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

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

convenes the

WORKING GROUP MEETING

ADVISORY BOARD ON

RADIATION AND WORKER HEALTH

DAY THREE

The verbatim transcript of the Meeting of the Advisory Board on Radiation and Worker Health held at the NIOSH, Cincinnati, Ohio, on June 2, 2005.

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June 2, 2005

TASK 1, MALLINCKRODT SITE PROFILE REVIEW NIOSH/ORAU ARJUN MAKHIJANI, SC&A HANS BEHLING, SC&A

TRANSCRIPT LEGEND

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In the following transcript (off microphone) refers to microphone malfunction or speaker's neglect to depress "on" button.

PARTICIPANTS

ABRWH MEMBER:
GRIFFON, Mark A.
President
Creative Pollution Solutions, Inc.
Salem, New Hampshire

OTHER ATTENDEES:

DR. HANS BEHLING, SC&A

MS. CINDY BLOOM, ORAU

MR. GREG MACIEVIC, NIOSH

DR. ARJUN MAKHIJANI, SC&A

DR. JIM NETON, NIOSH

STAFF/VENDORS

STEVEN RAY GREEN, Certified Merit Court Reporter

1	PROCEEDINGS
2	(9:00 a.m.)
3	DR. NETON: Okay. We're ready to start. Oh, let me
4	bring Mark Griffon I just hung up on Mark. Hang
5	on.
6	(Pause)
7	DR. NETON: Okay. We're ready to go then. Mark
8	Griffon is is participating via via telephone
9	connection and today we've got Cindy Bloom and Hans,
10	Arjun, myself, and and Greg Macievic. Where we
11	left off yesterday was we were now going to get into
12	the external dosimetry questions that SC&A had, so I
13	guess I'll turn it over to Arjun and he can start
14	the questioning.
15	DR. MAKHIJANI: Yeah. Hans, should I should I
16	just go through the questions
17	DR. BEHLING: Yeah, yeah.
18	DR. MAKHIJANI: and then maybe you take over the
19	discussion
20	DR. BEHLING: Well, we'll we'll yeah, I have
21	some questions here. I haven't even looked at the
22	questions you submitted but
23	DR. MAKHIJANI: Okay. Should I run through my
24	questions first
25	DR. BEHLING: Yeah.

1 DR. MAKHIJANI: -- and then you can run through 2 yours? Should we do it that way --3 DR. BEHLING: Yeah. 4 DR. MAKHIJANI: -- since they -- they have it in 5 writing already? Okay. I'll just read the question so it's in the record. 6 7 What proportion of employees have no external 8 monitoring data? 9 DR. NETON: Right. This is -- this is similar to 10 what we discussed yesterday for the internal. 11 don't have the answer at this point but we'll 12 certainly try to get that fleshed out soon, 13 hopefully well advance of the board meeting. DR. BEHLING: Is there any indication to whether or 14 not people were cohort badged or badged on the basis 15 16 of likely to be exposed? And there's a difference 17 obviously. 18 DR. NETON: It's -- it's -- exactly. It's our --19 the indications that I have from looking at the 20 files are that people were individually badged. And 21 in fact I think there's a memo we can point to in '49 that indicated that the badge was actually part 22 23 -- that the film badge was part of the security 24 credentials. 25 DR. BEHLING: Okay.

1	DR. NETON: And I guess I we do know a little
2	more than what I just indicated about percentages
3	that were monitored. If one looks at the SEC
4	evaluation report, there is a table on page 17 I
5	don't have copies to distribute that indicates by
6	year the approximate total number of employees at
7	the site during that year, and this is based on the
8	the epi study that was done by the Center for
9	Epidemiological Research, which in traditional
10	epi style they they only looked at white males.
11	But we believe that's it's a good indication of the
12	magnitude of the workforce at that time.
13	And then we have a listing of the number of
14	employees who were monitored during that year. And
15	for external monitoring after '49, the numbers
16	appear to be well over the majority. In 1949 we
17	have 506 employees monitored out of an estimated
18	workforce of 676 and it appears that the percentages
19	go up from there. And in the last year, you have
20	virtually well, 90-plus percent of the workers
21	being monitored, so a very large large percentage
22	of the workers had film badges
23	DR. BEHLING: So you
24	DR. NETON: after 1948.

DR. BEHLING: Yeah, the majority of people were

1 likely to have some radiation dosimetry records. 2 Now --3 DR. NETON: Now -- now the other question which is 4 more relevant, do we have the --5 DR. BEHLING: The records. That's true. DR. NETON: -- records. Now I have looked through a 6 7 sampling of the records and it seems to be that the 8 majority of the people have some records, but I 9 don't know if that tracks with this table. Now I'm 10 assuming -- and we need to do a little homework here 11 -- that this information came from somewhere and 12 presumably it has to do with the records that we 13 have. But again, I have not, you know, followed 14 this thread all the way through. But a very, very 15 large percentage of the workers have monitors --16 were monitored in those time -- in that time period. 17 DR. MAKHIJANI: Now, Jim, when you say 90 percent do 18 you mean 90 percent of all the workers in the AEC 19 area or 90 percent of the production workers? 20 DR. NETON: Oh, I would assume it's AEC area. Ι 21 mean is that -- Mallinckrodt of course was a 22 chemical factory and I -- there'd be no reason to monitor workers who were working with chemicals. 23 24 DR. MAKHIJANI: I misstated my question. I meant 90 25 percent of the production workers in the AEC area or

1 90 percent of all workers in the AEC area? 2 DR. NETON: This -- well, I would say this is 90 3 percent of all white male workers in the AEC area, 4 not just production workers. Those other ten 5 percent then would presumably be people such as 6 clerical folks and --7 DR. MAKHIJANI: Yeah. 8 DR. NETON: -- and such. 9 DR. MAKHIJANI: Of course a lot of the clerical 10 folks were women. 11 DR. NETON: Right. 12 DR. MAKHIJANI: So that would not fall in this 90 13 percent. 14 That's correct. In fact, those women DR. NETON: 15 would not be represented in this total number of 16 employees value because, again, this was a -- this 17 was taken from an epi study and they typically only 18 -- for statistics purposes pick all white male 19 workers at the facility to get the large bulk of the 20 population. But, you know, we will -- we will go 21 through and look at the individual cases that we 22 have, because as I indicated before, I believe we 23 only have about a 130 or so cases that -- that 24 initiated employment between 1949 and 1957 currently

in our possession.

1 DR. MAKHIJANI: Yeah. In -- in the context of this 2 men/women thing, I -- I interviewed a -- a woman 3 employee in whose records I found two urinalysis 4 samples. And she was a clerical employee, so I kind 5 of found that a little odd because I did not find --6 they were low, in the two micrograms per liter 7 range, so they --8 DR. NETON: (Unintelligible) limit of detection --9 DR. MAKHIJANI: -- not within the detection limits, 10 so one doesn't know what to make of that. But I 11 wondered why there would be urinalysis samples in a 12 clerical worker's --13 MS. BLOOM: We found at some sites that 14 stenographers frequently followed people into areas 15 to take dictation while somebody was doing a report. 16 That was in the records at Windy* we saw that. 17 DR. NETON: And I -- I don't --18 MS. BLOOM: They also might have taken them as 19 blanks if they wanted to see, you know, what -- what 20 are other people looking like. It's hard to say. 21 DR. NETON: And also, I don't know what time frame 22 this was or what, but I know for instance at the 23 Fernald site not all workers were monitored, but 24 everyone when they started a new hire, 25 (unintelligible) that's part of the physical, have a

1	urine sample. And actually every year part of the
2	annual physical was a urine sample, whether you
3	worked in the plant or not.
4	MS. BLOOM: And that could have been true, the
5	initial and termination samples there.
6	DR. NETON: I I don't know. We do have to look
7	at the specific case
8	DR. MAKHIJANI: Yeah.
9	MS. BLOOM: (Unintelligible)
10	DR. NETON: and if you've got a number I'd be
11	interested in
12	DR. MAKHIJANI: I'll try to bring it up later. I
13	think I have it in my notes. Maybe not. The I
14	looked at the medical records sections in some of
15	those large DOE files, and there wasn't a column for
16	so they have routine medical type of urinalysis,
17	whatever they do in the medical side
18	DR. NETON: Right.
19	DR. MAKHIJANI: but there wasn't a column in the
20	form for uranium.
21	DR. NETON: That that would not normally show up
22	in the medical form itself.
23	DR. MAKHIJANI: It would not show up
24	DR. NETON: It would be like I know at Fernald, I
25	don't know that this holds true at Weldon Springs,

1 but -- or Mallinckrodt, but they would just split 2 the sample and half would go down to the bioassay 3 laboratory and, you know, we would analyze it and 4 then keep it in the dosimetry record file. 5 early on, though, some of the medical files had 6 dosimetry records but they weren't typically on the 7 same form. 8 DR. MAKHIJANI: Okay. 9 DR. NETON: Physicians -- physicians really didn't 10 know what to do with it. 11 MS. BLOOM: I would say they might not even be the 12 sample at some of these sites --13 DR. NETON: Yeah. 14 MS. BLOOM: -- that you'd collect the urine during a 15 physical --16 DR. BEHLING: I mean (unintelligible) --17 MS. BLOOM: -- (unintelligible) you'd take the --18 DR. NETON: (Unintelligible) 19 DR. BEHLING: -- pre-employment requisite which 20 obviously would preclude the need for doing 21 urinalysis for isotopes, so it's not likely to be a 22 split sample. 23 MS. BLOOM: Uh-huh. 24 DR. BEHLING: I mean it's -- as part of your 25 employment that you submit to a physical, that

1 includes a urinalysis but there would be no reason 2 to at this point assess you for internal exposure --3 MS. BLOOM: Sometimes they did baselines --4 DR. NETON: Oh, yeah. Baseline --5 DR. BEHLING: Baseline? MS. BLOOM: Yeah. 6 7 DR. NETON: -- workers coming in, sure. I mean 8 there's -- there's issues -- and I don't know how up 9 these people were back then, but -- but people 10 coming in who have well water that has high uranium 11 values or for whatever reason would show positive, 12 you'd like to know that up front. DR. BEHLING: Yeah, well, I know in contemporary 13 14 times you use baseline --15 DR. NETON: And I don't know exactly what --16 DR. BEHLING: -- but in those days I'm sure they 17 were concerned --18 DR. NETON: And I don't know that this is even part 19 of the medical. We -- I'd like to look at the file. 20 DR. MAKHIJANI: Yeah. I -- I'll -- during a break I'll just come up with a name. Maybe we can 21 22 pull up the (unintelligible). 23 DR. NETON: That's fine. 24 DR. MAKHIJANI: Okay. Hans, any other follow-up on 25 the number of records question? Okay.

1 Next question is are there claims where zeroes were 2 entered into the records but no monitoring was done 3 and no back records exist? 4 DR. NETON: I guess I'd like to get a little more 5 clarity on this. We've talked among ourselves about this and there is a -- and I know Mark I think has 6 7 brought this up, where zeroes were entered in lieu 8 of no monitoring. I -- I'm not that familiar with 9 this issue. Mark, can you --10 MR. GRIFFON: Oh, I don't -- I -- I'm trying to 11 remember exactly how -- it -- it was actually 12 brought up by one of the -- the petitioners, I 13 believe, initially. 14 DR. NETON: Right. That's true. That was in the 15 evaluation report. MR. GRIFFON: Yeah. And I -- I've just been 16 17 following up on it as to whether we've resolved 18 anything on that because I think they're going to 19 raise it again. But the question of -- of, you 20 know, sort of putting zeroes in for entries that --21 DR. NETON: Right. See --22 MR. GRIFFON: -- there was actually some question of 23 whether they were actually putting zeroes in for 24 values that -- that were a positive value. 25 DR. NETON: Well, see that's what I was going to

1 raise is I don't know that we have any indication 2 that that happened. 3 MR. GRIFFON: Right, right. 4 DR. NETON: I think they're -- you're right. 5 you refreshed my memory, there were assertions by 6 petitioners that if a person weren't monitored, they 7 would put zeroes in there. And I -- we've discussed 8 this with Janet and she's not, as far as I can 9 recall, aware of this happening. But we don't have 10 any more to answer. This is sort of proving a 11 negative-type situation. 12 MR. GRIFFON: Right. I don't know how -- yeah, how 13 do you prove it, that's --14 DR. NETON: Right. 15 MR. GRIFFON: -- the problem I think. 16 DR. NETON: And I guess the worst case I could --if 17 -- if we -- accept the fact that that happened and 18 they -- and we can -- we can hopefully get 19 comfortable that they didn't take high values and 20 make them zeroes, and maybe that's part of this 21 validation thing I'm trying to do. But if they 22 entered zeroes where there was no monitoring, what 23 could conceivably happen is we would --24 DR. BEHLING: Assign dose --25 DR. NETON: -- compute missed dose --

1 DR. BEHLING: Yeah, because there was no dose. 2 DR. NETON: -- well, but -- but worse I think --3 MR. GRIFFON: Well, they were unmonitored --4 DR. NETON: If they were unmonitored we would assign 5 missed dose versus unmonitored dose, and that --MR. GRIFFON: Right. 6 7 DR. NETON: -- that would -- I suspect could make a 8 difference, albeit this would be on the low end of 9 the -- of the dose reconstruction spectrum, but --10 and honestly I'm not sure how we would deal with 11 that. If -- if we could find evidence that it 12 happened, we'd have to deal with it somehow. 13 guess is not unlike the situation where people are 14 saying well, I had a badge but I never wore it, 15 because then that's, you know, monitored dose --16 unmonitored dose when we're assigning zero missed 17 dose. 18 MR. GRIFFON: But I don't know if there's any kind 19 of -- of record that you have that show who was 20 assigned dosimetry. I don't think you've seen those 21 kind of records, have you? I mean you just have the 22 cards with their film data --23 DR. NETON: Right. 24 MR. GRIFFON: -- you don't have -- yeah. 25 DR. NETON: Yeah, and if there's a zero in there --

