UNITED STATES DEPARTMENT OF AGRICULTURE

FOOD SAFETY AND INSPECTION SERVICE

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NATIONAL ADVISORY COMMITTEE ON MEAT AND POULTRY INSPECTION

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May 23, 2006 8:30 a.m.

USDA South Building Cafeteria 1400 Independence Avenue, S.W. Washington, D.C.

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(8:30 a.m.)

Good morning. MR. TYNAN: It's a little past 8:30. So we're behind time already, and we haven't even started. So I apologize for that. want to welcome you to our Spring 2006 Meeting of the National Advisory Committee on Meat and Poultry Inspection. I know I appreciate all of you coming from the Committee, taking time out of your busy work schedules to participate in our meeting. important event for us, and we have an ambitious agenda for today's meeting tomorrow's as well.

So we have an ambitious agenda, and it will focus again on risk based inspection as we did in our November meeting, and since we're -- we have so much to do today, I want to be the first to introduce our first speaker, Dr. Richard Raymond, our Undersecretary for Food Safety, who will make a few welcoming remarks.

DR. RAYMOND: Thanks, Bob. Good morning.

I'm happy to welcome you all back to Washington, D.C.

and for those of you who live here, I just welcome you

to the South Building Auditorium.

Your work here today and tomorrow is going to be very critical to our efforts to move towards a more robust Risk-Based Inspection System as you know and to insure -- help insure that all federally inspected establishments have the resources that they need to bring HACCP plans into the 21st Century.

Last November, I challenged this Committee to identify the ideal working group to assist us as we approach the next steps needed to enhance our risk-based system. You answered that challenge. I had two requests in November. One, that the working group would have to help insure that the process we're beginning will be as transparent and as inclusive as possible. And, secondly, I wanted this committee to focus less on what has worked in the past. Instead, I asked you to focus on what would work in the future for employees, consumers and industry through this important process.

My monitor, when it comes to consumer groups, industry reps and our employees, is that everyone must move forward together.

I've tried to beat this point home in talks using a three-legged stool. Well, I use the metaphor, a three-legged stool. I haven't really tried to bet it with the stool itself but all three legs must help support this platform as we move forward.

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Your final recommendations have exceeded my expectations. The time, thought and effort that you put into that final report at the last meeting and in the interim, for those of you who continue to work with us, has been very easy to recognize the quality. I appreciate that dedication, and I believe actions have demonstrated just how seriously we've recommendations. Knowing taken your what this committee is capable of, I'm here again asking for your input on two critically important areas.

The first, we are asking for guidance on how to most meaningfully measure, how well meat and poultry processing establishments control the inherent risks in their own operations. Everyone here knows that I want to focus our time and valuable resources on prevention rather than on response and recall. Command and control is the old agency mantra. What we

are now after is the common sense of public strategy to best serve the American consumer by prevent human illnesses. That is why it is so important that we begin to lay the foundation for a more robust Risk-Based Inspection System for the 21st Century.

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The key to this effort is an objective set of results orientated criteria that will allow us to better measure how well potential risks are being controlled in FSIS inspected establishments. This is not about increasing or decreasing the resources that we dedicate to inspections. It's about how best to use currently available inspection currently -- to related data and resources to improve the effectiveness of our hardworking inspection program This will allow them to better protect personnel. consumers.

Later today, you're going to be hearing our preliminary ideas of what we want to measure as well as how we might measure it. We're very interested in hearing your own thoughts and ideas on this important subject. We look forward to a lively discussion. It is not finalized by any stretch of the imagination.

In addition to continue to host open forums with our employees on these topics, we also are going to follow up this Committee's meeting next week, on Tuesday, with a joint meeting of consumers and industry to hear their thoughts on this important topic and our proposal. We've been meeting monthly with industry and consumers since Barb and I took our This will be the first time we've roles last summer. met jointly with them on this topic.

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And then, after that, a neutral third party, as you recommended, will be the one tasked with facilitating and soliciting input from all stakeholders for a finished -- for a recommendation for a finished product. You'll hear more about the third party when Barbara gives her your comments.

I firmly believe and still believe and will always believe, that an open and honest approach which takes the time to address everyone's concerns is in the best interest of all of our food safety partners.

The second thing we're going to ask you to do these two days is provide us input on FSIS' efforts to insure that every establishment that it affects or

is under State inspection is using a HASA plan that's designed to meet the demands and challenges of the 21st Century.

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Everyone here knows this marks the 100th year anniversary of the Federal Meat Inspection Act. That Act was a watershed event in the history of food safety and public health in the United States. that Act was based on visual examination for visible The future demands that we be able signs of diseases. to focus on things which the human eyes cannot see, the nose cannot smell or the fingers cannot feel. plan of every federally That HACCP establishment must take this and other realities we face in the 21st Century into account.

We do realize the small and very small plants have unique needs when it comes to HACCP plans and they do not have the resources that the large plants do. That's why I've made it an absolute priority for us to increase the communications between FSIS and the small and very small plants so that we can identify and respond to those typical needs faster and more efficiently.

We began last summer by holding listening sessions. We held three sessions in Montana, California and Pennsylvania and have followed up with two more since then. These sessions did help us better understand what was causing the gaps between plant performance and our expectations for them to operate under HACCP.

In December, Barbara and I attended a two-day brainstorming session hosted by the International HACCP Alliance on the key needs of small and very small plants. Some of you were there, and you know that at that meeting, six key needs were identified and an internal FSIS Task Force immediately formed to pinpoint how the Agency could better meet these needs.

We're here today asking for your comments and advice concerning the report that the Task Force produced. It's called the FSIS Strategic Implementation Plan for Strengthening Small and Very Small Plant Outreach, and it represents a dramatic shift in the approach of our outreach efforts and what has been done in the past.

Our new approach will be focused on insuring

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that small and very small plants can easily access accurate and consistent answers to their questions whether it has to do with HACCP compliance, food defense or food safety.

More importantly, the plan details concrete actions for FSIS to take in order to implement this new mindset. We will also be providing more educational resources and workshop opportunities for the small and very small plants on a variety of important subjects, concentrating on helping the plant employee who wants to participate but has too many demands on his or her time to be able to do that.

I know that it's often just not practical for owners of small and very small plants to take the time, to take a day off to go attend a workshop. Sometimes one person taking a day off for a workshop means one-third of the workforce is gone. That's why we're dedicated to providing small and very small plants with numerous options designed to make it easier to receive this training than we have in the past. These will include web tests, more night and weekend workshops, as well as making sure that all of

our workshops are available on DVD and CD-ROM for those who want to access them so they can actually attend these courses at the time that they choose and in the environment that they so choose.

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We'll also be working with Dr. Gayle Buchanan, the just recently confirmed Undersecretary for Research, Education and Economics, on ways to reenergize Extension programs and expand the reach of In fact, I'll have to leave our educational courses. this meeting for a little bit this morning to talk with Dr. Buchanan about that very issue. He's very interested in working with us and see Extension become part of our course again.

This is important to use them because I believe that education facilitates a greater understanding and helps close the performance gap that we're seeing in come of the implementation of the HACCP plans in small and very small plants.

This hopefully will achieve the goal of HACCP compliance without taking a lot of regulatory and enforcement actions. I'm much happier with the solution that calls for increased education rather

than increased regulations. However, we will do whatever it takes to make sure that a robust HACCP system is implemented and is maintained in each and every plant that is producing meat or poultry products for human consumption no matter how large or how small that plant may be. We must have safe products no matter what the size of the plants.

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When we eat a hamburger in this cafeteria, you don't know if it came from a small or very small or very large plant. When you go to a barbecue that someone's holding, you don't know where the meat or the poultry product came from, what size of plant. We must make sure they all have equally robust HACCP plans and equally safe.

realize also that implementation of We robust HACCP plans is not always just a matter of education. Sometimes it's а simple matter of That's why we'll also be working with economics. Undersecretary Thomas Dorr for rural development to distribute important information to small and very small plants concerning the loans and grants that are This is actually a little bit about rural available.

economic maintenance or rural economic development. A lot of the NRs what we write address shortcomings in a plant's physical environment and sometimes the money is just not there to correct that physical environment problem. But with low interest loans and with grants that we can make available, those plants can come into the 21st Century.

These are just a few examples of the new approaches that we're going to be taking for small and very small plant outreach and the concrete steps outlined in the plan. You're going to hear a lot about this from Karlease and Bobby later today, right after lunch I believe.

But the biggest change is unwritten. It's not in the plan. It's not in the six steps. And that's a change of mindset at FSIS. This may be the most difficult task that we have. Dr. Mann, who is my Deputy Undersecretary for Food Safety is using an analogy once in a while about when you go by a police car, you often see the logo, we're here to protect and serve. Well, FSIS is going to change our thoughts, and we're going to be -- we are here to regulate and

we're also here to educate, and if some of our employees can't make that mindset change, we'll have to help them along the way, but we will create mindset change of being helpers rather than just regulators.

As you learn more about this plan, don't think of it as a culmination of our efforts to improve outreach to small and very small plants. Please think of it just as the beginning. This issue will remain an important priority for the Office of Food Safety as long as Dr. Mann and I are here.

Before I finish, I do want to thank the Committee again for your valuable service. These are very important issues. Your wealth of experience and knowledge will be critical to insurance the successful outcome on these two projects. We know from past experience that we can improve food safety and protect public health by relying on sound plans and more effectively mitigate risks. That's why we're asking for your input on how we measure how well meat and poultry processing establishments control those risks in their operations.

We also know that we cannot move forward

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1 unless all of our food safety partners, 2 small and very small plants, consumers, employees, industry, all work in cooperation. 3 It's critical. 4 That's why we're dedicated to using resources to help 5 small and very small plants meet their unique needs. 6 We want to insure that we remain a strong and vibrant 7 partner as we move in through the next 100 years of 8 food safety. After all, the state of public health is We cannot afford the risk of 9 constantly evolving. 10 letting ourselves, our partners or our nation's food 11 safety system stagnate. Standing still in public 12 health is just a polite way of saying that we're 13 really moving backwards. 14 So Ι hope everyone has а successful, 15 I'll look forward to reviewing productive meeting. 16 your final report. I will be here as much as I can to 17 listen. I get more from listening than I do reading a 18 I look forward to the exchange report. So 19 dialogue. Thank you. 20 (Applause.)

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introduce our next speaker who has

Thank you, Dr. Raymond.

I'd

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MR.

like to

TYNAN:

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opening remarks, Dr. Barbara Masters, who is the Administrator of the Food Safety and Inspection Service. Dr. Masters.

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DR. MASTERS: Thank you, Robert, and thank you, Dr. Raymond.

On behalf of FSIS, I also want to welcome everyone to this important meeting. I always enjoy these meetings, Ι appreciate and everyone's Committee's work participation. This and their recommendations are certainly vital to our ongoing efforts at FSIS. In particular, I want to thank the committee for your hard work in recommending that we use a third party facilitator to assist us in getting input from all of our stakeholders.

I especially want to thank the Subcommittee for your input as we develop a statement of work to select our third party, and I'm please to report that we have selected a third party company, Resolve, Incorporated, to solicit input from our stakeholders. We have some of our representatives from Resolve, Incorporated in the audience today, and I'd like to recognize those individuals, Cathy Grant and Brad

Spangler. Cathy and Brad, can you stand up please. We certainly look forward to working with you and welcome you today to listen to the dialogue and tomorrow we also hope to be joined by Abby Dilly and Paul Demorgan. So we look forward to working with your company. So thank you for being here.

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We have a great agenda for this two-day meeting, and I want to point out that we're trying a slightly different format from meetings in the past. We're making some progress as we move forward on this road in our journey towards a more robust Risk-Based Inspection System, and again we're moving forward with this meeting.

As Dr. Raymond discussed, two of our presentations and the issues that you'll discuss, relate to a more robust Risk-Based Inspection System. The first, measuring establishment risk control for risk-based inspections and our strategic implementation plan from enhancing outreach to small and very small plants. Both of these issues are critical to the success of a more robust Risk-Based Inspection System, and are issues that your

Subcommittee will deliberate on later this afternoon.

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But in addition to these two issues, we have two additional presentations that we want to actually focus on related to our more robust Risk-Based Inspection System.

One of these is public health our communication infrastructure in a more robust riskasking based environment. actually are Wе the Committee to provide us input on this presentation but giving you until June 30th to provide we're We're hoping that you'll take into account comments. of the discussion and recommendations some emanate from the discussion today and tomorrow. Wе recognize that a reliable and integrated public health communications infrastructure is а foundation for protecting public health. We must get the right information to the right people at the right time to make the right decisions. Having the same data from the border, the field and laboratory personnel at the same time is essential so that everyone can connect the dots and proactively respond to that data rather than waiting to react after the problem has surfaced.

As we improve our public health communication infrastructure, we need your input. For example, we would like your input on the type of public health data and information that will help us make critical risk-based decisions. We would like to know what experience others have had in using business intelligence technology to better manage food safety or food defense risk. Based on the input that we receive from you, we anticipate we will bring this topic back as an issue for future meetings for further dialogue.

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The other presentation that we're actually going to spend the time to talk about from the podium is our FSIS Employee Focus Groups. The purpose of this presentation is to update you on how we're involving our employees in our discussion on our more robust Risk-Based Inspection System.

You may recall at our last meeting, that I described the risk-based inspections as a major structure built on a strong foundation with three pillars providing support. The three pillars taken together maintain the System's integrity. Those

three, industry, FSIS employees and our consumers. We wanted to make sure that as we solicit input from our stakeholders, that our employees were also well represented. After all, they're certainly critical to the success of our more robust Risk-Based Inspection System.

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We certainly heard your comments at the last meeting and wanted to insure that we had Agency employees involved in our discussions on a more robust Risk-Based Inspection System. It's probably a good time to mention that we actually have three employees today to listen to the discussion and also tomorrow, and I'd like to introduce them and have them First, we have Ms. Alfreda Dennis, who is stand up. National Joint Council representing our of Food Inspection Local. have Mr. Martin We Hickman, Association representing our of Technical and Supervisory Personnel, and have Dr. Patricia we Bennett, representing our National Association Federal Veterinarians, and I certainly appreciate their participation in the meeting, and they'll be sitting in during the discussions and the breakout

groups this afternoon.

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In closing, it is certainly important communicate openly and often with all of our food safety stakeholders to gain input and support. So again, I'd like the Committee to know how much we appreciate your commitment, not only at these meetings but again as we had conference calls from the time of last meeting up until this meeting. our appreciate your recommendations, and I want to thank you in advance for your hard work. These two days are always very action packed and very busy and very fulfilling. So thank you very much for all the work I know you're going to do, and I look forward to a productive meeting. Thank you very much.

(Applause.)

MR. TYNAN: Thank you, Dr. Masters. There's a couple of things -- I'm on the Agenda right now to talk about the charge to the committee, and normally at this point, I go through the rules of order that have fondly become known as Robert's Rules, but I'm not going to do that today. I think we've been through it enough that, that everybody's probably

memorized it. I would refer you though to Tab 3 and the Rules and Committee responsibilities, the Subcommittee information, how we're structuring the Subcommittees today is all included under that tab.

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Τ would like to mention couple а of logistical things. For those of you in the public that have come in and represent employee our organizations and Resolve, there should be registration book out on the table, outside of the door, if you could register there, I would appreciate it very much so that we can have an accurate record for the transcriber as well in terms of making your sure we get your names spelled correctly when we have the transcript and so on.

If you intend to make a short presentation during the public speaking portion which I think is at 2:00 this afternoon, I would be very grateful if you would put your name down there as well. It'll make it a little bit easier to recognize you and be aware of who's speaking and perhaps how long we need to dedicate to that portion.

What I would like to do though, at this

particular point, is briefly walk through the agenda so we can talk a little bit about how we're going to structure the meeting. As Dr. Masters mentioned, we are changing the format just a little bit and after I get through here, I'm going to turn it over to Janet Stevens and Marcelo Olascoaga.

MR. OLASCOAGA: Olascoaga.

MR. TYNAN: I'm sorry, Marcelo. I struggle with that a lot. A very talented guy but I can't pronounce his name.

(Laughter.)

MR. TYNAN: And they're going to talk a little bit about public health communications infrastructure as Dr. Masters mentioned.

At 9:45, you're going to see my bright and shiny face again and we're going to talk about the focus group that Dr. Masters mentioned as well, and I know this will be a disappointment to you, but at 10:15, we're going to have a break, and at 10:30, we're going to start talking about some of the briefing and issues papers that are included in your book. These are sort of updates from previous

meetings. We're not going to spend anytime making presentations as we will on the two previous topics. So it will just be a matter of if you have questions, we'll have people in the audience from the Agency that can hopefully respond to the questions you have, and if there is no one here that can do that, then we will take the question and get you an answer after the meeting.

So that will take us until probably around 11:00, at which time we'll introduce the first issue and that will be Measuring Establishment Risk Control for Risk-Based Inspection, and Mr. Don Anderson of our Program Evaluation and Improvement Staff in OPEED, I beg your pardon, OPEER, will be, will be here to make that presentation.

We'll have a break for lunch and at 1:15, we'll get back together to talk about the Strategic Implementation Plan for Enhancing Outreach to Small and Very Small Plants. And at 2:15, tentatively assuming that we're running exactly on time, we'll have the public comments and adjourn. And at that particular point, we'll ask the Subcommittees to take

the issue papers and the questions that have been posed to them, and the room numbers are there, and we'll get you to those rooms since this building can be a little complicated for those who are either new to it or don't traverse the halls very often.

