VIDEOCONFERENCE MEETING

STATE OF CALIFORNIA

ENVIRONMENTAL PROTECTION AGENCY

AIR RESOURCES BOARD

SCIENTIFIC REVIEW PANEL

ON TOXIC AIR CONTAMINANTS

CALIFORNIA ENVIRONMENTAL PROTECTION AGENCY

1001 I STREET

SACRAMENTO, CALIFORNIA

THURSDAY, JULY 9, 2020 9:08 A.M.

JAMES F. PETERS, CSR CERTIFIED SHORTHAND REPORTER LICENSE NUMBER 10063

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APPEARANCES CONTINUED
REPRESENTING THE DEPARTMENT OF PESTICIDE REGULATION:
Edgar Vidrio, Program Manager, Environmental Monitoring Branch

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1. Welcome and Introductions

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- 2. Developing provisional health values to support the AB 2588 Air Toxics "Hot Spots" Program.

The California Air Resources Board (CARB) and local air districts are responsible for implementation of the Air Toxics "Hot Spots" Act (Health and Safety Code section 44300 et seq.; AB2588, Connelly). The goals of this program are to compile information on toxics emissions; identify facilities having potential for localized impacts; evaluate their health risks; notify nearby residents about significant risks; and ultimately reduce the risks below a health protective threshold.

For the past year, the Panel has provided feedback and approved preliminary findings on CARB's proposed updates to the list of air toxics that must be reported by stationary sources. The SRP-reviewed air toxics list is an important component of the Hot Spots program, but many of the proposed and existing air toxics chemicals on the list do not have OEHHAapproved cancer potencies or noncancer reference exposure levels. In order to contextualize the emissions from chemicals that have not been assigned an approved health value, staff propose to assign provisional values to these chemicals. Given the SRP's role in reviewing health values for specific chemicals and chemical classes, staff from the Office of Environmental Health Hazard Assessment (OEHHA) will present an overview and CARB staff will give a brief presentation on their proposed approach for assigning provisional health values, with the objective of seeking Panel Members' feedback on ways to improve the process.

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	The AB 617 Consultation Group includes individuals representing environmental justice organizations, air districts, industry, academia, public health organizations, and local government. Its meetings provide an opportunity to discuss various aspects of Community Air Protection Program implementation. The Panel's representative in the group will provide an update on the February 26, 2020, AB 617 Consultation Group meeting. OEHHA staff will provide a brief overview of SRP's potential role with respect to AB 617.	
	Part II: Update from Department of Pesticide Regulation regarding 1,3-Dichlorpropene mitigation pilot studies	58
	Staff will provide the Panel with a brief overview of proposed mitigation pilot studies for 1,3-Dichloropropene (Telone) in the AB 617 community of Shafter, as an example of a topic for which SRP might provide future input.	
	Following these updates, the Panel will have an opportunity to discuss potential ways in which they might support air quality and health protection at the community level.	
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PROCEEDINGS

CHAIRPERSON ANASTASIO: Okay. Good morning everyone and welcome to the meeting of the Scientific Review Panel. My name is Cort Anastasio. I'm Chair of the Panel and I'm a professor at UC Davis. I'd like to first start by welcoming everyone on the broadcast, including our Panel. So Panelists, if we could just go around, if you could very briefly introduce yourself, that would be great.

PANEL MEMBER HAMMOND: This is Katharine Hammond from University of California, Berkeley. I'm a Professor of Environmental Health Sciences at the School of Public Health.

PANEL MEMBER RITZ: This is Beate Ritz. I'm a

Professor or Epidemiology and Environmental Health at UCLA

School of Public Health.

PANEL MEMBER MILLER: This is --

PANEL MEMBER KLEINMAN: This is Mike Kleinman.

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PANEL MEMBER MILLER: Go head, Mike.

PANEL MEMBER KLEINMAN: Okay. Mike Kleinman, University of California, Irvine. I'm an inhalation toxicologist.

PANEL MEMBER MILLER: Okay. This is Lisa Miller.

I'm a professor at the UC Davis School of Veterinary

Medicine.

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PANEL MEMBER BESARATINIA: Good morning. This is Ahmad Besaratinia. I'm an associate professor of preventive medicine from University of Southern California.

PANEL MEMBER LANDOLPH: Hi. This is Joe
Landolph. I'm associate professor of molecular
microbiology and immunology pathology and molecular
pharmacology and toxicology, and a member of the USC
Norris Comprehensive Cancer Center. My specialty is
chemically-induced cell transformation and carcinogenesis.

CHAIRPERSON ANASTASIO: Any chance that Paul is on the line. Paul?

Yeah. Okay. Guess not. Well, thank you, panelists. Appreciate that.

First to let everyone know who's attending, there are five handouts available through GoToWebinar. You'll see there's a -- on the menu on the right, the -- at least on my screen, the next to last component on the menu are the handouts.

A couple of administrative items before we begin. Everyone will be muted except for the panelists and the presenter. We will be accepting oral comments at the end of the meeting. This is only on the AB 617 items, which will be our last agenda item, so we're not accepting

comments on the other items.

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If you would like Spanish interpretation for the final item, which will be a presentation from Edgar Vidrio from the Department of Pesticide Regulation on the plan for Shafter, if you would like that, please let us know when we get to that item when we have a translator prepared to provide that service.

Let's see, Panel members, I've already mentioned please mute yourself. If you want to be called on during this discussion, you know, either wave your hand or put in an exclamation point in the chat.

Yeah. Okay. So let's get to the agenda items. So we have three major items today. The first will be a discussion of the developing provisional health values for CARB's proposed updates to the AB 2588 air toxics list. So we'll have a presentation about items for that.

The second piece will be an informational update on the February 26th, 2020 AB 617 Consultation Group meeting, which Mike Kleinman attended, so he'll be discussing that.

And actually before that, John Faust will give a brief reminder of ways in which ARB thinks that the Panel might be able to assist with AB 617 items.

And then finally, we'll end with Edgar Vidrio's presentation about 1,3-Dichloropropene pilot mitigation

measures proposed for Shafter. And that's the item that we'll be taking public comment on.

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Okay. So without further ado, let's begin.

First major agenda item, developing provisional health values to support the AB 2588 air toxics hot spots program. So, Panel, you'll remember that for the past year, although it seems longer, CARB has been working with the SRP to update us about the list of proposed air toxics that must be reported by stationary sources.

Now, as we've discussed in the past, you know, this -- we're talking about adding hundreds of items to the Appendix A. And many of these don't have approved health values. And so because it's such a huge number, rather than going through individually as we've been normally doing, the idea is we have some kind of interim provisional health value. So we're going to start today with an overview of this risk evaluation process from the Environmental Health Hazard Assessment OEHHA. And then CARB is going to give a presentation about their -- their current thinking on how to develop provisional health values. And hopefully, the Panel will have some suggestions about how to work with that.

Okay. So John Faust is going to start us, give us an introduction. So I turn it over to John.

(Thereupon an overhead presentation was

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presented as follows.)
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             DR. FAUST: Okay. Thank you. So we need to --
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             MS. KLEIN:
                         John, I just handed you control of
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    your screen.
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             DR. FAUST: Okay. And is that showing full
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    screen?
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             MS. KLEIN:
                         It is showing not in presentation
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   mode, but we can see the slides. And if need be, I have
    your slides and I can do them for you.
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             DR. FAUST: Okay. For some reason, it's showing
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    up in presentation mode on my second screen, but not my
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   main screen.
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             MS. KLEIN: On your -- do you see a Button
13
    dropdown that says display and that says swap presentation
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    and view, by chance?
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16
             DR. FAUST: On the dropdown?
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             MS. KLEIN: Yeah on your -- in the view where you
    can modify things not on the viewer screen. You can swap
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    view. But if you want, I can just pull up your slides
    quickly.
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             DR. FAUST: Okay. Maybe that would be better.
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    Thank you.
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             MS. KLEIN:
                         Yes. Okay. So just say next slide
    and -- when you're ready for me and I'll switch slides.
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DR. FAUST: Okay. Great. All right. So just on

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the first slide. So good morning. I'm John Faust. I'm Chief of the Community -- Community and Environmental Epidemiology Research Branch in OEHHA, Office of Environmental Health Hazard Assessment

So as part of an introduction to today's first agenda item, I'm going to make some comments on potential ways to address the issue of unassessed chemicals. So I'm going to give a little background on the nature of the problem, some of the ways we can address it, some things to think about as we move forward, and also some next steps.

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DR. FAUST: So this slide describes some of the reasons and ways that the issue of unassessed chemicals comes up. As the Panel is well aware, establishing health guidance values, such as reference exposure levels or unit risk factors by traditional approaches can be very time and resource intensive. And because of this, we and other entities have only established values for a fraction of chemicals in use or, for example, on the AB 2588 hot spots inventory.

So we encounter chemicals without values in a number of different situations. Environmental monitoring data, for example, can reveal the presence of chemicals in

air, water, soil, or food. Community air monitoring, for example, can identify pollutants that people are likely exposed to at home.

Other specific studies like sampling and analysis of synthetic turf, an OEHHA program, can also identify sets of chemicals that may produce potential exposures. Emissions inventories like the air toxics hot spots list leads to information about emissions of chemicals from facilities. And other databases also provide information about how and where chemicals are used in commerce or industry, such as disclosures about fracking chemicals or ingredient reporting.

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DR. FAUST: So one way to address the gap in health guidance values is to establish provisional values. And here, I describe this as a mechanism to provide information in a more expedited manner on the potential for health risks from exposure to a given chemical.

So while we adopt numerical values for traditional RELs or unit risk factors, adopting qualitative characterizations of toxicity, such as categories, is also a possibility.

So since this approach carries a greater level of uncertainty than traditional approaches, it's important

that we match the decision context that we're using to be -- and the level of confidence. And it's also possible but in some context, the level of uncertainty in developing or adopting a provisional value is going to be too high to be acceptable.

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DR. FAUST: So this slide describes some of the key options in establishing provisional health guidance for unassessed chemicals. They fall broadly into two categories. One, using work from other authorities when it exists, and two, using alternative approaches when no values are available from other authorities or they're not considered reliable.

And here, I've identified two options when there -- when there are other established values. One, option is to adopt their values. For example, recent values from U.S. EPA's IRIS Program could be considered for adoption. Alternatively, we can adapt other authorities' health guidance values. For example, we can apply uncertainty factors or adjustments consistent with our established California values to a point of departure from a key study identified in another agency's assessment.