1 MR. GRIFFON: Right. 2 DR. NETON: -- it would normally be concluded by us 3 that, well --4 MR. GRIFFON: That they (unintelligible) --5 DR. NETON: -- that particular person wore the 6 badge. 7 MR. GRIFFON: -- dosimeter. Yeah. 8 DR. NETON: So... 9 MR. GRIFFON: Yeah. 10 DR. BEHLING: Couldn't you, on the basis of job 11 description, determine whether the person should have been monitored and -- and realize that the job 12 13 description itself would almost mandate the issue of 14 monitoring. If he worked in -- in --15 DR. NETON: Right. 16 DR. BEHLING: -- Building 6 and he was a certain 17 assigned job and there's no records, you can clearly understand that either the records are missing or he 18 19 was not monitored but should have been monitored. 20 DR. NETON: Right. I -- I think that's a good 21 point. We -- based on the job description, I would 22 -- I would guess that if a person was not monitored 23 and our professional opinion was they did not need 24 to be monitored, then -- and they had zeroes, the

missed dose would be larger than the ambient dose

1 that we would have assigned them. So --2 MR. GRIFFON: Right. 3 DR. NETON: -- in -- in most situations, that would 4 end up giving them a little more dose than we 5 otherwise would have. But the -- the worst 6 situation, though, is if we made the judgment that 7 they were monitored -- or should have -- did not 8 need to be monitored and should have been. 9 DR. BEHLING: Yes. 10 DR. NETON: But then that's an area where -- I don't 11 know. You know, there's --12 MR. GRIFFON: I -- I think -- Jim, I think what 13 you're saying is, you know, you don't have evidence 14 that this happened but, you know, if -- if, you 15 know, the worst case would be that you could 16 consider individual claims or verify on a case by 17 case basis maybe, I don't know. Because it seems to 18 me that you're right, that if their -- there were 19 zeroes but they say I never was monitored and you 20 look at them and -- and it turns out that they were, 21 you know, administrative or whatever --22 DR. NETON: Right. 23 MR. GRIFFON: -- then -- then you probably are --24 are going to give them the higher of the two doses,

coworker versus -- versus a missed dose and the

1 missed dose is likely to be higher anyway, so... 2 DR. NETON: Right. 3 DR. BEHLING: Just a quick question of the 4 approximately -- I think yesterday you said there 5 was about 120 claims that have yet to be processed, is that correct? 6 7 DR. NETON: For this time period. 8 DR. BEHLING: For this '49 to '57 time period. 9 those 120 claims, any idea how many of those 10 individuals are alive or being -- claims being 11 submitted by survivors, which allows you at least to 12 interrogate the claimant himself and sort of assess whether or not he worked where and under what 13 14 circumstances and what the probability was that 15 these uncertainties can be resolved by a direct 16 interview. MR. MACIEVIC: Well, see, I think a 17 misinterpretation comes to people from -- I mean 18 19 over time you have a person -- and I noticed this in 20 several files of different sites, and that a person 21 would be assigned a gamma dose and -- this is not 22 exactly what's happening here, but they'll be

assigned or have a dosimeter that they're checking

for gamma. They never had neutron dosimetry, but a

zero will go into that value --

23

24

1 DR. BEHLING: Okay. 2 MR. MACIEVIC: -- but it -- so technically they were 3 never monitored for neutrons, but they throw a zero 4 in there for the record-keeping purposes. Now that 5 you can see by, you know, looking at different files 6 and how it's laid out. But yeah, this -- this 7 question is --8 MR. GRIFFON: A little different, yeah. 9 MR. MACIEVIC: -- yeah, different than what that is, 10 yes. 11 DR. NETON: To answer to Hans's original question, I think -- I don't know exactly, but if -- if it holds 12 13 true for the rest of the sites, it's about 50 14 percent of the cases are --15 DR. BEHLING: Are survivors? 16 DR. NETON: -- survivors. 17 DR. BEHLING: Yes. How about also --18 MS. BLOOM: I think we also find, if you look at the 19 interviews, that somebody will say I didn't wear a 20 badge. And you go to the records and you go yes, 21 you did. 22 DR. NETON: Now see, this is when it's part of 23 security credential, they don't know. 24 MS. BLOOM: Yeah. 25 **UNIDENTIFIED:** Right.

1 DR. NETON: I mean they might not know, but --2 DR. BEHLING: The other thing I was just thinking of 3 as a cross-reference would be looking at if there 4 are -- and again I don't know because I haven't 5 looked at the records, but are there occupational 6 medical exposure records and would you have given a 7 person who's not a radiation worker an occupational 8 medical. Is that the criteria? In other words, 9 were people given occupational medical exposures who 10 were not radiation workers? And if that's the case, 11 then any time you see an occupational medical 12 exposure with no dosimetry records, you say chances 13 are you're missing records. 14 MS. BLOOM: I think it changed over time and at a 15 lot of the sites there was a pre-employment physical that included routine X-rays, I think. 16 17 DR. BEHLING: So that's not a distinguishing factor. 18 MS. BLOOM: I -- I think it changes and --19 DR. NETON: I don't think we can hang our hat on 20 that, no. 21 MS. BLOOM: -- finding documentation of the exact 22 criteria is really tough. 23 UNIDENTIFIED: (unintelligible) 24 MS. BLOOM: The other thing that I -- I did find in 25 the records when I was looking at one claim

1 yesterday, some letters that said oh, by the way, 2 you haven't turned in your badge, we've sent these 3 in. And so there is some indication that they were 4 tracking badges, they were following up and you 5 might be able to go back to records and see what was entered for that time period, did it come in later, 6 7 did it come in at all, was it a zero between two, 8 you know, large numbers. 9 MR. GRIFFON: Hey, Jim --10 DR. NETON: Yeah. 11 MR. GRIFFON: -- the other thing I'm trying to 12 remember is I'm not sure that this claim wasn't 13 partly based on -- on some Mont Mason memos that 14 they were referring to. I'm --15 DR. NETON: I think you're right, Mark, that there 16 was some issue about --17 MR. GRIFFON: There was some kind of claim in one of 18 those memos that there could have been, you know, 19 and -- and I think the -- the petitioners picked up 20 on that, so --21 DR. NETON: Yeah, I need to maybe go back --22 MR. GRIFFON: -- so I think we need to track that 23 back, too -- yeah, and I forget what the issue was. 24 DR. NETON: Right. I -- I think that I agree with

your -- your -- your statement earlier, though,

Mark, that, you know, if -- if we can't find any evidence that it did happen, we do have zeroes in the record and we do a case by case evaluation of the -- of the job title and either could make a determination to assign either -- well, missed dose, which would -- would probably -- we would probably assign missed dose at a minimum since they --

MR. GRIFFON: Yeah.

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DR. NETON: -- were monitored.

MR. GRIFFON: And I think you -- you -- I mean I don't know. It's your decision, but you might offer that, you know, if -- if there's a claim made by an individual and in that particular case -- I mean I can -- I can foresee a situation where -- it doesn't seem like it because you've got 90 percent of the (unintelligible) monitored here, but I can foresee -- on some sites I've been on there's been situations where maintenance people kind of fell through the cracks because they were assigned to a maintenance building, but they would go in other areas where they -- they should have had a badge but they just kind of fell through the cracks and they did work in those areas and -- and never were monitored. you saw a case like that, then you could say well, in those cases we'll give the higher of the two,

1 coworker or missed dose, you know --2 DR. NETON: Right. 3 MR. GRIFFON: -- if there's a claim made by -- and 4 we'll -- we'll handle that on a case by case basis. 5 So we don't have any evidence, but we will be claimant-favorable in -- in those situations if --6 7 if people make those allegations. 8 DR. NETON: Then maybe --9 MR. GRIFFON: I don't know. 10 DR. NETON: You know, Mark, I'm -- I'm looking at 11 this -- the SEC evaluation report and it looks like 12 -- I can't tell exactly, but it looks like they're 13 referring to the zero recorded for site breath radon 14 results. 15 MR. GRIFFON: Oh. 16 DR. NETON: The TBD indicates technique 17 (unintelligible) for internal exposures for other 18 isotopes based on uranium. Site breath radon 19 results indicate a 0.000 will not affect the ability 20 to reconstruct doses to individuals because 21 surrogate information is available. At least that's 22 what we've said, so unless there -- it appears 23 somewhere else in here -- I'll -- I'll -- I'll go --24 I've got to go through and -- and address this. 25 MR. GRIFFON: We should check back on that because I

1 recall them saying about the film, as well, that 2 maybe -- maybe that was more of a --3 DR. NETON: Yeah --4 MR. GRIFFON: -- a personal --5 DR. NETON: -- you're probably right. MR. GRIFFON: Yeah. I don't know. 6 7 DR. NETON: I'll -- I'll look through it and, you 8 know, I think the approach we were talking about 9 here is as best we're going to do and -- and in 10 fact, if it was a person such as like a chemical 11 operator and they were assigned zeroes -- well, see, 12 I -- I find it hard to believe with that many people monitored that --13 14 MR. GRIFFON: Right. 15 DR. NETON: -- someone like a chemical operator 16 would not have -- have a badge result. 17 MR. GRIFFON: But -- but -- yeah. It -- it does --18 it's a little tricker than that 'cause your 19 percentage of people monitored is based on all those 20 zeroes counting as real -- real monitoring, right? 21 So anyway... DR. NETON: Well -- well, you've got a point there 22 23 if -- if that is indeed true. 24 MR. GRIFFON: But I -- I tend to think you're right. 25 I mean it's like the majority of these people were

1 monitored, so... 2 DR. NETON: I -- I think if we go through and pull 3 out the ones that weren't monitored, sort of a 4 sampling, and get a feel for the -- the job titles, 5 vou know --MR. GRIFFON: I think that would be --6 7 DR. NETON: -- I'm envisioning something like a 8 little histogram or something. Yeah. Okay, I -- I 9 think we've -- we've got the thread here. 10 DR. MAKHIJANI: A couple -- couple -- couple other 11 things in this area is I think -- from my worker 12 interviews, it seems that guards were not monitored. 13 DR. BEHLING: Well, I think they were --14 DR. MAKHIJANI: And guards may have --15 DR. BEHLING: -- in -- in some instances. DR. MAKHIJANI: -- some of the guards may have 16 17 fallen through the cracks, but that -- I don't know 18 whether that's true or not. 19 (Unintelligible) DR. NETON: 20 DR. MAKHIJANI: (Unintelligible) the guards were 21 monitored? I mean this is just --22 MS. BLOOM: I believe guards were monitored. 23 MR. MACIEVIC: And I just read that this morning in 24 here --25 DR. MAKHIJANI: Okay.

1 MR. MACIEVIC: -- somewhere and I can't quite find 2 it. 3 DR. MAKHIJANI: All right. And I -- it's in the 4 TBD. I'll check it. 5 MR. MACIEVIC: Yes. DR. MAKHIJANI: I will check it. The other thing is 6 7 that, you know, as you said earlier, Jim, that 90 --8 that 90 percent of the white male workers were 9 monitored --10 DR. NETON: Well, at the very least. 11 DR. MAKHIJANI: -- but you may not have -- you may 12 not -- in the recent -- in the last years --13 DR. NETON: Yeah. DR. MAKHIJANI: -- but you may not have all the 14 15 records. 16 DR. NETON: Right. 17 DR. MAKHIJANI: And so I think in a way the zeroes question and the lost records question is sort of 18 19 tied up with, you know, your ability to -- to define 20 a job category and make an assessment of what that 21 situation is. And so this raises sort of -- one of 22 the questions that we brought up in our review is, 23 in the case of the survivor claimants sometimes you 24 have a tougher time with the job history if you

don't have the records because they may know only

1 the last one. I've -- I've looked at lots of 2 interviews and really it's --MS. BLOOM: Uh-huh. 3 4 DR. MAKHIJANI: -- it's a lot of don't knows, and 5 you can -- you can understand that. And so in -- in those cases I think -- I think it's kind of 6 7 important to know what -- what fraction of the --8 what portion of the universe you're dealing with 9 here in terms of unavailable records as well as 10 records that we think where the data might not be --11 DR. NETON: Right. 12 DR. MAKHIJANI: -- of the integrity --13 MS. BLOOM: I think when you're talking about job 14 title and it -- I don't think it matters whether you 15 have the employee's recollection or the survivor's 16 recollection, you should take it with a grain of 17 salt. And I -- I think one of the claims you were 18 talking about yesterday, I went back and looked at 19 There was a strike in 1963. That person who 20 became an administrative worker went back and in 21 1963 during the strike was the foreman in 22 maintenance again and that's in the record. 23 DR. NETON: Interesting. 24 MS. BLOOM: I've got a page number for you so you can take a look at that.

