on the distance and the time and stuff
14	from Mallinckrodt. Then the question is, are
15	the other facilities, do the workers spend,
16	are they so identical and the process is so
17	identical that worker A in nuclear spends the
18	same amount of time afoot from whatever it is,
19	and
20	MR. ALLEN: Like we said from the
21	start, TBD-6001 is surrogate data, that's
22	pretty much the purpose of the document and

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then the applicability, as you're saying, is essentially a site-specific thing. You can't really judge it on the generic document basis.

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CHAIRMAN ANDERSON: Okay. We will come back. Your job is to remember. We're going to come back at 4:30 and you're going to be stuck here.

THURBER: Okay, finding number five. with This again deals inhalation exposures and the issue here is this. Christifano and Harris provide data from 1948 through 1954, Ι believe. Some of these operations began as early as 1942. So the question is, and certainly there is a lot of anecdotal evidence that suggests that in the very early days things were much worse than they were at the time that Christifano and Harris began their measurement protocols.

And clearly during the course of the time that Christifano and Harris did their air sampling, beginning in 1948, there was continuous improvement, continuous lowering of

the dust levels.

So, in the absence of data prior to 1948, NIOSH is -- well, the NIOSH approach was this: they took the average value for that period where they had data, 1948 to 1954, and they said the data in the first year where we have data was 6.8 times higher than the average, so 1948 was 6.8 times, roughly, higher than the average for the period `48 to `54.

So we are going to take that 6.9 factor and we are going to use that going backwards in time and we are going to hold it at that level back to 1942. And we make some arguments in our review saying that that is not claimant favorable, that if you develop some kind of a regression line for the period, you will find that the numbers are higher than that approach would yield. So that is what underlies this finding.

DR. MAURO: Any chance we could put this up?

MR. THURBER: I don't know.

CHAIRMAN ANDERSON: I can see it becoming --

DR. MAURO: Yes, you have probably seen this if you took a look at the report. You can see when you go before 19, let's say, '46 and all the way back to 1942, we deemed that the correction factors may be too low by a factor of 10 so for those years, the generic approach used by TBD-6001 to account for the early years being worse. But we think that correction factor is too low and I guess this is a question worth posing to NIOSH, do you think that -- there is a correction factor that can be derived but we think it's about 10 times higher, if you want to go back to 1942.

MR. ALLEN: I believe we had, when the document was created, was you had essentially a war effort in `42 to `45, about the last thing on anybody's mind was health and safety, it was a very low priority. That same process is pretty much continued until it

was, the weapons program was turned over to the Atomic Energy Commission once it was created.

And one of the first organizations to start worrying about health and safety -- not that there was no worry about it -- but one of the first ones to put some significant effort into it was the Health and Safety Lab and that's kind of what created them were these AWEs, I think the big seven that Naomi Harley talks about.

And you can see that very clear in the data, that when they started up, 1948, you see the data as considerably higher than even 1949 and just steady improvement right down the line.

The individual reports from 1948, it's very clear there was no controls to speak of at these facilities. They made a lot of recommendations at least some of which got implemented and then more and more as they continued to try to improve.

1	DR. NETON: Yes, I don't quite see
2	the basis for a straight-line extrapolation
3	back if the processes were similar. They only
4	generate so much dust with certain types of
5	equipment, and we agreed, this was manual,
6	shoveling, lifting.
7	DR. MAURO: Let's just circulate
8	this is my copy let's send this around the
9	table, it's self-explanatory.
10	DR. NETON: I understand what
11	you're saying, that you're going a straight-
12	line extrapolation back.
13	DR. MAURO: The question is what do
14	you do?
15	DR. NETON: The question is why
16	would you believe that a straight-back
17	extrapolation is appropriate, other than the
18	fact it's more conservative, because of
19	process similarities. I mean, you can look at
20	the equipment being used and say this process
21	is very similar to what was being done in

1950, 1948, going back, and there's no local

capture ventilation	all	those	sort	of
parameters, you know,	I'm h	aving t	trouble	
other than the fact it	's more	e conse	rvative	, I
just don't know that the	hat is	necessa	rily go	ing
to make it right.				

DR. MAURO: I agree with that but right now we are in this funny place. You see, you picked 6.8 as your maximum multiplier that goes all the way back to 19 -- and if you were to extrapolate according to the graph, it would be 10 times higher. Now, which is the right number? I don't know. And you know, in the world we are in, unless you can make a case, as you are starting to make right now, why 6.8 is the right number all the way back to forty --

CHAIRMAN ANDERSON: It couldn't have been any worse than it was in `48 is what you are saying.

(Simultaneous speaking.)

DR. MAURO: And we are not convinced of that.

1	MR. ALLEN: In most cases the
2	improvements, the slope of that graph is
3	caused by Health and Safety Lab starting to
4	sample in that year.
5	DR. MAURO: So you would argue that
6	starting in wherever the break point is on
7	that graph, that before that, that at that
8	point in time, that's how bad it always was.
9	And then it got better after that.
10	DR. NETON: Especially since you
11	have no controls for six years, that's about
12	as bad as it's going to be. It's not like they
13	
14	DR. MAURO: Well still, I remember
15	talking to Bill about this
16	DR. NETON: Well, we will have to
17	look at
18	CHAIRMAN ANDERSON: Well, I mean,
19	the other issue would be you've got data from
20	`48 to `56 and you know, will you go back
21	further than that?
22	You could simply say we aren't

going to assign, you can't assign, there's no data from there. All that we know is it was bad. But we don't know, could it have been worse? In some of the place, it could have been better. You never know.

DR. NETON: When you look at the processes, I mean, with that logic I don't know why you wouldn't do an exponential curve.

I mean, what difference does it make? But I think we owe it to justify those, that --

MR. THURBER: We are not saying that this is the way to go. As we indicated, we are merely saying this is an alternative and I would point out that one has essentially the same problem in Electro Met when we get into that, because there was very little data early in the process and so NIOSH took the approach that since the process was the same, that that was not an unreasonable thing to do.

We argue in our NIOSH, our review of Electro Met, I'm sorry, that there was some evidence to support that assumption and there

is also some evidence that refutes that assumption. So it's clearly a question that needs some more attention.

We are the first to admit this is not the right answer but we don't think a horizontal extrapolation without good justification is the way to go.

CHAIRMAN ANDERSON: Do we have from any of these sites workers' testimony? I mean is there anyone saying, boy, you know, that was bad in `48 but I was there in `44 and it was, you know, you couldn't be in there.

MR. ALLEN: You are kind of getting into the issue of this generic document and you know the applicability to an individual site. Like we said before, Electro Met, United Nuclear kind of had their own data. We are not even really using values out of those, so you now, you don't get a lot of specific comments from workers on a generic issue, you know, it is something to be addressed on an individual site specific TBDs.

1	DR. MAURO: I guess a good question
2	is, you know, when, I guess, when the
3	measurements were made, they go back to 19
4	I guess they started in `48, `46? `48.
5	MR. THURBER: Christifano and
6	Harris.
7	DR. MAURO: Christifano and Harris.
8	Now whether or not their data, air sampling
9	data that go earlier than that, I bet you
10	there are. I mean we are talking about, I
11	mean, my guess is they took
12	DR. NETON: I am not sure.
13	MR. ALLEN: Mallinckrodt you can
14	find some, other than that, I mean, that's
15	MR. THURBER: There's a little
16	Electro Met data in the early period and as I
17	recall, there was a change where they put in a
18	vacuum cleaner or something and so on a month-
19	to-month basis they got a big improvement so
20	that's some of the kind of information that
21	may prove relevant to developing a position on

this question.

DR. MAURO: And this is an ongoing discussion. You know, when you were talking about, okay, what's a reasonable default dust load. Now, we are not asking the question, what's reasonable that represents typical discussions occur across the complex of AWE facilities at that time.

We are really asking the question, what's the upper end that it is likely to represent so that if we are going to pick a number and we have a guy, and we know a guy who worked there in 1942 and we are going to assign some dust load onto him, you don't want to assign to him your best estimate. You are going to say listen, I want to assign to him a number that I am feeling pretty sure it was not higher than that. You know, we want to place, we want to give him the benefit of the doubt.

And in our opinion, one of the recurring themes that we are going to see over and over again is, there's a difference

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between saying what is a reasonable number that represents typical conditions at the facility in a given year and a typical operation, and what we believe to be the right number to pick, to assign to a person that you are sure probably didn't have much of an exposure that was higher than that.

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So it's, you know, extrapolation is sort of a way to say, you know, even the numbers themselves, the actual numbers that are plotted, are we plotting the 95th percentiles of here, these concentrations? You know what do you -- I am always concerned that -- this project is not concerned with coming up with realistic estimates of typical workers. It's concerned with coworker models that provide a level of assurance that when you assign a number to a real person, that you guy, а are not underestimating him.

So you always want to pick a number that you can argue, it's very unlikely

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this guy had a higher exposure than that. And this curve, I think, that's why this curve troubles us.

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DR. BEHLING: John, this is Hans Behling. Can I make a comment here?

DR. MAURO: Sure.

DR. BEHLING: One of the things that I looked at when I looked at MetLab, which is really time zero for this whole program, and that is, and you'll referenced throughout some of the TBDs or Site Profiles in the early years, and that is the tolerance level and when you look it tolerance level gives you some understanding how high the air concentrations could have been without concern for the protection of workers.

And in my reports for the Met Lab, and it was part of the 250 day issue, I cited tolerance levels that were astronomical by today's standards and clearly define a very, very different set of acceptable limits for

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air concentrations, body burdens et cetera.

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DR. MAURO: So, I think, to close this down, when you addressed this issue of extrapolation back in `42, you can make your case whether it's flat or it goes up, I think it's important also to, the overarching thing is that when we go back in time, you want to do it in a way that we can say with a degree of confidence, it's unlikely that anyone would have experienced an annual exposure higher than this dust loading -- not the best that would be claimant estimate, because neutral. If you picked the best estimate, you are being claimant neutral. You want to be, you want to make sure that all workers that worked at that time are unlikely to experience concentrations higher than that over the course of a year.

We realize in any given day the numbers can be very high but over the course of a year, you know, the exposure a person might have experienced because of his job,

1	unlikely to be higher than this.
2	DR. NETON: Well, I think we've had
3	the action
4	CHAIRMAN ANDERSON: Yes.
5	DR. NETON: in this document,
6	why we believe our values are appropriate.
7	MR. THURBER: So are we good on
8	finding five?
9	CHAIRMAN ANDERSON: Yes.
LO	MR. THURBER: Six. This relates to
11	the fact that radon was not included. I think
L2	that given what David said earlier, that we
L3	can probably pass this on by, if that's
L4	agreeable to everyone?
L5	MR. ALLEN: That was going to be my
L6	response so
L7	DR. MAURO: If there's no water,
L8	there's no radon.
L9	DR. NETON: So that's close.
20	MR. THURBER: Now we have a number
21	of observations. These should go more quickly,
22	hecause the more generic issues were findings

	the	first	observat	ion d	leals	with	the	fact
tha	t it	would	be helpf	ul if	some	basi	s for	the
sel	ecte	d time	e period	was	prov	ided.	It	just
app	eare	d as `4	12 to `58	. Why	· .			

And we noted that in another place in the document it said `44 so this is a -- I don't think it requires much more discussion than that, but if anybody has any clues?

CHAIRMAN ANDERSON: Fifty-eight end is --

MR. ALLEN: That's I believe the end of the data was `58. And the beginning was essentially meant prior to `58 and you can see where we went to some effort to try to back-extrapolate prior to the data showing and whether you agree with that or not, the basis was prior to `58.

MR. THURBER: Observation two.

Actually, we already touched on this. The TBD-6001 said that data on the solvent extraction step was under development and that's probably worth an indication of what is going to happen

1	with regard to that piece of the operation.
2	MR. ALLEN: Again, frankly, we
3	haven't found any situations where we needed
4	that information. I mean, this solvent
5	extraction pretty much happened at the bigger
6	sites where we have data from that site
7	instead of needing the generic numbers. So
8	MR. THURBER: Mallinckrodt
9	attributable. Yes. Okay.
10	MR. ALLEN: Yes, Fernald and
11	Mallinckrodt. I don't know if we actually
12	intend to fill that in until it's needed.
13	MR. THURBER: Again, if you revise
14	the document, you might say that.
15	MR. ALLEN: Yes.
16	CHAIRMAN ANDERSON: Well, yes, I
17	mean if you're saying it's under development
18	but it isn't, you could simply change that to
19	say, do you recognize this is an issue but up
20	to this point, it doesn't seem to be a generic
21	assessment that's needed.

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MR. THURBER: Perfect.

1	CHAIRMAN ANDERSON: That way, it
2	would lay it to rest.
3	MR. ALLEN: If we were to revise
4	the document.
5	CHAIRMAN ANDERSON: Again, if it
6	says it's under development and you say well,
7	yes, we are working hard on it, we don't need
8	it, but we don't need it, we don't need it.
9	Okay.
10	MR. THURBER: Observation three
11	relates to ore handling and radon exposures
12	and that's been addressed.
13	DR. MAURO: That's like a radon
14	problem. Radium-226 and thorium-230 go away if
15	you don't have water.
16	MR. THURBER: Okay.
17	CHAIRMAN ANDERSON: So
18	MR. KATZ: It's a non-issue.
19	CHAIRMAN ANDERSON: We need to
20	state up front though, that these facilities
21	don't contain these?
22	DR. MAURO: No, this TBD

1	MR. THURBER: This document doesn't
2	cover
3	DR. MAURO: Doesn't cover that.
4	MR. THURBER: Doesn't cover floor
5	handling.
6	MR. KATZ: That's handled in the
7	finding one, already handled by the finding
8	three decision.
9	CHAIRMAN ANDERSON: Yes.
10	MR. ALLEN: There's some action
11	necessary but it's covered in the finding.
12	DR. NETON: I frankly wonder if
13	there is a little more than that, though.
14	MR. KATZ: This is handled by that.
15	DR. NETON: It is, but maybe these
16	could be like, marked held in abeyance until
17	the document is revised to state that.
18	MR. ALLEN: Well, they're covered
19	under finding one and two and
20	DR. MAURO: That's what I was
21	saying earlier.
22	MR. ALLEN: Where, you know, we
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1	just need to go on without that.
2	DR. NETON: Well, what I'm saying
3	is that these can be specifically closed right
4	now, if we just say we are going to revise the
5	documents and say it's not applicable. But you
6	can't close finding one just by saying that.
7	That's my point.
8	DR. MAURO: Oh, okay, I see what
9	you are saying.
10	MR. THURBER: No, I understand.
11	MR. ALLEN: What is the process on
12	observations? Are we closing these? Are we
13	treating them like findings or are they just
14	noted?
15	MR. THURBER: I think some of these
16	require some action as we'll see.
17	DR. MAURO: In the past, some of
18	our authors feel that it's important to
19	separate things that we really think that are
20	important from things that are less important.

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author's judgment

1	be designated as an observation, I would not
2	automatically say that this need does not to
3	be addressed.
4	In fact, when I was looking at
5	these observations, some of them I would have
6	called findings. So I think that though they
7	have been categorized, some clearly are
8	observations that are really not that
9	important and in fact we have a whole section
10	where we found typos and inconsistencies, all
11	of which have to be cleaned up, so, house
12	cleaning.
13	But I think some of these, I
14	wouldn't discount some of these yet so I would
15	like to go through the observations.
16	CHAIRMAN ANDERSON: Yes, I want to
17	go through them
18	DR. MAURO: And they need to be
19	addressed.
20	CHAIRMAN ANDERSON: I'm also

thinking in terms of the rest of the Board,

when we come back to them they are going to

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1	have this matrix and somebody is going to say,
2	well, what about this?
3	So somehow we need to have at
4	least a mention in here that, you know,
5	observation three, that because this, 6001
6	address doesn't address ores that contain or
7	processes that
8	DR. MAURO: Let me ask you, when we
9	reissue this matrix, there's a column called
10	"NIOSH initial" the next column over you
11	see is the response. Basically we deferred the
12	response.
13	CHAIRMAN ANDERSON: Yes.
14	DR. MAURO: Are you folks going to
15	fill that in?
16	MEMBER FIELD: Yes, we'll fill that
17	in.
18	MR. ALLEN: Yes, we can, I was just
19	having a hard time with that very first
20	finding on the, telling you this was, ore was
21	not associated with it, to address these is
22	more of a big White Paper rather than a

paragraph	or	а	sentence	and	

DR. MAURO: Well, I can see on the radon and the thorium-230 and the radium-226, simply say this issue is being addressed under finding one.

MR. ALLEN: Yes, I think most of the observations are going to be a one sentence --

MR. KATZ: For finding one, in the matrix, you can just write, "White Paper addresses."

MR. THURBER: Yes. But recall now, as Jim said, as we expanded our discussion on finding one, we pointed out there were several places that went beyond the ore question and that's why, again, as Jim said, we shouldn't pile a lot of these things into finding one. I think a simple statement that TBD-6001 was not intended address ore processing or something to that effect, does that job.

DR. NETON: Because that way you don't have to rely on closing finding one to

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1	close all these other things.
2	DR. MAURO: You could close this.
3	I understand what you are saying. I am fine
4	with that.
5	MR. THURBER: Absolutely.
6	Absolutely.
7	DR. MAURO: I'm fine with that.
8	CHAIRMAN ANDERSON: Okay, why don't
9	you
LO	DR. MAURO: Because of one item,
L1	right.
L2	MR. THURBER: Observation four
L3	points out that there is no discussion of
L4	handling recycled uranium or enriched uranium,
L5	the ores we have already talked about. It
L6	seems at a minimum there should be a statement
L7	of intent that this is not intended to address
L8	sites that handle recycled uranium or enriched
L9	uranium or some provision as to how those
20	should be handled.
21	MR. ALLEN: Yes, it actually, there
22	actually is a table in there with the, on page

five	of	the	TBD	that	has	the	re	ecycli	lng,
recyc	led	ura	nium	comp	ponen	ts,	pl	utoni	Lum,
neptu	nium	, tec	hneti	um et	cet	era.	It	says	to
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MR. THURBER: I missed it I'm sorry. Yes, I mean it's, it should be, you know, tell us that.

MR. ALLEN: I know. That part is in there. The enriched uranium it doesn't mention, and just like it needs to be revised for -- but you know, make it clear that it doesn't cover ore, I think it also needs to mention that about the enriched uranium, I think you're right on that one.

MR. THURBER: Good. Observation five. The document says that you should use a default air concentration of seven dpm per cubic meter for non-operational areas.

That requires some further discussion I think, because if you look at some of the available data for clerical people and others in the Mallinckrodt documents,

you'll find that nurses and whatever are exposed to higher levels than seven dpm per cubic meter.

So it seems to me that either you need to think a little bit more about whether that is an appropriate default value, or somehow explain what non-operational areas of the plant are, or both.

MR. ALLEN: Yes. I am --

CHAIRMAN ANDERSON: Let the record show he's nodding and shaking his head at the same time.

MR. ALLEN: I understand the comment. It's difficult in a generic basis to address this kind of thing rather than site-specific. I mean I'll come up with some sort of response for this matrix at least but I'm not clear where I go at this point.

MR. THURBER: Well, if there's, if there's data that says that seven dpm per cubic meter is not a good number, then you can think about revisiting that number.

1	CHAIRMAN ANDERSON: What's the
2	basis of the number?
3	MR. ALLEN: I don't know. I was
4	going to say I don't recall off the top off my
5	head.
6	MR. THURBER: It somehow is tied in
7	with one percent of 100 MAC or something like
8	that.
9	CHAIRMAN ANDERSON: But it isn't
10	based on measurements or anything?
11	MR. THURBER: No. It was based on -
12	_
13	DR. NETON: I think a lot of this
14	depends on what we define as a non-operational
15	area. Maybe that needs to be clarified.
16	MR. THURBER: Well, that's why I
17	say, that's a possibility and, but I did
18	notice somewhere that
19	CHAIRMAN ANDERSON: So it's a
20	calculate
21	MR. ALLEN: I mean, it gets
22	difficult

CHAIRMAN ANDERSON: You could call it a bottled number.

MR. ALLEN: I mean, when you're having a hard time with them sampling operational areas and you know they are not sampling all of them. You can see there's not a lot of data for office buildings and that sort of thing.

DR. MAURO: These rules of thumb, we run into this seven dpm per cubic meter and on a number of occasions we have seen it as being, we are going to assign this -- and their rationale originally, as I recall, was that, there was an observation that when you were measuring numbers in an operational area, then you went off some place where there wasn't an operational area, in general the concentrations in the non-operational area are about one percent of the concentrations in the operational area.

Now, the implications being that operational areas in these kinds of facilities

typically have 700 dpm per cubic meter. So, therefore seven dpm per cubic meter is probably a good number for a non-operational area.

So I think there are two aspects to this issue. One is, is 700 a good number to represent a bounding concentration for an operational area at these classes of facilities, and yes, we agree that the one percent, because we see the data, that yes, on many occasions we see non-operational areas are much, much lower than operational areas and the issue becomes, all right, from an implementation point of view, you have got to be careful when you do that.

And Bill, you gave lots of, several examples on page 20 of this report where it shows that there are places where 175 dpm, per cubic meter, this was a dispensary at a facility. Another location where some numbers as high as 50 were observed.

Now, whether or not the average

1	over the course of the year, which is what you
2	can do when you do these dose reconstructions,
3	or whether, you know, what do you, the 1.5 to
4	175 for example for the dispensary, were those
5	annual average or those individual
6	measurements? I'm not sure.
7	DR. NETON: Also, were those at a
8	facility that only processed uranium and not
9	uranium ore? That can make a big difference.
10	DR. MAURO: Yes, okay.
11	DR. NETON: We measured some short-
12	lived daughters in there
13	DR. MAURO: And I will be the first to
14	admit I'm not
15	DR. NETON: I would be surprised if
16	it would be several MAC
17	DR. MAURO: I understand but when
18	you see these numbers track
19	DR. NETON: If you get airborne
20	though, I mean, this sounds to be
21	DR. MAURO: I would, again, this
22	is, I would agree, the fact that you went into

1	a dispensary and at one point in time picked
2	up 175 dpm per cubic meter in a given day,
3	that doesn't mean that's what you're going to
4	experience over the course of a year.
5	And I think that, you know, so if
6	you're go with the seven, you have got to make
7	your case a little stronger.
8	DR. NETON: I guess we are not
9	arguing, we just prepare some kind of
10	response.
11	DR. MAURO: And this crosses a lot
12	of sites, where I've seen this number at least
13	10 or 12 times.
14	DR. NETON: You have got to come up
15	with some kind of lower bound, I mean,
16	sometimes we have used like 10 percent of the
17	air concentration and one percent of the
18	actual value in the plant. But
19	MR. ALLEN: Well, many of these
20	smaller sites, we don't have the information
21	to place somebody in an office versus an

operational area and you end up essentially

1	putting everybody in the operational area.
2	DR. MAURO: You give them the big
3	one anyway.
4	MR. ALLEN: It's moot in most of
5	these facilities.
6	DR. NETON: We don't know whether
7	the secretary had walked through the plant.
8	CHAIRMAN ANDERSON: Yes, I mean,
9	the practical reality
10	DR. MAURO: We don't use it.
11	CHAIRMAN ANDERSON: It would be
12	nice to know, when we argue over something
13	like previously has never been used, if this
14	is, if we are just kind of doing this for
15	completeness sake, well
16	MR. ALLEN: Honestly, I think
17	that's the situation. I can't say as we had
18	actually ever used that number but it is
19	there.
20	CHAIRMAN ANDERSON: Okay.
21	MR. THURBER: Observation six. The
22	comment is that the document doesn't tell you

1	how the doses are apportioned from between the
2	operators, laborers and supervisors and so
3	forth. Now, I presume that if you read on to
4	the next section of the document, it tells you
5	how it's done there, which was the section on
6	internal dose. So this is kind of perhaps kind
7	of a cheeky comment, but it's just one of
8	those things that makes the functionality of
9	TBD-6001 a little troublesome.
10	You know, you don't tell me,
11	anybody, a viewer, and this is a problem, some
12	of these are problems more for the reviewer
13	than the user. The information is not
14	provided.
15	MR. ALLEN: Well, not conveniently
16	anyway.
17	MR. THURBER: I mean, it's only
18	provided by inference, you know, you tell me
19	what you are going to do with the internal
20	dose.
21	DR. MAURO: When I was looking at

this, it seemed to me that how, the sorting

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of, we're going to, supervisors we are going to assign this distribution, and laborers we are going to assign this distribution. It wasn't apparent how you got there from the Christifano and Harris data, the data behind that. How did you do it?