So that will take us through the end of the day. So will the Subcommittees deliberate on the questions and the issues during the afternoon, and then as usual, on Wednesday morning, we'll start with a quick recap. Dr. Masters will talk a little bit about sort of reviewing what we did today, and then we'll have the Subcommittee report outs, and that will take us through we hope until about 11:15, we'll have another public comment period, and then we will be finished for the day. And I see a tent card is up already. Dr. Masters.

DR. MASTERS: Robert, since we do have some individuals, particularly our employers who are new to the audience, I'm going to force you to go through Robert's Rules of Order to make sure the public does understand their ability to at least listen in on the Subcommittees' deliberations and interact as much as

the Committee Chair allows them to listen in, et cetera, to make sure they know all the public is welcome to listen in on those Subcommittee deliberations.

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MR. TYNAN: And as I mentioned earlier, that I was going to hold the Robert's Rules until the very end before I spoke about them.

Robert's Rules under Number 3, the Rules of the Chair, which the FSIS Administrator Order, going to be the meeting conductor. The Chair is going the meeting and will be to open the person recognize those who want to speak. We're going to pose some limits in terms of time. Obviously we have a lot to cover. So we'll have to -- perhaps if you go on too long, we'll have to say time out, and we'll save you for the public presentation portion. A lot of the time Dr. Masters, and I assume she's going to do that again today, will delegate to Robert.

All questions, request to speak, will be addressed to the Chair. People must be recognized by the Chair before speaking. Normally, as we always do, we ask you in order to make it an orderly process,

that we take the tent cards, stand them up and then we can go around the room in an orderly fashion and make sure that everybody has an opportunity to comment.

Presentations on the issue papers will be followed by a short question and answer period so that you can clarify any point that the speakers made. In the interest of time, again we'll have to -- we may have to limit the length of time people speak at that particular portion.

Speeches or statements of opinion again should be made during the Subcommittee discussions or during the time set aside for public comments. Committee members and members of the public will be recognized by the Chair during the public comment periods of the meetings. So a request to speak may be presented to the Chair in advance. So that's why I ask you if you have some particular comments, separate and apart from the discussions we have today, if you have a little speech that you want to make or comments that you want to get on the record, please sign up outside, and we'll recognize you at 2:00 or 2:15.

Committee members are expected to attend the

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plenary sessions of the meetings and the Subcommittee meetings to which they are assigned. Committee members who do not attend the presentation of the issue will participate basically in a Subcommittee meeting, are going to be restricted in terms of their participation in the plenary session. That's simply in fairness to those folks on the Subcommittee who took the time to develop the recommendations. It would seem a little unfair to have somebody kind of come in the following morning and not really have deliberated or contributed to the development of the report.

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The Subcommittee Chair is designated by the Chair, and as I mentioned, that's in Tab 3 of who the Chair and the Subcommittee members are, and the Chair is allowed wide latitude to control the Subcommittee Members of the public may attend those session. Employee organization people and the folks sessions. from Resolve may attend as well, and it will be at the discretion of the chair how much access to the discussion they want to permit. And again, we will allow wide latitude in terms of doing that.

And then as always, the Rules of Order are

1 subject to review at each of our Committee meetings, 2 and if there are some changes that anyone would like to make, we can entertain that at -- whenever you'd 3 4 like. So are there any questions regarding the 5 6 Rules of Order? I see somebody in the back. Were 7 you -- okay. Just an animated individual. Are there 8 any other comments or questions? Let's get into the substance then of 9 Okay. 10 the discussion. It looks like we're making a change 11 in the -- as moderator of these meetings, you have to 12 be very flexible. 13 Does anyone have any questions or comments 14 at this particular point in our meeting? 15 We'll wait just a moment until we get Okay. 16 Thank you. Can everyone see that in our screen set. the back? I know it's a little bit lower but it seems 17 18 to be a little bit clearer in terms of seeing it on 19 the screen as opposed to on the white background that 20 we had. So I apologize. I think, Janet, you had PowerPoint handouts -- no. 21 Okay.

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I'm going to turn it over to

All right.

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1 Janet Stevens and Marcelo, Marcelo O, to talk a little 2 bit about the infrastructure. 3 MS. STEVENS: It's my dream to be on 4 American Idol. I just wanted to hold the microphone. 5 Well, good morning, everyone. Thank you for 6 allowing us to budge into your meeting a little bit 7 this morning. Again, we'll be talking about working together to protect public health, what that means to 8 the public health communications infrastructure. 9 10 First, I want to remind everyone of the FSIS vision, which is protecting public health for food 11 safety and food defense. Really what does this vision 12 13 mean for the Agency data and communications systems? First of all, it means giving control of the 14 15 data and the information, which means controlling the 16 Why it's important is because we need to access. 17 create a single source of the data and of the 18 information because if that single source is 19 traceable, then the data is not reliable. We also need to provide our stakeholders 20 with the data and information that they need 21 22 protect public health, and as Barb Masters put it so

well this morning, we need to get the right information to the right people at the right time to enable to them to make the right decisions to protect public health.

We also need to insure that the date and information are available when needed, because we may have the right information, but if we don't get it to our stakeholders in order for them to make that right decision at the right time, that window has passed.

We also need to better target our Agency resources to manage the data and the information to support this more robust risk-based system. The way we want to do this is to have a more efficient and effective management of strategic alliances with you folks. We also want to best target our resources to protect a more customer driven risk-based public health communications infrastructure.

I think this pretty much goes without saying but we need to make sure that the data is secure and is accessible only to the staff and to the stakeholders who should have access to it.

And this is mostly our focus for today which

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is where we want to be able to use leading business intelligence technology to allow for more predictive analysis because, otherwise, what we're doing is we're looking at the past and we can look at the past in real time, but if we're not using that data to be predictive so that we can be less reactive and more proactive, this will enable us to prevent and mitigate risk.

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One of the ways we're looking to do this, I'll get into this a little bit later, something called Assurance Net, and what this tool will do is allow us first of all to look at data in near real time in a more visual way. The way we're looking to do this right now is through more of a dashboard system, just like being in your car, you need all those monitors to know your speed, your oil, We need that instant feedback in a the temperature. more easy to see way instead of trying to look at all of the data and the numbers. We need it to assemble together into a dashboard system to make quicker and better decisions.

We also need to look at things that would

help us do some trend analysis. It's not enough to know how we're doing right now. We need to understand where we're trending. Are we not too good but getting better? Are we better but it's starting to turn? Where is that little, you know, turning point, that key point that will allow us to go in there and say it's right here, we need to focus our efforts and our resources on because that way it better targets our resources to make informed decisions.

We need to remember that business needs drive technology decisions. We need to always remember that so that we build the system and we provide the data in the way that helps support the mission.

And how do we get there? One is through web services. I'm sure most of you are familiar with our website. That was redesigned April of 2004. Based on our customer analysis, we are doing very well. We have been nationally recognized for the improvements that have been made. We are currently tops in the Department, and we are actually better than CNN.com and ABC News. I know. We were surprised as well.

We also launched Ask Karen and Ask Karen is a consumer-based tool that allows people to ask questions in a natural way like how long can I leave my leftovers on the counter before you can't eat them in the morning? So it's more of a consumer driven tool.

We also put out an e-mail subscription service that has been wildly popular. It's about 23,000 subscribers in about a year and a half and that helps people to only get the information they need when they need it instead of thinking I need to go check the website, how does this come up, when does the press release come out, things of that nature.

We also do this through strategic planning, which is where the management controls and performance measures come into play, and that's the tool you just saw which was Assurance Net. What that tool will do when we launch in June of 2006 is take the first set of management controls and there are a little over 50 performance measures of inspection based data and these are combined into what you saw of the dashboard to allow us to monitor what's going on in your real

time and able managers to make these decisions and target those resources for the Risk-Based Inspection Program that we see.

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I'll pass it over to Marcelo who will discuss the enterprise architecture vision.

MR. OLASCOAGA: Thank you. It's a pleasure to be here this morning.

going to give two-minute you а definition on what enterprise architecture is. an industry tool of methodology that's been used widely throughout the industry and now the Government taken it and really developed on comprehensive framework. What it allows us to do is document all of our business prophecies and then understanding that, we can reengineer our prophecies if we need to, streamline them when we need to, and also develop a vision, and that's something that we have embarked on and so far have had a lot of success. In fact, as a new person or a new employee less than a year here at FSIS, I think that the steps that we have taken were just incredible with the support of the Administrator, Deputy Administrator and the Chief

Operating Officer, and we have been able to already come up with a vision which is presented right here. It's very high level, but what it does show is that everyone is included. Everyone in the Agency, which is the middle box that you're looking at, is part of this vision, and it's going to be a very integrated system. We are going to get rid of a lot of our stovepipes, and we're going to be able to respond to any issues that come out in the industry or whether it's an economic problem that we may be facing or by closing down plants or some kind of problem that may be widespread.

Another thing is that we do show on the top layer, how everyone can interact with us because it's not just us working with the plants. We do have customers. We do have the public that we have to address, and we do have other agencies within the Government that we have to share data with. So by working together, this enterprise architecture is going to allow us to very transparent and very open with our data so that we can work together with any entity, throughout really the world, and that's what

we're aiming to do.

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Now this very complicated spot is very easy but what it really means for all of us is that we've already created at governance process, basic -- this slide, and the one after it, are sort of a trimmed down version of what you just finished seeing but it's a capital investment process. It's what every business does and it's what we need to do in the Government as well, and we have implemented this now.

So any idea that may come up, a risk-based going system, is something that you're be addressing later maybe in your Subcommittee, well, this idea has to be taken through this process to see how we're going to develop it, to really define our requirements and to insure that it aligns with all of our strategic goals within our strategic plan. So everything that occurs, even changes to the system that we do create, have to align to certain goals and we will measure every step that we take to insure that we're meeting everything that has to be met.

Now one of the things that we are targeting, which is the low hanging fruit that we will in the

next 12 months really capture is eliminating all the stovepipes that exist as far as reporting systems. So we will create what is called the data warehouse. We will create a vast recovery site in case -- so we can have fail over, and what this means is that we're going to have data in one repository that would allow to basically run complicated algorithms to us understand what the trends are, what will happen, to measure all of the performance measures that were established in the strategic plan.

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То really comprehensive be more and integrated, is really what you're hearing all the time in the Government. get rid of We need to the stovepipes. Well, these are the first steps that we're going to take to do that. And, although it's very high level and almost simplified in this slide, it's a very complex process but something that we did through the governance process and being supported by everyone within the Agency because all of these stovepipes are created within each of the individual programs throughout FSIS and now we will be able to eliminate that uncertainty by integrating all of this

data into one repository.

And, I think you're going to take over from here. There you go.

MS. STEVENS: Thank you. Again, we just wanted to remind everyone of our focus which is protecting public health, which is getting the right information to the right people at the right time to make informed decisions.

And again, this doesn't only support the mission. This really equals trust, and I think that's a lot of what this is all about, is just that trust and having a single source of data, having things be reliable, being more proactive and less reactive will get us there.

And now the fun part, which is how you can help. And basically we understand that you folks have discussed data needs before at previous meetings, and we wanted to kind of refine this a bit to make it a little bit more targeted response. So the first set of questions we have really revolve around the business intelligence technology such as what type of public health data and information will help us make

critical risk-based decisions? And, for example, how can the business intelligence technology, which are those applications that I was talking about that hit that data warehouse, pull from multiple databases, pull together in those understandable dashboards, how can they help to provide more predictive analysis? Ιf you're currently aware of technology that you'd like to bring up and discuss, we welcome your experience or knowledge on that. And, if you're currently using it, how are you using that tool to better manage or food safety and food defense communicate risk internally? We do have a deadline for feedback of June

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We do have a deadline for feedback of June 30th, and Robert, as he so wonderfully calls himself, will be issuing an e-mail address where you can provide us that feedback.

Right now, we'd like to ask if you have any clarifying questions for us. It's a lot to inhale at 9:15 in the morning. Yes, sir.

MR. ELFERING: Yes, Kevin Elfering from Minnesota. What actually -- what kind of data do you want to collect? Do you want to collect

microbiological data, inspection data, and what all -- what are you all trying to accumulate?

MS. STEVENS: Marcelo.

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MR. OLASCOAGA: Thank you. I can start out by saying that everything we need and really go to the questions we're posing now, we need to find out or get advice on what data we need to be more responsive to the industry. So those -- that's the reason for the questioning. The other parts we can handle on our As you can see, we are going to be classing a own. lot of the systems and creating this repository but by receiving advice from the Committee, then we'll be able to understand what additional algorithms we need to develop to answer whatever it is that's taking place in the industry.

MR. ELFERING: Perhaps one thing in addition you could provide to us, what data you are gathering right now. Are you getting data from PulseNet for example? Anything that you're collecting will be nice to know what you're already collecting so that if we can think of any ideas of additional data.

MR. OLASCOAGA: Sure, we can do that.

1 MS. ESKIN: Sandra Eskin -- what are you 2 collecting now? Well, Ι think 3 MR. OLASCOAGA: there's 4 somebody better to answer that question, but we are 5 collecting all the information on the samples that we 6 take from all of the plants. We are collecting in on 7 performance measures, what our inspectors are doing, 8 logging in all the time that it takes for them to 9 perform an inspection. We are logging in information 10 that may be related to law enforcement if we need to 11 So, for example, the NRs that were take action. 12 mentioned earlier. all of that is being electronically as well, so it's not just paper based. 13 And just to follow up on that 14 MS. STEVENS: 15 Assurance Net which launches next week which has those 16 50 performance measures, the information that it's currently analyzing is going to be information on the 17 18 HACCP, antemortem, postmortem inspection, food safety 19 assessment, things of that nature pulled together. Dr. Harris. 20 MR. TYNAN: I just had a very simple 21 DR. HARRIS: 22 question. I was going to ask if we could get a copy

1	of the slides as a handout?
2	MS. STEVENS: Yes, we'll provide that.
3	MR. TYNAN: We'll do that.
4	MR. OLASCOAGA: And the questions are in the
5	back of second page of your paper in your binders.
6	MR. TYNAN: I think it's Tab Number 4. I'm
7	sorry. I should have called that to your attention
8	before.
9	Mr. Elfering?
10	MR. ELFERING: Maybe one more comment.
11	Maybe one of the things that I think you may want to
12	look at, as far as collecting data, is things like NRs
13	are almost more opinion in a lot of cases, and I don't
14	know if that's always the best data to be looking at
15	rather than something that's a little bit more
16	scientifically sound. That's probably going to be
17	able to give you better predictability or at least
18	reactionary.
19	MS. STEVENS: Thank you for the comment.
20	DR. MASTERS: This is Barb Masters, and then
21	we'll get to you Sandra. I just wanted to comment
22	that a lot of our discussion later today on the

measures to control risk will discuss a lot of types of the data the Agency is already bringing in, and that's why we thought it would be difficult to have a full discussion on this particular topic and likely will be issue we address later but an thought it would be useful, that's why we thought it would be useful for you to give us some ideas after we have that full robust discussion on the measures to control risk because I think you'll see that when we get into that issue, that most of that is data driven, and it's based on data the Agency is currently collecting. So I think that you'll see once you get into that topic, a lot of the data the Agency is currently collecting drives where we want to go on the measures to control risk and establishment. think a lot of that will help you understand why we thought this would be useful to take as a homework So I think that will answer a lot of your assignment. questions.

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MS. ESKIN: Sandra Eskin. Following up on the question before about what data you're currently collecting, again what you identified is data that the

Agency collects, you know, again that we grappled with before, we've discussed in previous meetings and we'll discuss today is the stuff that industry collects. Let me back up. It's not only Agency but if you get public health data from various Public Health Departments, PulseNet whatever. How -- have you thought about what types of industry data you need to include in this system, how to get it, how to present it?

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MS. STEVENS: Yes, we have. We had a pretty robust I think discussion for several weeks, and we also invite you folks to let us know of the data you find of the highest priority to include first because there a large amount of data out there but we would also ask you to focus on the priority information that is needed to protect public health first because there is a lot of data out there in a lot of different formats. So it would all need to come together in something like a data warehouse.

MR. OLASCOAGA: Does that answer your question?

MS. ESKIN: Yes and no. Again, the

1	discussions we've had in the past and that we'll
2	continue to have is again the tension between needing
3	data to see to identify problems in the priority
4	nature of this data, individual enforcement and that's
5	something that's probably the stickiest part of this
6	whole discussion. So I'm not sure how to proceed.
7	Again, we've discussed it before but I don't know what
8	type of thinking you all have as far as how to get
9	access to that data?
10	MS. STEVENS: I think what we're interested
11	in first is trying to collect the information from you
12	folks so we can get that documented down so that we
13	can right now there's a lot of discussion and we
14	want to get this into a format that we can actually
15	start
16	MS. ESKIN: Meaning here 10, 12, 15, 20,
17	whatever types of data and things we need, and here's
18	where it comes from. Is that what you're suggesting?
19	MS. STEVENS: Correct.
20	MS. ESKIN: Okay.
21	MR. TYNAN: Mr. Govro.
22	MR. GOVRO: Thank you. Mike Govro from

1 One of the things you might consider as you 2 collect data is linking this to your training and education efforts in house, and one of the things 3 4 we've done in Oregon is we gather the data on all the their 5 violations that inspectors our note on 6 inspection reports and then run them in a spreadsheet 7 to see where individuals or areas of the State are 8 different from others, and it allows us to address problems of consistency and internal training so that 9 10 the program is administered more evenly throughout our 11 State. 12 STEVENS: I think that's an excellent MS. 13 point to have that direct nexus to the training. Again, there is homework. 14 MR. TYNAN: There 15 are questions associated with this topic that we'd 16 like to have some response from you all by the 30th of 17 June? 18 MS. STEVENS: Correct. 19 MR. TYNAN: Is that correct? For the 30th 20 of June. So we'll be setting up a location for you to 21 submit those comments, and we'll be moving along. 22 I'll try to remember to send you a reminder so that it

doesn't get lost. I know all of you have other duties besides being on a committee. So we'll try and make sure we help you with that to remind you that the questions and the responses are due.