1 DR. NETON: Excellent. 2 MS. BLOOM: So I think as you pull the threads and 3 you find data, what you realize is that you can 4 probably figure out how to put your arms around 5 things, but you should never think that your data is 6 all, you know, that -- that you know everything you 7 need to know, because I think every time I pull on 8 those threads, I find out I missed something. Also 9 although that sounds really good, the dates are off. 10 And they're off in a way that makes that not exactly 11 the answer, but just a compounding factor to the 12 information so... 13 DR. MAKHIJANI: Okay. Yeah -- no, I mean I -- I 14 raised the question --15 MS. BLOOM: Uh-huh. 16 DR. MAKHIJANI: -- yesterday about I think Mr. B it 17 was --18 MS. BLOOM: No, I think it's a good -- I think it's 19 a good question and a good example. 20 DR. MAKHIJANI: -- purely, truly, as I don't know 21 what's going on here --22 MS. BLOOM: Uh-huh, yeah. 23 DR. MAKHIJANI: -- because we know -- well, you 24 know, if you take Fernald -- which is a facility 25 that I know perhaps best of all the ones that we

1 talk about because I've studied it for the longest -2 - in 1955 they had enormous emissions of 3 radioactivity. And if you -- if you were across the 4 street from Fernald rather than, you know, two miles 5 away, you could have gotten pretty big doses. MS. BLOOM: Uh-huh. 6 7 DR. MAKHIJANI: If you were in an office building on 8 the Fernald site, you could have gotten pretty big 9 doses. And so not to prejudge what goes in Fernald, 10 but just from the -- the stack records and the 11 scrubber records and so on, you can say quite a lot. 12 So I just raised that as a question without knowing 13 the answer because we don't -- we haven't done a 14 source term evaluation for Mallinckrodt. I don't even know whether it's possible to do such a thing 15 16 in terms of what went up the stacks. I haven't 17 looked at any of the records so I -- I'm not prejudging that answer. So this is actually very 18 19 useful information --20 MS. BLOOM: Uh-huh. 21 DR. MAKHIJANI: -- because people forget after --22 it's true, people do forget after so many years. 23 MS. BLOOM: And it was an odd situation but I think, 24 you know, that's certainly something that you 25 probably wouldn't necessarily remember.

1 DR. NETON: Right. 2 DR. MAKHIJANI: Yeah. 3 DR. NETON: I forgot where we were going here 4 (unintelligible). DR. MAKHIJANI: Okay. So is there a kind of -- how 5 6 do -- is there -- are you going to come back to us 7 with some kind of information about the proportion 8 of records -- roughly? You know, I realize that you 9 cannot --10 DR. NETON: It's just like with the internal 11 monitoring, we're going to come back with some type 12 of a -- of a distribution. I mean with 120 claims 13 it's, you know, it would be some -- worth of work, 14 but it would not be that hard to go through each one 15 16 DR. MAKHIJANI: Right. 17 DR. NETON: -- and -- and just check a box. know, it's going to be a little harder so it's going 18 19 to be a rough cut. We're going to say some or none because, you know, just because there's some does 20 21 not necessarily mean that it's complete monitoring 22 history but at least it's an indication you've got 23 something on the guy and the monitoring status

(unintelligible). It may be instructive to

determine what percentage of those are zeroes, yeah.

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1 DR. MAKHIJANI: Right. But if in that same thing 2 you could give us an idea of --3 (Whereupon, an unrelated discussion ensued off 4 the record.) 5 DR. MAKHIJANI: If in that same thing you could give 6 us an idea of job titles, you know, whether you have 7 -- not what the job title was, whether you have the 8 job title information. 9 DR. NETON: Right. 10 DR. MAKHIJANI: Because I think then your ability to 11 assign missed is obviously improved --12 DR. NETON: Right. 13 DR. MAKHIJANI: -- a great deal. 14 DR. NETON: Sure. Now again, you know, we have to 15 be careful in generalizing this for all cases. 16 you don't have a job title and as -- as we went over 17 that case yesterday where the lady -- I think it was 18 a uterine cancer -- we assigned her the highest --19 average of the highest ten doses at the facility for 20 each year without knowing anything about her job and 21 demonstrating that it was not likely that her 22 uterine cancer was caused by her exposure at work. 23 So, you know, these things -- yeah, I -- I'm very 24 reluctant to generalize and say if we don't have job

titles, this is how we're going to do it.

1 it depends on the case. I think that was an 2 instance of a short latency period possibly 3 (unintelligible) -- there's other factors that come 4 into play. 5 MR. MACIEVIC: And when you're talking about missing 6 data, too, I mean it depends if you're talking a 7 person who worked ten years and you're missing eight 8 of those years or a person who missed a few months 9 out of those years and you can interpolate in 10 between. So I mean missing data is --11 DR. NETON: Yeah. Well, this will be a little rough 12 and we're going to -- we're going to provide an idea 13 of are there bioassay, are there external results 14 (unintelligible) -- I know there's some with none. 15 I mean clearly we had one yesterday, but I don't 16 think we knew the job title of that person. We just 17 knew she worked here. I'm sure by -- by all accounts -- I mean she could have been a secretary; 18 19 she could have been a chemical operator. 20 DR. MAKHIJANI: Okay. I think that --21 DR. NETON: Okay. 22 DR. MAKHIJANI: -- the next question is how is NIOSH 23 addressing the issue of organ versus badge location 24 geometry for workers such as those who scoop 25 residue, shovel pitchblende into digesters, stamped

ID numbers on ingots. I think here the stamped ID numbers on ingots -- I wrote this before I went to St. Louis, I think -- may not be as big an issue as maybe at Fernald because I don't think they were doing it in the same way. But -- but the others do appear to be somewhat -- I don't know how you address the geometry problem. It was -- it was there at Iowa, we brought it up where the pits were in front -- in the pelvic area and we all estimated a factor of 2.5 or something like that. But I don't know how you would do that, approach that here, what the magnitude of the problem is.

DR. NETON: Well, we -- you know, we -- we've looked -- we've looked at this issue some since you raised it and we need to do some analysis. I mean we can't just out of hand reject it and say that it's not important, but it's our opinion that for area -- for functions like shoveling, it's not going to be a huge difference. I don't think we're talking as large a difference as the pits holding up at the abdomen. But, you know, it may be -- you know, I'm speculating here but, you know, something that's a 25 percent or something of that magnitude. So we're not -- I don't -- I don't think it's going to be as large an issue. But it was not addressed in the TBD

and is something that we have to answer.

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DR. MAKHIJANI: And is there kind of an approach that you've thought of -- thought of to developing an answer to --

MR. MACIEVIC: Well, one of the things that I'm going to be looking into is we have a new software. It's a transport -- radiation transport software called Attila that -- it's a deterministic model as opposed to the probabilistic like the MCNP, and we're going to try to do some calculations using that for different scenarios for body position with the band badge and organ position with respect to the source to get -- see what kind of limits there are. And you probably will see that as a person is moving off from that source by a certain percentage, as long as that distance between the badge and whatever organ are, you know, pretty much the same, there's not going to be that much difference. going to be in the cases where -- you know, like you're saying, where the person's definitely got it close to one place and the badge is now distinctly different that you'll see it. But this, hopefully will be able to generate some numbers for that and get a good feel for the kinds of distributions (unintelligible).

1 DR. NETON: Right. 2 DR. MAKHIJANI: One -- one of the things -- I have a 3 picture that I'll show you at the break from 4 Fernald. But one of the things that seems to be an 5 issue -- and Hans, correct me if I'm not on the mark here. But the -- the -- the angle -- the geometry 6 7 of the radiation source where sometimes the badge is 8 kind of dangling down and when the source is beneath 9 you, you know, you -- you don't have a perpendicular 10 incidence of -- of the radiation on the badge. 11 DR. BEHLING: The issue of angular, angular 12 sensitivity. 13 DR. MAKHIJANI: Angular -- so the angular -- this question came up in my mind reading the TBD and 14 15 trying to study the operations actually first from 16 an angular dependent point of view, because you've 17 obviously got the work beneath you --18 MR. MACIEVIC: Right. 19 DR. MAKHIJANI: -- and you're -- whenever you're 20 bending down, as you'll see in the picture that I 21 show you, the badge is dangling vertically and 22 you've kind of lost your near-perpendicular 23 incidence. 24 MR. MACIEVIC: Well, yes, and I think that's part of

the thing because some people wore badges like that

where you had a strap and the dosimeter was on this, and as you move forward that would also swing out as opposed to having it attached to the clothing at the chest level. So --

DR. BEHLING: Right.

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MR. MACIEVIC: But that kind of thing I think you could model up relatively easy with the Attila. I say relatively easy, but I think to -- to get some different types of scenarios, and if the -- the -the effects of geometry get even more important with -- as the energy or the beta -- like with the beta particle and the -- the mean free path across. you've got photons and that which have a high enough energy, the -- it will interact with the film on the -- the angularity effect is a lot less for something like that as opposed to a -- a beta particle which now you're hitting it at different angles so that penetration through or not hitting the filters is different. So you can mock that kind of scenario up a little bit better with -- using this Attila software and try to come up with some kind of factors in -- now as far as what you make as an assumption as to how many people wore straps where the dosimeter hung -- swung free versus how many would have kept it to their chest, I don't know.

1 DR. BEHLING: Well, it's not even the swing. I mean 2 angular dependence is something that you have to 3 worry about if you deal --4 MR. MACIEVIC: Sure. 5 DR. BEHLING: -- with an isotropic source because 6 you're getting simultaneous radiation from all angles other than normal. 7 8 MR. MACIEVIC: Sure. 9 DR. BEHLING: And of course, especially the deep 10 dose where you go through 1,000 milligram of fill 11 dirt material, whether it's cadmium or silver, the -12 - the thickness obviously is a function of -- of 13 deviation from normality in terms of deviance in 14 radiation. So you know -- and I've gone through --15 I think in some of my write-ups regarding -- was it 16 Iowa? -- as well as the discussion in -- under task 17 3, I provided some data that comes straight out of the classic textbook (unintelligible) that measured 18 19 the angular dependence of early film dosimeters. 20 And it's clear that any deviation from normality is 21 -- is going to affect the -- the response of the 22 film, so --23 MR. MACIEVIC: There's also an energy 24 (unintelligible) --

DR. BEHLING: Oh, yeah --

DR. NETON: Yes.

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DR. BEHLING: -- of course there is, but in fact in some instances if you're actually at 90 degree angle, your -- your film dosimeter will -- will not even be -- I think if -- if -- and it's an unusual case which would never happen, but I think in the early day they actually had the little lead marker that says if it didn't show up then that exposure wouldn't even be registered because they would assume that it's a false positive at 90 degrees. would give you a high dose because you're obviously avoiding the actual shield or the cadmium shield, but that would also be indicated to the reader because the -- the -- the lead marker wouldn't be seen on the film and therefore he would say ah, this is -- this is an artifact and it would essentially be recorded as zero when in fact it was a positive Those are all issues that are obviously limitations when you talk about film dosimeters, angular dependence.

DR. MAKHIJANI: Is it -- is it -- Hans, is it -- is it sort of more with the two-element thing, two-element dosimeter as opposed to the other ones where you might have --

DR. BEHLING: Well, it's -- it's -- it's -- all --

1 all dosimeters have that problem. Think of this as 2 -- as the filter that overlies your -- your -- your 3 deep dose portion of the badge. If you go at right 4 angles, it goes through basically one millimeter of 5 silver or cadmium, whatever it has. If it goes 6 through at an angle, you realize that --7 DR. MAKHIJANI: Yeah. 8 DR. BEHLING: -- the path is considerably thicker 9 and -- and so therefore you -- you see some 10 attenuation effect. And there are some data in the 11 early measurements that were done, empirical 12 measurements (unintelligible), that tell you exactly 13 at -- you know, at angle of 45 or 30 degrees, 90 14 degrees, et cetera, what -- what the reduced 15 response for a -- for a mono-energetic beam would be 16 17 MR. MACIEVIC: But you would also have to look at --18 I mean for that kind of thing -- and that's true if 19 you were under a certain condition all the time --20 DR. BEHLING: Yes. 21 MR. MACIEVIC: -- it will do that. But the 22 assumption is -- I mean you're moving around 23 continuously (unintelligible) --24 DR. BEHLING: Yeah. You're basically dealing with 25 one --

MR. MACIEVIC: -- (unintelligible) geometry --

DR. BEHLING: -- isotropic source --

MR. MACIEVIC: -- at -- at all times.

DR. BEHLING: Either the source is truly isotropic or your body motion makes it an isotropic. If you spin on your own axis, even a point source essentially appears to the dosimeter as an isotropic source.

DR. MAKHIJANI: So some idea of an approach may be with an illustrative calculation or two?

DR. BEHLING: Well, I think it's part of the uncertainty that's normally introduced, although I think from the uncertainty -- and this is one of the things that I've always taken exception to because I've been in the utilities where you -- you do your uncertainty by taking obviously several dozen badges and you put it in -- in a circular fashion. You rotate about a point source and then you essentially determine what the average value is and you find your sigma value. But in most instances you're dealing with a controlled exposure. It's acute exposure, it's mono-energetic exposure, and all badges are always normal to the incident radiation. So you get a sigma value that is an artificially low value. It doesn't, for instance, take into

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consideration many of the other issues, including the -- you know, the -- and if you remember this -the -- and I wrote up about this, the National Research Council report on film dosimetry and atmospheric testing, and they go through all of the different types of contributions due to laboratory, radiological and environmental as being contributors to the uncertainty. And in most instances there, you only deal with one uncertainty as opposed to the environmental and -- and laborat-- not laboratory, the radiological uncertainty that includes, for instance, angular dependence, which is never captured when you do that sigma value under controlled conditions because you don't rotate the badge or you don't necessarily subject it to high temperatures. In fact in some -- one of statements here involved a very high false positive read that was ultimately interpreted as being temperatureinduced. And, you know, for -- for TLDs you have so many factors, everything from chemiluminescence, (unintelligible) luminescence, you name it, they can all contribute, which is usually not captured when we deal with badges under controlled exposures. know -- you know, in a field you -- you put people into environment that are hot, humid --

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MR. MACIEVIC: The -- the thing is -- is -- I mean the assumption there is that you're not backing it up by some kind of --

DR. BEHLING: Yeah.