I mean I would like to be able to go in and say, here's the 20,000 measurements. I could reasonably sort them by people, say okay, you know, 1,000 of them were people that we could say were supervisors and we know that, pull them out, then I could say and look that number, the numbers for the at supervisors that in that 20,000 are measurements and oh that say, yes, distribution, that is a sign you can use for supervisor, it looks pretty good. We can't do that. Right now, we can't do that and I think that's needed. That's the front end of the problem that clearly goes towards this report. And then of course the more difficult problem is when you eventually use -- let's say it

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turns out that is good, let's say the numbers you have picked for supervisors, the distribution that's in there, the dust loading, is -- you go back and you check it and you find out yes, those are good numbers.

Then you have the headache of saying all right, now I have a real site, I know this guy worked at the site and he was a supervisor, sometimes you have them, yes, he was a supervisor. But we always run into this situation. The name of a person's job, for any individual, troublesome given is always because we find out when we interview workers that yes, I was a supervisor but I used to spend my time all over the place.

And it almost becomes, when we do

-- I do a lot of dose reconstruction audits

and I see that they took a guy, looked at what

his job was and he described himself, and they

might drop him into a category when it's not

apparent that that's the most claimant

favorable assumption.

So you know, that's the back-end of the problem, that is, even if you have a good distribution for supervisors, when you get a real good case, do you use that or do you say no, I'm just going to give this guy the benefit of the doubt and assume that he also, he might have been exposed to something higher?

So I, this matrix, when it's time to implement it, is a problem, especially if you get a guy who ends up getting a 45 PoC and the reason he got a 45 PoC is because you assigned him as a supervisor and there's some question of well, how do we know he always was in that mode and maybe he spent a lot of time someplace else. And this is not unusual. You talk to these workers and you find out these labels, these designations mean very little.

MR. THURBER: On that subject, I would mention that TBD-6000 specifically says if you have doubt as to what the worker's job description was, use the maximum values and

that is very good guidance and it would be nice if that same kind of guidance was in TBD-6001 to kind of clarify, demystify that question.

MEMBER FIELD: That sounds like a different issue though, I mean, it's like, it sounds like you are sure but you question how reliable that category is, versus you are not very sure, I understand what you are saying but it seems like two different issues.

CHAIRMAN ANDERSON: Well, I mean, if this, as a generic guidance for surrogate data, the question is, okay, now you are at a facility where the guy -- it's a plant that would be a 6001 and he says he's a laborer but there's no data from the site. And now you are going to go to 6001 and say he's a laborer, go to the table and this is how we are going to assign his exposure.

So really the question is, how confident are you that a laborer in this facility is the same as a laborer, I mean, you

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the example of а supervisor, probably more problematic because somebody doesn't show, it's а small work force, supervisor steps in because he knows all the jobs, those kind of things, where once you are a mechanic or you are a whatever, you are shifting back and forth maybe not as much.

So, I mean, how confident are we that the variability between what a laborer is in one place versus another in a different part the country, in different unions, all of that. Is that something we can use or do we need a default, let's say, that if you don't have site-specific --

MR. ALLEN: I think people do get kind of hung up over the titles in this thing. But essentially what we try to do is is somebody that's routinely operator operating with the material, or the equipment that is containing the material; supervisor or laborer is usually somebody that is routinely the but not necessarily hands-on in area

know that kind of a --

working v	with t	he	mater	rial;	th	ne oth	er		
	CHZ	AIRI	MAN	ANDE	RS	SON:	You	7	will
probably	need	to	then	have	a	defin	nition		you

MR. ALLEN: Yes, I agree. I agree.

CHAIRMAN ANDERSON: Because I think, I mean, when they say it to me, I tend to place their job title into a union -- or something more, you know, what you're saying, it really isn't, you are looking at what the work was and now you are, instead of having everybody being unique, you are now saying this person fits this work profile, this exposure profile of a --

MR. ALLEN: Our dose reconstructors are well aware of that. The term I always use, the job title I always use is clerk. That can mean anywhere from a payroll clerk in an office building to a materials control and accountability clerk that is stamping numbers in uranium ingots.

So yes, we realize we have got to,

we do do a telephone interview on the individuals, as much information as we can get there, from what we know of the site, and job titles there, and we try to, you know, case-by-case, put it all together and put them in the right category. That's, I think, way off the topic.

DR. MAURO: Our primary concern is that we could not track the distributions for supervisors, laborers and clerks, the data distribution that is in the look-up table back to the original 20,000 measurements and how they were created, you know, like -- this is what we could say, yes we looked at this.

Because one of the things we try to do, especially when we are doing SEC-related issues, I mean, you know, you are saying, wait a minute, let's look at the data, and right now, you know, that hasn't been done and I think it's important.

This is the rock you are going to stand on for many sites, at least five anyway,

1	and we have to look, we have to see that data
2	and a case has to be made why that
3	distribution is the right distribution for a
4	person that you know was a supervisor. Whether
5	or not you are going to assign it to that
6	person, that's another question.
7	CHAIRMAN ANDERSON: Okay. So you
8	know
9	MR. ALLEN: I think I know what the
10	question is.
11	CHAIRMAN ANDERSON: Whoever is
12	going to be next filling out the matrix, they
13	need to, you know, have we made what we want
14	clear enough for you? Okay.
15	MR. THURBER: Are we okay?
16	CHAIRMAN ANDERSON: Yes.
17	MR. THURBER: All right.
18	Observation seven. In using the Christifano
19	and Harris data, what NIOSH does is they take
20	the raw data by the various categories, like
21	denitration, and they convert the raw data to
22	an assumed log-normal distribution and they

1	calculate the geometric mean and the geometric
2	standard deviation and the 95 th percentile, if
3	it's appropriate, or whatever.
4	So for each of the data tables
5	which come, the raw data tables which come
6	directly from Christifano and Harris, NIOSH
7	prepares a companion table where they show the
8	log-normal statistics that they have derived
9	from the raw data.
10	One of the problems is that in
11	several cases, the tables show the geometric
12	standard deviation as less than one, which it
13	cannot be.
14	CHAIRMAN ANDERSON: Well, we don't
15	have statistician here, so we can't argue
16	with you on that.
17	MR. ALLEN: That would be a tough
18	argument anyway.
19	DR. NETON: It's a calculation
20	error.
21	CHAIRMAN ANDERSON: Or a typo.
22	MR. ALLEN: I think it's a typo. I

1		am	still	digging	into	the	numbers
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MR. THURBER: The observation eight, as John mentioned, as we went through this, we just as an aside made a list of typos and fuzzy statements and things that might or might not be helpful to NIOSH when and if they do revise the document so I don't think that requires any further discussion.

CHAIRMAN ANDERSON: What is the process for fixing these kinds of things?

MR. ALLEN: Well, I mean, we can revise the document, ideally we would settle or come to a resolution on all the findings and issues and revise it one time rather than piecemeal put a whole bunch in there.

CHAIRMAN ANDERSON: I mean, when we do it, not on this kind of a thing, you fix it and now you put it up on your database as the revised and give it a new number and it's not as though you have to go through any kind of real laborious approval and 17 sign-offs.

DR. NETON: Well, we have our

1	review process but it's all internal.
2	CHAIRMAN ANDERSON: Okay.
3	DR. NETON: We don't submit these
4	for external review at this point. It's not
5	that difficult, it would be Revision 1 at this
6	time.
7	CHAIRMAN ANDERSON: Yes.
8	DR. NETON: And we keep the old
9	revisions so there's a paper trail. But we
10	would want to get some resolution on this
11	issue.
12	CHAIRMAN ANDERSON: I mean, at some
13	time, we'll have accumulated enough of these
14	that, you know, rather than wait for final
15	resolution, we got all of this stuff kind of
16	rolling around there.
17	DR. NETON: The good news here
18	though I think is not many dose
19	reconstructions, if any, have been done
20	against the findings that we are seeing.
21	CHAIRMAN ANDERSON: Well, that was

going to be my next question.

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1	DR. NETON: And that's the good
2	news, so when we revise it, we are probably
3	not going to have to go back and redo a lot of
4	dose reconstructions. That might not be true
5	100 percent, but I think it's true as far as I
6	can think.
7	CHAIRMAN ANDERSON: All right.
8	Moving right along.
9	MR. THURBER: All right.
10	Observation nine, we noted what appeared to us
11	to be a calculational error in calculating a
12	specific geometric standard deviation.
13	MR. ALLEN: I don't disagree. It
14	goes along with observation seven, there are
15	some typos or calculational errors.
16	MR. THURBER: Yes, and actually
17	there are a number of other ones and David and
18	I have discussed some of these and we will get
19	into a couple more of them but probably there

ought to be an audit of the data, the data

tables in general. As we have pointed out in

our review, we did not go in and look at every

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number in TBD-6001. We just did a few spot checks and some of these observations are really a result of things that we have been doing since then we have tried to use it or apply it to some other site-specific things.

DR. MAURO: I have to say that when both Bill and I looked at this, we felt that this document was rushed, that its errors, it is not -- it is as if they tried to rush it out real quick. Too many inconsistencies, too many typos, we have a whole list, two pages of it. And I think we could slow the train down, take a look at this thing and clean it up. This is not one of the better TBDs. There's a lot of problems.

MEMBER FIELD: Can I ask you a question? For the geometric standard deviation, did you have data to figure out if any of them are correct?

I am just curious if it is a systematic error in whoever calculated the GSDs or it's just anecdotal or just every now

1	and then you see one. I just wonder if you
2	found any correct ones.
3	MR. ALLEN: They I have a
4	spreadsheet on what they used for most of the
5	actually, several spreadsheets and I am
6	missing pieces of the data here and pieces of
7	the data there and I am trying to get a hold
8	of the original author to
9	MEMBER FIELD: That would be
10	interesting to see.
11	MR. ALLEN: Yes. Yes.
12	CHAIRMAN ANDERSON: But this is
13	three, four years old?
14	MR. ALLEN: Yes. If it was a
15	contractor who no longer works for us
16	CHAIRMAN ANDERSON: You were
17	dealing with this two-and-a-half years ago and
18	probably took two years to put together so
19	we are looking at history here at a time when
20	the program was rushed and hurrying.
21	MR. ALLEN: The effective date is
22	December `06.

1	MR. THURBER: Procedurally, there
2	is a companion document which we mentioned
3	here called Strom 2006 or something, which is
4	a statistical compendium as how to apply log-
5	normal statistics or whatever to various
6	situations and presumably the authors of this
7	document used that compendium, which gives you
8	some straightforward cook-book procedures, if
9	you will, on how to calculate these statistics
10	from very limited data.
11	DR. NETON: This document is under
12	contract with Battelle.
13	MR. THURBER: Okay? Again, probably
14	a minor point, but some of the confusing
15	instructions that TBD-6001 provides, it says
16	in one sentence, it says, "Use the default
17	values above" and then in the next sentence it
18	says, "Don't use the default values above, use
19	ICRP 66."

And in looking further, we don't think that ICRP 66 is the correct reference. We think it should be ICRP 68.

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1	MR. ALLEN: Yes, I actually
2	remember that one. It was the first three or
3	four parameters are definitely 66 and then it
4	mentions the solubility at F1 value which then
5	comes from 68 and that is why somebody decided
6	you cannot say use defaults from 66 and
7	changed it to use the table above and
8	apparently didn't get a sentence deleted.
9	MR. THURBER: Anything further on
10	that?
11	CHAIRMAN ANDERSON: No. Pretty
12	straightforward.
13	MR. THURBER: Observation eleven, I
14	think we can pass on because that again
15	relates to the settling assumption, which, as
16	we have discussed earlier, has been resolved.
17	Observation twelve is virtually
18	the same as the previous observation, dealing
19	with the basis for the seven dpm per cubic
20	meter air exposure in non-operation areas. I
21	take it you covered that.

And observation thirteen.

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	CHAIRM	AN AN	DERSON:	So	back	to
seven, that	t's kin	d of a	default	valu	e, but	. on
a site-spe	ecific	basis?	Would	you	use	the
percentage	issue d	or not?				
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I mean let's say you had a facility where this is based on the 700 value in an operational, now if you had one where the operational value was 1,000, would you now use the one percent increase from seven to 10 or would you only use the seven or is it at all tied to any data at the site or is the 700 the max at all of these facilities?

MR. ALLEN: It's kind of the catch22 question, you know, if you have the data at
the site you generally don't use the default
values in here that we are talking about so
it's --

CHAIRMAN ANDERSON: Okay. But if you had, I mean you wouldn't have a non-operational result typically in any facility, so would you then use a percentage basis thing?

1	MR. ALLEN: Typically, we would use
2	some percentage of the operational value then.
3	DR. NETON: I just looked up the
4	pedigree of those values where they were high
5	and that was a Mallinckrodt value data which I
6	think are most suspect because of the radon.
7	There are hundreds of MAC radon
8	air, hundreds of picocuries per liter of radon
9	there and even if you are left with some decay
10	you are going to get some high concentrations.
11	CHAIRMAN ANDERSON: Okay, sorry.
12	MR. THURBER: No problem. Look the
13	final point, I'm sorry there's a typo, it
14	should refer to section 8.5.2, and section
15	8.5.2 deals with resuspension of dust when
16	operations are not going on, either they are
17	suspended between campaigns or after the
18	processing has been finished.
19	And the calculational approach in
20	that section is different from the
21	calculational approach in the prior section
22	3.4.2 in the same packet. And in part, it

deals with the number of days that the deposition occurs, it appears in the section 8.5.2 that the deposition occurred over some strange number of days and to your earlier section, it was over 365 days. So there's something that seems amiss there.

MR. ALLEN: I didn't think that was the case. I thought it was the starting air concentrations that was the difference. I am trying to look it up right now real quick while I am reading. I thought they both used 365 days. Yes, I would also, rather than bore everybody here I need to respond to this particular point.

DR. MAURO: It is what it is.

MR. THURBER: And I would, I note that the same information is included in section 8.3 as in 8.5.2. This is the duplication one.

MR. ALLEN: Yes, I agree, I mean -
MR. THURBER: Okay I think that

covers our comments on TBD-6001 except for

1	this other story on the transparency problem.
2	DR. MAURO: The Hooker example?
3	MR. THURBER: Yes, the Hooker
4	example. Do we want to go on to that or
5	CHAIRMAN ANDERSON: Do you want to
6	go back now to that one? Yes, I would like to
7	kind of finish up on the 6001 and then decide
8	on what our next steps are, all right? One is
9	going to be in a time line of, you are going
10	to put in the responses, there's going to be a
11	White Paper. Those seem to be the main
12	MR. ALLEN: Yes.
13	CHAIRMAN ANDERSON: Most of the
14	responses are the next step will be to
15	close those out.
16	MR. ALLEN: Yes and then we car
17	probably start closing things out, narrow it
18	down to
19	CHAIRMAN ANDERSON: It seems that
20	finding one is where we may have to have some
21	more discussion. But we'll see on that.
22	MR. ALLEN: Do you have any

1	objection to a comfort break at this point?
2	CHAIRMAN ANDERSON: Oh, well, how
3	much time do we think this next, to go back,
4	and
5	MR. THURBER: Fifteen minutes.
6	CHAIRMAN ANDERSON: Okay, let's
7	take a quick
8	MR. KATZ: Five-minute comfort
9	break?
10	CHAIRMAN ANDERSON: Break, I would
11	say, then break for lunch before we
12	MR. KATZ: Twenty of? Twenty of, we
13	will be back?
14	CHAIRMAN ANDERSON: Yes.
15	(Whereupon, the above-entitled
16	matter went off the record at 11:34 a.m. and
17	resumed at 11:42 a.m.)
18	CHAIRMAN ANDERSON: I think we lost
19	Mark.
20	MR. THURBER: What I would like to
21	do is give you of a real world example, if you
22	will, of how TBD-6001 is used for a specific

site and in this case it's Hooker Chemical. We were recently tasked by the Board to do a review of Hooker Chemical. The review is in progress so I am not going to comment particularly on the review but I wanted to give you an example of how it interplays with the TBD-6001.

What went on at Hooker Chemical was they received some uranium-bearing slags, they treated the slag with hydrochloric acid to increase the grade of the uranium from about a pound per, I'm sorry, 0.2 percent up to two percent, on that order.

So, and the reason they did that at Hooker was that Hooker was engaged in some other work for the AEC and they had some excess hydrochloric acid and so they thought it was a good way to use that excess acid.

The specific example I am going to discuss is the external exposure during a residual period and, in particular, the exposure that a worker gets from standing on

the contaminated surface that we talked about earlier this morning.

And if you look at table AA-3 in the Hooker Appendix, and I will show you an excerpt from that table in just a second, you see that the external exposure is 0.376 millirem per day per an operator and this is based on the scrap recovery operation from TBD-6001.

So, quickly, this is the excerpt from, this is what you would see if you looked at table AA-3 of Appendix A or a part of what you would see and we are talking about, as I say, an operator called in this case Plant Floor High. The period is the residual period from 1946 to `76, the whole body dose is 0.376 mR per day and we are referred to table 7.3 of TBD-6001 and this is one of those big tables that I mentioned earlier and said we would come back to.

So let's go to table 7.3 of TBD-6001. We have the scrap recovery operation.

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Again this is an excerpt from a very large table. In this case, we are talking about the operator, the external dose pathways that are examined and you will see that this operator is assigned that dose of 0.376 mR per day and that's based -- and that is his median exposure.

geometric The assumed standard is five. deviation This is standard а assumption that is provided in this other document I mentioned earlier, prepared by 2006. Battelle, Ιt is default Strom а assumption when you don't have enough data to actually calculate the geometric standard deviation from the log-normal distribution.

One of the things that is very confusing is this table is, assumes a calendar day and a lot of the data is reduced to a calendar-day basis for the convenience of the dose reconstructor. But the problem is that TBD-6001 seldom tells you whether you are talking about the work day or a calendar day

so that adds to the difficulty in trying to understand what the document is purporting to tell you.

Here's what TBD-6001 says in terms of, well, the question is where does this number of 0.376 mR per calendar day come from? And the guidance here in TBD-6001 says we use the deposition velocity that we talked about this morning, 7.5 times 10 to the minus four meters per second. We allow it to deposit for 365 days and then we multiply that number by the air concentration and then we get the floor concentration in terms of dpm per square meters.

So we need two things, then. We need to know what the air concentration is and we need to know what the dose conversion factor is, and really it's the exposure conversion factor because we are talking about exposure in terms of mR per day and not dose in terms of millirem per day.

So, and TBD-6001 says well, to do

this, use the dose factors in table 6.1 of TBD-6001. So let's look at what that -- oh, I am sorry, we will come to that in a second.

First of all, let's talk about the air concentration number. TBD-6001 doesn't tell you where the air concentration came from. I talked to David about this. He says you go to table 8.23, which is the scrap recovery table and you take the data for scrap recovery operators and you average those numbers and that's where you get the air concentration.

Again, here's an excerpt from the table in TBD-6001 and you can see that the general area samples for the furnace area are 900 dpm per cubic meter and 200 dpm per cubic meter. Those numbers were averaged with an assumed geometric standard deviation of five.

The geometric mean of 151 dpm per cubic meter was calculated and using that and the deposition velocity, we get a value for the floor concentration. Again, none of this

is revealed in the document. You have to find a helpful person like David to get to this point.

Now, recall that the guidance we looked at says, "Go to table 6.1 to find the dose conversion factors." You go to table 6.1, you will find the dose conversion factors are in terms of millirem per day, not mR per day.

The dose conversion factors in table 6.1 are based on seven dpm per cubic meter, not the 151 dpm per cubic meter that we calculated and the data in table 6.1 don't tell you whether you are dealing with calendar days or work days. So this is kind of useless.

So then we go further back into the document and we go to table 3.10 to seek out these exposure conversion factors. Table 3.10 gives us a conversion factor and of 5.6 times 10 to the minus 10 mR per day per dpm per square centimeter, the surface concentration.

We had independently, in our

review of TBD-6000 commented on this conversion factor and provided some calculations that showed it was too low by, I forget, do you remember John? A factor of two or more?

Because it did not include Bremsstrahlung nor did it include the direct contribution from electrons, which would cause exposure in the skin and other organs close to the surface. So, that basic number has been questioned elsewhere, so let's, but let's just leave it at that for the moment.

Now, if you do the calculation, then you find out that the exposure you calculate is 0.002 mR per hour. And again, in talking with David, David said well the spreadsheet shows 0.2 mR per hour, a factor of 100 different and apparently there was an error in the spreadsheet and, which is propagated through the calculation.

So then what one needs to do is to take this value, and we will stick with the

0.2 mR per hour number, the number that is too high by a factor of 100, and see how that gets converted to this number of 0.376 mR per day.

So first thing we have to do is convert it to exposure per calendar day and for some reason, the calculation assumed that there were 350 days in a year. You know, it's not a big deal, is whether it's 365 or 350, but it just leaves the whole thing kind of suspect.

The second thing is that they then took this number, 1.37 mR per calendar day and said this is the geometric mean of a log-normal distribution, which you will recall, that we had already assumed that the original data was for a log-normal distribution and so it's not clear why you take a log-normal of a log-normal, which obviously reduces the values you are dealing with.

So, to summarize, TBD-6001 doesn't contain enough information to trace the values to the summary tables. There are errors in

calculating the external exposure. It incorrectly converts exposures to a calendar day basis and for reasons not obvious, it then takes a value which is based on a geometric mean and further reduces it to another geometric mean.

So that kind of summarizes some of the practical problems that you get into when you try to apply the TBD-6001 data to a site-specific situation.

CHAIRMAN ANDERSON: Comments? It seems to follow. I didn't quickly do all this math to see if your math was off as well.

MR. THURBER: Yes, as I say, you now, we could never have sorted this out at all without David's help.

DR. MAURO: I mean that's the first problem. The problem is we sit down and read it and we can't figure it out so we call David, and say can you help us out. So now we understand -- problem one, is it shouldn't be like that. The document should be complete

enough that an independent reviewer should be able to go through it and not say -- in fact Bill called and he says, "John, I can't figure out what they did."

And I said, "Well, you now, call David." He says, well, you know, I don't know, should we doing that? I say, call David, you know, so we call, and because we, again --

CHAIRMAN ANDERSON: You block caller IDs.

DR. MAURO: The funny thing is that you know, we are allowed to get clarification. The ground rule is that as long as we are not resolving an issue and we just want to figure out, please explain to us what you did here, whereabouts did you do that, and we do that and, but it shouldn't come to that.

You know, we are often in this very uncomfortable position of saying, listen, are we stupid, maybe we should be able to figure this out. Why are we having so much trouble figuring this out?

And so, we keep digging to try to figure it out, you know, and then two or three days pass and we still can't figure it out and I said listen, you know, if we could put a couple of people on it for a couple of days and we can't figure out what they did, this is not our problem. It should not be that hard.

So we call and we get help. The reality is it shouldn't be that way. It should be something where another health physicist with some experience should be able to figure out what was done and David explained it to us. Now, problem number one is just being transparent, and maybe even put an attachment to the report that walks you through this is how we got the numbers.

The second problem of course is once you've understood what was done, you think it was done incorrectly. Well, we could never have known that if we didn't call David to find out what they did.

So we think that, you now, as

applied to Hooker, and that this is -- now, so, in a way it applies to TBD-6000 in terms of being transparent. It also applies to TBD-6000 in terms of we think there might be some problems here in the numbers and they need to be fixed, and of course it has some real-world implications for Hooker, which is right now before us as an SEC, what we are doing right now.