If we're all covered with that topic, we'll go onto the next one.

MS. STEVENS: Well, thank you all very much, and we look forward to your comments.

(Applause.)

MR. TYNAN: I'm back again, and I want to talk about a topic that is a very important one as Dr. Masters pointed out in her opening remarks. We're working very hard to get our employee stakeholders involved in the process and get comments and information from all of them.

So I want to talk briefly about a series of focus groups that we conducted with our employees. It's a beginning point in the effort to insure that our employees, as well as our external partners, had an opportunity to be aware of and comment on this RBI process.

Because this is the first in a series of

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sessions where employees, it's sort of a broad look at the concept, and basically it's an overview. We enjoyed doing the sessions. We had an interested and engaged group of individuals that participated with us. In fact, actually Alfreda Dennis, who is here representing the National Joint Council, participated in the Beltsville session. So I'm going to put her on the spot, and at some point during the session, if I miss something important, Alfreda, I'm going to ask you to remind me or raise your hand. She may have some additional thoughts on some of these topics.

We did get a number of very good comments and thoughts and I wanted to go quickly through to highlight some of the things that we came up with.

Now the focus group format, we had four sessions. One in Atlanta, one in Beltsville, and we did two by a net meeting process where we projected the slides and the audio out to our employees at different locations around the country, and I have to say the net meeting worked very, very well. We had about 10 people in each of the sessions, and that's usually about optimal for a focus group kind of a

session, and we devoted about 2 1/2 to maybe as much as 3 hours for each of the sessions.

So we started with a PowerPoint on risk-based inspection, and it was a version of the presentation that Mr. Derfler did for you during the November meeting, and I believe that there should be a copy of that PowerPoint presentation with the audio in your notebooks, and you should also have a copy of my slides in there as well so that you can -- if you wanted to take some notes as we go along.

The CD provided some basic information for our employees. Obviously, this is the first opportunity they've had to hear anything about it. So we presented the CD sort of as background information for about 25 minutes, and then we had a question and answer period that ran approximately 2 hours.

We addressed six issues, and the issues are there on the slide. I won't take you through each. We have incorporated also a wrap up question where we asked the employees to give us the most important thought that they heard during the session or maybe that we did not talk about that they thought was

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There were two issues that came up on the CD when we were preparing it which had to do with food security and retail issues, and we did not do those in any of the focus group sessions because -- due to time constraints.

On this topic, there was the first issue that we dealt with which was anticipating problems, and that's the question we asked and there's a couple of points there that were brought up. We introduced this topic by telling the people that we believed that risk-based inspection is going to allow us to identify problems that are likely to occur in the process before they actually become serious. So as part of risk-based inspection we will capture and analyze data across plants. We also look to see if there are any patterns, breakdowns, findings or even hints of major And after that introduction, we asked this problems. question about what information should we be capturing so we can easily identify emerging problems.

I think the four groups were pretty varied in their responses. There was a general consensus

though that FSIS needs to capture more plant records and data. I think sanitation information, both from FSIS and plant records, was stressed by all of the groups, and it was stressed as being underused for identifying problems early.

There was also a discussion on getting more facility information, that that was under reported in the view of some of the participants. And they talked in particular that the PBIS has a task for facilities and equipment. That's OCB01 for those of you who are knowledge in those things. And that was mentioned as pretty broad. So I think there was some interest in having that become more specific or that additional instructions be provided on that particular task.

There was discussion about obtaining in house lab testing from plants. That was stressed in the focus groups as well. So I think in addition to the testing that's done for FSIS, there's also lab testing that's done independent of that, and that recordkeeping would also be helpful to us as well.

I think the last, probably the important point, there was a lot of discussion about in transit

shipping and temperature records. Temperature reviews on transportation from the plant to the location in retail where there might be some loss of temperature causing a problem and not being aware of it.

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We asked a second question in regard to anticipating problems, and the question there is, what risk do the establishments, in the establishments do you believe are not adequately addressed when it comes to food safety. The second question, I think foodborne illness and consumer protection, sort of umbrella term that some of the folks used were the top two answers of all of the four groups. Food-borne illness meaning -- included sanitation issues, such as employee hygiene, product distribution issues, pathogens such as Campylobacter and salmonella, well as farm-to-table issues in general. There was a number of comments that talked about trying to improve across the board, not only in the plants but making this more comprehensive as sort of a farm-to-table approach.

One of the participants indicated that they felt that salmonella was not adequately addressed, and

one of the examples given was an emerging problem was found in ground beef causing food-borne illness, and that it's difficult to regulate since salmonella is not considered an adulteration in raw products.

Another person described the problem of salmonella and Campylobacter are being primarily addressed in their poultry plants by regulating the presence of visible fecal material.

And again, distribution issues came up as important to a number of the participants.

We also talked about the risk-based factor that will be inherent in our Risk-Based Inspection System and if you remember Mr. Derfler's presentation, there were a number of factors that he proposed. I won't go through each of those, but they are listed here on the slide.

We obviously asked the question what do you think of those factors and I think in general, all four groups were fairly comfortable with those factors? Do you think that's correct, Alfreda? Do you remember?

Yeah, I think everybody pretty much agreed

with the factors as they were, and so we went onto the second question which had to do with looking at other factors that we should consider. So the participants expressed concern over how the new roles. So we got into a discussion of our inspection roles in this particular topic, and it was concern that we needed to have clear definition of the roles, the methods of assignments and who makes decisions and that in the employees' minds needed to be in place essentially before risk-based inspection can really get up and be successful.

It was also suggested that we need a lot of flexibility and speed in this new system, and that there may be much more -- it may, in fact, require additional staffing as opposed to less.

So it was a common concern also that the inspectors were talking about their own resource, but they were also talking a little bit about the plant employees, and I think they thought that a factor that needed to be considered in the new system had to do with the importance of insuring that plant employees are given adequate background screens. So this is

sort of a food security issue. And it was pointed out that it is critical for protecting the food supply, that plants must insure that employees are honest and, and have high ethical standards. That came across pretty much I think in all of the focus groups, and the employees also must be trained properly in food safety procedures. So we had sort of an issue of our inspection force as well as the plant employees as well.

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We also talked a little bit on the third issue which had to do with the work to be done, and we introduced this topic in a risk-based system, foresee flexible allocation of much more an inspector's time. So we try to free up our inspectors to spend as much time as they need to fully explore inspectional findings that relate to food their safety.

We are proposing a decision tree to help inspectors in their work. And so we asked the question, are there additional ways to guide inspectors besides a decision tree? And I think some of the participants stated that they didn't feel that

there was anything wrong in the current system, that they were fairly comfortable with the system, that it works well, and in short, we like the system we're in.

Many of the participants expressed need for inspection personnel in plants to have more freedom to use their discretion, and they agreed that more flexibility in training to empower the inspectors was needed. I think again the topic of lack of resources or perhaps the need for additional resources came up concerning how we can adequately cover assignments when you have three or four facilities that people are working in.

And data again was a recurring issue and it came up in this, and so this ties a little bit with what Janet was talking about. I think the employees have concerns about the data sources and data needs, and I think the folks in the net meeting particularly, felt that there was a need for the best data systems, and they talked a little bit about stovepiping here and another question as well.

So some of the participants didn't feel that our systems communicated across the programs, that

there needed to be more -- a more integrated data system, more communication and cohesiveness, and also a participant noted that the database system's success in making good decisions is contingent, as we all know, on good data. Do you agree with that, Janet?

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We also talked on issue number 4, the design of the inspection system, and here we introduced the subject by asking participants to focus on the aspects of the plant's process, where -- control is likely to occur. And we talked here about using a recently developed hazards control guide to help inspectors identify these critical aspects of an establishment's operation.

And Ι should have pointed out, the composition of the group was varied. We had in plant inspection personnel as well as some headquarters personnel and various programs, and I think not many, and I have to say when we asked this question, not many of the participants really had prior knowledge of this hazards control quide in the first two sessions. We described it a little bit better in meetings, and I think there was a general agreement

that this hazards control guide, which I believe is part of our 5000.1 or 2, Bobby?

MR. PALESANO: 5100.2.

MR. TYNAN: 5100.2, I'm sorry, was part of that, and I think once we talked a little bit more and explained that particular hazards control guide, it was a little bit more apparent and employees felt it was a little bit more worthwhile and would be helpful to them. That, of course, assumes that they would have a little bit more training and support in using that guide.

We also talked as issue number 5, on how risk-based systems should respond to findings, and here we introduced the topic a little bit on -- in terms of how we traditionally do things, evidence of compliance or noncompliance doesn't necessarily affect the intensity of inspection. So noncompliance is always potentially led to enforcement activities, a NOIE or something along those lines, and that would certainly continue under RBIS, but under the system, noncompliance would also lead to a greater level of inspection. So we posed this question about how

should risk-based systems respond to the finding, and basically is the approach satisfactory, and I think generally the answer varied a little bit. I think the participants thought the approach was satisfactory, and that RBI would certainly improve inspection overall from their perspective.

One of the participants agreed that the approach was satisfactory but they wanted to know a little bit more about what constituted good control versus bad control. So they wanted some clear definitions of how you make decisions on what more intensified inspection was required.

One of the participants thought that laws and regulations needed to be revised, and I think that was the question that was brought up when Mr. Derfler was making his presentation back in November. So they have some of the same issues and concerns that you all did.

Data came up again. So many of the participants were concerned that RBI was dependent on the quality of the data that's available. So we're back to garbage in, garbage out. As one person

stated, a data driven system depends on accurate data with checks and balances.

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Now this varies just a little bit from the technical aspects of RBI, but it is a very important question from -- I know from my perspective and I think Dr. Masters, certainly from her perspective. We have done a great deal of work with our employees in trying to improve our internal communications to make it a more cohesive workforce, get the information that people need. So we added this question on how we could do continuing communications with our employees, and certainly we have a number of things that we do internally. We have news and notes. I think some of you are probably familiar with that. We have Internet access, intranet access for our employees. We have a Beacon newsletter. So a lot of the information gets out, and I think everybody agreed that those were good, adequate ways for employees to stay informed.

But a number of the employees asked for more listening sessions, not listening sessions, but perhaps focus groups that they participate in focus groups as we were doing on RBI, so that they could be

kept involved, that there be more town hall meetings perhaps, all hands meetings and, in fact, we're going to be planning an all hands meeting in early June for all of our employees that can access that. So we'll be working on that as well.

They pointed out that obviously on the kill floor, they don't have really access or, you know, time to walk off the kill floor to go to a computer. So they were looking for direct mailings as an option there.

And they pointed out, too, that we have some work to do internally in terms of our computer system, to make them more accessible to our field employees. We have a dial up system out in the field, and it has been a source when we're doing focus groups on this topic and others, that caused a problem and, in fact, when we did the net meeting, we had a couple of employees that tried to log on and we were 15 minutes waiting, and they were only at 34 percent downloaded of the material that we needed them to be looking at. Fortunately it worked out fine, because it had an audio portion that they could listen in on. So we had

to ask them to take notes. So that's an important area.

So in short, we were asking for ways to make communications about RBI. I think there was a general agreement, and I think, Alfreda, correct me if I'm wrong, but I think everybody seemed to want more information than less, and I think they all pointed out that while they may not agree with the process, that certainly their ability to help us get there, as well as buy into the process would certainly be more aided by more information than less.

I think the last question we had was the wrap up question, and we basically said to everyone on each of the focus groups, what idea or thought that they had heard today, during the session, or maybe that we did not talk about, that was particularly important to them that they wanted to bring up and sort of get on the record if you will, and actually four areas came out.

We had inspection resources and we talked a little bit about that. The response is focused on how resources might meet the demands of RBI.

1 We also talked about the communications. So 2 we went back to talk a little bit more about how they 3 would be heard and more listened to by upper 4 management. There was concern the employees be 5 informed on an ongoing basis for the communications 6 was the second one. 7 We also went back to data quality and 8 integrity. So I think what Janet and Marcelo talked about earlier is becoming increasingly important not 9 10 only probably from your perspective but from our 11 employees' perspective as well. 12 last, but not least, I think 13 employees wanted more empowerment of the in plant 14 inspection personnel which should be enhanced by RBI. 15 It gives them a little bit more flexibility in the 16 future. So that's basically the outcome of our focus 17 18 groups, and with that, I'll turn it over, and if you 19 have any questions. 20 Mr. Elfering. 21 MR. ELFERING: I've got a couple 22 On the first issue, anticipating problems, questions.

1 when you talk about they're not adequately addressing 2 issues when it comes to food safety, who adequately addressing issues? 3 4 MR. TYNAN: I think the focus of the 5 adequately addressing question was us in our 6 inspection system. The way our current system is 7 working, what is not adequately addressed? 8 MR. ELFERING: And then as a follow up, what is their solution to adequately following up on those? 9 10 If you don't have a solution, you become part of the 11 problem. Well, I think that was probably 12 MR. TYNAN: 13 getting a little bit more detailed than we anticipated I think what we were asking is 14 for the focus groups. 15 isn't addressed so that the folks what that are 16 developing the system and the concept will incorporate those things and solve some of those problems. 17 So our 18 intent in asking employees was just to see if there 19 are areas that we had overlooked, not to necessarily 20 get the solution at the same time. Does that help? 21 MR. ELFERING: Yeah, I guess maybe the same 22 thing you'd answer then with having more inspectors on

issue two. I don't necessarily think that more inspectors are going to increase food safety. Again, no solution on how -- where you would be employing those additional inspectors? Would it be on line inspection, processing inspection?

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MR. TYNAN: No, but I think that the issue came up a number of times that really is an area that the Agency needs to think through, particularly when you're talking about if we're intensifying inspection and we're asking an employee to go from say Plant A where they're doing well to going to Plant B and perhaps having somebody at Plant C going to another plant and passing on the way. So they were simply saying to us that we need to really consider the resources that are going to be invested in this, and some of them felt that it perhaps would require additional resources.

Dr. Masters, I can see you have a --

DR. MASTERS: This is just our first opportunity to introduce the information to our employees, and our qoal would be to take this information and provide it to our third party

facilitator and allow them to try to take the raw data and then take it back with new questions to our employees through whatever they determine the right process to try to then ask the follow up questions because it was the first two and a half hour type sessions that we got the initial responses, and we will use our third party facilitator because this was first attempt even prior to having a third party facilitator in place. So we would envision our third party facilitator to then say, ah-ha, this is an area that we need to go back to our FSIS employees and ask some follow up type questions, to say you believe there's a lot of responses that said you feel like you How would you envision using need more employees. those employees if that was a follow up question that felt was appropriate and whatever, if they they determined the right way to go back to our employees.

MR. TYNAN: Does that help, Kevin?

Mr. Finnegan?

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MR. FINNEGAN: Yes. On issue number 6, I would ask you to keep in mind how Agency employees keep informed and ask you keep in mind that the State

1 programs are 28 States. So there's a good access to 2 keep those people informed either through the State 3 Director. 4 MR. TYNAN: And I think there were questions 5 about getting data with the States, working more 6 closely with it. Some of the more -- we tried to 7 capture the more universal type comments but there 8 were other things like that. So that was a concern of 9 our employees as well. 10 MR. FINNEGAN: Sure. I just wanted to 11 comment on that. 12 All right. Thank you. MR. TYNAN: 13 And also back to a couple of MR. FINNEGAN: the other -- I think the first slide talked about 14 15 records and things. Are we talking about NRs there, 16 written on plans, part of the data? 17 MR. TYNAN: No, I don't think they were 18 talking about NRs because I think that's sort of 19 universally accepted. I think there was a sense that 20 needed additional plant data either the sanitation. think there 21 facilities, the I was 22 discussion about the additional lab testing that was

1 necessary and things of that nature. Alfreda, is that 2 sort of an accurate --MS. DENNIS: 3 Yes --4 MR. TYNAN: Okay. Thank you. Does that 5 help, Michael? 6 MR. FINNEGAN: It does. I just wanted to 7 make a comment. If part of the data is NRs, there are 8 different NRs. Some are -- take for example, a plant has a positive for E. coli O157:H7, that's a little 9 10 different than a NR written because somebody forgot to 11 put initials in, you know what I mean? We've got to 12 weigh those NRs, ones that have some meanings to them 13 and other ones that are clerical errors. 14 MR. TYNAN: I think that was brought up in a 15 couple of the sessions as well as I recall, and I 16 don't know if it was in the Beltsville session, but 17 there was a distinction between NRs, that they weren't 18 all equal. 19 MR. FINNEGAN: Right, right. 20 MR. TYNAN: So some of the same thoughts 21 you're suggesting were brought up by our 22 employees as well.

