NUCLEAR REGULATORY COMMISSION



COMMISSION MEETING

In the Matter of: PUBLIC MEETING

BRIEFING BY EPA ON DRAFT GENERAL STANDARDS (OCCUPATIONAL EXPOSURES, MILL TAILINGS)

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	1	UNITED STATES OF AMERICA
	2	NUCLEAR REGULATORY COMMISSION
	3	Public Meeting
	4	BRIEFING BY EPA ON DRAFT GENERAL STANDARDS
345	5	(OCCUPATIONAL EXPOSURES, MILL TAILINGS)
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REPORTERS BUILDING, WASHINGTON, D.C.	10	Wednesday, September 3, 1980
VASIII	11	The Commission met, pursuant to notice, at 2:04 p.m.,
ING,	12	BEFORE:
G III D	13	JOHN F. AHEARNE, Chairman of the Commission
TERS	14	JOSEPH HENDRIE, Commissioner
S.W., REPOR	15	VICTOR GILINSKY, Commissioner
	16	PETER A. BRADFORD, Commissioner
EET,	17	NRC STAFF PRESENT:
H ST	18	SHELDON TRUBATCH
300 TTH STREET,	19	MR. HOYLE
	20	HCWARD SHAPAR
	21	WILLIAM DIRCKS
	22	ROBERT MINOGUE
	23	ALSO PRESENT:
	24	David Rosenbaum, EPA Dan Egan, EPA
	25	Allan Richardson, EPA Stanley Lichtman, EPA William Ellett, EPA William Mills, EPA

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PROCEEDINGS

CHAIRMAN AHEARNE: The Commission meets this afternoon to hear an eagerly awaited, highly regarded, and interesting presentation from a gentleman well known to many of us.

Mr. Rosenbaum has graciously agreed to come and talk, rather than just directly send a missive. We're delighted to hear from the illustrous representative of the EPA.

David?

DR. ROSENBAUM: Thank you.

I am delighted to be here. I always feel like I am coming home, partly, when I come to the NRC. First I would like to apologize for the slowness with which these documents are appearing. That is largely due to me.

(Laughter.)

CHAIRMAN AHEARNE: Such frankness.

DR. ROSENBAUM: They would have been out much faster,

I'm sure, if I had not been there. I hope that the slowness

is in part --

CHAIRMAN AHEARNE: You undercut my argument for a single administrator.

DR. ROSENBAUM: -- made up for by the quality of the document. That is for everybody to decide on their own, but at least we have made an effort to write them in language that people can understand; and in that way, to make the arguments clearer, so that people who want to support them or attack them

D.C. 20024 (202) 554-2345 300 7TH STREET, S.W., REPORTERS BUILDING, WASHINGTON, can at least be talking about the same thing.

Fortunately for all of us, members of my stars know a great deal more about most of these than I do. Therefore, they will do most of the calking. It is my practice to let the people who know the most do the talking, but I may break in from time to time if I feel it is productive.

Let me emphasize a few things: One, we are not sure that everything in these documents is right. We are not certain of hardly anything. This is a hard business to be certain of things in, and we have done the best we could. That's all I can say. There is still "Jom for discussion. We have not closed our minds.

In that regard, I feel very strongly that it is our job to make the decisions in so far as they are within our purview. Therefore, we look on such distinguished groups as the BEIR Committee, which we paid for, including the BEIR III Report and the BEIR II Report, and the publications of the ICRP, in much the same light as we have looked at scientific literature in general. The people who worked on those things, in general do not have access to any information that is secret or that is not generally published in the scientific literature. We try as best as possible to keep up with the literature in our field, and therefore we don't feel bound by any of these things. We do consider them seriously.

We also consider other important sources. In this

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guidance there are numbers. These numbers, I would like to emphasize again, are not set in concrete. They are proposals at this level.

When I say "this level," I mean "my level."

CHAIRMAN AHEARNE: Would you review briefly, before
you turn over to your co-workers, the procedures that you
see occurring on both of these documents?

DR. ROSENBAUM: Yes. I would be glad to.

As a start on that, I would like to say that the primary things we are going to talk about are the occupational guidance, and the uranium mill tailings standards for inactive piles. We will be willing to talk a little bit about the high-level waste, but that hasn't even gotten through me yet. And in any case, I don't think there would be time to go through all three of those very important documents with any depth in one meeting. We would be glad to come back another time and talk about high-level waste in more detail.

Neither the occupational guidance nor the uranium mill tailings have been approved by David Hawkins, who is the Assistant Administrator and my boss. And even after they are approved by him, they would still have to be approved by Douglas Costel, who is the head of the agency. It is not that they don't know what it's about -- you know, what they are about and have been briefed on them, but they have not approved them. So that this is not an official EPA position

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even as a proposal.

What has to be done, the pathways are somewhat different. The occupational guidance is guidance under the old Federal Radiation Council authority. It is not issued by EPA. EPA recommends it to the President. If the President so chooses, he issues it.

The occupational guidance which is now in effect was issued by President Eisenhower in 1960. Therefore, it is done -there is always interagency consultation on anything, but in the case of guidance there is much more, because the President will want to have the advice of all the different agencies concerned presumably before he issues something under his name.

Therefore -- and in the case of guidance, because it is guidance we are under no legal compunction to have any hearings at all, and could in fact just send the guidance to the President, and if the President so chooses he could sign it and put it in the Federal Register. In fact, that is not what we are going to do.

We plan to have widespread hearings in this case, and as in the other standards and guides we will be issuing we are going to be taking a great deal of our travel money so that we can have hearings around the country and not just in Washington. Many of these hearings will be held -- all of these hearings will be held in each place in part on weekends and at nights so that those who have jobs and who are not paid

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to go to the hearings can come and tell us what they think. There will of course also be hearings in Washington.

We have an arrangement with NRC and OSHA to have joint hearings. They actually will be paid for by EPA, and we already have a contractor who is doing the administrative workup on the hearings.

CHAIRMAN AHEARNE: That sounds like a good arrangement.

DR. ROSENBAUM: Yes. Somebody else. And we're happy to do that. So that at such time that the guidance is signed by Mr. Costel, it will be put in the Federal Register. We will have a comment period, and after the comment period is closed and we have time to digest the comments, there will be set some hearings at different places around the country including Washington.

CHAIRMAN AHEARNE: It will be signed out as a proposed document?

DR. ROSENBAUM: As a proposed document. Even after the whole process of the hearings and we have considered the comments, and Mr. Costel signs it again, it will still only be a proposal to the President. It will be a recommendation to the President that he sign it.

I would hope, if everything goes smoothely, that we could have hearings early in the spring. I don't think, given the procedures that have to be gone through, that it is

realistic to think that we could have them much earlier than that. Therefore --

CHAIRMAN AHEARNE: Which year, David?

DR. ROSENBAUM: This year -- next year; next spring.

That's a reasonable comment, given my record.

CHAIRMAN AHEARNE: No, not given your record; given the record of this issue.

DR. ROSENBAUM: Well, I see no reason why we shouldn't be able to have them next spring, at the moment, unless things come up that I don't now know about.

And if that happens next spring, and everything else goes smoothely, we should be able to get them to the President by perhaps a little more than a year from now. I would say sometime the fall after this fall we're entering.

What the President would then do, I have no idea.

But I think at that point, ordinarily the President asks for advice from other affected agencies, and perhaps from the public. And now that we have consolidated government by forming several new independent radiation bodies such as the Radiation Policy Council, some of these things may be referred when they get to the President, to those bodies, and that may take longer -- therefore, the process of the President signing might take longer than it would otherwise me; I don't know.

As far as the uranium mill tailings goes, in that

20024 (202) 554-2345 D.C. 300 THI STREET, S.W., REPORTERS BUILDING, WASHINGTON, case it is a standard--it doesn't have to go to the President-issued by the Administrator of EPA. We have already put in
the Federal Register the cleanup standards for the inactive
piles.

Me have now sent to Mr. Hawkins the rest of the package, which is the disposal of handling the piles themselves for the inactive piles. And if all goes well, we might be able to get that proposed in the Federal Register within a month or so. I think that won't take very long -- I can't guarantee it.

And then we have to have hearings, and go through the normal process of making a standard. We have to answer the comments -- and I don't have any way to guess how voluminous the comments will be.

CHAIRMAN AHEARNE: Did you say you would have hearings on that, also?

DR. ROSENBAUM: Yes. In the same way, we'll have them around the country, particularly in appropriate spots where they have uranium mill tailings. We probably won't have them in Boston.

We have contractors already working on both of those -- administrative contractors -- to do the administrative work to set up the hearings. And we will in these hearings not only hold them in cities, but we have given the contractors an advertising budget so that the people in that

area will be well aware, through the public press and the other media, that the hearings are being held and that we are indeed interested in their opinion.

Is that adequate for what you meant for the procedures?

CHAIRMAN AHEARNE: That's excellent.

COMMISSIONER HENDRIE: Dave, before you go on, the mill tailings standard is cited in the mill tailings bill of a year-odd ago. And as I recall the language of the bill, it becomes a controlling general standard for us, I think, doesn't it?

DR. ROSENBAUM: Well, you're dealing with the active piles, I believe, primarily; and the standard we are putting out so far is the inactive piles.

COMMISSIONER HENDRIE: I see; okay.

DR. ROSENBAUM: The inactive piles were due by law to be promulgated last November.

COMMISSIONER HENDRIE: But in due time, there will also be some active piles standards?

DR. ROSENBAUM: Yes.

COMMISSIONER HENDRIE: And those we follow because the mill tailings law says EPA does the general ones, and you do the license conditions to conform.

DR. ROSENBAUM: Yes.

COMMISSIONER HENDRIE: But on the occupational

Branch and are controlling for all of the Executive Branch agencies. Nominally as an independent agency, it is not -- well, I am not sure whether we are bound by that or not.

If I look to your end of the table, Sheldon --

MR. TRUBATCH: We have stated publicly in the past -the Commission has stated that it would consider itself bound
to follow the standards. That doesn't really answer your
question as to whether we are legally bound.

COMMISSIONER HENDRIE: Well, I know what we're going to do, we're going to follow them. But I was just curious --

CHAIRMAN AHEARNE: Joe, one of the reasons we are going to conduct hearings is to develop our own understanding of the issues.

COMMISSIONER HENDRIE: Well --

CHAIRMAN AHEARNE: But perhaps Bob Minogue might want to answer your question as to what the Standards Office plans to do.

Don't worry, David, we will get to your presentation.

DR. ROSENBAUM: We are here to serve.

MR. MINCOUE: Guidance of this type has to be implemented by a specific agency. So the activity that we would look to do within NRC would be more aimed at implementing standards to take that guidance and apply it to what we regulate.

COMMISSIONER HENDRIE: Yes.

MR. MINOGUE: On the other question, I believe that the matter of conforming to FRC guidance is one of policy. What the law is on the matter I'm not clear, but it certainly has been the past policy to conform as if it were mandatory.

COMMISSIONER HENDRIE: I was just curious as to how the law lay on the matter. There is the general proposition of are we compelled by Executive Orders, because this would be in effect an Executive Order of the President. I think in general we claim not to be.

On the other hand, in this radiation protection matter, there is Reorganization Plan No. 3 a number of years ago which split off the responsibility for establishing the general health standards from a radiation standpoint in EPA, and I wonder if there isn't a tie that binds legally back through Reorganization Plan No. 3.

I was just curious --

MR. MINOGUE: I think it is quite complex, and I am not aware that any detailed analysis has been done, because it is a matter of policy that we have always conformed to FRC guidance. I really don't know the answer to the legal question.

CHAIRMAN AHEARNE: Howard?

MR. SHAPAR: I believe the answer is that if this is a Federal Radiation Council function -- which I believe it

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is -- then that function stems from Section 274 of the
Reorganization Act -- I mean, of the Atomic Energy Act. It
is specifically mentioned in our Act. Now the Federal Radiation
Council was transferred by Reorganization Plan No. 3 to the
EPA. I think the answer to the question is: If that function
is performed pursuant to Section 274 of the Atomic Energy
Act and the recommendations made of the President, and the
President signs it, it is binding on everybody including the
NRC.

COMMISSIONER HENDRIE: You have more muscle than you thought you had, Dave.

(Laughter.)

COMMISSIONER HENDRIE: So what do you want us to do?

DR. ROSENBAUM: Do you have any extra money?

(Laughter.)

DR. ROSENBAUM: Just let me introduce the people who are going to give most of the briefing. First of all, let me say, please feel free to interrupt at any time. We have really set up in such a way that we would give some of the information in response to questions.

It is not only a question of questions, we would very much welcome your suggestions. We don't have all wisdom, and we would be very happy to hear your ideas and suggestions.

The last thing I want to say is about the uranium mill tailings standard. That is, that we were bound by law to

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issue on May 8th -- to promulgate last May 8th, the standards for active piles. We have just begun to work on them. There may be some feeling on some people's part that these standards which we are now proposing for inactive piles will set a procedent for the active piles, and that we will necessarily make the active piles conform to them in some way, or be more stringent or less stringent.