MR. MACIEVIC: -- dose investigation because many of these cases that you're talking about, like with the film and my working at Landauer for about five or -five years over there and doing research on different types of dosimetry, film, TLD, track etch types of things and all that, you can define what you -- you are -- the one nice thing about film is that you do have a picture and you can determine that there's something wrong when you're monitoring it. And several of these places, if there's something wrong with the badge, they'll put it in a code that there was something wrong with the badge. So in a case like that, you would have some kind of estimate or there'd be something to state that there is a problem here out of the ordinary. So I don't think you would -- you could say that all these offconditions were a routine practice that it would account for some huge variation in the badge. mean there's the -- the motion of the person and things like that which will cause some variation in the badge, but some of these things -- like in

chemiluminescence and that on a dosime -- on the TLD, one of the things that you do in having a heating ramp is that you basically burn off all the crap that might do that when before the photomultiplier even starts reading the number. If you read out a dosimeter and you anneal it -- because when you read it, you anneal it -- if that still has another read and what your process should be and has -- is, in reading several of these documents on rereading a dosimeter, if there's a residual of a certain percent left in the thermoluminescence, that again indicates that there's a problem with this badge and then you would go back. And having done that kind of thing at Fernald, too, is that you see people will take their dosimetry through the shower and get soap into the material --

DR. BEHLING: They put it in microwave ovens --

MR. MACIEVIC: That's right.

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DR. BEHLING: -- all kinds of things.

MR. MACIEVIC: -- and it will -- but it will show that that kind of duplication comes up each time. You'll see a dose for a person one month; they get the next dosimeter, there's zero; then the next month they're getting a reading again. And when you go investigate you find out that dosimeter is messed

up so you have to go back and do an investigation on those. So I -- the assumption is is that -- or in what you're saying is that when a reading has gotten on a badge that no one would have gone back and looked if something stood out as strange on it. And I think in many cases if the badge is operating normally, you're not going to have the investigation. But if something were -- if -- if you can see it --

DR. BEHLING: Yeah, of course. No, I -- I'm fully aware of it, but invariably it would -- would never been identified as an artifact or a -- a critical problem, it's still an issue of underresponse due to such things as angular dependence.

DR. NETON: All right. I -- I think we're getting far afield from the film badges at Mallinckrodt here, and let's focus on that I think. And I think what Greg is saying is true that, you know, you -- I don't know that we've got such an exaggerated sequence as you're suggesting where a guy is at 90 degrees to the source --

DR. BEHLING: No, no --

DR. NETON: -- and you've got a worker who's shoveling. And I think our -- our contention here is that we can do some bounding estimates using

1 Attila to demonstrate that when a person is three 2 feet away with a shovel from a -- from a vat of 3 something, that the -- the response of that film is 4 going to be probably -- I'm -- I'm guessing here, 5 but plus or minus 25 percent or something of that nature. And so I don't -- I don't think we have a 6 7 huge issue here that is unsolvable. We need --8 DR. BEHLING: In fact --9 DR. NETON: -- to do some sort of demonstration that 10 we believe it's probably within the uncertainty of 11 the whole process, so... 12 DR. BEHLING: But if there's an uncertainty, it's 13 probably in favor of the claimant, and that is film 14 badge contamination. That turned out to be a major 15 problem in --16 DR. NETON: Sure. 17 DR. BEHLING: -- specific testing place because of 18 fallout. You know, people do things, they touch 19 things, and they place their hands on it, and that 20 contamination is going to contribute to dose until 21 the moment you read out the film. And so you 22 realize --23 DR. NETON: Right. 24 DR. BEHLING: -- badge contamination is a major 25 problem.

1 MS. BLOOM: In the -- in our program for the other 2 sites, we've started moving towards assigning 100 3 percent AP exposures. That's been our assumption. 4 I'm not sure that that's what we're doing on 5 Mallinckrodt right now. I know just for the AWEs in general, though, that's the direction we've headed. 6 Does that change any of your concerns? 7 8 DR. BEHLING: Well, I have problems with the -- the 9 whole issue of the DCF because one of the things 10 that I believe all of the Appendix B and others are 11 -- are wrong because they make assumptions that I 12 think start out as an air dose and then they convert 13 it into tissue doses, which is not correct. For 14 instance, you know, you can tell in Appendix B that 15 for low energy photons if you have the PA geometry 16 exposure, the DCFs are virtually the same as in AP. 17 The problem is you're always wearing a badge up 18 front --19 MS. BLOOM: On your --20 DR. BEHLING: -- and so you realize those numbers 21 are off the wall. They don't -- they're not 22 correct. 23 DR. NETON: And that's -- that's an issue we need to 24 address in that document itself. I don't know that 25

1 DR. BEHLING: No, no, I'm not saying it is here. 2 DR. NETON: It's captured in another review 3 (unintelligible). 4 DR. BEHLING: But -- but when -- when -- when the 5 dose -- the doses are calculated that are not organ-6 specific and you convert a recorded dose into an 7 organ dose, you still have to defer to the -- the 8 Appendix B DCFs. And for instance, I -- I look at 9 the numbers and -- and for all of them -- if you 10 look at, for instance, the eye or the thyroid and 11 you have a PA geometry, well, you know, you realize 12 that all those tissues have to have a DCF greater 13 than one because you're dealing with an exit dose. 14 MS. BLOOM: Uh-huh. 15 The dose is here. Okay? And if the -DR. BEHLING: 16 - the source is behind you, what you're measuring on 17 your -- on your TLD or film is an exit dose, which 18 means that --19 MS. BLOOM: Right. 20 DR. BEHLING: -- any tissue that is in between the 21 source -- and that's starting on your back, the skin 22 on your back throughout your torso -- is going to 23 have a higher exposure than what's recorded on that film --24

MS. BLOOM: Right.

1 DR. BEHLING: -- badge by definition. So --2 DR. NETON: We're aware of that --3 DR. BEHLING: -- when I look at those DCFs I know 4 for a fact --5 DR. NETON: We have that comment and we will address 6 that comment. That's in procedures review and I'm 7 aware of that. I don't want to get off on -- on 8 that issue here. 9 DR. MAKHIJANI: And so basically, just for my 10 clarity, what -- what's in the procedures review 11 pipeline automatically get reflected in the dose 12 reconstructions --DR. NETON: Everything, across the board --13 14 DR. MAKHIJANI: (Unintelligible) 15 DR. NETON: -- those will all be reworked 100 16 percent. 17 **DR. MAKHIJANI:** Okay. DR. NETON: Any -- anything that is of a broad, 18 19 sweeping -- such as that, we would go back and redo 20 every single dose reconstruction that used that 21 concept. 22 DR. MAKHIJANI: Okay. So that's not an issue in 23 terms of reconstructability --24 DR. NETON: Right. 25 DR. MAKHIJANI: -- it's just a procedures thing that

1 2 DR. NETON: It's a matter of interpretation of the 3 existing data. 4 DR. MAKHIJANI: But what I can expect in terms of my 5 producing a draft on this particular question is that Greg will do a little exploration and then --6 7 MR. MACIEVIC: Yes, right. 8 DR. MAKHIJANI: -- you'll -- you'll send us 9 something? 10 MR. MACIEVIC: Yes. 11 DR. MAKHIJANI: So I can look at it and I could call 12 you. Presumably you'd have some (unintelligible). 13 MR. MACIEVIC: (Unintelligible) have the right 14 number, but yes, you can call me. 15 DR. MAKHIJANI: Okay. 16 MR. MACIEVIC: Yes, I'll (unintelligible). 17 DR. MAKHIJANI: Thank you. My -- my aim is to 18 produce a rough draft at least by the 15th and 19 closer to a final by the 20th so we can have our 20 internal --21 DR. NETON: Need to get something. Can you do 22 something like that fairly quickly, do you know, 23 Greq? I mean --24 MR. MACIEVIC: Yeah, I'm already talking with the 25 people who do the software about ginning-up some

scenarios like this. I'll -- I'll call them and talk today --

DR. NETON: The software is very nice, actually provides -- one of the features of it provides some very nice graphics. I mean, you know, images that you can show, you know, the source strength and all this stuff in relation to the -- you know, magnitude of the exposure at different positions relative to the person and badge and things so --

DR. MAKHIJANI: So, Hans, do you -- do you have -- are you familiar with this?

MR. MACIEVIC: It's pretty much brand new. I mean this is -- they've used this transport software. What this does is you model up -- whatever your universe is that you're going to create, if you have the person, the source, you'll model an area. It meshes this area and you calculate the radiation transpoint (sic) at all points within the entire area that you have. So what it's going to do is give you isoflux lines; it'll give you dose lines and all that through different materials and through all the particles. It's a -- it's a very -- it's quicker and more -- it's not -- it's just as accurate as Monte Carlo. But with Monte Carlo you end up picking a few points and do the calculation.

This will compute for the entire area and you'll get doses at all points, which is why this will be nice and you get nice graphics to show. If you have something here, it'll show you the dose and the flux distributions through the entire body at different organs, and if you placed a dosimeter here, you'll get to see what the lines are that pass through this point and all that. It's a really neat software for -- for doing this, and it's a -- I think people are just starting to use it. I mean it's been around a lot. I mean radiation transport using this method has been around a long time, but the computer capacity -- it's had to have so much to crank these numbers to follow every photon through that, it just took too much. Now it's starting to come into its own light so...

DR. MAKHIJANI: Greg, could I make a request that the -- the -- that we get the assumptions that you're going to put into this in very simple language that I can understand --

MR. MACIEVIC: Sure.

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DR. MAKHIJANI: -- so that we can do some back of the envelope checks? Because whenever there is a new complex model that's in the computer, it makes me very nervous and I like back of the envelope

1 checks because it makes it sense. And of course,
2 Hans is our point person on this and --

MR. MACIEVIC: Well, Bob Anigstein would probably be the person that looked at the computer.

DR. MAKHIJANI: -- and Bob, Bob Anigstein. And I will call Bob also and convey this to him and get him ready for, you know, whatever you have to say because this is an area, you know, in which in -- in our team basically it's Hans and Bob who look at the issues (unintelligible).

Mark, shall we move on to the next or did you have something?

MR. GRIFFON: No, that's fine.

DR. MAKHIJANI: Okay. Next question, external dose data did not provide job categories for -- in the five -- six boxes did not provide job categories for personnel whose badges had doses below 200 m-rem and in some cases below 300 m-rem, hence external dose data do not appear amenable to being grouped into job categories in ways that will enable the construction of external dose distributions for various job categories. How is NIOSH going to construct surrogate worker cohorts given the lack of job categories for data applying to majority of workers?

DR. NETON: This is just of course, referring to the five or six boxes. I think if you look through the list of the 12,000 TLDs or whatever, most people have a job title or category associated with them. So I think -- I'm not -- and I don't know what -- I'm not familiar with what you're talking about in the five or six boxes, but I -- I do know that people have individual badge readings with job categories -- in the CER database, at least.

DR. MAKHIJANI: Right. So -- so I guess what you're saying is that you're going to construct the -- this may be a more straightforward -- the question was long and maybe the answer is more straightforward, is that when I looked at these records the way they were, was -- they were simply identifying the most exposed --

DR. NETON: Right.

DR. MAKHIJANI: -- personnel, and they were calling them out by job category. And for most of the people -- the vast majority, 90-plus percent of the people -- there was no job category. But I don't know why they were collected in that way, but -- so I didn't -- I don't think that that data can be used for --

DR. NETON: Right.

1 DR. MAKHIJANI: -- coworker analysis. 2 MS. BLOOM: I would say that that's probably just a 3 partial set of data, and as you go through records 4 you find you've got lots of partial sets that you 5 need to pull together and make sure they match and 6 that they -- you know, you've got an issue of 7 zeroes, sometimes you find out the worker wasn't 8 here and that's what the zero means when you -- but 9 you find it in another record. And so that would be 10 similar with that, that that's just supplemental 11 data that we need to pull all together to make the 12 big set of coworker data. 13 DR. MAKHIJANI: Yeah. So -- but my -- my -- my 14 feeling is, looking at that data in, you know, more 15 detail than were able to do before the Iowa meeting, 16 it seems to me that -- that pretty much when -- when 17 you're constructing coworker data you have to do it from the individual records. 18 19 DR. NETON: Right. 20 DR. MAKHIJANI: It would not be possible to use those aggregate -- at least the aggregate record 21 22 that are in --23 MS. BLOOM: You -- you cannot --24 DR. MAKHIJANI: -- those boxes.

MS. BLOOM: What I've found is you cannot use any

set of records by itself, whether it's the original or the summary or anything else. And part of the -one of the main reasons is illegibility. You can't read names or numbers or dates, and sometimes you can find that in the summary when you can't read it on the card. Sometimes you find it in the card and you -- you know, so you need to look at it all and pull it together. That's why some of this takes time.