I guess this is the example. I think there's a lot of problems with TBD-6001 that have to be cleaned up so that another -- to get errors corrected and once they're corrected, it should be transparent, exactly what was done so that another person can read it and check the numbers and say yes, everything was okay.

MR. KATZ: Just for the record, you should never spend a couple of days trying to figure out what was -- get on the horn and spare yourself two days of useless labor.

DR. MAURO: I agree.

MR. KATZ: Even if it's useful at the end it's two days wasted.

DR. MAURO: Well, let me take it a step further and while we get a chance, let's -- I call it a couch trip. When we are doing a dose reconstruction audit, you folks haven't seen these, but they are very, very sophisticated.

They are at a point in the process where some very sophisticated spreadsheets and workbooks are being developed and it takes quite a bit of time to get to the point where we understand the mechanics and the rationale behind these workbooks.

These are not self-evident and I believe you folks probably have a pretty extensive training program for your dose reconstructors to get to the point where they can function.

So the reality is, this is, it does take time for independent reviewers to figure out what is going on and very often you

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find out everything is fine.

But it is not always apparent what was done and this goes, this is a longstanding issue, is that when you guys are putting products whereby your together your reconstructors for example giving are instructions on a spreadsheet or a workbook, the product that is used, the degree to which you can make it understandable to someone independent, like us, so we can figure it out and right now we are spending too much time trying to figure things out.

This is one example where you're right, we shouldn't have spent two days. But it's not unusual for us to find ourselves in a position of trying to understand what was done here. We could certainly use a little help on these products that come out, a little bit more explanation so that, you know, we don't have to call.

CHAIRMAN ANDERSON: Just kind of a process question here. Is this kind of a, I

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you went through it and found this, is of problem. Is one your risk assessors, when they are doing this, or dose reconstructors, would they have identified this problem or are they more rotely going to the, quickly going to a table, pulling out a number, putting it into the formula calculating and not working this through?

MR. ALLEN: A dose reconstructor normally wouldn't have seen this. They are going to with the appendix that says to assign this external dose per day and that is what they are going to assign. I mean, if they are curious and start digging into the basis they might see an issue, but that's something that's supposed to be caught during the review of the documents.

CHAIRMAN ANDERSON: Yes. Because I'm just wondering how much of this is because 6001 has not been really applied that much so that a lot of these facets aren't picked up or is it simply that now that we have, somebody

really has gone through and reviewed it, we are finding these and the question then, really, as you raised, is you haven't done this exhaustively for all of the tables so what is your, what is the likelihood that this is --

MR. ALLEN: This particular error is essentially, is in the Hooker Appendix but it's carried forward from TBD-6001 so somebody reviewing the Hooker Appendix, you know, probably went as far as to say, okay, yes, that's the number that's in TBD-6001 and I mean that would have been -- the review of 6001 is where that should have been caught but that's the point where it was missed and it's up to the reviewer. They could carry it back further if they wanted to or they could stop with an already-approved document, you know, and saying that's where it came from.

DR. MAURO: But if TBD-6001 had an appendix which held the reader's hand and listed, this is where we can go right back

through the original data, to the database of 20,000 measurements sitting someplace on the site query database and it's explained that they sorted the data into a spreadsheet, which I'm sure they did, took the data and this is what they did and just walk the reader -- and what would happen if that type of description is provided as an appendix, you would have picked it up.

It's, you know, it would not have gotten to the point where we have a product out here that has some errors in it and contradictions in it. So in a funny sort of way, one of the things we find out when we are writing our reports, when we are writing our reports, when we are writing our reports, we try to write it in a way that someone could read, say, oh okay, and all the calculations are there, and in the process of writing it, you catch your errors. That's when we catch our errors.

I think that in this case, this is just a recommendation that if the final

1	product contains a hand-holding description of
2	
۷	exactly how we got the numbers that are in
3	this report, the errors would have been caught
4	and they wouldn't have to come to here to be
5	caught.
6	CHAIRMAN ANDERSON: How many TBDs
7	are there?
8	DR. NETON: Lots.
9	CHAIRMAN ANDERSON: Well, I mean,
10	to go back, it seems to be pretty massive.
11	DR. NETON: Well, I think this
12	might, I am not going to say this is an
13	isolated example but TBD-6001, I think was
14	rocky. I will admit that. So, hopefully, you
15	know, this is not
16	CHAIRMAN ANDERSON: So I guess
17	DR. NETON: indicative of a
18	whole problem.
19	CHAIRMAN ANDERSON: I am looking
20	for some reassurance.
21	DR. NETON: We have been through an

in-depth review of many, many TBD documents.

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1	DR. MAURO: Oh, we've reviewed
2	hundreds, hundreds of procedures, you know,
3	literally hundreds of documents.
4	CHAIRMAN ANDERSON: So, is the, the
5	kind of the bottom line, is NIOSH going to go
6	back through this and do another internal
7	audit of all of this?
8	DR. NETON: We need to go back and
9	refine it.
10	DR. MAURO: Six thousand is much
11	better.
12	Well, 6000 is a simpler document
13	since you're only dealing with uranium
14	machining. In other words, this is a more
15	complex document. You've got all these
16	measurements that you're working with as, you
17	know, so this is a tougher nut to crack but it
18	shouldn't be that difficult for us to check
19	numbers and you know
20	DR. NETON: And you look at what
21	happens here, I mean this is no excuse, but
22	you know you are getting down into what I call

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about 0.3	mR per day	exposure,	millirem	per
day for a	guy standi	ng it doe	sn't make	e it
right that	it's wrong	, but I'm say	ving when	you
get down i	nto these	-		
	CHAIRMAN	ANDERSON:	It's	the

CHAIRMAN ANDERSON: It's the impact.

DR. NETON: -- the impact of these very small, calculated doses, it doesn't make it right but possibly the attention to detail is not as great as if it is the big, bigticket exposure items and that is I think what has happened here.

ALLEN: And in all honesty, MR. with 6000 they were able to take the description that, you know, what we can claim we did, you know, run some numbers, verify that, you know. It might not have had the hand-holding as nice as you would want but you can verify the numbers from the description pretty much.

The reason they were not able to

1	in 6001 was there was calculational errors
2	behind the scenes so you couldn't verify what
3	they were saying. So it'd kind of indicative
4	of this document not, you know, a systemic
5	problem or anything.
6	CHAIRMAN ANDERSON: But it does
7	seem to me, before we take up Hooker, this is
8	probably going to need to be redone, or
9	DR. NETON: I am not sure, I mean,
10	Hooker is a very
11	MR. ALLEN: It uses a very small
12	piece of this.
13	DR. NETON: I think you can sort of
14	investigate the little piece we use in some
15	detail and get your hands around it fairly
16	quickly.
17	CHAIRMAN ANDERSON: Okay, well I,
18	you know, I just
19	DR. NETON: I would prefer not to
20	wait until we resolve all these issues that
21	are best taken up elsewhere.

CHAIRMAN ANDERSON: Well, I mean,

1	that's kind of what my question is, is this
2	sufficiently, that you know, we go back to the
3	Board and somebody is going to raise, well,
4	gee, until you have done that we have got to -
5	- or, and then, or can we basically
6	MR. THURBER: No. I don't think
7	that, from our perspective, we need to have
8	these questions answered. They are adequately
9	answered so that we understand them. As you
10	say Jim, this is peanuts in terms of exposure.
11	The question is how endemic the problem is
12	from a TBD-6001 perspective.
13	But in terms of what we need to do
14	for Hooker, I think we are okay with some, any
15	clarifications here.
16	CHAIRMAN ANDERSON: So that's all I
17	needed to hear.
18	DR. MAURO: The big issue on Hooker
19	is going to be Hooker uses TBD-6001 as a
20	surrogate. It's going to become a surrogate
21	data issue and the heart of the matter is

going to be, assuming that the numbers are

corrected, you know, see, I look at this as what I call a Site Profile issue. There's some bad numbers in the report; we've got some contradictory diagrams, all of which can be fixed, right?

Now, the real question is going to be, okay, once that's fixed, the question then becomes all right, at Hooker, and you brought this up before, at Hooker, let's say they used this particular number or distribution from TBD-6001 and we know what they did to get that.

And then the question is going to be, does that really, is that a good surrogate for this particular guy or any particular area at Hooker? And it becomes a surrogate data issue that's going to -- and that's where the SEC is going to comply.

In the end, the real, you know, the judgment calls, and this is the tough one and these are things that 16 Members of the Board are going to say, whether or not -- does

this represent a reasonable surrogate for the guys that worked in Hooker or not?

And you know that's going to be a difficult discussion to have but that's where it's going to end up, you know. The technical issues will be worked out. It's the judgment issues that are going to be the tough ones.

CHAIRMAN ANDERSON: Okay, so next steps on this.

MR. ALLEN: We owe responses on all these -- I think I can get you some short responses quickly and some others, we'll be saying, this will be addressed in a White Paper to come or I can wait until the whole White Paper is done and it will be a little longer.

CHAIRMAN ANDERSON: What, when we, what kind of a time line, holding your feet to the fire, when could, I mean a White Paper seems to be something that is really going to be helpful for us.

MR. ALLEN: Yes, I agree. I don't

know if you wanted the one-word answers, you
know, the one-sentence, two-sentence answers
first or just wait for the whole, you know
CHAIRMAN ANDERSON: While it's sort
of fresh in your mind I would like to try to
fill in short things
MR. ALLEN: That's what I was
thinking.
CHAIRMAN ANDERSON: pretty
quickly then
MR. ALLEN: That's what I would
like to do.
CHAIRMAN ANDERSON: I would like
to, by, you know, the meeting out in Idaho, be
able to at least report back when, what the
time line for the
MR. ALLEN: Okay.
CHAIRMAN ANDERSON: I would think
by then we ought to have the short answers and
hopefully in time for you to look at them and
say, yes, well, it's kind of, we can look at
it and say, yes, that's what we remember and

1	then what would be a time line for a White
2	Paper and I don't have a sense of how much
3	time and effort that will take.
4	MR. ALLEN: I don't think I have a
5	good sense on that at this point but I can
6	come up with short responses
7	CHAIRMAN ANDERSON: Yes I would
8	like to hear some kind of a target date you
9	know, that we can begin to look at and again
10	as the surrogate issues, I would like to be
11	able to report back some confidence level on
12	the some of the 6001 components so we are
13	arguing more on the policy and the other
14	issues than we are, oh well, this is still you
15	know, I don't want to add any more uncertainty
16	to the picture than we need.
17	DR. NETON: Can you put the date in
18	the response, I mean, would a date for the
19	White Paper, would that be in the response?
20	MR. ALLEN: Yes, that's what I was
21	saying.

ANDERSON:

CHAIRMAN

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Okay, that's

1	fine.
2	MR. ALLEN: I can give you the
3	short responses
4	CHAIRMAN ANDERSON: Yes. Great.
5	Yes. Good.
6	MR. ALLEN: and in the email I
7	am sending those I will give you a target date
8	for White Papers.
9	DR. MAURO: And SC&A has no action
10	item, so we are clean, right? So the only
11	action item I guess, Ted, once that White
12	Paper is issued, are we authorized to move
13	immediately or do we wait until you authorize
14	it? So the action item on our end is when the
15	White Paper becomes available, we are
16	authorized to immediately go ahead and take a
17	look at it and get it back to you.
18	CHAIRMAN ANDERSON: And you will,
19	when you get the initial responses, you will
20	fill in the
21	MR. KATZ: Updated matrix.
22	CHAIRMAN ANDERSON: Yes.

1	MR. KATZ: There will be an updated
2	matrix.
3	DR. MAURO: The first cut at the
4	updated matrix is from the review folks.
5	MR. ALLEN: Yes, the short
6	responses will be in the, essentially in the -
7	_
8	DR. MAURO: And then we will look
9	at that.
LO	MR. KATZ: Is your matrix a PDF or
11	a
L2	DR. NETON: It's in Excel, isn't
L3	it?
L4	(Simultaneous speaking.)
L5	MR. KATZ: A lot of your stuff is
L6	in PDF so you need to send it to them in a
L7	writeable file form.
L8	DR. MAURO: Can we get them to send
L9	this this probably went out as a PDF.
20	DR. NETON: But it's probably a
21	Word table.
22	DR. MAURO: Certainly we could send

1	it out as a Word version.
2	MR. THURBER: Okay, well, I am
3	sorry, what do you want?
4	MR. KATZ: So, SC&A's item is just to
5	send a writeable file.
6	MR. THURBER: A Word version of
7	that.
8	(Simultaneous speaking.)
9	CHAIRMAN ANDERSON: Well, and there
10	were a few typos which you identified too.
11	(Simultaneous speaking.)
12	DR. MAURO: Don't worry about it,
13	Bill, they got it in Word.
14	MR. THURBER: Well, we will, in due
15	course we will fix those typos.
16	DR. NETON: I mean normally I
17	understand most people send PDFs because then
18	you don't want people to start changing things
19	around.
20	DR. MAURO: Right, but not in this
21	case.
22	CHAIRMAN ANDERSON: Okay, do we

1	have anything else on overview 6001? Shall we
2	take a lunch break and hustle back and then we
3	can hopefully get through the other two, at
4	least the start of it? How much
5	MR. KATZ: Normally
6	CHAIRMAN ANDERSON: I suppose this
7	is where we eat. We could take the van over to
8	the airport and eat at the airport, I suppose,
9	otherwise. What is your plan?
10	MR. KATZ: Normally, we take ar
11	hour break.
12	CHAIRMAN ANDERSON: Okay. I have
13	got to check out anyway.
14	MR. KATZ: So it's 12:15 now, we'll
15	break until 1:15 and then reconnect the phone.
16	CHAIRMAN ANDERSON: Anyone on the
17	phone?
18	DR. NETON: Mark is.
19	DR. BEHLING: Yes, we are still or
20	the phone.
21	CHAIRMAN ANDERSON: Okay.
22	MR. KATZ: Okay so we will back at
	11

1:15.

2 CHAIRMAN ANDERSON: Be back at

3 | 1:15.

4 DR. BEHLING: Thank you.

5 MR. KATZ: Thank you for hanging in

6 there.

(Whereupon, the above-entitled matter went off the record at 12:14 p.m. and resumed at 1:16 p.m.)

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1	A-F-T-E-R-N-O-O-N S-E-S-S-I-O-N
2	(1:16 p.m.)
3	MR. KATZ: This is the TBD-6001
4	Work Group. We are just reconvening after a
5	lunch break. Let me check on the line and see
6	if we have Mark Griffon back.
7	MEMBER GRIFFON: Yes, I am here,
8	Ted.
9	MR. KATZ: Great. And then we can
10	proceed. I think we are ready to address
11	Electro Met.
12	CHAIRMAN ANDERSON: Yes.
13	MR. KATZ: So first on the agenda
14	is a brief presentation by DCAS, by Sam Glover
15	of the evaluation and petition.
16	DR. GLOVER: Does everybody have a
17	copy of the Evaluation Report or it's a
18	little different format than we have done in
19	the past so it's sort of presenting the piece
20	so
21	CHAIRMAN ANDERSON: All I have is
22	the
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1	DR. GLOVER: You have the ER, do
2	you have the presentation that we gave? I
3	don't have the detailed
4	CHAIRMAN ANDERSON: No.
5	DR. GLOVER: from you guys.
6	MR. KATZ: You don't have to re-
7	present it in its entirety the way you would
8	have for the Board, but just to refresh
9	everybody.
10	DR. GLOVER: Sure.
11	MR. KATZ: And get the ball
12	rolling.
13	DR. GLOVER: Let me just pull it
14	up, though, because that is probably the
15	simplest thing, to start where we left it at
16	in 2009 when we presented it to the Board.
17	So this has not been taken up in
18	front of the Committee yet, right? You got to
19	realize, I picked this site up after it was
20	initially presented so this is new to
21	MR. KATZ: This Work Group has not
22	met before

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1		DR.	GLOVER:	This	is	brand	new.
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MR. KATZ: This is brand new.

DR. GLOVER: October 2009, we presented the Electro Metallurgical Corporation SEC Report. We concluded that we could do all aspects of dose at the facility. Just very quickly, they began in 1943. SC&A correctly points out we had some discrepancies in some start dates but in the very beginning the MED, the Manhattan Engineer District, they began producing uranium metal and so they ran 53 intermittently, basically going through from uranium tetrafluoride to uranium metal using a thermal reduction process that was done, I believe, at Iowa. Is it Iowa? What's Iowa State. that? Yes. Ames. Yes. Ιt was developed there.

So it was basically a single facility that was carved out by the AEC to produce it all. It sounds like there was some preparatory work that may have been done at some other facilities ahead of that. They

have gone through a series of successions. They were acquired by Union Carbide. They had a couple of stand-by periods. Essentially, we concluded they only had one uranium, one nuclear component which was uranium but SC&A had some other concerns that they identified.

The petition was started on November 17, 2008. We qualified it in March of 2009 and we issued the report July 23, 2009. The proposed Class was all workers that worked in any area of Electro Met from August 13, 1942 through December 31, 53.

We evaluated the Class from April 1, 43 through June 30, 53, in part I believe based on our premise that work did not actually start there until April 1, 1943.

I won't go through all the different sources of information other than to say that we have continued to conduct searches of the facilities, including -- we have been working quite a bit with the Army Corps of Engineers, so we have gone up there and

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conducted site research for all AWE facilities up there.

We have also realized that Hanford had not gueried their full data sets, databases and so we searched on Electro Met and all other AWE facilities that we have active and so we have even some classified reviews and some documents and I have shared those with Kathy Robertson so she complete listing of all of our Hanford terms searched. So those that in the were are process.

And also, as you guys participated in NARA, the National Archives, there have been a lot of ongoing data access, data retrieval.

So just to give you an idea of the number of claims for the site, we have 98 claims. This is back on August 1, 2009. Ninety-two of those were completed for dose reconstruction; 44 percent of with a PoC greater than 50 percent.

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So the petition basis was that a few workers were monitored for external exposure and also the efficacy of the health protection and industrial hygiene programs.

It was evaluated. Our evaluation concluded that they intermittently monitored workers. They did issue some dosimeters in 44 and then later, 48 through 49, for which we have results, and we only limited bioassay from 44 and 49.

Now we did have breathing zone and area monitoring from 44 and 47 through 49 and we had basically production processes. Now we concluded that the production processes were essentially the same for the entire MED in AEC period when we did this report. There clearly documents in the thing that, in the SRDB, that we have that mention that health -that changes were done to the health practices.

So we made a conclusion here that the production processes hadn't changed. So

that was the basis for how we did our review. We know that they were exposed to extremely high levels of internal -- there were some very high -- very high air concentration data from 1948. And it is not replicated in the earlier time frames, way back 600 MAC air. These are near 800, 900 MAC I think, very large, very very high.

And the, basically, personal protection was, as we all agree, that it was much less than modern standards.

We conducted a number of sample dose reconstructions. The basis for these is that we essentially used the highest operating conditions for the worst-case person in 48 through the whole time frame.

And so we generated very, very, very high dose assessments for all the people who were evaluated and so the dose reconstructions are available to you guys but let's see if they summarized them on this piece here.

Even if you look at the external photon dose to the skin, it's 25 rem, 126 rem electron, X-rays are almost -- are very small: 15 rem of dose, rem from uranium, this would just be from the external standpoints. So very high doses are used to evaluate it. Not that very high is meaningful but I'm just trying to give you guys a feel for the magnitude of doses that were used at Electro Met.

I am trying to see if we have the internal; from 43 to 53, we used 60,000 dpm per day intakes during operational periods and 473 dpm per day for stand-by periods. In addition we used an ingestion of 1,178 dpm per day of natural uranium.

So obviously with those kinds of intakes, respiratory tract cancers, unless it's something, you know, unless it's a latency issue, are going to be almost always paid.

So, but very, very large intakes. However they also had very -- they had some

1	potential during the reduction operations for
2	some very large airborne measurements that
3	were taken. So anyway, based on our our
4	feasibility conclusion was that we can do
5	internal, gamma, beta, neutron and medical X-
6	rays. That's where we left the report and
7	hopefully that's at least a little bit of an
8	overview and a kick-off.
9	CHAIRMAN ANDERSON: How many staff
10	you said you really only had measurements
11	from 44 and 48, is that it?
12	DR. GLOVER: There are, let's see,
13	I would have to pull up the specifics in the
14	report but it's
15	CHAIRMAN ANDERSON: How many
16	samples do they have?
17	DR. GLOVER: You mean for bioassay
18	or for well actually the SC&A summarized
19	that on table one. I'll just go ahead and use
20	theirs because it's readily available right to
21	me.

For bioassay, for air samples what

1	do you need? What are you asking me?
2	CHAIRMAN ANDERSON: It's just
3	DR. GLOVER: Air samples
4	CHAIRMAN ANDERSON: Just a sense of
5	what's
6	DR. GLOVER: Let's see, they
7	mentioned having four air samples in 43, 15 in
8	44, less than 10 in 47, a much larger,
9	obviously, in 48 when the AEC begins
10	operations, with 154 in 48 and 215 in 49.
11	CHAIRMAN ANDERSON: In the SEC?
12	DR. GLOVER: The AEC taking over.
13	CHAIRMAN ANDERSON: Yes, I know, I
14	know, but I mean these, there, the SEC
15	petition ends when?
16	DR. GLOVER: Well, the building was
17	actually destroyed, or what do you want to
18	call it, it was removed. Let's see.
19	MR. THURBER: Fifty-seven, I think,
20	Sam.
21	DR. GLOVER: Fifty-seven?
22	MR. THURBER: I think that's when
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1	the demolition was done.
2	CHAIRMAN ANDERSON: Okay.
3	DR. GLOVER: So.
4	CHAIRMAN ANDERSON: I got you.
5	DR. GLOVER: Right, but the active
6	period was, I think 53 is when all of the
7	activity on site other than the demolition
8	would have
9	CHAIRMAN ANDERSON: And how much
10	biomonitoring do you have?
11	DR. GLOVER: With the bioassay it
12	looks like we had 67 in 44 and nothing again
13	until 49. Now in our report, we only relied on
14	air monitoring data. We did not use the
15	bioassay to try to, we just took the air
16	monitoring data and ignored any application of
17	bioassay.
18	So that's where we came.
19	CHAIRMAN ANDERSON: Okay. Okay.
20	Shall we move on to the are there any other
21	questions you have?
22	MEMBER FIELD: Yes, what was the

1	percent PoC again?
2	CHAIRMAN ANDERSON: Forty-four
3	percent.
4	DR. GLOVER: What do you mean, how
5	many were above 50 percent? Forty-four
6	percent.
7	CHAIRMAN ANDERSON: Which is higher
8	than the overall
9	DR. MAURO: You used the type M,
10	type S, in other words, depending on the organ
11	of your intakes, in other words did you use
12	uranium intake, or did you automatically
13	assume it was all S, or?
14	DR. GLOVER: It said S here. That
15	does not necessarily make sense but it
16	probably because, I mean
17	CHAIRMAN ANDERSON: Is Mark on the
18	phone?
19	MR. KATZ: Mark is on the phone.
20	DR. GLOVER: I would have to
21	double-check to see what the reality is.
22	Usually we use the most claimant favorable

1	DR. MAURO: That's what I was
2	asking.
3	DR. GLOVER: We all and S is the
4	standard default. It said type S in the ER
5	evaluation. We always would have used the most
6	favorable version. So
7	CHAIRMAN ANDERSON: Mark, do you
8	have any questions?
9	MEMBER GRIFFON: Not right now, no.
LO	CHAIRMAN ANDERSON: Okay.
L1	MEMBER FIELD: I just have one
L2	question. You based it all on air monitoring,
L3	is that what you said?
L4	DR. GLOVER: At this time it was
L5	all based on air monitoring, that is correct.
L6	MEMBER FIELD: And you have
L7	bioassay data. I just wonder if the bioassay
L8	data would reflect what the air monitoring
L9	showed. Would you have already looked at that?
20	DR. GLOVER: That is something that
21	I think NIOSH would partially because we
22	would have back-extrapolated from 1948 into

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So what we have done, just to give you a bit of a -- I know SC&A is going to provide a matrix on some of what they have done. But we are going through everything, all all the documents right now, of material we have, and getting a comprehensive listing of all of the data, external, internal, bioassay, air monitoring, so we can actually see, what do we have to fill in for gaps and so that's where we are.