1	MR. FINNEGAN: Thank you.
2	MR. TYNAN: Dr. Carpenter.
3	DR. CARPENTER: David Carpenter. I want
4	Robert, I want to turn or tie issue one where you have
5	in house lab results to the wrap up where it says data
6	quality and integrity. Does in house lab results mean
7	data that are generated in a consistent type of
8	manner, like using AOAC method or a method that has
9	been sanctioned by Pat Makasky (ph.) and his group or
10	is it just right now pretty nebulous as to where those
11	data come from?
12	MR. TYNAN: It was pretty nebulous. I don't
13	think any of the employees really thought about data
14	in some sort of a certified kind of a format or with a
15	standard operating procedure. I think they were
16	simply saying that that's an area that we need to
17	explore as we develop the concept. Does that help,
18	David?
19	DR. CARPENTER: Yes.
20	MR. TYNAN: Ms. Eskin?
21	MS. ESKIN: Sandra Eskin. I think
22	Dr. Masters actually addressed the point I'm going to

make, but I think it bears repeating. I understand the role that focus groups play generally in any sort of policy development. I am concerned that because there wasn't a third party leading these focus groups that the employees were maybe not as forthcoming as they otherwise might have been. So I had that concern initially but when Dr. Masters explained that, in fact, was preliminary, that in fact our third party facilitator will go back in a systematic way. I think that's critical here because again focus groups are closer to anecdotal information than they are to any sort of statistically significance.

MR. TYNAN: You're exactly correct. What I didn't point out when we talked about the format though was to be sure that we got honest responses, we told everyone in the groups, and I'm sorry I've violated that with Alfreda, but we told everybody that their participation was anonymous. So we captured —for the face-to-face sessions, we captured everybody by number as opposed to by name.

MS. ESKIN: Sure.

MR. TYNAN: So that, so that they were

1	welcome to say what ever they wanted and when we were
2	developing a summary and that will be final hopefully
3	this week.
4	MS. ESKIN: Where were they done though?
5	Where were they conducted? In
6	MR. TYNAN: One was conducted in Atlanta.
7	MS. ESKIN: No, no, I mean
8	MR. TYNAN: In the District Office.
9	MS. ESKIN: Okay. Again, to get the larger
10	context there. I understand the anonymity but when
11	it's an FSIS employee who is doing the questioning and
12	it's in an FSIS facility, there always is the concern
13	that they're not going to be as forthcoming but again
14	since this is really a preliminary discussion, I jut
15	want to make sure that we all understand that and,
16	it's the first step before the third party
17	MR. TYNAN: No, no, I understand. Well, we
18	did try to make it as objective as we possibly could
19	but I recognize what you're saying.
20	MS. ESKIN: Inherently subjective
21	MR. TYNAN: That's why Resolve is back
22	there. They're going to help us with that.

Dr. Masters.

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And I appreciate Ms. Eskin's DR. MASTERS: I just want to reemphasize that there was a comment. time lag between the November NACMPI meeting and the time we were able to get Resolve involved, and we felt it critically important to get some information out to our employees. And so we felt like if we waited until we got Resolve involved, we would have had a second NACMPI meeting and would have had no information for employees. So that was why we felt it critical to provide the information from the first, and I think Robert addressed most of the information on the first NACMPI meeting, and so I want to reemphasize what we will handle all you're saying which is this information, the raw data over to Resolve and turn over our process to Resolve to get all further input from the employee groups.

So we should have stated that up front to say this is preliminary. It was all pre-third party facilitated, and so everything from here forward with our employees will be done by a third party. So I think it was just an important effort on our part to

get our employee groups involved, and it was our attempt to at least catch them up on the information that had been shared with this group so that they at least would say why is there all this information out in trade association journals --

MS. ESKIN: Right.

DR. MASTERS: -- and why are we being left out. You know, my third pillar is Dr. Raymond's third leg to his stool and, you know, those -- that structure is going to collapse, the stool is going to collapse and we just felt that strongly about getting them engaged, and so we should have said that up front and I appreciate your comments.

MS. ESKIN: Yes, and again, I have no problem with obviously involving employees. They are probably the ones most directly affected by what we're talking about. Maybe it's something as simple as not calling them focus groups. I don't know what else to call them because focus groups is a term of art, and whether it's listening sessions or discussions or whatever.

MR. TYNAN: Well, I can't say that I made it

1 an art form. That would not be true. But we did have 2 a good session, and I do think we do think we got a lot of very candid comments, and I think everybody 3 4 understood that there was not going to be any reprisal 5 for not liking something about the system. So I think 6 it was fairly objective and as Dr. Masters says, we're 7 going to leave Resolve to do the rest. Other questions regarding focus groups? 8 Oh, I'm sorry. Ms. Grondahl. 9 10 MS. GRONDAHL: Thank you. I'm just wondering if you have any plans for additional focus 11 12 sessions, and if so, if there's 13 consideration to include State inspection employees? At this point, the work with 14 DR. MASTERS: 15 all of our stakeholders will be done through Resolve, 16 and as -- I don't think you were at our last meeting, Andrea, and -- were you, and all of that work will be 17 18 done through Resolve and, yes, the State employees 19 were certainly on our list of those stakeholders that we think critical to be involved. 20 The approach that Resolve takes is yet to be 21

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We'll be meeting with them later this

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determined.

week, and I think that they are looking at a variety of ways to reach out to our stakeholders but certainly State employees will be one of those fruits that we're looking at, making sure we have them involved, and I appreciate Mike's comment to make sure that we make sure that any information we share we give to the State Director.

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And as Dr. Masters mentioned MR. TYNAN: earlier, we have two of the Resolve folks here, Ms. Grant and Mr. Spangler, and you're welcome at the break, which we're going to take in about two seconds, to chat with them and voice some concerns at this We will be trying to meet with them, I think point. the NACMPI Subcommittee to help us with the original statement of work. We're going to meet with Resolve probably at the close of business tomorrow, and then we're also going to have them meet with our Steering Committee later on in the day. So we're going to try and get them lined up as quickly as we can so that they can get off and running on this process.

With that, if there's no other questions regarding the focus groups, I'm going to suggest we

take a break, and if we could get back together, since we're now a little bit ahead of schedule, maybe about 20 after 10:00, and I will call you back to order when that occurs.

Okay. Thank you very much.

(Off the record.)

(On the record.)

MR. TYNAN: I'd like to get started with this portion of the meeting. If everybody could finish up their coffee and come on in if you're still out there snacking away.

In the Agenda for this morning, we're at the point where we're going to talk a little bit about the briefing and update materials, and unlike the two sort of presentations we did before, this is material that we send you. It should have been in your notebook and is here simply for just open discussion and questions if you have any regarding any of these particular topics. We should have folks from the program that should be here to answer questions. As I mentioned earlier, if for whatever reason they weren't able to make it at the appointed time, we'll try and get an

I'm on

answer to your questions after the fact.

I think the first one that -- briefing paper
that I have and would ask if you had any questions

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5 applying the mark of inspection to product tested for

regarding, has to do with the mark of inspection and

6 an adulterant, and I think I -- I'm sorry.

page -- on Tab 8. I apologize.

Mark, do you have a question?

MR. SCHAD: Yes. It might be a little bit too early just because the guidelines were issued in September 2005, but just looking at the bar graphs, it looks like more small and very small plants are holding product than they were in the past. And I was going to ask the question, is the Agency satisfied at this point with the progress that has been made?

MR. TYNAN: Let me see if we have somebody in audience who is going to address this issue? Come on up. This gentleman told us when he came up that he has laryngitis. So I may have to interpret for him.

MR. LEHMAN: I'm sorry. What was the question?

MR. SCHAD: I know the guidelines, the

1 industry guidelines were just sent out to the plants 2 in September 2005 but I was looking at the 3 graphs --MR. LEHMAN: 4 Yes. 5 SCHAD: -- and I know this wasn't a MR. 6 statistical analysis or anything, at least not what I 7 see in the report, but it looks like the small and 8 very small plants are holding more product than they 9 have been in the past. 10 MR. LEHMAN: Yes. 11 SCHAD: And I wanted to know if the MR. 12 Agency was satisfied with the progress so far? 13 MR. TYNAN: Before you do that, Excuse me. Mark, would you like to introduce yourself? 14 15 Oh, yeah. MR. MICHAEL: Matthew Michael. 16 Director of Evaluation I'm the the Program and 17 Improvement Staff. I believe T']] defer to 18 about whether Dr. Masters or not the Agency is 19 satisfied. I'll agree, there has been a trend in 20 every category where product has been held by every size plant, every year since 2003, with one exception 21

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which we think is a data error and not really a break

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in the trend, and as we said in our summary points, the vast majority of product is being held with between 80 percent and 100 percent of product being held in 2006, and really 100 percent of all the product held by large plants every since 2003.

MR. TYNAN: Mr. Schad, do you have --

DR. MASTERS: At this point, the Agency will continue to watch and insure those trends continue and insure that small and very small plants in particular, since the large establishments have consistently been holding product, to insure that trends continue and insure that products are held because that is certainly the goal of the Agency, to insure that products are being held.

MR. TYNAN: Mr. Schad, do you have a follow up question? I can see one forming?

MR. SCHAD: No, that's fine. I just thought of one other question I had. This data is from the sample form that's filled out when the sample is taken because I know there is a question on there, is the product being held?

MR. MICHAEL: Yes, yes.

1 MR. TYNAN: Dr. Harris? 2 As I recall, one of the key DR. HARRIS: concerns that the Agency had was, was the number of or 3 4 incidents of recalls attributable to product not being 5 held when a sample was pending. Is that data been 6 looked at? Is that improving as well or can you 7 update us at all on the trend or if there is a trend 8 there? I believe that trend has been 9 MR. MICHAEL: 10 improving. There's been very few recalls this year so 11 far for FSIS deposit product. I don't have the number 12 at hand. 13 Dr. Harris, MR. TYNAN: okay? you 14 Mr. Kowalcyk? 15 MR. KOWALCYK: Yeah. I have one question 16 about the date you're presenting here. Will the 17 Agency be reissuing this data once a year is completed 18 because right now we're looking only at the first 19 quarter, and it's compared years -- full years against 20 first quarter, and I quess to kind of really get a true measure, will the Agency be coming back to this 21

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Committee with results at year end as well?

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1	MR. MICHAEL: Yes, we're planning on
2	continuing looking at this data. We wanted to
3	establish a baseline or context to look at the overall
4	trend and then next year see if we could find an
5	impact correlate an impact of the guidelines on
6	holding, holding product and that's why we looked at
7	the first quarter of 2006. We didn't see any break in
8	the trend there, but it's very early but, yes, we will
9	continue looking at this data and applying it, yes.
10	MR. KOWALCYK: Thank you.
11	MR. TYNAN: Mr. Govro?
12	MR. GOVRO: Yes, I have a comment on the
13	update on training and outreach if we're finished with
14	this subject. If there's other comments, I can wait.
15	MR. TYNAN: Okay. Hang onto training and
16	outreach for just a moment. Any other questions on
17	this particular paper? Yes, Mr. Finnegan?
18	MR. FINNEGAN: Yeah. On graph 2, you have
19	percentage of ready-to-eat products sampling 0157:H7.
20	The Agency samples ready-to-eat products for E. coli
21	0157:H7 due to cross-contamination or I mean if

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you have a -- step, the E. coli is dead. I can see

1 listeria, possibly salmonella, but I --2 MASTERS: DR. We sample ready-to-eat fermented sausages for example for E. coli 0157:H7. 3 4 That was Barb Masters. 5 MR. Other questions this TYNAN: on 6 particular paper? Thank you, Matthew. 7 We're going to proceed through these 8 order if you do not object to that. If you have 9 another way you want to approach it, that's fine. 10 We have Legislative Update on the list and that should be under Tab 9, and I saw Ms. Picard there 11 12 before. Yeah, come on up. We have questions on the 13 Legislative Update. I brought Lisa all the way up here for nothing. 14 15 Mr. Elfering, did you have a question? 16 MR. ELFERING: I guess I'd be remiss if I didn't at least bring up the issue of the funding for 17 18 State inspection programs. As probably all of you 19 know, State inspection programs in the past have been 20 funded up to 50 percent of their inspection programs and we found out kind of midyear that that's not going 21

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to be happening in this Federal fiscal year, and if

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that is going to be looked at, as that will be the future, that 50 percent funding will not be available for State inspection programs or is that something that the Agency is working very diligently So that at least the State programs will rectifying. be able to predict how much funding they will have. For example, this year not finding out until half the year is over, they're going to be scrambling and probably have to make some significant cuts from their programs this year by not finding out until really half the year is over.

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The Agency as Mr. Elfering has DR. MASTERS: indicated held State level in funding as to what they provided since 2005, and so we recognize and appreciate the difficult -- difficulties that they're having in trying to look to where they're going to have to have fiscal restraints in their spending as FSIS is also going to bat this fiscal year. we are going to be working diligently as Mr. Elfering suggests to make sure -- make certain that in future years, we'll be working with the States as we tried to do this year, to try to make sure that we honor their

requests for additional funding. So we'll continue to work and hope that that is not the case in future years.

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MR. TYNAN: Mr. Elfering, do you have a follow up question?

MR. ELFERING: I do have a follow up. Dr. Masters, I've just been hearing rumors on budget issues that there was just an article that I read the other day that there is some discussion of furloughing federal inspectors. Could you maybe put our concerns to rest on that issue?

Absolutely. This is Barb DR. MASTERS: The Agency at this time has absolutely Masters again. no plans to furlough our employees. We also face a very tight budget year, and we are working very diligently through a lot of responsible budget actions to insure that public health is protected, but we are looking at hiring freezes and those kind of positions outside of the front line inspection force as well as restrictions on non-front line inspection activities but obviously if we were not able to meet our needs at the end of the year, we would have to

1	look at more drastic measures, but at this time, we do
2	not see that in the future. So at this time, the
3	Agency has no plans to furlough our employees.
4	MR. TYNAN: Dr. Carpenter, you had a
5	question? I know the spotlight is on you apparently.
6	DR. CARPENTER: Only for what? The next
7	half hour. On the second page, we talked about
8	laboratory. Obviously I have to bring up laboratory
9	issues. You talk about increasing funding for FERN
10	and eLEXNET and also a repository of analytical
11	methods. Is that going to be the responsibility of
12	the Athens Lab to create this repository or is that
13	going to be more diffuse than that?
14	MS. PICARD: Let me find out if I can get a
15	specific answer for you on where and who is going to
16	be handling that repository.
17	DR. CARPENTER: Okay.
18	MS. PICARD: I'll follow up with you later
19	this morning.
20	DR. CARPENTER: Thank you.
21	MR. TYNAN: Mr. Govro?
22	MR. GOVRO: Can you describe the nature of
	Free State Reporting, Inc.

1	the \$105 million in new user fees?
2	MR. TYNAN: I'm sorry. Say the first part
3	again, Mike?
4	MR. GOVRO: The \$105 million mentioned here
5	that are new user fees that are proposed. What, what
6	types of fees are those and who will they be assessed
7	to?
8	MS. PICARD: Those users fees are not
9	currently in the bill that is before the House this
10	morning. So they are not.
11	MR. GOVRO: So then let's get to the numbers
12	here. \$862.9 million is what's requested and \$105
13	million of that is proposed user fees. Do you
14	understand that?
15	DR. MASTERS: The 105 in the President's
16	Budget that went forward, Mike, the user fees would be
17	those above and beyond the first shift that's provided
18	by FSIS, that would be paid by the industry for
19	inspection beyond that covered by the first shift by
20	FSIS.
21	MR. GOVRO: Okay.
22	MR. TYNAN: Other questions on the
	Free State Reporting, Inc.

Legislative Update? There being one, thank you, Lisa.

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Now we're onto issue number 10 which Okay. has already been raised which has to do with the Update on Training and Outreach, and we have two individuals that can speak to that. Mr. Govro, you have question? Ms. Kelly or Dr. Kelly а and Ms. Cutshall, did you want to?

DR. KELLY: A question?

MR. GOVRO: Actually I just had a question. Mike Govro, Oregon. In Oregon, a professor at Oregon State University, Dr. Donna Beegle, did a study on communication differences between people different cultures, specifically oral and written cultures, and talked about how they perceive messages that are given to them, and the difficulties that people from oral cultures have in receiving messages that were designed and delivered by people from written cultures who think quite a bit differently, and that's particularly the case in a workforce with a lower level of education, and it's particularly important in the food service industry where there's a lot of turnover, and it's a fairly low paid work

force, and I just wanted to direct your attention to that, because I think it would be fairly significant in communicating with line employees in a processing facility. And I can get you the name of the study. I don't have it on me at this time, but I'd be happy to get that for you.

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MS. CUTSHALL: This is Mary Cutshall, and I think that's a very good point, one that needs some looking at and some of the things that we've worked on this year has been trying to reach out to employees. As you know in past years, with small and very small plants in particular, one of the things that we've tried to do is translate a number of materials. One of the things that's different, particularly to your issue, is that we're developing materials both orally and in writing for a population that's growing among working in plants and that's the -- population. So we've actually started down that road to attempt to do that, and I think the point is a very good one.

DR. KELLY: Yes, and I would be very interested in a copy of that article. Anytime we're preparing workforce training materials and I'm sure

we'll be working together on the outreach materials, the small and very plants, we think about the way people learn which is some people learn better through reading and visual representations. Some people learn better through what they hear. Other people actually have to get their hands on it and do it before they actually really understand it, and try incorporate all of those features into all of the training that we do, but training is always a work in progress, and we welcome any idea about things that we could do to reach people better. Thanks.

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MR. TYNAN: Mr. Elfering?

MR. ELFERING: Just in looking at this -all of the issues that we have to deal with, it's ever educational materials changing, have you put any for your employees Avian together Influenza, on knowing the differences between high path, non-H5, non-H7, low path Avian Influenza? You know, poultry industry has been testing for Avian Influenza for a number of years. I know our poultry industry, the turkey industry rather, has been testing for probably more than 25 years. We find Avian Influenza

every year, but it's all low path, actually probably typically swine influenza that is being picked up by poultry, and before too long, the broiler and egg industry is also going to be testing for Avian Influenza, and I think it would really be important that your employees -- they're going be to the communicators out there, anybody who works for USDA is perceived to be an expert in everything as it relates to animal disease and human health issues, and I was just curious if you put any information together for employees on AI?

