
Safety Assessment of Diatomaceous Earth as Used in Cosmetics

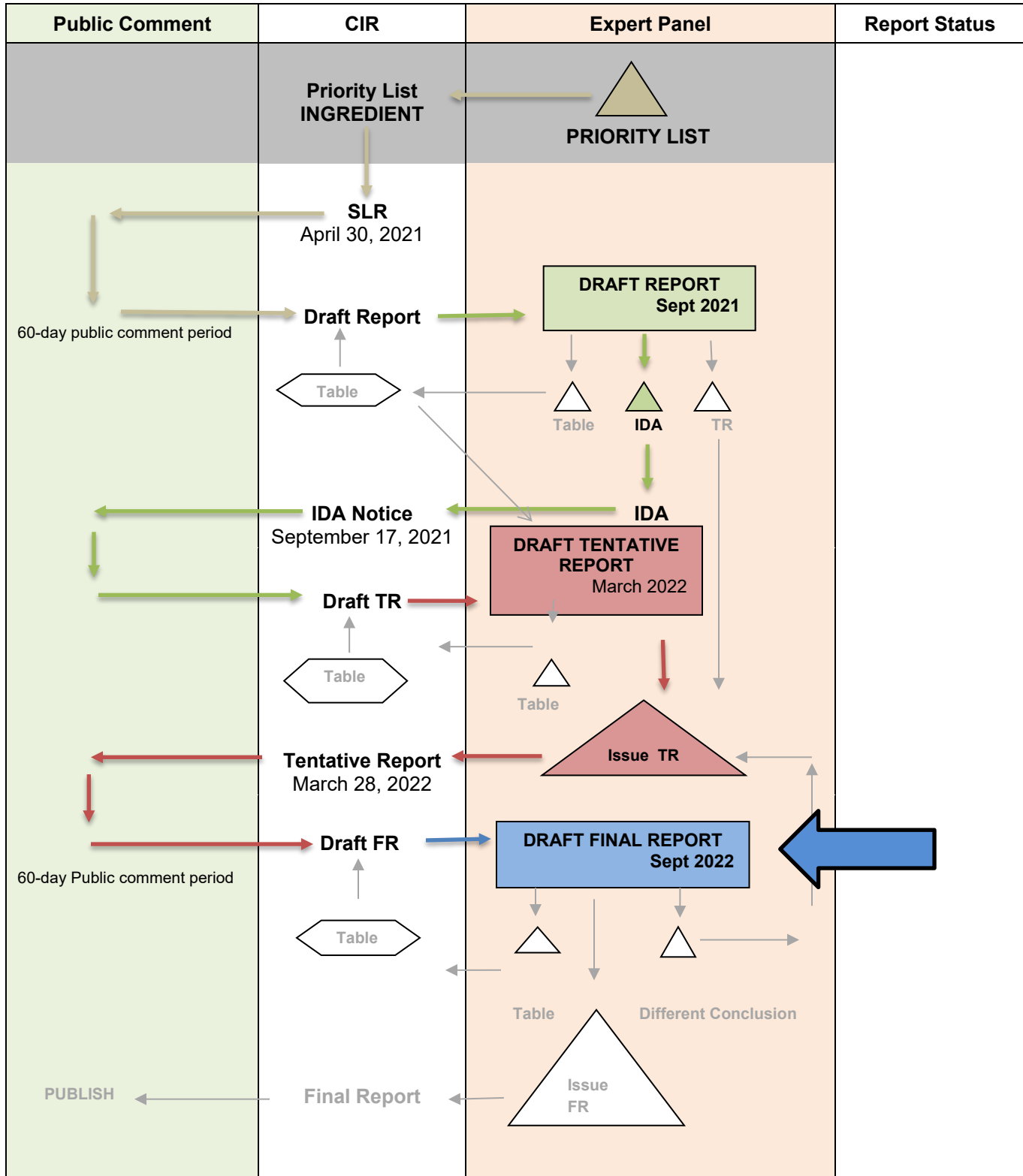
Status: Draft Final Report for Panel Review
Release Date: September 1, 2022
Panel Meeting Date: September 26-27, 2022

The Expert Panel for Cosmetic Ingredient Safety members are: Chair, Wilma F. Bergfeld, M.D., F.A.C.P.; Donald V. Belsito, M.D.; David E. Cohen, M.D.; Curtis D. Klaassen, Ph.D.; Daniel C. Liebler, Ph.D.; Allan E. Rettie, Ph.D.; David Ross, Ph.D.; Thomas J. Slaga, Ph.D.; Paul W. Snyder, D.V.M., Ph.D.; and Susan C. Tilton, Ph.D. Previous Panel members involved in this assessment: Lisa A. Peterson, Ph.D. and Ronald C. Shank, Ph.D. The Cosmetic Ingredient Review (CIR) Executive Director is Bart Heldreth, Ph.D. This safety assessment was prepared by Christina L. Burnett, Senior Scientific Analyst/Writer, CIR.

SAFETY ASSESSMENT FLOW CHART

INGREDIENT/FAMILY Diatomaceous Earth

MEETING September 2022





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Memorandum

To: Expert Panel for Cosmetic Ingredient Safety Members and Liaisons
From: Christina L. Burnett, Senior Scientific Writer/Analyst
Date: September 1, 2022
Subject: Safety Assessment of Diatomaceous Earth as Used in Cosmetics

Enclosed is the Draft Final Report of the Safety Assessment of Diatomaceous Earth as Used in Cosmetics. (It is identified as *report_DiatomaceousEarth_092022* in the pdf document.) At the March 2022 meeting, the Panel issued a Tentative Report with the conclusion that Diatomaceous Earth is safe in cosmetics in the present practices of use and concentration as described in the safety assessment.

Since the issuance of the Tentative Report, CIR has received no new unpublished data. The attached Council comments on the Tentative Report have been addressed (*PCPCcomments_DiatomaceousEarth_092022*), as noted in the check sheet immediately following the comments (*response-PCPCcomments_DiatomaceousEarth_092022*).

Additional supporting documents for this report package include a flow chart (*flow_DiatomaceousEarth_092022*), report history (*history_DiatomaceousEarth_092022*), a search strategy (*search_DiatomaceousEarth_092022*), transcripts from the previous meetings (*transcripts_DiatomaceousEarth_092022*), FDA VCRP raw data (*VCRP_DiatomaceousEarth_092022*), and a data profile (*datapofile_DiatomaceousEarth_092022*).

The Panel should review the Abstract, Discussion, and Conclusion, and issue a Final Report.



Memorandum

TO: Bart Heldreth, Ph.D.
Executive Director - Cosmetic Ingredient Review

FROM: Alexandra Kowcz, MS, MBA
Industry Liaison to the CIR Expert Panel

DATE: April 4, 2022

SUBJECT: Tentative Report: Safety Assessment of Diatomaceous Earth as Used in Cosmetics (release date March 28, 2022)

The Personal Care Products Council respectfully submits the following comments on the tentative report, Safety Assessment of Diatomaceous Earth as Used in Cosmetics.

Key Issue

The Toxicokinetics section states that no studies were identified. Although there may not have been any studies specifically about kinetics, the repeat-dose studies that measured silica in organs following oral exposure, and in the lungs following inhalation exposure should be mentioned in the Toxicokinetics section (or at least refer to the repeated dose sections). Currently, this information is only presented in Table 4 and in the Discussion. If the information is used to “help mitigate concern” in the Discussion, it would be helpful if it was mentioned in the text elsewhere in the report.

Other Considerations

Introduction – The second paragraph of the Introduction discusses the 2019 amorphous silica report (1 report), but the last sentence of this paragraph states “reports” twice. Either the “s” needs to be removed from “reports”, or more than one CIR report should be mentioned in this paragraph.

Non-Cosmetic Use – Please revise: “anticaking material foodstuffs” (add “in” between “material” and “foodstuffs”)

Short-Term, Subchronic, and Chronic – Please state the effects observed at the lowest concentration (0.7 mg/l) in the 28-day rat inhalation study.

Genotoxicity – It is misleading to call Diatomaceous Earth “unprocessed”. The method of manufacture section indicates that it is “crushed, dried, ground, purified and alimented”, saying uncalcined would be clearer.