In the next slide, I'll describe a little bit

more about the things that we need to think about in using that type of existing work.

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So if there are no values from other authorities, there's another range of options, and these include producing expedited values in-house. This can be done if there's a set of readily available studies that clearly establish a point of departure for toxicity like a no adverse effect level, and then uncertainty factors can be applied to that, for example.

On the other hand, there is also so-called read-cross approaches, where we apply knowledge about chemicals that are more well studied to other potential analogs that are -- that are data poor or less well studied. And to be an analog, different types of bioactivity can be considered, such as structural, metabolic, or toxicological similarities. And I'll describe a little bit more about this shortly. And then there may be other possible approaches as well.

So if we could move to the next slide.

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DR. FAUST: So this slide shows some of the things that should be considered in either adopting or adapting the values from other entities. First is to understand how consistent the methodologies used to establish the value are with California's health risk

assessment guidelines. For example, were they developed with the purpose of predicting sensitive populations or, for example, were they assessments to support occupational standards.

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Second is the methodology consistent with ours, for example, with respect to the use of uncertainty factors or how the dose response is evaluated and was the value derived from an appropriate route of exposure.

Another consideration is whether the assessment is comprehensive, that, is whether all health endpoints were assessed versus, for example, a health guidance value based on developmental only or other more limited set of endpoints.

And finally, we want to also consider whether the assessment was peer reviewed, whether there was public -- it was publicly reviewed and is currently available and well documented, as well as how recently the data were evaluated to produce the assessment.

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DR. FAUST: So for alternative approaches, I'll briefly describe the two I mentioned earlier, developing expedited values in-house, and considering read-across -- so called read-across approaches. For expedited values to be established in-house, we generally need to work from a

reliable data set that can be evaluated relatively quickly. The dose response information would also need to be straightforward allowing a clear identification of the point of departure, for example. So this would be, you know, work that we would do in-house only.

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And on the other hand, read-across is a method of filling a data gap where a chemical with existing data is used to make a prediction for a similar chemical. I drew this definition from a presentation by Grace Patlewicz of the U.S. EPA who discussed this topic at an OEHHA symposium last spring.

So an example of a workflow broadly would be to include identifying the decision context that the read-across information is to be used, identifying chemical analogs that are under consideration, identifying the critical data gaps between the more and less well studied chemicals, evaluating analogs for read-across opportunities, and then characterizing the uncertainties before completing the assessment.

This general approach can be adapted to different levels of confidence, completeness, and speed. And there are other possible approaches as well that can be considered. For example, thresholds of toxicological concern is a concept originally proposed by the US FDA for food additives in which human exposure thresholds are

established, below which appreciable risk to human health is very unlikely. And these would be based on specific types of toxicities, such as presumed carcinogenicity based upon structural inspirations.

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DR. FAUST: -- I have a few next steps relating to ongoing work at OEHHA. So in April of 2019, OEHHA held a symposium entitled, *Understanding and Applying*Read-Across for Human Health Risk Assessment. And as a follow-up, we continue to evaluate different existing read-across platforms for their potential usefulness, but for single chemical assessments and for groups of chemicals, such as perfluoroalkyl substances, or PFAS.

In collaboration with academic partners, OEHHA is also supporting the development of methods using in vitro studies and in silico molecular docking data.

And then regarding the SRP, OEHHA also wants to bring a more robust discussion in the areas that I've introduced, including evaluating existing non-California health guidance values, as well as applying alternative approaches. And, of course, we'll continue to coordinate with the Air Resources Board on their efforts to establish provisional values for the Hot Spots Program that you'll hear about shortly.

So I hope that background has been helpful. And at this point, I can either take questions or we can move forward with the -- the Air Resources Board's presentation.

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CHAIRPERSON ANASTASIO: Yeah. This is Cort, I think in the interests of time, let's move forward with the ARB presentation. John, thank you very much. That was very interesting and I look forward to hearing more about that at a future meeting. And Panelists, if you have questions on John's material, we'll have a little bit of time after the next presentation.

So next, Melissa Traverso of CARB's Air Quality Planning and Science Division is going to give us presentation about the proposed approach for assigning provisional health values to the chemicals that are lacking values currently.

All right. Take it away Melissa.

(Thereupon an overhead presentation was presented as follows.)

AIR POLLUTION SPECIALIST TRAVERSO: Thanks you, Cort, for the introduction. Good morning, members of the panel. Today we are back before you to begin the discussion of a new topic related to the evaluation and implemen -- implementation steps of the AB 2588 Air Toxics Hot Spots Program.

Today's topic is the development of the provisional health guidance, which will be non-regulatory in nature, and which is needed to support the data evaluation steps of the Hot Spots Program. We would like to thank our colleagues from OEHHA for their help in setting the stage for this discussion.

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AIR POLLUTION SPECIALIST TRAVERSO: Today's presentation will be provide a brief background on the current status of the AB 2588 Air Toxics Hot Spots Program updates and how this new topic, the concept of non-regulatory provisional health guidance, is anticipated to fit into the program. We will discuss the need and purpose for provisional health guidance in the context of AB 2588, as well as some prior history of the program related to default health values.

Since we are still at the beginning of the discussions regarding this topic, we will be providing some framing questions as a way to guide today's discussion on how to best to address the more formal process just described by our OEHHA colleagues to meet the needs of the Hot Spots Program.

Lastly, we will discuss the proposed process and anticipated timeline for developing the provisional health

guidance, including how it relates to the overall timing of our AB 2588 emission inventory guidelines regulation update, which is expected to go to the Board in November 2020.

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AIR POLLUTION SPECIALIST TRAVERSO: Overall, the goals of the program are to collect air toxics emissions data and make it available to the public, identify facilities that may have localized impacts, assess the risks to public health, and notify nearby residents about significant risks, and reduce these risks to levels that are more health protective.

Our previous meetings with the panel have focused on compiling the list of chemicals for the emission inventory and criteria guidelines regulation, which we'll also refer to as the EICG regulation.

Compiling and updating the chemical list is part of the regulatory process. We appreciate the panel's valuable review and input in that process of updating the substance list. To date, we are proposing to add roughly 700 substances to our chemical list for which emissions must be quantified by the facilities subject to the AB 2588 program.

Updating the chemical list allows us to address

new and emerging chemicals since our last major update, which was 1996, and to include chemicals that have been recently recognized by international, national, State, and other experts as having potential health concerns. It is important to collect information on where, how, and how much these chemicals are being emitted from California facilities and operations. This is a necessary first step in understanding the relative potential for public health impacts.

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At this stage, most of the new substances we are proposing to add into Appendix A list do not yet have an official OEHHA-approved health value. In fact, learning about the nature and extent of emissions is one of the things that later helps OEHHA with their health value prioritization process.

In the absence of OEHHA approved values, it can be very helpful to utilize any and all available basic information regarding the relative toxicity of new substances to help support the hot spots process. For example, in the early days of the AB 2588 Hot Spots Program, when there were not yet official cancer or noncancer health values for some important substances, the California Air Pollution Control Officers Association, or CAPCOA, prepared default health values for a number of chemicals that had some available data. These default

health values were found to be useful to facilities, as well as districts, as a means to screen what types of emitting processes and chemicals were likely to be of either minimal concern for the facility or alert them to potential instances warranting more careful consideration.

With the Panel's guidance, we would like to propose to develop non-regulatory provisional health guidance for as many of the new substances as feasible where OEHHA acute or chronic REL and/or cancer potency factors are not yet available.

To clarify, the provisional health guidance we are proposing would not be explicitly listed in the EICG regulation nor would it be used for official health risk assessment purposes. Instead, it would be provided outside of the regulation as useful, technical information for facilities, air districts, and others. Its purpose is to help guide reporting and preliminary evaluations, which could potentially lead to voluntary emission reductions buy facilities.

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AIR POLLUTION SPECIALIST TRAVERSO: Our purpose for developing provisional health guidance aligns with the goals of the Hot Spots Program to identify facilities that have localized impacts, assess the risks to public health, and reduce these risks to levels that are health

protective.

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Provisional health guidance will continue to help CARB, OEHHA, and facilities better understand the relatively importance of processes and emissions at the facility level. Facilities can use this information to make more informed decisions regarding voluntary reductions with assistance from CARB and districts. Additionally, OEHHA can allocate their resources for development of new health values based on the quantity of emissions being reported for each new substance under the AB 2588 program.

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AIR POLLUTION SPECIALIST TRAVERSO: We'd like to provide a little more context about the concept of the provisional health guidance and also contrast how it might be developed and used compared to the other program elements of the AB 2588 Program, such as compiling the chemical list itself and assigning the values we call the reporting degree of accuracy values, which specifies how accurately the emissions of each listed chemical must be reported by a facility to ensure the emission data will be useful.

First, we utilize information about the nature of potential health concerns as one of the factors we are

considering in identifying substances for inclusion on the chemical list.

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The chemical list is part of the official EICG regulation and is developed as part of that rulemaking process, which we anticipate taking to our Board in late 2020.

Second, most AB 2588 facilities are required to report their emissions on a four-year cycle. And that reporting is subject to a reporting degree of accuracy that we assign to every reportable substance and chem -- in the chemical list and it's included as part of the EICG rulemaking process.

This value is meant to serve as a practical value associated with emission reporting and it communicates the facility how much and how precisely a particular substance needs to be reported. For example, a highly potent metal like hexavalent chromium must be reported out to several decimal places in pounds per year, in order to have the reported emissions be useful enough to evaluate the possible public health implications for that facility. By contrast, the emissions of benzene are sufficiently accurate when reported to the nearest two pounds per year, and the emissions of toluene to the nearest 200 pounds per year.

These reporting degree accuracy values are

typically guided by some combined consideration of the basic toxicity information about the chemicals and the levels and types of their usage.

By contrast, the idea of provisional health guidance, we are proposing today would not be a part of the EICG regulation or its rulemaking process and timeline. The provisional health guidance would be intended to be available as technical supporting information likely posted on our website to help inform and support the later implementation stages of the AB 2588 Program consistent with how OEHHA-approved risk factors are currently documented.

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AIR POLLUTION SPECIALIST TRAVERSO: We would like to emphasize our plan to work closely with OEHHA in the development of AB 2588 specific health guidance. I will now go into detail on some framing questions around our approach. And some of these topics John Faust touched upon in his presentation.