All I can say is that that is not true. We intend to look -- the active piles are quite a different situation, and we intend to look at them de novo, and to try to do as best as we can to make sensible standards.

It may indeed come out that they will be very close -
COMMISSIONER GILINSKY: Would 't you expect that

you would do better on the active piles than on the older

ones?

DR. ROSENBAUM: What does "better" mean?

COMMISSIONER GILINSKY: To be able to enforce a stricter standard?

pr. ROSENBAUM: I don't know, because we have not just gone through the staff work. I don't know. There is no point in my speculating. I don't have any basis on which to judge. I am just saying that philosophically we do not feel bound in any way to do anything but the best we can; and that these standards are not going to be sort of based on the inactive standards. It is a different kind of problem

with different opportunities to do things, and different financial responsibilities. We are just going to look at the problem and see what we think is in the public interest.

COMMISSIONER GILINSKY: Fine.

DR. ROSENBAUM: It might come out that way, but it isn't prejudged to be that way.

The first person who will begin the briefing is

Mr. Allan Richardson, somewhere. Why don't you come on up

here. He is head of the General Standards Branch and who has

been in charge of developing both the guidance for occupational

activity, and also for the uranium mill tailings standard.

Is somebody manning the slides?

DR. RICHARDSON: Yes.

(Slide.)

I have put this slide up, which you don't have, so that I can tell you what I am not going to talk about in my prepared presentation. It will give you some idea of what we are prepared to talk about, if you want to ask extra questions.

I am going to skip Items 1, 3, and 4, which are rather major items, and get directly to talking about the recommendations -- which is what the balance of the list consists of.

A lot of the recommendations are based on findings from those -- especially Items 3 and 4 -- and I will have to

ask you to take those on faith, and then we can go back to them later. So can we go to slide one?

(Slide.)

This is a summary of the proposed changes. I will just read through it quickly, so that you will have a feeling for what we are up to; and then when we go to each individual one you will have some idea of the total context that we are talking about.

In any system of radiation protection, there are three basic principles. The first is that the exposure needs to be justified. The second is that you need to minimize the exposure. The third is that you impose some kind of a limit on individual risk, because the first two principles don't automatically ensure that. That is what these first three requirements do.

We are not proposing very much in the way of changes in those. The 1960 guides required that exposure be justified; it's not always clear how that is done, but it is still a requirement.

We are adding some language which points out that alternatives to radiation exposure should be considered explicitly. That was implicit before.

Under the "minimization of exposure," which has been, in some radiation protection circles, called the "optimization of exposure," which sometimes has an unfortunate

inverse connotation. That was required by the 1960 guide.

That is what we have known through the years as ALAB or ALARA.

We are now explicitly saying that this is an operation which is carried out on the collective dose, which is the only way you can sensibly do it.

Item 3 is the limitation of exposure to individual workers. It is split there into three parts. We are proposing some changes there. We are proposing to abandon the 3 rems per quarter, and 5(N-18) cumulative limits, which have been broadly interpreted as being 5 rems a year on the average; and replacing it with a single 5-rems-per-year limit -- or proposing to do that.

DR. RICHARDSON: Later on I will talk about, there will be an admonition to keep lifetime doses within a limit, a 100-rem limit, but that comes in a different part of the

CHAIRMAN AHEARNE: Without any cumulative?

guidance.

The partial body limits, just as an aside, all exposure is of course not to the whole body. There are cases in which you breathe or ingest radionuclides and get higher doses to individual organs, so that there is a need for limits for the individual organs, some of which are more radio sensitive than others are.

In the past, this has been done by setting limits for individual organs, and limiting the dose to the organ

D.C. 20024 (202) 554-2345 300 7TH STREET, S.W., REPORTERS BUILDING, WASHINGTON, which came closest to its limit, and ignoring the rest of the organs. It's called the so-called "critical organ" approach.

We are proposing a change to something called a "summation of risk" approach, in which each of the organs is given a weight which is proportional to the fraction of the whole-body risk which comes from that organ.

We are changing some of the names of the organs that are listed specifically, and dropping some that don't seem to be needed any longer. I would say that on the average the -- and we will talk about this in more detail later; this is a rather controversial part of the guidance -- that this approach leads to more restrictive limits, and a far more rational system. I won't say any more about it now.

Another aspect of current guidance is that internal and external exposure -- "external exposure" being governed by the whole-body limit, the 5-rem limit; and "internal exposure" being governed by the individual organ limits -- have been limited independently. So somebody could get 5 rems of whole-body exposure, theoretically, and also get the lung limit.

We are proposing now, since we now have weighting factors for the individual organs under the summation-of-risk approach, that the sum of the risks be limited from both internal and external exposure.

Item 4 there is new. In the past, there have been

no specification and guidance of minimum radiation protection requirements. We are proposing that there be three ranges of increasing stringency of instruction on the risks of radiation and radiation protection principles, on supervision of radiation protection and the application of radiation protection; of monitoring and recordkeeping -- and that is where the lifetime dose gets covered. I will talk about that in a lot more detail in a few minutes.

Item 5, we are recommending to the agencies -proposing to recommend. Pardon me if I don't always say
"proposed."

CHAIRMAN AHEARNE: That's an understandable problem.

DR. RICHARDSON: We are proposing that the regulatory agencies establish limits lower than 5 rems for specific classes of workers or work situations when they think this is appropriate. This was not done in the 1960 guide.

Item 6 under the intake guides, we are proposing a methodological change, a minor one here, from radioactivity concentration guides, to radioactivity intake factors. I am not going to talk about that very much. It is tied up in the means for carrying out Recommendation 3(b).

The exposure of miners, we are proposing no change.

Item 8 is a potentially extremely controversial one.

We are proposing that exposure of the unborn be limited. We are proposing four alternatives for public consideration, and

D.C. 20024 (202) 554-2345 300 7TH STREET, S.W., REPORTERS BUILDING, WASHINGTON, we will talk about those in a few minutes.

Item 9, and the last one, the only change -- the old guidance permitted agencies to exceed the guidance. We are now making it explicit that they ought to tell people why they are doing it when they do it.

The next slide, please.

(Slide.)

DR. ROSENBAUM: Let me say one thing about the exposure of the unborn, in utero exposure. In the 1960 guidance, there were two classes of people, or "pre-people," who were protected. In the first place, the people themselves, the workers, were protected against cancer, supposedly, by limiting the dose which they could get. But in addition, restrictions were put on a gonadal dose. The reason for that is to protect against mutations, and therefore the protection that is being offered is not for the worker, but for people not only yet unborn but yet unconceived.

Any time you put restrictions on a worker, it restricts his ability to earn a living. Therefore, the precedent was set already in 1960 that workers could be restricted not for their own benefit only, but for the benefit of people yet to be born.

However, there was a class which wasn't covered at all. That is, people who were conceived but not yet born.

We know a great deal more about health effects now in 1980

than we did in 1960. The largest single source of data on radiation health offects is the Japanese survivors. The data from the Japanese survivors shows little, if any, evidence of any genetic damage at all. I don't mean by that to suggest that there's any proof that there isn't any, but certainly not any proof that there is.

On the other hand, it shows very high effects on children who were in utero at the time. Perhaps the most prominent of those effects, but certainly not the only one, is microcephaly. "Microcephaly" is literally very small head size, and it is very, very highly correlated with severe mental retardation -- very severe mental retardation.

There are other effects from in utero radiation -childhood lukemia, for example, after children are born, and
other sorts of genetic effects. The summation of these
effects, it seems from the data, seems to be a very much
larger problem than the genetic effects which seemed very
serious to the people in the late '50s who were drawing up
what became the 1960 guidance.

So when I came to this job a little over a year ago, this was not part of the occupational package, but was scheduled for a further action after this package. It seemed to me, in conscience, that we ought to bring it cut in the public and have a public debate about what, if anything, ought to be done about it. What we have done is to propose -- which

we will show you -- four alternatives. These aren't the only possible alternatives, and no doubt other people will suggest other reasonable possibilities at the hearings.

That's all I had on that.

CHAIRMAN AHEARNE: Go ahead, Allan.

DR. RICHARDSON: Let me move on now to some considerations that shaped our thinking about how to put the guidance together as a whole, not just the question of whether the number should be "5 rems," or "9," but how should the guidance operate.

There are a couple of striking facts about radiation exposure of the work force that are important, I think. One of them is, we did a survey of radiation workers for the year 1975. I think we looked at something like 350,000 records out of a little over a million estimated numbers of workers.

We find that about 95 percent of workers get less than 500 millirems -- less than one-tenth of the maximum permitted by the guide. We also have concluded, after talking to -- I haven't put up a slide which shows all the agencies on our interagency working group and how this thing has moved forward, but we have a group of about 10 agencies that have been working on this for a number of years. We have become convinced that there are justified tasks that exist that require doses on the order of a few rem. All right, that's a dichotomy.

Another observation, after looking at the risks associated with average exposures, average exposures by the way for the roughly 1.2 million workers in 1975 were about 120 millirem, a very low number.

The risk that is associated with that is quite small.

It is actually less risk than the risk of accidental death in the safest of all trades. We can show you some information about that later.

On the other hand, the maximum permitted lifetime radiation risk -- that is 5 rems per year from age 18 to age 65 -- is not small. It is on the order of something between 5 and 10 percent chance of a premature death due to cancer, radiation-induced cancer. It is an exceptional circumstance. I don't think any of us could find a worker who had gotten 5 rems every year of his working life, but it is theoretically allowed by the guide.

It i a risk which is from one to three times, depending on which risk model you use, higher than the risk of accidental death in the most hazardous of occupations --

DR. ROSENBAUM: The average.

DR. RICHARDSON: The average risk. That is, miners, construction workers, agricultural workers. So it is something which has to be dealth with.

Could I have the next slide, please? (Slide.)

COMMISSIONER GILINSKY: "Average" means simply you are averaging over a large group?

DR. RICHARDSON: Averaging over all miners, averaging over all agricultural workers, and so on, and comparing that to the maximum risk permitted under the guidance. That's something like 235 rems over a lifetime.

DR. ROSENBAUM: We don't have the information to compare the highest possible risk under the guides with the most dangerous miners' occupations, or the most dangerous -- because there aren't such statistics.

COMMISSIONER BRADFORD: But how meaningful is it to say that the highest possible exposures exceed the average risk of death than others? I mean, the average itself obviously is made up of the number of highs and lows.

DR. ROSENBAUM: It is not all that meaningful, I quite agree. It certainly hasn't been a very important part in the decision-making. It is just an attempt to give some perspective in terms of other industries. It would be much more sensible to compare the average nuclear risk with the average in the most hazardous industries, and the highest possible nuclear risk with the highest risk for miners and people like that; but we simply don't have the data for the higher risk. There are no statistics compiled, so there is no way to do that.

DR. RICHARDSON: It may be more pertinent to look

at the absolute risk, which is on the order of 5 to 10 percent, or I guess it is actually 4 to 7.

COMMISSIONER GILINSKY: Well, it is not an irrelevant number for someone who is engaged in this occupation -
DR. ROSENBAUM: No, it's not irrelevant at all.

COMMISSIONER GILINSKY: -- and might be subjected to these sorts of numbers. I don't know what the risk is.

DR. ROSENBAUM: It is very important.

DR. RICHARDSON: All right, what do we conclude from that? We concluded that individual and collective risks should be minimized. That's a motherhood statement. They should be limited to values, at worst, no greater than those from other occupational hazards.

In order to do this, we concluded that we should design the guidance so that workers are permitted doses up to 5 rems, which appear to be necessary in some cases, only when clearly necessary; and that there be some kind of administrative controls -- some kind of a structure -- that would keep the vast bulk of workers operating under limits which are, in effect, for them much lower.

Now there are different ways to do -
COMMISSIONER GILINSKY: May I ask a question at this point?

DR. RICHARDSON: Yes.

COMMISSIONER GILINSKY: I know you said you were

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going to skip the important background sections, but -
DR. RICHARDSON: You wanted to talk about that some
more?

COMMISSIONER GILINSKY: I would like to ask one question.

DR. ROSENBAUM: We are willing to go into anything you want.

COMMISSIONER GILINSKY: I gather this is all based on a linear model?

DR. ROSENBAUM: Yes, sir.

COMMISSIONER GILINSKY: Yes.

COMMISSIONER GILINSKY: How does one get from
limiting risks to annual dose limits? Why would one impose an
annual dose limit, as opposed to a cumulative dose limit?

DR. ROSENBAUM: Why would you want to impose it?

DR. ROSENBAUM: First of all, imposing an annual dose limit automatically imposes a cumulative limit.

COMMISSIONER GILINSKY: I understand that.

DR. ROSENBAUM: It's true that it makes a difference when you get the dose, because if you get the dose -- all this is very model dependent. Ordinarily peopl consider most cancers to have a latency period sometimes of 10 to 15 years say for lung cancer.