Occupational Exposure Limits – The OSHA limit should be cited to an OSHA reference rather than a NIOSH reference. As (80 mg/m³/ % silicon dioxide), is an unusual format for a standard, an example of a value would be helpful (see: <https://www.osha.gov/laws-regs/standardinterpretations/2007-03-22>).

Summary – Please correct “for examples”

Discussion – Calling Diatomaceous Earth “unprocessed” is misleading as the raw mined product is at least ground before use, it is also likely that drying involves heating. If the Discussion is trying to distinguish between uncalcined, and calcined Diatomaceous Earth, it would be clearer to use those terms rather than “unprocessed” or “heat-processed”.

Table 4, 28-day rat inhalation study – Please correct: “OECD TA” to “OECD TG”

Table 4, reference 15 – Please identify the “tissues studied other than the lungs”. If the same tissues were studied for all species, this can be stated once for the rat study and the descriptions of the studies in other species can refer to the rat study.

Table 5 – The cell proliferation assay (reference 16) does not belong in the genotoxicity table (or in the genotoxicity section). It should be in a cytotoxicity section.

Diatomaceous Earth - September 2022 – Christina Burnett	
Comment Submitter: Alexandra Kowcz, Personal Care Products Council	
Date of Submission: April 4, 2022	
Comment	Response/Action
Key Issue: The Toxicokinetics section states that no studies were identified. Although there may not have been any studies specifically about kinetics, the repeat-dose studies that measured silica in organs following oral exposure, and in the lungs following inhalation exposure should be mentioned in the Toxicokinetics section (or at least refer to the repeated dose sections). Currently, this information is only presented in Table 4 and in the Discussion. If the information is used to “help mitigate concern” in the Discussion, it would be helpful if it was mentioned in the text elsewhere in the report.	Additional language regarding findings in repeated dose studies added to the Toxicokinetics section.
Introduction – The second paragraph of the Introduction discusses the 2019 amorphous silica report (1 report), but the last sentence of this paragraph states “reports” twice. Either the “s” needs to be removed from “reports”, or more than one CIR report should be mentioned in this paragraph.	Corrected to the singular.
Non-Cosmetic Use – Please revise: “anticaking material foodstuffs” (add “in” between “material” and “foodstuffs”)	Corrected as suggested
Short-Term, Subchronic, and Chronic – Please state the effects observed at the lowest concentration (0.7 mg/l) in the 28-day rat inhalation study.	Detail added.
Genotoxicity – It is misleading to call Diatomaceous Earth “unprocessed”. The method of manufacture section indicates that it is “crushed, dried, ground, purified and alimented”, saying uncalcined would be clearer.	Changed “unprocessed” to “uncalcined”
Occupational Exposure Limits – The OSHA limit should be cited to an OSHA reference rather than a NIOSH reference. As (80 mg/m3/% silicon dioxide), is an unusual format for a standard, an example of a value would be helpful (see: https://www.osha.gov/laws-regs/standardinterpretations/2007-03-22).	Both references no cited. Both entities cite each other’s limits in their guidance. Value is written as stated in the reference. It also is described as millions of particles per cubic foot of air (mppcf).
Summary – Please correct “for examples”	Corrected
Discussion – Calling Diatomaceous Earth “unprocessed” is misleading as the raw mined product is at least ground before use, it is also likely that drying involves heating. If the Discussion is trying to distinguish between uncalcined, and calcined Diatomaceous Earth, it would be clearer to use those terms rather than “unprocessed” or “heat-processed”.	Updated paragraph
Table 4, 28-day rat inhalation study – Please correct: “OECD TA” to “OECD TG”	Corrected
Table 4, reference 15 – Please identify the “tissues studied other than the lungs”. If the same tissues were studied for all species, this can be stated once for the rat study and the descriptions of the studies in other species can refer to the rat study.	Details added under protocol for each the rat, guinea pig, and dog studies as details were slightly different.
Table 5 – The cell proliferation assay (reference 16) does not belong in the genotoxicity table (or in the genotoxicity section). It should be in a cytotoxicity section.	Summary moved to cytotoxicity section.

Diatomaceous Earth History

April 30, 2021 – Scientific Literature Review issued.

September 2021 - the Panel issued an IDA. The additional data needed to determine safety for this cosmetic ingredient are:

- Clarification on the type(s) of Diatomaceous Earth that is used in cosmetic products (i.e., natural, calcined, and/or flux-calcined)
- Method of manufacturing for the type(s) of Diatomaceous Earth that is used in cosmetic products
- Composition and impurities data (including crystalline silicate content) on the type(s) of Diatomaceous Earth that is used in cosmetic products

November - December 2021 – Unpublished data received by CIR staff.

March 2022 - The Panel issued a Tentative Report with the conclusion that Diatomaceous Earth is safe as used in cosmetics in the present practices of use and concentration described in this safety assessment. Diatomaceous Earth is a polymorph of silica, or silicon dioxide, and is naturally-occurring. The Panel understands that Diatomaceous Earth, whether unprocessed (natural) or heat-processed (calcined or flux-calcined), can contain crystalline silica, a known respiratory carcinogen. However, the Panel noted that chronic inhalation studies of flux-calcined Diatomaceous Earth (which may comprise up to 60% crystalline silica) were negative for fibrosis or tumors in rats and guinea pigs. This data, coupled with the fact that Diatomaceous Earth is used as relatively low concentrations in cosmetics, mitigated concerns about use in products that may be incidentally inhaled, including face masks which may flake during drying.

Diatomaceous Earth Data Profile* - September 2022 - Christina Burnett

				Toxicokinetics			Acute Tox			Repeated Dose Tox			DART		Genotox		Carci		Dermal Irritation			Dermal Sensitization				Ocular Irritation		Clinical Studies	
	Reported Use	Method of Mfg	Impurities	log P/log K _{ow}	Dermal Penetration	ADME	Dermal	Oral	Inhalation	Dermal	Oral	Inhalation	Dermal	Oral	In Vitro	In Vivo	Dermal	Oral	In Vitro	Animal	Human	In Vitro	Animal	Human	Phototoxicity	In Vitro	Animal	Retrospective/Multicenter	Case Reports
Diatomaceous Earth	X	X	X			X		X	X		X	X			X			X	X		X		X	X	X	X	X	X	X

* "X" indicates that data were available in a category for the ingredient

Diatomaceous Earth

Ingredient	CAS #	PubMed	FDA	HPVIS	NIOSH	NTIS	NTP	FEMA	EU	ECHA	ECETOC	SIDS	SCCS	AICIS	FAO	WHO	Web
Diatomaceous Earth	61790-53-2; 68855-54-9	√	√	√	√	√	√	√	√	√	√	√	√	√	√	√	√

Search Strategy

PubMed

Diatomaceous Earth = 938 hits, 73 relevant

Diatomaceous Earth toxicity = 75 hits, 30 relevant

Diatomaceous Earth cosmetics = 20 hits, 6 relevant

Diatomaceous Earth dermal = 0 hits

Diatomaceous Earth sensitization – 3 hits, 1 relevant

Search updated July 2022 - 0 new relevant hits.

LINKS

Search Engines

- Pubmed (- <http://www.ncbi.nlm.nih.gov/pubmed>)