The first question is what data sources should be consulted for health guidance? We are proposing a hierarchical approach relying on available data and other information, and we would value your input on any additional sources we should look into further.

Currently, we're giving priority to health related databases, such as Proposition 65, which can provide an indication for provisional health guidance. For example, the cancer potency of styrene could help us estimate levels that might be of concern, even though there isn't a formal AB 2588 adopted potency yet.

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Additional health values available under U.S. EPA's Integrated Risk Information System, or IRIS, quantitatively characterizes health hazards of chemicals found in the environment and can cover a chemical, group of related chemicals, or a complex mixture. IRIS would help provide a more quantitative approach. Whereas, for example, PubChem or more related to HSDB, may provide qualitative information to support the development of provisional health guidance.

Another source of information would be workplace limit values such as Cal/OSHA's Permissible Exposure Levels, or PELs, and the American Conference of Governmental Industrial Hygienists, ACGIH, threshold limit values, or TLVs. We would propose several adjustments to these workplace values to be more protective of vulnerable populations and lifetime exposures instead of accounting for only healthy adult workers during an eight-hour work shift in a career duration.

Third in the hierarchy of available data sources

is the structure activity modeling or relationships. The idea here is that looking at similarities in chemical structures could help provide information on how to treat a general class or group of similar compounds and provide a benchmark type of approach for health guidance purposes.

Lastly, for substances on our list where there is not enough supporting data from the other sources previously mentioned, we are considering looking into limit or guidance values from other media, for example, water programs. We are aware that these values would have to be adjusted to consider the potential for water-borne pollutants to become airborne and would like the Panel's guidance and thoughts on using this type of source.

The second question we'd like to pose to the Panel is what is the most appropriate approach to establish health guidance for a functional group class and its members. As you may recall, we are proposing to add three categories of substances defined by broad chemical functional groups to the chemical list. We would like to get the Panel's thoughts on possible approaches. For example, could we consider using ranges of toxicity values or treating subsets of the functional group class together?

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alr Pollution Specialist Traverso: The third question in this framework is what types of adjustments should be considered? I mentioned some of these adjustments when discussing the various data sources available in the previous slide. However, we would like to get the Panel's perspective on any additional criteria that should be considered and how much weight to be applied to factors, such as structure activity relationships, the quantity and quality of available health information, and the severity of health effects, and relative toxicity, and other factors. Examples of other factors we might consider could be persistence, bioaccumulation, and multi-pathway effects.

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AIR POLLUTION SPECIALIST TRAVERSO: I will now provide an overview of the process and anticipated timeline. Subsequent to today's discussion, we plan to engage with our OEHHA colleagues to closely coordinate the development of a methodology for reviewing and developing provisional health guidance. We propose to return in early 2021 to update the Panel on the proposed methodology and request your feedback on any modifications that might be necessary. We anticipate that draft health guidance will be released for public review by mid-2022 and

available for use by mid-2023.

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AIR POLLUTION SPECIALIST TRAVERSO: It should be slide nine. That concludes today's presentation. Thank you, Panel members for your time. Anyone listening in on this call can feel free to contact Gabe, Greg, or myself using the contact information provided here, if you have any additional follow-up questions.

PANEL MEMBER BLANC: Sorry to interrupt. Paul Blanc here. I had some difficult, but I'm on now.

CHAIRPERSON ANASTASIO: Great. Welcome, Paul. Glad you could join us.

PANEL MEMBER BLANC: I apologize.

CHAIRPERSON ANASTASIO: Oh, it's no problem.

Thank you very much, Melissa.

Questions from the Panel for Melissa or for John on his presentation, prior to Melissa's?

All right, Melissa --

Go ahead.

PANEL MEMBER KLEINMAN: Oh, sorry. I just wanted to get some clarification on Melissa brought up some framing questions. And are these things that we want to be addressing on this call or is that for the later discussion?

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CHAIRPERSON ANASTASIO: Yeah. I think we'd like to discuss them today, to the extent that we can. And then I imagine that some of them will come up in the future as well.
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So Anne, could you go to the framing questions slide?

MS. KLEIN: Yes.

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AIR POLLUTION SPECIALIST TRAVERSO: It starts on slide six, Anne.

MS. KLEIN: Okay. Do you see that?

AIR POLLUTION SPECIALIST TRAVERSO: I see slide nine.

MS. KLEIN: Weird. Yeah, there's some -- plenty of stuff going on with the slide transitions for some reason today.

16 CHAIRPERSON ANASTASIO: Okay. Now, we can see 17 it.

MS. KLEIN: Okay.

CHAIRPERSON ANASTASIO: Okay. So data sources, most appropriate approach for functional group classes. And then, Anne, could you go to the next slide for the last question? Okay. Types of adjustments to be considered.

Okay. Well, let me -- let me start with a question that's actually not related to the framing

questions. Melissa, I had a question. You talked about provisional values being used -- I think used by 2023. Can you explain what you mean by that? Are you suggesting that you will have provisional values by then or that this process will be producing provisional values by then?

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AIR POLLUTION SPECIALIST TRAVERSO: The process will be producing provisional health values by mid-2022 and available for use by mid-2023. So there's kind of like a phase-in approach to first kind of reporting the chemical substances that are new on our list and then using those provisional health values for reporting purposes.

CHAIRPERSON ANASTASIO: I see. Thank you. And do you think you're going to have all of the compounds with provisional values by then?

AIR POLLUTION SPECIALIST TRAVERSO: That is the plan.

CHAIRPERSON ANASTASIO: Wow. That would be great.

PANEL MEMBER RITZ: So I -- this is Beate.

CHAIRPERSON ANASTASIO: Go ahead, Beate.

PANEL MEMBER RITZ: So I actually don't understand. If you're not -- if you're not asking facilities to do anything, or is that a wrong understanding, what are these provisional health values

giving you?

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AIR POLLUTION SPECIALIST TRAVERSO: So the provisional health values will provide facilities with an idea for how toxic that substance might be. A lot of these substance on our chemical list are, as you know, very new. So facilities may not know they are reporting those substances, or if they are reporting them, they'll have a better idea of how much of a concern they should give weight to those substances, if that makes sense. And, Beth, if you want to chime in, feel free.

PANEL MEMBER RITZ: Yeah. I really don't get that, because, you know, if we don't know much -- if they're new, we won't know much. And then to give them an uncertainty about a possible health effect will leave it open to doing nothing.

AQPSD STAFF AIR POLLUTION SPECIALIST SCHWEHR:

This is Beth Schwehr. Let me jump in for just a
moment here. Can you hear me okay?

CHAIRPERSON ANASTASIO: Yes.

AQPSD STAFF AIR POLLUTION SPECIALIST SCHWEHR:

Okay. So one of the things that happened in the very early part of the Hot Spots Program is that as facilities began to look at their overall operations and processes with an eye to what chemicals they had and what they emitted, they began to learn more about what their

operations -- which of their operations were posing more risk, for example, to the public than others.

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But to really get at it, it's not just the amount of emissions. They have to weight the amount of emissions by the relative toxicity of those. For example a very small number hex chrome doesn't look like much. But, of course, if you realize it's very potent, then you realize it might actually be your risk driving operation.

And so by having even a semi-quantitative number available for some of these newer chemicals, as they began to evaluate their whole suite of operations, they can start to say, oh, which of my operations are actually causing more risk to the public than others, right? They can begin to identify what is their risk-driving operations?

And what happened in the early days of the Hot Spots Program, is that -- that understanding resulted in facilities very quickly making a lot of voluntary emission reductions and process changes. So they identified that maybe a solvent they were using was their risk-driving process and they decided to substitute a less toxic solvent, for example, or they identified that there was a process that had a lot of fugitive releases and that that was creating concerns.

So then they would make efforts to tighten up

that process and have less fugitive emissions. But to do that, you really do have to have at least a relative sense of how toxic some of the chemicals are to be able to put that in context along with their respective usage.

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So that's what we're hoping to do here is to have kind of some guidance. Even if it's not a formal perfectly fleshed out health value, it can often be helpful to a facility to understand whether this is a chemical that's going to end up being really important for their overall impacts to the public or a chemical that's just going to be petty minimal for their impacts.

CHAIRPERSON ANASTASIO: So related to that -PANEL MEMBER KLEINMAN: Hi, this is Mike. To
what extent is this information going to be communicated
to the communities that are potentially impacted? I see
this as -- you know, this is one of the factors that I'm
going to be talking about later in terms of uncertainty in
the AB 617 communities about other toxic chemicals that
were not on their initial watchlist.

So I see this as a -- you know, another way that pressure can be brought from the community level back up, but it's got to be done very carefully and correctly. So will there be any attempt to put this in context -- these emission data in context with potential community exposures?

AQPSD STAFF AIR POLLUTION SPECIALIST SCHWEHR:

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This is Beth again. So when the health values become -- or health guidance type of approach become available, we would anticipate posting something for these chemicals on say a website. It wouldn't be regulatory, but it would be something that people could use, and it would be similar to the kinds of ways that adopted health values can be looked at now. So someone could look at the emission inventory that was reported for a facility. There are some formulas, if you will, that the California Air Pollution Control Officers Association, CAPCOA, have come up with to give kind of a score approach to -- to tell whether a chemical and that amount of emissions will likely be of high priority or not high priority. So someone -- members from the public could do that same kind of thing.

What we anticipate is that it might be used by air districts, for example, when they prioritize their facilities as well. And, of course, by the facilities themselves, they can do those same calculations to assess what's important and what's not.

But, yeah, these things would be -- provisional information would potentially be posted on a website made available to anyone who wanted to do that kind of analysis.

PANEL MEMBER KLEINMAN: It might be very useful. I'm sorry. It might be very useful to have a brief presentation of this to the consultation group that -- if and when they have another meeting, so that they're aware that this is available and that they should be looking at that website.

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AQPSD ASSISTANT DIVISION CHIEF EDWARDS: Yeah. Mike, thanks for the comment. This is Dave Edwards. Yeah, we can definitely circle back internally to talk to your AB 617 counterparts to see if we can get that on the -- the next agenda.

PANEL MEMBER KLEINMAN: Great. Thanks, Dave. CHAIRPERSON ANASTASIO: Thank you, Beth.

PANEL MEMBER RITZ: Yeah. This is Beate again.

I have a bit of a concern doing this chemical by chemical.

