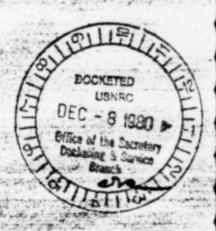
ORIGINAL

Transcript of Proceedings

UNITED STATES NUCLEAR REGULATORY COMMISSION

40-CFR-190



Silver Spring, Maryland

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PROCEEDINGS

MR. MILLER: Thank you for coming. We called this meeting with the purpose in mind of informing our licensees of the NRC of the steps that we are taking, and the steps that you will have to take to implement the EPA's environmental fuel cycle standard 40 CFR 190 which goes into effect for uranium mills on December 1st.

It is a meeting intended primarily to inform our licensees. There is, of course, broader interest so we have consented to others being present.

The meeting grows out of an awareness on our part that as we get nearer and nearer to December 1st that more and more questions come up. Concerns get raised about implementation of the standard, and we thought it most appropriate and most effective to have you together as a group to lay out the program, both what we will be doing in the near term, December 1st, and over the long term on an ongoing basis to implement the standard.

In laying this out for you we hope to show how the program we have established or will establish recognizes and accounts for the concerns and questions that have been raised.

There are several petitions before the NRC which effectively request that the standard -- that the enforcement and implementation of the standard be stayed.

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The theory is that because there are also petitions before the EPA requesting reconsideration of the standard as it applies to mills, that would NRC stay implementation.

on December 1st. The lawyers of NRC, several of which are present, have concluded, or are about to make a recommendation to the Commission—and Sheldon, you can help me out—the gest of it is that it's a valid Federal regulation and, therefore we are duty bound to enforce it.

I am sure I didn't get it exactly right, Sheldon, but I will let you, later, give a more fine tuned answer.

What I am trying to say is that we are not here to discuss the merits of the petitions before the EPA. We want to talk about our implementation program, and I will avoid answering questions that relate to that petition.

My name is Hubert Miller, and I am the head of the New Uranium Mill Licensing Section. I will be giving a presentation, a briefing, of the program. To help me on that is Greg Eadie of my staff who is the lead person in implementation of the standard; and also John Linehan who I am sure you all know is the Section Leader of the Operating Facilities Section.

Also in the room to assist if they are needed are members of the Office of General Counsel of the NRC, the

Office of Executive Legal Director. That's Sheldon
Trubatch, and Bob Fonner respectively; INE, I believe Doug
Sly. Doug, are you here? If he is not, he should be here
shortly.

The Office of Standards Development, John Hickey,
Don Harmon; and other members of the Uranium Recovery
Licensing Branch Staff to help us on this presentation.

What I had in mind as far as an agenda was to give a presentation. It takes a few minutes to describe the program. I will beg your indulgence while I am giving the briefing.

Unless you have a question where you just don't understand what I am saying, it's just not clear, I would ask you to hold questions until I can go through the presentation.

Then I would like to entertain questions and discuss the program with you.

I want to back up one minute and thank the

American Mining Congress for helping us with getting notice,
getting such short notice to people.

I should mention right here that the American
Mining Congress has requested that the meeting be transcribed.
We are a public agency. We have no problem with that.

Before we progress any further though, I want to see if there is any overwhelming objection to transcription

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of this meeting. If there is, we won't allow it.

(No response.)

MR. MILLER: I don't see that there is any overwhelming objection.

QUESTION: Do you have any arrangements to get some more of the handouts?

MR. MILLER: Yes. I was going to get to that in a minute. Apparently you don't have enough. More copies are being made.

There are two handouts, one of which is thethe material has some text. It's the description of
procedure for compliance determination I think. We have
limited copies of that... If you could share copies.

If I just talk through the briefing it would take about one-half hour, but I expect there will be some discussion and question. So, perhaps it could take several hours.

I hope that the ventilation holds up long enough, if it's not already unbearable. I don't know what we can do to make it any better except to just go quickly.

After the meeting I believe what we present will be all you'll want to know about how we are going to implement 40 CFR 190 and what you have to do. There are a few facilities where we would like to review site specific points with you, and we can meet with you afterwards.

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After our meeting there are a number of licensees, and we will seek you out--perhaps we will have a break and we will contact you--but we want to make some time to meet with a few of you, and then anybody else who has questions beyond what we present here today are welcome to meet with us.

One word about security. I have been asked to say that you have got to have this badge that you have been given visible at all times, and you can be on the first floor and you can be on the basement level without an escort.

If you come up to see one of the project managers or come up to our offices for some reason, somebody has to be escorting you.

There are two pass-outs. Hopefully you will have a copy. If you don't -- There is a copy of procedure for compliance determination and then a copy of the viewgraphs that I will be talking from.

MR. TRUBATCH: My name is Sheldon Trubatch from the Office of General Counsel. I just want to emphasize, especially because this meeting is being transcribed, that this is an informal meeting with the staff and I believe most of you now have a copy of a letter which reflects the Commission's views on the Commission's contemplated enforcement of this regulation.

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MR. MILLER: On December 1st the standard does take effect at uranium mills, and we will essential begin the implementation through orders which amend licenses, and I will get into more about what those orders are in a minute.

At the same time that we are issuing the orders we will be issuing a report which will be the supporting regulatory document for the orders.

Sometime in the near future--it probably won't be December 1st -- but sometime in the near future 10 CFR part 20 will be revised to incorporate the requirements of 40 CFR 190. That was proposed a while back, and it is near the point where it will be finalized.

As with any new standard, it is not possible to immediately be to the point where you are operating your determine and compliance. You are carrying it out in the way that you, after a period of time, would be doing.

The program that we have outlined is-- it involves a phased implementation of the standard.

Just to briefly give you an overview, we have been, for the past several years, begun the process of evaluating through our licensing action new mill licensing renewals, the performance and operations of mills in terms of how they would meet 40 CFR 190.

For those facilities that that was not done on

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our four months completed radiological assessments much like those we have done in licensing cases, with the objective in mind of identifying potential problem areas, highlighting those areas requiring environmental monitoring to essentially establish a base upon which to begin implementation of the standard.

On December 1st we will issue orders which formally invoke the standard. Then there will be a phase, the compliance determination, the problems of implementing the standard will be worked out.

A second phase will then be ultimately reached or achieved, and I will go into more of this in a minute, where we reduce the problems of compliance determination to hopefully simplify procedures.

There is going to be a period of time when there is more intensive monitoring, and then later there eventually will reach a phase where it will be not a large commitment.

Let me say a few things about the conclusions of the report that we are working on that will essentially kick off the standard.

As I mentioned, we formed objective assessments of each mill. The conclusions are that with the existing requirements that has been established at mills through the renewals and through new licensing actions, the

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commitments made under the interim tailings performance objectives, controlled blowing of tailings and whatnot, should assure that 40 CFR 190 is being met at each facility that no specific additional measures beyond the requirements already committed to appear at this time, or have been identified as being necessary.