appropriate qualifiers are used as necessary

search results are reviewed to identify relevant documents

Pertinent Websites

- wINCI - <http://webdictionary.personalcarecouncil.org>
- FDA databases <http://www.ecfr.gov/cgi-bin/ECFR?page=browse>
- FDA search databases: <http://www.fda.gov/ForIndustry/FDABasicsforIndustry/ucm234631.htm>;
- Substances Added to Food (formerly, EAFUS): <https://www.fda.gov/food/food-additives-petitions/substances-added-food-formerly-eafus>
- GRAS listing: <http://www.fda.gov/food/ingredientpackaginglabeling/gras/default.htm>
- SCOGS database: <http://www.fda.gov/food/ingredientpackaginglabeling/gras/scogs/ucm2006852.htm>
- Indirect Food Additives: <http://www.accessdata.fda.gov/scripts/fdcc/?set=IndirectAdditives>
- Drug Approvals and Database: <http://www.fda.gov/Drugs/InformationOnDrugs/default.htm>
- FDA Orange Book: <https://www.fda.gov/Drugs/InformationOnDrugs/ucm129662.htm>
- (inactive ingredients approved for drugs: <http://www.accessdata.fda.gov/scripts/cder/iig/>)
- HPVIS (EPA High-Production Volume Info Systems) - https://iaspub.epa.gov/opthpv/public_search.html_page
- NIOSH (National Institute for Occupational Safety and Health) - <http://www.cdc.gov/niosh/>
- NTIS (National Technical Information Service) - <http://www.ntis.gov/>
 - technical reports search page: <https://ntrl.ntis.gov/NTRL/>
- NTP (National Toxicology Program) - <http://ntp.niehs.nih.gov/>
- Office of Dietary Supplements <https://ods.od.nih.gov/>
- FEMA (Flavor & Extract Manufacturers Association) GRAS: <https://www.femaflavor.org/fema-gras>
- EU CosIng database: <http://ec.europa.eu/growth/tools-databases/cosing/>
- ECHA (European Chemicals Agency – REACH dossiers) – <http://echa.europa.eu/information-on-chemicals;jsessionid=A978100B4E4CC39C78C93A851EB3E3C7.live1>
- ECETOC (European Centre for Ecotoxicology and Toxicology of Chemicals) - <http://www.ecetoc.org>
- European Medicines Agency (EMA) - <http://www.ema.europa.eu/ema/>
- OECD SIDS (Organisation for Economic Co-operation and Development Screening Info Data Sets)- <http://webnet.oecd.org/hpv/ui/Search.aspx>
- SCCS (Scientific Committee for Consumer Safety) opinions: http://ec.europa.eu/health/scientific_committees/consumer_safety/opinions/index_en.htm
- AICIS (Australian Industrial Chemicals Introduction Scheme)- <https://www.industrialchemicals.gov.au/>
- International Programme on Chemical Safety <http://www.inchem.org/>
- FAO (Food and Agriculture Organization of the United Nations) - <http://www.fao.org/food/food-safety-quality/scientific-advice/jecfa/jecfa-additives/en/>
- WHO (World Health Organization) technical reports - http://www.who.int/biologicals/technical_report_series/en/
- www.google.com - a general Google search should be performed for additional background information, to identify references that are available, and for other general information

SEPTEMBER 2021 PANEL MEETING – INITIAL REVIEW/DRAFT REPORT

Belsito's Team Meeting – September 13, 2021

DR. BELSITO: Boy, you got some good ones here, Christina.

MS. BURNETT: You know it.

DR. BELSITO: Okay. This is the first time we're looking at this. The SLR was released on April 30. Concentration of use survey, Council provided some dermal irritation, sensitization, and phototox. Comments from the Council were addressed. We also got some comments from the International Diatomite Producers Association, which are in here. And it's used in 116 formulations. Leave-on products and about a quarter of rinse-off paste masks. They have the 2019 concentration of use data to 5 percent face and neck, 20 percent hair, 62.2 in rinse offs. So that's what we got. So, under chemical properties, Christina, it just talks about the variety of shapes from which it's formed. Do we have any idea? I'm thinking in terms of inhalation issues.

MS. BURNETT: I'm sorry, can you repeat that again?

DR. BELSITO: So, under chemical properties for particle size distribution, medium and fine grade materials less than 90 microns. Do we have a lower limit? I mean, so --

MS. BURNETT: Right. Let me see if I can pull up that reference. Looking from ECHA data. It's going to take me a few minutes to figure that out.

DR. BELSITO: Okay, well, so the next thing is, on PDF page 11, we're going to get into the silica issue. It says in commercial products a large proportion of the amorphous silica in Diatomaceous Earth is converted into crystalline form during thermal processing, up to 40 to 60 percent.

MS. BURNETT: Correct.

DR. BELSITO: So, how are we going to deal with that, team members?

MS. BURNETT: We have a hand raised.

DR. BELSITO: Who has raised their hand?

MS. BURNETT: Mr. Ellis.

DR. BELSITO: Okay, can we allow him in?

MS. FIUME: He should be able to unmute his mic and speak. He is the person that sent us the IDPA information.

DR. BELSITO: Okay.

MR. ELLIS: Okay, can you hear me?

DR. BELSITO: Yes.

MS. BURNETT: Yes.

MR. ELLIS: Okay, great. Well, thank you, I realize that you're typically operating as the Expert Panel, but I had some communications with the CIR staff prior to this meeting to try to add some clarification that may be helpful to you as you evaluate Diatomaceous Earth.

I'm Mark Ellis, I am the executive director of the International Diatomite Producers Association, or IDPA, and we represent the major global manufacturers of Diatomaceous Earth. We became aware of what the Expert Panel and CIR were doing through our due diligence and learned about the scientific literature review. We discussed positions and we developed the comments that were filed that were shared with you.

And part of that due diligence looked at your consideration of the silicates because crystalline silica is an issue for Diatomaceous Earth, but part of what you need to appreciate is that there are different grades of Diatomaceous Earth. There's a natural form that's basically just dried and then there are calcined and flux-calcined grades that are thermally treated and in that thermal treatment the amorphous silica is converted to crystalline silica. And there are higher concentrations of crystalline silica in both calcined and flux-calcined Diatomaceous Earth.

The issue really comes down to how the CIR approaches some evaluations of chemical substances used in cosmetics. My understanding is that part of what staff refers to initially is the International Cosmetic Ingredient Dictionary and Handbook. And that staff is pretty much restricted to address what is in that dictionary and handbook.

And CIR staff shared with me the excerpt for Diatomaceous Earth, and it lists two chemical abstract service registry numbers. One is for natural Diatomaceous Earth, and the other one is for flux-calcined Diatomaceous Earth. And the general description also raises a question of calcined versus non-calcined, but they are wholly different animals in terms of their crystalline silica

content. And our members would prefer that CIR focus its scientific literature review, its draft report, or any tentative report moving forward solely on natural DE because it's essentially amorphous silica with very low, if any, crystalline silica content.

I'm going to offer another observation too because I think it's somewhat relevant to the discussion that you've had on the silicates. I'm a lawyer, but I've worked in the area of industrial minerals for about 40 years in a variety of settings. And a lot of it has focused in on occupational safety and health, so I have a lawyer's appreciation for toxicology.

One of the things that is difficult to appreciate is that these are not reagent-grade chemicals that you're dealing with. They have impurities in them because they are natural, so you may look at a chemical abstract service number and say that's what it is, but in reality what you have to look at in the evaluations you potentially are doing are, what is in the -- I used to call it the Material Safety Datasheet. It's the safety datasheet that lists the components in the product that the manufacturer of the cosmetic is using.

So, let's just say, if they list DE on there, and they have a chemical abstract service number for flux-calcined, it is going to have a much higher silica content than the natural DE. But there will also be other contaminants, potentially clays, and those aren't accounted for in the chemical abstract service number. They're only going to be revealed in the safety datasheet.

And I think that's part of what the issue you're dealing with here on the silicates, is that you're looking at one-tenth of one percent, which we know is the limit for listing a carcinogen on a safety datasheet. But it gets into limits of detections, limits of quantification as opposed to what's actually in the substance.

So, our sense of it is, is if you can depart from the strictures of that Cosmetic Ingredient Dictionary and Handbook and focus exclusively on natural DE, that your evaluation of the safety of DE in cosmetics will be tremendously improved. So I'll stop there.

DR. LIEBLER: Well, Mark, this is very helpful and your memo, which we reviewed when we received this draft report was also very helpful, particularly your offer to provide us additional information on the characterization of the Diatomaceous Earth ingredients that may be used in cosmetics. And I think that that would be very helpful to us. Some more information, more data would be really helpful to us.

MR. ELLIS: Well, we're interested in maintaining this dialogue with CIR to try to get a safe use determination for natural DE. I think part of it is that we're coming to you as outsiders without an appreciation for your procedures and policies. You know, I know enough to jump in and try to read what's there and respond to it, but much of the data, as I understand it, is coming from the manufacturers of the cosmetics themselves. And it's difficult for us to parse out from our members how much goes into cosmetics. I could tell you it's a small amount.

The principle use of Diatomaceous Earth is used in filtration and all of the processing that we do typically is geared towards making sure that beer that goes through it, wine that goes through it, oils that go through it are filtered appropriately. But it has many other uses because of its unique micro and macro characteristics as being the skeletal remains of these diatoms. We're happy to do what we can, but it's not a chief part of what we do, so I can't promise that we'll do everything that you want.