I heard something about groups, but more or less in terms of assessing the relative toxicity of several chemicals in a group. But what about mixtures? I mean, we might have ten different neurotoxic agents spewing out in emissions from the same facility, and, you know, ranking them. And giving relative toxicity does not help when you want to assess the overall toxicity of this kind of mixture. Is there any -- is there any sense that mixtures will be evaluated?

AQPSD STAFF AIR POLLUTION SPECIALIST SCHWEHR:

There are a couple of approaches for mixtures. Sometimes things are actually used in a mixture and we would apply a profile to understand what is the composition, the relative proportions in that mixture.

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Alternatively, I'm thinking maybe what you're getting at, is that when a facility as a whole is evaluated, all of these would be looked at together. And these -- this CAPCOA prioritization guidance document, which I referred to a moment ago, does look at that, and it does say what -- you can add up various things and look at the facility as a whole. So you could look at the overall -- say, look at all the carcinogens and look at that as a -- as a whole.

And then for the -- the non-cancer effects, as a -- at a screening level, sometimes those are looked at across all endpoints. But if you want to do a more, you know, correct assessment, you can look within endpoints, for example. But, yes, the CAPCOA document is intended to be looked at as a facility overall, you know, impact.

PANEL MEMBER RITZ: Great. Thank you.

 $\label{eq:chairperson} \mbox{CHAIRPERSON ANASTASIO:} \quad \mbox{Can we go back one slide,} \\ \mbox{Anne.}$

CHAIRPERSON ANASTASIO: Panelists, any thoughts about these framing questions, additional data sources that should be consulted for health guidance? I know we

had a lot of discussion of data sources when we were discussing the AB 2588 list last year. Are there additional things they should be looking at in terms of health guidance? Anybody have any thoughts?

Beate.

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PANEL MEMBER RITZ: Definitely. I think all the IARC documents should be looked at for cancer -- for carcinogens. And I'm sure there are other European documents that would give you some insight as well.

CHAIRPERSON ANASTASIO: Great. Thank you. Other suggestions on health document or health guidance data sources?

PANEL MEMBER KLEINMAN: Yeah. On the workplace data, the more recent ACGIH TLV, you know, workups, they are really calling out things that are either respiratory sensitizers or skin sensitizers. And it would be very -- yeah, looking at the notations that go with the guidance that they provide for workplace data could be extremely important, because sensitizers, after an initial exposure, can have a much more exaggerated effect in continuing exposures. And so some of that kind of information could help shade what you'd put down as a potential risk factor. So, you know, certainly look at that.

And the other thing in the newer ACGIH data, they're taking into account potential developmental

effects. And even though those might not be put in as the basis for setting a guidance level, they are important in terms of our being aware of chemicals that do have potential developmental effects. And then carry that forward to the general population with a little more sensitivity.

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PANEL MEMBER RITZ: There's also the National -- CHAIRPERSON ANASTASIO: Beate.

PANEL MEMBER RITZ: There's also the National Toxicology Program, NIEHS, that puts out documents. You should look at those. And I would even suggest to look at the latest meta-analyses or summary analyses in PubMed and see, you know, what -- what the scientific community thinks about certain agents and groups, in terms of health effects.

CHAIRPERSON ANASTASIO: Great. Thank you.

Anyone else have other -- Joe.

PANEL MEMBER LANDOLPH: Yeah. I would also look at the EPA documents, as long as they reflect current scientific understanding, and all the authoritative bodies that we gather the data from, and are used by OEHHA that go onto the Proposition 65 list.

CHAIRPERSON ANASTASIO: Right. Thank you, Joe.

I do think the last bullet -- sorry, I was going to say the last sub-bullet under this main bullet,

threshold values from other media, I do think that's a good idea, for example water. Obviously, the root of exposure would be different, but in terms of setting relative values, that could be very helpful.

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Was somebody else going to say something?

PANEL MEMBER BESARATINIA: Correct.

CHAIRPERSON ANASTASIO: Yes, Ahmad, go ahead.

Yeah. I was wondering there were -- there was a slide regarding the timelines and some milestones were mentioned. I'm wondering with the current situation with COVID, there would be probably some challenges with regard to field work and data selection. I am wondering if this was taken into account when they were setting up all these timelines or not?

CHAIRPERSON ANASTASIO: Melissa or Beth, you want to speak to that.

AQPSD ASSISTANT DIVISION CHIEF EDWARDS: This is Dave. I could do that.

CHAIRPERSON ANASTASIO: Or Dave.

AQPSD ASSISTANT DIVISION CHIEF EDWARDS: So, yeah, we definitely are -- are cognizant of what's going on right now. And as sort of Beth and Melissa alluded to in some of the responses, we are looking at a much more phased-in approach than maybe we were even three to four months ago and that will be rolling out in the next month

or so with our -- with our hot spots materials for our regulatory update.

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But we do understand that there are going to be some limitations moving forward. And, yes, the timeline that we do have out to 2023 right now, that is sort of our sort of ideal path, but we also are aware that that may get pushed out as things -- as new things develop, so...

CHAIRPERSON ANASTASIO: Thank you, Dave.

PANEL MEMBER BESARATINIA: Thank you.

CHAIRPERSON ANASTASIO: Okay. Let's move on to the second framing question then. People have thoughts about this question, the most appropriate approach to establish health guidance for functional group class and its members. I mean, it seems that one part of that would be gathering data that we know about whatever members of that class are. And then you look at the variability within the class to try to get some sense of whether that's narrow or very large. And if it's large, what seems to be influencing the toxicity of individual members?

But I don't know if people have any experience with this or any thoughts on this?

PANEL MEMBER KLEINMAN: Well, this is a really new sort of issue in terms of the kind of chemicals that

are coming out like the PFAS and PFOAs. There are literally hundreds of them. And it seems like when they set regulations for one, somebody tweaks a molecule, and now you've got a whole new chemical.

CHAIRPERSON ANASTASIO: Right.

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PANEL MEMBER KLEINMAN: So I think it's really important to set up exactly what you said, Cort, understanding what are the driving factors for initiating toxicity and focusing more on that than on a specific chemical, so that we could use some sort of hazard index, or, you know, group approach to a class of or family of chemicals, because we're not going to, you know, be able to keep pace the new chemicals being generated.

CHAIRPERSON ANASTASIO: Yeah.

PANEL MEMBER RITZ: Yeah. And I'm really afraid that we have to worry about the tendency to go after one -- one criterion, such as the lengths of the molecule or how many fluorides, or fluoro compounds, or bromides or -- brome is part of the chemical and making it more or less persistent, right, because persistence is one of the criteria that we are worried about, but then the toxicity is another one. And it doesn't mean that when they are less persistent, they're also less toxic.

So I think these are the kind of things that probably chemists who know more about this have to think

about, and toxicologists, how to approach that.

CHAIRPERSON ANASTASIO: Thank you, Beate.

Joe.

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PANEL MEMBER LANDOLPH: Yeah. As Mike mentioned, there are just a plethora of certain things like polycyclic aromatic hydrocarbons, and then there's TCDD and its congeners, and dibenzofurans, et cetera. They've set up this toxic equivalency factor usage. And that seems to work okay, but, of course, they keep discovering more and more, and there's a large number of them. But the TEQ approach seems to have worked out okay.

Maybe we'll use something like that until we get better guidance for dibenzofurans, and TCDD and its congeners.

CHAIRPERSON ANASTASIO: Great. Thank you, Joe.

Any other thoughts on the second framing

17 question?

Okay. Anne, could you go to the next slide for last framing question?

So types of adjustments to be considered.

Melissa, can you clarify this, types of adjustments to be considered under what circumstances?

AIR POLLUTION SPECIALIST TRAVERSO: Yeah. So I mentioned some of these adjustments when I was discussing the various data sources available in the previous slide,

but we'd like to get the Panel's perspective on any additional criteria that should be considered, and like how the weight -- how weight should -- how much weight should be applied to factors, such as like the structure activity relationships, the quantity and quality of available health information, and the severity of the health effects and relative toxicity.

And then like what you mentioned previously, the persistence bio -- bioaccumulation and multi-pathway effects. So, you know, it sounds like persistence and toxicity are two very separate things. So trying to figure out how much weight to give one factor over another for these provisional health guidance.

CHAIRPERSON ANASTASIO: Okay. Thank you, Melissa.

Thoughts. Okay. Joe. Sorry. Hod on, Joe.

I -- Joe, it says you're self-muted -- well, and muted by an organizer. Okay. Joe, you're self-muted. Joe, you need to unmute yourself before you talk.

Still muted, Joe.

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Oh. Now. Okay. Hold on one second. Okay. Go ahead, Joe.

PANEL MEMBER LANDOLPH: Okay. Yeah. Certain chemicals we have an enormous amount of mechanistic data on how they act for carcinogenesis. And I'm thinking of

polycyclic aromatic hydrocarbons. And many physical chemists have made calculations of the stability of the carbonium ions generated. So they can use those calculations to predict which ones would make the best carbonium ions to attack DNA covalent and covalent adducts, which would make mutations, et cetera. So that data can be used to a certain extent.

And to the extent we have data like that for other compounds, like aromatic amines, where they have carbonium ions, which resonate with nitrenium ions, you can make those calculations or simple binding to DNA, et cetera. You can make predictions as to which out of a whole group of chemicals would be carcinogens. But some of the other chemicals, of course, we don't have such great data on.

CHAIRPERSON ANASTASIO: Yeah. Thank you, Joe.

Yeah. So definitely there's some modeling approaches that may give insights into toxicity of chemicals where we don't currently have values. And, yeah, that would be very -- a more rapid process to estimate some health values.

Anyone have any other thoughts about this last framing question?

Beate.

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PANEL MEMBER RITZ: Yeah. I think you need to

consider how much you want to use the modinomics technologies to infer health data or health information, because we will not -- or we shouldn't be waiting for long-term health outcomes that may occur in 10 years, 20 years, 30 years. We might need to consider whether we are okay with looking at a chemical and finding that it disturbs a certain metabolic pathway, or changes a certain methylation patterns in tissue. And we know that these metabolic pathways or tissue disturbances have been related to health effects downstream and how much that could be used as a warning sign for later health effects, because otherwise we're going to be waiting for 20 or 30 years for health information, at least for the newer compounds to exactly accumulate.

CHAIRPERSON ANASTASIO: Yeah. Thank you, Beate. That's a good point. Right. With the old mix approaches, there's a wealth of data. And perhaps some of that can be used to understand relative health values.

Yeah.