COMMISSIONER GILINSKY: I guess I was trying to get a better idea of what your model is.

DR. ROSENBAUM: Well, the point is that if someone gets -- if someone would have had just a lifetime dose and they were to get it at age 20, it would be much more serious than if they would get it at age 60, for two reasons. One, if they were to get cancer from it, they would be likely to get cancer earlier, and therefore lose much more of their life.

Two, the chances of somebody getting it when they were 60, if there's a 10- or 15-year latency period, there's quite a sizeable chance that they will die from other causes since most of it would show up 20 or 25 years later, before they ever got the cancer, and therefore the chances of them getting cancer from it are much less, at all. And if they do get cancer from it, then they lose much less of their life. So on both accounts, it makes a great deal of sense not just to have an overall lifetime limit, but to have a yearly limit.

Is that a satisfactory answer?

COMMISSIONER GILINSKY: Yes.

DR. RICHARDSON: You mentioned in part of your question: How do we calculate these risks? We will talk about it more later, but briefly we use a life table analysis and we take into account the latency period, and the risk of competing death, and the length of the expression period for cancers; and we have used the BEIR I risk estimates up until now.

COMMISSIONER GILINSKY: But even taking those scrts of things into account, it wouldn't really make a great deal of difference if you doubled the dose one year and had zero the following year.

DR. ROSENBAUM: It wouldn't make so much of a difference if those two years were together. In other words, it wouldn't make much difference if you had a two-year limit instead of a one. And indeed, that's sort of the thinking that we did to do away with quarterly limits. At the present time, there are quarterly limits of 3 rem per quarter, which allow 12 rem a year.

people did in Hiroshima and Nagasaki, then you may have a very serious dose-rate effect. That's quite possible. But there is no evidence that I know of that there is any difference in dose-rate effect between spreading it out over three months or a year, or two years for that matter.

CHAIRMAN AHEARNE: At these low levels?

DR. ROSENBAUM: At these low levels, yes. So the year seems to be a convenient period for administration.

COMMISSIONER GILINSKY: Also, we have been using it for a long time.

DR. ROSENBAUM: And it seems -- and for the reasons

I've just mentioned to you, one doesn't want to spread it out

over too many years, because the risk does change from year to

year as you get older.

DR. RICHARDSON: Well, moving on, the guidance proposes two different mechanisms for basically imposing ALARA on doses above and beyond the 5 rem limit. One of them is a tiered system of minimum radiation protection requirements, which I will talk about next; and the other is an admonition to the regulatory agencies to set limits that are less than 5 rem when it is justified to do so.

We chose to leave -- one of the other alternatives would be to set such limits ourselves as part of the guidance for different classes of workers. We decided that that was not an appropriate thing to do in guidance; it should be left to the regulatory agencies.

I have talked about limiting excessive lifetime dose, and I will talk more about it in the minimum radiation protection requirements. We have also concluded that there is no need to continue the 12-rem annual dose limit, which is the 3-rem quarterly Limit as it is now expressed.

Next slide, please.

(Slide.)

All right, these are the first two recommendations and I think they speak for themselves. The first requires for justification of exposure; and the second, for minimization of the collective dose.

Next slide, please.

25 .

(Slide.)

This is a summary of the minimum radiation protection requirements. I just put it up here so you could get an overview of what they contain. We are proposing three ranges, A, B, and C, that are respectively up to one-tenth of the radiation protection guide, from one-tenth to three-tenths, and from three-tenths up to the radiation protection guide.

In the first range -- there are four different types of things covered by the requirements: education requirements, supervision requirements, monitoring, and records.

Education requirements are what is appropriate for the range. Supervision is interesting. There we propose that in Range A only through generic requirements. An example of workers in Range A are dental technicians or transportation workers. Packaging requirements on radioactive materials are a generic requirement for dose limitation. Shielding requirements and the like on X-ray diagnostic machines are an example of generic requirements.

At the level of Range B, we propose that there should be professional radiation protection supervision on the job.

And at the level of Range C, that this be extended to individual tasks for high-dose jobs.

CHAIRMAN AHEARNE: Now over in the far right-hand lower column, you mention you are going to include Range B doses in C.

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(202) 554 2345 20024 D.C. S.W., REPORTERS BUILDING, WASHINGTON, 300 7TH STREET, DR. RICHARDSON: Once a worker has been exposed in Range C, we propose that his Range B doses from that time forward only be kept track of.

I think the rest of that is self-explanatory. Can we have the next slide.

(Slide.)

Here is the statement of the minimum radiation protection requirements for Range A. I don't think I need to say very much about them.

The monitoring requirement there is simply a requirement to know enough about the exposure conditions to ensure that the doses are in Range A.

Next slide, please.

(Slide.)

Range B has two important requirements. First, that the professional radiation protection supervision be present. Second, that individual monitoring and annual recordkeeping take place.

CHAIRMAN AHEARNE: Now by your "professional" provision, do you go into further detail, or do you intend to, as to what that would mean?

DR. RICHARDSON: No. We've avoided doing that.

There are going to be some medical applications where there are very few employees where it would not be appropriate to hire a health physicist, and where there is going to have to

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be provisions for either additional training of employees in health protection or consultative services. But there are only approximately 2 to 3 percent of all workers in Range B.

Next slide.

(Slide.)

Range C, which is the highest doses, which are a little bit less than 2 percent of all workers, calls for close supervision on a task-by-task basis of high-dose jobs, and lifetime dose records once you enter the range.

Finally, that every reasonably achievable effort be made to maintain doses to less than 100 rem over a lifetime to workers.

Next slide.

(Slide.)

And here is the recommendation that agencies establish lower limits when it is appropriate. And a clarification in the last sentence, that the ranges specified by A, B and C are not intended to specify those lower limits.

Next slide, please.

(Slide.)

Partial body exposure is a complex subject, and I don't want to take any more of your time with it than is appropriate and necessary. There are several -- there are many issues. I've put four of them up here. I've already mentioned the question of whether internal exposure should be

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limited by individual organ limits, or by the sum or risks to all organs.

There is the further issue of whether somatic and genetic risks should be treated separately or together. For instance, in ICRP current guidance they are treated together; we have not done that.

CHAIRMAN AHEARNE: Are you in debate with ICRP on this?

DR. ROSENBAUM: Let me answer that. We don't "debate" with ICRP. They have a different role to play than we. We read what they do and take it into consideration.

CHAIRMAN AHEARNE: But you don't attempt to change what they do?

DR. ROSENBAUM: Well, we have no mechanism to change what they do. We don't have anybody on ICRP, and in fact when we were offered -- essentially offered -- the ability to put somebody on the ICRP, I decided we couldn't possibly do it because of limitations of travel, among other things, which we don't have the luxury of either staff or travel money to do that sort of thing.

DR. RICHARDSON: I have informally discussed these proposals as they have been developing with the President of the ICRP, and they have no difficulty with them, for what that's worth. They believe that they are consistent.

All right, I think the rest of that speaks for

itself, and then we have already talked about it a little bit.

Can we have the next slide?

(Slide.)

This is a complex slide, and I apologize for it.

This shows most of the system for limitation of individual risk and partial body exposure.

There is under "A," a list of individual organ limits.

We have chosen to continue the 5 rem limit to the gonad and

to the lens of the eye, because there seems to be no good

reason to change it.

We are proposing that the limit to the hands be lowered to 50 rem. And we are proposing that non-stochastic and non-lethal cancer risks be limited by a single limit for any organ of 30 rem.

We are proposing that the risk from partial-body exposure of individual organs be also limited -- and this will usually be controlling -- by the sum of the weighted risk to the individual organs, with the weight taken in proportion to their contribution to whole-body cancers.

I have a table of those weights. It is basically the same set of weights that the ICRP uses, with the genetic weight taken out and the rest of them renormalized.

And then finally, the proposal at the bottom that internal and external doses be -- that the sum of the risks be limited.

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Next slide, please.

(Slide.)

The Unborn. Here are four of the considerations that affect the choice of recommendations, the possibilities -- the first being whether they should be voluntary or mandatory limits. That is intimately tied up with the right-to-work and equal opportunity questions.

The next question, which is a difficult one, is whether the guidance should apply to a period of known pregnancy, to periods of suspected pregnancy, or simply to all fertile women. It is particularly important, because the most sensitive period is early in pregnancy when, although pregnancy may be suspected, it probably is not known.

That leads to another question. That is: Should the limitation be expressed in terms of something like some short-term dose rate, like a monthly dose rate, which would serve to protect the unknown fetus? Or should it be done with an annual dose as all the other limits are?

And then finally -- and this is the equal opportunity question -- should the limits for male and females arbitrarily be made the same?

Next slide, please.

(Slide.)

This lists four alternatives. The first two are voluntary; the second two are mandatory. Of the first two,

the first one is an annual limit -- I'm sorry, it's a limit during any known or suspected pregnancy. Since it is a voluntary limit, it can be applied to a suspected period of pregnancy also.

The second one is the same, but it adds a voluntary limitation on dose rate.

The third one is essentially the second made mandatory.

And the final one is the least restrictive limitation that we could think of that applies equally to male and female, and provides the same level of protection to the unborn.

CHAIRMAN AHEARNE: But as you say, it essentially greatly reduces the maximum exposure.

DR. RICHARDSON: It would affect the whole guidance.

DR. ROSENBAUM: Let me talk to that point a bit, because I think it is important. I think the fourth choice there, while it is there, it really begs the question. The real question is: Because women bear children and men do not, what does that imply about setting a different level of dose limits for women, or at least women who can bear children, than men? And lowering the whole structure doesn't change the fact that whatever level you set the structure, at least within the models that we use, the risk will still be greater for the children.

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By the way, I should point out that the risk to the unborn is far greater than the risk to the mother, or to a male in the same situation, far greater. The most sensitive time of a human being's existence to radiation is early in its fetal development.

CHAIRMAN AHEARNE: By "early," is the Japanese data sufficient that you can pin that down? Are you talking about months? Weeks?

DR. ROSENBAUM: Yes. We would be happy to -- Bill, could you bring up -- I'd like to interpose that slide.

This is Dr. William Ellett, head of the Bioeffects Branch.

Do you have the slide there with you?

DR. ELLETT: Yes. Could we have that second stack of slides, the one marked 8-A?

(Slide.)

This is the record for the single endpoint microcephaly, small heads, that Dr. Rosenbaum mentioned, 52 cases observed at Hiroshima.

DR. ROSENBAUM: That's the right slide.

DR. ELLETT: It is the right slide, but just --

DR. ROSENBAUM: Upside down.

(Discussion off the record.)

DR. ELLETT: Let's get X-Zs, first (indicating).

This (indicating) is increasing dose, going back in the chart.

DR. ROSENBAUM: In red.

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DR. ELLETT: This is actually the air dose, if you will (indicating). This is the percent in each interval of time of gestration here. These are the weeks, post-conception, from zero to 36-plus.

of newborn that showed microcephaly (indicating). It turns out that it is very much concentrated in exposures to the curve of somewhere from the 5th to the 11th week. There is a window for the embryo that is starting those developmental processes that lead to the brain and the skull.

The small skull is probably more a result of the small brain than a direct end point. Both the BEIR III report has this, and we did it ourselves independently. We looked at the dose response for this most sensitive period from the 5th to the 11th week. Could we have that next slide?

DR. ROSENBAUM: Before you go on, I just wanted to point out one thing about that: That a number of those humps go up to 100 percent, a fairly wide range. That is to say, it no longer is a probablistic thing at all, but 100 percent of the children were born with microcephaly and therefore almost certainly severely retarded. "Severely retarded" means something like you can't write your name when you are 18. It is really severe.

DR. ELLITT: And they didn't look for "less severe" degrees.

DR. ROSENBAUM: The Japanese only monitored very severe retardation.

DR. ELLETT: Could we have Slide 8-B, please? (Slide.)

Could we have that twisted 90 degrees?

DR. ROSENBAUM: I knew it.

(Laughter.)

CHAIRMAN AHEARNE: That's Murphy's law.

(Laughter.)

DR. ELLETT: Air dose is here (indicating), but pay more attention if you would, please, to these doses here (indicating). This is the interuterine dose (indicating), the dose to the fetus (indicating), and this is the frequency -- these are different dose ranges here (indicating); it's a bar graph, really. These are dose ranges.

This (indicating) is the frequency of the cases.

Now once you got up to something like 32 rad gamma, 3-1/2 rad

neutron, you had 100 percent. This is for the most sensitive

period.

Here (indicating) are the controls. They were seen in both studies, about 3, 4 percent down here (indicating).

This is a little bit of data here for this first step, 0 to 10, but the average doeses there are quite small. It's 1.2 rad gamma, .1 rad neutron. The neutron doses were not particularly high inside the body. This is something that we had missed

before in previous analyses.

This first step may not be statistically significant.

Actually you get 11 percent incidence here. If it had been 12,

it would have been statistically significant. It is borderline;

it's pretty close.