DR. MAKHIJANI: Yeah. Okay. I -- I think -- I think that's fair because -- because I've looked at

DR. MAKHIJANI: Yeah. Okay. I -- I think -- I think that's fair because -- because I've looked at a fair number of individual dose records and I do know that -- actually as -- as we said yesterday, the job title information at Mallinckrodt is pretty good.

DR. NETON: Yeah, it is. It's actually -- I -- I -- it's not in the exact dosimetry file, now that I'm looking at this, but there are work history information tied to all of the film badge records, and that was what I was going to end up sending Mark. Yeah, we do have a lot of job -- job titles, categories for Mallinckrodt workers, and that's -- that's clearly what we'd use.

DR. MAKHIJANI: Okay. I think that -- that (unintelligible) can consider it taken care of.

1 Table 33 has only scattered data for external dose. 2 How is NIOSH going to construct claimant-favorable 3 and scientifically-defensible values for surrogate 4 worker cohort external dose? I guess this is a different -- different incarnation of the same 5 6 question. 7 DR. NETON: Yeah, I think so. I was going to look 8 at Table --9 MS. BLOOM: Table 33 is the workplace exposure rates 10 11 DR. BEHLING: Yes. 12 MS. BLOOM: -- and that's -- that's to provide 13 people information on the kind of exposure rates 14 that did exist at Mallinckrodt. It's not 15 necessarily to reconstruct any specific job. 16 not meant at this time to reconstruct doses but it's 17 a supplemental information table to orient you to the site. You know, on a case by case basis it's 18 19 possible that it might be useful for somebody to 20 say, you know, look at these dose rates and look at 21 the badges, and this makes sense or it doesn't make 22 sense --23 DR. NETON: Right. 24 MS. BLOOM: -- but it -- it's not meant to be a

stand-alone, we're going to assign doses from this

1 table. 2 DR. BEHLING: Yeah, that was my question, how will 3 this table be used --4 DR. NETON: Right. DR. BEHLING: -- if at all. 5 6 MS. BLOOM: It's informationally and a case by case basis. 7 8 MR. MACIEVIC: And it does help to fill in the holes 9 10 MS. BLOOM: Yeah. 11 MR. MACIEVIC: -- where you've got data over here 12 and now you have some pieces here and see that it --13 it makes sense what you --14 DR. BEHLING: Any idea what instrument was used to 15 measure these dose rates? 16 DR. NETON: That's a good question. I was just 17 looking --18 DR. BEHLING: Something like an R02 or something? 19 DR. NETON: Yeah, I'm sure --20 MS. BLOOM: I don't -- did the RO2 exist at that 21 point? 22 DR. BEHLING: Probably not. 23 MS. BLOOM: Junos were very common at that point. 24 -- I'd have to go back to the records and find the 25 individual information. A lot of times you will

1 find some information, but typically it was an 2 ionization chamber. 3 DR. BEHLING: (Unintelligible) unit or something, 4 ANPDR --5 MS. BLOOM: Sometimes --6 **DR. BEHLING:** -- 37. 7 MS. BLOOM: I have not heard that instrument model 8 number. 9 DR. BEHLING: I used it in the field a lot, the 10 ANPDR-37. 11 MS. BLOOM: I've not seen that in the older records. 12 It might be there. Juno was the typical one, 13 Victoreen, Nuclear -- Nuclear Chicago was another 14 common instrument --15 DR. BEHLING: Yeah. MS. BLOOM: -- the 20 -- I can't remember if the 16 17 2650 was both a exposure rate measurement instrument 18 as -- I think it may have been. Sometimes 19 (unintelligible) detectors were used. 20 DR. MAKHIJANI: The -- I think there's an 21 intersection there between Table 33 and the data in the five, six boxes. And of course, you know, it's 22 23 not possible for me to go and check through, but in 24 terms of dose rates it may be -- I think there's 25 kind of quite valuable information in those boxes

1 that may be useful in modifying Table 33 and 2 updating it because I think some of the dose rates 3 indicates in -- in that collection of data may be 4 higher or may be more useful as a guide for job 5 titles because Table 33 is organized by job titles and areas, if I remember it correctly. 6 7 DR. NETON: Right. You know, my -- my thought on 8 this table --9 DR. MAKHIJANI: No, so I -- I just -- the data 10 seemed very, very sketchy in terms of years and --11 even as a guide. and it seemed to me that -- that 12 what there is in terms of the -- not -- it's not a 13 criticism of what's there, obviously --14 DR. NETON: No, sure. 15 DR. MAKHIJANI: -- you know, a very -- Janet did a 16 monumental job of compiling all of that. We've said 17 that I think a number of times, but -- but I think 18 there is some information in those boxes that could 19 be used as a complement to that data in particular. 20 But that may not be so because I made a -- SC&A made 21 a partial compilation of the data in those boxes and 22 23 DR. NETON: Right. 24 DR. MAKHIJANI: -- if you take a look at it, it 25 might be useful.

1 MS. BLOOM: And I think that's -- because this is 2 supplemental at this point, I think that's still 3 something to look at and we should look at it, but -4 5 DR. NETON: All right. You know, I'd like to point 6 out --7 MS. BLOOM: -- I don't see this as a primary --8 DR. MAKHIJANI: Okay. 9 DR. NETON: Right. You -- you need to look at the -10 - you know, the hierarchy of data usage. 11 clearly in cases where we have all these film badges 12 and we can validate them, then we would 13 preferentially use that, then followed by these area 14 results which are supplemental. And in the case 15 where you have zero information, these of course 16 would become very valuable. But I -- I think that 17 the second level, though, would like -- more likely 18 be coworker dose distributions rather than these 19 area badges. 20 MS. BLOOM: That seems unlikely that you'd use this 21 22 DR. NETON: Right, but they do --23 MS. BLOOM: -- unless you saw an incident or 24 something --25 DR. NETON: I -- I think they do sort of provide

1 some kind of a sanity check, though. If you have a 2 worker who spent like all year in one of these 3 places where you're seeing 50 mr per hour and his 4 CATI says I -- I held these boxes, you know, for 5 hours on end and -- and you're showing zero result, 6 you might question that and do a sanity check on 7 what you're -- what you're proceeding with. 8 DR. BEHLING: There are a couple of good ones here 9 at the feinc filter and that you talk about some of 10 those people who spent a lot of time handling these 11 filters --12 DR. NETON: Right. 13 DR. BEHLING: -- and on page --14 DR. NETON: Right. 15 DR. BEHLING: -- 232 you'll see some values here in 16 terms of what the dose rates would have been --17 DR. NETON: Right. DR. BEHLING: -- in front of the filter -- at one 18 19 foot, 210 milli-r. 20 DR. NETON: Right. 210? DR. BEHLING: No, I'm sorry. I'm sorry. No, no. 21 22 It's -- it's expressed in percent tolerance. 23 **DR. NETON:** Okay. 24 DR. BEHLING: So it's 210 percent, meaning that what 25 the tolerance dose was defined here as what -- 100

1 mr per eight-hour day. 2 DR. NETON: Right. 3 DR. BEHLING: That would have been then -- yeah, 210 4 mr for an eight-hour day, right? 5 DR. NETON: Yeah. So matter of fact, that crossed my mind when we were talking about these raffinate 6 7 workers. If you had a person with almost no 8 recorded dose --9 DR. BEHLING: Yeah. 10 DR. NETON: -- you've got a fairly good feeling that 11 this person was not working with these raffinate 12 streams where there are, you know, 50 mr per hour So that, in -- in my mind, is one approach 13 fields. 14 that we may take in this and to defining -- defining 15 some of these people at Plant 6 that we talked about yesterday. I used the external to help bracket the 16 17 internal potential for exposure. 18 DR. MAKHIJANI: Well, I -- I -- I'm not sure that 19 you can actually go there, because the main issue 20 with the raffinates, apart from that -- for that 21 small group of workers I think you could do that, 22 but the main issue with the raffinates that came up 23 yesterday was on the reprocessing of the raffinates, 24 which is a bigger issue --

DR. NETON: Right, right.

1 DR. MAKHIJANI: -- at Plant 6. 2 DR. NETON: Right. 3 DR. MAKHIJANI: It's not an issue --4 DR. NETON: Right. 5 DR. MAKHIJANI: -- just where those filters, so --MS. BLOOM: But -- but you'd still have those high -6 7 - much higher dose rates --8 DR. NETON: I mean the radium is still --9 MS. BLOOM: -- from handling the --10 DR. NETON: -- there, right? I mean --MS. BLOOM: 11 The radium and the progeny. 12 DR. NETON: -- it depends on which -- which stream. 13 DR. MAKHIJANI: Yeah. I -- I'm not -- yeah, you 14 know, if you're talking about the digesters, you 15 know, the -- the external dose (unintelligible) 16 shielded by all the acid in the tanks and very --17 pretty far, so --18 DR. NETON: Right, but that means that you're not 19 having much particulate exposure if it's in a tank. 20 See, in my mind, these raffinate workers -- the --21 the highest potential for exposure is the people 22 that are scraping the filters and drumming the 23 material. 24 DR. MAKHIJANI: Yes, I agree. 25 DR. NETON: At that point it's completely

1 unshielded, or almost unshielded, and you've got a 2 very large source term sitting right in front of 3 you, concentrated material. So I don't know where 4 I'm going with this, but it just -- it just --5 thought crossed my mind that we could use that to 6 our advantage to bracket these things. 7 DR. MAKHIJANI: Okay. I think we're done with that 8 question. Hans, do you have anything more on that 9 question? 10 DR. BEHLING: No. 11 DR. MAKHIJANI: Okay. Okay. Now here's -- here's 12 your question, Hans. Hans/NIOSH addressing the 13 nonlinearity and the optical density and dose at low exposures. Specifically it appeared that this could 14 lead to systematic underestimates of dose. 15 16 developing a correction factor to address this 17 problem? Do you want to clarify that question? I'm 18 not sure --19 DR. BEHLING: Yeah --20 DR. MAKHIJANI: -- I got it exactly right. 21 DR. BEHLING: On page 116 -- and this is commonly 22 done here and I'm not sure to what extent that error 23 is -- is going to amount to a -- a value that is 24 significant, but bullet number 7 -- and I think it's

stated elsewhere here on I guess page -- let's see,

where are we here.

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MS. BLOOM: Is that page 92, the --

DR. BEHLING: Yeah, page 92 is the -- the use of simple subtraction to segregate out beta from gamma components. And it's not something you can just look at and say okay, the open window is obviously a shallow dose or that it was responding to both photons and -- low energy photons and betas, and the shielded portion is obviously likely to be a response to higher energy photons only, and simply subtracting the two gives you an understanding of the beta components. And -- and that issue is discussed very -- in detail in the National Research Council, the 1989 report of atmospheric testing and film badge dosimetry. And they were very adamant in those days to try to identify what part of that exposure in the open window was due to betas as opposed to photons, and you will read in that description the difficulties -- and they finally quit in trying to make that distinction. reason being is that the film is not a linear response (unintelligible) in terms of optical density. When you plot net optical density as a function of exposure usually it's a sigmoid curve and -- and in essence simply subtracting optical

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density from the shielded portion from the open window is not necessarily the approach. In fact it gives you a false reading. And what they tell you do is -- and it has to be calibrated properly -- is to convert each value first into a dose, and then subtract the dose as opposed to the optical density. And that apparently is exactly what is done here and this is something that's -- at least in -- in that report -- was identified as a difficulty that was not easily overcome. You have to go back to the report and -- and again here, I -- they used basically the same film badges here, the Dupont 502 and the Dupont 510 for the low dose/high dose so that you could capture even doses in the, you know, in the tens of rads or even hundreds of rads. And I realized the same problem would probably prevail here in trying to assess the component, the beta component from the -- from the gamma component. MR. MACIEVIC: Well -- well, you're absolutely right. I mean if you have a two-filter badge you've got the -- the -- the thick shield, the -- that'll -

DR. BEHLING: Yeah.

- the -- that'll --

MR. MACIEVIC: -- wipe out all the low energy photons. You're not going to have that overresponse

under that particular filter. But yes, in the open window, if you've got beta and you have low energy photons, trying to pull out which is the low energy photon and which is the beta when all you have is one other filter -- that's why the multi-filter badges do much better because you can get that intermediate energies in there to go and see ratios between different filters. But I believe what we do on there -- I mean that is addressed in the OCAS Imp. guide as far as how to deal with these kind of -- you -- you're going to make -- what is it, calculation based on -- I believe that it's a photon exposure as opposed to the beta because the photon is going to give you the most conservative -- DR. BEHLING: Yeah.

MR. MACIEVIC: -- number. So when you're in the state of not knowing, you're going to go with the most conservative and say it's a photon exposure and that the overresponse is in there and you're going to compute that number. I think I'm going in the right (unintelligible).

DR. BEHLING: But I'm -- I'm not sure that necessary is the issue here. This is basically the -- the methodology of subtracting the optical density under the shielded portion of the film from the net

optical density on the open window. And according to that study -- as I mentioned, this is Frank

Massey's* report -- that is something you should not be doing. You should first convert each of those portions of the -- if you have a two-element film badge -- into dose and then subtract the dose from each other rather than subtracting that optical density from -- one from the other.