DR. LIEBLER: Well, Mark, we normally get nothing. We always get almost nothing from industry, so what you're offering to us is much more than that. And I realize there's some uncertainties about specifically defining which of these go into cosmetics.

On the other hand, if you can provide us information on the -- you know, if the overwhelming bulk of the production goes to DEs that are used in filtration for food and beverage, that's relevant to our thinking about this, even if you don't have hard numbers on what are the steps for cosmetic ingredients.

And the most useful thing, as I see it, is going to be a definition of the differences between the manufacturing processes that are used, the flux-calcined and the, sort of, natural that you mentioned, is briefly mentioned in our current draft report, but I think we'd benefit from better characterization that we can get from you and your team.

So, normally, the Expert Panel doesn't directly communicate with industry. It's CIR and the Council, or the Council communicates with industry, gathers data, passes it on to the CIR, and then the Expert Panel evaluates that. So that's the normal process. And what you're offering will be very helpful in that process.

MR. ELLIS: Okay, and we can work through Council to provide that information to CIR, and through CIR to the Expert Panel. Part of it is just determining what information you seek and what we are able to provide. But we do have a board meeting coming up in November, which is before your December meeting. Hopefully we'll be able to turn some of that information around and get it to you.

I plan on making a similar statement into the other breakout session. That was one of the things that was suggested because of how you're approaching your evaluation of these different chemical substances. So I hopefully will share this same kind of discussion with Dr. Cohen's group.

DR. LIEBLER: Don, you're muted.

DR. BELSITO: So I guess the major issue, from what I'm hearing, is the naturally sourced we're really not worried about crystalline silica, but with calcined and calcined flux that could be an issue, but we don't know what kind of Diatomaceous Earth is used as a cosmetic ingredient.

MR. ELLIS: Right, and I'm assuming that the only reason that flux-calcined and natural are in the same report is that somebody reported using that chemical abstract service number for flux-calcined, and we would counsel that that probably is not appropriate. We would suggest that they use natural DE.

DR. LIEBLER: At the time the definition was created in the dictionary, the flux-calcined got in there through whatever, you know, whatever source. That and the natural were the two that were in the dictionary. That's what we're bound to pursue because it's in the dictionary. So that's our starting point.

MR. ELLIS: Right.

DR. LIEBLER: If --

MR. ELLIS: And as I understand it, Dr. Heldreth said that, you know, you have some latitude as to what you consider and where staff is -- I'm going to use the word -- bound to address what's in the dictionary in the first instance that you can limit your evaluations to part of that if you feel is appropriate.

DR. LIEBLER: That's correct.

DR. BELSITO: Or a mixed conclusion.

DR. LIEBLER: Right, if need be, that's also correct. We get into these mixed conclusion situations, Mark, when we just don't have enough data to exclude. For example, if we have a significant amount of information that all leans towards natural, not flux-calcined in cosmetic products, then we can, in the discussion of our report, explain why we focused on the naturals, and that's what our conclusion is based on.

But, if we had insufficient information, then no. Then we kind of have to consider both as possibilities, and then we have to craft our conclusions accordingly based on the data. So more information helps us make a more informed assessment.

MR. ELLIS: Yep, I understand. Thank you.

DR. SNYDER: Mark, I have one question. So you mentioned the one-tenth of one percent as a requirement for listing as a potential carcinogen on any product. Where does that apply across? Would that apply to cosmetics?

MR. ELLIS: Well, it's part of the globally harmonized system of hazard communication. And I typically operate in the occupational realm rather than in the consumer product realm. But, typically, in a safety datasheet that would be required by OSHA, you would have to list as a carcinogen anything that has one-tenth of one percent of something that's been identified by IRAC in their monographs.

But, for instance, crystalline silica is identified as a group one carcinogen, which is a known human carcinogen. And natural DE is identified as a group three, which is unable to classify, and that's because there's limited evidence in humans, limited evidence in animals.

So, you know, that one-tenth of one percent is an artificial cut point, if you will. It's probably not toxicologically related.

I mean, I work with much higher concentrations of crystalline silica in an occupational setting and people that mine, people that work on construction sites, people that work in foundries, lifeguards on beaches, we all have crystalline silica in our lungs. And that's just because it's ubiquitous in the environment. But it probably has no toxicological effect because the body, over time, has developed mechanisms to deal with it, where the macrophages attack the crystalline silica and prevent fibrosis from happening. It's only when those biological responses are overwhelmed that you start to see the toxicological effects.

DR. ANSELL: The tenth of a percent is a threshold for disclosure.

MR. ELLIS: Yes.

DR. ANSELL: It isn't tied to --

MR. ELLIS: That's exactly it.

DR. ANSELL: -- any toxic event. Anything over one percent has to be disclosed, except for carcinogens, where the threshold is one-tenth of a percent.

DR. LIEBLER: Don, you're muted again.

DR. BELSITO: We also have the repeated dose toxicity study for inhalation on PDF page 12, specifically looking at flux-calcined Diatomaceous Earth.

DR. SNYDER: Yeah, that's all tabulated on Table 3, Don, on pages 18 to 21. And I read through all of those, and, to Mark's point, in many of those studies the only finding was aggregates sub-alveolar macrophages. Which would suggest that that is

reaching the lungs and it's being dealt with. It's just a matter of what levels exceed the capacity of the macrophages to engulf it and not let it drive any toxicity.

MR. ELLIS: Yeah, you mentioned read across earlier on in your discussion, and this is not a situation where a read across between flux-calcined DE and natural DE is appropriate.

DR. LIEBLER: We do not use read across for anything inorganic, Mark.

MR. ELLIS: Oh, okay. Thank you. That's something I didn't know.

DR. LIEBLER: Yeah. Sure.

DR. BELSITO: So, Mike, and to get back to the repeated dose tox study, would that clear the use of calcined Diatomaceous Earth and flux-calcined Diatomaceous Earth in respirable cosmetic products?

MR. ELLIS: I think that would be relevant. Again, I'm a lawyer, I'm not a toxicologist. But, you know, it seems to me that if you're looking at a repeat exposure type situation, that's something that ECHA, for instance, has evaluated and has it as a (inaudible) for inhalation. That's for the flux-calcined, so I just don't think it's appropriate to take that same moniker and attach it to natural DE.

DR. LIEBLER: So, Mark, just to be clear, if I was operating Ajax Cosmetic Company and I wanted to make a product that I wanted to use Diatomaceous Earth in, I could go to order Diatomaceous Earth and I would be presented with options of flux-calcined and natural and so forth, is that correct?

MR. ELLIS: That's true and it depends on who you buy it from because, if you buy it from a manufacturer, a manufacturer will typically work with a customer to try to meet a product that serves their specification. But, if they're buying it through a distributor, that product's already in commerce, and they may not have that same technical expertise that the manufacturer themselves might have.

DR. LIEBLER: Yeah, so in other words, there is the opportunity to have a selection based on properties, whether you customize it or wherever you get it, at least it's defined.

MR. ELLIS: Yes. I mean, you have a choice.

DR. LIEBLER: And the reason, where I'm going with this, I think if there's a substantial difference in the composition of flux-calcined versus natural and we can establish the degree to which natural versus flux-calcined is probably used in cosmetic ingredients, then I think we can deal with this. If the flux-calcined is not really used, then we don't need to be trying to assess the toxicity of flux-calcined.

MR. ELLIS: I think that's correct. The consultations that we've had with our members seems to be uniformly that they're only providing natural DE for cosmetic applications.

DR. LIEBLER: Yeah. I mean, anything we could get that documents that would be very helpful.

MR. ELLIS: Okay. Very good. I'm making a note there. Well, thank you again. I appreciate it.

DR. LIEBLER: Yeah, this is a big help. We appreciate it too, Mark, thank you.

DR. BELSITO: Thank you, Mark. Okay. I think that was really very helpful. To go back to the document, just --

MS. BURNETT: I was able to find the different particle size distribution information if you'd still like that.

DR. BELSITO: Yeah, sure, I think that should be incorporated too, obviously, but what was it, Christina?

MS. BURNETT: Okay. So you would like it for under ten microns?

DR. BELSITO: Yes.

MS. BURNETT: Coarse, approximately four percent or below is at ten microns. For the fine grade, it would be about 50 percent or less.