CHAIRMAN ANDERSON: Okay. You're up.

MR. THURBER: All right. Let me talk a little bit about what we understand to be the physical situation of Electro Met.

Electro Met was a ferro-alloy manufacturing plant. They made ferrochrome, ferrosilicon, silico-manganese; products that they sold to the steel industry which were

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used as alloy additions to make stainless steel for example, ferrochrome, whatever.

So that was their business. This business was done in large electric arc furnaces. When they undertook their contract with the Atomic Energy Commission, they built a small facility in one corner called the Area Plant and the Area Plant was basically where - well, the Area Plant was where all the uranium production was done.

Now there may have been done some work done in the laboratory. We'll that. It was probably small, but -- so they had this facility called the Area Plant, where they did the operations that Sam described. the green salt, the uranium They qot tetrafluoride from Linde, they mixed it with magnesium, put it in a thermite reduction bomb, converted it to uranium metal, cleaned up the uranium metal, remelted it in a vacuum induction furnace and shipped the finished billets off to somebody else to be fabricated.

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So that was the operation that they did there. One of the concerns that we have, and it is reflected in finding one, is really the definition of the exposure cohort. You can see that, as Sam mentioned, the exposure cohort was designated to be all of the employees that worked at Electro Met.

And we think that this needs to be examined further because there was a cadre that worked in the Area Plant, based on the interviews that we have conducted and some of the documentation originally that NIOSH uncovered, there pretty significant was separation between the Area Plant and the commercial facility.

There was a fence, it was guarded, people had to have badges to get in and out. There was some evidence that some maintenance workers might come in a couple of days a month, but it was basically a separate, self-contained operation.

And so we think it is important to

determine whether it is appropriate to consider everybody at the plant as part of the exposure cohort when most of them didn't work in the Area Plant.

And so that is what this first finding speaks to and I would note that if you look at the original petition -- so there were actually were two petitions made to consider the possibility of a Special Exposure Cohort for Electro Met, and one of the petitioners said on behalf of her husband, I believe, that it should cover the Area Plant.

The other petitioner said that it should cover all of the employees at Electro Met. The two petitions were merged by NIOSH into a single petition which formed the basis of their Petition Evaluation Report.

Now if you look at the information available on the two petitioners, it's not clear that either of the petitioners worked in the Area Plant. In one case there is a letter that was provided to the petitioner who I

believe was the spouse of the deceased employee from Union Carbide and it said, on the basis of this -- what Union Carbide told the woman -- on the basis of our employment records, it looked like this man worked his entire career at the carbon products division of Union Carbide, which is a totally separate thing even from Electro Met.

I mean Electro Met was a subsidiary of Union Carbide, the carbon products division was a subsidiary of Union Carbide. Linde was a subsidiary, but their records showed that this guy didn't even work for Electro Met.

Now their records may be wrong.

I'm not saying that all corporations keep good records, but that is the evidence that is available.

With regard to the second petitioner, if you look at the information provided, it's clear that he worked for Electro Met, it's reasonably clear that he did

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not work in the Area Plant. So I just offer that as background.

But anyway, the stupid computer is gone. I talked too long.

GLOVER: It may be worthwhile DR. to iust briefly discuss that because we obviously don't set the covered facility. And the Department of Labor, we can, if we find things we can offer them, you know, what we find. It's their job to determine if a person is at this facility and right now if you go to the covered facility, it says Electro Corporation. Electro Metallurgical is the facility that is covered.

So clearly there is a building where this work was done at. We have intermittent records, you know, as far as occupational, we have to apply some kind of an analysis as to what kind of dose these guys could have received. So --

MR. KATZ: But you can specify, if there are no radiological exposures, you can

1	certainly, you don't have to define a Class
2	for individuals who have no radiological
3	exposure whatsoever, even if the covered
4	facility includes that.
5	So, I mean, as they are saying
6	DR. GLOVER: There is no record
7	support that they have
8	MR. KATZ: Whether they can
9	implement such a Class is a whole other
10	question
11	MR. HINNEFELD: The issue would be
12	whether the Class could be administered.
13	MR. KATZ: Right.
14	MR. HINNEFELD: And absent evidence
15	that people were in one place or another.
16	MR. KATZ: Right.
17	MR. HINNEFELD: You know, if you
18	have maybe the security clearance records,
19	which apparently don't exist, they apparently
20	had a destruction schedule shorter than 50
21	years.

And so, ask them for some piece of

information that would say this employee worked in this part of the plant, all we get is employment verification that they worked somewhere then.

We don't know of any other record that provides a more precise placement of the employees. We don't know, we don't have a segregation of the work force, a record of the segregation of the work force that remain today.

MR. THURBER: But isn't it true,
Stu, that there are records of who the
monitored employees that worked in the Area
Plant are, by name, by badge number?

MR. HINNEFELD: Well, there probably are, but it's always a little tricky from our standpoint to say that people who were not monitored were not exposed. Yes, there could have been, I mean, I would think that people were assigned there all the time. We have this discussion all the time. You know, people who were assigned there all the

1	time, well sure, they were probably monitored.
2	But what about maintenance craftsmen who
3	perhaps had the necessary requisites to get in
4	but they didn't monitor them.
5	I mean we don't have unless we
6	have the evidence of the monitoring program
7	being sufficiently robust, that they always
8	caught people, people were always monitored,
9	you know, we have really been hesitant to try
10	to delineate from that much, from monitored
11	versus un-monitored. Anyone want to help me
12	out here?
13	DR. NETON: I guess, I was
14	distracted here for a second but I am not sure
15	what the issue is. We are saying we can do
16	dose reconstructions.
17	DR. GLOVER: We didn't offer a
18	Class. The finding one, however, is that we
19	didn't confine, I guess, our
20	CHAIRMAN ANDERSON: But your dose
21	reconstruction

DR. NETON: But the point is that

1	we are saying we can dose reconstructions in
2	the area where these exposures occurred.
3	Right? That's what we are saying? So it
4	doesn't really matter
5	MR. THURBER: But no, you are
6	saying more than that. You are saying we can
7	do dose reconstructions for the entire Electro
8	Met facility.
9	DR. NETON: Right, and you are
10	saying the exposure only occurred in a certain
11	part of the facility.
12	MR. THURBER: Right. And we are
13	further saying that the access to that
14	facility was constrained by a fence and badges
15	and so forth.
16	DR. NETON: I mean, the question
17	here is, where the exposures occurred, can we
18	do those dose reconstructions with sufficient
19	accuracy. And I guess what you are saying is
20	we are assigning dose to everyone as if they

chimed in, which was true, that this is very

in that facility and Stu

all worked

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1	typical of our abilities to administer, to
2	define Classes.
3	We are hard-pressed, I don't think

we are hard-pressed, I don't think we have ever been able to go back, with some minor exceptions, and define Classes more narrowly than the entire facility, generically.

MR. KATZ: This almost, this predates the Class. You are saying you are doing dose reconstructions for individuals who worked outside of the area where there were radiological exposures. So you are saying you couldn't even, in their interviews and so on, place them in- or outside of the radiological area?

DR. NETON: This is an issue at General Electric Aircraft Engines at Cincinnati, this 8,000-person is а huge employer.

DR. GLOVER: Its exposure said in 1942. This is 70 years ago and since it was classified --

1	DR. NETON: Usually what happens is
2	you have people, workers will say, actually,
3	yes, they might have had access controls, but
4	as you already alluded to, maintenance staff
5	were common for those areas, they would go in
6	there, administrative support services would
7	go through those facilities. We typically end
8	up taking a very expansive view of this.
9	MR. THURBER: Well, I point it out
10	for the record, that's all, you know, I mean,
11	it is not in a sense a technical question, but
12	I think it's
13	DR. NETON: Well, what I was trying
14	to get out of this is, were you trying to say
15	that, given that we don't know who went in
16	there, it should be a Class because we are
17	assigning these large doses to all members,
18	because I think would that be
19	MR. THURBER: No.
20	DR. NETON: If that's not your
21	point then this

THURBER: No. I am not saying

MR.

1	that. That's not the point at all. No.
2	CHAIRMAN ANDERSON: Is there
3	anything in the record that, when they did the
4	biomonitoring, was everybody
5	DR. GLOVER: It was very small. I
6	mean, if you look at the
7	CHAIRMAN ANDERSON: I mean, we had
8	67 people
9	DR. GLOVER: In 1948, there were
10	1,156 external dosimetry results. There were
11	164 bioassay results. So, and you know, we
12	also, how do we know we have everything in
13	these records? We would have to have a record
14	to compare.
15	I don't know if we have ever gone
16	to Union Carbide and said, do you have a list
17	of people who only worked in that facility. I
18	would have to find that out. We would have to,
19	that would be the only way it could be more
20	restrictive. And the Department of Labor is
21	really responsible for putting them in the

building.

1	MR. HINNEFELD: Well, the fact that
2	they were employed in the covered facility
3	DR. GLOVER: In the covered
4	facility.
5	CHAIRMAN ANDERSON: The question is
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7	MEMBER GRIFFON: Henry.
8	CHAIRMAN ANDERSON: if it comes,
9	is that
10	MEMBER GRIFFON: I just wanted to
11	ask you a question for clarification.
12	CHAIRMAN ANDERSON: Go ahead.
13	MEMBER GRIFFON: Is this for 42
14	through 53 and is there another part from 53
15	to 59 or something like that?
16	DR. GLOVER: There is a shut-down
17	period from 1953 to 57.
18	MEMBER GRIFFON: Okay.
19	DR. GLOVER: That would be kind of
20	a residual contamination time frame.
21	MEMBER GRIFFON: So there was
22	another petition submitted for that later

1	period, is that correct, or because I am
2	looking on the website. It looks like there
3	were two.
4	DR. GLOVER: I will pull up our
5	report. We actually would have described those
6	in the summary.
7	MEMBER GRIFFON: Okay.
8	MR. THURBER: Both petitions were
9	for the same period, I believe.
10	MEMBER GRIFFON: Oh they were? All
11	right.
12	MR. THURBER: I think so.
13	MEMBER GRIFFON: Maybe I am
14	misreading that. But it looked like two
15	petitions and it looked one went up to 1959.
16	The proposed covered period, anyway, by the
17	petitioner.
18	MR. THURBER: Yes, and I believe
19	that was modified.
20	MEMBER GRIFFON: Oh, okay. Okay.
21	MR. THURBER: But let me see if I
22	can be more precise about that.

1	MEMBER GRIFFON: Because that
2	might, my other question would be, I guess, if
3	it's only going up to 53, it's pretty easy to
4	believe the assumption of no recycled uranium
5	issues or anything like that.
6	MR. THURBER: Yes.
7	MEMBER GRIFFON: If it goes to 59,
8	then I guess I was wondering but it sounds
9	like there were no, it wasn't operational; it
10	was residual period after 53.
11	MR. THURBER: And what actually
12	happened here is that the one petitioner who
13	originally proposed that it cover the period
14	from 1952 to 1959 but then NIOSH went back to
15	her and she agreed that it was acceptable to
16	change the covered period to be from 1942 to
17	1953.
18	MEMBER GRIFFON: Well what is the
19	covered period established by DOE?
20	MR. THURBER: That.
21	MEMBER GRIFFON: That is, okay, so

yes, it wasn't really NIOSH

she

had,

determining that, it was, okay. Okay.
DR. GLOVER: The covered time
period is 1942 through 53. So that is all we
can evaluate.
MEMBER GRIFFON: Okay. And there
was no thorium work done at this site at all,
right, it was just?
MR. THURBER: No.
MEMBER GRIFFON: Okay. All right.
Thank you.
MR. THURBER: Should we go on?
CHAIRMAN ANDERSON: Well, I mean,
what kind of an answer are we looking to, to
that first one?
DR. GLOVER: I think we can
summarize
CHAIRMAN ANDERSON: Is finding one
an observation or is it a
DR. NETON: No, I think we can
provide a response
CHAIRMAN ANDERSON: Okay.
DR. NETON: which I think we all

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agreed to, is that this is a covered facility and this is no way to apportion that --

CHAIRMAN ANDERSON: Yes. Yes.

DR. NETON: -- that work at anything less than the whole facility.

MEMBER FIELD: At most, you could probably ask DOL to ask if they information for who worked in that building. Is that something that is part of the process? DR. NETON: That would be process if we were deciding to add a Class, but we are saying that we could do the work, reconstruct exposures for all workers at the

ALLEN: One technicality MR. this is it's labeled as a Department of Energy facility and it only goes through 53 and that usually means there's some proprietary interest, which would be that Area Plant you are talking about, the AEC built that plant or owned, had some ownership in it, and with that the being covered facility, I think that

NEAL R. GROSS

plant, we could.

essentially means the Area Plant is the covered facility.

So any claimants we are getting technically, DOL has decided are part of that Area Plant, not part of the commercial plant that is adjacent to it, whether they are doing this accurately or whether, you know, how they are doing that, I don't know.

MR. THURBER: As they say, one petitioner said the Area Plant, the other one said Electro Met and when the petitions were merged, it's my understanding that whoever makes those decisions said that it will be the entire Electro Met plant.

DR. GLOVER: I will commit to clarifying the covered facility definition, what they really are doing. How does that sound? I will make sure we respond to that.

CHAIRMAN ANDERSON: I mean, the issue, if there are so many workers that weren't even exposed there but you are going to -- it certainly could be viewed as claimant

favorable, but certainly is not an accurate reflection of what their exposure was for the rest of Electro Met.

So you are way over.

DR. GLOVER: Only if I can put him there.

CHAIRMAN ANDERSON: I know, but I mean if the whole thing is, if you worked at Electro Met, even in their commercial facility, during this period of time, but they weren't -- see, that, --

MR. KATZ: You are not estimating those. You are not estimating doses beyond the AEC doses. So that's not --

DR. NETON: Dave Allen's point is a good one. By the time we get these, the Department of Labor has determined they worked at the covered facility. I mean, they worked at an AEC facility. That's what they are saying, and so that exposure is covered and so we reconstructed the exposure for the AEC period.

1	CHAIRMAN ANDERSON: So you would
2	only, you would have to put them into the
3	MR. KATZ: DOL is putting them in.
4	In effect, DOL is saying these people are part
5	of that building, even whether they are
6	accurately doing that or not, they are.
7	CHAIRMAN ANDERSON: And that
8	building is not in that whole commercial
9	plant.
10	MR. THURBER: No.
11	CHAIRMAN ANDERSON: That's where I
12	am confused.
13	MR. THURBER: Yes. If an employee
14	of Electro Met in this relevant time period
15	applies for compensation, NIOSH will do a does
16	reconstruction and decide on the basis of the
17	procedures here, the guidance that
18	the dose reconstructor uses, whether the man
19	is compensated or not, regardless of where he
20	worked within that facility. Is that correct,
21	Jim?
22	MR. HINNEFELD: Well, there's one
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step between their application and our dose reconstruction, and that is that the Department of Labor verifies that they were employed at the covered facility. And so -- at the end the Department of Labor sends it to us for dose reconstruction.

So what Dave's point was -- now, let me, here I am talking again. Dave's point was that this site is characterized as a DOE site, not as an Atomic Weapons Employer. What that means is that there was some piece of Electro Met that the DOE built. I think the contract even said they essentially had built or AEC had built, or MED, for this purpose.

And so it was essentially their facility. building, their And so realistically, then, the people who worked in the AEC part of Metallurgical are the ones who the covered employees. And are so, by extension, by the fact that it is in -- by considering it a DOE site, then our conclusion logically is that the Department of Labor, in

verifying the employment of that person, has essentially reached the decision that that person worked in the DOE site of Electro Met.

Now, in reality, are they doing that? I doubt it. I imagine they are writing to Union Carbide, Union Carbide says, they worked at Electro Metallurgical and that's all we know. I don't know if that's the case or not, but it, very often that is what happens.

So I mean, we can ask DOL what they are getting, you know, what they are asking for and what they are getting, we can do things like that. But, you know, from our standpoint, in terms of the logic of the law, to the extent that there is logic in this law, the logic of the law is that the decision has been made, essentially, by the Department of Labor that these people work in the DOE site at Electro Met.

And so we can proceed with, you know, we can, you know, it's to our advantage to just say, okay, then we are doing the dose

1	reconstruction. I mean, that's the logic of
2	the process.
3	MR. KATZ: Right. The only
4	distinction I would make to that though is
5	when it comes to an SEC petition, I think,
6	unless there's someone who has already been
7	through a dose reconstruction process, it
8	falls in our laps to determine that they are a
9	qualified petitioner.
10	MR. HINNEFELD: A qualified
11	petitioner.
12	MR. KATZ: Petitioner, right. So
13	that we would then have to verify that they
14	were employed by the proper AEC site. But if
15	they already came through the dose
16	reconstruction process and they are a
17	petitioner, we would assume that DOL had done
18	that.
19	MR. HINNEFELD: Okay.
20	DR. MAURO: But, if the definition
21	of the Class that's been qualified is Electro
22	Met, not the Area Plant but Electro Met, by

1	definition, there is that's it's
2	presumed that if you because by definition
3	
4	DR. NETON: But that's the
5	definition of facility. You can't qualify
6	anything else.
7	MR. HINNEFELD: All we did was
8	DR. MAURO: And that is fine, I
9	mean, all we are doing is pointing out that
10	there's this very large facility with lots of
11	people where clearly, based on the research we
12	did, most of the people that worked at Electro
13	Met were not involved in any of this activity.
14	The degree to which that can be
15	demonstrated, proven to be, to the
16	satisfaction of, I guess, the Department of
17	Labor, but the best we can tell, this was a
18	very large facility involved primarily in
19	metallurgical, commercial activities. A very
20	small part of it was involved in this
21	particular area.

The way in which the Class is

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1	defined is Electro Met. So by definition, if
2	you just go by the Definition of the Class,
3	anyone that worked at Electro Met
4	DR. NETON: Well there is no other
5	option to define the Class.
6	DR. MAURO: And that's fine. We
7	are just letting everyone know that we've got
8	the situation.
9	CHAIRMAN ANDERSON: And my point
10	being is if 90 percent of the people working
11	at Electro Met had no exposure, how reasonable
12	I mean we have to look at it, is
13	reasonable, are you
14	DR. MAURO: That's the only reason
15	we bring it up.
16	CHAIRMAN ANDERSON: dose
17	reconstruction for those people. How, you
18	know, can you really dose reconstruct for
19	everybody at Electro Met?
20	MR. KATZ: This is becoming
21	circular. But, again, the facility definition
22	is actually just the DOE portion of that

1	Electro Met. That is the facility definition
2	is what I am hearing from David.
3	MR. HINNEFELD: Well, it depends
4	on whether you are choosing the words or
5	whether you are choosing the designation
6	category. The words say Electro Met.
7	CHAIRMAN ANDERSON: Yes.
8	MR. HINNEFELD: The designated
9	category says it's an AEC facility or DOE
10	facility. So it's, you know, that's where the
11	
12	CHAIRMAN ANDERSON: I don't have
13	any problem
14	MR. HINNEFELD: departing from.
15	CHAIRMAN ANDERSON: with the
16	facility where it was work. My concern is
17	when you now open the Class far broader than -
18	- and is what if DOE is saying it's just
19	the facility at Electro Met where they did,
20	where they owned the building, whatever it is,
21	that's quite different. Now how they, if they

qualify everybody regardless of where they

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worked, if their intent was just that facility, then the dose reconstruction for that facility --

MR. KATZ: What I am saying to you is if this were an AWE facility, you would then have concern because the entire facility would be covered and you would have concern about any radiological exposures anywhere in the facility. In this case, because this is a DOE facility, the only radiological exposures that DCAS has to be concerned about are those the building of occurred in because those the only are covered radiological exposures.

MR. THURBER: In principle, but, as Stu said, in practice, it's not happening.

No, but you are missing MR. KATZ: my point. You are missing my point. The only radiological exposures that have to be reconstructed are the ones that occurred under the AEC operation. That's the obligation.

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Ιf people are pulled into Class because they worked elsewhere in Electro Met by the way this is administered, it's not a concern for DCAS because they only have to accurately estimate with sufficient accuracy the radiological exposures that are covered. And those are the ones within the DOE facility.

It's not -- it's a non-issue whether they are capturing any other exposures because they are not covered under EEOICPA.

CHAIRMAN ANDERSON: But you are assigning exposures.

But that's not the --MR. KATZ: they don't have be accurate for to exposure outside as long as they are capturing the exposures within -- that are covered, it does the job. And if DOL is funneling individuals in that didn't work in an issue for DCAS building, that's not terms of estimating doses with sufficient accuracy because they are not estimating any

1	other doses.
2	CHAIRMAN ANDERSON: But is the
3	dose that you are estimating, based on your
4	assumptions, accurate?
5	MR. KATZ: If DOL is saying we
6	can't distinguish who worked inside the DOE
7	facility so we are going to funnel everybody
8	through this, that's sort of a given. You are
9	given this individual, you have to assume this
10	individual did work in the building because
11	DOL cannot distinguish. So that is just a
12	given assumption.
13	CHAIRMAN ANDERSON: Moving right
14	along
15	MEMBER FIELD: Is the next step to
16	say you will clarify with DOL?
17	DR. GLOVER: I submit and I will
18	clarify
19	MR. HINNEFELD: we can find out
20	what they are doing.
21	CHAIRMAN ANDERSON: I think that's
22	all, that's really all I want because it

sounds	like,	you	know,	their	intent	is	there,
which i	s real	l√ w	hat we	are a	fter.		

MR. KATZ: Just, my point is, you could not -- the Board could not decide to create an SEC Class for the individuals who did not work in the building because these are too high exposures because there is no coverage, there's no other covered exposures to have an SEC Class outside of the people who worked in that building. There's no coverage. You can't create an SEC Class for people who aren't covered.

CHAIRMAN ANDERSON: Right.

MR. KATZ: Or for operations that aren't covered. That's all I'm trying to say.

CHAIRMAN ANDERSON: Okay. Number two. We are going to check.

MR. THURBER: We do the hardest ones first. Okay. There is some evidence that a limited amount of work may have been done outside of the Area Plant, particularly in the research laboratory.

1	I suspect that the amount of work
2	was small, but I think it's important that
3	this aspect be addressed and it be
4	demonstrated that whatever went on in the
5	laboratory doesn't really affect the ability
6	to reconstruct the dosage, or it would change
7	them in any way. That's all. Second finding
8	in the process.
9	DR. GLOVER: I didn't re-read your
10	piece when I re-read that but, so, since we
11	just had a discussion of facilities, do you
12	mean outside the covered building?
13	MR. THURBER: Outside the Area
14	Plant. Everything is covered.
15	DR. GLOVER: Yes, well, outside
16	the Area Plant.
17	MR. THURBER: Outside the Area
18	Plant, yes, there's a research laboratory
19	right next door, and there's some evidence
20	that some work was done there prior to the
21	start-up of the Area Plant.

GLOVER:

DR.

That would not be a

1	DOE facility, perhaps.
2	MR. KATZ: Not covered.
3	MR. ALLEN: You would have to go
4	to DOL and get them to recognize that as an
5	AWE.
6	DR. GLOVER: It would have to be a
7	separate designation.
8	MR. THURBER: Okay. Fine.
9	DR. GLOVER: I am not, I am just -
10	- okay, I just want, since we are clear, to
11	figure out, that's a, the finding two, I can -
12	- we can begin follow-up on that.
13	MR. THURBER: Yes. Sure.
14	CHAIRMAN ANDERSON: Did did
15	they know about it?
16	DR. GLOVER: Well, I saw some of
17	the things, we can look at that and see if
18	there's a reason, we can give them what
19	information; we do not create a covered
20	facility.
21	CHAIRMAN ANDERSON: Right.
22	DR. GLOVER: We can give them the
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1	evidence, and it's their choice.
2	CHAIRMAN ANDERSON: No, I mean, if
3	they are saying all of Electro Met, then it
4	would be covered. If they're not, it isn't,
5	that's really what would you do?
6	DR. GLOVER: Right.
7	CHAIRMAN ANDERSON: I assume you
8	haven't gotten any lab techs who have applied
9	yet, so they haven't made it through the
10	clearance site.
11	MR. HINNEFELD: I wouldn't
12	guarantee we didn't.
13	CHAIRMAN ANDERSON: Well, I mean,
14	if you did, then it would mean there's a
15	moderate likelihood, but moving right along.
16	MR. THURBER: Good.
17	CHAIRMAN ANDERSON: Go ahead.
18	MR. THURBER: Finding three is
19	kind of related. It says that there is some
20	evidence that a little work was done prior to
21	the start-up of the Area Plant. Was it
22	significant and does it change the start date?