Here (indicating) it isn't significantly above expected all the way out. What is a little bit spooky about this is, if you draw a line here (indicating) from the zero dose non-exposed controls to up here where they had the 100 percent, it falls awfully close to the middle of these steps.

Now a linear response isn't what you particular expect for this sort of thing. In animals they see it sometimes; more frequently they see a non-linear response. It depends on the end point you're talking about at the time of exposure. For this data and some other things, we have some real concerns about in utero exposure -- more so than we have had before.

The BEIR I Report is rather sanguine about this, as long as you're under 5 rem per year. The BEIR III Report is not that way at all. It says: Caution, except when mother's health is involved for exposures more than a few times background.

COMMISSIONER GILINSKY: What is the date of this work?

DR. ELLETT: It's in your handout. The update of

that is in a Terogenics -- Teretology, and --

(Pause.)

DR. ROSENBAUM: While he is looking that reference up, let me say a couple of things about this.

One, you can see why we are concerned. At doses in which people would have only a very small chance of cancer, in utero exposure means 100 percent chance of very severe mental retardation, so it's a very severe effect.

I think that there is just no statistical, or otherwise, doubt about this data at all. The one difference that one can say is that the Japanese exposure took place almost instantaneously, and therefore it may be to some extent doserate dependen. We have looked at other data from animals and other things in making our own guidance; but in looking at this data myself, I couldn't see how we could ignore bringing up this issue in the occupational guidance.

I don't consider it a matter of women's rights, at all.

In all fairness, half the children being born with microcephaly

and retardation will be women. So it's not the issue at all.

DR. ELLETT: The data itself is from Teretology.

It's from a paper by Miller and Mauhill, Occupational Institute of Health, 1976. They have a whole series of papers going back to at least 1972 and 1968 on this topic.

COMMISSIONER GILINSKY: Why did it take so long to understand this, or at least observe it?

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DR. ROSENBAUM: I think it had been observed scientifically. I don't know why it took so long to be brought up publicly. I just did it as soon as -- I just put it into the process as soon as I saw it. I had a number of --COMMISSIONER GILINSKY: What about the scientific

papers in 1976?

DR. ROSENBAUM: I don't know why. That was the latest. They go back quite a long way.

DR. ELLETT: I think the real difference was, when we started the BEIR Study, we insisted that the BEIR people use the organ dose rather than the air doses for this. At Oak Ridge they did a magnificent job in getting doses to different organs of the body, including what the interuterine dose was.

These doses, we always said: Well, gee, the neutron dose is pretty high. This must be due to neutrons. And that is pretty much stated in the papers going back to '72.

The problem is that the neutrons were much more absorbed than the gamma rays. The neutron doses were only one-thirteenth of the gamma doses. So it is very hard now to abscribe all this damage to just the neutron component. I think that is where we misjudged the thing.

DR. ROSENBAUM: We have also --

DR. ELLETT: In looking at the dose in air, and not the dose at place of entrance.

DR. ROSENBAUM: We have also done careful statistical analysis of the difference between Hiroshima which had such a large neutron component, and Nagasaki, which was largely gamma, to see if in fact most of it came from the neutrons. That's not the way it looks when you do a careful analysis.

COMMISSIONER GILINSKY: Well, are these expressed -Oh, they're expressed in rads. There isn't any factor that
takes account of the different --

DR. ROSENBAUM: No, no. You could adjust them by whatever RBE, multiply the neutron. The neutron dose is given explicitly. You could convert it to rem by multiplying the neutron number by any Q that you like.

COMMISSIONER GILINSKY: It looks as if the gamma and the neutron components are roughly comparable in rem.

DR. ELLETT: Yes, roughly I think so.

DR. ROSENBAUM: If you look at the difference between Hiroshima and Nagasaki as an experiment, the different effects, it is difficult to make a case that it's primarily neutrons. I can just tell you that from the data. That doesn't prove anything, but it is very difficult from that data -- which was most of what we have -- to make such a case.

Let me point out that this same issue has been brought up very strongly in the medical field partly by the work of Alice Stewart, such that 10, 15 years ago women in their first pregnancies were routinely X-rayed to see the size of

20024 (202) 554 2345 WASHINGTON, D.C. BUILDING, 300 TIM STREET, S.W., REPORTERS the pelvic canal and the position of the baby. It was a routine thing, without any second thought about it. Now, women who are pregnant are never X-rayed unless there is a compelling reason to do so, even though the doses to the fetus are quite small and much smaller than these that we're talking about here, and also that the doses are given -- and the X-rays are almost never given in the most sensitive period, but are often given in the 7th or 8th month where the sensitivity is much less.

Nevertheless, the medical community has decided that women who are pregnant should never be X-rayed unless there is a compelling medical reason to do so, and I think it is part of the same thing.

COMMISSIONER GILINSKY: Wouldn't there be records for women who had been X-rayed in the early months of pregnancy that one could use to develop a corresponding set of results for the U.S. population?

DR. ROSENBAUM: Let me discuss that. Alice Stewart has done some of that in Britain. I have not gone over her papers myself, and therefore I don't know how good they are. But certainly they have had an enormous effect.

We are negotiating a contract right now with the Mayo Clinic which has an enormous --

CHAIRMAN AHEARNE: The entire health records -
DR. ROSENBAUM: -- the entire health records of

almost a million people. What's more, they have birth-to-death

(202) 554-2345 D.C. WASHINGTON. BUILDING, 300 77 H STREET, S.W., REPORTERS records of a large part of it. People are born in the Mayo Clinic, their pediatrics is done their, and they grow old and die in the Mayo Clinic.

COMMISSIONER BRADFORD: And they get out.

DR. ROSENBAUM: Excuse me?

COMMISSIONER BRADFORD: They get out.

(Laughter.)

DR. ROSENBAUM: In between yes, they get out.

But epidemiology is a very difficult thing to do in a way that will prove anything. An enormous number of studies have been done at great public expense -- epidemiology studies, which are a waste of public funds. They don't show anything. And I want to be very careful not to add to the list of things that are a waste of public funds.

We are looking very carefully right now at the backup data to the Mayo Clinic Study which was published already. Was it in Science? Or the New England Journal of Medicine?

DR. ELLETT: The New England Journal of Medicine.

DR. ROSENBAUM: In the New England Journal of Medicine, on leukemia, which said essentially there was no evidence of any harm below 300-rem lifetime dose. I don't know whether -- we haven't done enough analysis of that in great detail. We went out to the Mayo Clinic, several people, Bill Ellett and I, among them, and talked to them in

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great detail, and we now have received, or are about to receive, their backup data, and we are going to make a careful analysis of that paper.

We are also making an analysis now of the possibility of using their records to look at this in utero dose exposure.

If it seems that we can get an answer one way or another out of the records, then we will fund such a study.

COMMISSIONER GILINSKY: Have there been any animal experiments on this effect?

DR. ROSENBAUM: Yes. There's been a lot of animal experiments on the effects, and I would be glad to have Dr. Ellett discuss them in some detail. But it is difficult to extrapolate numerically from animals to people, though, to look at types of effects.

COMMISSIONER GILINSKY: But is the effect observable?

DR. ROSENBAUM: Yes; strongly.

DR. ELLETT: I would like to add that there have been some studies -- I won't swear how good they are -- on diagnostic X-rays, and they did not observe the sort of thing that we see in Hiroshima and Nagasaki. So we don't understand the reason for this. They weren't looking for this as a particular end point. They observed microcephaly, but not at the prevalence you see here; but these studies were not designed for this sort of thing, so I don't want to jump to conclusions one way or the other. It may well be worth the

554 2345 20024 (202) D.C. BUILDING, WASHINGTON, 300 7TH STREET, S.W., REPORTERS time and money to do something more definitive on this.

DR. ROSENBAUM: Let me just add one thing. We have just gone through a reorganization in my little kingdom over there. One result of the reorganization is that we have formed a Statistics and Applied Math Division -- something akin to what I tried to do at the AEC, which eventually became the Statistics Group at NRC.

I believe that we will have a very considerable capability in that regard to do work -- path-breaking work -- which is what is needed to handle these kinds of questions, and we will be looking very carefully at the data.

As an example, the person who has been chosen, though not yet certified, by whatever the Civil Service

Commission is called these days — they keep changing names on me — the Office of Personnel Management, is Dr. Donald Ruben who is to be the Director, who is the current Editor—in—Chief of the Journal of the American Statistical Association, and now head of the statistical activity at the Educational Tenting Service in Princeton, which is one of the largest such practical statistic apply programs in the world, and we have already begun to bring in statisticians not only from the United States but from around the world. And I think because of that, we will have the capability to go into these sorts of statistical questions in a great deal of depth, and to try to make as much of the data as we can. But it is very difficult

to get hard answers from epidemiology, except in overwhelming cases like shipyard workers who spend most of their lives working with asbestos or something like that. But at low effects, it is very difficult.

I don't want to make any promises, except that we will try very hard to get every bit of information there is out of the data.

CHAIRMAN AHEARNE: Dr. Ellett, you were going to comment on the animal experiments.

DR. ELLETT: Well, I don't really think there is that much more to add from what they have. There has been a lot of animal work that's reviewed very well in the latest UNSCA report.

What happens is, an embryo progresses in a sequence of various stages in pregnancy. Now this is speeded up in animals as compared to man, but when you look at the time-different organ generation starts, organ genesis, and give you the dose at that time, you get effects in those organs which are startling.

In fact, they use radiation as a tool to study embryology, to find out what the sequence of events are.

I will say that in animal studies they have been unable to demonstrate a threshold, and nervous tissue seems to be a particularly sensitive end point.

DR. RICHARDSON: If we could go back to the first

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stack of slides, it is number 14 next.

(Slide.)

Perfect. This is a set of miscellany. I am not going to talk about No. 6. I will give you a moment to read it.

(Pause.)

Number 7 continues the existing recommendation for miners.

CHAIRMAN AHEARNE: That's not a change?

DR. RICHARDSON: That's not a change.

Number 9 is not really a change except for the last line. By "miners," we mean -- well, it says "younger than eighteen" up there.

The next slide, please.

(Slide.)

This lists the things that are not covered by the recommendations, although some of them are mentioned in the recommendations in passing.

"Emergency exposure limits" are not covered by the recommended guides. The same for "accidental overexposures."

We feel that that is a matter for regulatory agencies to deal with.

"Medical exposures of patients" is a very interesting item. One could make the case that it is possible that there is as much medical exposure of workers from required X-rays as there is from occupational exposure, because when you

consider that there's about a million radiation workers out
there getting an average of about 1/10th of a rem a year, it
doesn't take very many of the much larger -- perhaps 50 to 100
million workers in the non-radiation workers in the work force
as a whole getting required back X-rays, or whatever, or food
handling X-rays, to add up to a dose that might be bigger than
the total occupational work force. That is a comment in passing,
and we also comment in passing that there is already federal
guidance on the use of diagnostic X-rays for non-medical
purposes and for screening purposes.

"Normal background exposure" of course is not covered.

COMMISSIONER GILINSKY: No recommendation.

DR. RICHARDSON: We're not recommending that background be changed.

"Radon decay product exposure of miners" is already addressed by guidance which was updated in 1970. We're going to look at it again after we get finished with this exercise.

"Guides for time periods other than one year" are only mentioned in passing.

COMMISSIONER GILINSKY: Would you say a word about the exposure to miners? Are the rules or the standards in some way comparable in terms of risk?

DR. RICHARDSON: They're probably higher, perhaps as much as a factor of 2.

(202) 554-2355	2	DR. RICHARDSON: The four working level months per								
	3	year, depending on what risk estimate you use								
	4	COMMISSIONER GILINSKY: Higher than								
	5	DR. RICHARDSON: 5 rems per year.								
	6	COMMISSIONER GILINSKY: than the 5 rems per year.								
	7	DR. ELLETT: I could quantitate that a little bit,								
20024	8	if you'd like.								
W., REPORTERS BUILDING, WASHINGTON, D.C.	9	DR. RICHARDSON: Yes, go ahead.								
	10	DR. FLLETT: If you use the BEIR III risk model,								
	11	a miner that is employed for 30 years at 12 working level								
	12	months per year has about a 9 percent chance of death due to								
	13	lung cancer. Somebody getting 5 rems a year for 47 years has								
	14	about a 6 percent chance.								
	15	COMMISSIONER GILINSKY: The 9 percent was for how								
	16	many years?								
EEI, S.	17	DR. ELLETT: 30, which is probably as long as most								
SIR	18	miners are employed in the mines.								
300 711	19	COMMISSIONER GILINSKY: And how did you come up with								
	20	the 47 years?								
	21	DR. ELLETT: That is 18 to 65 for the radiation								
	22	worker.								
	23	COMMISSIONER GILINSKY: I see.								
	24	DR. RICHARDSON: So the annual risk is about twice								
	25	on that hasis. The lifetime risk is roughly comparable								

COMMISSIONER GILINSKY: Which are higher?

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a re-examination?