DR. LIEBLER: Yeah.

MS. BURNETT: I can type that up and put it in a table to help show that.

DR. BELSITO: But I still think that the repeat dose inhalation covers that, do we not?

DR. LIEBLER: What do you mean by that, Don?

DR. BELSITO: Well, I mean, so when Christina incorporates that, there's obviously going to be respirable particles.

DR. LIEBLER: Right.

DR. BELSITO: We know that the flux-calcined Diatomaceous Earth is going to contain a larger amount of crystalline silica, and we have that repeat inhalation dose toxicity study where there was no fibrosis over 2.5 years, so I think that covers the inhalation, correct?

DR. LIEBLER: I think so.

DR. BELSITO: Okay. Paul, do you agree?

DR. LIEBLER: Paul's muted.

DR. SNYDER: Sorry. The one study there, the hundred percent flux-calcined, where there was no observable effect concentration could not be determined. I do agree that one big study where they went out two and a half years and there was nothing, but I was trying to see at what concentration that was at. There's a one and a half year (audio gap).

DR. BELSITO: Well, they have a guinea pig study that was 1.5 years and --

DR. SNYDER: The dog study was two and a half years.

DR. BELSITO: Right.

DR. SNYDER: Yeah.

DR. BELSITO: And the particle range in the guinea pig study was 0.45 microns to greater than 10 microns, which I think is pretty good. I mean, you have multiple different species. Okay. And then the repro study, we don't have any developmental or repro --

DR. SNYDER: No, no DART. We have no DART, yeah.

DR. BELSITO: So do we need a 28-day dermal for absorption? I mean, we really can't go with GRAS status, can we? We don't know that all Diatomaceous Earth is used to filter wine, beer, et cetera.

DR. LIEBLER: I don't think we have any information that would suggest that these are absorbed. I mean, the little toxicokinetics suggests no absorption in the -- and that's in a dietary study. Livers, kidneys, spleens. Analyzed for residual silica, no difference between treated and controls. I can't imagine that the constituents of Diatomaceous Earth would be dermally absorbed.

DR. BELSITO: Right. So no oral absorption.

DR. LIEBLER: Yeah, no oral absorption, which would be easier.

DR. BELSITO: Right.

DR. LIEBLER: You know, we don't even have that. I mean, there's none of that. So I don't think a 28-day dermal is a reasonable request.

DR. BELSITO: Okay. So, in the discussion, we should point out that we don't have DART data, but there's no oral absorption making dermal absorption unlikely, or something to that affect?

DR. LIEBLER: Correct.

DR. BELSITO: Okay. And, then, just address, well, the subcutaneous exposure is really not pertinent to cosmetic use, so do we need to discuss that, the intraperitoneal too?

DR. SNYDER: No, those aren't relevant to cosmetic use.

DR. LIEBLER: Yeah. And I would point out under the non-cosmetic use, it says Diatomaceous Earth is GRAS as a filtering aid in food and beverages, and it's also, GRAS is the substance migrating to food from paper and paperboard products. In other words, if I read that correctly, it's understood that that could get into food from paper and paperboard products, and it's considered GRAS in that context. So I think that mitigates the systemic toxicity concerns.

DR. BELSITO: Right, okay, so then I can go into the discussion and -- then I just had a question, Christina, on the dermal irritation and sensitization studies. It says a cosmetic product containing 9 to 11 percent Diatomaceous Earth was not sensitizing in HRIPT, nor was it phototoxic in a human single application study, but, per our concentration of use table, I thought it was 0.9 to 1.1 percent.

MS. BURNETT: Yes, this data that was presented to us was a little tricky. Let me see if I can find the -- we were given a statement saying that this one trade name contains 9 to 11 percent and then they presented data afterwards using that trade name. Whether they reported it to the Council is how they use it. I know it doesn't match up with what the Council provided us in the survey, but I'm not sure how I can rectify that.

But on PDF page 33 is where we're giving a statement saying what these two trade names contain. So it could be from that they get diluted down into the formulation, but I don't know.

DR. BELSITO: Okay, and the max leave-on is 20 percent, but, I mean, I'm okay with the HRIPT at 9 to 11 percent. So we're going to have to have the heavy metals boilerplate and the discussion concerning calcined and flux-calcined and crystalline contaminants, but the chronic respiratory tox studies clear that. No repro or developmental, but no oral absorption, dermal absorption, not likely. Do not have the highest, or don't have data on sensitization at the highest, but I don't think we need it. So, for a conclusion, I would say safe as used, but, Dan, Paul?

DR. LIEBLER: I agree.

DR. SNYDER: I agree.

DR. BELSITO: Okay. Any other comments?

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DR. COHEN: Okay. May I move on to Diatomaceous Earth?

MS. BURNETT: Sure.

DR. COHEN: Okay. This is a draft report. This assessment is for a single ingredient, Diatomaceous Earth. It's the first time we're reviewing it. It's used as an abrasive absorbent, anti-caking agent, bulking agent, and opacifying agent in cosmetics.

We have frequency of use reported at 116 products and a max use of 5 percent on face and neck, and up to 20 percent in hair tonics and dressing at 62 percent in rinse off products, a paste mask.

It's also GRAS, as a substance, migrating to food from paper and paper board products. There's a discussion of crystalline silica content of 0.1 to 4 percent. We have some irritancy data on it, as well, and an HRIPT up to 11 percent.

I just had one question on PDF 24. It said, "A cosmetic formulation containing 0.9 to 1.1 percent Diatomaceous Earth." And it referenced 44 on PDF 29 which said 9 to 11 percent. I just wasn't sure if I was reading that wrong.

MS. BURNETT: I'm sorry. You broke up from me there.

DR. COHEN: On PDF 24, it said, "A cosmetic formulation containing 0.9 to 1.1 percent Diatomaceous Earth."

MS. BURNETT: Yes.

DR. COHEN: And it referenced 44 on PDF 29 that said 9 to 11 percent. And I wasn't sure if they were two separate things or the same thing off by a decimal point.

MS. BURNETT: I believe there's a dilution there.

DR. COHEN: Oh. Okay.

MS. BURNETT: Let me look which one that is, though.

DR. COHEN: So the question also here is, are we going to have a silicate discussion with this?

DR. PETERSON: Yeah, I think we have to because the silicate can be pretty high in some of these. If you look at the composition --

DR. SHANK: Yeah.

DR. PETERSON: -- it's the presence of the crystal and silica can be pretty high in some of the products. So I didn't have any data needs. That was the highlight is that there was a possible presence -- there is presence of crystalline silica which is going to be a concern for some of the products that are inhaled.

And that the inhalation studies, when they were done inhalation, they didn't see fibrosis, but when there was the intratracheal installation, there was evidence of lung issues. So I think we need to have a statement about considerations.

MS. BURNETT: We do have a hand raised. Before that, though, Dr. Cohen, that was a ten percent dilution on that in that table.

DR. COHEN: So it's one-tenth. Okay.

MS. BURNETT: Yes.

DR. COHEN: Thank you. Thank you.

MS. BURNETT: No problem. And Mark Ellis has his hand raised.

DR. HELDRETH: Yes, Mark is the executive director of the International Diatomite Producers Association, so he may have some insight about these ingredients.

DR. COHEN: Please proceed.

MR. ELLIS: Okay. Great. Thank you, Dr. Cohen. Let me please introduce myself. I'm Mark Ellis, I'm the Executive Director of the International Diatomite Producers Association or IDPA. We represent the major global manufacturers of Diatomaceous Earth products.

When we were made aware that the CIR was considering Diatomaceous Earth, we brought our members together doing due diligence, and we discussed difference positions and filed comments on this. And those have been part of the material that have been shared with you.

One of the things that we were aware of in that due diligence, was that CIR has been looking at silicates and the question of crystalline silica. And that really is an issue with Diatomaceous Earth because Diatomaceous Earth is more than just one thing. It is not a chemical reagent. It is a naturally occurring product that, in processing, is done in three different grades.

Natural DE is just lightly dried, but there's no thermal properties really attached to it, whereas calcine DE and flux-calcined DE purposely are calcined at higher temperatures. And, in the process of doing that, the amorphous silica is converted to crystalline silica. And it's done for a variety of reasons mostly linked to the principle purpose of Diatomaceous Earth which is as a filtration product.