That's something that is given to you, you know, we understand.

But, and on the other end of that, there was no provision for residual exposure in your Evaluation Report, and there was several years, three or four years, I guess, between the time that the AEC contract ended and the building was demolished, so there is a question of whether there was residual exposure or not. And so that all ties in with being sure that the --

DR. GLOVER: We could only do the covered -- because DOL closes it in 1953, that ends when we can actually legally evaluate it so -- but anyway, that is why it had to be terminated at that point.

MR. THURBER: Okay.

DR. GLOVER: But the start date I think, you know, you mentioned the start date, and I think we may have started in April of `43, and there is discussion there about them doing work before that in November of --

1	MR. THURBER: In the fall,
2	November of `42,
3	DR. GLOVER: `42.
4	MR. THURBER: You know, that sort
5	of thing.
6	DR. MAURO: You lost me a little
7	bit. So you are saying by definition, any
8	exposure workers might have experienced after
9	the covered period are not going to be
LO	included in the dose reconstruction?
11	DR. GLOVER: That is absolutely
L2	correct.
L3	MR. HINNEFELD: For a DOE facility
L4	there is no residual aspect. The residual
L5	contamination part of the law
L6	DR. MAURO: I've been spending too
L7	much time working on AWEs.
L8	MR. HINNEFELD: applies to AWEs
L9	MR. THURBER: Okay. We kind of
20	talked a little bit about this finding four
21	this morning. And the basic assumption that
22	NIOSH made regarding their ability to

reconstruct the doses was that even though there was very little information prior to 1947 or 1948, that one was dealing with the same process the entire period and so it was not unreasonable to take the `47, `48-era data and back-extrapolate that, if you will, to the beginning of operations in 1943.

We provide a number of arguments in our review paper that both support and refute that position. One of the arguments is cited here. There was a report from the AEC to Congress, and it says that many changes were made in the end of 1947 to improve the process.

So if that is the case, is it reasonable to say 1942 through `47 were the same as `48 and on, and we think that that point needs to be addressed more vigorously. And we provide, as I say in our document, a number of other arguments that speak to both sides of that coin.

DR. GLOVER: I think we also

1	agreed back-extrapolating in that time frame.
2	We are getting all the data together, as I
3	previously mentioned. We will look at how
4	bioassay affects our decision, how we best
5	move forward, maybe we compare that event to
6	the `48 data.
7	I will say that we are, I believe,
8	just off the top of my head 1,000 MAC
9	numbers which are near what we consider the
10	boundable, sustainable dust-loading from these
11	operations, I think. So, because we need to
12	look at all the different
13	DR. NETON: I just talked to Sam,
14	we have some 1948 bioassay data, and one could
15	balance that against the intakes that were
16	projected back in the 1,000-MAC era and sort
17	of see if they make sense.
18	DR. GLOVER: If the same people
19	work.
20	DR. NETON: If you have the same
21	people working, if they had worked during the

-- early years.

1	DR. GLOVER: We will certainly
2	pursue that.
3	MEMBER FIELD: The way I'm reading
4	this comment here, this second part, it sounds
5	like there are changes
6	DR. GLOVER: Yes.
7	MEMBER FIELD: in, maybe, the
8	amount of bioassay or other factors, but it
9	doesn't look like process.
10	DR. GLOVER: It's unclear. They
11	said that they made a survey and they made
12	changes to the operations, if you read the
13	full text, so it's possible that there may be
14	some changes that occurred. I think most of
15	it I believe they perhaps did more
16	respiratory protection, and they, at a later
17	time they do begin speaking about making these
18	guys wear respirators in that bomb reduction
19	room. So
20	MR. THURBER: And on the subject
21	of bioassays, we did an analysis in our review

paper where we compared the bioassays from

1	1943 and, what was it, `48. Whichever, and we
2	couldn't show that they were statistically
3	different. So as I say, there are arguments
4	on both sides of the question as to the
5	reasonableness of this assumption.
6	CHAIRMAN ANDERSON: Why don't we
7	just look it over, make a decision.
8	DR. NETON: Thanks for helping
9	out.
10	(Laughter.)
11	CHAIRMAN ANDERSON: Which way do
12	you come down?
13	MR. THURBER: job description.
14	DR. GLOVER: We did not fully
15	scope that because we relied on late-term data
16	and back-extrapolated. We are now fully
17	scoping all the data and seeing what our best
18	options are to evaluate our path forward,
19	whether that's bioassay, whether that's so
20	we have captured new documents since we left
21	this so
22	DR. NETON: Did you say there was

1	documents at NARA that you picked up recently
2	or
3	DR. GLOVER: There was NARA, there
4	was also, there are a number of new reports by
5	the, in the Buffalo Corps, so there's some new
6	there's like, just 50 or 60 that we have
7	seen just since I had Cheryl do a quick
8	search. There's other documents. There's
9	also some stuff at Hanford's. We want to make
10	sure that nothing in that affects our
11	decisions.
12	MR. THURBER: And we haven't
13	looked at any of that documentation either, or
14	not either, we haven't looked at that
15	documentation.
16	CHAIRMAN ANDERSON: When do you
17	expect this to be done?
18	DR. GLOVER: With the current
19	documents we have in house, I guess, I should
20	have a draft of what they put together by the
21	end of July

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CHAIRMAN ANDERSON:

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Okay.

1	DR. GLOVER: early August. And
2	so we will if anything else comes with the
3	other documentation. So, in August, I think we
4	should have our hands around what data we
5	have.
6	CHAIRMAN ANDERSON: Great. Okay.
7	MR. THURBER: Finding number five
8	is minor. There is an inconsistency in the
9	report regarding whether there is some
10	available data for the stand-by period or not,
11	and there is some data, as we indicate here,
12	that should be, that should be tidied up in
13	the report.
14	MR. KATZ: So is that a
15	clarification that needs to be made in the
16	report, is that what you are saying?
17	MR. THURBER: Yes. It says
18	original stand-by data we provided a
19	reference, or we have cited the reference.
20	It's one that we, everybody has, of x , that's
21	the

MR. KATZ: Thank you.

MR. THURBER: Finding number six speaks to basically a generic problem with sampling, air sampling, as to whether it is breathing zone samples, whether the device is attached to a worker's lapel, whether it's a general area sample away from the operation, or whether it's a general area sample that is quite close to the operation.

And in a lot of the sampling that was done, we believe that the sampling was -the fixed head samplers that were probably
fairly close to the operator but this -- it's
important that this point be addressed because
it does make a big difference.

DR. GLOVER: When we look at the date we will certainly make sure if we are using BZ process, GA --

DR. NETON: Were these the HASL, -DR. GLOVER: Well, `43 would have
been military, some of those guys, so we did
have some HASL measurements, so it would be
time-dependent. We will make sure that we --

1	we will make sure, and I'll we'll respond
2	to it, we will make sure we properly
3	MR. THURBER: Because, you know,
4	some of the `48 measurements, as you say,
5	there is some very high measurements in the
6	green room where the workers were exposed to
7	the uranium where they are putting the bomb
8	together, the uranium tetrafluoride being
9	mixed with the magnesium, and that was
10	terribly high there were terribly high
11	exposures in that area.
12	It appears that a lot of those
13	samples were taken four-and-a-half feet off
14	the ground and presumably at a fixed-head
15	sampler and it is important to establish
16	whether that measurement is the kind of
17	exposure the worker actually got or not. I
18	think that's what kind of underlies that
19	finding.
20	CHAIRMAN ANDERSON: Yes. Okay.
21	MR. THURBER: Finding seven, you
22	know, relates to job titles and whether job

1	titles really relate to job descriptions or
2	whether things were highly interchangeable,
3	and it is important to provide guidance that
4	if you, for example, if you don't know the job
5	descriptions, you assumed worst case or
6	whatever.
7	But it is a point that is not
8	adequately we didn't feel it was adequately
9	addressed within the Petition Evaluation
10	Report.
11	CHAIRMAN ANDERSON: Are you using
12	job titles though? I thought you were just
13	assigning everyone basically the same
14	DR. GLOVER: I didn't think we
15	were
16	CHAIRMAN ANDERSON: I didn't think
17	
18	DR. GLOVER: we weren't trying
19	to break down this in my review. We were
20	using a pretty maximizing addition to I
21	won't say a maximizing addition we were

using -- because we couldn't put people in

1	places, we took that exposure which represents
2	the 95th percentile or whatever, this is the
3	condition which would be bounding, and that's
4	what people were evaluated against.
5	DR. NETON: If you look at what
6	Sam said earlier, 90 claimants, if 44 of them
7	were over 50 percent PC. Clearly we haven't
8	been doing much in the way of triaging these
9	people by job title.
10	CHAIRMAN ANDERSON: But that if
11	it doesn't say that, that's a clarification
12	you can just
13	MR. THURBER: In Appendix C, you
14	do provide job categories, so the answer is
15	that
16	DR. GLOVER: Appendix C has some
17	information.
18	DR. NETON: We probably aren't
19	using the or
20	MR. THURBER: And they may not be,
21	but it's a question that needs to be

addressed.

1	DR. NETON: That's a good point.
2	MR. THURBER: Yes, what the dose
3	reconstructor does is different than what the
4	review talks about in some cases.
5	DR. GLOVER: We will verify
6	against what
7	MR. THURBER: Okay. In doing our
8	review of the Electro Met exposures and we
9	found that the inhalation intakes were quite a
10	bit higher than those in the generic document,
11	TBD-6001, which then raises the question about
12	TBD-6001 as to whether those numbers are
13	appropriately conservative and claimant
14	favorable or not.
15	DR. MAURO: So this is more a
16	comment on TBD-6001
17	MR. THURBER: Yes.
18	DR. MAURO: than it is on
19	Electro Met.
20	MR. THURBER: Yes, it is.
21	CHAIRMAN ANDERSON: Yes, does 6001

Electro Met --

DR. MAURO: It almost begs the question, okay, if you are going to use TBD-6001 and -- for some dose reconstructions at another facility, I guess there has to be some assurance that the conditions at this other facility are not such that, yes, that you have bounded it with 6001.

Clearly there are circumstances when even TBD-6001, which is designed to be bounding and trying to capture most AWE facilities, may not capture most -- some AWE facilities, and there's an example.

Obviously from this, if for some reason, Electro Met did not have the data, and you were going to assign a reconstructor -- you understand -- you would have missed the dose. You would have underestimated the dose for this person. So it goes toward TBD-6001.

MR. KATZ: So can we really relabel this because this is really a finding for the TBD-6001?

1	MR. THURBER: And we say that in
2	the bottom line
3	MR. KATZ: I just mean for future
4	tracking, we should really shift this over to
5	be a comment on 2001 TBD rather than Electro
6	Met comment, that's all.
7	DR. MAURO: Matrix.
8	MR. KATZ: So then David can knock
9	it off with his White Paper or whatever.
10	DR. GLOVER: How about we don't
11	just don't re-number that because we can still
12	use the same finding numbers, and we don't
13	but they just move somewhere else.
14	MR. KATZ: Sure.
15	DR. GLOVER: Not Sam's problem.
16	(Laughter.)
17	MR. THURBER: We will, that's the
18	initial response, this is moved to
19	DR. MAURO: You transferred it. It
20	sounds like
21	DR. GLOVER: Transferred it,
22	that's nice.

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MR. THURBER: Okay. Okay. Finding ten. One of the problems that people who were doing this bomb reduction experience was it's almost uncontrolled explosion of the bomb and a lot of contamination.

And so this clearly was an important issue which was examined in Petition Evaluation Report. NIOSH concluded that they had no evidence that these occurred at Electro Met. We looked at the same information, we said yes, that's what it says.

But we were troubled by the fact that these had occurred everywhere else that this process was practiced, and it was a fairly common occurrence. And so we felt that it needed further examination.

It wasn't clear to us what magic Electro Met had that they were avoiding this. Now when we prepared our review we said we are going to try and do some more interviews of people who worked there to see what we

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could find. At the time we did the report, we had talked to one worker who said he had no knowledge that blow-outs occurred, supportive of what the Petition Evaluation Report says.

Since that time we have talked to,

I believe, four more people, and three of them
said blow-outs did not occur. One guy said
well, sometimes we got a minor release but,
you know, nothing kaboom.

One of mу concerns with the additional people that we have talked to was they were primarily chemists who would only go out into the production area sometimes. Ιt clear that they wasn't were totally knowledgeable about this subject, not that it was a big place, because it wasn't, and you would think that if there was an explosion that everybody would know about it.

But we are trying to find some additional people who were actually working in production to see if we can put this to bed.

DR. GLOVER: Do we have your

1	interviews by the way, the additional
2	interviews that you have conducted, the ones
3	that you are discussing now?
4	MR. THURBER: I don't know. I
5	will have to talk to Kathy about that. You
6	should have them, yes.
7	DR. GLOVER: Because it's not
8	automatic that we get them. Let us make sure
9	that we
10	MR. THURBER: I will take care of
11	it.
12	DR. MAURO: It is part our
13	procedure that you get everything we get, and
14	I think it first goes through a DOE clearance
15	and then it goes to you. If you haven't
16	received it yet, you will.
17	MR. THURBER: But I will be sure
18	that that happens.
19	DR. GLOVER: Because in certain
20	time frames that is true and maybe more
21	recent, because I like anyway, I appreciate
22	it.

I	
1	MR. THURBER: Yes. We will take
2	care of it.
3	DR. GLOVER: That would be good.
4	Because I want to make sure we are not talking
5	to the same people.
6	DR. MAURO: This is pretty recent,
7	right?
8	DR. GLOVER: Because we may want
9	to follow up on some stuff, so we will make
LO	sure we continue to talk to people and make
L1	sure we are both aware of what we are doing
L2	so, aggravate the same people, since they are
L3	very old. They are knowledgeable about this
L4	process.
L5	MR. THURBER: Some of these
L6	interviews are really quite interesting, the
L7	people's attitudes about the situation. They
L8	are refreshing. Sorry.
L9	MR. KATZ: So SC&A is still are
20	you still hunting down some
21	MR. THURBER: We are trying to get
22	somebody that was in the production area

1	MR. KATZ: Right.
2	MR. THURBER: rather than in
3	the lab. Because most of the people were
4	chemists, you know, and I think kind of one
5	chemist says why don't you talk to this other
6	chemist, but we do have some other names we
7	are trying to locate.
8	DR. GLOVER: So is this still an -
9	- should we consider this an SC&A issue still
10	at this time, rather than passing it on to me
11	because we are still in, it sounds like you
12	are still
13	MR. KATZ: It sounds like they are
14	still trying to confirm that, so yes, it
15	sounds like it's still on your plate.
16	DR. MAURO: Yes, I mean, Kathy is
17	trying to schedule interviews with selected
18	people, yes.
19	MR. THURBER: I am quite happy to
20	leave this on our plate.
21	DR. GLOVER: I mean, I would, I
22	would love to be, you know, when an interview

1	happens, just so we could, you know, because
2	that way I can hear what they
3	DR. MAURO: Well, I mean, we
4	DR. GLOVER: say.
5	DR. MAURO: when we do that, I
6	mean, protocols require us to inform you that,
7	you know, we are making certain we have
8	identified certain people, we are scheduling
9	certain interviews, and we are required by our
LO	procedures to inform you of that, that this
11	process is moving forward.
L2	DR. GLOVER: We had a change of
L3	people, so since now it's my site, we may have
L4	an so we just want to make sure that
L5	DR. MAURO: Okay.
L6	DR. GLOVER: But that would be
L7	great.
L8	MR. THURBER: I will take care of
L9	it.
20	MEMBER FIELD: If we do some more
21	interviews and everyone says no, what's the
22	difference? Probably never know.

1	DR. MAURO: In the end, we are in
2	a situation where we are building a weight of
3	evidence that in the end, again, becomes a
4	judgment by the Work Group and then the Board,
5	whether or not the weight of evidence is such
6	that no, we can discount these explosions,
7	these blow-outs.
8	MR. ALLEN: I was going to say the
9	blow-outs happen when you build up a pressure
10	in a sealed container, and based on the
11	airborne you are seeing at Electro Met, I am
12	not so sure it was that sealed.
13	(Laughter.)
14	CHAIRMAN ANDERSON: How much
15	higher would it be if you had a blow-out? It
16	was leaking all the time.
17	DR. NETON: It's a got 1,000 MAC
18	air going all the time and in a blow-out
19	it's an acute event. It happens; it generates
20	some high airborne for
21	DR. MAURO: For a short time.
22	DR. NETON: short period of

time.

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MR. THURBER: But those, a lot of those measurements were made by HASL, and, you know, I would -- you would think they would have commented when they were making the measurements if this was an extraordinary occurrence rather than what they are just, what the sampler is sitting there sucking in. I don't know.

DR. MAURO: I mean, once the, you know, you have -- the way I always had it in my head was, you know, you have this bomb. That, when you reach that temperature, the conversion happens very abruptly, an extremely exothermic reaction. You know, and either the thing goes or it doesn't, and, you know, so, were saying that maybe where you it leaking all along, I -- once that switch turns, that an exothermic reaction, either you have an explosion and the thing falls apart and you've got a real serious problem, I mean, walls comes down, or you don't.

1	MEMBER FIELD: You would think a
2	worker would remember that.
3	MR. THURBER: That's exactly the
4	point, exactly the point, you know. Are we
5	done with this finding? Okay.
6	Finding eleven, I think we have
7	already talked about, and Ted advised that
8	that is a definitional issue that is outside
9	the scope of the technical review.
10	MR. KATZ: Right, it's not covered
11	exposure
12	DR. MAURO: Yes, I guess we should
13	withdraw that since
14	MR. THURBER: Yes, we will
15	withdraw it on that basis, not covered
16	exposure.
17	DR. GLOVER: By the time that this
18	facility was I mean, it's not our call.
19	MR. ALLEN: Well, now when you are
20	looking into, this is the residual one, it's -
21	_
22	DR. GLOVER: Right, I mean, but it

1	closes in `53 though. It closed, you know, we
2	could say, here's the information we have
3	about `53 to `57. It's Department of Labor's
4	call, though, if there's any covered activity
5	there.
6	MR. ALLEN: You are digging into
7	some of that
8	DR. GLOVER: Yes.
9	MR. ALLEN: and you are going
10	to find whatever you find and
11	DR. GLOVER: Right.
12	DR. NETON: contract, it's not
13	covered.
14	DR. GLOVER: That's true.
15	MR. THURBER: Okay. Finding
16	twelve. As Sam indicated, the bioassay data
17	was not used, but they obviously looked at it,
18	and they said that the calculated excretion of
19	uranium was less than what you would calculate
20	from the air samples so the air samples were
21	bounding. And we did some calculations which

we provided that suggested that might not be

1	the case, and so there's something that needs
2	to be sorted out between NIOSH's calculations
3	and ours.
4	DR. NETON: You used the same
5	urinalysis data to conclude that it
6	MR. THURBER: Yes, we took the
7	same data, you know, there's the
8	DR. NETON: Assuming what, acute
9	exposure scenario, something of that nature?
10	MR. THURBER: Yes. Yes.
11	DR. NETON: We will look at it.
12	MR. THURBER: Yes.
13	MR. KATZ: Sounds like you may
14	need to share your actual calculations.
15	MR. THURBER: Yes, and I
16	MR. KATZ: Yes.
17	DR. GLOVER: I mean it's here in
18	the report, so
19	MR. THURBER: I think that the
20	detail is pretty much in the report, but if
21	it's not, we can expand upon it, no question.
22	CHAIRMAN ANDERSON: How different

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DR. NETON: It would be hard to imagine, given the high airborne intakes that we are finding that the urine data would show that it could have been higher than what we are finding. I just, it seems --

MR. THURBER: Well, let me see here.

DR. NETON: -- hard for me to fathom how that could happen. But, you know, it all depends on your assumptions, and I guess we'll just have to take a look. I haven't looked at those assumptions so I can't speak to that.

MR. KATZ: So if you run through those and you have any problem identifying all the assumptions you need, you can just write - contact SC&A and get details to be able to confirm or refute the -- okay, so that's a DCAS follow-up.

DR. GLOVER: I think some of those are related to different exposure categories

1	like supervisors, if I'm reading your text
2	here correctly. So, yes, we will be happy to
3	we will make sure that we
4	DR. NETON: That ties in with the
5	previous finding which is what are we really
6	using for exposure scenarios.
7	MR. THURBER: Right. We
8	understand that it wasn't used, but,
9	commenting on your comment.
10	DR. NETON: That makes some sense.
11	DR. GLOVER: I think that's right.
12	Okay.
13	MR. THURBER: Finding thirteen.
14	This in a sense relates to let me back up a
15	second here. We were specifically tasked by
16	the Board to review the Petition Evaluation
17	Report for Electro Met. We were not tasked by
18	the Board to review Appendix C, which is the
19	Electro Met appendix to TBD-6001. But
20	obviously, in the course of reviewing the
21	Petition Evaluation Report we had to look at

some parts of Appendix C to -- because they

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tie together to a limited extent.

this relate And comment may Appendix C than to the Petition to Evaluation Report, but we said that important to provide specific guidance in the Appendix C as to what to do if you don't know what the operator's description is. We talked about this this morning. There was excellent language in TBD-6000 providing that kind of guidance. We are suggesting that that kind of quidance would also be appropriate in Appendix C.

CHAIRMAN ANDERSON: Okay.

MR. THURBER: The question of what you assume for the exposure from medical X-rays, we raised this before, and the document on medical X-rays suggests that you ought to use photofluorography, but the language in the document is a little hazy as to whether that guidance applies only to DOE sites or it applies to AWE sites as well.

And we think that point needs to

1	be clarified as to the intent, and if the
2	intent is that it should be applied to
3	everything, then the X-ray dose rates should
4	be revised to assume that the technique was
5	photofluorography rather than more
6	conventional X-rays.
7	DR. MAURO: This is a DOE site.
8	DR. GLOVER: This is DOE.
9	DR. MAURO: This is a DOE site.
10	MR. THURBER: Well, okay, then
11	DR. MAURO: Even more so.
12	MR. THURBER: Then the comment is
13	that it should be photofluorography.
14	DR. MAURO: Unless you have
15	evidence that the X-rays are the size, the
16	small ones versus the big ones.
17	DR. GLOVER: Five by sevens.
18	DR. NETON: I am not sure we have
19	any evidence of that
20	CHAIRMAN ANDERSON: One way
21	DR. NETON: one way or the
22	other.

1	MR. THURBER: There is some
2	language, I think, from one of the doctors
3	talking about X-rays, but whether he is
4	speaking generically or I don't know.
5	DR. NETON: My it's been a
6	while since I've gone through this, but it
7	seems the photofluorography was used when
8	there was mass screens because it was cheap.
9	You could take a picture, put the guy up
10	there, take a picture, and move on.
11	And that's why we thought when we
12	meant DOE sites I really think we meant to
13	imply there the larger sites, where there
14	would be a need for mass screening, where
15	these smaller, mom and pop type AWEs would not
16	benefit from that type of a procedure.
17	So I am not sure where Electro Met
18	falls. We will have to look into that a
19	little closer, but that DOE guidance is a
20	little misleading. What it really means was

MR. THURBER: Okay.

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larger sites.