I think what I am trying to say is on December 1st the order won't say as of today you've got to do this, that or the other thing in addition to what you have already committed to in the past.

However, our assessments have involved both predictive modeling as well as consideration of what environmental monitoring data we had available.

We have identified the need for additional information in many cases before a further conclusion about compliance can be made.

In many cases it involves us screening out extraneous sources. In other cases it is incomplete data, and many cases it's uncertainty about the effectiveness of the controls that have been previously committed to.

Let me say a few words about what the standard covers and what it doesn't. First, it's 25 millirem limit to a whole body or any other organs, and there must be reasonable assurance, and I underline those words, that that is being met.

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It includes routine releases from normal operations. It doesn't include releases from an accident.

It's an annual average. It isn't the sort of thing, at least in our implementation program, where if you got in one quarter greater than the 25 millirem limit you are in violation. It's an annual average. I will get to more in a minute how that works out.

Cumulative mill sources. It involves the exposure to the public to not only your mill, but one that might be nearby. What it excludes, of course, is radon and its daughters, natural background radiation, mining operations and associated activities. This could be stockpiles and whatnot that are around the mines.

It does not include transportation, it does not include decommissioning and decontamination. IT does not include releases prior to 1980 and the ground contamination that's associated with that.

As I mentioned before, we were talking about phase implementation. It is going to take some time to get to the point where it is reduced to simple operating procedure.

The objectives of our compliance determination program is to achieve at each mill a simple standardized assessment procedure with correspondingly simple reporting requirements.

The objective is to assure consistency between

mills and to have consistency over time. We don't do
it differently from this month, or this six month period
to the next.

Eliminate uncertainty. The minds of the public are saying are the mills operating within the limits or not; do the mill operators reduce exposure on the part of the operator and any kind of uncertainty that might exist in the public, in the minds of the regulatory agencies which would be the NRC; and uncertainty on the part of our Federal agencies and State agencies that are interested in the way the mills operate.

Another objective is to facilitate prompt identification of problems where they exist. You avoid a situation where you have to go through a complex procedure in order to get to the point where you determine that there is a problem; and to minimze the costs and staff time for you and for us.

The primary basis for the compliance determination will be actual monitoring data, not computer codes. The emphasis will be on measurements at the nearest residence, the location of concern.

Codes. Let's take Mildos for instance. The kind of code that I'm talking about is one that includes estimation of cource terms, atmospheric transport and dosimetry.

As anyone who is familiar with these codes knows

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that there is a good deal of uncertainty about estimates of source terms; considerable uncertainty about atmospheric transport, particularly if there are terrain effects.

For these reasons it is most appropriate to not go through the process of computing that concentration at the nearest residence, but to measure it.

Now, while the code isn't our primary method of compliance determination—of course there is value and necessity for codes. We obviously have to use them in licensing action because we have an obligation to predict the potential impacts of the licensing action.

So, on a prospective basis protective codes are a must. They can also -- they are also helpful in developing environmental monitoring requirements and identifying potential problem areas.

As we talk more you will see them actually aiding in the interpretation of the environmental monitoring data that will be gathered in the programs that are established.

Very simply, the program involves having an environmental monitoring program in place, and these are the environmental monitoring programs that we have established for ALARA and NEPA reasons over the past three years as documented in staff positions several years ago, and most recently in our reg guide 414, to gather the data at the nearest residence, to subtract out the contribution from

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background and other sources.

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In most cases that is going to be one of the things that causes us to spend some time in phase one.

I will get to that in a minute.

To compare the dose with the standard and determine compliance, identify any additional controls that may be necessary or if appropriate, possibly some additional monitoring to supplement or confirm a problem if it is first identified. Then to report, of course, periodically to the NRC the compliance determination or compliance assessments that you do as mill operators, because the burden is primarily with you for that determination.

This is part of the package, the figures on procedures that was handed out. Once we get it our project managers will be reviewing your reports.

Of course, if compliance is determined it's no nevermind. Some action would have to be taken obviously where there is a potential problem that arises, or if there is unusual conditions under the provisions that were established at EPA in setting the standard for variance. That, too, is another pathway that can be taken.

We expect to annual be issuing a brief report that summarizes the status of the industry.

Let me talk about where we are headed as to what we have in mind, or what our objective is. It is to

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establish that simple standarized procedure which would involve establishing some point of receptor concentration or dose action levels. I am not sure what it would be.

It may be 25 or higher than 25 millirem, but it will be developed based upon the experience of going through phase one. It will be, of course, combined with a commitment of specific control measures and to acknowledge that there were no significant changes in the surrounding environs—that there wasn't a new mine that opened up nearby and so on.

We expect to establish eventually some point of receptor concentration or dose action levels. We would do that—we would enter from phase one to phase two with some license amendments, because your initial orders or licenses would not be set up in such a fashion that you have such levels.

Back to the pre ious slide, the trick is going to be in subtracting out the other sources that are not covered by the standard.

Greg Eadie, I will have him describe to you one case that we have recently gone through where we have, in fact, calculated doses of higher than 25 millirem, but through observation and through evaluation of environmental monitoring data and so on we have been able to determine that it is most likely that it's not the mill tailings that are causing the exceeding of the 25 millirem.

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During phase one we expect to be completing the implementation or the installation of environmental monitoring programs that have been committed to by and large up until now.

I think there may be only one operator who does not right now have an approved environmental monitoring program, that is one that is consistence in performance with the reg guide and that will be done shortly.

During that period of time we are going to be sorting out-or you will be sorting out other contributions to the dose at that nearest residence.

We may, in certain cases, be working with you to establish some short-term limited sampling and in studies, but the outcome will be, or it is our objective and our hope of simplified procedure.

Now, the orders. On December 1st they will, where there is not an environmental monitoring program in place, it will effectively establish one.

In some cases we have committed, or we have nailed down our worked-out environmental monitoring programs over a period of time. It has been over the past three years.

There may be a few that need some minor modifications to be consistent with the regulatory guide.

We will specify where there are not environmental monitoring programs in place and operating right now, we will

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specify some time frames, dates, certain by when the program has to be installed. We will be envoking quality insurance requirements where they are not now envoked.

We will be committing operators to the dose assessments that are described in the one handout, or the procedure that you have a copy of, and it's Appendix A of that document which effectively takes the dose conversion factors from the regulatory guidance we have out on that.

The number on that is 802-4. I think most of you are familiar with that.

The objective there was to try to make it as simple as possible to compute these doses from the concentration, and to establish reporting requirements.

The next major category of conditions that would be in these letters are call for short-term studies. Greg will give you an example of what that might involve.