But when we filed our comments on the scientific literature review, we made the point that we felt that the CIR should focus exclusively on natural DE because of its amorphous silica content rather than the calcined or flux-calcined.

In subsequent communications with Dr. Heldreth, we learned that, when the CIR staff approaches a chemical substance, they make reference to the *International Cosmetic Ingredient Dictionary and Handbook*. And that as they present the scientific literature reviewed view and the draft report, they have to address what's listed in that dictionary and handbook.

And that dictionary and handbook, Dr. Heldreth shared with me the excerpt there. And it spoke to not only natural DE with a chemical abstract service registry number, but to flux-calcined DE with a chemical abstract registry number. And then there's also in the definition a discussion of natural versus calcined.

And we think that this is an important point because it really confuses the issue to talk about Diatomaceous Earth as one substance. And, particularly, if we're going to be talking about the safe use of DE in cosmetic products, our members believe that you need to be focused in on natural DE and not on the higher silica content products.

And, from what I understand from your procedures, you have the ability to address these separate grades as opposed to dealing with them collectively. And I think that might facilitate you moving forward with a draft tentative report on the natural DE, whereas, it may be more problematic for the calcined or flux-calcined.

DR. PETERSON: Do we know which one is used in -- is it primarily the natural one that's used in cosmetics? I mean, is there some statement somewhere?

MR. ELLIS: Apparently, there's not. But what I received anecdotally from my members is that it is the natural DE that's used. And we had some discussion about that in Dr. Belsito's working group, and we'd be happy to talk to you about it, as well.

DR. COHEN: There's a comment in Composition and Impurities, that the crystalline silica content of un-calcined Diatomaceous Earth, is 0.1 to 4 percent. So is uncalcined DE the same as natural DE?

MR. ELLIS: Yes.

DR. COHEN: So is that specific sentence in there inaccurate?

MR. ELLIS: No, it's not. The problem that we're -- it's not a problem; it's a fact. The fact that we're dealing with is that this is a natural product. And a Chemical Abstract Registry number identifies it as being natural Diatomaceous Earth. But that assumes that it's basically a hundred percent pure, and it isn't.

And where you see that differentiation of what really is in the product is not in the Chemical Abstract Service number, but in a safety data sheet that might be produced by a manufacturer that discloses that it's 98 percent natural Diatomaceous Earth and lists the Chemical Abstract Service number. And it may have two percent crystalline silica and so it would list that Chemical Abstract Registry number for the amount of crystalline silica that might be in there.

But there also could be clays and other material that are in that natural deposit.

DR. BERGFELD: Is it forthcoming that something could be generated from your group to officially be included in our documentation?

MR. ELLIS: Yes, we talked about that. And I think a big part of it is determining what is most relevant to your deliberations. I think that part of what would help is to know what Diatomaceous Earth products are being used in cosmetics. And I know that the data that apparently goes into that dictionary and handbook is based on surveys done by the Personal Care Products Council.

I don't know whether we can trace, as producers, what material may be going in to cosmetics, but that would be potentially one thing we could do. If we could demonstrate that the bulk of Diatomaceous Earth going into cosmetics is natural DE, then you may be able to focus on what's really being used as opposed to what's out there as an alternative.

DR. COHEN: So it sounds like the crystalline content of natural DE is going to be the lowest of the three forms you've described, right?

MR. ELLIS: Yes.

DR. COHEN: But even in that situation, it could be up to four percent crystalline silica?

MR. ELLIS: Right. But I'm a lawyer; I'm not a toxicologist. I work in the occupational realm, and I do understand toxicology and epidemiology. You have to think about what the delivered dose is. It really doesn't matter what the percentage might be, although the percentage might cause you to believe that a higher-delivered dose might be there.

But we all have crystalline silica in our lungs. I mean, it's an ubiquitous thing. It's in dirt. It's in sand and dust. And our bodies over millennia have learned to adjust to that. We have different clearance mechanisms. Our macrophages can take that. It's only when the body's defense mechanisms get overwhelmed that we see indications of disease, most typically, silicosis.

But looking at a consumer product, from my perspective as a lawyer working occupationally, the exposures in a consumer product are going to be much less. I mean, California has done safe-use determinations on kitty litter or on flat latex paint. And the amount of silica that consumers might be exposed to in those instances is relatively small. And I might tentatively offer that it probably would be the same in cosmetics.

DR. COHEN: I think our dilemma stems from our silicate discussion, as a whole, right?. I mean, this is an off-shoot of that which is this quandary of taking PELs or occupational exposures or lifetime exposures and producing advice or our opinion on the manufacturing of a product. How do you take lifetime exposures, or exposures that have a time variable in there as they do in occupational medicine. It's over a number of hours per day, per year worked. How do you take that and say, this is how much you could put in a cosmetic agent for personal use? I personally have a hard time figuring that.

MR. ELLIS: Well, you should. And my guess is that there probably has not been a lot of transcription between what is a safe occupational -- or take it conversely -- a hazardous occupational exposure and how that translate to a safe consumer exposure.

There's scaling, obviously, involved. And, in the occupational setting, it's typically safe to work for a 40-hour work week for a 40-year working lifetime. So people like miners, people that work in foundries, those kind of people are exposed to crystalline silica at much higher rates than any consumer would be. And those occupational exposure limits tend to be protective.

DR. COHEN: Can I ask your opinion? Were you attending to the meeting when we spoke about silicates and our conclusion about the detection about crystalline silicates?

MR. ELLIS: I didn't attend the last meeting, but I did review the material on silicates. And I mentioned this point to the other working group. I think that you're confronted with talking about apples and oranges in many contexts here. This notion of one-tenth of one percent or one percent, those typically are related to the presence of a carcinogen.

Let's just take the case of crystalline silica. The International Agency for Research on Cancer has identified that as a Group 1 carcinogen, a known human carcinogen. But IARC also has looked at natural Diatomaceous Earth and it rates it as a Group 3, insufficient evidence in animals and insufficient evidence in humans. So it's not classifiable.

I think that what everybody focuses on -- and we do, too. If you look at those one-tenth percent or one percent numbers, they relate to specifically addressing the presence of a carcinogen and inhalation route of exposure.

And, again, a working lifetime because that's what the globally harmonized system classification deals with. But as you're dealing with consumer products, how do you get to what the delivered dose is and whether or not the body's mechanisms are sufficiently active to overcome that insult?

DR. COHEN: Yes, you've summarized the issues we've gone through on the silicates and how we're trying to understand Diatomaceous Earth.

DR. BERGFELD: Well, it sounds to me like we've been directed to using only the natural Diatomaceous, which has the lowest silicon. And discussing the other two as probably should not be included in cosmetic products until further evidence is presented.

DR. COHEN: So I think for purposes of having aligned messages. Two chemicals ago, we were talking about crystalline silica, and now we're talking about crystalline silica again. We don't quite have evidence that just uncalcined DE is used in cosmetics. We think maybe that's possible, but we don't know that for sure. And I think, Wilma, your comment about having that in the discussion is important.

And we have max use of 5 percent in a face and neck product, 20 percent in a hair tonic and dressing. And then we have this rinse off face paste. So that's a very interesting issue of 62 percent DE applied to the face to dry, right, perhaps, and then being in close proximity to the nose and mouth as it's drying. And then you rinse it off which is probably perhaps a lower risk.

Where do we align the level of detection of crystalline silica into this assessment?

MR. ELLIS: Well, I think that IARC specifically looks at crystalline silica as a Group 1 lung carcinogen. So inhalation is really the route of exposure that you would have to work with. But, obviously, something that's wet is not biologically available in that situation, so you probably can move away from concerns there.

I do think that the pursing out of this 0.1 percent and 1 percent is problematic because it doesn't get to the issue of delivered doses. It's more a limit of quantification or a limit of detection as opposed to a determination of what the dose is.

I think that one of the things that I will try to do -- because we are interested in working with CIR to get a safe-use determination from natural DE. I will try to work with our members to determine, sales into the cosmetic market, to what extent are they natural DE, to what extent are they calcined and to what extent they are flux-calcined?

And what I would propose to do is to communicate this information to the Personal Care Products Council, and I'll copy Dr. Heldreth on this. But the notion would be then that PCPC would share that information with CIR and CIR would share with the expert panel.