1	CHAIRMAN ANDERSON: It would
2	depend on whether they had a program for the
3	general plant. We are on the same these
4	guys
5	DR. GLOVER: The PFGs would be
6	claimant favorable.
7	MR. THURBER: Oh yes, absolutely.
8	And
9	DR. GLOVER: Oh, we are not doing,
10	okay.
11	MR. THURBER: No, they are not
12	being done, and it is quite a big difference.
13	DR. GLOVER: Oh yes, it's very
14	large.
15	DR. MAURO: Three rem
16	DR. GLOVER: A couple rem.
17	DR. MAURO: versus
18	DR. NETON: As far as we have
19	no idea whether they were even doing annual X-
20	rays as part of the condition of employment,
21	and then you say okay, well not only do we not
22	know they were getting them, we are going to

1	assign this photofluorography dose, I mean,
2	it's sort of
3	DR. MAURO: But that's what TBD 6,
4	you know, Ron Kathren's TBD says that you
5	should do for DOE sites.
6	DR. NETON: For DOE. I think the
7	jury is still out on the AWE and the AWE-like
8	DOE sites.
9	DR. MAURO: And you will see that
10	comment on all our reviews on AWEs because
11	really there is nothing right now in your
12	guidance that says explicitly for AWE sites,
13	you know, assume it's just X-rays.
14	DR. NETON: Well, see, we did have
15	good evidence that it did occur at DOE sites.
16	DR. MAURO: Right.
17	DR. NETON: That's the key there.
18	We have no evidence that it occurred at these
19	smaller AWEs.
20	MR. THURBER: I understand that.
21	The language is ambiguous
22	DR. NETON: You're talking

1	we'll	address	it.

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MR. THURBER: Item fifteen finding fifteen. Again, this relates to We took what we thought was the Appendix C. same data set or what was close to the same data set and calculated the log-normal distribution parameters for the beta found electron exposures, and we agreement with the values developed by NIOSH in table C5 of Appendix C.

We had not such good agreement for the supervisor labor category and the other category. Obviously these are less significant in terms of the total exposure that a person would receive, but we ought to try and reconcile why these differences --

DR. NETON: I am not clear, did you do a different type of analysis, like a rank order versus a curve fitting?

DR. MAURO: I don't know.

DR. NETON: That could be why there would be a difference if it's based on -

1	- it's either a math error or a different
2	technique. I am just trying to figure out
3	which.
4	DR. GLOVER: We have got to see if
5	these classes even exist in our current
6	methodology. So we'll clean this up. We will
7	look and see what the final number
8	CHAIRMAN ANDERSON: It's kind of a
9	
10	DR. NETON: It would be
11	interesting to know why there's a big
12	difference like that because
13	DR. MAURO: Did Harry do the work-
14	up on that?
15	MR. THURBER: Yes, Harry did it.
16	DR. MAURO: It's probably it
17	may be a good idea to make sure that we did it
18	the same way you did it. That might be the
19	reason for the difference.
20	DR. NETON: I've got to well, I
21	would think it might be.
22	DR. MAURO: And then

1	CHAIRMAN ANDERSON: Should agree
2	at the fifth, at least. But the 95th, I mean
3	
4	DR. NETON: Again, I think if it's
5	a rank order fit versus a linear fit for the
6	data and then picking off the curve that could
7	make a difference. We have seen that before.
8	DR. MAURO: But usually it's the
9	95th where we really deviate, when you do a
10	rank order versus
11	DR. NETON: Not the fifth
12	DR. MAURO: I know.
13	MR. THURBER: Yes, and I
14	DR. NETON: I don't know how
15	sparse this data is though.
16	MR. THURBER: I don't remember. I
17	Harry typically uses the calculational
18	method rather than the rank order graphical
19	method, and that may be the cause. As we have
20	pointed out in some other areas, you can get
21	some fairly significant differences actually,
22	depending on what the tails look like and

1	things.
2	DR. NETON: How you treat non-
3	detects.
4	DR. GLOVER: We are also making
5	sure that we have as much data as we have
6	available. I don't know if we had the same
7	data set to work with so
8	CHAIRMAN ANDERSON: And this is
9	table C in the
10	DR. GLOVER: Well, they compared
11	the values that were generated, they didn't
12	say whether they used the same data. They went
13	and independently looked through the data.
14	MR. THURBER: Yes.
15	DR. GLOVER: So the numbers may,
16	you know, they may have pulled additional data
17	together since we put Appendix C together.
18	Appendix C was not updated when we did the ER
19	so that's several years old. So it could be a
20	number of, you know, a number of years out of
21	date.

THURBER:

MR.

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Finding

Okay.

1	sixteen again ties in with Appendix C and we
2	note that table C5 and Appendix C, which deals
3	with external exposure, has a category of
4	"other skin" which is the skin other than the
5	hands and arms.
6	But there is no information on
7	hands and arms and we think the data should be
8	added to specifically address the hands and
9	arms.
10	DR. NETON: So this is the top crop
11	issue with the surface contamination? Did
12	Electro Met actually re-melt already cast
13	ingots or derbies or did they
14	MR. THURBER: Yes. They did vacuum
15	induction melting of the bomb reduction
16	derbies.
17	DR. NETON: That's only important
18	issue if you have aged product, correct?
19	MR. HINNEFELD: Did they recast or
20	did they just do bomb reduction?
21	MR. THURBER: No, they recast.
22	MR. HINNEFELD: They did recast?

1	MR. THURBER: Yes.
2	DR. NETON: Otherwise a freshly
3	separated metal won't have this product.
4	MR. THURBER: Well, it depends on
5	how long since, well, whether there was any
6	time line. They also handled scrap. They
7	recast scrap.
8	They did some scrap recasting
9	where they did receive scrap from outside
10	sources.
11	DR. NETON: And then it's a matter
12	of what percentage of that versus the total.
13	We will look into that.
14	MR. THURBER: We have really kind
15	of discussed this question earlier. The
16	question is if, how appropriate is the back-
17	extrapolation approach to the period prior to
18	1948, when the bulk of the data is available.
19	We have touched on that already did that
20	again.
21	CHAIRMAN ANDERSON: Okay. Any,
22	Bill? Mark are you there? So as far as the

1	whole Electro Metallurgical, you are still
2	reviewing some of the data, right?
3	DR. GLOVER: I will be at Hanford,
4	Sandia, probably Simonds Saw and Steel. I have
5	three weeks of travel so in that time SC&A or
6	ORAU will be completing putting the data
7	together, hopefully end of this month,
8	beginning of next. We will see what additional
9	data captures flow out from Hanford.
10	So I would, hopefully in August we
11	will have our hands around the data and then
12	we can make some decisions about the best way
13	to, you know, approach this.
14	CHAIRMAN ANDERSON: Okay. And we
15	have some action items in terms of
16	communicating some of the calculations we made
17	
18	DR. MAURO: Yes.
19	CHAIRMAN ANDERSON: And sharing
20	them with you, so
21	DR. MAURO: Yes.
22	MR. THURBER: And I think that to

1	the extent that what we have done is not
2	transparent, we will do whatever we can to
3	help you, provide you with whatever you need.
4	If what's there is sufficient great, if it's
5	not
6	CHAIRMAN ANDERSON: Are we going to
7	do a query to DOL?
8	DR. GLOVER: That's certainly part
9	of the we will have to I will, me and my
10	boss, we will work
11	DR. NETON: Are we going to fill in
12	the matrix though, in the interim? Or are you
13	guys, I mean what's the
14	DR. GLOVER: We can certainly,
15	because we both agreed, John and I, or
16	whatever, we will list that through our thing
17	and we will make sure who owns what action
18	items and that will be on this. It will show
19	who is doing what.
20	DR. NETON: I think it is important
21	to get this matrix filled out.

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(Simultaneous speaking.)

1	DR. MAURO: Sometimes it is helpful
2	to get the transcript, but right now I know
3	that we have relatively little to do.
4	MR. KATZ: You really have only
5	responsive things.
6	DR. MAURO: Right.
7	MR. KATZ: If they need your help
8	clarifying how you calculated certain things
9	then they are going to come to you for that?
10	DR. NETON: Well, with the
11	exception of the additional interview
12	MR. KATZ: With the exception of
13	the interviews
14	(Simultaneous speaking.)
15	MR. THURBER: We will provide that
16	information. I have that and
17	DR. GLOVER: It may be as we talk
18	about the areas up there, I'm likely to go to
19	Simonds Saw and Steel again and if we, at the
20	same time Steve Buskay of the Army Corps says,
21	if it would be useful, I could actually get on

site if I wanted to see what, you know, where

1	this place was and how it fits into the big
2	operation
3	DR. MAURO: They are close to each
4	other?
5	DR. GLOVER: They are on Niagara
6	Falls.
7	MR. THURBER: Yes, everything up
8	there, cheap power, that's why everything
9	DR. GLOVER: And if that is, if
10	that turns out to be the case I will make sure
11	you guys understand when we are going to do
12	that.
13	MR. THURBER: One of the things I
14	look for, and I couldn't find it, just for my
15	own perspective on this, how big is Electro
16	Met is compared to how big was the area plant?
17	I couldn't find any information on how big the
18	total work force was at Electro Met.
19	DR. GLOVER: It may be worthwhile,
20	if we got on site and talked to the folks
21	there.
22	CHAIRMAN ANDERSON: That's why I

	was asking about now many
2	DR. GLOVER: to see if they could
3	actually, you know, the guys at Union Carbide
4	have a better feel for it, so
5	DR. MAURO: And in the plot plan,
6	is there, how, just like, the area was
7	MR. THURBER: The plot plan was
8	only, shows the corner of the facility. It
9	doesn't show the furnace area or anything like
10	that.
11	DR. MAURO: Oh, this is the one
12	dealing with the residual period, in other
13	words, we were not aware that the residual
14	period is not within the scope of the dose
15	reconstruction for DOE sites and that was one
16	of our comments. I forget what number it is.
17	MR. THURBER: Yes
18	DR. MAURO: But we withdraw that.
19	DR. NETON: I would mark it closed.
20	DR. MAURO: Closed, yes. Closed is
21	the right answer and we will put the reason.
22	MR. KATZ: It was another one where

1	their work was outside of the covered facility
2	possibly, but that depends on what they find
3	out from DOL.
4	MR. THURBER: And there was another
5	one that is really assigned to TBD-6001.
6	DR. MAURO: Transferred to
7	MR. THURBER: And that too.
8	DR. NETON: Well, my only concern
9	was just closing them and then if they show up
10	in these roll-offs of numbers
11	CHAIRMAN ANDERSON: Yes.
12	DR. NETON: And it inflates some of
13	the magnitude of the issues that are I mean
14	it's trivial in a way, I mean, very few
15	DR. GLOVER: Usually when they're
16	withdrawn though, it's with concurrence by the
17	Board.
18	DR. NETON: Oh yes. Oh yes.
19	DR. GLOVER: So I don't know if you
20	guys have to do anything formal, but you guys
21	have to

(Simultaneous speaking.)

1	DR. GLOVER: the Working Group
2	should, yes.
3	CHAIRMAN ANDERSON: Should we take
4	a break?
5	(Whereupon, the above-entitled
6	matter went off the record at 2:44 p.m. and
7	resumed at 3:01 p.m.)
8	MR. KATZ: Okay, this is the TBD-
9	6001 Work Group and we are just reconvening
10	after a short break. Let me check on the line
11	just to make sure we have Hans because I think
12	we need him for United Nuclear.
13	DR. BEHLING: Yes, you have me.
14	MR. KATZ: Oh great. Thank you
15	Hans. Happy to have you.
16	DR. BEHLING: Actually I was on the
17	line when you asked before the break and I
18	couldn't hit my mute button, I kept pushing
19	the wrong button.
20	MR. KATZ: I was faster to hit this
21	mute button than you were to hit yours, I
22	guess.

CHAIRMAN ANDERSON: Take it away.

DR. BEHLING: Okay. I was talking to John during the break and he said we will be working with the matrix so I will simply follow that but I hope that for any particular finding or discussion that may ensue you can ask questions that may divert from the limited discussion that we see on the matrix.

Let me just briefly go over the chronology of events here because they are somewhat unusual here. The first Appendix D, Revision 0 was issued by NIOSH on March 14, 2008 and we in turn were asked to look at that Appendix D and our audit in our initial report regarding our review of Appendix D was issued somewhat about a year-and-a-half later, in September 2009.

Unfortunately we never received any actual response from NIOSH regarding our review of Appendix D and it wasn't I guess until early this spring that NIOSH elected to revise Appendix D with Revision 1 and at that

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point I believe Ted had asked SC&A to forego any discussion of our initial review of Appendix D, Revision O and focus on Revision 1 and basically roll up this audit into a final review.

And so what we ended up doing was we issued a supplemental report on June 11, 2010, just a couple of weeks ago, a few weeks ago, that reiterates our initial findings and then looks at the Revision 1 of Appendix D to see to what extent our finding still stands.

And I think our matrix pretty much reflects that evolution of events, that is we identified the initial findings and then in again, response now, it's question to whether or not NIOSH as actually looked our original at audit Appendix D, Rev O, in rewriting it, or whether these things independently came about.

But regardless I think they can comment on the issue if they choose. But we will simply identify each of the six findings.

In addition to the six findings we did have one single observation which was a generic one and I think we will understand what the issues are when we go through the findings.

So let me go through the findings. In our review of Rev 0, we had identified that our review of the medical dose was somewhat too brief. I think, in fact, let me just quickly read what it says so that we can get an understanding of the issue.

In the TBD, the guidance was that there are no diagnostic medical X-rays to which workers may have been exposed and there was no information regarding this. And so in essence, the recommendation was to look at the guidance in OTIB-0006 as a way of establishing medical exposure doses.

And my feeling was that that is somewhat overly brief because OTIB-0006 really provides various options and I think we need to be a little more definitive, not so much in a sense where this will potentially add a

significant dose to anyone.

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But the fact is we need a certain amount of consistency. I think we've been adamant of consistency. So that when you have a half dozen dose reconstructors doing a dose reconstruction for UNC, you will hopefully not have one who will take liberty in using exposures that perhaps are within framework of OTIB-0006 in a conservative way and another one using obviously the least conservative approach.

And so this is our first finding, is to assume that we need a little bit additional guidance with regard to how to assign medical dose because of the fact that right now all they do is refer you to OTIB-0006.

And I think this is something that reasonably easily be corrected because we all do need to give is either we will agree to assign only one PA X-ray per year or perhaps in a more liberal way, we can talk about

1	photofluorography perhaps, in the early years,
2	or in addition to photofluorography, lumbar
3	spine radiographs, all of which perhaps
4	augment the assigned dose from medical X-rays.
5	So when I looked at one I looked
6	at the issue of the medical dose and there was
7	no additional changes, so apparently, from
8	what I gather, that issue was not addressed in
9	Rev 1, either because ORAU or NIOSH never
10	really looked at the initial finding or
11	decided that the guidance in OTIB-0006 was
12	adequate.
13	Would it be appropriate to ask
14	NIOSH to respond at this point, or
15	MR. KATZ: Yes, Hans the way we
16	have been doing it is going finding by finding
17	so, thank you.
18	DR. BEHLING: You want me to just
19	continue?
20	MR. KATZ: No, no, no, in other
21	words so yes, in other words NIOSH will
22	respond to finding one and then you will

1 present finding two and so on.

DR. BEHLING: Okay, so it's NIOSH's response that we are waiting for.

MR. KATZ: Yes.

MR. ALLEN: And we agree with you that the Appendix could benefit from a more clear discussion and as far as what the exact procedures are and stuff, I think we could have brought that up with Electro Met, too.

So I do apologize for not addressing it in the revision. The revision was undertaken when we wrote the Evaluation Report and we found some new data in looking for that and we decided to revise the Appendix to incorporate the new data and what was in the Evaluation Report and this initial review of the Appendix got lost in the shuffle somewhere.

CHAIRMAN ANDERSON: So for United Nuclear we have a Site Profile.

MR. ALLEN: Yes, it's an appendix to TBD-6001.

1	CHAIRMAN ANDERSON: So the Site
2	Profile, as far as documents, the only
3	document we really have is Appendix D?
4	MR. ALLEN: Yes.
5	CHAIRMAN ANDERSON: It does both.
6	There's no SEC petition?
7	MR. KATZ: No, there is SEC
8	MR. ALLEN: Yes. There is an
9	Evaluation Report.
LO	MR. KATZ: And it is an Evaluation
L1	Report of the petition.
L2	CHAIRMAN ANDERSON: Okay, I was
L3	just, because the matrix
L4	MR. KATZ: We sort of skipped over
L5	that because
L6	CHAIRMAN ANDERSON: Yes. Yes.
L7	MR. KATZ: They were going to
L8	present
L9	CHAIRMAN ANDERSON: Yes.
20	MR. KATZ: something on the
21	petition initially before Hans went
22	CHAIRMAN ANDERSON: Yes, okay.

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	DR.	BEHLI	NG:	With	rega	ırd	to	tha	t,
and I am	sure	that	we	will	discu	ıss	or	NIO	SH
will discu	ss th	ne SEC	! pet	tition	n and	the	ER	th	.at
they have	gene	erated	l, j	ust	as a	sur	nmaı	ſΥ	up
front, I	see	none	of	the	find	ings	tł	nat	I
identified	as S	EC is:	sues	. So					

CHAIRMAN ANDERSON: Okay.

DR. BEHLING: I might as well make that statement up front. I think all of the findings that I have are solvable with a certain amount of additional information or data or guidance.

So are we prepared to then discuss finding two?

CHAIRMAN ANDERSON: Yes.

DR. BEHLING: Yes. In Rev 1 of Appendix D, the initial assessment for dose reconstructing external exposure was really confined to a couple of summary reports that were generated by the Atomic Energy Commission in 1960 and from that data it was concluded extrapolate that could for the full we

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duration of the operational period.

And Ι looked at the data and realized that that was probably not something that you would want to necessarily do, given the fact that there were many process so changes over the period of operational times and therefore the single summary data that were presented in the AEC report, the limited, would report, had а very questionable approach to satisfying all of the questions that we did have regarding latter years of operation and process changes.

And so that was my finding number two and in looking at Revision 1, as it turns out, NIOSH was able to identify a significant amount of additional data from various time of that data frames, some was external exposures defined units of dose as well as beta skin doses for some years. For other years, the data was lumped together so that a shallow dose and deep dose was combined.

But based on the fact that for

many years, or for a good number of years, doses were in fact separated, they used a fractionation process by which the data for the years where the photon and beta doses were segregated, they would apply that ratio to those years where the dose was combined.

And even though it was somewhat less than the most desirable form of data, I think the data are adequate for filling in the gaps for all the years during the operation.

So as far as I am concerned the issue of finding two as being inadequate has been reasonably well resolved by the use of additional data that allows for both deep dose as well as skin dose, in addition to answering certain questions regarding the frequency of badge exchange which occurs early on, on a weekly basis and subsequently on a monthly basis.

DR. MAURO: Hans, this is John. On the matrix I noticed that it sounds like you did take a look at the new data to some degree

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but on the matrix it indicates that we really haven't take a close look at it.

In other words, it sounds to me that, based on your preliminary review of the new data, it looks like that they have got quite a bit of data that fills in the gaps. Do you feel that SC&A needs to do a little bit more investigations into the completeness of that data, whether it covers all the workers necessary, all the time periods -- in other words, I guess what I am looking for is whether an action item here or not, because it does appear that there is an action item.

DR. BEHLING: Yes, you're absolutely right John and I wrote that into my supplement to the initial audit but as turns out, because Ted had requested SC&A to do just a very initial review of the SEC petition and the ER report, as it turns out, it wasn't until I looked at the ER report that actually identified the Site Research Database that was used to supplement that

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data, which were not identified in the Revision 1.

So as it turns out, when I did a review of the SEC petition and the Evaluation Site Research Report, where the Database identified numerous documents, I was able to actually look at dosimetry data and conclude that on a fairly substantial review process, I don't say it was exhaustive but I looked at enough to convince myself that supplemental evaluation of the data question is a reasonable approach and I think they did a reasonably good assessment of that available data in filling in the gaps.

So as I said, it wasn't until I looked at the Evaluation Report that I had a chance to identify those documents that were used to fill in those gaps, and that occurred after I had actually submitted the audit for Revision 1.

DR. MAURO: So what I am hearing is we don't have an action item here. You have

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reviewed the data to the extent that you are satisfied. Now the only thing I want to bring up, and this goes, really a question to the Work Group, to the Board Members, is when we do an SEC, the work we did here was what I call as Ted requested, let's take a look at it so that we could come to this meeting and inform you of what we did and where we are right now in the process.

Our SEC reviews, when formal review, is quite a bit more exhaustive. We will go, I mean, we will go into the data, download it all, load it into spreadsheets, do statistical analysis, look the time at covered, look at the different types of work activities, whether everyone was covered or not, everyone was not covered with the film badge.

We will look at, is there a coworker model, is the coworker model robust?
We will also include some interviews. I don't know whether or not any interviews have yet

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been performed for United Nuclear to do some data capture.

So in other words what Ι am getting at is I don't want to leave the Work Group with the impression that we did what I We did would call a comprehensive review. clearly a review to the extent that Hans is feeling pretty good about it, but I think it would be inappropriate for me to say that we have done the things that we normally do in terms of, for example, for an SEC petition. Right. And that being said it's really up to the Work Group as to whether there's any more formality you would like.

Because right now you can see by the report itself, relatively brief, and you know, if you would like something more formal -- and also there's a cycle of reviews within the company, within our group, where two or three independent people will check the work also.

So I think you are getting sort of

like a preview of where I think we are right now in our understanding, but I think I would not refer to this as a complete, formal SEC review as we have done in the past for other sites.

DR. BEHLING: And I agree with you John, as I said, this was not an exhaustive review. I did review a whole series of Site Research Database references. One of the things that did strike me was that while there was a large number of people with nominal exposures, and they are consistent with the doses that will be assigned for non-monitored coworker model.

But there were instances, and I am looking right now at, especially for 1966 for individuals who were exposed. In one case I am looking at nearly six rem for the year, for 1966 there was some, a couple of other people here, I highlight them, with 2.5 rem, another one at 6.3 rems and so on.

So it looks like a distribution of

doses that is marked with large numbers of people with modest doses, but then there were some people in perhaps selective job categories or locations, like at the red room, whose exposures were fairly high for the year.

And that might impact certain assumptions about the use of 95th percentile values, although I would assume obviously in these cases, where people had high exposures, these people were all monitored, obviously, otherwise we wouldn't know about it, and that those high exposures were, in fact, committed to people who were obviously identified as high-risk workers and therefore they were monitored, so we do have this data.

DR. MAURO: Is there a coworker model in the Rev 1?

DR. BEHLING: Yes.

DR. MAURO: There is. And I guess, it sounds like you did take a look at that and you feel that it covered all the time periods and different -- the data that does exist, is,

1	cuts across the time periods and the job
2	categories and locations to the extent that
3	the coworker model can, you know, be used, to
4	assign exposures, to workers who perhaps were
5	not monitored but should have been?
6	DR. BEHLING: Well, to look at
7	that, the external doses were defined in table
8	D-2 and they were modified considerably
9	between Rev 0 and Rev 1.
10	DR. MAURO: They went up by about a
11	factor of 10 as I recall.
12	DR. BEHLING: Yes, they were
13	substantially raised, yes.
14	DR. MAURO: Although, Hans, what I
15	am trying to do is get a sense here of whether
16	you feel that you, that SC&A, given what you
17	have done so far, whether or not, you know,
18	there is more to be done to make this a formal
19	review.
20	I have to say, as SC&A project
21	manager, I would like to do something where we
22	can say we went through the full procedure. We

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have a procedure. SC&A has a procedure to do SEC reviews, and there's a whole bunch of things that we do, which includes interviews and it includes data capture and site visits, all of which really haven't been done. So I --

MR. KATZ: Let just also, me though, explain, I mean, the Board has done several things with SECs and in some SECs it's done focused reviews where it has had particular issues that said SC&A go dog these issues down and in other cases it's had SC&A just sort of do, like John is saying, whole A to Z review of an SEC petition and it really depends on what is in front of the Board and what the Board's concerns whether it unloads an entire review of all the issues comprehensively or it has particular concerns that it feels it needs buttoned up.

So that is a judgment that this Work Group will make as to how extensively you want to use SC&A to dig into issues, any issues that you may have concerns about.