We will be looking for at each site where there
is not already a year's worth of meterology and met data
or an accumulation of a year's worth of met data, identification
and characterization of all significant nearby extraneous
sources. That is part of that sorting out that I talked
about earlier.

Over the past several years the commitment was to a program for controlling the blowing of tailings and dusting from diffused sources.

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We will do what we have done in the recent cases, and that is to formally require written operating procedures. I think that is standard everywhere. We will formall require the written operating procedures and weekly inspections to confirm that those procedures are being following.

The EPA standard is a strict one for mills. I don't think I would have anybody in this room argue that with me.

However, for the past several years we have upgraded programs. We have, under the performance objectives, required the control of the blowing of tailings and dust control.

Our final regulations issued on October 3rd.

Again, state that requirement flexibility as to how you do it. But, whatever you do, document it in some way to give it a discipline.

I'm thinking primarily of the tailings pile. You can't put an automatic control device on the tailings pile.

It takes management attention. It takes discipline, and that is the reason for the written procedures.

The primary emphasis must be on a mission control and secondarily on controls such as moving the fence, or moving the residence; although, that may be, in some cases, called for.

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I will review reporting requirements before I ask Greg to say a few words. I think during phase I you've got the picture, or hopefully you have the picture of having to sort of work our way through the process of implementation. We are going to be learning and during that period of time we are going to want more frequent reports than the semi-annual report that is called for in the procedure that you have.

Eventually when we get to phase II that will be the requirement. Routine reporting would be as you get the quarterly data that is required to be analyzed by your environmental monitoring programs, that that would be submitted to us.

The non-compliance reporting requirement that is in the part 20 regulation change that John Hickey is now finalizing would not apply. That would not apply during this period.

However 1 an ongoing basis in phase II it would be a requirement 30 days after identification of a problem that you would report that.

What I will ask Greg to do is take a case, a real-life case. We won't name who it is, but you can probably figure out. The calculation alone of taking monitoring data at the nearest residence, or even via the code, you come up with doses that are in excess of 25

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millirem.

I should mention something about the code.

We have done what is prudent in licensing, and that is to use conservative assumptions and simplifications on the predictive analyses that we have done.

There have been many cases where we have computed in the licensing action a dose that is close to 25 millirem, and in some cases over. But we recognize that there are no simplifications in the use of the predictive code, that there are these conservatisms and so we find that it's still reasonable to go and making a finding that is likely that the mill is meeting 25 millirem.

Greg, why don't you run through this case.

MR. EADIE: I have to work from the hand-out, so I would like to start the discussion with the map. It is about half-way through the first handout.

This procedure I am about to describe we have gone through for all our license facilities. I'll take the time and explain the procedures.

First of all, the map of the site is a typical site that has many influences, some of which contribute to the nearest residence exposure potential which are not covered under 40 CFR 190. For example, mining activities ongoing in those areas.

When they are around the mill site, even close

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to the nearest receptor, you may have mine overburden, poor storage pads and so forth. The mill influence and the tailings piles under reg guide 4.14 will have nearby particular monitoring stations. We are also working to the closest receptor. In this diagram we call it a camp to help particular monitoring stations set up and operate continuously.

So, this is a typical diagram of a mill site that we are looking at.

The next page, mildos computer code generated

50 year dose commitments. Let me try and describe our

initial assessments based on the computer code. It is the

mildos computer code. It is a 50 dose commitment.

the remaining 50 years what the dose would be. We break it down into two pathways. We call it the direct exposure pathway due to the actual inhalation of the air-borne particulates, and also external gamma radiation due to cloud immersion and the deposition of this material on the ground and the direct exposure to the gamma.

We focus on the uranium, radium, lead. We are not dealing with radon and its daughters.

The second pathway we talked about is the ingestion exposure pathway, and this is the pathway, as you can see in the table, has the greatest potential exposures.

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However, this is where the code seems to break down and we have the poorest accurate assessment of input parameters. For example, the met data is very limited We don't have, for example, in many cases four years worth of met data.

Also, we are finding out that the closest receptor, one might have to go in and get additional met data at that point.

We have made assumptions, for example, on the vegetable intake pathway. If there is a potential for locally grown vegetables, that all vegetables would be grown there and consumed at that point.

The same for meat ingestion where you have cattle grazing. You pick the highest potential grazing area that the animal has the greatest up-take. The individual at that local consumes that meat for the entire year.

So, this is the mildos of the predictive model that we go through initially to give us an idea of what the potential exposure at the nearest receptor is.

Any questions on the mildos assessment?
(No response.)

MR. EADIE: Okay, on the next page, for this particular facility and several others we have been able to review actual environmental monitoring data. This is where the instrument and air sampler has been set out.

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It has been running continuously. We have a quarterly analysis of the samples reported.

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We have taken a look in this particular location at two off-site residences. For the air-borne particulates, uranium, radium, thorium and lead, these are excellent reported data now, concentration on pCi/m3.

The next one gives you the dose factors that we have applied. These were also given to you in the other handout which describes the procedures, that is the inhalation pathway of those conversion factors.

You simple multiple the uranium natural, the air-borne concentration, times the dose conversion factor for uranium whole body and the volume is .0656. It is a 50 year dose commitment for one year of exposure.

The point here to make is that if one looks at the bone or lung for example at the nearest residence south, we see a value like 61 for bone and 124 for lung. This is based on actual data.

Looking back at your map you can see it is more than likely that this exposure is not caused solely and not predominately by the new operations.

The most influential exposure pathway is probably for mining activities, from transportation activities throughout that area.

So, what we've had to do is go back to the licensee

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and, if you will skip about two pages it is example E, we have gone back trying to look at the actual data.

MR. MILLER: Let me back up just to talk for a minute on this map. You can see that the prevailing wind is out of the south, southwest, and the point of receptor here is over to the right. I believe there are sections there, so you are talking about a mile or more to the east of the tailings pond and the mill, and on the other side of a topographic high or anticline type of formation.

It doesn't take an expert to conclude from this that the primary source for concentrations at that camp are from mining activities and not from the tainings pond.

In addition to that, we know the conditions of the tailings and by and large they appear to be--I believe they are in pretty good condition. I mean, we know that there is not a lot of blowing going on.

So, it's a case where if you calculate and take the data, the raw data from the nearest residence, and you calculate a problem. You can see it through this process. Any reasonable person would conclude that not necessarily, and most likely not.

Let's have Greg talk more about how we are going to confirm that conclusion.

MR. EADIE: This is the example EMP modification.

First of all, to go in there and do an inventory of materials

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in the area. For example, identify the mining areas, any areas of mine over-burden or storage pads. Perhaps go into characterizing by giving the volumes, the heights, the concentration of radioactive materials in that area.