DR. COHEN: Tom, thoughts?

DR. SLAGA: Should we wait for that? Table this? Or what are we --

DR. COHEN: I'm a little stuck on -- let's say it comes back and it's natural DE -- it's all natural DE in cosmetics, right?

DR. SLAGA: Right.

DR. COHEN: We have a sentence in composition and impurities that the uncalcined is up to four percent crystalline silica, right?

Now, that's just the uncalcined DE. It gets diluted down once it's in products, you know, down 20 times. But, in this face mask, it's two-thirds DE. It's basically a DE mask. That's applied and then I don't know if it's dried or rinsed off before it dries. I just don't know.

How do we deal with that four percent crystalline silica issue even under the best of circumstances? And we don't have to make a final decision, here, but where are we generally going to work with this?

DR. PETERSON: So the problem is inhalation. So if you put a mask on, it's wet, you dry it off. There's not that much that you would necessarily inhale from putting on a mask unless it's sprayed on, but mostly you don't spray on a mask. I think you have to worry about the products that are inhaled.

DR. HELDRETH: There's also a 20 percent max concentration of use incidental inhalation sprays.

DR. COHEN: So that would get us down to 1 percent crystalline if we use the max of 4.8 percent. If we use the max of 4 percent crystalline.

MS. BURNETT: It looks like we have another hand raised.

DR. COHEN: I can't see who it is. Can you see, Christina?

DR. BERGFELD: Shripal Sharma.

MS. BURNETT: Shripal Sharma?

MR. SHARMA: Yeah. Good afternoon, everyone. My name is Shripal Sharma, and I work with Imerys as Director of Product Stewardship. And we are one of the suppliers of Diatomaceous Earth that Mark Ellis just mentioned. So I'd like to clarify a couple of issues here for the benefit of this expert panel.

One is that, based on our industry knowledge, we are not aware that any calcined DE or flux-calcined DE are used in cosmetics. We are only aware that only natural DE are used in cosmetics. Again, this is the knowledge -- that's the information that we have based on our own understanding of the market and based on the survey that Mark carried out with ITP companies.

The second point I would like to make is that, even though natural DE may contain some level of natural quartz as crystalline silica up to four percent, not all of those natural DE products are suitable for cosmetics. So, in general information, we produce and sell into various markets natural DE.

And only a fraction, maybe less than one percent of that natural DE goes into cosmetics. And even natural DE, as you say, from zero percent crystalline silica to four percent, based on our understanding, the natural DE going into cosmetics are relatively very low on crystalline silica, not in the range of the four percent, rather, on the lower side.

DR. BERGFELD: How can you document that?

MR. SHARMA: We document that based on the work that we do with our customers who are cosmetic customers, as an example. We work with them to provide them with samples they determine what the specification of the product will be. And, so, we document based on our own testing and the product that goes to the customer.

DR. BERGFELD: So you have a product line that is only cosmetic, is that right? And has certain classifications?

MR. SHARMA: No, we have hundreds of product lines for natural DE. And based on the specification that a customer is looking for, we sell one of those hundred products to those customers. And, based on our testing regime in our plant, we know what that product is and what its composition looks like.

DR. SLAGA: Well, we should be able to get that documentation then. Right?

MR. SHARMA: Yeah, the testing that we carry out in our lab, that's our documentation.

DR. BERGFELD: But the problem I see is who purchases it? Is it the cosmetic industry or others? We're interested in the cosmetic.

MR. SHARMA: Yeah, so, what I'm saying if we know which products are used in cosmetics. And we have the test results for those.

DR. BERGFELD: I'd like to see that.

MR. SHARMA: Okay.

DR. COHEN: Yeah, that would be useful.

DR. HELDRETH: Would that be something we could cite from the document?

MR. ELLIS: I think that there might be an issue there of confidential business information, and I can't speak for Imerys which is one of our member companies, but I know that others might have that concern.

If PCPC or CIR has provisions for receiving data that's confidential business information, we probably can get that information to you.

DR. HELDRETH: Yeah, CIR procedures don't allow for us to receive confidential information. Everything that we receive and the Panel relies on gets published on our website for anybody to see.

Now, you can submit information to the Personal Care Products Council. In the past, I've seen them scrub out company details, but the data needs to still be completely visible so the Panel can make their evaluation.

DR. BERGFELD: Bart, could we get a letter of summary? Just a general summary that is sent to us by the company who is the supplier based on their information without sharing in great detail?

DR. HELDRETH: Yeah, if the Panel feels comfortable with that I see no issue with that, if they're willing to provide such.

DR. COHEN: I think it will be helpful. I still think we need to harmonize our silicate manuscript with this one. So I agree. The method by which it's applied really will speak to risk. So, with the silicates we have, I think, a notation when there's a respiratory risk. When there's a risk of respiratory exposure, this is what we ask for. If there wasn't a respiratory exposure, we didn't really comment on the amount of crystalline silicate. Did I remember that correctly?

DR. BERGFELD: Yes.

DR. HELDRETH: Yeah.

DR. COHEN: So for DE, many of these exposures are -- some of them are not respiratory; some are. And when they are, wouldn't we ask for the level of crystalline silica in the final product? Not the DE itself, but the final product would be less than 0.1 percent crystalline silica.

DR. BERGFELD: That can be done.

MS. BURNETT: Yes, this is a draft report, so this is the first stage. You may issue an insufficient data notice with whatever needs you would like. So, if that is a need -- if you would like the Council, you know, you can ask the Council to ask their members about the calcined versus non-calcined. You know, what they use in their products. You can do that, too.

DR. COHEN: Yes, that was one of the IDA questions, the use of uncalcined versus calcined or flux-calcined DE in cosmetics.

DR. SHANK: The silicate report puts the limit on the silicate as less than 0.1 percent in raw material, not in the final formulation. Or it says, "Or if there is repeat dose inhalation data, but there is no adverse effect."

DR. COHEN: Right, but the silicate is the target chemical of that report. This is a contaminant or a component of a different thing, right? So the silicates are a component of the DE. That other report is purely for silicates.

DR. SHANK: But, as a raw material not as an ingredient.

DR. BERGFELD: Final formulation.

DR. SHANK: What the conclusion says that's a raw material not in the formula.

DR. COHEN: So, Ron, you're suggesting this raw material should have less than 0.1 percent crystalline silica in it?

DR. SHANK: Yes, that's what we said. But, the silicate report, the 0.1 percent is based on limited detection.

DR. BERGFELD: That's a good route to go.

DR. COHEN: It's the logical route to go, right, because it's the same day we're talking about this issue, right?

DR. BERGFELD: I guess so. Yeah, consistency.

DR. COHEN: From the industry, could you comment on that kind of conclusion just so we have some perspective?

MR. ELLIS: I'll take a shot at it. One of the things that I did most recently was I served as president of the Industrial Minerals Association North America, and I did that for 17 years. And I'm moving towards retirement, which is not coming soon enough. But, again, I have a 4-year career working as a lawyer but working principally in the area of occupational safety and health.

And I've worked with toxicologists and epidemiologists. And I'm convinced that you can use any product safely including nuclear radiation or coal if you use it wisely and use it prudently and you take precautions in how you use it. And I think that's part of the problem that we're dealing with here with crystalline silicate. I don't know how you deal with anything else that's been identified by IARC as a carcinogen. Because now you're translating to a strict number which is -- I hate to say it -- I believe it's arbitrary. One-tenth of one percent? It's something, as someone else mentioned, you can quantitate.

But as a toxicologist or medical doctors, you know the doses that poison, so how much is likely to cause an adverse reaction in a human? Not symptomatic, but I'm saying disease. And I think that the numbers we're talking about here are infinitesimally small compared with what you deal with in the occupational setting.

And I know that there's a precautionary principle, and that you're dealing with a consumer product. There may be certain reservations about what may be a safe exposure, but I do think that the literature on occupational exposure is a guide to take a look at what consumer exposures might be.

Again, translating is another issue I can't touch, but I think that when we're dealing with this one-tenth of one percent or one percent, those are really arbitrary numbers that don't relate to a delivered dose that may have anything to do with an adverse effect.