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I can at least begin by saying DR. KELLY: that we have contributed people to a working group that have begin to put materials together. We have worked with people outside of FSIS and, in fact, we have even worked internationally to get digital images and a variety of kinds of things to put together, to together. Ι don't believe everything pull is finalized but I think we're well on our way.

DR. MASTERS: Yeah, this is Barb Masters. I think it's important to recognize that all of our public health veterinarians are trying to recognize

1 the symptoms and signs of Avian Influenza 2 poultry flocks, and what we're looking at is what materials we need and what we can do to enhance and 3 4 update that material for them. 5 So, yes, our veterinarians, public health 6 veterinarians have been trying but we are looking at 7 what we might be able to do to provide some additional materials to them to enhance and update and refresh 8 their learning and education on that information. 9 10 MR. ELFERING: Maybe as a follow up, this is 11 Kevin Elfering again. We are developing some -- we 12 actually have put together a couple of informational 13 brochures for veterinarians and companion pet owners and we're also -- for small, small scale flock owners, 14 15 I'd certainly make those available to you as nothing 16 more than a resource. So I'll get copies of those to you, or else we're just putting the final touches on 17 18 one related to food safety as well. 19 DR. KELLY: Thank you. 20 MR. TYNAN: Thank you, Kevin. Other 21 questions regarding the training update? 22 Thank you, Dr. Kelly. Thank you,

Ms. Cutshall.

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Under Tab Number 11, we have an update paper on the Cooperative Agreements Assisting Small and Very Small Plants and, Dr. Syed, are you here?

DR. SYED: Yes.

Are there questions on the update paper on the Cooperative Agreements?

Dr. Syed, that was a very well written paper. Fine job. Thank you for coming up.

Under Number 12, I beg your pardon -- yeah, under Number 12, we have the National Advisory Committee for Microbiological Criteria for Foods, and we have Gerri Ransom here that can perhaps answer your questions. Mr. Elfering?

-- a number of food-borne MR. ELFERING: cases associated with some chicken entrée illness product and the Advisory Committee made some suggestions, and companies who produce these products had to submit new label sketches to USDA by May 1st, and I'm just curious to know where that's at, you've received a lot of labels and have any new labels been approved and when would we expect to see

1 new product labels out on the retail shelves? 2 up to 17 cases now of food-borne illness associated with various species of salmonella just in Minnesota 3 4 associated with these products? 5 MS. RANSOM: I can update us on the Okay. Advisory Committee but not on the policy aspects. 6 7 Maybe Dan can --8 We also have Dr. Englejohn who MR. TYNAN: 9 has joined us and maybe can respond to that question 10 for you, Kevin. 11 DR. ENGLEJOHN: Thank you. And again, I'd 12 just like to say that it was the Minnesota Department 13 of Health and Agriculture that was instrumental in 14 getting this issue before the Agency and before the 15 Committee, and so we -- it was something that was 16 timely. It needed to be dealt with. To answer your specific questions, we did 17 18 set some dates by which we needed the industry to 19 to modifying labels for entrée products respond 20 particularly those of poultry that appear to be ready to eat but that can contain raw poultry product. 21

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There are three dates by which industry is responding

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right now in terms of the type of labeling. Labeling directly for products going directly to consumers had to be at least identified and, and establishments need to make information known about how they were going to modify their labels to address whether or not the products are, in fact, not ready to eat, and explicitly what they're going to be doing about validating their cooking instructions by May 1.

Another set of labeling relates to those labels for products that are -- for products going to this school lunch program. These would be products involved in the Food and Nutrition Services Commodity Purchase Program, and there's a different approval process by which they need to review those labels prior to them being submitted to FSIS. So we set a date of June 1 for those labels.

And then for those labels going direct to products that go to institutions, such as institutions that prepare the food for the consumer, we set a date of July 1, and it turns out that that is the largest segment of the industry with regard to product labeling, and we expect there to be roughly 5,000 of

those labels that will be coming in.

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To go along with this, the Agency issued letters to everyone that knew produced these we products, identifying to them that they needed to modify their label to make the name of the product explicitly identified as a non-ready-to-eat product, to identify that for safety, the product should be cooked to the minimum temperature of 165 degrees as measured by thermometer, and that the cooking instructions would have to be validated.

We did issue temporary labels for many of these products, and that temporary label by regulation is allowed for up to six months. And so from the perspective of the industry, the Agency will be issuing or expects to issue an FSIS notice in early September that would provide verification instructions for inspectors to verify that the labels have been, in fact, modified and addressed the labeling features that we issued in the letter.

So to make a long story short, there are in excess of 5,000 labels that are affected. We will be issuing specific verification instructions to our

employees to verify that all the labels for these types of products have been, in fact, modified and addressed, product name and validated cooking instructions.

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MR. ELFERING: This is a follow up. This is Kevin Elfering again. FSIS still has a fact sheet on their website that gives recommendations for cooking poultry, and one of them is to not use a microwave for cooking raw poultry, especially frozen. That seems to be in conflict with validating cooking instruction microwaving still would where be acceptable practice on the label. What are your thought about, you know, in one, in one regard you're saying don't, don't use a microwave to cook raw product but yet still allowing the label to say that this product would be considered safe if it's microwaved.

DR. ENGLEJOHN: The National Advisory

Committee for Microbiological Criteria for Food

specifically addressed that issue in the sense that it

was determined by the Committee that microwaving could

be safely accomplished and deliver a safe product,

that it had to be done in very specific ways and there

had to be specific information provided to the consumer. It was included in the report to say that it's not advisable to microwave products in a frozen state, a raw product in a frozen state. So they did not explicitly state that it's not feasible from a scientific perspective.

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From the Agency's perspective, we have made it known that we think that the burden on industry to demonstrate that they have validated cooking instructions is far greater for an establishment who is going to use and recommend that a microwave be used for these types of products that appear to be ready to eat but contain frozen or uncooked product in it that are in a frozen state. Part of the validation for the cooking instructions as recommended by the Committee and as the Agency is pursuing, is that there has to be information available to t.he manufacturer t.hat. demonstrates that the manufacturer has some knowledge that the consumer is aware of the type of product they're producing, that the cooking instructions are, in fact, practical, and I think that's an important word.

As you pointed out in the information that you shared with the Committee, cooking product from a frozen state in a microwave has varying degrees of results, depending on the power of the microwave, depending on the user. In many cases, it's children or elderly or individuals who are looking for a particular palatability or organoleptic quality as opposed to truly using a thermometer to measure for safety. And so we did make clear that our expectation for any label that would come in on these types of products that would include a recommendation that the product could be microwaved would have to have some fairly compelling data on file demonstrating that they have practical, feasible cooking instructions.

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We'll assess that. If we find that the product is -- that these issues are not be adequately addressed, then the Agency would further contemplate other policies that we would need to pursue with this type of product. One of which we made known to the industry was we that clearly do not want any additional outbreaks associated with this kind This was the third outbreak that we know of

1	that was rather major associated with the product, and
2	that an alternative could, in fact, be considered,
3	that raw product can't be used, that it would have to
4	be precooked. And so that would be our some of the
5	thinking that we would go into in terms of policy for
6	the future if, in fact, we find that there isn't a
7	good response on industry on addressing validated
8	practical cooking instructions.
9	MR. TYNAN: Additional questions regarding
10	the Micro Committee paper?
11	There being none, I want to thank Gerri.
12	Gerri, thank you for coming up.
13	MS. RANSOM: Thank you.
14	MR. TYNAN: I think on Tab 13, we'll move to
15	Tab 13, which has to do with Post-Harvest
16	Interventions to Reduce Salmonella in Poultry. We
17	have Dr. Patty Bennett who is here and who is going to
18	talk about that or respond to any questions you have.
19	We're getting off easy at this meeting I
20	think. Questions on Post-Harvest Intervention?
21	Okay. There being one, I'm going to let
22	Dr. Bennett go. If you have any questions, you can

1	catch her at lunch. Okay. Thank you.
2	Okay. We have before I bring the next
3	speaker up, I'm going to ask the question first. Do
4	we have any questions on State Reviews under Tab
5	Number 14? An Update on State Reviews. Dr. Harris?
6	DR. HARRIS: No one needs to come up. This
7	is a very quick straight forward question. Are we
8	still on schedule to be completed for all 28 by the
9	end of '06?
10	DR. MASTERS: Barb Masters. Yes.
11	MR. TYNAN: Dr. Harris, do you have a follow
12	up? That was a any other questions regarding State
13	Reviews?
14	I think last but not least on briefings and
15	updates, we have the Update on the Technical Service
16	Center. Mr. Elfering, and I'll see if Lynvel here?
17	DR. MASTERS: No.
18	MR. ELFERING: Actually, just a very quick
19	question. We still her periodically that there are
20	discrepancies from people in District Offices and the
21	Tech Center, that information that has come in out of
22	the Tech Center is not always in agreement with some

of the interpretations done in the District Offices, and I'm just wondering if you're hearing anything on those issues, and if there's been anything done to rectify it.

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While Phil's figuring out the DR. MASTERS: microphone, I will comment that one of the things you'll be seeing in the Strategic Implementation Plan on Small and Very Small Plants, I think is Phil is trying to provide some leadership over that. Certainly from the time I started, at the Tech Service Center, consistency is always an issue when you have that number of employees trying to answer questions, and so that's always a leadership challenge when you're trying to provide leadership for that number of employees but under Phil's direction, he's working with Dr. Roth and Matthew Michael who you met as a leader of the Program Evaluation Staff, and they are doing an evaluation of our Technical Service Center, including interviewing our employees in the field, also the folks at the Technical Service Center, and also they're going to be doing a Federal Register notice to get public comments on the Technical Service

Center services.

Additionally, another component of that is they're starting to post commonly asked questions on the website and that helps to see in one location all the commonly asked questions and that helps to get that consistent answer by posting those questions in one location. Additionally, they're starting to do a process of collecting questions and answers at the Tech Service Center and collect those in a database at the Tech Center. So I think Phil's trying to address that in multiple ways at the Technical Service Center.

DR. DERFLER: I don't really have anything to add. The only thing I'd add to that is among the commonly asked questions are questions having to do with recently issued directives and notices, and so we're collecting the questions that we get about them and we're posting them. There's a link on the Tech Center website and they're also listed underneath the specific directive or notice that they address.

MR. TYNAN: Other questions on the update on the -- I'm sorry. Mr. Link.

MR. LINK: Charles Link. Just on the IKE

1 area, the comment, I'm just curious as to are you 2 getting comments and how is that whole system working o United States? 3 4 DR. DERFLER: We are getting comments. What 5 draft, Interactive Knowledge do is we post a 6 Exchange, to provide an opportunity for comment on it. 7 We then considered those comments in final and post a 8 final version. There are some comments that we get, 9 and we use them in putting together the final version. 10 MR. TYNAN: Other question regarding Tech 11 Service Center? 12 folks, if there's Again, some οf our 13 questions that come up during the breaks lunchtime that you want to pose, if the folks are not 14 15 here, we'll try and get the answer to you at a later 16 time. I think it's important to probably the meat 17 18 of our meeting, which has to do with the actual issues 19 for discussion that you will be doing in your Subcommittees this afternoon. 20 And there are two. the first is measuring establishment risk control for 21 22 risk-based inspection. There is a paper under Tab 6,

and I have Mr. Don Anderson from our Office of OPEER that will be -- that's the Office of Program Evaluation, Enforcement and Review, that will be doing the presentation to kind of pose the discussion just a bit. And we're getting some light on the screen. Can everybody still see it or do we need to move for just one moment.

MR. ANDERSON: I hope you can see that okay.

I know it's getting some sunlight on it there. Thank
you very much, Robert.

I'm Don Anderson. I believe that FSIS has made some good progress on several fronts in advancing risk-based inspections. We're going to talk about a specific area today of risk-based inspection, but I think an extremely important one, and that is how FSIS can measure how well establishments control the risks in their operations.

Last November FSIS reminded the Committee that under traditional inspection, you have two responsibilities that we have to meet. We have to conduct carcass-by-carcass inspection on slaughter lines and we also have to conduct inspections in

processing operations at least once per shift. And also as explained in November, okay, there is no plan to change either of those requirements.

Rather, we intend to add a new dimension as to how we do our inspections and that is to do inspection on more of a risk basis. So it's an additional dimension.

We are going to be considering, or we think we'll be considering several different types of risk factors to make our resource allocation decision.

We'll look at hazards. We'll look at likelihoods of hazards. We'll look at the exposure potential in establishments.

What we want to focus on today is how well establishments control the risk in their operations. So it's a particular part of these risk-based inspections.

As Dr. Raymond said this morning, we want to make sure that we're using data the best we can to make use of our inspection resources in the most effective ways that we can. So today's discussion is about risk-based inspection in both slaughter and

process establishments and processing only establishments.

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Today we have approximately 5500 active federally inspected establishments that are under So there's about 5500 plants that HACCP requirements. we're talking about today, and approximately a quarter of those combination slaughter/processing are establishments and approximately three-quarters are what we refer to as processing only establishments. And we are talking about how we allocate resources in all 5500 of those establishments but we're not -- we talking about how we allocate inspection resources to the slaughter line inspection. That's not a topic for today. That doesn't change under what we're talking about today. Rather, we're talking about risk-based inspection of processing.

Furthermore, we're not talking about how we do inspection in egg products establishments. We're talking about federally inspected meat and poultry processing establishments only.

You see on this slide what we consider to be the several important components of what we anticipate

to be a Risk-Based Inspection System. We're going to talk a little bit about each of these now, food safety implementation, food safety system design, system pathogen control, in-commerce findings, enforcement actions and then we've got sort of a catchall other considerations that we don't think necessarily fit nicely into one of those components. We understand, we acknowledge that there may be a little overlap between some of these components. We think that's probably okay. We think it's more important that we capture all of the data and all of the information risk control and if there's goes to overlapping factors, that's okay. We want to make sure that we don't want to miss anything important, and we'll have a question for you about that later.

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Let's talk first about the first Okay. component, food safety system implementation. Our measurement objective here, what we're trying to do, is measure how well and maybe how consistently if you want to think of it that way, an establishment controls risks in its operation. own Okay. Noncompliance records or NRs, which have been referred

to several times in the discussion today already, document all noncompliance that inspection program personnel observe, whether they're food safety or other consumer protection. Risks will not change under risk-based inspection. We will continue to document all regulatory noncompliance.

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However, we do not believe, the Agency does not believe, and this is what was brought up earlier, NRs are equally indicative of how that all establishments control risks in their operations. so what we're currently doing is discussing what types of noncompliance, what types of regulatory deviations do we think are more important or better indicated than others of how well establishments control risks We're trying to look at what in their operations. types of noncompliance are more important than others, and if you think some are more important than others, how can we weight the additional weight to those noncompliances that we think are more important than measure others in producing а of how well establishments control their risks.

The second component, which was also talked

a little bit about today, is food safety system design. Dr. Raymond again emphasized earlier this morning how important it is to some of the Agency initiatives we have to try to improve this, how important it is for all establishments, regardless of size, to have robust designs. Okay. Effective food safety risk management again is essentially embodied in prerequisite programs, sanitation standard operating systems and HACCP systems.

All else equal, we believe that establishments that have better, more robust system designs, are going to be better able to control the risks in their establishments.

Currently today, FSIS' best window or best look at how well establishments -- how robust the establishments design -- food safety system designs are, are what we call food safety assessments. We conduct food safety assessments and establishments on an ongoing basis, and we think that they are the best way to get a good look at how well designed food safety systems actually are. So what the Agency is discussing now is how best you can capture the

information from food safety assessments or from other sources to get at the robustness or effectiveness, kind of intrinsic effectiveness of the design of the food safety system.

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The third component is pathogen control. There's also been a little bit of discussion about this this morning. The Agency has several pathogens microbiological testing and testing programs or Ι should in federally inspected programs say, establishments. Wе do conduct pathogen testing programs of several different types of ready-to-eat We conduct listeria testing of ready-to-eat We conduct salmonella testing of ready-toproducts. eat products. And the question came up, do we conduct E. coli 0157:H7 testing in ready-to-eat products, and the answer is yes, in certain of those beef and -products we do. I think the last time I checked, there were several hundred establishments in which we're conducting E. coli 0517:H7 tests in ready-to-eat products. There are several thousands establishments actually that have -- that are subject to some of our ready-to-eat testing requirements.

FSIS, of course, also conducts Ε. 0157:H7 testing in certain types of raw ground beef currently about 1600 products, and there are establishments that are subject to those testing requirements.

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And finally, we conduct salmonella testing, salmonella verification testing in approximately 2,000 establishments that do combination of some slaughtering boilers, cattle, market hogs, soon to be turkeys and also establishments that product ground beef, ground chicken and ground turkey. So there's a lot of data from t hose 2000 odd establishments on how pathogens well they control think in that we salmonella testing data.

So what FSIS is doing is trying to solicit different ways that we can factor the results of our testing programs into our measure of how well establishments control risks. We consider that to be a very important component of risk control.

The next factor that we want to talk about are in-commerce findings. Products that are potentially deleterious to public health food

sometimes in our commerce, and when they do, we sometimes see, frequently see product recalls and/or consumer complaints, or there may be other events or incidents that will bring to the Agency's attention, okay, that there are potentially deleterious products of in-commerce.

So let's see. In about the last year there were approximately 40 health related recalls and there were several thousand consumer complaints. We have a recall database. The Agency has a consumer complaint monitoring system, and again we're looking at ways at a minimum to bring data in from recalls and bring data into our system from consumer complaints so that we can include both types of findings in our measure of how well establishments control them.