Also, we have asked the licensee to establish the supplemental air monitoring program. Not an additional station. Move one that is in existence for a brief period of time to a location between the mining activities and the closest residence as a short term sampling, five weeks, maybe two months, limited analyses perhaps on uranium and radium.

In this case since we are trying to identify contribution from mining activities, or we want to look at the ratios of the radionuclides, so we have asked the licensee to look at all parameters.

As a matter of fact, if you go back to the actual environmental monitoring reported data, you would see that uranium is quite high in comparison to the other. It seems to indicate that it's not tailings. It is probably an actual material, for example, ore.

Back on what we have asked them to do for a short period of time to help us sort out the contribution at the nearest residence.

Correlation of met data. Here again, with the intervening terrain, the mill site and so forth, we have

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asked the licensee to put portable met station for a limited time to correlate met data, wind speed and direction with the actual measured air-borne concentrations at the nearest receptor.

Of course, all of this information that we will get in, we will review it and reevaluate the dose estimates. That is pretty much the procedure we're going through.

MR. MILLER: It is awfully warm in here. Let's take a break for ten or 15 minutes. This will give you some time to think about what we have just presented.

We may have a few more things to present when we come back, but after the break we will open it up.

(Whereupon, a short recess was taken.)

MR. MILLER: I apologize for the warmth of this room. It is not a tactic to cut down on the number of questions.

What I would like to do is entertain questions.

There is a sign-out sheet that wasn't at the door when you first came in, but it is now near the door. Would you please sign in on your way out or sometime before we break up.

I don't know how long the questioning is going to go. That's really up to you. We are here to make sure that you understand what it is we are doing to implement this standard.

So, as long as you desire, if it runs on for a while, I think we ought to take another break, but we can

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decide that later.

With that, are there any questions?

QUESTION: Are there any cases where the NRC will take meterological data and look at the site and say, based on a mildos calculation, they show that the nearest residence, wherever it is, is like 10 or 15 millirems a year?

In that particular case will you still request that we go out there and put a sample at that location in order to determine the fact that it is?

MR. MILLER: We began about three years ago.

As I said before, for reasons of ALARA and for reasons of our NEPA responsibilities to require environmental monitoring at mills.

One element of that is to determine at the point of nearest receptor where the potential greatest impact would be, what kind of impact is, in fact, happening.

so, the answer to the question is that you've got to have that kind of program anyway. It happens to serve the purpose of determining compliance with 40 CFR 190. Those are programs which were being instituted two or three years ago.

So, the answer is we've got to have an environmental monitoring program, and that includes, of course, the point of receptor sample.

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QUESTION: I notice in your handouts that you are listing lead 210 which is a radon daughter. I infer from that that you are going to count lead 210 for the compliance of 40 CFR 190, and I wonder how that can be since 40 CFR 190 excludes radon and its daughters.

MR. EADIES: It is also in the natural material ore. As a particulate we will include it, of course recognizing you cannot distinguish the particulate versus the lead grown in from radon and decay.

QUESTION: 40 CFR 190 excludes radon and its daughter, and lead 210 is a daughter. How can you attempt to enforce 40 CFR 190 with respect to lead 210?

MR. EADIE: It is also in the ore, is it not?

QUESTION: That may well be the case, but the regulation doesn't apply to radon and its daughters.

MR. EADIE: Lead 210 is in the ore.

QUESTION: So is radon, but the regulation doesn't exclude radon and its daughters except for radon and its daughters which are in the ore.

MR. HICKEY: I'm John Hickey from NRC Standards.

I would suggest that we not give a final answer, because

I don't think that there is unanimity of an agreement that

lead 210 is included in 40 CFR 190.

Unless we could attempt to come to an agreement on that, the way I would like to leave it is we expect

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you to monitor for lead 210. My personal opinion is that lead 210 is not 40 CFR 190, but it is of concern as far as protecting the public from the 210.

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Paul, would you say that lead 210 is covered by 40 CFR 190? I would suggest that we leave that question open-ended.

MR. MAGNO: It is clearly ambiguous the way it is stated.

QUESTION: I don't see that it's ambiguous. It says excluding radon and its daughters. If lead 210 is excluded from the calulations --

MR. MAGNO: I say it's ambiguous with respect to the intentions.

MR. MILLER: In a plain reading it seems to be excluded. I think that is one we will have to, as John said, not give you a definitive answer right now, and we will consult with the EPA as we have done all along.

We have laid out this program and we have kept in touch with Allen Richardson and Paul Magno and the folks at EPA who were responsible for the standard initially. We will touch that base.

QUESTION: The differentiation between background and particulates arising from the mill, how did you arrive at that? To begin with, what was the background level and what kind of variation did it have?

MR. MILLER: Let me attempt a general answer and then, Greg, why don't you add to it.

Environmental monitoring program is more than just a point of receptor. I'm talking now about air sampling. It includes a background monitor. It would be judicious in selecting a location, but you want to try by that background monitor to pick up all those sources that would be present in the environment near the mill, but would not include mill tailings. So you have to look at it on a site specific basis to determine where is the best place to put that.

In addition, there are monitorings around the tailings and around the boundary of the mill. It is a combination of looking at various monitors, and knowing prevailing winds on an annual average basis, on a basis that's short. It is not a grab sample. It is continuous monitoring.

You should be screening out the short-term kind of effects that make grab samples not appropriate for our purposes.

I don't know if Dan or Greg could say something more specific about the general trend in the monitors in this case here. Greg or Dan, do you want to attempt -- Wasn't there some falling off in the direction going from several monitors down wind from the mining areas?

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If it were the tailings pond or the mill, you wouldn't to have the monitors further away from the miller. You would have higher readings than the ones near the mill.

It is a process of looking not just at a point of receptor, but a number of different points and not just the environmental monitoring data. It is looking at other things. It is an assessment of all of the circumstances that are given. They are going to be required to sort out and portion out the contributions to the dose.

QUESTION: 40 CFR excludes the contribution prior to December 1980; is that right?

MR. MILLER: That's the way we are going to implement it.

QUESTION: How do you determine the vegetation and the livestock to what they eat? Do you take it to what they ate prior to 1980, or do you go out and sample the cattle now and then sample it again after December 1980?

How are we suppose to determine what impacts we have before and after?

MR. MILLER: That is part of this process that

I'm talkingabout in phase I through step by step figuring

out how to sort these things. I can't give you a real good

answer right now. I haven't thought that one through, but

perhaps somebody else has an answer on how to do that.

QUESTION: During the rule making proceeding I think

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NRC indicated that physical monitoring was not possible.

Has something come up that has changed this? I thought

NRC had indicated this was impossible to do, and I'm trying

to find out if something came up in the meanwhile to change

your mind?

MR. MILLER: I won't attempt to speak for the people who are speaking for the NRC three or four years ago.