2	DR. ELLETT: Seeing as how these numbers were set
3	pretty much without risk estimates, it is amazing that they
4	turned out to be within a factor of two, really.
5	DR. RICHARDSON: The last one is of significance.
6	We have said
7	COMMISSIONER GILINSKY: Let me just ask you again
8	about that. Do miners typically come close to that limit?
9	You were saying that So really the numbers are more
10	different even than was suggested here a minute ago.
11	DR. ELLETT: That's true, and the miners probably
12	get more gamma rays than the average radiation worker, or the
13	average nuclear worker for that matter.
14	DR. ROSENBAUM: Mining in general not only
15	uranium mining is quite a high-risk occupation.
16	COMMISSIONER GILINSKY: That is covered by your
17	guidance, ultimately?
18	DR. RICHARDSON: Yes.
19	DR. ROSENBAUM: Yes.
20	DR. RICHARDSON: The guide was established in 1970,
21	and it is for four working-level months per year. That is
22	a measure of exposure to radon decay products.

Roughly. 50 percent more, but not --

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COMMISSIONER GILINSKY: It sounds pretty important.

CHAIRMAN AHEARNE: And you say it will turn into

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DR. ROSENBAUM: Yes.

DR. RICHARDSON: The last item is the question of dosimetric conventions -- how deep do you measure the dose? What metabolic models do you use for ingestion -- swallowing or breatning radionuclides -- and figuring out where they go and what organs get dosed.

What parameters do you use to express the average worker, called "reference man" here. What quality factors do you use? And so on and so forth.

We have said that the factors in these recommendations. the factors recommended by the ICRP may be used; and that if we felt that there was a need to issue further guidance on any specific issues, we would do so. But we have not done so for anything specific in these recommendations.

The next slide, please.

(Slide.)

This is the last one. David --

COMMISSIONER HENDRIE: Does that last line take us all into the new radiation exposure units, then?

DR. RICHARDSON: We didn't say anything about units. "Units" was not on that list.

COMMISSIONER HENDRIE: Oh. Okay.

DR. RICHARDSON: I said "conventions," and "metabolic

models."

DR. ROSENBAUM: I have been trying to discipline

myself, but I've been unable to do that.

(Laughter.)

DR. RICHARDSON: We are leaving that to the Bureau of Standards.

CHAIRMAN AHEARNE: Recalibrating.

DR. RICHARDSON: David tells me that this schedule is optimistic and he's probably right, because I have always been optimistic in the past and proven wrong. But this does correspond roughly to what he said to you at the beginning of this presentation.

I think that if we don't reach any real snags, we should be able to meet that schedule. That's all I have.

CHAIRMAN AHEARNE: In the Interagency Work Group, do you have to have agreement?

DR. RICHARDSON: No.

DR. ROSENBAUM: Under the guidance authority we don't have to have anything. We could simply give it to the President, who could sign it if he wants to put it in the Federal Register.

As a matter of practice, in fact it is a good idea to try to get as much agreement as you can.

CHAIRMAN AHEARNE: Certainly, but obviously on some of these issues there is going to be some significant contention, so you are just going to have to go out --

DR. ROSENBAUM: Yes. Mr. Costel can sign the thing

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and send it to the President any time he so chooses. He doesn't need anybody else's agreement to do that, to recommend it to the President.

We have prepared now to go either into the technical bases of any of these things -- including the health effects data -- to whatever degree you would like; or, to the worker exposure data, how the population breaks down, who is exposed and by how much and so forth; or, to move on to uranium mill tailings; or to talk a little about the high-level waste, whatever your pleasure is.

CHAIRMAN AHEARNE: I would, I guess for myself,

propose -- unless some of my colleagues have burning questions

on this -- recognizing the importance, but our staff will be

looking at it and we will want to think through it more

carefully, and there is a long period as you have pointed out

for going through it -- to move on to the mill tailings.

DR. ROSENBAUM: All right.

CHAIRMAN AHEARNE: Theme you very much. And as you know, I am delighted to see this.

(Pause.)

DR. ROSENBAUM: Let me say one word of apology to the world.

CHAIRMAN AHEARNE: They're not all here.

DR. ROSENBAUM: But they're all listening.

(Laughter.)

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DR. ROSENBAUM: We are very late with these standards. We were, as I said, required to promulgate the ones we are discussing today, which are the standards for inactive piles, last November 8th, and we were required to promulgate the standards for active piles last May 8th. And it will be quite some time -- we'll try to give you a schedule for this, but the active piles are just beginning, and it might take a year-and-a-half or two years to actually promulgate them.

DR. LICHTMAN: I would like to remind you that EPA's program for developing --

DR. ROSENBAUM: I didn't introduce you. This is Dr. Stanley Lichtman, who has been in charge of the standards.

DR. LICHTMAN: Our program for developing these standards was being carried out at the direction of the Uranium Mill Tailings Radiation Control Act of 1978, which directed EPA to write standards for remedial action programs at a number of inactive processing sites, almost all of which are uranium processing sites. There is one at Cannonsburg of a slightly different character.

In addition, writing standards for active processing sites, those which are operating now and those which may operate in the future.

Initially I will talk only about the program for inactive sites. As David said, we have made more progress on that.

The remedial actions for these sites -- and 25 of them have been designated by DOE to date -- remedial actions will be selected and performed by DOE according to our standards, and with the concurrence of NRC. So NRC has a considerable role in this program, and in fact it has very recently -- NRC has -- undertaken a role with respect to a uranium mill at Edgemont, South Dakota, which is comparable --

CHAIRMAN AHEARNE: "Undertaken" isn't probably quite the complete description of it.

DR. LICHTMAN: Well, I guess you're in the evaluative stages.

CHAIRMAN AHEARNE: Well, I was more -- it wasn't sort of that we --

DR. LICHTMAN: Oh, it's not that you undertook it; right.

CHAIRMAN AHEARNE: -- eagerly sought this role.

DR. LICHTMAN: In any case, it is a very similar job there to what DOE will be doing around the rest of the country. And in both cases, our standards will be the touchstone according to which the actions are carried out.

Now we have conceived of a standard, or two conceptual kinds of standards. One of them are cleanup standards for contaminated open lands and buildings. These are places where tailings may have blown or been conveyed and left on open land -- land which is not occupied by a building;

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or else tailings have been used rather extensively in some places, as you probably know, in the buildings themselves, in the foundations or in the building materials.

The second conceptual element are disposal standards for disposal of the tailings piles themselves, or the bulk of the material. So first I would like to -- Can I have the second slide, please?

(Slide.)

I would like to address the cleanup standards, which we have already proposed in the Federal Register, and you have copies attached in the handout I gave you. And for reasons explained in those Federal Register notices, we also declared the cleanup standards immediately effective as interim standards, primarily so that cleanup could begin as soon as possible.

These cleanup standards, the specifications that we list as the cleanup standards, we are asking that the remedial action program provide reasonable assurance that these conditions are met. These words are carefully chosen, they are something other than "proof"; and the point here is that we expect -- we expect, we urge -- in every way we can say it, we will use that word "reasonable," that as we develop the standards to be highly protective, but it would only be feasible to apply such standards if one didn't have to prove that every square inch of the world were decontaminated to that

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level, but rather that one could use reasonable procedures.

We are encouraging all the agencies involved to approach it that way. So the first standard for cleanup of open land, the key figure there is 5 picocuries per gram. We want contaminated open land, the tailings contribution, cleaned to a level of 5 picocuries per gram of radium 226. There are certain thicknesses specified there.

COMMISSIONER GILINSKY: I don't understand why you phrase it this way. What is it in terms of how much radon is escaping from the surface?

DR. LICHTMAN: Well, that would be an alternative, but we feel that's a much more complicated kind of determination. That a more direct determination is in terms of a property of the material itself. The radium --

COMMISSIONER HENDRIE: It's easier to take a sample of the dirt in the lab and account for an activity level, and infer from that.

COMMISSIONER GILINSKY: But this is under some covering layer?

DR. LICHTMAN: Not necessarily. As is indicated here, it is the same 5 picocurié per gram number but we consider -- but the standard applies only to thicknesses of material that are greater than 5 centimeters, if they occur near the surface; or 15 centimeters' thickness if it occurs below one food from the surface. So it is the quantity of

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the material, as well as the concentration that counts for something in determining the hazard it presents.

DR. ROSENBAUM: Let me say something about that.

It is difficult enough in the field, which is what this has to be done for, to make radium measurements that will have a great deal of credibility at such low levels.

If one were to ask in such a large number of circumstances that this will represent, for radon emanation measurements, it would be difficult to get credible measurements at all. In our own field measurements which we do at ORP out of our laboratories by putting a box over the ground and carefully monitoring what comes out, it is very difficult to duplicate the measurements with two boxes beside each other even apparently doing the same thing. And there are very many techniques about to do this.

There are lots of compounding things that make technically field measurements of radon emission very difficult. So I don't think it would be a workable standard to do that.

DR. LICHTMAN: Let me say further that that would only address one hazard of the material, which would be the radon emission. There is also gamma radiation to consider.

Moreover, the standard doesn't say how it will be implemented. One could work out various surrogates for determining that.

GOMMISSIONER GILINSKY: Well, but behind this there's got to be some standard or objective for how much radon you are willing to put up with leaving the ground at that point.

DR. ROSENBAUM: Sure. But, after all, the radon is a decay product of the radium. It's a direct decay product, and therefore it is not very difficult to go from a radium concentration in some average way through the thickness of soil to a radon concentration with some error, of course.

CHAIRMAN AHEARNE: Well, underlying this is there, then, some backup calculation that says: Here is what is an acceptable radon level above the ground?

DR. LICHTMAN: Not quite. Not quite. Yes, there is a backtround -- of course there have been estimates of what kinds of radiation hazards material of this character and concentration presents, but it's a continuum. There is no clearcut place at which one deals with an acceptable -- one finds an acceptable level. So one also has to consider a difficult it is to find the material and to clean it up, how extensive a problem that is.

COMMISSIONER GILINSKY: Well, you must have started by saying: We're willing to put up with so much radon leaving the ground.

DR. ROSENBAUM: Let me say, that's not the way things work -- not the way things work with us. That's not how we start.

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One would like -- in the first place, every one of these actions, or at least many of them, are taken under different laws. In each case we try to interpret the intent of Congress in making that law -- and they are not necessarily consistent with one another, nor are the laws, in terms of intent.

In this case we were dealing with the Uranium Mill Tailings Radiation Control Act of 1978, which deals only with material -- in this case, the inactive piles -- material which comes from piles which provided all, or virtually all of the material to the Federal Government. And perhaps because of that reason, Congress seemed willing to have the things cleaned up very well at public expense. The expense here is 90 percent borne by the Federal Government, and 10 percent by the state; except in the case of Indian Reservations where it is 100 percent by the Federal Government.

It was our having read the law and the history of the law, it was our belief that if Congress had intended that we should make the lowest standard which was reasonably feasible and say we should clean it up as much as it could be cleaned up without getting to the point where you couldn't find whether the material was there, or whether -- you wouldn't be able to believe your measurements as to whether it was cleaned up -- and it's much more driven that way than it was the other way.

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CHAIRMAN AHEARNE: Are you saying, Dave, that this is driven by measurementability?

DR. ROSENBAUM: It is driven partly by the ability to find the material in the field and measure it, and to verify that you've done it, and that it's feasible to actually carry out the standard.

The cost was not such a driving factor in this case, because Congress said they wanted to pay for whatever was necessary to clean these things up very well.

But let me say something in general about that, because it bears on the whole way radiation standards are made, at least by us. There isn't an acceptable level of risk.

The various guides, standards, advices to states, and other such things we've put out to the public vary enormously in the risk that they subject people who are covered by those documents to. The tightest one that I am aware of, the most protective, is 40 CFR 190, which limits the most exposed person around the nuclear fuel cycle facility to 25 millirem a year.

For example, our advice to the State of Florida about cleaning up homes on phosphate lands which had radon in them, if it were followed, roughly would subject the people to probably 100 times the risk -- and a great many more people; this isn't the most exposed person -- than we allow people living near nuclear fuel cycle facilities.

So that there isn't any such path that one decides what is a tolerable level of risk, and then demands that things meet that. That is not what has happened at all.

If we had the power to and were to demand that homes be free of radon so that the people in them would suffer no .

more than the equivalent of 25 millirems per year, it would --

CHAIRMAN AHEARNE: Devastate the conservation program.

DR. ROSENBAUM: -- radically change the -- it would do away with the conservation program, and radically change the nature of housing in the United States. It would probably call for demolition of millions of buildings -- at least millions.

So there is no consistency at all in the level of risk, even in our own little shop, of the level of risk that we allow or suggest, or recommend that people be subjected to. Therefore, we don't go about it that way by saying this is a tole able level of risk.

COMMISSIONER GILINSKY: You are saying that you have driven these standards to the limits of measurability of radium 226?

ments. That's not what I meant at all. I meant in terms of practical application in the field. There's no sense in making a standard in which people who have to carry it out in a

New Mexico or something, I'm going to be able to in good faith do a reasonable job with the sort of people they're going to have there and the sort of measuring instruments they have, and to be able to tell within reason whether they're meeting the standards or not.