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Any other --

AIR POLLUTION SPECIALIST TRAVERSO: Yeah, these are really great recommendations. Thank you.

CHAIRPERSON ANASTASIO: Yeah, you're welcome Melissa.

Any final comments from the Panel?

Mike, you're muted, I believe.

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PANEL MEMBER KLEINMAN: You're right. I'm just going to riff off of what Beate had said a little while ago about groups of chemicals that have the same target and also compounds that will come up under structure activity modeling. I think it would be very useful to have a formalized and -- you know, I know we have, but to kind of consider that in this framework to look at something like a hazard index as opposed to individual things for the individual level compounds. When you have things that target the same, you know, biological system or are structurally similar to chemicals that target a given system, there should be a way to add those into a framework with what we know about all the other things that are out there.

And, you know, in some cases, we use the hazard index where you look at the relative toxicity levels and have a scheme for adding them up, so that the net result gives you some idea of the toxicity of the -- and potential health affects of the overall group.

The other thing I did want to bring up, this approach sounds like it's facility by facility. And there should be, as an overview, are there other facilities that could impact a given area a given community that are producing some of the same chemicals? Because you may

have one facility producing almost nothing, you know, very small amounts. But if you have a lot of those, the total exposure to the community doesn't really care whether it came from smoke stack A or smoke stack B.

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CHAIRPERSON ANASTASIO: Yeah. That's a good point. This will be our final comment. Melissa, or Dave, or Beth, you want to talk about this cumulative exposure question.

AQPSD STAFF AIR POLLUTION SPECIALIST SCHWEHR:

Hi. This is Beth. Yes, I really appreciate that comment. We definitely want to be cognizant of cumulative exposures. We're doing things elsewhere within the emission inventory criteria and guidelines regulation to strengthen that as a criteria for districts to consider, when they're looking at impacts under the Hot Spots Program. And, of course, it's a big component of the AB 617 Program. So, yes, we're -- we're very aware of that. We appreciate that comment.

CHAIRPERSON ANASTASIO: Great. Thank you, Beth.

All right. Well, that will conclude the

provisional health value because discussion. I look

forward to talking about it at a future meeting. Thank

you very much, Melissa. Thank you, Beth. Thank you,

Dave.

We're going to move on now to our second major

agenda item, which is -- first, we'll have a report from Mike Kleinman on the AB 617 Consultation Group meeting that he recently attended. So these consultation group meetings are -- provide an opportunity to discuss implementation of various aspects of the Community Air Protection Program.

And we're going to actually start with Dr. John Faust from OEHHA who's going to give us a brief overview to remind us of three suggested areas where the Panel can hopefully assist in AB 617 efforts. And then we'll have Mike Kleinman give his presentation.

So, John, it's over to you.

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(Thereupon an overhead presentation was presented as follows.)

DR. FAUST: All right. Thank you. Yeah. So if you could just turn to the second slide here.

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DR. FAUST: So in the -- in the March 2019 SRP meeting, we presented three different concepts that we thought could potentially serve as a role that the SRP could play in moving -- moving things related to SB -- AB 617 forward. And the three are described on this slide.

So the first of them health risk values for contaminants in AB 617 communities. So this is around providing guidance on the identification of emerging

contaminants of concern, including recommending priority substances for OEHHA to develop or update health risk values. And these may include contaminants identified in communities through air monitoring or emissions inventories.

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So this -- this, you know, reflects a little bit back on the previous discussion, since we had introduced both the idea of identifying priority substances in communities to move, you know, through a very similar process as the -- as, for example, REL development, as well as thinking about expedited methods for establishing health guidance to be able to provide information. So this -- this has some -- some commonalities with -- with the work on provisional values.

The second topic area addressing cumulative exposures in communities. So here, this is about providing guidance on assessing potential health risks from combined exposure to multiple contaminants, especially where individual pollutant exposures are below current standards, for example, reference exposure levels or ambient air quality standards. And then risk assessments should consider sensitive populations and vulnerabilities as well. And then the third area is tracking community health benefits through indicators, providing input on identifying the kinds of health

benefits from reductions in localized air pollution that are most amenable to measurement. So those are -- those are the three topic areas.

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CHAIRPERSON ANASTASIO: Great. Thank you, John. Just a note to the Panel and also agency staff, you know, our next meeting in October, right now the agenda is a little light, so if we wanted to discuss one or more of these topics at that meeting, I think that would be great if -- to get us started. So I'll just leave that to the agency to discuss amongst themselves and we can talk about it offline.

So, Panel, these are three of the ideas that the agencies have come up with where we might be able to assist. So just keep this always in the back of your mind. And now, we're going to segue into Mike Kleinman's presentation from the February 26th meeting of the AB 617 Consultation Group.

Mike, the floor is yours.

(Thereupon an overhead presentation was

presented as follows.)

PANEL MEMBER KLEINMAN: Okay. Thank you. If we can have the next slide, please.

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PANEL MEMBER KLEINMAN: So this is the agenda of the consultation group meeting. And the first part of the

meeting dealt with a survey run by UC Davis looking at how the communities felt about basically giving a report card to the process so far. And that took up, you know, a fair amount.

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The -- the next part is a discussion of the blueprint that was provided. This was a joint effort to develop a blueprint for how the AB 617 communities would be working, what the guidelines were, what the time limits were, and what the communities felt about where we were in that process.

The last part, the Governor's office provided their overview from the Office of Planning and Research, and where we were going. And this was done just prior to a discussion of the actual AB 617 budget, which it turns out the Governor's office is reducing some of the support for that.

So if I could have the next slide.

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PANEL MEMBER KLEINMAN: From the survey, there have been several really important pieces of progress.

One is that the community air monitoring plans have been moving forward very well in some communities. And so deploying various inexpensive monitors, so that more granular data could be obtained, then you would get from the ordinary, you know, ARB and district monitoring

central site data have been implemented and are beginning to produce useful data.

In some -- you know, some areas not all of the air districts are created equal. And their relationships with the communities are also not equivalent in some cases. So there are some very strong groups that have really focused on using the AB 617 information to eventually come up with community emissions reduction plans, which can take into account land use, use of pesticides, and things like that.

And they -- there is -- there has been a lot better engagement working with the cities, counties, and the Department of Pesticide research. And so there was some real improvement over the initial beginnings of this.

And to some extent, projects and -- and plans have been allowing communities to have various levels of engagement in the process.

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PANEL MEMBER KLEINMAN: So there are still Challenges. The aggressive timeline, a lot of the communities felt that because they had a specific timeline to start implementing emission reduction programs, they didn't have enough time to get warmed up to -- especially if they weren't technically adept when they started up,

there's a fairly steep learning curve to doing effective environmental monitoring. And then even more so to being able to understand what that data is saying. So there was a feeling of pressure to move forward with emission reductions and/or plans for that before they felt comfortable with some of the data acquisition side of it.

There was also, you know, some discussion that in some cases, there was unsatisfaction with the goals of the process where it was more a cut-down engagement, where the districts took the lead and the citizens were invited to sit at the table, but they really didn't have a whole lot of power in terms of allocating budgets. So they -- that was their perception.

Just this kind of partially stems from tensions between the districts, and the industries, and the communities that have been long-standing. And some of the districts, as I said, were less involved with the community than others. But overall, one of the things that they felt was a challenge was they -- the communities felt that there should be a more proactive role for the Air Resources Board, and also for the SRP. And so input from us, OEHHA, and CARB, will -- you know, is really encouraged.

Next slide, please.

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PANEL MEMBER KLEINMAN: So this is -- the blue panel on the left is coming out of the blueprint. So these are things that the steps that went through to set up the different communities that were chosen for the first wave of communities that were done. You can see that there are ten communities that were selected and they had leadership programs set up. They established their meeting structures and processes.

And if you look at the right, you can see that each of the communities had different models for how this was led. So, for example, West Oakland has a very strong co-leadership relationship with the Bay Area Air Quality Management District and they are working very closely with them, the same as in Richmond. And in Richmond, they brought in an outside facilitator to help work with communications.

Some of the communities, the process is more district driven and some of them are less -- the communities have less direct input in terms of how the money gets spent, but some of it has led to, you know, some actual programs and discussions as in Shafter, which we'll be hearing about later.

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PANEL MEMBER KLEINMAN: So this is my take on the

highlights of the meeting. Most of it dealt with process. Very little dealt with what I consider, you know, the substantive discussion of actual results. But two things came -- you know, really floated to the top. One was the Bay Area and the West Oakland community had a source apportionment project, where they drew upon years of data that AQMD had been collecting and working with, and had done a lot of preliminary work, but they helped explain this data to the community representatives who could then bring these results back to the community. So they had a real understanding of what the sources of their exposures actually were in the area.

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The other thing that was very innovative was the district also funded and implemented a street level monitoring program, where instrumented vehicles drove every street and freeway in the West Oakland area and collected air sampling data all along that track, integrated that data, and came up with very extensive exposure-related information, which again they could provide back and explain to the community. And now the community has this information and they have a much better feel for what they're being exposed to, where it comes from, which then leads to the plan to how do we cut this out, and reduce exposures?

The other success that was brought up was the

South Coast Air Quality Management District. They took on the program of evaluating the various sensors that are being used and deployed in the different communities. And they've set up programs to compare these sensors, their reliability, their accuracy, calibrations, and things like that. And so this is an ongoing effort the AQMD in the South Coast District has been working on that standardizing, the monitoring protocols, and also standardizing how the data are reported, so that this can be used to really provide useful information for communities and for the regulatory community as well.

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PANEL MEMBER KLEINMAN: So the areas that they raised concerns were a lot of the communities don't have the training or expertise to really interpret and integrate the information that's coming out. And so they really want the community partners that are working with the districts to have access to better training and understanding, so that they can more effectively communicate with the rest of the people in their areas.

One of the things that several communities brought out was that there were very incomplete data on pesticide exposures that they could use. And there was some jurisdictional problems in that the agricultural

usage of some of these things is not under ARB control, or -- but under the agricultural division control.

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And so the -- there were apparent conflicts on which pesticides were allowed, what amounts were allowed, and things like that. And so some of that I think this is an administrative issue that could be worked on and improved.

And then monitoring at the community level for pesticides is not viewed as really practical. So it's really important I think going back to our discussion of what the sources of exposures are, I think, including pesticides is part of our emissions inventory somehow would be, you know, important.

And then air toxics, which are not part of the general air quality monitoring programs, the data on these things are really very localized and very hard to integrate over an entire community.