Based upon our licensing experience and based upon the expertise that we have on staff and with our consultants, we conclude that it is feasible.

You can see this program is one that is phased.

It is not one where we have all the answers right away. The objectives are to streamline, to standardize it, simplify it.

There is going to be some time during which we are sorting things out to some extent.

QUESTION: I would like to explore for a moment the question of your licensing experience in the past two or three years.

How many applications for renewal of existing new licenses has NRC examined in the past two or three years?

MR. MILLER: Ross, did you hear that question?

MR. SCARANO: We have gone through all but one
of the milling operations. It is quantified at the five
to somewhere around eight.

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MR. LINEHAN: It is probably about between six and eight that we have been in the process of reviewing. We have two right now that are pending renewal. I'd say it's about six to eight of the mills.

QUESTION: It is NRC's contention that it has actually monitored or required the mill licensees to monitor to detect how well they are presently in compliance with part 190, the license renewal applications?

MR. LINEHAN: We have monitoring data from a few sites over a period of a year or greater now on which we are making this determination.

MR. MILLER: What are you driving at?

QUESTION: All I'm trying to do is explore the foundation for the statements that have been made earlier that NRC has -- well, you have expressed confidence that the mills will be able to obtain the 40 CFR 190 standard on the basis of your experience with license renewals.

I was interested in seeing how much experience

NRC had had with license renewals and how much your confidence
was resting on things like mildos?

MR. MILLER: I think what I said is with the controls that have already been committed through renewals and through renewal licensing, In many cases it is just simply put as the control of blowing tailings and other control requirements that are consistent with ALARA under part 20,

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and consistent with NEPA that those requirements should be adequate. There are cases in our states where we have analyzed the situation and should be adequate to meet the standard. We can't make it as a firm statement.

As I said, some cases we are looking at are addition studies to be done to confirm that preliminary conclusion. It will require strict control of blowing the tailings. It will require the short of things that have been at some mills where clearing materials have been placed over embankments, embankments were made of tailings, chemical stablizers had been used, flexibility in the way you do it.

What we are saying is with the commitments that were already made, those were followed through and it should be doable.

QUESTION: So, the NRC's position is that if this licensee implement the controls that NRC is posing on the license renewal process, then NRC believes that they will be able to comply with part 190?

MR. MILLER: Subject to what I said before.

That's what it looks like to us. It's likely that they will.

The environmental monitoring programs will tell us for sure.

QUESTION: I would like to ask a technical question as to what extent NRC has done to document the confidence limits they can put on these measurements? They seem to

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be very down in the range where they may be very difficult to make up with a high degree of accuracy.

MR. EADIE: I will help out and I will ask John Hickey to help me. I think one of the big things that we would like to stress in the environmental monitoring program is establishing a good quality assurance program, not only on simple collection, but particularly on sample analysis. We do have a reg guide on that, 4.15.

Perhaps if John could help me out.

MR. HICKEY: I will answer the questions in two parts. We looked very hard at whether you could take a sample and analyze it to a certain degree of uncertainty within reasonable uncertainty limits. We feel that samples can be collected and analyzed using good analytic techniques within reasonable certainty.

Now, as far as our certainty that that result will be translateable into a dose that a real person is getting with a degree of certainty, there is much more uncertainty associated with that.

So, when you talk about uncertainty you have to differentiate between the technical analysis and the conversion to a dose.

MR. MILLER: What we are doing is essentially establishing in that Appendix A of the procedure that you have which, again, is derrived from or taken from our

reg guide on dose modeling, the dose conversion factors that we are going to use.

So, that in a sense eliminates the uncertainty as far as getting from those concentrations you measure to what we're going to be looking for. Those are dose conversion factors that have been in regulatory guidance and have received scrutiny.

We feel it is a very good regulatory basis upon which to be using those factors.

QUESTION: What I am asking for is, in fact, a numerical number that you people say. When you say 25 millirem, based upon these measurements, to you feel it's accurate plus or minus 5, 10, 15 or 100 percent, based on your experience?

MR. HICKEY: I don't feel comfortable answer that question. The issue of uncertainty has not been dealt with. It is a generic issue. It has not been dealt with by the commission. It is not dealt with in the regulations.

They say you have to meet these limits and they don't say you have to meet the limits with 50 percent uncertainty, or 95 percent uncertainty.

Because the conversion of a concentration to a dose depends on assumptions as to how the material gets into the body and how it behaves in the body, there are uncertainties associated with that. It is difficult to

answer either in generic terms or specific terms.

MR. MILLER: In a sense our procedure mutes that.

In any kind of standard, any kind of radiation protection

standard that's in term of the dose, you always have

a question of what models to use to get from concentrations
to dose.

What we have done by our procedure is to establish the ones that we are going to use, we expect you to use, that we can defend.

There is uncertainty and I guess we could do studies to try to precisely determine that. By the procedure you are not going to have uncertainty as to use of those and how we are going to react to that, nor do I think you have to worry about anybody else.

We are the regulatory agency. We are responsible for implementing it. So, I think I would feel comfortable in that sense.

QUESTION: This goes to the confidence that you expressed in the regulation. Again you said you were a responsible agency. I talked briefly with your counsel in the hall, and I had some question as to the confidence you have in 40 CFR 190.

If you have such confidence, I would see no particular reason to revise 1020. Would you like to address the confidence you have in EPA's rate?

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MR. FONNER: From the strictly legal point of view, the amendments of 10 CFR 20 is not a very great one to persons who are licensees of the NRC, because under the Atomic Energy Act we can proceed by order with respect to our own licensees.

The 10 CFR 20 amendment is significant; however, with respect to enforcement of the EPA standard in agreement States.

QUESTION: If a facility turns in to you a report that says they cannot comply with that standard, what would be your procedures? Are you going to issue a shut-down order? What will you be doing there?

MR. MILLER: First of all, I will repeat something I said before. The primary burden of determining compliance resides with you. You've got to meet the limit.

In the interest of trying to be sure of consistency and reduce uncertainty, we have prescribed the formulas that can be used, our calculating doses.

As I mentioned on one of the viewgraphs, when you see that you've got a dose higher it is up to you to sort out the extraneous sources. Once you have isolated an idea of what's coming from the mill, you determine you have, in fact, taken steps, reasonable steps, to control the source.

After that kind of study the is provision for

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variance, but it is under unusual type of situation. I forget the exact words, but we would not entertain a request or submittal that did not clearly indicate that a serious good faith attempt was made to control the emissions.

I don't know if I'm answering your question, but I can't give you an answer, quite frankly, on a generic basis anymore than I have.

DUESTION: Your proposed regulations would have been 10 CFR 20 on coal for the licensee to ascertain the extent of exposure of individuals, levels of radiation and concentration of radioactive materials involved, the cause of the exposure, and the plan to implement the course of cor the exposure, and the plan to implement the course of cor the exposure. That's all that is required for this 30 day report.