MR. MACIEVIC: I -- oh, I (unintelligible) --

DR. BEHLING: You have to go back --

MR. MACIEVIC: -- in there. I think what -- see, what you -- what you would do and -- and how I computed the doses in working with film is that you -- you're going to subtract off from all of them a blank which is a control --

DR. BEHLING: (Unintelligible)

MR. MACIEVIC: from that film. Now there -- then you're right, you convert it to dose and then you do your analysis between ratios and that between dose, not with densities, because you work with the dose numbers. Because yes, you don't -- don't work with the density values. I don't recall them doing that kind of thing where they're -- they're working in density units and then the end result is where they convert it. I think they are working with -- you

1	are you're subtracting off a blank in the
2	calibration. You have dose numbers under the filter
3	and in the open window and then you're doing the
4	subtraction there.
5	DR. BEHLING: Does anybody have a copy of that
6	report? The
7	DR. MAKHIJANI: NPPR?
8	DR. BEHLING: Yeah.
9	UNIDENTIFIED: (Unintelligible)
10	DR. BEHLING: I think it's online. I have a hard
11	copy but I
12	MR. MACIEVIC: Yes, that's
13	DR. BEHLING: can point to you the exact page
14	number
15	DR. NETON: I have the quote in our
16	DR. BEHLING: Yes.
17	DR. NETON: you know, but what Greg Greg, it
18	does say in our profile, and I mean looking at it
19	here, that Mallinckrodt did subtract the optical
20	densities.
21	DR. BEHLING: Yes, and that's
22	DR. NETON: So the relevant question then is
23	DR. BEHLING: something they don't want you to
24	do.
25	DR. NETON: Right. Now this of course would only be

1 relevant to skin dose. 2 DR. BEHLING: Yes. 3 MR. MACIEVIC: That's right. 4 DR. NETON: This does not have anything to do with 5 full body. DR. BEHLING: Yes. 6 7 DR. NETON: And that's -- that wasn't clear from the 8 way the question was phrased, so --9 DR. MAKHIJANI: Yeah. 10 DR. NETON: And that's fine but --11 DR. MAKHIJANI: I didn't --12 DR. NETON: -- I just want to make clear that this 13 is really a skin dose issue --14 DR. BEHLING: Yes, yes. 15 DR. NETON: -- not a deep dose. DR. MAKHIJANI: And a -- it's a little more than a 16 17 skin dose issue. Right, Hans? DR. BEHLING: Well --18 19 DR. MAKHIJANI: It would be --20 MS. BLOOM: Shallow or --21 DR. BEHLING: -- if you also convert that into the 22 breast and the testes, then it becomes -- and -- and 23 on that issue I even wondered to what extent -- why 24 -- for instance, under the DC9 code you do have the 25 eye as one of the potentials -- organs of -- of

concern, and also the thyroid. If you're going to consider testes and breasts as being part of that problem with a potential low energy photon or energetic beta, clearly the thyroid also would qualify. For -- for one, it's not covered by additional shielding such as clothing, as you would in terms of testes and the female breast. The thyroid is in fact an unprotected area. And especially for -- for females and -- and thin females, the overlying tissue of the thyroid is about 300 milligrams of -- of tissue, so an energetic beta could contribute to at least part of the thyroid dose. But anyway --

MR. MACIEVIC: I'd like to check into -- I mean I know that's what it says, but in my -- in the six months I've been here and reading Technical Basis Documents which are -- turn out to be mostly in seven different languages and you have to interpret what's being said in those documents -- I have a feeling they are not -- they do not mean that they are actually subtracting, 'cause that is not a process that I have seen in any of the other facilities where -- at -- at other sites and that in doing that process where they work straight with the densities. The only part where they're working with

1	the densities is you're subtracting off a blank, do
2	a dose conversion for filters, and then work with
3	the numbers.
4	DR. BEHLING: That's not
5	DR. NETON: Okay. Then what
6	DR. BEHLING: if that's the case
7	MR. MACIEVIC: that has to be checked into. I
8	I have a feeling that's they're saying it, but
9	that's not what they're doing. I think I I can
10	check into that and try to check some background
11	DR. NETON: Yeah, let's
12	MR. MACIEVIC: documents
13	DR. NETON: let's get it
14	MR. MACIEVIC: because yeah, I agree. That would
15	that that just doesn't seem as a process
16	that I've seen any other places. I've never done
17	that and I
18	DR. NETON: I believe early on these were done by EM
19	HASL, right, or
20	DR. BEHLING: It was an in-house processing.
21	MR. MACIEVIC: Well, they started outside and then
22	they went in-house.
23	DR. NETON: Yeah, I thought they did, also, but we -
24	_
25	MR. MACIEVIC: Right.

1 DR. NETON: -- we need to check into that --2 MR. MACIEVIC: Let me check in that because --3 DR. BEHLING: And as I said, it may not be a major 4 issue but I -- I noticed -- I mean it jumped out on 5 me when -- and I'm quite familiar with the film 6 dosimetry because of my work in the Marshall Islands 7 8 DR. NETON: Sure. 9 DR. BEHLING: -- and of course that was a -- a 10 direct report that corresponded to dose 11 reconstruction involving the Pacific testing period. 12 And -- and I remember distinctly that as a major 13 issue because there was so much interest in 14 understanding the different radiation components in 15 the badges and they -- they apparently gave up and 16 say we really don't have the means to do it. 17 MR. MACIEVIC: And when you're working with -- yes, you're right. The process should work with the dose 18 19 and if you're working with -- it -- it's not good --20 two -- two-filter badges leave a lot open. 21 DR. BEHLING: Yes. 22 MR. MACIEVIC: And using those when you have a very 23 good handle on the photon distributions and what 24 you've got, you can use that film and know what the

overresponse is and work with it. But if you're

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working under conditions where other things are
happening, two-filter badges don't cut it as well --

DR. BEHLING: Yeah, yeah.

MR. MACIEVIC: -- and that's why there is --

DR. BEHLING: Yeah.

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MR. MACIEVIC: -- but we do the --

DR. BEHLING: On -- on -- on that issue and -- and it's only peripherally similar to -- to the concerns here, what will happen in terms of assigning -obviously IREP demands us to identify the type of radiation that is potentially recorded under the shallow dose or open window as either being a beta component or less than 30 keV. And yet it certainly makes a big difference to -- to -- to distinguish between the two of them. One has a choice in saying it's either very low energy photon radiation that separates the deep dose from the shallow dose, or it's a beta component that separates the deep dose from shallow dose. And yet for IREP input it's a significant difference in terms of the relative effectiveness factor because when you look at, for instance, electrons greater than 15 keV which would correspond to beta particles -- and as I said, I --I've done a calculation that compares the two in terms of POC versus the less than 30 keV photons --

1 the POC goes from 14.81 percent to 37 for skin. 2 so it's important to know how will this be treated, 3 because for -- for skin exposures -- and you will 4 probably encounter some squamous cell carcinomas --5 the -- the interpretation of the shallow dose is going to be heavily affected by -- or the -- the POC 6 7 will be heavily affected by your assignment of a 8 shallow dose based on either less than 30 keV 9 photons versus greater than 15 keV betas. 10 MS. BLOOM: Were you saying that the less than 30 11 keV photons are giving you the higher POC? 12 DR. BEHLING: Much higher. 13 DR. NETON: Oh, yeah. 14 DR. BEHLING: Much higher. And it's up like at two 15 and a half (unintelligible) --16 MR. MACIEVIC: That's why I thought we --17 DR. BEHLING: -- higher. 18 MR. MACIEVIC: -- defaulted to that as if there's 19 not a known... 20 MS. BLOOM: I think we've actually gone the other 21 way. DR. NETON: It depends -- it depends on -- on the 22 23 facility. I mean --24 UNIDENTIFIED: (Unintelligible) 25 DR. NETON: -- we know for -- for plutonium

1 facilities it's going to -- you know, the low energy 2 dose is going -- it's going to be less than 30 keV 3 photon. 4 MS. BLOOM: For uranium we've --5 DR. NETON: At a uranium facility --6 MS. BLOOM: -- typically gone the other way. 7 DR. NETON: -- you need -- yeah, you need to look at 8 the -- the relative magnitude of the contributions 9 of the different spectra and the protactinium 234 10 admittedly is --11 DR. BEHLING: That's true. 12 DR. NETON: -- going to dominate -- dominate the 13 shallow dose. 14 DR. BEHLING: Yes, yes. 15 DR. NETON: In fact it's not unusual in uranium 16 facilities to get ten to one ratios of skin to deep 17 dose. I've seen that at -- consistently at Fernald 18 and we've seen that in the Mallinckrodt records --19 DR. BEHLING: And I think there's --20 DR. NETON: -- but most of that dose is going to be 21 due to the beta. 22 DR. BEHLING: Yeah. 23 DR. NETON: I -- I think you'd be very hard-pressed 24 to demonstrate that the predominance of those 25 shallow dose is from less than 30 keV photons. Ι

1 don't think we would --2 DR. BEHLING: But there should be some guidance so 3 that we don't have different people selecting one as 4 opposed to the other --5 MS. BLOOM: We have been putting --DR. BEHLING: -- because I think it's important --6 7 MS. BLOOM: -- that in the site profiles. 8 sure if it's in this one, but we have been selecting 9 10 DR. NETON: It's a replay of the first comment, that 11 we need to make sure that people don't arbitrarily -12 13 MS. BLOOM: Verify that. 14 DR. BEHLING: Use one or the other. DR. NETON: -- use one or the other because then you 15 16 get into consistency problems. 17 DR. BEHLING: Yeah, it's a two-and-a-half-fold 18 difference. 19 DR. NETON: I -- I think we've been doing these at 20 many uranium facilities, though, and maybe we've 21 just sort of gotten loose in our -- our write-ups. 22 But in uranium facilities -- I think it's -- it's in 23 general going to be the beta dose. 24 DR. BEHLING: Yeah. 25 DR. MAKHIJANI: Would you agree with that Hans?

1 DR. BEHLING: Yeah. 2 DR. MAKHIJANI: Okay. 3 DR. NETON: So -- yeah. I don't know if we have a 4 problem but I think you're right, for consistency 5 purposes we should -- we should point that out and make sure that we do it that way. 6 7 MR. MACIEVIC: Look and see what I (unintelligible). 8 DR. MAKHIJANI: So that -- so that if -- well, I'm 9 just trying to make the issues clear in my head so I 10 don't get off in the wrong direction. So Greg, you 11 -- you agree that if -- you don't think that they 12 actually were subtracting optical densities --13 MR. MACIEVIC: No, I don't. 14 DR. MAKHIJANI: -- in reading the -- because we 15 don't have the film badges. 16 MR. MACIEVIC: Right. 17 DR. MAKHIJANI: Right? So it's very important to 18 determine -- Cindy, do -- do we have the film 19 badges? 20 I -- I don't believe we have the film MS. BLOOM: 21 badges. I don't know that we don't have the optical 22 density readings. We may have that --23 DR. NETON: It may be possible to go back --24 MS. BLOOM: -- on some of them. 25 DR. NETON: -- and look at this but --

1 MS. BLOOM: I don't -- I don't think we have them 2 all. 3 MR. MACIEVIC: Look at some of them and see a 4 general case --5 DR. NETON: Finding them and -- and doing it in a systematic manner but --6 7 DR. MAKHIJANI: Well, it would be useful to check 8 just a few if you have them because it seems --9 DR. NETON: Yeah. 10 DR. MAKHIJANI: -- to me that this is a critical 11 issue --12 DR. NETON: Well --13 DR. MAKHIJANI: -- because we agree that if they did 14 -- if they did do it as it says in the TBD, this 15 would be a problem. 16 DR. NETON: Well, it's a critical issue for skin 17 dose. 18 DR. MAKHIJANI: Skin dose for these --19 DR. NETON: For the shallow organs. 20 DR. MAKHIJANI: -- for the shallow dose or --21 DR. NETON: (Unintelligible) certain organs, right. 22 DR. MAKHIJANI: -- no, I got that. Okay. So for --23 for the shallow dose organs it's important to know 24 what they did, and you don't think they did that --25 MR. MACIEVIC: No, I --

1 DR. MAKHIJANI: -- but some -- it would be useful to 2 have some verification, either from this site or the 3 (unintelligible) what was the general practice at 4 the time at least. 5 DR. NETON: Right. DR. MAKHIJANI: If there could be evidence from the 6 7 time about that, that would be very useful. 8 DR. NETON: Some of the language --9 MR. MACIEVIC: Because I have a feeling -- yeah, now 10 whether I can get it from Mallinckrodt or not, but I 11 know practice was not done that way at several other 12 sites. I mean that approach --13 DR. NETON: Most of these people did not make up 14 their approaches. They all --15 That's right. They took it from one MR. MACIEVIC: 16 place where they knew where they were working with 17 it and they took it, so --(Unintelligible) we have the densities 18 MS. BLOOM: 19 there because I know I --20 DR. NETON: What are those two pages that you cited 21 earlier? Was it 92 --22 Yeah, 92 --DR. BEHLING: 23 DR. NETON: Down at the bottom --24 DR. BEHLING: -- bottom of page 92. 25 DR. NETON: And there was another one.

1 DR. BEHLING: And then on page 2-- 116 at bullet 7. 2 DR. NETON: That's what I was looking for. So what 3 bothers me --4 THE COURT REPORTER: Excuse me one minute, 5 please. 6 DR. NETON: I'm sorry. 7 THE COURT REPORTER: I'm sorry. 8 (Pause) 9 DR. BEHLING: Okay. On page 92 you'll see the 10 second full paragraph starting with "There was a 11 series of meetings", et cetera, where you talk about the beta shield and so forth. And then again on the 12 13 very bottom of that page, "Net window density from 14 beta exposure alone is equal to actual net window 15 density minus net window density from the gamma 16 exposure alone. 17 DR. NETON: Right. 18 DR. BEHLING: And so -- and then again on page 116 19 repeats that under 7 --20 DR. NETON: Right, and it --21 DR. BEHLING: -- and so it tends to --DR. NETON: Well --22 23 DR. BEHLING: -- you know, multiple -- at multiple 24 points in --25 DR. NETON: Sure.