Enforcement actions that occur are normally the result of either food safety assessment. We mentioned food safety assessment a few moments ago. Enforcement actions typically result either from food safety assessments or from kind of a culmination, if you will, the logical culmination of implementation problems that are documented through serious types of

noncompliances. However, there are occasional instances when enforcement actions are taken against establishments for reasons other than those, and although it's not a very common occurrence, we do believe that it may be important to enhance inspection in establishments in which those types of enforcement actions are taken, and if we can't -- if we're not capturing them somewhere else, we want to make sure that we capture enforcement actions in such instances, and so we'll be looking at ways to do that.

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Finally we have as I said kind of a catchall area that we refer to as other considerations, and these are sorts of findings or data elements that aren't necessarily part of our traditional data example, federally inspected system. For establishments that product and ship beef other intact beef components to other federal establishments that in turn produce raw ground beef, consider those types of establishments, and if we get a positive E. coli 0157:H7 sample from a raw ground beef product, and we find that suppliers, if there are certain perhaps suppliers of establishments that test

positive providing beef trim to those establishments. They show up in what we call a steps database which is our system E. coli O157:H7 positive supplier database. I know that's a mouthful. But basically we think if an establishment is showing up as a positive, as a supplier, the E. coli O157 grinder, then we would like to recognize that and bring that into our measure of how well those establishments are controlling risks.

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We may also want to enhance inspection, of in establishments that ship products that course, positive pathogen agricultural tests in marketing service school lunch program -- testing program for example. So there are just a couple of examples of the other types of data that we think we should of how well bring into our measure establishments control risks.

And now let's turn to the questions that we'd like each Committee to consider this afternoon. The first question is are these all the appropriate objective for measuring risk control? Should any objectives or corresponding features be deleted? Or,

should any be added? If you would jump the slide 4 for a moment.

These are the components we're talking about. We would like you to consider this afternoon whether these are the right things that the Agency should be looking at and considering in our measure of how well establishments control risks. So that's the first question that we'll ask you to consider. If you'll go to 12.

The second question, once you've identified what you think are the important components that we should include in our measure of risk control, then ask yourselves this please. Are some of these components more important, that is, are some of these factors better indicators than others of how well establishments control risks? And it's a two part question because if you think some are more important than others, should we give some of those components more weight in our measure of this control than other components? So that's question number two.

And lastly, we would like to know what the Committee thinks about including in our measure of

1	risk control, kind of acknowledging if you will on our
2	measure of risk control, that certain establishments
3	have very effective implementation or very effective
4	system design. If an establishment for example, a
5	food safety assessment indicates that an establishment
6	has an especially robust design, food safety system
7	design, perhaps we should acknowledge that in our
8	measure of risk control so that we don't do as much
9	inspection in that establishment as we might
10	otherwise. So that's the third question that we'll
11	ask the committee to consider this afternoon.
12	So I'll pause and see if anyone has any
13	questions at this time.
14	MR. TYNAN: Questions from the Committee?
15	Dr. Carpenter who was first.
16	DR. CARPENTER: David Carpenter. I want to
17	go back to pathogen controls. You talked about
18	salmonella verification testing. That means a number
19	of things to me. So if you could just elaborate.
20	Does that mean that you're going to look at the

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ability to detect salmonella in the product in the

environment or does it mean that the test is adequate

1 for finding salmonella in any product 2 standardized methodology, or does it mean that salmonella is an appropriate indicator organism to say 3 4 that you're really cleaning up this, this operation? 5 MR. ANDERSON: Well, I think those are all three good questions, and I know we're capturing 6 7 those. I think that's something that we should I do know that the agency does both product 8 discuss. 9 testing and environmental testing, at least we 10 environmental testing in certain establishments, and I think that we will discuss whether we should include 11 12 both of those. 13 Т this time can't speak at to the verification of methods and those sorts of things but 14 15 I think that's a very good question because what is 16 important is the data that we're relying on and I measure this default was validated. 17 We want to use validated, of course. 18 19 DR. MASTERS: David, I'll add to that, and I 20 think those are good questions and ones we would want I think those are important things for 21 your input on.

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us to talk about and consider, and at this point, what

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we had wanted some input on is are salmonella control in raw products, is our salmonella testing one we talked about, when Don talked about our salmonella testing in 2000 plants. He was referring to our pathogen testing that -- pathogen reduction testing that we do as an Agency, that we're doing in the products to meet our pathogen reduction to form a standard that we're doing as part of our HACCP regulations that we put out, and that was the testing that he was referring that we're doing in our own That does not preclude us from discussing laboratory. and talking about the types of questions that you're putting on the table. So I think there some types of discussion that should take place this afternoon if that's helpful.

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MR. TYNAN: Mr. Govro?

MR. GOVRO: Mike Govro, Oregon Department of Agriculture. I'm looking at what you've written down here from the standpoint of someone who deals more with FDA programs, milk, shellfish, food manufacturing and probably the greatest portion of my time is in the retail program, and in the retail program, it's fairly

focused, and we look at the five identified factors CDC has identified as the risk factors, and they actually said these five things are the ones that are most closely associated with food-borne illness. I think they've even got percentages and ranked them, and this kind of looks back to your question of wading the NRs, and I think that would be a good way to go about it, and I'm just wondering if FSIS has done anything similar, has information from CDC that has identified practices in meat plants that are specifically related to food-borne illness and final product particularly ready-to-eat products production of adulterated products and if you think that information would be obtainable and useful.

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MR. ANDERSON: I don't know that the Agency has. Maybe someone else can speak to that. I am somewhat familiar with the FDA's priorities or weight. One of the things I want to remind us is that what we talked about mostly is, is different types of data that we might bring into a measure of establishing risk control. If I remember FDA's criteria properly, they also look at some of the so-called inherent risk

factors, and those are things that we are talking about including in our Risk-Based Inspection System but what we talked about, and so maybe we're kind of thinking -- talking a little narrowly here, but we're talking about bringing into a measure of risk control certain types of data, but there will be other types of data that go to other aspects or risk, that will also factor into this risk-based inspection, and I know that some of the FDA factors are --

DR. MASTERS: And, Mike, I can certainly tell you we've been working with CDC closely and I'm not aware of this specific document, but I can assure that we've met with CDC. Dr. Raymond happens to be bringing with him public health background and has been meeting with Dr. Gerberday (ph.) and so it's something that we can certainly put on his agenda, and so we'll certainly pursue trying to get our hands on that document. So I appreciate your bringing that up.

MR. TYNAN: Mr. Kowalcyk?

MR. KOWALCYK: I have a couple of questions.

One's related to the ranking of the or weighting of the NR report, and it also involves the pathogen

control measures. What resources has the Agency put up against these tasks? I mean what is the Agency's thinking in the way of who in the Agency is going to be charged with looking at this data to see if an appropriate wading scheme can be developed as well as applying what's really regulatory sampling for pathogen control measures in a risk-based system like What are the Agency's plans? I mean these are this? big issues to tackle. So I'd like to get an idea of what the Agency's looking to do internally to answer these questions.

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DR. MASTERS: To jump in here a little bit,
Don. We currently have a Steering Committee and a
workgroup that has begin to work on these issues which
is where this particular document has come from, and
so as Don indicated in the beginning there, there are
a few issues that came out of the last NACMPI meeting
and this is one from which he is the lead on the
Steering Committee and a lot of the work that will
come forth and come to this Committee as well as to
our third party will come from. Obviously the Agency
has to generate ideas so we can bring them forward,

and so there's a Steering Committee that is currently
generating these ideas so that we can bring them
forth, and that's kind of the division that we have at
this point, is having this Steering Committee that's
made up of predominantly our Headquarters employees
out of our policy group but there's representatives
from all eight program areas that sit on the Steering
Committee, that they would generate ideas and when
they're ripe enough to bring forward to this Advisory
Committee and to our third party group that we will
bring them forth so that we can get input before we go
too far that they're not able to step us back for
those kind of things. So we felt like we had reached
the point at which we could bring this one forth to
the Committee and to the third party. So this is the
first one that we've brought forth to the Committee
and will be moving forward with.
MR. TYNAN: Michael, did you have a followup

MR. TYNAN: Michael, did you have a followup question?

MR. KOWALCYK: No.

MR. TYNAN: Mr. Finnegan?

MR. FINNEGAN: Yes, I have a couple of

questions. Here on the components of risk control measures consider only significant food safety NRs. Are the very nature of NRs going to be changed like a major/minor critical type thing or how do you -- do you know what the true food safety NR like O157:H7 positive from a clerical, or do you have some kind of a system to determine that?

MR. ANDERSON: I think that -- again, those are good questions but I don't think what we're talking about at this time would involve actually changing the way that inspection personnel document noncompliance. Regulatory noncompliance, as I say, will continue to be documented.

What we're, what we're trying to do in our various working groups and what we're asking the Committee to actually consider this afternoon, is whether indeed some types of regulatory noncompliances are more indicative of risk control problems than others, and if so, which ones those are or how can we go about identifying which ones those are so that we can, can give greater weight into our measure of risk control, those types of regulatory noncompliances.

MR. FINNEGAN: Okay. And how about those food safety assessments? I've seen EAIO reviews that were I mean very small plants, the one I'm thinking in particular, slaughter only one day a week and all they did was, you know, cut and grind raw product. I mean that was like 26 pages. You know, is there some kind of a system where you can just highlight the food safety, food safety critique rather than have all them pages that somebody's going to have to go through? Just to narrow it down, simplify it, streamline it.

MR. ANDERSON: Yeah, I think the answer's yes. We, we -- food safety assessments are documented I believe on Forms 5000-8 I think is the current documentation form. Mr. Palesano is affirming that. So we have a form that captures the results of food safety assessments and a lot of that information is in narrative, but there also are some sort of findings and categorical findings at the bottom of the 5000-8. Again, we're considering how we can use information either from the existing 5000-8 or from some modified version of that but we think it's very important to do what we need to do to capture the salient information

1	from the food safety assessment process. If we're
2	capturing the salient information today, that's fine,
3	and we'll use that information as appropriate in our
4	measure of risk control. If we're not capturing that
5	information or all the information that we need to,
6	then we want to make sure we devise a way of doing
7	that. So that's something that we would like your
8	suggestions on, suggestions on.
9	MR. FINNEGAN: Right, I agree that the food
10	safety assessment is a very good tool. I was just
11	thinking of some way to streamline it a little bit to
12	get to the pertinent facts.
13	MR. ANDERSON: Absolutely. We think that's
14	critical, yes.
15	MR. TYNAN: Ms. Eskin?
16	MS. ESKIN: Under the list of risk control
17	measurement objectives, you have a heading other
18	enforcement actions.
19	MR. ANDERSON: Yeah.
20	MS. ESKIN: I'm looking at the document in
21	our folder, and it says prior enforcement actions
22	resulting from causes not captured by the other

components. Could you elaborate? I'm assuming those all have some -- these enforcement actions that have some food safety implication, but can you give examples of what that might be?

MR. ANDERSON: Examples of such food safety assessment?

MS. ESKIN: No. Of --

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MR. ANDERSON: Excuse me. Of --

 $\mbox{MS. ESKIN: $--$ other enforcement actions,} \\ \mbox{what would be an example of $--$ }$

I think there are a couple of MR. ANDERSON: examples that have come up. One is that enforcement actions are sometimes taken when inspection personnel observe extreme humane handling violations in establishments, we'll take enforcement actions There are other examples that other -- there that. have been cases where enforcement actions have been taken against the establishments because there's been cases of inspector intimidation if you will. What we'll have to do, one of our challenges will be to identify what types of enforcement actions -- what we're trying to isolate are those type of enforcement

actions that have a bearing on how establishments control this to make sure we have captured them.

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DR. MASTERS: Sandra, to get to your point, we were looking for those that relate to food safety, and in this case, we're looking for those, as Don mentioned earlier. Most enforcement relates to either failure to execute the program over time which would have been documented NRs or to the design of the program which would have been documented through a food safety assessment.

However, because we're not consistently in every plant every day on the design of a program, there might be a circumstance that a plant shipped product that had E. coli O157:H7, for example. in to do the food safety assessment and see that they cooking their product particular were not at а temperature. We go ahead and suspend the plant before we complete the food safety assessment. So we don't have a documented food safety assessment. They were implementing their product as it was designed. We don't have a documented food safety assessment in the system, and so we don't have that particular

component. So there's an example where we would not We don't have NRs have a food safety assessment. because the plant was doing a good job of executing the program as it was written but because we found initially went not well documented when we in а go ahead and take the we would have to program, enforcement actions for the safety of public health, and over time we would ultimately get that food safety assessment but for public health and welfare we felt the need to go ahead and suspend operations so they could get a well designed program in place.

MR. TYNAN: Mr. Govro?

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Mike Govro, Oregon. MR. GOVRO: Another different -- between question related to а the emphasis that FSIS and what happens at the retail level, an emphasis in the food code. And it has to do illness, exclusion of with employee hygiene, ill workers, and there seems to be a different. I looked up in I think it's 5001, Bill sent me a reference to in your briefing paper here, it talks about safety inspector performing a variety of consumer verification procedures which is was a little bit

unclear to me as to what that entailed. So he sent me this document, and in reading through it, on page 1it says if inspection program personnel have 20, questions about an employee having an infectious disease, he or she should discuss this with plant Inspection program personnel are not management. trained to diagnose infectious diseases which is quite a different from the emphasis at the retail level where there are four, and I think now five identified major illnesses, both managers and inspectors required to know what they are, what the symptoms are, and management are required to exclude employees who are ill with those illnesses. And so what I'm wondering is, is are these illnesses not a concern in particularly producing USDA plants, ready-to-eat Should we worry about it like we do at the products. food service level or perhaps should there be some more emphasis on that aspect of it?

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DR. MASTERS: This is Barb Masters, and I think that's a good point, Mike. However, from a FSIS perspective, we have traditionally put that burden back on the regulated establishments but I think it's

certainly something you should be welcome to raise at the Subcommittee, but we don't want to leave anybody with the impression that there's not a concern within FSIS or the regulated industry but typically this burden has, and by regulation, the industry is to preclude anyone with symptoms or signs of illness that would cause concerns and industry has taken that responsibility and borne that responsibility.

MR. TYNAN: Other questions for Don regarding the establishment risk control?

Well, with that, I would suggest we perhaps close out this particular issue. We're early on our agenda. I think we were pointing on lunch at 12:00. Maybe we could take a bit longer and go until 1:00 and come back and we will begin issue number two. Is that agreeable to everyone? Okay. I'm looking for a motion to adjourn for lunch? I didn't hear one, but I know you all are going to do it anyway. So -- all right. Thank you very much. We'll see you at 1:00.

(Whereupon, a luncheon recess was taken.)

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(1:00 p.m.)

afternoon portion of the MR. TYNAN: --Tuesday session. And at 1:00 on everyone's Agenda, we have the second issue which is the Strategic Implementation Plan for Enhancing Outreach to Small and Very Small Plants, and we have Dr. Karlease Kelly, our Chief Training Officer here, and Bobby Palesano, one of our Deputy Executive Associates with OPPED, and they're going to do the presentation. Dr. Kelly.

DR. KELLY: Good afternoon. I hope everybody had a good lunch, and you're ready for a busy afternoon. We're going to review with you our plan to redesign the FSIS outreach effort to meet needs of the small and very small plants.

To begin with, I want to give you a little bit of a background. I realize there are some of you who are quite familiar with this, but there are probably others of you who might appreciate a little bit of that background.

So as part of that background, obviously from our perspective HACCP is the foundation for

success for all meat and poultry establishments, and we believe that making the nations food supply safe is essential for public health but it also makes good business sense for the industry and practice. Effective HACCP implementation is critical to public health, and we have a role in our society to make sure that establishments have the tools that they need to maintain a safe food system. So with that as a background, another piece of information I'm sure you all are aware of it that there's a large number of small and very small plants, small and very small according to the definition of the Small Business Administration, they're scattered throughout the United States. This map shows you a current distribution and shows you where they're concentrated and where they are scattered.

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They're a very important part of the nation's food supply. As Dr. Raymond mentioned this morning, anytime you're eating a meat or a poultry product, you may or may not know where that product came from. You may not know, it may be coming from a small plant. Many small and very small plants are

suppliers or end up interacting with in some way with the large plants and products are intermixed and intermingled and goes many places. So they're a very key component of the nation's food supply.

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listening FSIS engaged in some sessions recently, outreach and listening sessions, with small and very small plants. Some of you participated in those at least from a point of view of being there to host the session or to join us, to listen to some of the things that small plant owners and operators had to say. Bobby Palesano, who is here with me today is the co-leader of the Task Force, attended a couple of We had sessions throughout these sessions. the country, Montana, California, Texas, Pennsylvania, We heard a lot of different things Massachusetts. from the small plant owners and operators. We're very interested in hearing about their needs, about their challenges, about the kinds of things that they would like to see from us in terms of assistance.

In addition to that, as background and somewhat as a catalyst to this, in December, the International HACCP Alliance hosted a meeting to do

some strategic planning to identify the needs of the small and very small plants. Dr. Masters attended that. Dr. Raymond attended that. Bobby Palesano and myself attended. Joe Harris was there. I may be missing some of you who are here who were there. Ιt of people who represented was group several Government agencies. They as well as academia and we had some very small plant owners who were also present to react and to provide some input to that.