There are also sources, such as certain storage areas that are not always included in emission inventories. For example, things that are unloaded from the ports, the ports have data on emission inventories. But once they are transferred and stored in off-site facilities, the ports really don't have control over that and they don't show up in their emission inventories. And so there was concern that -- about the completeness of the

inventories.

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And there was also the feeling that the SRP could provide more context for toxic air contaminants and pesticides, and really provide better information to the community on the potential risks and on the effectiveness -- the effectiveness of proposed mitigation plans.

I think that is it. Is there another slide? --000--

PANEL MEMBER KLEINMAN: Yeah. Okay. So a summary. Thank you.

The blueprint was written two years ago. And it's a living document, but it's really needing to be updated, especially with respects to what we've learned and what has now come out as best practices. And I think the potential role of source apportionment to identify targets have been applied in the West Oakland area, and that is an approach that could be used more, you know, effectively in many other communities, if the data are available. But it takes a buildup of looking at long-term data to really understand it.

There was definitely a call for a better assessment of potential impacts of TACs. And then budget issues were raised which is a real concern, because part of the funding, which is used for incentives, which

basically are incentives to have facilities reduce their emissions, that was cut 25 percent in the Governor's budget and the implementation activity funds were cut by 50 percent.

However, all of this predated COVID. And so at this point, things will -- you know, are in a changing landscape. And so there are uncertainties to say the least about how this is going to be prioritized for the future.

And that is it. Thank you.

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CHAIRPERSON ANASTASIO: Thank you, Mike. We appreciate the update. Before we move on to our final agenda item, I have one announcement and then I want to take a break. So the announcement is for our last agenda item, which will be the presentation from the Department of Pesticide Regulation on potential mitigation plans for 1,3-D usage in Shafter, if anyone would like Spanish translation, please put a question in the question box or put something in the chat. We have a translator who is able to provide that service for Spanish translation. So please identify yourself and then we will do that.

Claudia, could you please translate roughly what I just said into Spanish.

(Translated into Spanish.)

CHAIRPERSON ANASTASIO: Thank you, Claudia.

What I'd like to do now is take a five-minute break. I know our intrepid reporter, Jim, is on the line frantically pressing keys for the last hour and a half. So let's give him a break and this will give us a chance to run to the restroom, if we would like. It's 10:33 now, so let's reassemble in five minutes at 10:38.

(Off record: 10:33 a.m.)

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(Thereupon a recess was taken.)

(On record: 10:38 a.m.)

CHAIRPERSON ANASTASIO: Okay. Welcome back, everyone. First for Spanish translation, do we have any requests?

Patrick, maybe you could help me on that. I'm looking at the chat. I don't see anything about Spanish translation.

MR. GAFFNEY: I don't. Anne, maybe put up that, and then we could maybe have someone translate it to let people know to raise their hand if they would like Spanish translation using their control panel.

(Translated into Spanish.)

CHAIRPERSON ANASTASIO: Thank you, Claudia. So Lori, I'm not sure how to proceed in terms of Spanish translation. If we don't get any requests, should we go ahead and do it anyway or should we just provide the English version.

PANEL LIAISON MIYASATO: If there are no requests, I think we can go ahead and just do it English.

CHAIRPERSON ANASTASIO: Okay. So waits another minute.

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In the mean time, I would like to give a shout out to Jim Behrmann who is apparently joining us. Jim, it's nice to have with you us, although in a different capacity, and we hope you're enjoying retirement.

PANEL MEMBER KLEINMAN: This is Mike. Because this thing is being recorded and might be viewed later, you may need to do the Spanish translation.

CHAIRPERSON ANASTASIO: That's a good point.

Okay. So how about we will proceed with the Spanish translation.

Yeah. So Claudia, if you could translate Edgar's presentation, that would be great.

Which brings us to our last major agenda item.

This is going to be the overview from the Department of

Pesticide regulation regarding 1,3-dichloropropene, 1,3-D

or Telone, pilot mitigation measures in Shafter.

The Panel has been asked to accept or and written comment on this item, given the State's interest in transparency and community participation in AB 617. The upcoming staff presentation is informational only. We're not being asked to vote on this matter or take any action

today. So I just want to make that clear.

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If you do submit a comment today, we will make sure that agency staff -- appropriate agency staff gets the comment, so that they can follow up on that. We, the Panel, won't be doing any follow up.

And, of course, there are multiple points along the way where stakeholders and community members can provide input on AB 617 matters as they move through the agency's public process.

Okay. With that understanding, I'd like to turn it over to Edgar Vidrio of the Department of Pesticide Regulation for this 1,3-D presentation.

Thank you, Edgar

(Thereupon an overhead presentation was presented as follows.)

MS. KLEIN: Okay. This is Anne. I just want to interject. Edgar, I can show your slides or would like me to give you control?

MR. VIDRIO: Yeah. My sharing or allowing me to share my screen, I have some transitions that will be -- instead of saying, you know, next bullet, it will be easier on my end.

MS. KLEIN: Okay. I just handed you control.

CHAIRPERSON ANASTASIO: Yeah. And then Edgar,
just a reminder to pause for the translation service.

1 MR. VIDRIO: Will do.

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Let me make sure I share the right screen. All right -- oh, that's the wrong one.

Okay. Can everybody see that?

CHAIRPERSON ANASTASIO: Yes.

MS. KLEIN: Yes.

MR. VIDRIO: All right. So -- sorry, technical difficulties here. Are people still able to see the presentation screen?

MS. KLEIN: Yes.

MR. VIDRIO: Yes. Okay. So. All right. Okay. So for the Spanish translation before I begin, what I'm going to do is go over either a bullet or a few bullets that have to do with one another or read a paragraph that I have on the slide and then pause.

Does that work?

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THE INTERPRETER: Yes. That will work. If you keep the paragraph, then the interpreter can do a side translation into Spanish.

Thank you.

MR. VIDRIO: Okay. That would be great. Okay. So hello, everybody. My name is Edgar Vidrio. And today I'll be providing an update on the Department of Pesticide Regulations 1,3-dichloropropene, acute mitigation. I will

also give an overview of the upcoming 1,3-D mitigation pilot program.

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MR. VIDRIO: Specifically. Sorry go ahead.

Okay. For this presentation, I'll be providing essentially a very brief background on the pesticide 1,3-dichloropropene, which is also known as 1,3-D or Telone. I will be going over its uses, past mitigation, and the need for additional control measures.

THE INTERPRETER: I'm sorry. That was a little longer. Do you mind repeating it, point by point?

MR. VIDRIO: Not a -- not a problem. So for this presentation, I'll be providing a brief background on the pesticide 1,3-dichloropropene, which is also known as 1,3-D or Telone include -- so I'll be going over the uses, past mitigation, and need for additional control measures. I will also cover DPR's approach to mitigating short-term acute risks to fumigants, such as 1,3-D.

THE INTERPRETER: I'm sorry, 1,3-D you said?

MR. VIDRIO: 1,3-dichloropropene, but for the rest of the presentation, I'll be referring to it as 1,3-D.

I will then go over DPR's upcoming mitigation pilot program. And finally, I will go over the connection for the link between the proposed mitigation measures, the

pilot program, and the AB 617 community of Shafter.

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MR. VIDRIO: Okay. So let's start with a brief background on 1,3-dichloropropene or 1,3-D. 1,3-D is a pre-plant soil fumigant that is applied to control nematodes, insects, and disease organisms. As with any fumigant, the pesticide is directly applied into the soil, either via subsurface injection or through chemigation.

In California, 1,3-D is used in over 60 different crops, including fruit, and nut, trees and row crops.

Most of California's 1,3-D use is centralized to the San Joaquin Valley and central coast regions.

1,3-D is listed under the Title 3 of the California Code of Regulations section 6400 as a restricted material, which requires a permit from the local county agricultural commissioner to apply. And on top of that, it also requires that any application of this pesticide is only conducted by a certified pesticide applicator.

DPR has placed mitigation measures that control exposures to 1,3-D since 1995. These control measures include what we call a township use cap. This basically sets a use limit per California township with the focus of reducing long-term cancer risk. And for this, a township is defined as a six-by-six square mile area per the public

land survey system.

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So DPR, in part due to recent elevated 1,3-D ambient air concentrations that were measured at two monitoring sites, is proposing to add additional requirements that focus on the reduction of short-term acute risks to children and infants, which are the most sensitive populations.

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MR. VIDRIO: In general, DPR has several options available to mitigate unacceptable air exposures to pesticides.

These include buffer zone distance or setbacks.

THE INTERPRETER: I'm sorry, you said setback?

MR. VIDRIO: Yes.

Also, there's changes to the application method and the use of -- establishing use limits. Specifically, for 1,3-D in order to address a few exposures, DPR is considering the following options:

First is increase the distance between the application and the sensitive receptors, and this can be in the form of a buffer zone or setback. The next one is to limit the amount of 1,3-D that can be applied. Another possibility is to require the use of lower emitting application methods. And lastly is to increase the soil moisture requirement of the treated field.

So DPR combined the use of computer models with vast air monitoring data to develop these appropriate mitigation options. Specifically, DPR used two models. One of them is HYDRUS. And HYDRUS is a industry standard solute transport model that has been used by DPR to simulate fumigant transport and volatilization. The use of HYDRUS is very important, because it allows us to estimate the flux for emissions that result from 1,3-D either during and following an application.

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The second model that we used is called AERFUM, which stands for Air Exposure and Risk model for Fumigants.

THE INTERPRETER: I'm sorry. That one went very fast. Would you mind repeating?

MR. VIDRIO: Yeah. No. No worries. Air Exposure and Risk model for Fumigants.

AERFUM was actually developed by DPR in order to simulate fumigant air dispersion. So AERFUM uses a commonly used model that's called AERMOD, to -- we use that as a simulation engine. And then AERFUM allows the pre- and post-processing functions of fumigant applicants.

AERFUM is -- again, it's specifically designed for regulatory purposes and it actually takes into account the California pesticide use data, weather information, and GIS layers.

So DPR's views of both HYDRUS and AERFUM models went through an extensive peer-review process that was coordinated by the University of California in 2019.

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THE INTERPRETER: I'm sorry, could you repeat the second segment?

MR. VIDRIO: Sure. It went though an intensive external peer-review process coordinated by the University of California in 2019.