1 DR. BEHLING: -- in the --2 MR. MACIEVIC: No, I --3 DR. NETON: What I was going to --4 MR. MACIEVIC: No, I agree --5 DR. NETON: Greg, if you look at --MR. MACIEVIC: -- what I'm saying is you're right. 6 DR. NETON: -- but if you do look at page 116 I mean 7 8 it gives you the exact references that you need to 9 look at. 10 MR. MACIEVIC: Yes. 11 DR. MAKHIJANI: On page 92 also it has a 12 Mallinckrodt reference from the time. 13 DR. NETON: Right. 14 DR. MAKHIJANI: So I -- I presume it is on the O 15 drive (unintelligible) --There are three references --16 DR. NETON: DR. MAKHIJANI: -- I looked at that also. 17 There are three references in here and 18 DR. NETON: 19 then what -- what kind of looks suspicious is for 20 Mallinckrodt it is assumed that the beta readings 21 are subtracted so, you know, I don't know if these 22 memos were -- were bandying about the issue and --23 and thinking about it and how the effects are, so if 24 you look into those it'll probably give you --25 MR. MACIEVIC: Yes.

1 DR. NETON: -- a better feeling for what was 2 actually done. 3 DR. BEHLING: On that subject, can I make a comment, 4 and I -- I don't know who -- who writes these TBDs 5 but, you know, I did a -- I was curious. I did a word search on this TBD and the word "assume" --6 7 "assumes" comes up 21 times. The word assumed, past 8 tense, 175 times, and the word "suppose" comes up 14 9 Which leads you to question how much is 10 there that is really of substance. In fact it -- if 11 I point to the page 43, there were -- just in one 12 paragraph -- the word "was supposed to be worn", "was supposed to be vacuumed", "was supposed to be 13 14 installed", et cetera, and it's --15 MR. MACIEVIC: I think you're --16 DR. NETON: Well, you know --17 MS. BLOOM: I think we want to represent --18 DR. BEHLING: No, I understand. 19 MS. BLOOM: -- what we know and what we don't know -20 21 DR. BEHLING: I understand. 22 MS. BLOOM: -- and I think that you'd all have our 23 heads if we put it in black and white and --24 DR. NETON: We're trying to do reasonable estimates 25 of doses here. A reasonable man would take those

1 things and say --2 DR. BEHLING: Yeah. No, I understand but --3 DR. NETON: So I mean to be fair to the -- to the 4 writer, you've got to put that kind of language in 5 there, otherwise our lawyers would have our heads. DR. BEHLING: The stakeholders will read some of 6 7 that data and -- and sort of wince every time they 8 hear the words "were supposed to". 9 MR. MACIEVIC: It's equivalent to when you see on 10 the news when they say "the alleged killer" does 11 this, "the alleged" --12 DR. BEHLING: Well --13 MR. MACIEVIC: -- I mean you -- you're not going to 14 go and commit to saying -- because there is a 15 possibility there's more data on other things --16 DR. NETON: The bigger -- the bigger issue is with 17 "assumed", and I've gotten called to task on this at 18 several -- several meetings, public meetings --19 DR. BEHLING: So I'm not the first to --20 DR. NETON: No, no. The claimants will get up there 21 and say this thing is fraught with assumptions. 22 DR. BEHLING: Yeah. 23 DR. NETON: They assume my exposure was this and 24 this. And they're really looking at it from a 25 different perspective, which is -- may be correct

1 because they're the claimants. But when we make 2 these assumptions in general, we will insert 3 assumptions that we believe are generous --4 DR. BEHLING: Yeah. 5 DR. NETON: -- and claimant-favorable. I hate to 6 use that word so much anymore but -- so they are 7 assumptions, but that's what science does. 8 science makes certain assumptions that bracket the 9 truth and reality. You -- you never know anything 10 with 100 percent certainty so -- I hear what you're 11 I'm sensitive to it, but I'm not sure -saying. 12 DR. BEHLING: It struck me odd to see that many 13 words that assume, assume, assume --14 DR. NETON: I think there are some cases --15 DR. BEHLING: -- were supposed to --16 MS. BLOOM: That's less than one per page though. 17 That's not too bad. DR. NETON: Although --18 19 UNIDENTIFIED: Cindy --20 DR. NETON: -- I would -- I would say that in 21 certain cases like this last one I just read, you 22 could probably get by with different language. 23 DR. BEHLING: Yes. 24 DR. NETON: You know, it's not -- it is assumed, but 25 based on the evidence provided, we will use this,

1 you know, that sort of -- but yeah, I -- I agree 2 that, you know, it's difficult when you have these 3 words, these -- these --4 DR. MAKHIJANI: And sometimes there is a disconnect 5 between how would you use the word "assumption" in a scientific --6 7 DR. NETON: Yes. 8 DR. MAKHIJANI: -- tract and in a scientific context 9 as opposed to a general sort of literary context in 10 which --11 DR. NETON: Right, exactly. 12 DR. MAKHIJANI: -- you imagine that when you don't 13 know anything you make that unfounded assumption. 14 There -- there might be some kind of --15 DR. NETON: Right. 16 DR. MAKHIJANI: -- implication that you're making an 17 assumption because you don't know anything --18 DR. NETON: Well, exactly --19 DR. MAKHIJANI: -- as opposed to making an 20 assumption in a scientific context. 21 MS. BLOOM: Based on data. 22 DR. NETON: So --23 MS. BLOOM: And I think we do both, and we get 24 called to task when we leave it with that very open, 25 you know, what I -- this is my best guess.

1	that's one kind of assumption and that's a very wide
2	open one. And then you have your assumption where
3	you say okay, they said this and they said this and
4	they said that
5	DR. NETON: Well, right
6	MS. BLOOM: so we have to go somewhere with it.
7	DR. NETON: I mean, you know, oftentimes you'll read
8	I don't we assume Class Y, Type Y solubility.
9	Well, that's a great assumption for the claimant. I
10	mean
11	MS. BLOOM: Uh-huh.
12	DR. NETON: but they they read that, like you
13	say, in in the general parlance and say geez,
14	they had to assume all these things.
15	MS. BLOOM: They didn't know what it was, so
16	DR. NETON: They didn't know.
17	MS. BLOOM: Yeah, but we gave you ten times the
18	dose.
19	DR. MAKHIJANI: Mark, I think
20	DR. NETON: Mark, did you have something to say?
21	MR. GRIFFON: No, I'm I'm assuming we're done
22	with this topic.
23	DR. NETON: I think we are. We've gone off and
24	Okay. We just have a couple more questions
25	MS. BLOOM: Which which

1 DR. NETON: -- and then we're going to get into 2 other issues. Do we need to take a break yet or 3 should we finish up with these two? I -- I think 4 that we'll just take a ten-minute break, if that's 5 okay --6 DR. MAKHIJANI: That's fine. 7 DR. NETON: -- for comfort and --8 MR. GRIFFON: What is left, Jim, because I may have 9 to bail out at this point. 10 DR. NETON: Okay. 11 MR. GRIFFON: What topic --12 DR. MAKHIJANI: What is left, has NIOSH -- the two 13 questions on my list, Mark, are has NIOSH evaluated 14 importance of issue of highly variable response of 15 films for photons at energies less than 100 keV for 16 uranium facilities. It's a kind of a continuation of this --17 18 MR. GRIFFON: Yeah. 19 DR. MAKHIJANI: -- film badge thing, and then the 20 last question, is NIOSH using the open window dose 21 as shallow dose for skin testing in breast dose 22 estimation. So both of them are kind of 23 elaborations of what we've been talking about. 24 MR. GRIFFON: Okay. All right. I -- I think -- if 25 it's okay, I think I probably won't -- won't come

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1
             back on.
2
             DR. NETON: Okay.
3
             MR. GRIFFON: You guys have got it covered and --
4
             DR. NETON: All right. Then we'll --
5
             MR. GRIFFON: -- I've got work to do.
6
             DR. NETON: Yeah, well --
7
             MR. GRIFFON: Okay.
8
             DR. NETON: All right, Mark.
9
             MR. GRIFFON: All right. Thanks.
10
             DR. MAKHIJANI: Thank you, Mark.
11
             DR. NETON: We'll take a -- we'll take a short break
12
             here.
13
             (Whereupon, a recess was taken from 10:25 a.m.
14
             to 10:45 a.m.)
             DR. NETON: We're back from our break and we're
15
16
             continuing on. I think we have questions 7 and 8 to
17
             complete --
18
             DR. MAKHIJANI: Yeah.
19
             DR. NETON: -- at least on my list --
20
             DR. MAKHIJANI: Right.
21
             DR. NETON: -- external (unintelligible).
22
             DR. MAKHIJANI: Okay. Question 7, has NIOSH
23
             evaluated the importance of the issue of the highly
24
             variable response of the film to photons at energies
25
             of less than 100 keV for uranium facilities?
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1 this is kind of a continuation of how you read these 2 film badges. Hans, did I --3 DR. BEHLING: Yeah, because there is -- there are 4 two portions of it. For very, very low energy, the 5 film badges underrespond; for somewhat higher, they overrespond due to the photoelectric effect that 6 7 obviously for -- for silver bromide and you realize 8 that obviously is an issue here. And I think in 9 other instances NIOSH has basically generously said 10 no, we'll -- we'll -- we won't make a correction. 11 We will accept the overresponse and deal with the --12 the dose as-is. Is that correct? 13 MS. BLOOM: Uh-huh. MR. MACIEVIC: Uh-huh. Yeah --14 15 DR. NETON: Yeah. I think it's not -- it's not 16 unlike what we've done in other facilities. 17 MR. MACIEVIC: Exactly. Yes, it's -- it's true. 18 DR. BEHLING: Okay. 19 DR. NETON: Yeah, that was -- well, when I saw this, 20 I -- my -- my original impression was well, yeah, 21 we're going to be overestimating because of the 22 overresponse --23 **DR. MAKHIJANI:** Yeah. 24 DR. NETON: -- you know, of low energies and --25 yeah. It's -- that's what we're doing.

1 DR. MAKHIJANI: Okay. Well, that takes care of 2 that. 3 DR. BEHLING: And -- and the next one is basically 4 what we've already touched on and --5 DR. NETON: Right. DR. BEHLING: -- but for instance if -- if -- I want 6 7 to draw attention to page 48 of the TBD. It does 8 make some statements here that again goes to this 9 issue of the shallow dose. And -- and -- and I 10 guess sort of --11 DR. NETON: Where on 48, Hans? 12 DR. BEHLING: Top of the first paragraph but middle 13 of that paragraph that starts out with "Dose rates 14 measured with instruments were combined with time 15 measurements", and it says that the gamma doses were 16 said to agree well with film badge reading, but not 17 the beta doses. And -- and I guess this is -- goes 18 all back to the issue of how do we deal with beta 19 components. 20 DR. NETON: Right. 21 DR. MAKHIJANI: Which -- which paragraph are you in, 22 Hans? 23 DR. BEHLING: I'm at the first --24 DR. NETON: First major paragraph. 25 DR. BEHLING: -- paragraph on page 48 that starts

1 out with "As a result". 2 DR. MAKHIJANI: Okay. 3 DR. BEHLING: And middle of that paragraph it starts 4 a sentence, "Dose rates were measured with 5 instruments" --6 DR. MAKHIJANI: Okay. 7 DR. NETON: So --8 DR. BEHLING: -- and it just --9 DR. NETON: -- you're saying that the instrument 10 readings for beta did not measure -- do not agree 11 well with film badge readings. DR. BEHLING: Film badge data and -- and it goes 12 13 back to the same issue that we've been discussing. 14 DR. NETON: Yeah, I think we need to go back and 15 Greg needs to review the protocol for --16 MR. MACIEVIC: I will, yes. DR. NETON: -- looking at the beta doses --17 18 DR. BEHLING: Yeah, yeah. 19 DR. NETON: -- and seeing what -- what we have. 20 DR. BEHLING: I mean maybe due to the way the -- the 21 film was processed that we discussed earlier, maybe 22 due to other factors, I don't know and I -- I -- get 23 24 DR. NETON: Well, it's not surprising that if 25 instruments -- survey instruments for beta would --

1	DR. BEHLING: Yeah.
2	DR. NETON: not agree with a a badge that may
3	have been calibrated theoretically properly
4	DR. BEHLING: With uranium slag or something.
5	DR. NETON: uranium slag, which is what I think
6	they've used here.
7	MR. MACIEVIC: Yes. Uranium slag is what's used for
8	the (unintelligible).
9	DR. NETON: Standard uranium slag, so
10	MS. BLOOM: Uh-huh.
11	MR. MACIEVIC: And I mean yes, the pics and other
12	things, too, don't agree well with the dosimeters
13	and it's
14	DR. NETON: Particularly for these high energy betas
15	that we're talking about from from uranium, so
16	DR. MAKHIJANI: So we'll get something on two issues
17	
18	MR. MACIEVIC: I will
19	DR. NETON: Right.
20	DR. MAKHIJANI: I think. If I might summarize,
21	though
22	DR. NETON: Sure.
23	DR. MAKHIJANI: we dealt with question 8, for the
24	record. One will be this geometry question using
25	this Attila.