1	MR. THURBER: Well, John, do we
2	need to separate the Appendix C issues as
3	compared to the SEC issues? I mean, just, you
4	know, fundamentally we started out here
5	talking about Hans's review I'm sorry
6	Appendix D, not Appendix C Appendix D
7	issues, and then we have kind of deviated a
8	little bit into the SEC issues and are they
9	two separate things and should we be careful
10	to keep them separate?
11	MR. KATZ: Hans' statement was that
12	he didn't see any SEC issues among the TBD,

above all the review that he has done at this point.

But certainly if you, as you go through these findings, I think we are still on finding number two --

DR. MAURO: Well, maybe, put this issue in the parking lot for now. You have got a sense of what we did. Clearly, on this matter regarding external dosimetry, clearly going from Rev 0 to Rev 1, there was

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1	substantial improvement in the database, which
2	gave us a good degree of comfort that yes, the
3	problem that we had originally seems to be
4	largely resolved because so much more data
5	came in.
6	Now, I would be the first to say
7	though, that to say that that data now is
8	sufficient to build a robust coworker model
9	that would meet all the criteria for
10	sufficient accuracy as required by Part 83,
11	that's a richer question.
12	And I think that maybe we put that
13	in the parking lot for now. Let's go through
14	the rest of these and then
15	MR. THURBER: Well, is it
16	appropriate then that we alter our response
17	here to this finding based on what Hans is
18	saying here?
19	DR. MAURO: Not now. I would say
20	let's SEC reviews, as Ted pointed out, have

evolved in a way that they take a form and a

level of detail that is on a case-by-case

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1	basis. In other words, we used to SEC reviews
2	according to a very comprehensive procedure,
3	there was a ton of stuff we did, and it was
4	expensive and time-consuming.
5	And it became clear that a new
6	culture developed this must be about two
7	years ago where once the issues are
8	discussed by the Work Group, a judgment is
9	made by the Work Group which issues they would
10	like us to look at a little more closely, so
11	what I am saying is right now I can't answer
12	your question.
13	Really, we are going to look to
14	the Work Group to direct us on whether or not
15	there is more they would like us to do or not
16	in light of what we are hearing from
17	CHAIRMAN ANDERSON: Did the SEC
18	petition come in before or after Rev 1?
19	MR. ALLEN: Before. The Rev 1 was a
20	response to information we found when we were
21	putting together that

ANDERSON:

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Okay,

1	just, I mean the petitioner didn't know you
2	had this data at the time?
3	MR. ALLEN: I am not positive we
4	had it when the petition came in.
5	CHAIRMAN ANDERSON: Well, yes
6	MR. ALLEN: We had it when the
7	CHAIRMAN ANDERSON: Whatever, I
8	mean, that is just a matter of their, they
9	didn't look at it and say we looked at this
10	data and they got problems with it, it was
11	simply there isn't the data and then you found
12	it.
13	MR. ALLEN: Right.
14	CHAIRMAN ANDERSON: Okay. That's
15	helpful.
16	DR. BEHLING: I would just like to
17	perhaps make one comment. As I have mentioned,
18	there was significant revisions to table D-2
19	that needs to be used for the reconstruction
20	of external doses. In the initial table D-2 it
21	was really for the entire period because it
22	was based on 1960 NRC survey report.

The Revision 1 incorporates new data and allows for assigned doses for every single year during this period and in addition it is segregated based on whether or not the person was an operator, a supervisor or other.

The recommended values are in fact geometric mean values and if I have to say one thing, it's that the data is pretty much comprehensive involving those people who were most likely exposed who were in fact part of that database and those data are to be used for dose reconstruction.

We have only a small minority of people who may not have been monitored or for whom the data is not available. Perhaps the values in table D-2 are adequate. But if it turns out there was a significant number of people who may have been operators who were in fact not monitored or whose exposures are not available, then perhaps the geometric mean, again, may indicate a recurring issue with SC&A and trying to come to grips with the fact

that a geometric mean may not always be claimant favorable if the person was not monitored but he was in a high exposure group.

And as I had mentioned I am looking at 1966 in table D-2 for operators the assigned dose is 382 millirem and as I had mentioned a few minutes ago, there were people whose 1966 exposures were upwards of six rem.

Now if among operators who may not have been monitored, the assignment of the default value would truly perhaps not be claimant favorable and so the assumption here is that if we can identify an operator for whom the data is not available, perhaps the geometric mean will not be claimant favorable and we should look at the 95th percentile value.

But that is something that only a careful review would indicate, whether or not people in the high dose area, such as operators, are people whose exposures we do not have for dose reconstruction, perhaps the

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L	geometric	mean	may	not	be	appropriate.

So this would be obviously something that John focused on and that is perhaps we need to look at data again a little more carefully that are now available in the Site Research Database to be sure that the doses that we do have access to are on cases of the most likely exposed individuals.

MR. KATZ: Finding number three, Hans?

DR. BEHLING: Okay, finding number three, potential exposures to neutrons are currently not addressed in Appendix D.

And when I looked at, in fact, at 1960 AEC compliance inspection report, there series of references about was а issues and forth, criticality so and clearly was an indication that the quantities of UF6 at the facility would have given rise to neutron exposures which were not addressed in Rev 0.

So in Rev 1, NIOSH elected to

1	include the approach and I looked at the model
2	and it does appear that it is claimant
3	favorable in its basic assumptions.
4	So the issue of neutron assignment
5	is something that can be resolved with the use
6	of this model.
7	DR. MAURO: Hans can you describe a
8	little bit about how the neutron doses are
9	assigned?
10	DR. BEHLING: Yes, let's see, it's
11	a model, it's let's see which assumptions
12	were used. They used a highly-enriched
13	uranium, 93.1 percent enrichment, and I
14	need to look at all these documents here that
15	are in front of me, but it also, I don't have
16	at my fingertips the quantities that were
17	used.
18	But I think in all the, our model
19	appears to be fairly
20	DR. MAURO: So it is not based on
21	NTA film. It's actually based on
22	DR. BEHLING: No, no. No, no. It's

1	a modeled approach.
2	DR. MAURO: Okay. I got it, so
3	basically a source of neutron enrich
4	neutron, ran appropriate codes, predicting the
5	neutron flux and energy distribution
6	DR. BEHLING: Yes. They used 93
7	percent enriched uranium. They had the workers
8	there at one foot for 1,000 hours per year and
9	what is the quantity but it appears that
10	a 50 kilogram quantity so given that all
11	of the variables that they could have employed
12	in coming up with a model dose, both in
13	quantity, the time, the enrichment factors, it
14	appears to be that the neutron exposures are
15	fairly considerable.
16	CHAIRMAN ANDERSON: Finding four.
17	MR. KATZ: So, no action on that.
18	CHAIRMAN ANDERSON: Well, that just
19	sounds like it is
20	MR. KATZ: That's one that is
21	CHAIRMAN ANDERSON: good to go.
22	MR. KATZ: Closed.

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CHAIRMAN ANDERSON: Closed.

BEHLING: Yes. Finding four. DR. Inhalation intakes recommended by NIOSH may not correlate with empirical urinalysis data. Again, when I looked at all of the secondary reports in the Site Research Database, and I looked at it actually on behalf of Rev 1, I focused on a number of people and I looked at people who were exposed to fairly high air concentrations and for whom we also urinalysis data and subsequently chest count data and sorted through the assigned doses from the coworker model that were initially identified in the Appendix D, table D-1 of Appendix D, are they claimant favorable.

And I can't, I found it too much of an effort to go through all the iterations that I went through. But I tracked several people who were probably outliers in this whole distribution. They were the people who were exposed during a very critical time period when it was realized that there were

high air concentrations and there were changes in processing, et cetera.

And what I ended up concluding was that at least in behalf of those individuals, the assigned default value from table D-1 was probably not necessarily claimant favorable. In fact in some cases they were off by factors of 15 to even greater values, depending on whether you, which solubility you assume.

Now, in the original Rev 0, the table D-1 really made no reference as to who these values should be applied, in other words they were to be used regardless of whether or not that person had bioassay data or not.

And so I realized that in case of people who were clearly monitored, and for whom chest count data, urine data and air sampling data was available, that these values should be used.

So even though table D-1 between Rev 0 and Rev 1 remains the same, a major change was that in Rev 1, the option was to

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obviously default to real data rather than the coworker model, which is really what we are dealing with.

There are a substantial number of bioassay data available for individuals who were likely to have been operators and for whom the exposure was maximal and of course, one should choose that data as the highest exposure to the target organ.

And that was, that stipulation was incorporated into the Revision 1. So as a the table, default table minimum, a coworker internal exposures is no longer just a generic value that applies to all people, but it only applies to those people for whom the data either doesn't exist or it is insufficient for dose reconstruction.

For anyone who has read the original audit of Rev 1, you will see the effort I went to in trying to needs identify for outlier people, people whose exposures were clearly going to be high based on air

sampling data, urinalysis data and for chest count data, the default values in table D-1 would not necessarily be claimant favorable.

DR. MAURO: So Hans, with regard to the new paradigm that's been adopted, what I am hearing is there is this internal dosimetry data for individual workers. You feel that the workers that do have data, whether it's chest count or bioassay, that those workers, it's fairly clear that those are the workers that had the highest potential for exposure.

DR. BEHLING: Yes.

DR. MAURO: Okay, that's important, because what that means is it puts NIOSH in the position to know what the high-end exposures are. So if there is a worker out there that does not have any bioassay data and needs to be assigned some exposure, NIOSH is in a position to assign an exposure that is at the upper end if they feel that's appropriate.

You see, usually you run into an SEC issue when you have bioassay data but you

don't really know whether you have captured the high-end individuals or not. And all of a sudden you have a coworker model that you are really not sure -- we have seen that before -- whether or not you have got the high-end individuals.

But in this -- and Hans has been quite frankly one of the strongest critics of other SECs when he felt that you have got a lot of data but it is not apparent that you caught the high-end people.

You are saying in this case however you feel pretty confident that the data that is available, does capture those people with the jobs that would place them at the high end.

DR. BEHLING: Yes, I believe they realize that during certain periods of operation, that they had a very, very high airborne level and they did monitor at work locations, particularly the work locations, and assigned specific airborne levels to

individual workers.

And then they had the option of also reviewing their urine data and in some cases chest data, and of course some of these individuals were tracked for a period of years because of the very, very high-end exposures that they received.

So it's reasonable to conclude that the highest exposed people were in fact identified. Now I have identified, in addition to the original issue, that says please use the real data when available because there were in some instances there was an assigned value from the default table.

But one of the things that I also have to say that I didn't really include was the likelihood that the uranium to which some of these people may have been exposed to, may qualify for solubility Super S.

And I say this because I looked at some of the Site Research Database documents that actually had hand-written notes and

extrapolated the actual chest burden as a function of time.

And there were some people who, in addition to having air monitoring data, in addition to having urine data, were chest counted at Y-12 over a period of time and several people were counted for a couple of years thereafter.

And I looked at the individual data and by simply looking at a couple of data points that were separated by six months or so, you come to the conclusion that the effective half-life of uranium in the chest for a couple of these people in one instance exceeded three-hundred-and-twenty-some days. And I believe that would qualify for perhaps a Super S status for uranium.

DR. MAURO: Is the ER, the Evaluation Report and the Site Profile Rev 1 silent on this matter or is this --

DR. BEHLING: Yes, they are. They give options for assigning either solubility

1	type S or M and again, as always, NIOSE
2	usually says whichever yields the higher dose.
3	So clearly if it's an exposure
4	that involves, or target tissue involving a
5	lung you would obviously go for type S. For
6	others it might be type M. But Super S is not
7	identified as a potential.
8	Now I don't know exactly where the
9	dividing line between S and Super S comes in
10	at, but when you have an effective half-life
11	in the chest based on chest counting data from
12	Y-12 that exceeds 300 days, I believe that
13	would qualify for Super S.
14	I am waiting for comments from
15	NIOSH.
16	DR. NETON: No. No, I don't think
17	300 days would qualify that for Super S. I
18	think the long-term compartment, even in the
19	type Y was 500 days. So I don't know why 300
20	days would qualify that as Super S.

I mean we will certainly take a look at what you are pointing to but my first

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blush is I don't think so.

MR. ALLEN: If the absorption half-life for type S is 7,000 days half-life, that's only the absorption comment if the physical clearance is going to probably guide that down guite a bit.

But as I recall type M falls around 140 day half-life?

DR. NETON: I mean, we can certainly take a look at what the data set you are pointing to, but off the top of my head it doesn't seem like --

DR. MAURO: Is there anything, I guess, the type of operations that took place here, is there anything about those operations that would lead you to, I guess, to get a Super S out of the uranium, for some reason that's got pretty high temperatures. I am not even sure what they were going here, where there might have been -- in other words, you combine process knowledge with data, starts to build a case that no, maybe we don't have it,

or perhaps we do.

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But right now the report, it sounds like it's silent on this issue.

MR. ALLEN: Yes, Ι mean, Ι remember, I might be getting my sites wrong, but I am thinking that the big exposures that they ended up sending people to Y-12 for the whole body count et cetera were from the red room, and if I remember right, the red room was reduction process with green salt, if remember right, which would be а type material.

DR. BEHLING: All I can say is that in some of the communications between the health physicists at UNC and Y-12, they talked about having common problems of a much longer half-life that they would have expected in a lung of 120 days.

DR. NETON: Well yes, that was the old ICRP-2 model, I mean that only allowed for a soluble inside 120 days, it was sort of like a default value. I --

1	MR. ALLEN: That was Class W wasn't
2	it?
3	DR. NETON: Yes, W. I think, you
4	know, UO2 would be the most insoluble form of
5	uranium, which is type S right now. I am not
6	aware of anything out there right now that
7	would qualify for Super S, although I never
8	say never.
9	We will certainly be happy to take
10	a look at the
11	DR. BEHLING: I was looking at some
12	of the ICRP documents and when they talk about
13	it, they are very diffuse, and certainly when
14	you look at the older documents that are still
15	classified saying there's daily, weekly,
16	yearly, what does clearance mean? Certainly in
17	this case, and I counted it for one particular
18	individual and the effective half-life in the
19	lung was 327 days.
20	Well, that's only one half-life.
21	Okay, so it's less than one year, but what
22	does clearance mean? Is it just one half-life

removed or multiple half-lives where you reduce it down to some nominal level?

DR. NETON: Well, it depends on where you are in the clearance curve. I mean, there's all kinds of compartments with the ICRP 66 lung model as you know and there is no such thing as one half-life in the lung, there's a --

DR. BEHLING: No, I realize that, but Jim, I took, they were, I have data here for a bunch of individuals who were exposed and they were assessed in August of 63, September of 63 and March of 64 and I ignored the first one and I looked at September 30, which is about six weeks thereafter the initial exposure, minimum, and then they were reassessed on March 16, 1964.

So we have at least about 100, well actually 177 days during this exposure where you have the initial clearance from the lung, obviously, and using those two data points, I set the effective half-life in the

1	lung and for one individual was 262 days, for
2	another 327 days.
3	And I thought of not pushing the
4	issue of Super S, but I am raising the
5	question, where does the separation between S
6	and Super S come in? I mean
7	DR. NETON: It's a lot longer than
8	300 days. We can look at it. Again, I, you
9	know, I would like to look at the case but my
10	first thought, it doesn't seem to be an issue
11	but
12	DR. MAURO: One more question
13	before we move on to the next item. Hans,
14	everything is uranium here. Any reason to
15	believe there are other radionuclides?
16	DR. BEHLING: I didn't quite hear
17	you John.
18	DR. MAURO: We have been talking
19	uranium. Is there any reason, like I said, I
20	am not familiar with the processes. Is it
21	possible there's other radionuclides, thorium?
22	DR. BEHLING: Yes, in fact, one of

the things that was added in Rev 1 that one was the potential exposure to thorium, which was missing in Rev 0, and there was a single year in the process time period during which thorium was used, and so NIOSH added exposure to thorium. That is another change in Rev 1.

And when I initially reviewed Rev 0 I was not aware that thorium was even an issue until it was independently raised by NIOSH in their Rev 1.

MR. THURBER: Next.

CHAIRMAN ANDERSON: Fine.

MR. KATZ: Finding number five Hans?

DR. BEHLING: Okay, finding number five was doing -- and oh, one other thing that was changed. Initially the operational period was only extended to 1969 and apparently NIOSH reviewed some of the documents and realized that that date had to be extended to 1973. So that's another independent change that occurred from Rev O to Rev 1, an additional

four years during which operational exposures would have occurred and before the post-operational exposure would have occurred.

So finding five starts with the issue of information regarding the inhalation intakes from residual contamination and NIOSH had stated that their default value is 10.34 dpm per day with type S uranium.

MR. KATZ: Hans, could you just speak up a little bit more. You are fading a little bit.

DR. BEHLING: Okay. In Rev 1, and it remains unchanged, so it appears in Rev 1 and 0, the assumed default value, 10.34 dpm per day for uranium is a default value for inhalational internal exposure and that starts in 1974.

And I looked at the actual guidance in the document and I can't really understand how that came to be and I am not sure I will necessarily go through the details as to -- but for those who have my report, you

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can see what I did and I am hoping that if I am wrong, NIOSH can correct me.

But based on my assessment, the use of the maximum dose from table D-2 and the basic assumptions that were to be used for modeling the exposure, my value turns out to be 434 dpm per work day or 297 dpm per calendar day. That's 29 times higher than the value recommended by NIOSH.

And I had applied a certain modeling approach to that value that I believe interprets their recommendation but I certainly could not come up with their value and I think I am going to ask NIOSH to tell me what method they used to come up with their 10.34 dpm per day and why my number is perhaps not correct.

might MR. ALLEN: We have difference of opinion as far as how you got your number, but as far as what is in Appendix, verify Ι can there was а calculational there. The Appendix error

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going to need to be revised to change that value.

DR. BEHLING: It is basically a very simple calculation.

MR. ALLEN: Yes.

DR. BEHLING: You essentially define the air activity, the maximum air activity, and you need to apply a deposition velocity in terms of meters per second and you end up with a ground activity in terms of dpm per meter square and then I simply applied a resuspension factor that is defined by NIOSH and this is how I came up with my number that in fact, it's a very low resuspension factor, one to the minus six per meter, which we have questioned as part of OTIB-0077.

And so even using that unconservative number, at least my estimation is that that is an unconservative number, I come with a value that is 29 times higher and I think NIOSH should look at their value and look at my value and see which number is

1	correct.
2	MR. ALLEN: Agreed.
3	CHAIRMAN ANDERSON: Yes, agreed.
4	Okay. So there is a task. Yes. Six.
5	DR. BEHLING: Finding six. Again,
6	here this was based on an external
7	contamination residual, external contamination
8	dose rate and I looked at the default values
9	that are identified by NIOSH and I, their
10	default values were 11.6 millirem per year
11	whole body and 186 millirem for skin dose.
12	And again using the limited
13	guidance that was identified in Rev 1 as well
14	as Rev 0
15	MR. KATZ: Hans, your voice is
16	fading again.
17	DR. BEHLING: Okay.
18	DR. MAURO: That's good. That's
19	perfect. Okay.
20	DR. BEHLING: Am I coming through
21	now?
22	MR. KATZ: Very clear.

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DR. BEHLING: Ι looked at the default values for external exposure cited by NIOSH in both Rev 0 and Rev 1 and used their recommended approach in terms of how that was defined. And although it was somewhat brief, I federal guidance report used 12, the **EPA** federal guidance report 12 and used the data that they had recommended to use for external and they actually exposure came up numbers that were considerably lower than the values.

They had for external, as I said, for external they had identified 11 millirem per year, external whole body, and I ended up looking derived value of 2.8, at а approximately five-fold lower, or four-fold lower, and the 186 millirem for skin dose that NIOSH had projected, I ended up with only 10.2 millirem, and that using EPA federal was 12 principle quidance report as my conversion values for residual contamination.

Again, I don't know what

1	methodology they used in order to be able to
2	track their numbers, but I independently used
3	at least the limited guidance they provided
4	and came up with these values, which are
5	considerably lower than the default values
6	recommended by NIOSH.
7	MR. ALLEN: Yes, we used the
8	conversion factor we actually talked about
9	earlier today in TBD-6001 for external dose
10	from surface contamination and it was derived
11	from a MicroShield run. Didn't realize it was
12	that much higher than federal reg guide 12.
13	But I am not sure, Hans, I don't
14	think you included the thorium-234 and the
15	protactinium?
16	DR. BEHLING: No I did not, since

DR. BEHLING: No I did not, since the values originally were identical in Rev 0, where thorium was not identified, I have to conclude that the values, the default values in Rev 0 and Rev 1 did not address thorium.

MR. ALLEN: Yes, well, I am talking about the short-lived decay products of U-238.

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1	You can get some reasonable beta and gamma
2	from the protactinium-234m and the thorium-
3	234.
4	I think to use federal reg guide
5	12 they would have to include those two as
6	well.
7	DR. BEHLING: Okay, I have to look
8	to be sure that they weren't really
9	incorporated. You may be correct, which may
10	explain why my numbers were considerably
11	lower. In any case, especially the external,
12	penetrating dose of 11 versus three, we are
13	not talking about the doses, even though there
14	is a significant difference between two
15	values, in absolute terms they are still
16	nominal doses.
17	DR. MAURO: David, how did you
18	derive the build-up of uranium on surfaces for
19	the residual period?
20	MR. ALLEN: It was the same
21	DR. MAURO: Same thing?
22	MR. ALLEN: Technique that we

1	DR. MAURO: Using the airborne
2	activity in the deposition and for a year?
3	MR. ALLEN: In this one, actually
4	we used 2,000 hours. We used a work year.
5	DR. MAURO: One work year. It's
6	above the number that would be 27 days?
7	MR. ALLEN: Yes, it's above that.
8	DR. MAURO: Okay. All right. So I
9	guess the only issue we have, not so much with
10	the external, whatever dose it is, is this
11	resuspension factor. But again that is not
12	necessarily an SEC issue. We do have a problem
13	with the 10 to the minus six per meter number,
14	it's a longstanding issue.
15	MR. ALLEN: That's finding five
16	anyway. That's
17	DR. MAURO: Right. And, but that's
18	not an SEC issue.
19	MR. KATZ: So is there something
20	remaining on the table on this?
21	DR. BEHLING: Yes, there's only an
22	observation and

1	MR. KATZ: I mean for finding
2	number six. Do we have anything, is there
3	anything to do with finding six or is it
4	something that could be closed?
5	DR. BEHLING: Well, I can go back
6	and look at whether or not federal guidance
7	report 12 incorporates some of the short-lived
8	daughter products.
9	And whether or not that accounts
10	for the differences between my calculated
11	value and
12	MR. ALLEN: Essentially, we are
13	using two different models to model this and
14	coming up within a factor of four and it might
15	be closer by the time we reconcile some of
16	this. I don't know if you want to explore it
17	further or not.
18	DR. MAURO: Yes, federal guidance
19	report 12 only includes the uranium you have
20	got, so if you look at U-238, you get U-238.
21	You have got to include thorium-234, and
22	protactinium-234m so, and I think there is a

on the surface.

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1	one percent photon on that order from the
2	protactinium, which may be the major
3	contributor plus of course there's the
4	Bremsstrahlung issue.
5	But you are not going to get very
6	Bremsstrahlung here, I would imagine. You are
7	talking about just a little residual activity

MR. ALLEN: You have a high-energy beta but not so much --

MAURO: But nothing to knock DR. into, yes.

BEHLING: And John, now that DR. Jim Neton mentioned the issue of short-lived daughters, as I have said, I took only U-234 and 235 into consideration and you know, I I did not address the contribution, said, especially the beta component and that is very likely the reason why I ended up getting the much lower dose.

MR. KATZ: Okay. So do we need a report out on this?