Does NRC have any experience to believe that a mill could provide all that data and prevent a reoccurence in 30 days?

MR. MILLER: Go back to what I said about reporting requirements. We are trying to recognize—I think the program we have outlined recognizes the kinds of concerns that you are raising. During phase I there is a lot of sorting out to do and we don't think that you can get it down to a point where you can quickly determine whether you are incompliance or not.

During that period of time we expect -- Say the

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regulation that requires a 30 day notification would not be applicable during that period of time until we can amend the licenses and get very specific about it and get that more simplified procedure, it is not going to be possible to do that.

After that time we hope that the program will be tailored to allow for rapid determination of a problem if it exists. That is one of the objectives.

After that time we feel it is.

QUESTION: At present what level exposure exists which would necessitate a report to NRC of non-compliance?

Do you know?

MR. MILLER: During this phase I, as I said before, there will be none.

OUESTION: Is there anything in the present regulations which imposes on uranium mill a requirement to report over exposure at any particular level?

MR. HICKEY: Yes. There are requirements. First of all, there are requirements for various facilities depending on the facility. They might have to file a 30 day report at levels of a few millirem a year for nuclear power reactors for uranium mills. They are only subject to requirements that apply to all licensees that include short-term emission levels, that are more on the order of rates of 100's of millirem per year.

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The reason we have been going to the lower levels is because 40 CFR 190 talked in terms of doses of a few millirem a year. There are no radiation standards in existence anywhere that I know of other than 40 CFR 190, and some requirements for reactors that talk in terms of doses of a few millirem a year.

QUESTION: I guess -- It seems to me it's hard to put your finger on what the problems are, but we've got a set number, 25 millirem, to meet. We've got uncertainty in the statistical conference of the monitoring programs.

I don't think anybody is sure that you can separate out what comes. We haven't resolved the questions of what we are going to do with radon yet.

If there is no environmental monitoring program in place, we are going to use the predictive models which are recognized as inadequate. Yet, we still have to meet a 25 millirem standard.

It sounds like it is completely open-ended and you probably won't have -- I think it's not accurate to say -- You are asking us to meet a 25 millirem standard and you are stating whether it is statistically valid or not is irrelevant.

MR. MILLER: I don't agree with your characterization as being uncertaint as what you're saying. There is uncertainty in any kind of measurement you make, and you face it in meeting any standard, whether it's 40 CFR 190, whether

it's part 20, any of the EPA standards.

What I am saying is that always exists, and what we have tried to do in recognition of that is to lay out a program that defines this is what we, the NRC, is going to expect as being that reasonable assurance that is required in part 20.

QUESTION: What I am saying is right now that reasonable assurance is pretty much whatever you decide. There isn't any criterion.

Take for example radon citation from MSHA. They have a variability factor figures in. It's in policy statements and all the operators know when they are going to get cited.

You look at sampling methods with OSHA. They have variability factors built in. You are telling us here we don't have any idea what the variability is.

MR. MILLER: I can't say what that phase II
program is going to look exactly like. We try to characterize
its basic elements, but it very well may have factors of
uncertainty associated with it.

In the meantime we feel we have an adequate regulatory basis upon which to proceed through a phase ; . not to shut people down on December 1st, not to take precipitous actions. We feel we have that basis.

During that period of time there is not going

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to be a requirement for people, as soon as they run out and calculate 125 millirem from the monitoring data at the nearest point of receptor, to report themselves.

At least in our cases it looks like there are not mills exceeding the standard. I think our program, the phase I that I've talked about, accounts for the kind of concern that you are raising.

QUESTION: I'd like to address a question to

Mr. Fonner. I did not understand your real statement

you made relative to 10 CFR 20 amendment prior to implementation

of 25 millirems. Could you repeat that?

MR. FONNER: What I said was because we could proceed by order under the Atomic Energy Act, this program can be implemented with respect to NRC licensees without the necessity of amending 10 CFR part 20.

However, we stand at a different posture with agreement State licensees. They are not our licensees, and it is the expectation of EPA that the standard would be enforced in agreement States by the States.

However, we are, in a sense, an intermediary between the United States Government and the agreement States through the agreement, through the formal understanding, by which we relinquish regulatory authority.

The agreement States feel that they would prefer to have the 40 CFR standard incorporated in our regulations

before they would proceed with their implementation of the programs.

So, that is a matter of the understanding between NRC and the agreement States under the agreement.

QUESTION: Are you saying the next step is you would have to incorporate it as a change to 10 CFR 20?

MR. FONNER: Yes, Sir. Mr. Hickey is working on a rule to that effect right now. It also includes some reporting requirements which would be applicable to the implementation.

MR. HICKEY: I can elaborate on that. Licensing orders have already been written for fuel fabrication plants incorporating the requirements of 40 CFR 190 and slapping the 30 day reporting requirement on it.

If part 20 is never amended, all licenses are still going to be amended incorporating the requirements of 40 CFR 190.

Now, for mills reporting requirements are not going to be 30 days initially. The other reason for revising part 20 is that if we don't revise it, then you can read all the way through part 20 and never see 40 CFR 190 mentioned.

It's a help to somebody reading the regulations to have a cross reference to 40 CFR 190 just notifying them they are subject to those provisions.

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QUESTION: Can you talk more about the amount of time which you anticipate we are going to be in this phase I, and also is that going to depend upon each licensee and how they progress, or is it going to be a generic into that period of time?

MR. MILLER: I showed on the view graph it's going to be a variable time. They are far enough below the standard, they do not suffer source problem, and it should be a short period of time. At mills where there are more extraneous sources, it may take longer. Maybe a year, maybe longer than a year.

We have tried to and have an obligation to not let it string out forever. We are going to put into these orders dates by when these programs have got to be operational.

We are asking for this three month quarterly reporting requirement during phase I so that we can actively follow it and make sure that these programs are progressing, and that we get to phase II as soon as possible.

It is a variable kind of thing.

QUESTION: I would like to stop back on this reporting requirement point, because my reading of the current regulations is that licensees are required to file a non-compliance report if they see some 10 times level specificed in 10 CFR part 20. NRC is preparing an amendment to part 20 right now which would require a licensee to report

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non-compliance with the precise 25 millirem level specific in part 190.

That is an extraordinary departure from past practices, and I don't see that there are any health and safety reasons for requiring that kind of reporting requirement. I have not heard anything here today that suggests to me that uranium, in any way, would be compared to comply with that 30 day period, or it would be possible for them to comply with that 30 day period. Since part 190 itself nowhere requires this report requirement, I wonder what basis, in fact, NRC has, if any, for imposing it?

MR. MILLER; I don't think you did hear what I said before, because I did say during this phase I it will not exist.