COMMISSIONER GILINSKY: Well, still, I mean let's say portable instruments.

DR. LICHTMAN: Well, there is some controversy among the agencies as to exactly how to implement this, and whether it can be done with standard survey techniques and hand-held instruments. We are exploring that. We have formed an interagency --

COMMISSIONER GILINSKY: So this does not -
DR. ROSENBAUM: It's the other way. DOE has

complained that they may not be able to meet even this in

the field; that this is too stringent.

COMMISSIONER GILINSKY: They won't be able to measure?

DR. ROSENBAUM: They won't be able to measure accurately these levels, and therefore the standard isn't meaningful.

DR. LICHTMAN: Well using survey instruments.

Everybody recognizes that you can pick up a sample and process
it in the lab to much lower levels.

Let me say that we didn't only set the standards on those grounds, but also I want to emphasize, by examining the health consequences of these levels and determining that any hazard associated with them is certainly not unusual. That is to say, this is several times the average of concentration of radium in normal soils. It is something like 5 to 10 times the average concentration, but it is not unsual to find places with higher numbers than this. So there is no reason to think of the limited areas that would be cleaned up to the levels for proscribing places to be avoided. In fact, that is one of our goals, if it was practical to achieve, that these be places that one need not avoid and could be put to public use.

So it does seem to us that it is possible to achieve all these things with a reasonable cleanup program, and that is what we have tried to do.

DR. ROSENBAUM: Just to emphasize, I didn't mean that we didn't do a health-effects calculation. I meant that the way of thought wasn't what you suggested, that we first set an acceptable level and then saw what it would take to meet them. We did do health effect calculations and satisfied ourselves that the health risk was acceptable. Those are available.

COMMISSIONER GILINSKY: Well, let me just pursue this a little for a moment. If the means for cleaning it up is to

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be a covering of topsoil, or clay, or whatever, over a pile, how can you ever --

DR. LICHTMAN: Oh, no, no, it isn't that.

DR. ROSENBAUM: These aren't piles; these are clean up of open land.

DR. LICHTMAN: Let me explain. This is for cleanup.

I distinguish between "cleanup" and "disposal" standards. This
is the cleanup of contaminated land which is not the site of
a tailings pile.

COMMISSIONER GILINSKY: I see.

DR. LICHTMAN: This is other than dealing with the pile itself.

DR. ROSENBAUM: This is just windblown --

DR. LICHTMAN: This would be blown material. It might be -- you know, if you decide to move a tailings pile, you might then apply this to the ground that's left.

DR. ROSENBAUM: You can imagine, for example, if you had to survey 100 square miles and send samples back to the lab if you couldn't do it with a hand-held instrument. You wouldn't be able to verify it very well.

COMMISSIONER GILINSKY: If there are areas presumably in those neighborhoods where a natural concentration might be greater than this, what would you do about that?

DR. LICHTMAN: Well, the standard applies to the material from uranium mill tailings, as it says.

COMMISSIONER GILINSKY: And is that readily identifiable?

DR. LICHTMAN: Well, the question of "readily," I'm not sure that there ever really will be terribly --

COMMISSIONER GILINSKY: I mean, would you know that you're dealing with --

DR. ROSENBAUM: Theoretically there's a different isotopic mixture.

DR. LICHTMAN: If push comes to shove, you can tell the difference. I think normally that's not what the issue will be: There are some cases where a load of material exists somewhere, and one wonders what is it, where did it come from.

COMMISSIONER GILINSKY: It might be in the mining business.

DR. LICHTMAN: This is refined material. However, the uranium has been extracted from this. If one does an isotopic analysis, you find peculiarities. You can determine that it's tailings rather than something else.

CHAIRMAN AHEARNE: Okay, how about buildings?

DR. LICHTMAN: Standards for the cleanup of buildings address the two primary hazards of the material, which are the radon emitted into the building, and secondarily it turns out, the gamma radation.

CHAIRMAN AHEARNE: How does that radon background compare with the fluctuation that you might find in background?

DR. LICHTMAN: You will certainly find normal houses with no tailings that have these levels.

DR. ROSENBAUM: You find them in the East, too.

DR. LICHTMAN: We are applying the standards to places that have tailings, however. These standards apply to buildings that have tailings.

CHAIRMAN AHEARNE: I know David would be very interested in that.

DR. ROSENBAUM: Let me say that if one makes a reasonable effort to remove any tailings that are there and still the level is above the levels that are set, then you have already complied with the law. The law only applies to uranium mill tailings from designated sites. If the levels are higher for other reasons, it has nothing to do with that.

CHAIRMAN AHEARNE: I know you have looked at this a lot, but what is the rough average that you expect when you say the "East"?

DR. ROSENBAUM: The "average" is in great dispute, because there have only been many 15 or 20 homes that have been monitored carefully for a whole year in the world, but the average we use is .004 working levels as a typical house.

However, the Department of Energy, the MSOL Labs in New York City, have done some things in and around New York City and in New Jersey, not a place one ordinarily would mine uranium, and have found guite a few houses above .01, and some

even I think above .02. In places like Butte, Montana, one finds many houses above .1, and much higher than that.

There are -- and we have only begun to look around the country. There are houses in some of these mining districts, particularly if they are buttoned up, in which the miners are in more danger at home than they are in the mines, even with the current levels.

DR. LICHTMAN: Something to realize about radon is that it has a high risk factor associated with even normal, natural levels.

COMMISSIONER GILINSKY: What are the levels in the houses where tailings have been used for foundations or whatever?

DR. ROSENBAUM: Oh, there are some very high levels, as you will see.

CHAIRMAN AHEARNE: .47.

DR. LICHTMAN: Well, many such houses have been found in Grand Junction to fall below these remedial action levels. Of course many fall way above.

COMMISSIONER GILINSKY: What is "way above"?

DR. LICHTMAN: Well, easily 10s of times this number.

CHAIRMAN AHEARNE: I think .47 is the highest number.

DR. ROSENBAUM: I mean, there is a famous fire station in Salt Lake City which had 1.5 working levels, and that is a fire station people live in. Firemen sleep there.

	DR. LI	CHTMAN:	Well,	th	at	was	not	an	annual	average,
but they	did get	measure	ments	up	tha	t h	igh.			

COMMISSIONER GILINSKY: Could you define a "working level"?

DR. LICHTMAN: I'd rather not try to pop it off, but what it is is the amount of --

DR. ROSENBAUM: I'll pop it off. It's the amount of alpha energy — it has to do with the amount of alpha energy emitted from short-lived — he's going to lock it up and see if I'm right — short-lived rajon daughters. One working level is 1.5×10^5 MEV of alpha energy total emitted from short-lived radon daughters. It is the concentration that gives — the concentration in the air per liter of air that gives 1.5×10^5 MEV of alpha energy from short-lived radon daughters.

DR. LICHTMAN: It's 130 billion electron volts.

DR. ROSENBAUM: It's 1.3, then.

DR. LICHTMAN: And it comes out to exactly the same thing.

COMMISSIONER GILINSKY: It gives that position in some unit of time?

DR. LICHTMAN: It is any combination of short-lived radon decay products, which are the ones that we are concerned about, that the hazard is associated with in one liter of air that will result in the ultimate emission of alpha particles --

which are again the thing we are interested in -- but the total energy of 130 billion electron volts.

DR. ROSENBAUM: So it is 1.3 instead of 1.5.

DR. LICHTMAN: So it is a measure of the concentration of the hazardous radionuclides, mainly -- the shortlived ones that give off alpha radiation.

COMMISSIONER GILINSKY: So it's a measure of the concentration in air?

DR. LICHTMAN: In air.

DR. ROSENBAUM: In air, and these are short-lived radionuclides. They are what take you from radon down to lead 210 -- from radon 222 to lead 210, the alpha emitted in that chain. There are beta links, too, but --

COMMISSIONER GILINSKY: How do you describe the standards on someone? I suppose you're only allowed to be in that environment for a certain length of time?

DR. LICHTMAN: No. This is designed for buildings, and one can characterize what the occupancy of a building is likely to be, what fraction of the time people spend in buildings. So we have folded all that in in determining what is an appropriate standard.

COMMISSIONER GILINSKY: Can you convert that into a dose to the lungs in terms of millirems for a person who spends X number of hours, or half that time in that building?

DR. ROSENBAUM: Bill?

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(No response.)

DR. LICHTMAN: People have done that --

DR. ROSENBAUM: The answer is "yes."

DR. LICHTMAN: It is controversial. Bill Ellett, who is the head of our Bioeffects Branch, doesn't like us to even talk about it.

Bill, do you remember what the number is?

DR. MILLS: It is about .5 per rad working level month, and you will have to explain to them what a "working level month" is. But if you take .5, that is the National Academy of Science number. So one working level month will give you .5 rads.

COMMISSIONER GILINSKY: To the lungs?

DR. MILLS: It is to the lungs, but it is to the average lungs; not the lungs -- which makes a difference --

DR. ROSENBAUM: Let me say, that's not as important as you might think, because the health data on which standards are based is in terms of measurements -- measurements were made in terms of working levels. So the health effects are directly given from a certain amount of working level, and you don't really buy anything by converting those working levels to dose to the lung, or the epithelium of the lung, because the data themselves are in terms of working rads.

DR. LICHTMAN: We have a footnote on the subject in the Federal Register notice, if you care to refer to it.

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COMMISSIONER GILINSKY: Thank you.

DR. LICHTMAN: So that in determining the risk to the lowest degree reasonable, we have experience from the Grand Junction program, which was very useful for us in deciding what that lowest practical level is.

Moreover, we recognized that there will be exceptional cases that, try as hard as you may, you may not be able to get a house down to this level by removing tailings, let's say. And as I indicate later, we have built in a provision for exceptions.

Now the third element at the bottom of the page is kind of a catch-all standard. We designed the first two standards by considering the hazards of uranium mill tailings which, after all, are a fairly definitive kind of material. You know what's in it, by and large, which varies in concentration from place to place, but as I indicated the Cannonsburg, Pennsylvania, site has a little bit different kind of history. It wasn't a uranium mill; it was some sort of a processing facility. And on the possibility that there might be unusual substances there the hazard of which might not be dominated by radium 226 and its decay products, we wrote a standard -the third entry there -- which says, in effect, if you find other things, the total risk that you allow from the site should not exceed the risk you would get from radium 226 and its decay products, if only under the first standards.

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Now the status of our disposal -- next slide, please -- (Slide.)

I would like to address the disposal standards now, the status of which is somewhat different from the cleanup.

Cleanup standards have been proposed; they're before the public. We have a comment period which is remaining open on them while we continue to develop disposal standards.

Our objective is that we will propose disposal standards within a couple of months, announce a comment period at that time to run jointly for the cleanup and disposal standards, and then hold joint hearings on them and publish a joint EIS on both the standards. We split the package at one point in order to get the cleanup standards out as quickly as possible, but we intend to recombine the project and deal with them jointly as soon as the disposal standards are issued.

The draft disposal standards, which I am going to discuss here, are before the Commission Staff now for their comments, and they are before the other agencies in our Interagency Working Group. They are also before our Assistant Administrator.

COMMISSIONER GILINSKY: Now how do you -
DR. ROSENBAUM: We hope to get it out in October.

COMMISSIONER GILINSKY: How do you determine that
the annual releases will in fact not exceed those numbers?

DR	. LICHTMAN:	Well, to	wo ways	. Let n	me empha	size
those underl	ying words.	. We are	asking :	for a ":	reasonab	le
expectation"	that the c	conditions	below s	will be	met for	1000
years.						

COMMISSIONER GILINSKY: What I'm asking is: Can you measure it?

DR. LICHTMAN: Yes, you can measure it, but it is not clear to me that a measurement that you make at a given date is the ultimate way of providing that reasonable expectation, which is what the standard requires.

COMMISSIONER GILINSKY: I'm not trying to press you on whether you can --

DR. LICHTMAN: People have made flux measurements of tailings piles --

COMMISSIONER GILINSKY: -- going back to our earlier conversation --

DR. LICHTMAN: -- just as they make flux measurements off a piece of ground, as Dave talked about.

DR. ROSENBAUM: But they're not very reliable.

DR. LICHTMAN: They're not very reliable, and they change with the weather.

DR. ROSENBAUM: Let me say, yes, they do change with the weather considerably. They change with how wet the soil is, and the wind, and the atmospheric pressure, and everything.

When you're talking about something that's 1000 years

here, or in the case of the high-level waste standard perhaps much longer than that, 10,000 years maybe, then there is no way to enforce the standard in the sense that you can go to a house after you've cleaned it up and make some reasurements to see if the gamma levels are higher or lower than the standard to see if you have complied.

There is no certain way to proejct the future. The only thing that one can expect is that one takes measures which the best scientific examination would lead one to believe would comply with the standard in the future. There is no way we are going to know for sure, whatever we do, 700 years from now the standard will have been complied with, unless we wait 700 years and measure it.