1 MR. MACIEVIC: Yes, number 3. 2 DR. MAKHIJANI: Sort of like Attila the Hun? 3 MR. MACIEVIC: Yes. 4 DR. MAKHIJANI: As in the Hun? 5 MR. MACIEVIC: Yes. DR. MAKHIJANI: Because I want to tell Bob 6 7 Anigstein, you know, to maybe look into it. And the 8 other will be the shallow dose, this complex of 9 questions with --10 DR. NETON: Right. 11 DR. MAKHIJANI: -- with the shallow dose, how it was 12 done, optical density --13 DR. NETON: Right. 14 DR. MAKHIJANI: -- and so forth. Right, Hans? 15 DR. BEHLING: Yeah, yeah. 16 DR. MAKHIJANI: Do you have other --17 DR. BEHLING: Yeah, I just have a couple of other 18 probably insignificant issues, but on page 51 there 19 is reference to 100 millirem radium beryllium source 20 that might have been the source of neutron exposure 21 which I'm not sure are a -- a significant issue here 22 if it's, you know, a source that was used in the 23 radium and it -- on the bottom of page 51 it talks 24 about in the early years --25 DR. NETON: Oh, in the Shotgun Laboratory --

1 DR. BEHLING: -- Shotgun Laboratory. I don't know 2 if that's something that needs to be looked at. 3 It's probably something that was used by maybe one 4 or two people maybe and has no significance to the 5 workforce at large. DR. NETON: Well, it -- it talks about that it ended 6 7 by September 1944. 8 DR. BEHLING: Okay. In that case it's obviously 9 academic anyway. 10 DR. NETON: In that case it's not relevant for --11 DR. BEHLING: Yeah, yeah. 12 DR. NETON: -- this time period. 13 DR. BEHLING: I guess in internal exposures, we're 14 almost exclusively focusing on inhalation exposures 15 as opposed to ingestion. 16 DR. NETON: Correct. 17 Given -- but given -- yeah, I know. DR. BEHLING: Ι wasn't there yesterday afternoon so I may be 18 19 redundant in some of my questions, but with regard 20 to the possibility that people were using their 21 hands to shovel stuff and touching stuff and 22 obviously there was a significant amount of 23 contamination all over the place that people might 24 have transferred to their mouth inadvertently when 25 they touched their lips or took a cigarette break or

1 a lunch break, whatever it is. The issue is one of 2 solubility. It's generally assumed that most metal 3 oxides are relatively insoluble but that's usually 4 as a result of tests that are done in a neutral 5 aqueous solution. When you ingest it obviously these materials would encounter the acidity of the 6 7 stomach, which is basically one normal hydrochloric 8 acid which considerably changes solubility. Is that 9 an issue that needs to be looked at? It wouldn't 10 matter if we're dealing with urinalysis which 11 obviously doesn't care (unintelligible). 12 DR. NETON: Well, that's exactly right. We talked 13 about urinalysis --14 DR. BEHLING: Yes. 15 DR. NETON: -- being okay but --16 DR. BEHLING: If -- if it's --DR. NETON: But the models that are in the ICRP 17 18 default for insoluble at .002 F1 and .02 for the 19 moderately soluble material. And I don't think that 20 those were done in aqueous media. Those were done 21 based on a number of studies, including human 22 ingestion studies. 23 DR. BEHLING: Yes. I -- I remember doing a lot of 24 work in the Marshall Islands where the issue of

fallout and ingestion of fallout became an issue and

25

1 when you look at for instance some of the fission 2 products that are metal oxides in --3 DR. NETON: Right. 4 DR. BEHLING: -- in fallout, there is a quantum leap 5 between the -- the solubility based on the pH of the -- of the solution in which they are being 6 7 dissolved, and of course --8 DR. NETON: Oh, sure. Yeah, but I don't -- I think 9 that the physiologic models in the ICRP for the GI 10 tract, though, are not -- are --11 DR. BEHLING: They account for --12 DR. NETON: -- we believe them to be representative 13 of -- of the absorption (unintelligible) -- I mean 14 these were done in animal studies. Now whether the pH of a -- of a, you know --15 DR. BEHLING: As I said, it's academic if we're 16 17 talking about urinalysis because it doesn't really matter how it came -- was transferred from the bowel 18 19 or the gut into the bloodstream, et cetera. Okay. 20 DR. MAKHIJANI: Yeah, yesterday -- if I remember 21 correctly -- we agreed that ingestion is an issue 22 only when you don't have bioassay (unintelligible). DR. NETON: Yeah, (unintelligible) urine data we 23 will assume it's all inhalation, which provides a 24 25 higher estimate than ingestion pathway.

1	DR. MAKHIJANI: Right. And so and the issue of
2	ingestion in when you don't have data sort of
3	belongs in how many this question of how many do
4	you have actually that don't have
5	DR. NETON: Correct.
6	DR. MAKHIJANI: data for Mallinckrodt, and then
7	generally you're addressing it in some broader way.
8	Right?
9	DR. NETON: Correct.
10	MS. BLOOM: Uh-huh, with with
11	DR. MAKHIJANI: Is that right?
12	MS. BLOOM: TIB-9.
13	DR. MAKHIJANI: Okay. Hans, ready for more?
14	DR. BEHLING: No, I'm through here. No, I I'm
15	sure we covered the other issues yesterday so
16	DR. MAKHIJANI: Make a very, very quick check of my
17	of our review
18	DR. NETON: Uh-huh.
19	DR. MAKHIJANI: and that table just to see that
20	I've gone through it. I think we've gone through
21	everything.
22	(Pause)
23	DR. MAKHIJANI: Oh, yeah. This Revision 01 of
24	the TBD says external dose calculations on hold. I
25	think you addressed this at the meeting. Right?

1 DR. NETON: Yeah, yeah. We talked about that. 2 DR. MAKHIJANI: About coworker. 3 DR. NETON: Right. It's only on hold for people who 4 we don't have monitoring data for. So again, we get 5 back to this relevant issue --6 DR. MAKHIJANI: Yeah. 7 DR. NETON: -- how many people do we have data --8 DR. MAKHIJANI: Right. 9 DR. NETON: -- and for those who don't, we -- the 10 coworker distributions are tended to be applied. 11 DR. MAKHIJANI: Right, so we've covered that. 12 DR. NETON: I believe so. 13 DR. MAKHIJANI: I think that we are -- we are done. We -- is there a -- okay, here -- one -- one maybe 14 15 last thing is, is there a difference between how we 16 might handle infrequent incidents like uranium fires 17 compared to the more frequent ones that we kind of 18 agreed probably we've taken care of? 19 **DR. NETON:** With urine monitoring data available? 20 don't think so. I think it's -- it's -- as the 21 incidents become more and more frequent, it becomes 22 a closer and closer approximation to a chronic 23 exposure. 24 DR. MAKHIJANI: Oh, yeah. That I agree. 25 DR. NETON: But for -- but for very infrequent

1 incidents, it's -- it's equally as valid that this 2 chronic exposure scenario brackets the --3 DR. MAKHIJANI: The way you -- the way you model it 4 with the highest point --5 DR. NETON: Right. 6 DR. MAKHIJANI: -- is that how you normally do it? 7 DR. NETON: Yeah. 8 If -- if there was information that MS. BLOOM: 9 showed that you had a peak in your data somewhere, 10 you would model that. And we've been looking at 11 that both in terms of the coworker studies --12 generally it's a small chronic that you use to model 13 an incident because there's -- there's a cleanup 14 period and there's a -- there's higher exposures 15 associated with a number of things, but the -- the -- I don't know about Mallinckrodt but I think 16 17 probably my sense is that the urinalysis data is --18 one of the significant exposure scenarios is from 19 fires and they tended to be fairly routine in the 20 early days. I think that it -- it was part of the 21 ambient atmosphere in the workplace. 22 DR. MAKHIJANI: Yeah, there were uranium fires. 23 know the -- we have some idea of what the frequency 24 was.

MS. BLOOM: I mean you had some bigger ones --

25

1 DR. MAKHIJANI: So very frequent --2 MS. BLOOM: -- but I think you have some small sort 3 of routine --4 DR. MAKHIJANI: Yeah. 5 MS. BLOOM: -- incidents at a number of facilities. DR. MAKHIJANI: Yeah. The -- the reason I 6 7 raise it, in looking at my list here, is that I -- I 8 think by now we know that -- that blowouts were 9 pretty -- frequent enough that they would fall into 10 this umbrella, you know, that it will be covered by 11 a routine exposure assumption. But I don't have an 12 idea about the -- I've not seen any documentation or 13 worker evidence about the frequency of fires, which 14 is why I raise that question. It may not be an 15 issue. 16 DR. NETON: But again, I think if you go the other 17 extreme where you have very infrequent incidents, 18 then it's an even stronger case that the chronic 19 exposure will bracket that because you have a very 20 short spike in an exposure for a period of -- of a 21 day. Yeah, we're giving this chronic that brackets 22 the entire, you know --23 MS. BLOOM: I've done a lot of modeling --24 DR. MAKHIJANI: I -- I think that that's --25 MS. BLOOM: -- where they have that coworker data

1	and I've got this chronic; I've got an incident here
2	and I've got some more data out here, and if I model
3	that as a chronic exposure, it pulls it up to this
4	later, more acute type of data. It pulls up my
5	curve. If I drop out that incident data and model
6	my low level chronic and then add my short term on
7	it, you can see that the area under the curve is
8	much lower and that gives me much lower intakes.
9	DR. MAKHIJANI: Yeah. Well, Dave said that he was
10	going to send
11	MS. BLOOM: Yeah. He's
12	DR. MAKHIJANI: me something along those lines
13	MS. BLOOM: working on that.
14	DR. MAKHIJANI: actually if there were a couple
15	of examples or specifically an example
16	MS. BLOOM: Well, actually
17	DR. MAKHIJANI: with one incident
18	MS. BLOOM: the Simonds data
19	DR. MAKHIJANI: that would actually be very
20	useful.
21	MS. BLOOM: or not the Simonds, the the
22	Bridgeport Brass data that's not out yet, but the
23	graphs on that in the draft site profile show that.
24	DR. NETON: Right. We we run into this and
25	this is going to be valuable to do for several

1 reasons. We've run into this with comments on 2 Chapman Valve. There was a fire at Chapman Valve. 3 We don't know exactly when the fire occurred --4 MS. BLOOM: Actually that's in there, too. 5 DR. NETON: -- but -- but we have a lot -- we have monitoring data well after the fire. But if you --6 7 you take these chronic intake scenarios, it -- it 8 does account for the fact that there may have been a 9 fire, and we believe that we had bracketed the 10 exposure -- because you need to look at the 11 integration of the curve in microcurie days, coming 12 out as microcurie per liter days. And a chronic 13 intake scenario will -- will, at the end of the day, 14 be equivalent, if not more claimant-favorable. 15 DR. BEHLING: Is there any evidence from your review 16 of the bioassay data that bioassays were conducted 17 in response to specific incidents as opposed to 18 routine? I mean does a bioassay tell you that this 19 is a routine versus in response to a radiological 20 incident? 21 DR. NETON: I think there are codes, yeah. 22 think -- yes. I think that's true but, you know --23 MS. BLOOM: Frequently you can see the frequency 24 change --25 DR. NETON: Right.

1 MS. BLOOM: -- or you see (unintelligible). 2 DR. NETON: But that really doesn't matter too much 3 for our purposes, whether it was a routine or an 4 incident. We're going to have a curve that goes 5 through all the data points. In fact, if it were a 6 response to an incident, we would be more generous -7 8 DR. BEHLING: You would be -- (unintelligible). 9 DR. NETON: -- because we would be assuming that --10 DR. BEHLING: Yes. 11 DR. NETON: -- it was all --12 DR. BEHLING: Yes. 13 DR. MAKHIJANI: No, I -- I tend to agree, it's just 14 that it would be nice to have the actual example --15 DR. NETON: We'll try --16 DR. MAKHIJANI: -- (unintelligible) cite that. 17 DR. NETON: -- and we -- I'm hoping Dave got the 18 message. We're going to try to tie it to a -- a 19 real case --20 DR. MAKHIJANI: Yeah. 21 DR. NETON: -- so that we're not doing a 22 hypothetical anymore. 23 MS. BLOOM: Uh-huh. 24 DR. NETON: Yeah. 25 DR. MAKHIJANI: A real case. Okay.

1 DR. NETON: Okay. Well, that's great. 2 DR. MAKHIJANI: I think we are done. Thank you. 3 DR. NETON: I think this was a very good discussion, worked out much better than I thought given that I 4 5 didn't think we were quite ready. But I think we -we did get through a number of issues and I think --6 7 DR. MAKHIJANI: Yeah. 8 DR. NETON: -- worked out well. 9 DR. MAKHIJANI: Thank you. 10 DR. NETON: Okay. 11 (Whereupon, the meeting concluded at 11:00 a.m.)

C E R T I F I C A T E OF COURT REPORTER

STATE OF GEORGIA COUNTY OF FULTON

I, Steven Ray Green, Certified Merit Court Reporter, do hereby certify that I reported the above and foregoing on the day of June 2, 2005; and it is a true and accurate transcript of the testimony captioned herein.

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

WITNESS my hand and official seal this the 20th day of July, 2005.

STEVEN RAY GREEN, CCR CERTIFIED MERIT COURT REPORTER

CERTIFICATE NUMBER: A-2102