So DPR -- via the use of these two models, DPR results -- modeling results indicate that 1,3-D applications -- it basically indicates that for 1,3-D applications in the -- with the use of what is called a totally impermeable film, or TIF, tarp that the actual use of the tarp can suppress emissions to below health protective values.

And for those of you guys that are unfamiliar,
TIF tarps are commonly used for some fumigant
applications. And they're used to suppress the fumigant
in the ground while also maintaining certain soil moisture
levels. However, while the use of DPR approved TIF tarps
have a long track history of being able to reduce
fumigants, gas emissions, the use of TIF tarps may not be
feasible for all crops grown in the San Joaquin Valley.

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MR. VIDRIO: And there's actually several reasons

why the use of TIF tarps may not be feasible for all crops.

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The first one is availability. There's not enough TIF tarp manufacturers or supply to cover the whole 1,3-D industry.

The second one is waste or recyclability.

Currently, there are no commercially available means of recycling the amount of TIF tarps that will be produced, if the whole 1,3-D industry were to use these.

And the last one will be price. The cost of using TIF tarps per acre treated may only be -- make sense for high-grossing crops. And, for example, this will include something like strawberries. However, lower cash crops, like sweet potatoes for example among others, may not be able to recoup their costs of using TIF tarps. And this is because the use of TIF tarps can be up to \$1,500 per acre.

THE INTERPRETER: I'm sorry, could you repeat the price again?

MR. VIDRIO: It's an average of \$1,500 per acre.

Therefore, DPR's approach, instead of focusing exclusively on the use of TIF tarps, we decided to take a -- basically, a different approach and look at alternative options that could potentially reduce the 1,3-D emissions to levels that would be comparable to

using these tarps.

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And essentially, currently both the United States Environmental Protection Agency and DPR offer a 60 percent buffer zone reduction credit when applicators use TIF tarps in certain fumigant applications.

On top of that, by using the models that I mentioned on the previous slide, DPR modeling results show that 60 percent emissions equates to a minimum a 60 percent buffer zone reduction that will be needed for most field sizes and most application rates.

Therefore, for this mitigation effort, DPR aims to reduce 1,3-D emissions by at least 60 percent as compared to an equivalent untarped application, which in this case is the 18-inch deep untarped method. And the reason why we chose this as a base method is because it is -- this application methods is the predominantly used method to apply 1,3-D in California. And with around 80 to 90 percent of all applications that occur in the San Joaquin Valley taking place via the 18-inch deep untarped method.

By doing this, or by taking this approach, it will allow the flexibility for -- to reduce the fumigant emissions by the required amount, which in this case is 60 percent. But at the same time, they will allow the grower the flexibility to choose an option that best fits their

needs.

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DPR has identified several options that will reduce 1,3-D emissions by at least 60 percent compared to the base standard fumigation. We posted this -- a document on our website, which is linked on the bottom, that actually lists all 13 plus different options that provide this -- these comparable results.

So every proposed application method in that document may be feasible in all situations, but each method -- so sorry, not every application method listed in there may be feasible for all applications. However, each method should be feasible in a limited number of situations.

And what I mean by that essentially to kind of break that down is that not all listed options meet the 60 percent emission reduction goal individually. However, they could either be used in combination with one another or they can be additional restrictions placed on those applications in order to meet the 60 percent goals.

And these additional restrictions that could be placed on these options can be anything from either limiting the size of the application area either -- or limit the application rate that can be used, or you could actually impose a specific setback distance.

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MR. VIDRIO: So considering that 1,3-D is extensively used in California with an average usage of around 12 million pounds annually, also considering the fact that there are no commercially-scaled alternative options currently available where people can transition from 1,3-D to something else, and also some of the mitigation measures that we're proposing can be very costly, and lastly, the fact that we -- proposed mitigation measures may not be feasible and may not necessarily achieve the desired emission reductions for all cases, therefore, before we roll -- we roll these proposed mitigation measures into a statewide regulation, DPR is planning on rolling out a, what we call, a pilot program to test these mitigation measures. And we hope to do it in a regional setting prior to statewide implementation.

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We hope to begin this program in the fall.

Obviously, that, you know, COVID-19 may have a saying on this. By doing this pilot program, it will allow the DPR to evaluate the proposed mitigation measures on a regional scale, again before we go statewide. And for the pilot, while we have chosen select high-use regions that are located near air monitoring locations in the communities of Delhi, Parlier, and Shafter, the applicators in these specific areas will have the option of using 1,3-D in the

new application methods or with the new restrictions in order to reduce air emissions by 60 percent compared to untarped methods.

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The lower emitting methods that we hope to try on will rely on either having the fumigant injected at deeper depths or we could also increase the amount of pre-application soil moisture. And, of course, we always have the ability to have the -- either complete or partial TIF tarping. And ultimately, if none of these are an option, the growers may be able to combine with other options, or have additional restrictions placed.

And the additional restrictions are application rates or application area, or like I mentioned before, increased distance between the application and a sensitive site.

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MR. VIDRIO: Specially, the pilot program will have the following three objectives.

The first one is the collection and evaluation of air monitoring data that result from the use of these new application methods to validate computer modeling estimates. The next objective will be to evaluate the feasibility of the -- all the proposed mitigation options that we are putting forth. And lastly, it will be to evaluate the effectiveness of these mitigations at

reducing the emissions of the 1,3-D to the levels that we have -- we hope to get.

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On top of the new emission reduction options that we're going to put forth, there's going to be some enhanced air monitoring efforts. So we will continue our weekly ambient air monitoring sampling that we already conduct in the towns of Delhi, Parlier, and Shafter.

THE INTERPRETER: And I'm sorry, would you mind repeating the areas again?

MR. VIDRIO: Yeah, no worries. The towns are Delhi, Parlier, and Shafter.

On top of these, the ambient air monitoring, we also want to conduct application site monitoring studies that are targeted to measure and validate emissions from the new methods.

So so far, I've talked about DPR's regulatory development process, what we can do with 1,3-D specifically, and what we hope to achieve in order to address the acute exposures of 1,3-D this fall.

THE INTERPRETER: I'm sorry, what was the last segment? Can you repeat it again?

MR. VIDRIO: Yeah. What we hope to -- basically to address the acute exposures of 1,3-D this fall. The pilot program will help us evaluate the effectiveness and feasibility of these proposals at a regional scale again

before we scale it up to a statewide regulation. But so far, I haven't really addressed AB 617 or how AB 617 fits into these efforts.

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MR. VIDRIO: So although pesticides were not specifically included in AB 617, DPR has been collaborating with CARB and various air districts in an advisory role to address pesticide exposures.

For example, in the community of Shafter, DPR attended multiple steering committee meetings and made various presentations on pesticide-related topics. In that same community, there were several pesticide related amendments that were included in Shafter's approved community emission reduction program, or CERP.

THE INTERPRETER: I'm sorry, can you repeat the second segment?

MR. VIDRIO. Sure. That's the Shafter's community emission reduction program, or CERP. These commitments included our department's commitment to continue pesticide air monitoring activities in the Shafter community, as well as DPR's commitment to develop statewide regulations to reduce exposures to 1,3-D in ambient air.

Therefore, ensuring that the Shafter AB 617 community boundary was actually included in this pilot

program was definitely a priority for the Department. And as you can see in the map that you have here in this slide, the pilot area here is highlighted with a dark border. And the actual border of the community of AB 617 in Shafter is included in the lower right box.

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The image on the right of that is what we call the windrows and it's showing the wind direction. So you can see that the use on the northern and the northeast -- or northwest part of Shafter is blowing into the community, so that we capture in this pilot program.

Therefore, through the pilot program, DPR will be able to better assess the efficacy and the practicality essentially of these new proposed mitigation measures in how effective they are at reducing 1,3-D emissions.

And finally, the pilot program will help DPR meet those commitments that were included in the AB 617 community of Shafter CERP.

THE INTERPRETER: I'm sorry, can you repeat that again?

MR. VIDRIO: That it will help us basically reduce the pesticide exposures or emissions in the Shafter community.

With that, I'd like to thank you for your time.

And I will ask if there are any questions that the Panel may have.

CHAIRPERSON ANASTASIO: Thank you, Edgar.

Any questions from the Panel?

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Edgar, I have a question for you. So you talk about weekly monitoring. This is at fixed points throughout the area?

MR. VIDRIO: Yeah, So DPR has been conducting weekly ambient air monitoring in the community of Shafter, for example, since 2011. So in that community, we've been collecting one air sample that we analyzed for over 36 different pesticides. 1,3-D is one of them. I don't know, is there a translation that needs -- that will happen here or --

CHAIRPERSON ANASTASIO: I think in the interest of time, maybe we should not do the translation for the questions from the Panel.

MR. VIDRIO: Okay. But yes, it's a fixed item in each community and we do it weekly and we use it for trend, and exposure estimates and so forth.

CHAIRPERSON ANASTASIO: So when you say weekly, that's one sample that integrates over the entire week or that's one sample per week at a shorter integration time?

MR. VIDRIO: Yeah, so it's a 24-hour sample collected every week in a random day.

CHAIRPERSON ANASTASIO: Okay. All right. But presumably a person's exposure at very high levels would

be on the order of minutes at a much higher level potentially. I'm just wondering -- I mean, are there health guidance values for 1,3-D?

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MR. VIDRIO: No. So unfortunately, there's absolute -- there's no State or federal agency that has any health standards for pesticides in the air, unlike some of the other criteria pollutants. So for -- because of this, DPR, when we first started the air network, coordinated with OEHHA to develop what we call regulatory standards and screening levels. So we use those as kind of a barometer to see how high or high low we are from those values.

CHAIRPERSON ANASTASIO: I see. And do you know, is there any interest on DPR's part on -- in developing health guidance values for some of these pesticides that don't have them?

MR. VIDRIO: Yeah. So essentially what we do for most of these pesticides is again we have a -- what we called a screening level. But for pesticides that have gone through the risk assessment process, which is a little bit more in detail, they look at all the health effects, the exposures, the sources. They develop what we call a regulatory health -- regulatory target, which is a little more stringent and has more impacts and triggers for the Department to take regulatory action if they

exceed it. So we do have that for a few of those pesticides, including 1,3-D.

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PANEL MEMBER RITZ: Is that -- this is Beate. Is that monitoring data available for research?

MR. VIDRIO: Yes. So DPR has every single sample that's ever been collected by both DPR and ARB since 2010 on our website. We used to have actually a very detailed very user-friendly interface. But because of different technology problems, we have a downloadable file that can be imported into any system that you would like to use.