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What we learned through the listening sessions and through the needs identification process that we went through in Dallas, is in general, I mean we could give you a long list of things we learned, but in general, what we learned is our strategy up to this point has been focused on helping small and very small plants with developing their HACCP plans, other words, with an execution side of the HACCP rule. However, the kind of things that people are needing today are in a different direction. They're in the direction of designing a HACCP plan. So what that we needed to focus our learned is the scientific strategy on supporting basis for

developing and maintaining a HACCP plan.

We all know that the environment we work in is one that's very dynamic. We continue to learn through science and through practical experience, a lot of different things, and that means that the plans and the ways that we're implementing our food safety systems are going to have to change as we get more information. We know that the Agency's policies have changed. Therefore, we know that plants are going to need to change to address new information, things that happen with pathogens and so forth.

So the Agency stepped back, looked at what we had gathered, the information that we had gathered and the things that we have learned, and in response particularly to the Dallas meeting, Dr. Masters asked us to form an internal Task Force. Bobby is coleading that with me. There are a number of people in the room who are on the Task Force who are here that you could talk with. Mary Cutshall, Cheryl Hicks I saw. There may be others that I'm just overlooking at the moment.

But the Task Force represents all eight

program areas in FSIS and our charge was to look at the needs that are posted on the International HACCP Alliance website from the Dallas meeting, and to develop a strategic implementation plan to outline the things that FSIS will do. At the Dallas meeting it was recognized that industry may be doing some things as well, but as a Task Force, our focus is on the things that FSIS will do in response to these needs.

Essentially in your materials, what you'll see is a copy of the plan and the plan is organized around seven basic themes. So first I'm going to run through those basic themes, those seven basic themes with you, and then I'll give you some examples of more specific action items associated with those themes.

The first theme that you see here if you can't see through the sun and the shadow is the strategy of a one-stop customer service. What we found when we met as a Task Force with all those eight program areas, is that each program area actually contributed some part in serving the needs of the small and very small plant owner and operator.

I will tell you that my dad is a small

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business owner. He's not a meat or poultry plant operator, but he's a small business owner, and I can only imagine how frustrating it might have been for him to have eight different offices to call to try to get information. So when we saw that as a group, we realized that that had to be a real key component of what we were doing. We know that the small plant owners and operators don't have a lot of time. We need to make it as easy as it can be for them to access information if they're reaching out and they're trying to learn and trying to improve their food safety system, and their food defense systems as well.

So we're working to draw together all the information so that someone in FSIS does the legwork. If a plant owner and operator needs information, they don't have to call eight different offices. They call one place or they e-mail one place and someone who intercepts the call or the e-mail does the legwork to get that information for the small plant owner and operator.

Technical resources is another key theme that came up over and over again. For example, those

of you who are really familiar with FSIS know that we of material that mountain sometimes is difficult to wade through, difficult to try to figure out. I have a need. How do I match my need with all of these things that FSIS has to offer. So technical resources, and in some cases we're finding that we may have a mountain of things, but there may be something There may be one thing that a small plant missing. owner or operator needs that we just don't have. So that's another theme.

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Education and training is a theme. It has to do with the fact that the small plant owners and operators may not be able to afford to give a lot of access to resources, and as Dr. Raymond mentioned this morning, we're going to try to reach out to them through a variety of methods to make sure that they have the type of education and training that would be of benefit to help them improve their food safety and defense systems.

The other four strategies, one has to do with partnerships. Wе already had in place partnerships with Extension, with State, with

academia, but we wanted to revitalize those partnerships perhaps those and even extend partnerships even further, to get people more actively engaged and engaged in this effort that focuses on scientific basis for this food safety system. I think it was mentioned this morning that we're looking at partnering with the World Development Agency, that we may be able to route small plant owners and operators to the Rural Development Agency to let them know that if they need to make facility changes or equipment upgrades, there may be some type of way that they could get low cost loans through that Agency.

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So we're looking at how we can expand our partnerships outside of FSIS, to help small plants owners and operators, to get people engaged in this more fully.

Another theme that we felt was an important strategy was needs assessment. Those of you who have done these assessments know that in some ways it's like peeling an onion sometimes. You get a big picture, and that's really kind of where we are right now. We have a big picture. We know that there are

needs, but when we actually try to go out and pinpoint some of those needs, what we find is that we need more information to actually really be sure that we meet that need. So we need to do those needs assessments regularly in an ongoing way so that we don't find ourselves, you know, five to seven years out again saying, oh, we need to, you know, look at this and make some changes again. So needs assessment is the fifth theme.

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The sixth theme was evaluation. We have a We have a lot of activities that lot of resources. are going on right now but we don't have a lot of data that shows this. Where are we really reaching people. You know, as I said, we have this mountain of materials but small -- very small plant owners in the listen sessions are saying, we need something. So we need something with our evaluation that defines where is it that we're hitting the targets, where are we off base, and how can we bring all that together better.

And last, levering resources. At this time, under the constraints that we're operating under, we decided we're not going to reorganize at this point.

We're simply going to work together in program areas. As we discuss some of the issues, one of the things is that learned different people might we have actually been working on some of the same kinds of things in different offices. So we're trying to leverage that, put that together, so that people are working together, ending up probably with better products. trying eliminate So we're to any duplication, and then bring some synergy together with all these different program areas.

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So those are our seven strategies. Here are some highlights, some features, some of the things that we're working on right now to specifically bring those strategies to life.

One of those strategies that Mary Cutshall and the CIPO (ph.) staff are working on is working on the web page, consolidating all the information that the Agency has that would benefit the small and very small plant. We found again that that information was scattered all over the website, and that some of the information on the website could be reformatted to focus more on their needs, and Mary's already begin

some of those efforts right now at low or no cost, and she's making plans for some of the things that have bigger costs for the future.

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The website is listed here, and later this afternoon when we get into our working groups, those of you who are working on this issue, if you're interested in going to the see that website, we'd be very interested in having your comments on how we might be able to make that better.

We also have a toll free number that you're going to call to get support. That is at the Technical Service Center. We have not actually gone through the long and bureaucratic process of forming a staff that's going to deal with this, but we have created what we call an ad hoc group. We've taken the There is a group of people that is first steps. devoted to meeting the needs of the small and very small plants, and we're working on that right now, so that they're going to be the ones that someone's calling to make sure that the caller doesn't have to They know where to call and they call eight places. can get the information or they know when the e-mail

comes in, how to get that information for the small plant owner and operator.

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We're planning at some point we're going to have some additional listening sessions. We would still invite people to share ideas, to share their It's important for us to continue with those, input. and some other examples of things that we're doing is action items. We're creating a clearinghouse for questions and answers. That was one of the things specifically mentioned at the International Alliance as one of the needs. People are saying there are questions all over the place, answers all over the place, let's be able to go to one place so we can look at that, and we're working on doing that to get it posted on the website.

We have some plans to begin with joint training that would include plants and inspectors. This summer, at some point, you'll be seeing us working into some regulatory workthroughs, to meet that action item.

In addition, we're looking for validation support to meet the specific small and very small

plant needs. We're hearing from small and very small plants that there may be some types of products that they don't have, the type of information they need for the processes that they have to provide validation support.

So in big picture terms, what are the outcomes that we're looking for? Well, in general, we're looking to make sure that the small and very small plants have a full range of scientific and technical support and assistance that they need to compete by demonstrating that they have safe and effective HACCP systems and that we have some results that show that we have enhanced food safety and food defense protection.

Now we'll just walk through the questions that we're going to cover in the breakout group this afternoon. We've got four of those questions.

Our first question is what suggestions do you have for how FSIS through the International HACCP Alliance could locate industry representatives who are willing to share expertise and other technical resources and assistance with very small plants? In

other words, how can we engage the industry through the International HACCP Alliance in helping with the small and very small plant needs.

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Our second question is what suggestions do you have for how FSIS could obtain data on the types of support that small and very small plants need for their food safety systems? This goes back to that statement earlier about peeling the onion. Right now we're hearing that people need some type of support and when we actually ask, we're having difficulty getting some specifics on that. So it would really be helpful to us if you could help us with that question.

The third question is what suggestions do you have for how FSIS could best work with a users group consisting of all partners to provide feedback on the usefulness of existing tools and services to pilot new activities or materials and to make recommendations improve on how to the outreach functions to better meet the needs. In other words, we would like to test run some of our things that we are getting ready to introduce, or we would like to test some of our modifications. Obviously since we

1 have a raft of resources and we think they're really 2 good, maybe it's something simple like people just don't know they exist or they don't know where to find 3 4 them, but maybe there's something more fundamental. 5 So we need some feedback from a users group like that. Our last question is what suggestions do you 6 7 have for how FSIS can obtain data on the types of 8 support -- we just went through that question. 9 Our last question, generic а very 10 question -- I just wanted to make sure everybody's 11 I think we are awake. What other suggestions awake. 12 do you have for FSIS in strengthening our strategy for 13 outreach to small and very small plants. So in other 14 words, just in general, we're interested in all of 15 your idea about this. 16 So with that, I will -- that's the end of 17 the formal presentation. Bobby and I will take some 18 questions. Dr. Elfering? 19 DR. ELFERING: Back HACCP implementation, 20 there was a system that was set up where every State had a State contact and a State coordinator for HACCP. 21 22 Did it work and would it work again to kind of re-

implement that system where you had some contact people in every State and maybe some coordinators? Or maybe it didn't work very well. I don't know. I know that, you know, pre-HACCP implementation, we used to get a lot of calls and maybe it was just not a very good system.

DR. KELLY: I will just start the dialogue, and I think that when you're -- the time period you're talking about, it did work, and I think that we are interested in revitalizing that network. As I mentioned, we are moving away from the strategy that we started with to something new which means that it's time to revitalize, time to make those contacts again, time to engage in that, and other people may have some comments.

DR. MASTERS: This is Barb Masters and,
Kevin, I would that there are still contacts in each
of the 50 States and those are still posed on our
website and we still utilize those contacts and
coordinators, and I think what Karlease is saying is
true. They were very effective at the implemented
HACCP, and I think we move through the process of

implementation and we became more comfortable and we were all more confident with implementation, we all kind of moved away on relying on those contacts and coordinators, and we are looking at re-energizing those contacts and coordinators and your input on how effective you thought they were in the beginning. We found it to be a very effective resource. They still exist and I think we're very open to the idea of re-energizing and re-utilizing those coordinators as they were utilized in the beginning of implementation.

MR. ELFERING: This is Kevin Elfering again.

One of the reasons I asked because I was one of the -- I'm the State contact, and I guess we don't know that it's still being utilized, and it would be great. I mean we used to get a lot of calls. I still get a call every once in a while. So I think it might be another resource for you to be able to increase the outreach to some of these very small plants.

MR. TYNAN: Mr. Schad.

MR. SCHAD: Yeah, I just wanted to make a comment on that idea. I think it is and can be very effective but I think that would have to be part of

the clearing house on the Qs and As, so the answers across the country were the same no matter what coordinator you spoke with.

DR. KELLY: Other questions?

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I don't have a question but I DR. RAYMOND: have a comment. One of the things that I kind of like reflect on when we did these listening sessions and did the HACCP Alliance, the International HACCP Alliance meeting, we found a lot of fingers were pointing at the Agency. Why are we having trouble with some of the very small plants not having robust HACCP plans and, you know, the diverse answers to the same question for instance, and it's fairly simple to correct by having a Q and A. There were things that we just weren't doing well. There were times that I think we felt that everybody -- maybe didn't feel, but we failed to remember that not everybody had a QA person full-time or half a person full-time, and just hadn't taken it into consideration.

So I guess the point I want to make is we're doing everything that we think we can do. We want you to tell us what we've forgotten to do to try to

revitalize this outreach, but I realize that there will still be some very small plants and some small plants that will not come to the table, that will not watch a DVD or a CD-ROM, and those are the ones that we will begin more enforcement, more food safety assessment but we will make this effort so no one can say we didn't try. So it has to be a two way street. The small and very small plants that aren't up into the 21st Century must make that effort. We'll do everything we can to reach out to them and work with them but we cannot do it alone.

MR. TYNAN: Ms. Eskin?

MS. ESKIN: Yes. Can someone in the Agency remind me, and I guess the rest of us, of the definition of a small and very small plant and give us some idea again of what percentage of the market, with slaughter or processing, whether it's volume of sales, whatever criteria are used just to understand what kind of impact they have?

MR. PALESANO: This is Bobby Palesano. I will start with part of your question at least. We -- actually the SBA definition of plant size is those

establishments with over 500 employees are large. So the small plants run from less than 500 down to 10, and the very small are less than 10. There are some dollar figures involved in that also, and I don't have those off the top of my head but I'd be glad to get those for you.

MS. ESKIN: Thank you.

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MR. PALESANO: To follow up a little bit more on some of your other concerns, is I think it is quite well known that the large part of the production products of meat and poultry come from large facilities even though the largest percentage small establishments are and very small plants, Sandra, and if that's not enough information, I'll be glad to give you more information. Another thing, Sandra, is a lot of the small and very small are under State inspection programs also.

MR. TYNAN: Mr. Finnegan?

MR. FINNEGAN: Yes. To me a good, a good source for the very small plants is when they come in and do an EIAO review. In doing that, the EIAO reviews will document that they lack supporting

documentation, for example, or a CPP which validates their HACCP plan. They know what they need. Can these EIAO guys, are they allowed to or capable of, to just give supporting documentation if they have it to fit their particular product?

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DR. RAYMOND: That's part of the mindset that I mentioned in my opening remarks. Prior to today, they weren't really encouraged to provide help. They were there to do an inspection, and the mindset will change. Karlease and others are going to spend what, three days next week training trainers who will then train the EIAO officers on how they can help facilitate the plants moving into the 21st Century. They will be there not just to do the EIAO jobs as we know them now, but also to help and assist them with the documentation or what is needed, and that will take sometime to change the mindset like that. That's difficult. And some people will need more help changing their mindsets than others.

DR. MASTERS: Let me clarify though a little bit, Mike. If they're actually in the middle of an assessment and find a specific CCP that's not meeting,

they will not personally go find the CCP information. they will do is they will point them resources that are available because it's the plant that will have to make the particular determination as to what applies to their process because they will not know the ins and outs of their process like the plant's owner as you well know, but they will be in a position to help point them. If they're aware of resources, for example, on E. coli 0157:H7, documents that exist, then they will help point them to those resources, and those are the kind of tools that we want them to help make available so that they can say, I'm aware that there are some documents available that I want to point you towards, and they'll help point them towards those kinds of resources.

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MR. FINNEGAN: Great. I think that will be an excellent resource, I really do.

MR. TYNAN: Mr. Kowalcyk?

MR. KOWALCYK: Yeah, I have one question about obtaining data from the small and very small plants. Has the Agency entertained the idea of possibly sending out not necessarily scientifically

designed, but some type of questionnaire to get a pulse for where the small and very small plants are struggling with complying with the regulations, to try to identify any common themes that certain plants of a certain size can meet the requirements and they're struggling with understanding them or just communication of the rule, as well as getting feedback from them with respect to what support they would want from the Agency as far as helping them learn the rules and regs. Has the Agency considered anything like that?

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DR. KELLY: I can at least start the answer, and that is certainly we have considered it but the Office of Management and Budget has some pretty strict rules on gathering data that make it somewhat difficult for us to actually go straight out with a survey of that kind.

The listening sessions I believe have been very fruitful and that's one reason why we want to continue those, and obviously we want to gather information from anybody who's accessing the services that we have. We try to be as creative as we can

knowing that to some extent our hands are a little bit tied with going out with a survey. Maybe some other people want to add to that.

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DR. MASTERS: Mike, a couple of things, this is Barb Masters, that you'll see in our Strategic Implementation Plan that came out of the International HACCP Alliance meeting that I think get at some of the same issues that you're suggesting is I think you'll see that we suggested that we would look at trends of pathogen testing results which would suggest that they're not meeting pathogen sampling. That would be an indicator that they might have a food safety system that's not designed to allow them to meet pathogen testing results.

suggested that would look Wе we at noncompliance records, specifically in small and very small plants, that might indicate that they're not meeting a particular area of food safety plans. So we said we would look at those trends. We would definitely look at EIAO, our food safety assessments in the small and very small plants, specifically to see if there were any trends coming out in that area

that might allow us to come up with those same kind of determinations. We indicated that we would look at recall data to see if there were any trends. So those are some of the trends that we thought we would analyze for small and very small plants, trying to get at some of the same issues.

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We also pointed out that it would be useful for some of our trade associations, and that's kind of one of the first questions that we were asking of the Subcommittee, is we were hoping that some of the trade particularly associations, and through the International HACCP Alliance, might take the objective of working with the small and very small plants to do some of these kind of surveys for us, and we were hopeful, and that was one of the discussions that surfaced at this meeting, that we might be able to partner with some of the trade associations to do these kind of surveys because it's all very necessary and essential. Thank you. Good question.

MR. TYNAN: Other questions from the group?

There being no other, we'll close this issue out. Thank you, Dr. Kelly. Thank you, Mr. Palesano.

And I'm going to ask our A/V guy if he could help me just a minute. We're going to go into the public portion, and we'll need that microphone back over here.

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What we're going to do now, on our Agenda, we have a public comment portion that allows folks that have been sitting in the audience to make some comments. As I mentioned earlier, we asked that those folks that did want to make a comment sign up on the I have a number of folks that are prepared to make remarks. I'll call your name. I'm not sure what order you signed up but we'll all have an opportunity to get our two cents worth in, and if at the end, there's somebody that omitted to sign up or has had thoughts about that, we'll second give you the opportunity as well.

And the first person I would call that signed up is Dr. Craig Henry. And, Dr. Henry, if you could identify yourself and your organization, that would be great.