1	DR. MAURO: Yes, let's fix that.
2	Yes, we should, you know what we can do? In
3	our matrix, you know I guess we will go back,
4	revisit that number and see, if we went back
5	in light of this I guess it will be a note
6	here, you know, in the matrix saying that SC&A
7	will revisit the exposure and that will be
8	what we put in and of course, I guess we will
9	put a memo out subsequent to that, yes we did
10	and you are right and whatever the answer is.
11	DR. BEHLING: Yes, I am writing
12	myself a note here. I will do that.
13	CHAIRMAN ANDERSON: And then the
14	observation?
15	DR. BEHLING: The observation that
16	I mentioned was really a generic one and it
17	was
18	DR. MAURO: Could you get closer to
19	the phone again Hans, you are fading away.
20	DR. BEHLING: Okay, I don't know
21	why, this mic usually works very well, I don't

know why it is not working. In my observation

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one, I stated that UNC site description is insufficient and I think, based on my comments regarding the individual findings, I was really not able to review some of the numbers and I think that was really a reference to how these numbers were derived for certain instances such as the internal and external exposures that we just mentioned where they provide default values without simply necessarily providing new information which would allow you to track those numbers.

So it's just a generic comment. But then again, the TBD-6001 and the appendices are not intended to be equivalent of a Site Profile in terms of the definitive information that is normally incorporated and so I understand why these documents are so much more brief.

But I still felt that perhaps we could have benefitted from additional data that would allow us to actually follow the methodology that was used to identify some of

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1	the default values that NIOSH had incorporated
2	in their tables.
3	CHAIRMAN ANDERSON: Okay.
4	MR. ALLEN: Well, it sounds like
5	you are basically saying we could use some
6	more detail on how we did the calculations in
7	the text of the Appendix.
8	DR. BEHLING: Yes, and what I
9	had problems with initially, and this is what
10	John picked up on, regarding the external

exposures, there were no references 11 12 documents cited various that in were 13 Evaluation Report to the SEC petition that would have allowed me to look at those numbers 14

15 up front.

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And sometimes even the so, bibliography would have perhaps benefitted from some of those particular SRDB references that were cited in the ER and include those into the actual Appendix D.

CHAIRMAN ANDERSON: Anything else?

MR. There KATZ: only was

1	observation
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DR. MAURO: Yes, that's it, we're done.

CHAIRMAN ANDERSON: Anything on SEC? I mean, since this doesn't pertain too much, do we need to, you do have your report on that?

DR. MAURO: Well, this is a little bit unusual. What you have here is review of Rev 1 of the Site Profile with some consideration, limited consideration of is there anything in the ER that sort of raises a red flag.

And what we are hearing is, based on, you know, a limited review of the ER, there's nothing there that really jumps out at you that looks like there's something that might be serious, a serious issue that could trigger an SEC concern.

It's unusual because we usually don't come to that conclusion that quickly, as you know. And you know, and if you feel as if

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there's more to be done in terms of taking a closer look at the external data, downloading it, if you feel that, I mean --

DR. BEHLING: As I said, John, biggest concern that I might have is assignment of default values to unmonitored people if they turn out to be high-end exposed individuals, operators. As I said, you know, I looked at 1966 and the geometric mean is 380 some millirem and yet for 1966 among a select group of people, exposures as high as six rem are part of the record. And if it turns out that they were operators whose either exposure records are not available, who may have been monitored, but they are not available, or they may not have been monitored in spite of the fact that they were operators and high-end exposed individuals, then I would assume that the geometric mean may not necessarily be a very good number for those individuals.

DR. NETON: Well, but that in itself is not an SEC --

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DR. BEHLING: No, it's not an SEC issue.

CHAIRMAN ANDERSON: NIOSH said they can do dose reconstruction. Do we have any indication where you would be concerned that in fact there are some exposures at this facility that aren't covered here, weren't identified, or you know, this really is, there are some coworker issues maybe, but doesn't seem to be any surrogate data, so this seems to be new data was found in response to the petition and that's, you know, are you comfortable that that in fact does fill sufficient gaps and you know, not -- we don't need to talk about how they go about doing the reconstruction, but can it be done?

Sounds like your conclusion so far is that yes, we ought to as a Work Group go back to the Board and say, you know, we have discussed it, hasn't been an in-depth review, but you know, if NIOSH will summarize more of what the new data is as part of a

presentation, we can recommend -- I don't know, Mark, are you still there?

MR. KATZ: We don't have Mark.

CHAIRMAN ANDERSON: I mean, I would want to query him but my sense is, you know other than -- this one doesn't have as comprehensive an evaluation but it seems to me the go, no-go kind of thing is this is very strong as it goes.

DR. MAURO: I mean, certainly the review that we just heard --

CHAIRMAN ANDERSON: Yes.

DR. MAURO: I mean, I am probing away to see if there is anything about it that gets me to gee, you better look at it. I don't see it. In other words, what Hans is saying regarding the -- for example, you always have a problem with neutron exposures, but what was just described to me, you say, okay where could there be a -- bottom line is SECs are triggered because of inadequate bioassay data and inability to reconstruct neutron doses.

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All right, so what did we just heard regarding hear? We neutron exposures that they used a very large, highlyenriched uranium source. I am assuming that the quantity that was used was in fact an upper-end value and was -- I don't know that to be certain, but Hans feels comfortable with that, and then went ahead without attenuation, assuming no went ahead calculated what the neutron flux and energy distribution would be and derived the doses.

Without any attenuation, and putting a person close for an extended period of time, certainly places an upper bound. Now I am not going to say now that I, myself, or there are other people who reviewed it, say yes I agree.

Because normally what would happen is that conclusion that you heard, regarding neutron, that would have to make the rounds. In other words, there would be other folks that would independently review it as part of

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our required quality assurance process.

I have doubt, working no with Hans, that this is in fact the case, you know. But are in strange territory. We we talking an SEC issue and I just, I am a little concerned that, given the importance of decision like this, where you would deny an SEC, the, and not have gone through another, doing, checking the other eyes, numbers, convincing themselves that, you know --

I would like to be able to stand in front of the full Board and say Hans did the initial work, these were the issues we looked at that we thought were important and the big ones I can tell you right now is the neutron exposure, probably you would want to make sure that the --

It sounds like what he is saying is that the film badge data did in fact capture the high-end people and the bioassay chest count data did in fact capture the high-end people, and be able to say once you have

that, you are in a position to assign high-end doses as necessary.

And that means all the major potential SEC issues don't -- there aren't any. It just becomes a Site Profile, how you implement that.

So, but I have to say right now, I would, you know, it would be unusual for SC&A to show up at a full Board meeting without having gone through a little bit more due process and -- Hans this is not, it doesn't bear on you, I am not questioning you.

But there is a process we are required to follow and especially when it comes to an SEC, I would not want to give anyone the impression that we, you know, we cavalierly came, you know, we did not follow our procedures.

I would like to, I think we have to have these analyses, have the independent checking process go forward.

DR. BEHLING: And, John, I didn't

want to come across having the last word, clearly, as I said, and I prefaced my comments by stating that it was Ted's recommendation that we only familiarize ourselves with the site, with the SEC petition and the Evaluation Report and so, this was at best, a very, very cursory assessment of the issues that might come into play here for an SEC.

And our findings, at least the ones that I have identified, are probably resolvable at least, but not necessarily the final word in defining whether or not the issue is totally resolved.

And then you mention, obviously, the neutron exposure and maybe we need a second opinion. Is 50 kilograms a bounding value? Is one foot for 1,000 hours exposure a bounding value?

In my estimation, it appears to be, but I think I will defer to a second opinion on that.

MEMBER FIELD: John, I guess, my

1	thought was I guess reflective of what you
2	have expressed a bit ago that you know, this
3	process has been going on for a long time but
4	what you would like is a consistent process
5	throughout.
6	Didn't you say a little while ago
7	that times have changed.
8	DR. MAURO: Well, you know
9	MEMBER FIELD: And there's a new
10	paradigm about how things should be reviewed.
11	So I guess what, where do we go from here
12	forward if there's a new paradigm? DR.
13	MAURO: I will give you an example.
14	MEMBER FIELD: Okay.
15	DR. MAURO: I will give you an
16	example. On the Mound Site that would be a
17	very good way to think about it three or
18	four issues emerged as being important:
19	tritides, neutron exposures and radon, okay?
20	Now, we quickly zeroed in on
21	those, so it's not that we did this big

report, in fact we didn't. We didn't deliver

one of these 200-page reports where we exhaustively look, we did what Hans did, basically, and said, wait a minute, we think that when all is said and done, if there's going to be a place where there's going to be a problem, it's going to be the tritides, the radon or the neutrons, and that's what to zero on in.

Once the Board or the Work Group heard that, they said okay, we have got to go vertical on these. We have got to go do a data capture, we are going to find out everything we can about whether or not you really can reconstruct the doses from those three areas.

And a large effort, a very large effort, ended up going into those three days. So even though we constrained ourselves initially, once we constrained ourselves and then we hit it real hard and we are in the middle of that right now, I mean there's going to be a lot --

Now I am saying that what we just

did is something even lighter than that here. What we really did here, quite frankly, is come to this meeting prepared to give you folks a sense of where we think we are on, you know --

Certainly we have completed our Site Profile review and it turns out we have some issues, as you heard, that need to be looked at, and they all sound like Site Profile issues that can be fixed.

But it did not, you know, we did not do some of the things that we always do. Like for example, the tritide issue at Mound. That issue has been hit by three or four people within our group and checked and rechecked, there's been site visits to find out more about it --

So, listen, I am ready to stop on this one if you are comfortable with it. But I don't know if that is in the best interests of this process. I think that those -- we now know what the three issues are, that probably

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we want to make sure we got this right.

And we have to be able to tell the full Board that we identified these three issues that were in our original analysis, presented our story to the Work Group and in my mind, we have got to check those a little more deeply and confirm that through additional data either capture, perhaps through some interviews, that in fact the approach adopted by NIOSH with regard to these three issues is in fact the rock we can stand on and those doses can be reconstructed with sufficient accuracy.

MEMBER FIELD: Okay. I guess my thinking is that without doing further review of this, if this comes to the Board for further consideration, say even for an SEC vote, that the Board will say possibly, well you didn't do due diligence.

DR. MAURO: And I am afraid of that.

MEMBER FIELD: For the initial

1	review and make a vote based on what was known
2	rather than at that time asking for further
3	review.
4	Well, going back and doing a full
5	review.
6	DR. MAURO: I am not saying a full
7	review. I am saying that right now see,
8	what we did was
9	CHAIRMAN ANDERSON: What can you
10	done between now and Idaho?
11	MR. KATZ: I mean this won't be
12	ready for Idaho anyway. It is not on the
13	agenda for Idaho.
14	CHAIRMAN ANDERSON: Well, we could
15	still put it on the
16	MR. KATZ: I guess it could go on
17	the agenda for Idaho. I mean, I just to
18	throw in my two cents for you to consider, I
19	mean this is different than Mound in that they
20	had concerns about the issues they went deep
21	on with Mound

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DR. MAURO: Right.

1	MR. KATZ: From the get-go.
2	DR. MAURO: Yes, that's true.
3	MR. KATZ: And so here they are
4	saying they don't think they have concerns
5	about the three issues. But I do think that it
6	makes sense for SC&A to do its QA on these
7	three concerns.
8	I don't know whether it
9	necessarily means doing a bunch of
10	interviewing and going out and doing field
11	research, but having a down-the-table review
12	of Hans's work.
13	DR. MAURO: Well, let's go with an
14	example. Let's say we want to convince
15	ourselves that the neutron model that was
16	adopted, is bounding. All right, now, what do
17	you do when you do that?
18	You go in and you see, okay, the
19	source. What was the source? And go into the
20	literature, go into your site query database,
21	and look at the history and see if there is
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enough information in there that leaves you

with a warm and fuzzy feeling that yes, whatever the number was that was assumed, and the enrichment level, quite frankly, 50 kilograms of 96 percent enriched uranium? I can't even imagine you could have that in the same place. That's --

MR. ALLEN: We used 100 kilograms for the 20 percent and cut it down to 50 because this is just --

DR. MAURO: So I mean, we are starting to solve -- see, I would like to be able to say yes, that particular scenario is certainly bounding because you would never have that, quite frankly you are saying that is bounding because you have a criticality situation which is not going to -- you didn't have it.

And then you say, okay, good. The source is that hotbed. Next. How close are they to it? And for how long? And also, I would say the dosimetry itself. Neutron dosimetry is not straightforward.

1	You know, make sure that the
2	energy flux, it's basically neutron energy
3	distribution at the receptor location and
4	DR. NETON: See John, I think those
5	are Site Profile issues that you are
6	describing. Those are refinements. I think
7	DR. MAURO: Yes.
8	DR. NETON: an SEC review would be
9	is the model appropriate? I mean, can the
10	model appropriately be
11	DR. MAURO: Can you bound it?
12	DR. NETON: Bound those things, and
13	do you now the source-term is sufficient. You
14	know that, that is my opinion, I mean,
15	otherwise you are doing refinements of a
16	calculation that you already actually agree is
17	okay, you know?
18	DR. MAURO: I mean, yes.
19	CHAIRMAN ANDERSON: I guess I would
20	say let's
21	DR. MAURO: Maybe you are right.
22	CHAIRMAN ANDERSON: Go forward with

1	doing I guess I just don't want to we
2	have got so many things
3	DR. MAURO: Yes.
4	CHAIRMAN ANDERSON: On the table.
5	We have got Hooker coming up and I don't want
6	to, you know, if there is one that we can move
7	forward on pretty quickly, I would just as
8	soon try to do that, but I do want to be
9	confident that we have done enough and I don't
10	want to, as a first-time Chair
11	MR. KATZ: I don't think you want
12	to
13	(Simultaneous speaking.)
14	DR. MAURO: It's just that this is
15	a little bit unusual, that's all. We haven't
16	come, we haven't achieved closure on an SEC
17	issue in two hours in a long time.
18	CHAIRMAN ANDERSON: Well, I mean
19	MR. ALLEN: The bottom line is you
20	are the one that has to give some kind of
21	report to the Board and what you feel
22	comfortable with and what you want us to do in

order to --

CHAIRMAN ANDERSON: Given what they will ask you and you will go through this, I would say we have got to move -- you need to do more.

DR. MAURO: And I think that, what I could do is work with Hans, lay out something that would represent okay, we are going to do the following, put that in, send them an email out, I mean there's an action item on us, of exactly what it is we think we need to do --

CHAIRMAN ANDERSON: Yes. Okay that's good.

DR. MAURO: and a time period. And I can tell you now, from what I heard, you know, it's really a matter of touching the bases that need to be touched so that we can stand up and look everyone in the eye and say, listen we checked these numbers. Here's the -- and you know, and more than one person look at it.

1	We are required to do that, quite
2	frankly, so we have got to go through this QA
3	process. I don't think it's going to be big. I
4	don't think we are talking about months of
5	work. We are talking a month, you know
6	CHAIRMAN ANDERSON: Let's do that.
7	MR. KATZ: Let's do that.
8	CHAIRMAN ANDERSON: And if you can
9	kind of give what your proposal is.
10	MR. KATZ: He will do that in a
11	memo.
12	CHAIRMAN ANDERSON: And then before
13	Idaho, an update on where we are, I can give
14	people some sense
15	DR. MAURO: It will be my action
16	item. I will put out a memo, which will be,
17	have a list of five action items that I wrote
18	down, and I will put it out.
19	But in this particular one, I will
20	also give you an indication of when we will be
21	able to deliver a report.

CHAIRMAN ANDERSON: Good. Don't we

1	usually deal with, denial things only at face-
2	to-face, not on the phone?
3	DR. NETON: Pretty much.
4	MR. KATZ: Yes.
5	CHAIRMAN ANDERSON: So we are
6	really looking at
7	MR. KATZ: So we are really looking
8	at November.
9	CHAIRMAN ANDERSON: Yes.
10	MR. KATZ: Because it is not going
11	to get done in I mean we really have only
12	about, I don't know how quickly you will do
13	this, but otherwise, we only have four-and-a-
14	half weeks or whatever before the Board
15	meeting and I don't think that is going to
16	happen.
17	So we are talking about November.
18	We are talking about November here, which is
19	fine.
20	CHAIRMAN ANDERSON: Yes. That's
21	good. Okay.
22	MR. KATZ: I guess if it does end

1	up going to a dominal of source up and in
1	up going to a denial, of course, no one is
2	hurt by the delay, I mean
3	CHAIRMAN ANDERSON: Right. Yes.
4	Yes.
5	DR. MAURO: The process, quite
6	frankly, let's say we were to put this on a
7	fast track, the process would be a month of
8	SC&A work and then it would have to go to DOE.
9	That's usually two weeks or so, to get
10	approval.
11	So even if we want, you know, full
12	press, it's six weeks.
13	MR. KATZ: We are looking at
14	November, but just an indication from you
15	that, a time line.
16	DR. MAURO: Yes, what that looks
17	like.
18	MR. KATZ: What you will do and
19	when.
20	CHAIRMAN ANDERSON: Do we have any
21	other issues? Now that I have got 20 minutes
22	to make my flight, I am going to push you

1	harder but I will I will have a nice, quiet
2	dinner tonight.
3	MR. KATZ: Me too.
4	CHAIRMAN ANDERSON: Go at 7 o'clock
5	to the airport, the rest of you who are
6	waiting over, you can go into town, party up
7	and
8	Anything, so, we have pretty well
9	got three different items.
10	MR. KATZ: Yes, oh, the other thing
11	that you want to, that is on the agenda to
12	discuss, is just the status of DCAS and SC&A
13	work on Hooker Electrochemical, to have a
14	sense of where we are going forward.
15	DR. MAURO: Bill, could you give us
16	the low-down on where you are on Hooker?
17	Because you have been working Hooker, right?
18	MR. THURBER: Yes. You know, we are

moving along. It's, we are well along. I think

that the conversation we had this morning

about how Hooker and TBD-6001 interplay is

illustrative of some of the problems we have

19

20

21

1	had in sorting out things.
2	But our report is reasonably far
3	along.
4	MR. KATZ: So, sort of a time
5	frame? Maybe when you reply on the other
6	things, you can give us a because we are
7	going to need to plan another Work Group
8	meeting. The time frames for these kind of
9	relate to when all these deliverables will be
LO	coming in.
L1	DR. MAURO: Once we have finished
L2	the technical analysis of Hooker, which it
L3	sounds like is pretty far along, we have to
L4	do, we are going to have a section dealing
L5	with surrogate data. Hooker makes quite
L6	extensive use of surrogate data. It uses TBD-
L7	6001.
L8	And we are going to have and
L9	there are the Board has a surrogate data
20	criteria document.
21	MR. THURBER: I thought we put that

1	DR. MAURO: For Hooker?
2	MR. THURBER: I thought that the
3	issue of including discussions of surrogate
4	data in all of these appendices was kind of ir
5	abeyance. Wasn't that the guidance from you
6	Ted?
7	DR. MAURO: I thought that was just
8	the opposite.
9	MR. THURBER: No, because you
10	recall that I had originally advised everybody
11	who was working on these appendices that a
12	discussion of surrogate data, and given that
13	the Bethlehem model was appropriate and then I
14	thought that based on a conversation that you
15	all had, that that was not going to be a
16	requirement of these appendices at this time.
17	DR. MAURO: I have to say, just to
18	be
19	MR. KATZ: I have not had any
20	discussion one way or the other on this
21	subject.
22	DR. MAURO: I have been operating

1	on the premise that the most important
2	MR. THURBER: Hans? Are you still
3	there?
4	DR. BEHLING: Yes?
5	MR. THURBER: Can you shed any
6	light on that comment?
7	DR. BEHLING: You know, I was just
8	looking at something else.
9	MR. THURBER: I understand.
LO	DR. MAURO: No, we know where you
L1	were. Hans, we have a number of SEC petition
L2	reviews and Site Profile reviews that deal
L3	with a lot of AWE facilities right now and
L4	Bill is very much involved in that work.
L5	What I had in my direction, which
L6	I believe I gave everyone was, one of the
L7	chapters of this report has got to be the
L8	degree to which the use of surrogate data
L9	meets the five criteria I think it's five -
20	- laid out in the Board's guidelines, just
21	like we did in Texas City, Bethlehem Steel,

Dow and others.

1	You are saying, now I have got to
2	tell you
3	MR. THURBER: Well I have got to go
4	back and review my emails. I don't, I can't do
5	it here because they're on my other computer.
6	DR. MAURO: I mean, to me that's
7	the biggest and most important question.
8	That's what turned Bethlehem Steel, that
9	judgment, you know.
10	MR. THURBER: Well, I know that.
11	DR. MAURO: Yes.
12	MR. THURBER: But I thought that we
13	had a change in marching orders, but I will
14	check my emails when I get home.
15	MR. KATZ: Yes remind me when you
16	find what you find because I don't recall
17	giving any guidelines about that question.
18	CHAIRMAN ANDERSON: For our next
19	meeting, we probably need to do that before,
20	what is it, December?
21	MR. KATZ: The next meeting face-
22	to-face is in November.

1	CHAIRMAN ANDERSON: November. So,
2	we probably, hopefully, will get your review
3	on
4	MR. KATZ: Hooker.
5	CHAIRMAN ANDERSON: Hooker and
6	United whatever.
7	MR. KATZ: Oh, that, well in
8	advance.
9	CHAIRMAN ANDERSON: But I would
10	think at that meeting is where we would want
11	you to, as a Committee we would make that
12	decision so it needs to be far enough advance
13	of the meeting so that you have got time for
14	your agenda.
15	MR. KATZ: Does DCAS have anything
16	ongoing related to Hooker? Is Hooker, are you
17	
18	MR. ALLEN: No.
19	MR. KATZ: Okay.
20	DR. MAURO: You gave a presentation
21	and
22	CHAIRMAN ANDERSON: You've got 6001

1	issues you're going to, and I don't know, will
2	you have a White Paper in that time frame?
3	MR. ALLEN: Before which time,
4	before November?
5	CHAIRMAN ANDERSON: Yes.
6	MR. ALLEN: I would think so, yes.
7	MR. KATZ: But we'll need a Work
8	Group meeting in advance of the Board meeting.
9	CHAIRMAN ANDERSON: Oh yes, that's
10	what I mean.
11	MR. KATZ: Absolutely.
12	CHAIRMAN ANDERSON: And what I was
13	looking at is when. It would be nice it would
14	seem, if we have your White Paper to look at.
15	MR. KATZ: Right.
16	CHAIRMAN ANDERSON: And we will
17	have your little notes.
18	MR. KATZ: So we will get time
19	lines from, both from Dave and from John for
20	all of this work so we will know, and that's
21	the point of the time line, so we can figure
22	out when to have another Work Group meeting.

1	Because so far it is as soon as possible,
2	of course in advance of November, because you
3	never know what comes up with the review
4	that's the work that gets done.
5	But new issues get unearthed or
6	whatever.
7	CHAIRMAN ANDERSON: I just want to,
8	we got a window we can start looking at
9	people's schedules.
10	MR. KATZ: I think until we hear
11	back from them, until we hear back from them
12	though, if we pull out our calendars now we
13	have no sense of when we
14	CHAIRMAN ANDERSON: Yes, no, no,
15	but if we
16	DR. MAURO: When is the, there's
17	the August, and then what's the next meeting?
18	MR. KATZ: After August there will
19	be another teleconference but the next face-
20	to-face is in November, some time before
21	Thanksgiving, maybe a week before Thanksgiving

or so.

1	DR. MAURO: We will be finished
2	with everyin fact our fiscal year will be
3	over by then almost I mean, end of December.
4	MR. KATZ: Well December.
5	DR. MAURO: The end of December, so
6	we are a month away from the end and in
7	principal, for all of our missions, tasks,
8	should be close to being done by the end of
9	December, so I mean we are going to be done
10	with almost everything unless you give us new
11	work by that time.
12	That's way out, yes, we will have
13	it well before that.
14	CHAIRMAN ANDERSON: Okay. Good. Any
15	other questions, issues?
16	MR. KATZ: For the good of the
17	order.
18	CHAIRMAN ANDERSON: For the good of
19	the order. Adjourned.
20	MR. KATZ: Adjourned. Thank you
21	everybody for a lot of hard work today.
22	(Whereupon, the above-entitled

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matter went off the record at 4:24 p.m.)

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