PANEL MEMBER RITZ: And have you compared or are you regularly comparing this with the pesticide use reports?

MR. VIDRIO: Yes, we have. So every year, we develop a -- basically an annual report that has every single pesticide sample that was measured as part of the network. And we compared that with again either a screening level or regulatory target. And we -- if there's any exceedance then we talk about what was done to follow it up.

On top of that, we also look at -- every three years we look at the pesticide use patterns in the area and compare that or try to coordinate it to air concentrations, as well as look at how mitigation options that were placed during that time frame how that impacted

the use in air concentrations in the area. So we've done that and most of these document that we've done are posted on our website as well.

CHAIRPERSON ANASTASIO: Thank you.

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PANEL MEMBER KLEINMAN: It would be great if you could actually append some of those website listings to your presentation, so it becomes part of the record, so that if people do want to go look those up, they would be able to do it without having to wind through a more involved menu.

MR. VIDRIO: Yeah, we can definitely make those available. They're all pretty much on our website pretty easy to find, but we can definitely email those over so they can be included in a single document essentially.

CHAIRPERSON ANASTASIO: Yeah. Edgar, maybe you could cut and paste some of that into the chat. I believe the chat is archived as part of the presentation, so that would allow people to get it directly.

MR. VIDRIO: Yeah. Unfortunately, I don't have access to those documents right now. I know they're on our website.

CHAIRPERSON ANASTASIO: Okay.

MR. VIDRIO: Some of these, unfortunately because the State has been transitioning to an ADA new form of website, some of these documents may not be clickable

essentially on our website, but they're available.

Basically, there's a name, a document, and where -- a

place where the user can request that document. So I

could actually do that and just send those documents over

to you guys, and that way, you know, you'll have the

actual document as opposed to just the link.

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CHAIRPERSON ANASTASIO: Okay. Maybe you could work with Lori on that and then Lori could upload it to the website link for today's meeting.

PANEL LIAISON MIYASATO: I'll do that. Thanks, Cort.

CHAIRPERSON ANASTASIO: Thank you, Lori.

Other questions from the Panel?

panel Member Kleinman: Edgar, I have a -just -- this is more of an informational thing, but you
said it costs about \$1500 per acre for the TIF tarps. How
much of the cost of that is actually the cost of the tarp
and how much is the cost of getting rid of it after it's
used?

MR. VIDRIO: That's what we don't have a lot of idea. There's not -- there's not enough recycling plans essentially for this type of plastic. It's a different type of plastic. It actually has alcohol embedded in the sheets to block the fumigant from escaping, so it cannot be processed in the same type of recycling plants as

normal plastic is at this stage.

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In California, to my understanding, there was only one recycling plant, and that may have closed either last year or the year before. So we don't have any actual information on the recycling costs at this stage. But because there's no local area to -- or place to recycle, it might be higher now than it was three, four years ago, for example.

PANEL MEMBER KLEINMAN: Because I know a lot of toxics can be gotten rid of in cement calcining plants, because of the high temperatures of that process.

MR. VIDRIO: Right.

PANEL MEMBER KLEINMAN: And that has been done before. But I don't know if again whether that would be more practical than the existing method.

MR. VIDRIO: Yeah, because right now the use of these TIF tarps is very common, especially for like the hydroseed crops like strawberries, and mostly in the central coast regions. And so there is a way of getting, you know, rid of those, recycling them. But I -- we don't have any information on those about the cost at this stage.

PANEL MEMBER RITZ: So just out of curiosity, once you're taking the tarps off, is there any Telone left in the soil that escapes?

MR. VIDRIO: So fumigants have a very short lifespan, I mean, compared to, you know, the other type of chemicals. We have -- basically, when the fumigant is injected into the ground, the use of tarps are, for example, will keep it in ground, because it wants to move through the soil and either come out the top or breakdown with other -- with other nematodes and other organisms.

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But we have done testing where we look at how long that fumigant takes to break down completely, and we have seen that it takes anywhere from -- you know, most of it goes out between one and seven days. But there's some residue up to 21 days, for example. These tarps usually are kept in the -- especially for the strawberry production, they're kept for the whole season, because they also use them as a soil moisture regulator essentially, keeping the soil moist and keeping the water in there. So by the time they do take them out, there's pretty much nothing left in them, because it's been a few months at that time.

PANEL MEMBER RITZ: Thank you.

PANEL MEMBER HAMMOND: I'm pleased to see that you'll be actually testing some of the computer models. My memory of the methyl iodide situation was that after a lot of modeling on this, they actually used some tarps in Florida. And some of us had been somewhat skeptical of

the computer models. And I think that in Florida they found the concentrations released to the environment were much higher than what had been predicted by the computer model. So I think this is excellent that you're going to be actually checking the models.

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MR. VIDRIO: Yeah. No, we definitely want to, especially we're going to place a lot of interest or a lot of importance on the use of models for this. We wanted to make sure that we have the confidence in these. So again, that's why we initiated that peer-review process that CalEPA - we're one of the departments - usually goes with a UC and they do a peer review where they look at experts in those areas to evaluate not just the models that we're using, but how we're using them.

On top of that, we -- even though we have a lot of emission data on 1,3-D, we don't have a lot of emission data for the new methods. So there's definitely some extrapolation that we're doing. And we just want to confirm that, you know, the models are giving the right values when these methods are actually used in the real world essentially.

CHAIRPERSON ANASTASIO: Further comments from the Panel?

Edgar, I have a question for you. So you said that you have a rough health screening value for 1,3-D and

you've got this monitoring data. So do you see that there are occasional or regular exceedances of what you think the health value is in your monitoring?

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MR. VIDRIO: Yeah. So we've done monitoring of 1,3-D again in the City of Shafter for going over ten years now. On top of that, we currently monitor for 1,3-D in over ten different communities, again during that same 24 hours per week basis.

So during those -- that whole time, what we do is we have three basically levels that we compare to. We have an acute value that is usually done in the order of 24 hours. So we compare 24-hour samples with that value, and we do it anywhere from 24 to 72 hours, depending on where that acute study came from for different pesticides. For 1,3-D, it's 72 hours.

Then we also have a subchronic target of usually on the order of a few weeks. And lastly, we do the whole chronic, which is the year. So we compare air concentrations of 1,3-D to these values throughout the whole time. And so far, we've exceeded a subchronic value in the city of Shafter, for example, which is one of the reasons why it costs the Department to look a little closer into the current control measures for 1,3-D, as well as what else, if any, things should be done to reduce those emissions.

In terms of the acute, we have gotten a high-value of 111 parts per billion, for example. That is the maximum that we've gotten. And the current 72-hour target is 110. So on a 24-hour basis, that exceeded that value. But again, there' two different time scales, one was 24-hour concentration compared to a 72-hour target. But because of that again, it just caused us to look a little closer, so that we will have a definitive, whether that value or any other value has been exceeded. But it basically create or trigger the Department to look closely into those.

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CHAIRPERSON ANASTASIO: And presumably, there are people who live closer to the fields than the sampling site within Shafter?

MR. VIDRIO: And that's what we use the application site monitoring study. So the application site studies, as opposed to the ambient, gives us more of a sense of those acute exposures, because it's actually capturing — the complete emissions are coming out the field. And we do that — we usually have the studies for a — you know, again up to 21 days following the application. You want to capture everything that came out of the field.

Ambient, that's why we're in the areas that we're at, because we actually want to see what kind of

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quote/unquote normal air concentrations or average air
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    concentrations are exposed for the whole community. So
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    that that's kind of the difference of our two different
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    sampling types that we do.
 4
             CHAIRPERSON ANASTASIO: Yeah, I see.
5
                                                    I mean,
   part of the reason I ask is because you have this 60
6
   percent reduction target to get as good a reduction as a
7
8
    TIF tarp would. But it seems at least possible that there
9
    are exposures from residents near site where a 60 percent
    reduction would --
10
             (Off record: 11:36 a.m.)
11
             (Webinar shut down.)
12
             (On record: 11:46 a.m.)
1.3
             CHAIRPERSON ANASTASIO: So --
14
15
             PANEL MEMBER BLANC: Okay. I'd like to move that
16
   we adjourn
             CHAIRPERSON ANASTASIO: Can I get a second?
17
             PANEL MEMBER MILLER: Second.
18
             CHAIRPERSON ANASTASIO: Second. Thank you, Lisa.
19
20
             Okay. All in favor?
             Raised your hand and if I can see it or say aye.
21
             (Hands raised.)
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23
             (Ayes.)
             CHAIRPERSON ANASTASIO: Any opposed?
24
25
             Okay. I'll take that as -- so what I'll do is
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I'll work with Lori and the agencies to figure out what can we do to transfer over the public comment to the October meeting, because there were a number of questions that people had. Sorry about the technical glitch. I thought it was going reasonably well until then.

So just to remind everybody we will be meeting in October. Let me see here.

PANEL LIAISON MIYASATO: October 9.

CHAIRPERSON ANASTASIO: October 9th and 9:30. So please make sure that's in your calendar, October 9th at 9:30. We're still working on the agenda. You know, we normally stop at 2:30 or. So, for now, please plan for that and then as we get a better sense of the agenda, we will let you know if we can end earlier.

I also just wanted to let the Panel know that we have approved -- I, as Chair, approved the revisions that OEHHA made to toluene para-chlorobenzotrifluoride and cobalt. So we've made great progress on those and those are all in the bag. So congratulations to OEHHA, and John Budroe and staff on all of that.

Okay. That's it. Thank you, everybody. Sorry about the abrupt ending.

(Thereupon the California Air Resources Board, Scientific Review Panel adjourned at 11:48 a.m.)

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CERTIFICATE OF REPORTER

I, JAMES F. PETERS, a Certified Shorthand
Reporter of the State of California, do hereby certify:

That I am a disinterested person herein; that the foregoing California Air Resources Board, Scientific Review Panel meeting was reported in shorthand by me, James F. Peters, a Certified Shorthand Reporter of the State of California;

That the said proceedings was taken before me, in shorthand writing, and was thereafter transcribed, under my direction, by computer-assisted transcription.

I further certify that I am not of counsel or attorney for any of the parties to said meeting nor in any way interested in the outcome of said meeting.

IN WITNESS WHEREOF, I have hereunto set my hand this 21st day of July, 2020.

1.3

2.2

James & Path

JAMES F. PETERS, CSR

Certified Shorthand Reporter

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