In this level that we specified, this action level would not necessarily be 25 millirem.

Now, I won't attempt to answer the other questions about 10 times on one standard and not here.

MR. HICKEY: I explained that 40 CFR 190 is an extraordinary regulation, and the Commission's response to regulations which were strict doses to a few millirem a year, we've been doing it for five years. It is our response to do it for 40 CFR 190.

The 40 CFR 190 is considered a public health and safety regulation and we are obligated to enforce it.

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If we don't have a reporting requirement we have no way of knowing whether or not the licensees are abiding by 40 CFR 190.

QUESTION: You requiring a licensee to monitor and under the regulations, as I read them, you have access to their books, their reports and records?

MR. HICKEY: That's right. We have 20,000 licensees that keep records.

QUESTION: You have about 25 uranium mills and those are divided between the agreement States and NRC?

MR. HICKEY: That's right. We are talking about uranium mills at this meeting, but we are talking about inspectors that have to visit 20,000 licensees.

QUESTION: Given the situation where you have several facilities that might be in close proximity to one another, how does a licensee in one particular facility consider the activities of the other facilities, if at all?

How does the NRC do it, and how does the NRC insure that they are in phase I, and later, that the actions that these several facilities considered together and without duplicating expenses and so on to the licensees?

MR. MILLER: Let me first of all say that most mills are not so close together that the contribution from other mills to its nearest residence, and correct me if I'm wrong here-- I believe that there is not -- The additional

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exclusively the contribution from the mill-- I don't think it is going to be a big problem. It doesn't appear to be a big problem.

However, the standard is such that you've got to consider it. I think how we sort that out is by the same kind of process that you will be sorting out other mining sources.

I think it's a hypothetical and I can't give any real clear answer on it right now.

QUESTION: My question is related. You stated that the burden is clearly upon the licensee to determine compliance and radiation is radiation.

There are 40 CFR 190 sources, and there are non-40 CFR 190 sources. By what standards are they to determine what is and what is not a 40 CFR 190 source? Something measurable that they can rely on. Do you have any idea?

MR. MILLER: The purpose for talking through this case is to show you how, on a site specific basis, that kind of sorting out process will happen. There are enough -- In cases where there is an extraneous source, it is normally quite a few sources and there are large area sources in many cases. It is not a thing that lends itself to some sort of generic formula. It's got to be a

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monitoring programs with the monitors around the tailings pile, and the monitor at the nearest residence should build a strong enough case so that it's not so uncertain that we would be able to meet that reasonable assurance burden.

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QUESTION: That poses a problem. We are talking about 25 millirem standard. He just said over here that this is a very difficult standard. We have a set number. You are talking about with a whole lot of sources. Just sorting all these things out and coming up with 26.5 or 24.9, I mean, I don't think anybody believes you can do that.

There are those sources of uncertainty that I talked about because we have talked about them here today, and there isn't any resolution to those. The operator has no idea, and you say we would be working for maybe a year with some guy trying to figure out a problem.

The example you gave may be an easy one compared to some of the others. Let's say meterological conditions are a little bit more variable, or something of that nature. What is going to happen to this guy if a year later you are still getting readings that are over 25 millirem based on all different kinds of samples?

If you are saying you are not going to shut him down, that's fine. Let's say that in the letter. Let's say

we're not going to shut people down while we are working with them until we have given them adequate time to go through all the reasonable testing procedures to see what the problem is.

MR. MILLER: I would be repeating myself if I tried to give much of an answer to that. The program is calculated and intended to minimize uncertainties to provide the kind of reasonable assurance that I talked about.

We feel on the basis of what we know right now, we feel it's likely that we will be able to do it. Frankly, we may learn differently a year or so from now, but from what we know right now it appears as though it is something that is practical.

QUESTION: To be more specific and get away from some of these variables, as I understand it absent some order affecting our license we can proceed on the assumption that we have an approved monitoring program?

MR. MILLER: John, maybe you can --

MR. LINEHAN: You indeed do have an approved monitoring program. Any facilities that don't before December 1st, we will be contacting you. This will be addressed in the audit if you do not have an approved one. If it isn't addressed, you've got one that is approved.

MR. SCARANO: If you don't mind, I would like to say a few words. I know that the people up front have

gone through this once. Let me try one more time, because

I hear words like, "What's going to happen when it's shown

ultimately that a mill may not meet that 25 millirem."

We feel confident with the control measures that are already required as a result of the ALARA and NEPA reviews. Those control measures are what we have the confidence in in controlling the mill to meet that standard. It is not the monitoring program, it is the control method.

Now, if indeed through the phase I that we wind up at the end where a few years down the path after sorting all this out there is a mill that has a contribution in excess to 25 millirem, what do you look for? You look for another control mechanism, an added control mechanism, not shutting down the mill.

You would be called upon to say okay, we have isolated the source. We know we can do this, and that will reduce the contribution to that 25 millirem.

The goal is to reduce the radiation hazard to the public, not shutting down the mill.

QUESTION: Am I to understand that what Mr.

Scarano has said that during the one year interim period while you are working with mill operators that no mill will be shut down?

Let's address ourselves specifically to the question of shut-down and what the effect of showing

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non-compliance through these variable measures will be?

MR. MILLER: The letter speaks for itself.

QUESTION: The letter says precisely that, "We do not expect during the period to enforce the standard through a shut-down or some other type of procedure."

I want to know specifically does that mean you will not, during that one year period, shut down any mill?

MR. MILLER: Our licensees are committed to the program of controlling blowing of tailings. I don't think in the past three years or so where we have had these requirements in licenses that we have shut down anybody.

There may be cases where we have in an inspection found it as a deficiency, in looking at that requirement, whatever procedure was developed by the operator to accomplish that.

My own opinion, and Ross can add to this, I

don't expect during this period of time that we are going

to have a basis necessarily to shut anybody down. It

doesn't mean that during this time we wouldn't, against

those license conditions which require an operator to be

carrying out these controls, cite him and list as a deficiency, in

an inspection and enforcement action, cite him on not meeting

those criterion.

Until we get to phase II it is going to be difficult to have basis upon which to shut somebody down

in 40 CFR 190 non-compliance.

MR. SCARANO: I would think we would say in working through our licensees and looking at the preliminary compliance, we would have no basis in the world for closing down any of our licensees.

There are certainly some mills that look like they may be on the borderline, and that is the process of sorting out, you know, where are these contributions coming from.

Again, let's look at what happens down the path.

We will be looking at the control measures, are they working,
and do we need more control measures to meet that standard.

That is the bottom line.

QUESTION: Are you going to be sending something to the agreement States on the same lines and suggesting similar procedures?

MR. MILLER: I am sure, consistent with our normal policy, we will be and have in the past worked on informing the agreement States how we are proceeding on this, and I would expect that we would do -- This has been somewhat of an evolutionary process.