So we have to make some kind of reasonable scientific projection, and that is the essence of what the "reasonable expectation" is.

COMMISSIONER GILINSKY: No; I understand that. I was just asking about today.

DR. ROSENBAUM: You can measure it, but I think that if one were, for example, to cover, as one alternative to cover a tailings pile with certain types of material to a certain depth, and take certain other actions to ensure it wouldn't blow away and stuff, and take care of water runoff and things, that that would be a better way to have a reasonable expectation that it would meet the standard for 1000 years, than going

out with a box whose measurement would depend probably on where you put it on the piles -- and these things, have you ever seen a uranium mill tailings pile? These are enormous things.

COMMISSIONER GILINSKY: But I am still curious to know whether it can be measured.

DR. ROSENBAUM: It can be measured, but not very reliably in the field. That's my impression.

COMMISSIONER GILINSKY: So it would be done on the basis of some calculation?

DR. ROSENBAUM: It will be done on the basis of models. One can get an estimate, certainly, of whether it is very high or very low; but I think that basically these standards will have to be satisfied by measures which, when examined carefully by technical people, seem to provide a reasonable expectation that they meet the standards.

One could then go out and measure after these measures had been taken to see if you got readings that were higher.

That might be worthwhile doing. But Stan's point is very important: The actual reading you get on a day in a particular place depends on the place, and very much on the weather.

Very much. And the weather that's been in the past few days --

COMMISSIONER GILINSKY: Well, if they take a lot of readings they ought to know it would be above this number.

DR. ROSENBAUM: That's right. If you take a lot of readings and they were all above the number, you would have to

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worry. That's right.

DR. LICHTMAN: Right. I think it can be literally carried out by measurements that one can determine for a given year, even, that the annual average won't exceed 2 picocuries per meter squared. But the essence of the standard, in my view, is the full language -- the reasonable expectation that for at least a thousand years that will prevail. And there is no way you can decide that aspect of it by making a series of measurements.

There is some novelty to this requirement. We'll all have to work hard to develop ways to implement it.

The reason we picked a thousand years, rather than some other number, is because it seemed the longest doable practical time for this program. I emphasize that. This is a remedial action program; it's not a start-from-scratch program.

The second -- the following two requirements are for protection. The first requirement protects releases to the air of radioactivity. We have no need to address gamma radiation releases, we think, because we believe that implementing these three standards will, in almost every way we can think of implementing them, also screen out the gamma radiation.

The remaining hazards to be addressed are water protection for radiological and non-radiological contamination. The standards which you have available to you now through

that package that we sent, I didn't want to write down all the numbers here, but they specify that for that same period there should be a reasonable expectation that releases to groundwater will not either cause the concentration in groundwater of certain specified substances to exceed specified levels; or, if they are already exceeding those levels, that the concentrations not increase because of releases from the tailings.

So these are conditions that apply to releases of tailings after they are disposed of. They do not try to rectify the past. But it says what conditions should be satisfied after one disposes of the tailings.

Similarly, releases of toxic substances or radioactivity to surface water should not degrade the surface water as the condition.

Now we recognized, in developing these standards, that we have a wide variety -- the next slide, please -- (Slide.)

-- that this is a remedial program for existing conditions which occur under a wide variety of circumstances in different climates and different quantities of tailings, tailings of somewhat different concentrations, in different water environments and so forth. And that if we are going to write what we believe are strict standards that are necessary to protect health, that there has to be some flexibility

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applied to them as well for those exceptional cases for which the standards would be unreasonable.

We have identified some exceptional circumstances and given criteria under which the standards need not be met. Among those are requirements where meeting the requirements of the standards would endanger public health or safety -- such as one, in deciding whether to move a uranium tailings pile in order to meet the standard, if that should be required anywhere, one should also consider the safety hazard of moving the material, which might be a factor that would lead you to make another decision. Or, in some instances the tailings are very precarious in their present location, and it is difficult to get equipment in, and the workers might be endangered, and one might decide then not to move them for that reason.

Another reason would be that if environmental protection itself would be better served in not cleaning up open land -- say if tailings were spread in not very high levels over a large amount of desert soil where vegetation is scarce and takes many hundreds of years, really, to restore, or if there might be some very scarce vegetation there, or some rare creatures, one should have the flexibility of deciding that environmental protection would be better served by not cleaning that material.

Thirdly, if the costs of cleaning buildings should be unreasonably high in some instances -- unreasonably high,

considering the context in which we made these judgments as to what would be unreasonably high, and compared to the benefit.

So, for example, if one had a house in which the indoor radiation levels were slightly exceeding the standards -- and I emphasize it must be "slightly" -- and the costs would be very high to remove whatever remaining material there was, or even to find it perhaps that was causing the excess, one might justify not doing it.

In any case, these actions should be documented.

DOE -- the next slide, please.

(Slide.)

perform remedial actions that come as close to meeting the standards as is reasonable under the exceptional circumstances. They should notify the private owners and occupants of affected properties, and ask them for comments if they have any; and they should notify EPA when they take exceptional remedial action so that we can track our own standards, and their practicality, and learn from the experience.

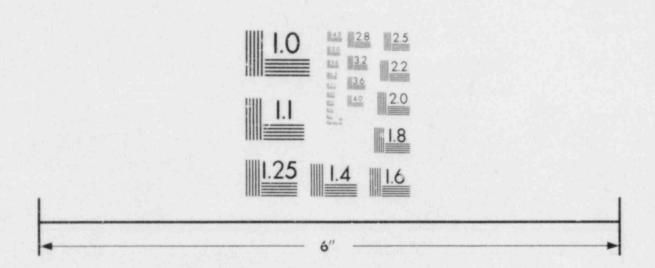
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(Slide.)

I would like to very briefly discuss the standards for active mills. I guess we've really done that. We need cleanup standards and disposal standards certainly for tailings at active mills; but there is an additional element to be

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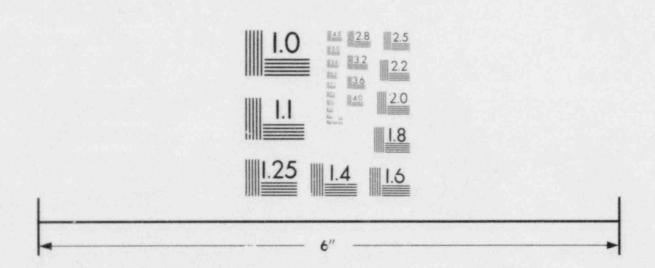
IMAGE EVALUATION TEST TARGET (MT-3)



MICROCOPY RESOLUTION TEST CHART

1.0 1.1 1.1 1.25 1.4 1.8

IMAGE EVALUATION TEST TARGET (MT-3)



MICROCOPY RESOLUTION TEST CHART

addressed, which is the appropriate limits, if any, on effects and emissions from tailings piles during the operational period of the mill. That is something that we will want to look at carefully, and it is a difficult technical problem, as you know.

CHAIRMAN AHEARNE: Do you expect to have any significant differences in the standards you are going to propose for the active sites versus those that you are here mentioning for the inactive sites?

DR. LICHTMAN: Well, as David indicated, we are trying not to have expectations. We want to look at the problem from the beginning.

CHAIRMAN AHEARNE: Well, except that you here have ...

DR. ROSENBAUM: Certainly a lot of the work, the scientific work that we have done on these will carry over to the others. But standards are not made in a vacuum; they are made depending on the actual circumstances and the ability to carry them out, and the cost to carry them out.

CHAIRMAN AHEARNE: And I gather -- did I interpret correctly the point you had made about the legislative history of the Mil. Pailings Act led you to a certain set of assumptions on how to approach the inactive that may not correspond and track?

DR. ROSENBAUM: Not necessarily. I don't know. It certainly wouldn't be surprising if they were very similar, but it's not -- we just have to look at the evidence carefully.

My staff has looked briefly at the proposals that will be shortly before you, I guess, for active sites. We don't, at the moment, have any disagreement with them, if that's any help.

DR. LICHTMAN: I would like to point out that the schedule for proposing the standards of September 1981 allows us to take advantage of the comments we receive on the inactive site package and benefit from that in approaching the active sites.

That is all I had to say.

CHAIRMAN AHEARNE: Vic?

(No response.)

CHAIRMAN AHEARNE: Joe?

COMMISSIONER HENDRIE: No, no questions. I am very pleased with the discussion. I am glad to see we are charging ahead. Keep going.

CHAIRMAN AHEARNE: Thank you, Dave.

DR. ROSENBAUM: Thank you.

CHAIRMAN AHEARNE: David, before you disappear, would you just say a few words about high-level waste and the status?

DR. ROSENBAUM: Sure. I don't know if Dan Egan is still here? Yes.

This is Daniel Egan who is in charge of that thing, and he has got a couple of slides which would -- this is only

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two slides, so it will be very brief. And it has the information I think you wanted.

(Slide.)

MR. EGAN: Okay, what I am just going to do, rather than get into any details of the action we are currently developing, is just to give you an overview of what it contains structurally and indicate where we are in our own internal process.

I apologize first of all for the title, but I haven't figured out any way to make it any shorter yet. As David indicated before, we had two authorities to work with in this are. One is to promulgate generally applicable standards like the mill tailings standards. The other is to propose better radiation guidance like the occupational guidance. This package has both types of proposals in it. It also covers both management disposal, and it considers spent fuel high-level waste, both as we typically used to think of it as reprocessed waste, and also spent fuel, if it is to be disposed of. The standards also apply to transurance waste above

The two parts of the environmental standards would be Subpart A and B. Subpart A would apply to waste management operations and storage of these wastes. 40 CFR 190, which are our uranium fuel cycle standards, specifically exclude operations at waste disposal sites from consideration in that

554-2345 300 7TH STREET, S.W., REPORTERS BUILDING, WASHINGTON, D.C. 20024 (202) regulation. It is also unclear as to what types of storage are addressed by 40 CFR 190.

What this action will do will just explicitly extend the same dose limitations that are in 40 CFR 190 to these other processes as well. It would include operations up to the implacement of waste in their repository, and monitoring a repository before it is finally sealed, and backfilled, and walked away from in the sense of being easily able to get back at the waste that is placed in the repository.

Subpart B, which is the standards for disposal, are then of course much different than standards we've developed before in 40 CFR 190, or in Part A of this standard. We are here discussing limits on projected releases over a 10,000-year period. Similarly to the mill tailings, you are talking about something that can only be implemented on the design basis and through analytical models. And perhaps unlike the mill tailings, in this case you have no recourse at all to measurement.

with the mill tailings, you could theoretically go
out to a pile which you have stabilized and measure to see if
the standard was initially met. You don't have the same
situation with these standards. The entire process will have
to depend upon the analytical exercise, and verifying
repository design projections.

And of course the other part of the requirements for

disposal which we propose to include as an appendix to the CFR language, the Federal Radiation Guidance containing general principles that should be followed for disposal sy lems.

This part of the action would be promulgated somewhat differently, as David explained earlier, when we finally make the action final, in that the Federal Radiation Guidance of course would be recommended to the President for issuance as guidance. The Administrator cannot issue it directly by himself; whereas the standards Subpart A and B, say, would in fact be issued directly by the Administrator.

DR. ROSENBAUM: Let me say one word about that.

This complication arose very late in the process when our lawyers, just a month or so ago, decided that we couldn't issue the whole thing as a standard. We had to separate out part of this and issue it as guidance.

CHAIRMAN AHEARNE: Lawyers are like that.

MR. EGAN: I guess I would pause for any questions here. My next slide will only cover the process that we are involved in, and has no substantive details of the standards, per se. So if there is anything you would love to get at here --

CHAIRMAN AHEARNE: Have you, on the seven general principles, could you say a few words on what approximately these are?

MR. EGAN: Sure. There are a couple of them that

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are fairly straightforward, and probably would be motherhood statements, if you will. Some of them can be quite controversial.

The simplest one is that releases from a disposal system should be reduced as low as is reasonably achievable. I don't expect there to be much quarrel with that.

Another one that is somewhat related but again different, is that the disposal system should use multiple
barriers to isolate the waste; and that each of these barriers
should be designed to provide substantial protection, even if
the other barriers don't work the way they're supposed to.
This gets to the question of whether systems analysis on a
whole system is necessarily reliable over this period of time.
It is a definitely a conservative principle. In other words,
you are saying that even if you have a good site, and say a
good waste form, you shouldn't use a terrible canister. You
shouldn't take that easy step.

Another one would be that we believe that active institutional controls to protect the disposal system should not be relied upon for more than 100 years. Now that particular criteria, if you will, was one of those we had proposed as a general radioactive waste criteria back in late '78. In fact, it arose from some of the workshops we have had in that program before we actually even proposed the criteria.

We have modified that to accentuate the word "active." A lot

20024 (202) 554-2345 300 TTH STREET, S.W., REPORTERS BUILDING, WASHINGTON, D.C. of people criticize us, I think fairly, that knowledge, per se, can be considered institutional control and can survive for a long period of time. What we are specifically saying is the idea of guards and fences that will forbid people from going on that site don't last forever, and you shouldn't rely on them to last forever.