DR. HENRY: Absolutely. Thank you, Robert.

Good afternoon. I am Craig Henry. I am the

Senior Vice President and Chief Science Officer for the Science and Regulatory Affairs Group the Food I'm going to read this verbatim Products Association. for the sake of transcription and time but first let the Food Product Association is say that the largest trade association serving both the food and beverage industry in the United States and worldwide, and our laboratories, scientists and professional staff provide technical and regulatory assistance to member companies and represents the food industry on various scientific and public policy issues.

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The FTA commends FSIS and USDA officials for again placing this most important issue on the Agenda of the National Advisory Committee on Meat and Poultry Inspection.

FTA is a founding participant in the Risk-Based Inspection Coalition, which is a broad-based industry coalition composed of 10 associations whose members represent the vast majority of meat and poultry products produced in the United States. The RBI or Risk-Based Inspection Coalition supports risk-based inspection as a means to enhance food safety.

FTA and the Coalition believe that it is proper for FSIS to focus the allocation of its inspection resources based upon risk.

We wish to reiterate our support for Agency involvement of the National Advisory Committee on Meat and Poultry Inspection in this open and transparent process to engage all stakeholders in the pursuit of our mutual goal of enhanced food safety by properly focusing obviously limited inspection resource as well as industry resources on the most significant food safety issues.

As the Committee deliberates on this important issue, we encourage the Agency to pursue an aggressive timeline with key benchmarks to achieve during the tenure of the current administration and reasonably complete Risk-Based Inspection Program that allocate resources based on risk.

The Coalition emphasizes again that to be successful, a risk-based inspection effort must focus on risk-based sampling and testing programs as well as risk-based allocation of the inspection resources.

The Agency continues to make significant

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strides on risk-based policies, to reduce the incidence of listeria monocytogenes in ready-to-eat products and more recently on policies to reduce the occurrence of salmonella in poultry. However, the allocation inspection proper alignment and of equally critical improving food resources is to safety.

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We compliment the Agency for making the Enforcement Investigation Analysis Officers, EIAOs, available to establishments to answer questions and provide input on ways to improve food safety programs.

As stated by the Coalition in earlier comments, measures in success should include the following: reduction in product recalls for food safety reasons, better compliance with food safety requirements along with fewer enforcement actions for significant food safety issues and reduction in foodborne illness outbreaks and sporadic cases to the extent we are able to measure them.

FTA compliments that Agency for pursuing a third party contractor, Resolve, to assist in obtaining input from all stakeholders. We look

forward to working with the contractor and providing stakeholder input on what risk-based inspection will need to be for a successful program to enhance food safety.

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the Committee deliberates on its Δs assignment to provide recommendations to the Agency on measuring establishment risk control, for risk-based inspection, trust that the recommendations we developed by the coalition and forwarded to the Agency December will be taken into consideration. Hopefully by now it is recognized by all that an unweighted use of a number of noncompliance records is not an appropriate measure for the intended purpose. coalition Indeed, the has recommended that modification to the NR system may be required to make it a significant element in any measurement system for food safety risks.

In conclusion, again we commend USDA for continuing the process that once successfully completed will benefit FSIS, industry, and most importantly, the consumer as it focuses everyone's efforts on the areas with the greatest potential for

positive impact on public health. Thank you.

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MR. TYNAN: Thank you, Dr. Henry.

The next individual we had sign up is Mr. Tony Corbo. Tony, if you'd come up and identify yourself and your organization.

MR. CORBO: Tony Corbo from the consumer group, Food and Water Watch.

I have a question as it relates to the FY 2007 Appropriations Bill that's being debated right now on the House floor. The language contained in the Report that the House Appropriations Committee sent floor to the indicates that Committee is going to provide the full amount to cover pay costs, an increase of \$16,625,000 for risk-based management and control of salmonella, an increase of \$2.6 million and information technology support inspection, an increase of \$1.886 million. And it has additional funding for food defense.

Following that, there's this statement. The Committee directs that within the amount provided for food safety and counter terrorism activities, priority should be given to maintaining existing personnel and

operations that are critical to insuring the safety of domestic and imported food rather than funding new functions, grants or agreements.

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Now the total amount that the house is, is proposing is \$853 million. It's \$9 million less than the total budget that you all went up to the Hill with. So that included the \$105 million requested for user fees.

So my question is, is the amount that is in the bill that's on the House floor going to cover both the existing inspection functions and any risk-based inspection activities that you all envision for this coming year?

I think all the deliberations, DR. MASTERS: Barb Masters, that we've been talking about, Tony, at this time and in this Committee as we mentioned at the last meeting, we envision doing within our existing I think we've been talking about gaining resources. doing things within our your input on existing allocations, within our existing resources and we're gaining input from this committee to try to do things more effectively and efficiently within our existing

resources and gaining ideas from this Committee. And if they suggest to us that we need to go further than that, then obviously we'd have to go back in future years to Congress with those future ideas. But ideas we're bringing forth through this Committee and through Resolve at this point, are with existing allocations and with existing resources.

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MR. CORBO: So in light of the duress that you're under right now, you don't, you don't foresee having to curtail any of the activities that you foresee with risk-based inspection?

Again, at this point, we're DR. MASTERS: looking at working with this Committee on ideas that we might be able to implement under existing allocations or existing resources or existing authorities, and if we can gain ideas that we could move forward with sooner rather than later, then that would be great. If they give us ideas that we need to go back to Congress with, then we would do that as well, but we're obviously at the beginning stages of moving forward.

MR. CORBO: Thank you.

MR. TYNAN: Thank you, Mr. Corbo. The next name that we have signed up for comment is Felicia Nestor. Ms. Nestor, if you wanted to say who you're with.

MS. NESTOR: I'm also with Food and Water Watch. I have a couple of comments.

The first that's in my mind, I think that the monetary cutoff for very small plants is \$250,000 if I'm not mistaken.

There's been a lot of concern here about the NRs and how well the NRs are going to reflect the food safety profile of the plant. I just wanted to mention a number of things that I'm aware of since I have so frequent contact with the inspectors in the plants that might impede the possibility of trying to figure out how well these plants are performing.

First of all, the Agency supposedly has a policy of NRs being written on every single noncompliance. I hear from the field that inspectors are frequently dissuaded from writing an NR until they spot a trend, and I recently saw a question about a BSE issue and an answer from the Tech Center saying,

you know, if it's the first time, forget about it, you know, you may want to wait until you see a trend.

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Also if I'm not mistaken, I could be wrong here, but someone can correct me if I'm wrong, prior to HACCP the 7s on the line were able to participate in writing the PDRs. So what the 7s would see would get recorded. Under HACCP, it's my understanding that 7s are no longer allowed to write the NRs. They have to inform a GS-8. So the foreign sector, only the foreign sector can write the NR, and the foreign sector can only write a NR on what or she sees. the foreign sector is not there at the time the kidney comes out of the animal that's 30 plus, and the GS-7 tells the 8 that it happened, there's not going to be Those are only two examples, but I think an NR on it. the Agency should look very closely at other things that impeding the recording of all the are noncompliances.

And another thing is we've got inspector shortages and we have had for years. Yeah, we're not going to have any furloughed, but there are other decreases in inspections. It's my understanding, and

I've heard from several District Offices, that they've been told for years that they're not allowed to hire the number of inspectors that are slotted for that district. There's an unofficial ceiling. So we don't -- we're not fully staffed in any district as far as I know. And I certainly invite any, you know, if I'm wrong, you know, please tell me and I'll go back to my sources and ask where the problem is here.

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Also Barb Masters says we have a hiring I don't know how long that hiring freeze is for, going to go on but we know that there's I mean the workforce is, a lot of them attrition. were hired in the seventies. A lot of them are eligible for retirement. So if we have a good number of inspectors retiring and no one being hired, you know, we're going to have less inspectors.

We know that for years inspectors have been doubled and tripled up in the metro areas. Previously FSIS used to record if they couldn't perform an inspection task, the inspector would record the reason why that task wasn't performed and one of the codes was inspector shortage or vacancy. In other words, I

couldn't get to the plant. So if we have inspectors doubled and tripled up, and they are just barely able to do some of the 03 tasks, the food safety tasks in these plants, you know, we should be able to confirm that from the data that, you know, this plant has not had full inspection and perhaps the reason that there are no NRs in this plant is because it hasn't really been inspected very frequently.

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wonder how you answer the question. Perhaps some of the plants that are performing very well, perhaps they're performing well because they've had a very strong inspection presence for years and what's going to happen if you pull the inspector out of that plant if the plant starts slipping, and then you don't have the inspector present to be able to So I think the Agency should address that determine. There's got to be some kind of a fail-safe way issue. of, you know, spotting the red flags as they go up.

The other thing I wanted to say is I'm happy that a member of the NJC was invited to attend this meeting. I mean I think it's really important that the frontline personnel who put these regulations into

place every day are here to hear what's being discussed and also to provide input about what's happening on the frontline. I'm hopeful that maybe in the next meeting or perhaps within the next few meetings, the NJC can find a place at the table.

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And, one final thing. Barb, you were talking about the three pillars, we've got industry, we've got consumers, we've got the Agency. And as a person who worked with whistleblowers for many years, I found it really invaluable to be able to talk to the frontline inspectors and find out how the regs are actually being enforced, and Ι think that that information has provided valuable input the I think the Agency is not always consumer community. hospitable to inspectors when they, when they want to let the consumers know something that the Agency is going to keep behind the scene. There's an inspector who is going to be negotiating a disciplinary charge within the next few days, and the reason he got the charge was because he forwarded me an e-mail from the Tech Center, and he's up on charges because he sent it Now that doesn't sound to a known consumer advocate.

like an Agency that wants to involve consumers and wants to pursue a transparent process. This was on the non-amenable species, and the e-mail was the one where the Tech Center said, yes, perhaps dogs and cats could be mixed in with amenable products and they wouldn't have to be inspected first. I think that's something that consumers would want to know. And so, you know, I don't think that this inspector -- I think that that e-mail probably was discussed between the Agency and the industry. I think consumers should be in on it as well. Thank you.

MR. TYNAN: Thank you, Ms. Nestor. I should point out, too, just as a reminder to everybody, that in addition to the National Joint Council, we do have representatives from our association of technical and supervisory professionals, and they have a number of people that work in field locations. We have Olga Morales who is sitting in this afternoon for ATFP, and she's a recent graduate of EIAO positioned in Puerto Rico, and we have Patty Bennett who is representing the NAFE, and they certainly have a large number of people that are in plant locations as well. So we're

trying to reach out to all of our employees in that regard but thank you, Ms. Nestor.

The last person I have that signed up is from our international community. It's Dr. Wolf Meyer (ph.), and I hope I pronounced that correctly, sir, with the European Union, and if you could be certain I caught that correctly.

DR. MEYER: Right. Thank you. Thank you very much. My name is Wolf Meyer. I'm with the -- Health and -- of the -- Commission.

I'm amazed to see how similar the questions are here as we are asking in Europe and it's certainly not a coincidence because the food trade goes over a large part, and the food industry is operated on a global scale and so are the risks we have talked in these authorities and worldwide.

So I think there's also scope of reach out to risk management on a global scale and follow this trend of USDA in the United States. And what I mention is in terms of the communication. I think there would be a lot to be gained in terms of risk management, increasing food safety -- risk, if we

1 could resolve issues around data exchange, information 2 exchange between the authorities, on market surveys, if inspection 3 findings, but we can allocate on 4 resource to where maybe somebody else has identified it. Secondly, we should be aware that in changes 7 in inspection procedures, even in -- we have communicate because the importing country might feel if these procedures are followed -- I think

11 changes in inspection procedure, jointly and keeping 12 dialogue and communications.

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said, we have basically the same As Ι questions in Europe as we do here. So I think that it's a true window of opportunity to, to exploit these sort of -- situation and opportunity of change and manage it in the way that we have communication with each other and dialogue.

there's a lot to be gained also if we try to develop

Maybe also there's a lot to learn from other countries like --

That's what I wanted to say. I thank you for this opportunity to be here, and I hope that we

can continue the dialogue.

MR. TYNAN: Thank you, Dr. Meyer.

I'm going to -- for those of you who may have come in a little bit late, perhaps did not have an opportunity to sign up, is there anybody else that would like to make a comment at this point?

DR. RAYMOND: Bob, if no one else is getting up, I'd --

MR. TYNAN: I think Dr. Raymond would like to make a comment.

DR. RAYMOND: Yeah. In response to a couple of the, of the public that have commented, Dr. Meyer and I have had a conversation aside over the lunch break, and I couldn't agree with him more that there's no point in every country taking a look at risk-based inspections by themselves and reinventing the wheel and going through the same -- mistakes and so forth, and we definitely can benefit. I have met with representatives of the European Union just last week, and we had this discussion, and I did tell them at that time, that our proposal that's on the table today is not ready for, you know, prime time, and compared

with other countries yet. That's why we're here today and tomorrow. We'll be meeting with consumers and industry next week, and then our third -- neutral third party, of course, will be taking this out on the road show and looking for more comments. And then at some point in time, we would love to share our plan with other partners including our trading partners, particularly our trading partners, and those who maybe have taken steps already towards risk based.

Barbara and I and others met with the Netherlands two weeks ago with our counterparts and listened to what they're doing in the Netherlands and, you know, there's some interesting things out there and there's some exciting things out there. So, Dr. Meyer, I appreciate your comments and they're duly noted, and we will follow up on this when this product is a little bit more ready for prime time, number one.

Number two, with the concerns of international trade, I don't think we're talking about anything here today that will influence a negatively or positively international trade. We're just trying to make the food products that we have in this country

safer. We're not talking about less than date inspection. That's been brought up by comments by the public today. No one has mentioned less than date inspections on the Food Safety Inspection Service Team. That's not where we're going. We're talking about resources currently available to us and moving them around to where they're most effective and to where we can continue to improve our, our product.

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And then lastly, the comment about not being open and transparent. I don't know the particular incident that was mentioned. if I did, I Even wouldn't comment on it in public but with Dr. Masters and I assuming our roles last summer, if you did not know, I'll let everybody here at the table know that we have instituted monthly meetings with consumers and monthly meetings with industry. Barbara and her team meet with consumers for an hour. Office Food Safety meets with consumers for an hour, both on the same A lot of those are by telephone calls so not day. everybody has to fly into D.C. We've done that.

We've also made Barbara's and mine and Dr. Mann's and Mr. Quick's schedules. They're on the

web. You can see our daily appointment schedules. done that. We've tried to be We've open I think, I think we're making an honest transparent. efforts. We'll make mistakes along the way. have to, you know, I apologize for those ahead of time, but Ι think for the most part, this Administration has been much more open than in the That's why the third party's coming on board to past. help continue that process thanks to your recommendations at your last meeting and, in fact, recommendations followed at the last meeting on who we should even entertain, you know, to submit applications from, and we appreciate that input from the Subcommittee of this Committee.

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So we'll continue to try to be as open and transparent. When you catch us not being, just let us know.

DR. MASTERS: This is Barb Masters. Just a couple of things before we move to Subcommittee. I know several of you in the audience as well as the Committee have asked for copies of all of the PowerPoints. We will get the Committee copies of the

PowerPoints to try to work with this afternoon. We will make all of the PowerPoints available on our website so that those of you in the audience can have access to those after the meeting. So we will make those available to you speaking of transparency, so they will be available.

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And also I just wanted to comment Felicia brought it up on the hiring freeze. I want to make it very clear, I used the term that we have a hiring freeze on non-frontline positions probably should make that very clear so understands what non-frontline positions are. got a hiring freeze on those positions outside of our So here in Headquarters, District Offices, so field. anything that is not related directly to inspection. So I want to be very clear on that so everybody walks away from the table on the same page. So I want to clarify that, so everybody leaves the table at the same place. So I appreciate your indulgence in that.

So I look forward to -- I'll try to walk back and forth between the two Subcommittees. Before Robert turns you loose, I appreciate all the good

input that we got during the public comment period, and we'll look forward to everyone's work this afternoon. So thank you.

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MR. TYNAN: Before we go, I would direct you back to Tab 3 in your notebook, and I think on the first page, we have the Subcommittee breakdowns, and I think the Subcommittee that will be looking Ms. Eskin establishment risk control, has been designated the Chairperson of that group, and they will be meeting in Room 0161 in the South Building, and we'll get somebody to guide you there. I can see you're relieved already.

And then the second Subcommittee will be chaired by Dr. Joe Harris, and he'll be looking at strategic implementation plan for enhancing outreach to small and very small plants.

I have on the clock on the wall a little bit after 2:00. I would suggest perhaps this be sort of a rolling break, and if the Chairpersons are both amenable, we would suspect that everybody should be in the conference room at 2:30 or later. No later than. We can -- excuse me a second while I sidebar. Okay.

1	2:30 is it.
2	So for Subcommittee 2, it's Room 1160, and
3	again if you need some guidance in getting there, Joe
4	will make sure you have someone to help you get down
5	there.
6	And with that, with that I move we adjourn
7	for the session and allow the Subcommittees to do
8	their work.
9	Are there any last minute questions or
10	issues before we go?
11	There being none, thank you very much for
12	spending the time with us this morning.
13	(Whereupon, at 2:10 p.m., the meeting was
14	concluded.)
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1	CERTIFICATE
2	This is to certify that the attached proceedings
3	in the matter of:
4	NATIONAL ADVISORY COMMITTEE ON
5	MEAT AND POULTRY INSPECTION
6	Washington, D.C.
7	May 23, 2006
8	were held as herein appears, and that this is the
9	original transcription thereof for the files of the
10	United States Department of Agriculture, Food Safety
11	and Inspection Service.
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