We started a year or so ago to go through these assessments to develop a reasonable rational scheme that addresses the kind of concerns that you raise. We have had the same ones, and we try to build that in.

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I can't say that there is any specific meeting in mind, but I am sure that there will be some presentation with the agreement STates to help them with the same -- or to ensure that they are taking the same kind of course, same, kind of approach.

QUESTION: When do you think that will take place?

MR. MILLER: I can't say. We have kept them informed through sending them copies of materials and whatnot. We don't have a date or anything.

MR. FONNER: The agreement State program is handled by another office in the agency, and I don't think there is anybody here from that office.

MR. KENDIG: I'm John Kendig, Office of State
Programs.

We have been tracking what is going on with regard to 40 CFR 190, and the people on the site keep us informed and we, in turn, provide that to the agreement States.

Whether or not Hub is going to provide this

pack of information or not I'm not sure. I suppose they

will. That information, the two documents plus the letter

from our legal department, will be going down to the

agreement States.

Whether or not there will be any separate discussions, I don't know at this time. I would imagine

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there would be.

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QUESTION: I'd like to know what you are going to consider your value, or how you would define the background. Is it an absolute, or is it a value plus or minus?

Second, I would like some references so we can determine how you determine the dose conversion factors.

MR. MILLER: The ones I mentioned before, it's the regulatory guide.

A VOICE: Those dose conversion factors have been changed. The new ones are in the code package.

MR. MILLER: In the mill dose code package that we documented and distributed.

The other question was about the uncertainty on background. I expect that it will be subject to about the same errors in background measurement as you will from the monitoring samples around the mill, and it will be a straight subtraction.

QUESTION: Are you talking about a 96 percent confidence level?

MR. EADIE: The determination is no different from the point of the nearest receptor or the monitors near the tailing piles. You simply collect the sample and analyze it.

QUESTION: This compliance determination procedure says draft on it. Are you passing this out to us now

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because it's almost a non-draft?

MR. MILLER: The reason I put that there is because there are some minor differences between what is in this package that I just presented in viewgraphs and what is in that. It is largely more just emphasis in the way it is presented.

I expect to emphasis, although it's in there, more of the phased process. So, I didn't want that to be taken as the gospel. That will be a reference in the orders that we make, and it will be final then.

QUESTION: Does NRC have any estimates of the cost of the monitoring and reporting requirements that it is imposing through the regulatory guide and through its imposed amendment to impose on the uranium mills?

MR. MILLER: Let me say a general statement and then I will ask John or Greg to make some statements about specific parts of your question.

We don't feel that the level of monitoring that was established under the part 20 ALARA and NEPA requirements of the past three years ago. The increment above that for 40 CFR 190 is significant.

There is, during this initial phase, some studies that will be conducted, primarily short-term studies. We expect to be looking to streamline the environmental monitoring and reporting requirements as best we can over time.

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determination of compliance, this action level, is a kind of simplication that we are looking to provide.

Now, the costs --

MR. HICKEY: There have been cost estimates made of 4.14 and those were provided to everybody that commented on 4.14 which included some new licensees. It was also provided to the American Mining Congress.

There are not requirements in 4.14 that were added because of 40 CFR 190. All of those monitoring requirements stood on their merits.

The reporting estimates were done as far as the actual cost of preparing the report, not the cost of identifying the causes of failure to comply, not the cost of assembling the information you need to comply with 40 CFR 190, but the cost of filing the actual report with NRC.was deemed to be insignificant.

The reason for that was that we don't expect after this phase--You know, we are not going to be having this after that. We don't expect very many of these to be filed, and those that are filed are nto going to be very expensive.

QUESTION: I would like to mention another

potential element of uncertainty. NRC has justput out its

proposed general statement of policy procedure for enforcement

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noting raises and penalties and all that available.

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It also mentions that they are going to emphasis prompt and vigorous enforcement in dealing with persons who are unable or unwilling to comply with NRC requirements.

It is the Commission's intend that an enforcement program be marked by an aggressive enforcement strategy and seek more use of stronger enforcement measures with particular attention to detail, high standard of complience will be achieved. Furthermore, licensees who cannot achieve and maintain adequate levels of protection will not be permitted to operate.

That sounds somewhat inconsistent with phase I.

MR. MILLER: I think I have given the basis for the approach we are taking during phase I. As I said, we are confident. The basis for our confidence will be laid out in the report that we issue on December 1, that there is a defensable regulatory basis upon which to go through a phase I.

I think it is a policy of INE, but

INE has a problem too. In order to enforce that regulation
they have to show non-compliance.

We have the same problem you have. To show non-compliance we are going to have to have some kind of a formula to go through.

So, when your problems are solved so are curs.

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In the interim it looks like what is going to happen is that we are going to inspect solely on the license additions that are put on your license by NSS. Those will be judged on our basic inspection. It looks like they will all end up as deviations or something like that. It depends on how serious we find that you are not following that particular addition.

We are not going to be shutting down mills. We are going to try to get this regulation incorporated into the mill.

MR. TRUBATCH: With regard to the general statement of policy, that is a proposed general statement of policy regarding enforcement.

Some of you today have received your letter which has been read by the Commissioners which address the particular enforcement policies for phase I for this program.

MR. MILLER: It is specific to this problem today and I think it should be taken as superceding the general statement of policy.

QUESTION: This is a follow-up question to the earlier one that I asked about cost of monitoring. I have regulatory guide 4.14 in front of me, the provision dated April 1980, and I note for the record that it expressly calls for pre-operational sampling to detect compliance

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with 40 CFR 190, and also operational sampling to detect compliance or non-compliance with that regulation.

I ask my question again, has NRC made any estimates to determine the incremental cost of that kind of monitoring?

MR. MILLER: Clearly that program serves the purposes -- Those monitoring requirements serve the purpose of compliance determination with 40 CFR 190.

We have contacted the various people that we wanted to talk to. Is there anybody that wants to meet with us in addition to those who have been contacted? If so, you don't have to raise your hand now, but seek either John Linehan or myself out and we will arrange a time to meet with you.

I appreciate your coming. If there are any follow-up questions, don't hesitate to call this branch.

Thank you.

(Whereupon, at 11:30 a.m. the meeting was concluded.)

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CASE TITLE:

DOCKET NUMBER:

Meeting: 40-CFR-190

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HEARING DATE:

November 14, 1980

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LOCATION:

Silver Spring, Maryland

REPORTER'S CERTIFICATE

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I hereby certify that the proceedings and evidence herein are contained fully and accurately on the tapes and notes reported by me at the hearing in the above case before Nuclear Regulatory Commission and that this is a true and correct transcript of the same.

Date: November 17, 1980

Official Reporter

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