Related to that -- and I go back to that one point that the 100 years we measure from after the waste has been disposed of. It's an introduction to the next one which says that we believe waste should be disposed of promptly once you've got a system that will do it. We do not believe, at this point, that it is appropriate to essentially enter into what amounts to non-ending storage that has to be continually monitored. There have been some proposals that in fact waste should not be disposed of at all; that in fact you should put them in a muscleum or some other such mechanism where man would commit future generations to watch them either endlessly or until they eventually decayed to be innocuous, which would be a tremendously long time for these materials. That is principle number four.

Another principle is that you should locate a site away from potential areas of resources -- both resources which are obvious that we now consider to be resources; but also away from areas where there are unique concentrations of materials that may be a resource in the future, even if they're

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not now. We don't presume that we can predict that we know what is going to be a resource hundreds of years from now.

DR. ROSENBAUM: Let me make a comment on that. The question always comes out. That requirement is not intended to, per se, exclude salt.domes, or salt deposits, even though salt is indeed in some sense a resource. It would be possible, perhaps -- obviously each site has to be looked at in particular to see if it satisfies the requirements -- but it would be possible, certainly, that a salt dome might satisfy, or a salt deposit might satisfy the requirements for the repository.

MR. EGAN: Yes. It is clear that salt deposits don't have a problem. There are maybe certain categories of salt domes that might be ruled out just because they are very often used for their storage capacity. Salt domes are very frequently used to store natural gas, an easy place to store things. We agree that certainly our standard criteria would not categorically rule out anything; but it would require that that principle be considered in your licensing of a repository.

Another principle is just that you should record, and mark, and otherwise warn the future about the repository as well as you possibly can. What we call "markers and records." When we started this, there were some people that were advocating that perhaps the safest thing to do with the repository was to hide it so that nobody could ever find it.

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We talked to some archeologists, and they said:

People are pretty clever about finding things. Somebody who
is looking would find some anomaly, maybe not knowing what it
was. So the idea of hiding the waste so that nobody could find
it wouldn't make any sense. And having agreed that you should
tell people, you should tell them well.

The last one, and the one that usually requires more explanation than the others, is that we feel the waste should be disposed of what we call "recoverably." We picked that word to avoid the word "retrievability," which has been used in a much different context. We mean that you can recover the waste. We don't mean that you leave a repository open, for example, so that people could walk in and monitor the progress or the status of the waste in the repository. We mean that you dispose of it in such a way that it is feasible to unearth it, or to recover it, if you will, some unspecified time in the future if somebody discovers that we did it wrong, that we should not have ever put it there.

Mined repositories, as we typically think of them, don't really have a problem of that particular type of general principle, but there are disposal methods that have been proposed that in fact once you implement them, the baby scream is gone with the bath water. The rock melting concept is one I don't even like to talk about. There was an idea that you could put it in little waste canisters that would get so not

they would melt their way to some depth, and therefore be isolated. Once you do that, you can't undo it. There are other examples of that type of disposal.

DR. ROSENBAUM: That was something I mentioned to the Commission in my first week on the job when I came to talk about another problem. I think it has to do with the limits of predictability.

It is certainly true that a lot of things we believe in good faith now are going to turn out to be wrong. It seems to me to be good sense to be able to undo your mistakes; to, if you make a mistake, to at least be able to go back and fix it. I don't mean that it should be easy to retrieve it, but just that it should be possible if somebody, for some reason we now can't understand, finds out that we've made some terrible mistake. So that's the idea behind that.

MR. EGAN: And having capped it at seven, I think it's all right.

COMMISSIONER HENDRIE: For how long?

MR. EGAN: We have been asked that question a lot, and we really don't specify. The principle says you should be able to cover most of the waste. What we are arguing of course is not that say a repository and some waste leeches, obviously some atoms could get away from you and you couldn't get them back. But it almost says that if it could be done the first 100 years, you would probably be --

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DR. ROSENBAUM: Well, there's one thing that we're almost certainly not wrong about, and that is the half-lives of the isotopes. So that certainly if you could get them back in the first 500 years, after that it is difficult. They would have degraded to the point where it's difficult to imagine that your mistakes would be catastrophic. I don't know exactly what number it would be, but that would be one thing I do want to say is fairly certain, the hel-lives of isotopes.

COMMISSIONER HENDRIE: Does that apply to canister design which is guaranteed say for 1000 years?

MR. EGAN: No. It doesn't apply to me, anyway.

DR. ROSENBAUM: I don't --

MR. EGAN: It applies, the canister in the waste form -- you know, if the canister may fail and the waste form may be subject to groundwater, and there may be some leeching, it doesn't mean, you know, that I'm assuming it's all going to disappear very rapidly or quickly. If the waste form leeches, most of it will stay in the general area of the repository, and only a small fraction of it will actually be washed away.

DR. ROSENBAUM: I understand your point, even if it came out of the canisters and was sitting there, you might not be able to get it back in any reasonable way. It may turn out after further thought and in the hearings that that's not a very bright idea, this retrievability, or whatever we're calling it, recoverability. In that case, we'll drop it.

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It seemed to me to be prudent, I think is the right word, to try to do that if it makes sense. If it doesn't, we'll take it out. I think there is ample time in the hearings and everything to explore the issue of whether that pages sense as a requirement.

COMMISSIONER HENDRIE: I guess I am more concerned about the level of proof that one would be required to present and establish in a litigation on this point than I am about the general requirement. As an engineering principle, you know, I am willing to say that if you vitrofy in reasonable ways as the best evidence at hand indicates, and put it in canisters and overpacks, as best-technique indicates, and keep the unit loadings low so the temperatures are not very high, and you stick it in a hole in the ground, why I have ultimate confidence that if you are willing to spend enough money on it and do it carefully and so on, why you can go down and mine out, you may end up with a great volume of material, some of which is highly radioactive and a great mass which is trace radioactive, and now what are you going to do with that? So I'm not sure it's a great idea to mine it, but the ability to mine it, I have confidence.

Now you say: Good. Take your confidence, even, and let us see you present your evidence, your testimony and your evidence, and stand up in litigation and prove it in order to meet a regulation on waste disposal. I'm not so sure how I go

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about that.

DR. ROSENBAUM: Well, let me say two things.

One, all the proof in this case is going to have to be done largely by models and by technical projections, which probably mean computer models or some sophistication into the future. So any kind of proof you are presented with is going to be of a nature and therefore complicated to understand how much you can believe it.

The other thing I would say is that one good reason to drop a requirement is that it is not implementable. And if it turns out that that's the sort of requirement that the Commission wouldn't find possible to implement and couldn't in any satisfactory way decide whether to implement it or not, that would be a reason not to do it.

COMMISSIONER HENDRIE: It is likely to go the other way, because there are certainly parties who are active in the field who have an interest in preventing the implementation of long-term high-level waste disposal. Now if they can ge: a standard established which you then can't meet in some appropriate level of proof in a hearing, by George, they've got it made.

So you are going to have at least some elements of the multi-party scene who will be attempting to induce unmeetable standards. I have some concern.

CHAIRMAN AHEARNE: But I trust it isn't us.

DR. ROSENBAUM: I would hope that we would have testimony among other people from the Commission -- that we would have people speaking to the issue of whether the standards will be implementable or be able to be done.

commissioner Hendrie: You will have people coming in and saying you have to implement something like this, and whether it is implementable or not is just tough luck. If you can't implement it, the whole system is bowed.

DR. ROSENBAUM: Well, that certainly is not our attitude. But I must say, in all fairness, that this wasn't developed by any conspiracy of people --

COMMISSIONER HENDRIE: No, I don't think it was.

DR. ROSENBAUM: -- with a particular interest. It actually was developed by me.

COMMISSIONER HENDRIE: I know what you mean, and if you and I were the only guilty parties, why I wouldn't have a problem. And I am sympathetic to the general proposition that to the extent you can provide a fall-back in case you make a mistake, or find later on your knoweldge gets better and you say: Gee, I should have bored them horizontally instead of vertically, or squared instead of round, or what have you, or in the salt instead of granite, whatever, if you can retrieve it, why that is fine.

But what I think would be unfortunate would be in the interests of trying to provide that kind of flexibility

and resilience in the system, one began to build in sets of requirements which become available to other parties then to use as, in effect, insurmountable barriers. That is one of the great difficulties in trying to move forward in this field.

DR. ROSENBAUM: I think that is a good point, and we will certainly consider that. I don't think that that will affect what we will propose -- which has already been turned over to David Hawkins -- but it is certainly something we will consider seriously in the hearings when we go towards a final.

CHAIRMAN AHEARNE: Why don't you now speak to this chart you have.

MR. EGAN: David asked me to put together a chart that is in substance an outline of our own internal review process, which I've done here. I want to emphasize that this is not a schedule. It is not really quite cricket to add up those little time periods on the right and say that it will come out as a borne child at the end of that period, because there are review processes, and David's own review process, that are in little bullets on this chart.

What we have just recently completed, the agency has a procedure that is set forth in its implementation of Executive Order 12044. One of the initial steps is an interagency, intra-agency work group composed of representatives of all the various offices within the agency that simply help us develop the standard and provide us with comments. And on this

issue, as you might guess, we get a lot of feedback, and a lot of discussion with offices that normally don't interact with us terribly heavily.

We just recently completed that process, and we are now getting ready to put the formal package together to send it to David and up to Mr. Hawkins for his consideration. And if he is reasonably happy with it, he will then start the next part of the process, which is the EPA Steering Committee review, which is a board with senior representatives from each assistant administrator's office that acts effectively as a super work group. They essentially get together to meet and discuss the standards, issue any action it ms that they feel should be changed in the package, and send it back to us to go further in the process.

Obviously at an stage in this process, there are lots of opportunities for endless iterations. We think now that is probably less likely because of the length of the work review itself.

Once the steering committee is finished with it, it then goes, again after Mr. Hawkins' approval, into what we call our "red border review," which is not a meeting, per se, but where each individual assistant administrator has to concur or nonconcur with the action.

Once that is completed, it goes for signature to the Administrator. The two time periods indicated, and which

we have discussions with our planning and management shop, the ones on the left are the book numbers for the way the procedure is supposed to work, which typically is vaat you might achieve with minor regulations.

We both guessed that at least a factor of two would be needed for this particular regulation, and I think the possibility for iterations between those staffs may increase that by some specified factor.

Once the Administrator has approved it and the standards and guidance are proposed in the Register, we will then of course start our public comment and hearing phase.

Now we plan to schedule that initially for 180 days. In that period of time, we will hold a number of hearings on this topic as well as we will on the other actions we've talked about today around the country.

I suspect of course that the comment period will in fact drag on quite a bit longer than that. A lot of issues will probably come up from the hearings that people will want to be pursued further.

Also, in parallel with all of this, at the request of Congressman Udall, we are also planning to establish an independent technical review of the material supporting the standards by our Science Advisory Board. It is similar in concept, although not necessarily similar in organization, to the Lewis Committee that reviewed the Reactor Safety Study.

That will be ongoing approximately in parallel with all of this.

After we get done with all of that, of course, we will be revising the package. We then go through the same internal review process again -- work group, steering committee, red border -- and I'm sure we'll find that all the people that were in the old work group have moved on, and we have new people in the work group; that's the way these things always work. Three months is probably what we might be able to achieve, if we all love the standards after public comment, which is not likely -- just because, if nothing else, the complexity of the issue.

And then after all of this, we will promulgate final regulations. And of course, since we have a split package here, this is a somewhat complicated route, because you're talking part of the package through a different process, through the White House approval, and the other part of the package would come through the Administrator.

That is where we are.

CHAIRMAN AHEARNE: Good luck.

Joe?

COMMISSIONER HENDRIE: If some of these waste bills go, why --

DR. ROSENBAUM: We would be relieved of our duties.

COMMISSIONER HE DRIE: We're going to sort of run

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late, I guess.

CHAIRMAN AHEARNE: David, thank you very much, and all of your people. I am just delighted to see these things coming through, and I hope that we can both provide whatever assistance is appropriate, and also continue to work with you. I know we have been working fairly closely on this thing, and I thank you very much.

DR. ROSENBAUM: Yes, sir.

COMMISSIONER HENDRIE: Very good.

CHAIRMAN AHEARNE: The meeting is adjourned.

(Whereupon, at 4:29 p.m., the meeting was

adjourned.)

NUCLEAR REGULATORY COMMISSION

	Commission Meeting
in the matter	of: Briefing by FPA on Draft General Standards (OCCUPATIONAL EXPOSURES, MILL TAILINGS) Date of Proceeding: September 3, 1980
	Docket Number:
	Place of Proceeding: Washington, D. C.
were held as h thereof for th	erein appears, and that this is the original to the file of the Commission.
	Jane N. Beach
	Official Reporter (Typed) Ane N. Sench Official Reporter (Signature)