DR. COHEN: I fundamentally agree and understand regarding your comments. And, again, with occupational exposure limits, they're contextualized with time, right, time and place of exposure. We don't have that here. And one other thing that sort of resonates with me -- and I'm not sure -- but I don't recall ever seeing a safe dose of crystalline silica. So we're stuck between a dose that doesn't cause any demonstrative clinical disease and some other dose.

I'd ask our team, Tom, Ron, what's the safe exposure to crystalline silica?

DR. SHANK: We don't know.

DR. SLAGA: That's right. We don't know.

DR. SHANK: That's the problem.

MR. ELLIS: So take hydrogen fluoride as an example. That is a chemical that I would not want to play with. And small amounts can be fatal. You don't find that with crystalline silica. You have a long latency period. The body has clearance mechanisms that address most exposures that you have every day. But hydrogen fluoride is a whole difference animal.

DR. COHEN: I think it's a lot for us to contemplate. I think we're issuing an IDA at this point. And, Ron, I don't think we have sensitization data on max use or anything close to that, right? Christina, it was about 1 percent, 0.9 to 1.1 percent in the HRIPT? Because it was a dilution in there? Or is it 11 percent? Table 5 --

MS. BURNETT: It's a dilution. It was 10 percent up to 11 percent.

DR. COHEN: So --

MS. BURNETT: So then that would be 1.1 percent.

DR. COHEN: Yeah, so we're at 1.1 percent for the HRIPT, and we have max uses of 20 to 60 percent.

DR. SHANK: Twenty.

DR. COHEN: Well, there's a max with 60 percent, Ron.

MS. BURNETT: That's a rinse off. Do you want the max rinse off use or the max leave on use?

DR. COHEN: What's the max leave on use again?

DR. SHANK: Twenty percent.

DR. COHEN: Twenty percent.

MS. BURNETT: Right.

DR. COHEN: You know what's interesting? What then constitutes a max? So it's sort of in between. I look at rinse off products as shampoos, conditioners, soaps that go on and go off. What if you leave a mask on for three hours? Or six hours? Or overnight? Is that a rinse off product or a leave on product? I'm not quite sure, right? It's a provocative question, but at least -- all right, say 20 percent. We're off by a factor of 20 on that. So we need sensitization on max use, right.

DR. SHANK: Yes.

DR. COHEN: What else do we need?

DR. SHANK: Should we drop the calcined Diatomaceous Earth from the report?

DR. COHEN: There's only one ingredient in the safety assessment. But, apparently, there's methods of manufacturing that are different. So would we not put that in a discussion?

MS. BURNETT: Do you want industry to clarify?

DR. PETERSON: Yes.

MS. BURNETT: You want something like a memo or something that details what their suppliers use? Their members?

DR. BERGFELD: Very important, yes.

DR. PETERSON: And then I think in the discussion we can say that we wouldn't support the use of these other -- the natural one is going to be the safest.

MS. BURNETT: Would you like them to detail their impurities? Or do you think what we have is sufficient based on what I found?

DR. COHEN: If this data that is different from this, it would be very helpful.

MR. ELLIS: Okay. Dr. Cohen, I just want to thank you and the expert panel for allowing me to make those remarks.

DR. COHEN: We appreciate them. They were very valuable in helping us get through this. Thank you.

From the team, any further comments? Wilma, any?

DR. BERGFELD: I didn't hear. Is it Mike? Did he say he would give us that memo? Mark. It's Mark Ellis. Yeah, Mark.

MR. ELLIS: I think that I'm a little uncertain as to what that memo might say, so perhaps, doctor, after the discussion tomorrow in the plenary session, that could be narrowed down.

DR. BERGFELD: Thank you.

DR. SHANK: Okay.

DR. COHEN: Any further comments from the team before we move on? Okay.

Full Panel Meeting – September 14, 2021

DR. COHEN: Yes, this was a source of considerable time and effort and discussion. So this is Diatomaceous Earth, which is a draft report, and it's the first time we're reviewing it, and the assessment is for this one ingredient. It's used as an abrasive, absorbent, anticaking agent, bulking agent, and opacifying agent in cosmetics. We have max use of five percent in face and neck care products, 20 percent in haircare products, and 62 percent in a rinse-off paste mask. And we have frequency of use reported. Diatomaceous Earth is also GRAS, as a substance migrating to food from paper and paperboard products.

Our group issued an insufficient data announcement with the request for clarification of the method of manufacturing for the three major types of DE, namely natural DE calcine and flux-calcine. Particularly since their crystalline silica content are different, we'd also like further information about the disposition of those three types as they relate to use in cosmetics.

We have irritation data but we wanted sensitization data at max use for a leave-on product. We also are faced with the information that even the uncalcined Diatomaceous Earth has a crystalline silica content of up to four percent. So, we couldn't help but interlace the conversation we had about silica earlier. That's the motion right now.

DR. BERGFELD: And it's an insufficient data announcement motion, correct?

DR. COHEN: It is.

DR. BERGFELD: Okay. Don?

DR. BELSITO: Yeah, so we had a slightly different take on this. And, we certainly appreciate, you know, the Cohen team's approach. We thought that we could potentially go with a safe as used conclusion and in the discussion limit heavy metals, crystalline contaminants, and that its GRAS status cleared systemic toxicity endpoints. We recognize that we didn't have sensitization at the highest leave-on of 20 percent, but again there's nothing in these ingredients that really would be sensitizers.

DR. LIEBLER: Well, Don, the only thing that didn't get highlighted in our discussion yesterday is this issue of the uncalcined Diatomaceous Earth having up to four percent crystalline silica. You know, it was right there in front of me and I didn't really flag it. That becomes then the same issue for incidental inhalation as with the silicates. And that's the only problem, other than that Diatomaceous Earth is devoid of systemic tox. So, I agree with David and his team that that's an issue we need to address.

DR. BELSITO: Okay.

DR. SNYDER: But I thought we learned yesterday that only natural Diatomaceous Earth is used. Those other two forms are not used in cosmetics so this would be a little bit like the silicates and we would say this report is only dealing with the natural Diatomaceous Earth. And it would have a similar type of discussion regarding the crystalline silica. And it would be expected that the flux and the calcinate would not be used in cosmetics and because of the presence of crystalline silica.

DR. LIEBLER: Well, but they --

DR. PETERSON: And I would add something similar to what's in the silicate that, you know, the expectation is -- because the conversation we had with industry was that they have some ability to manipulate the crystalline silicate that's in the Diatomaceous Earth. And, so, it seems like we could put a similar caveat to this one that we put in the silicate where it has to be formulated such that the crystalline silicate is below -- I forget how we worded it in the silicate, but that seems like it would be appropriate here.

DR. LIEBLER: The thing is with the silicates we came around to not being able to define a safe level of crystalline.

DR. PETERSON: Right, right, and so you come to the same thing here because, you know, you basically have another source of crystalline silicate, so the conclusion should be similar to what it is in the silicates in that there is no safe level of the crystalline silicates that we know of for inhalation. So (audio skip) that caveat.

DR. LIEBLER: Correct. Right.

DR. COHEN: We didn't have certitude that only natural DE was going into cosmetics. We had an assumption, I don't know, did we have that in a report that that was the case? We were sort of reassured, but we wanted more clarification from industry on that. That was part of our IDA.

DR. BERGFELD: I think we asked for a memo.

DR. COHEN: Yeah.

DR. LIEBLER: Yeah, I think that Diatomaceous Earth is only naturally sourced. It's not synthetically generated. And then it's maybe processed by this flux-calcination process for some applications. But it all comes from natural sources. So the unflux-calcined Diatomaceous Earth may have crystalline silica between .1 percent and four percent as it says in our report, and there's the rub.

DR. COHEN: That's exactly the rub, because our last iteration of silica had a .1 percent because that was the level of detection. And, I'm recalling that I think one of the manufacturers reported something like .11 percent. So, if we stick strictly to the last version of the silicate document, then it looks like we're knocking Diatomaceous Earth out of cosmetic use.

DR. LIEBLER: For inhalation?

DR. COHEN: For incidental inhalation.

DR. LIEBLER: Incidental inhalation, yeah.

DR. SNYDER: We're saying the data is insufficient to support safety, yeah.

DR. LIEBLER: Yep.

DR. COHEN: Well, we thought we would get further clarification about method of manufacturing, assurances about where each type of DE processing went where and specifically which one were in the cosmetic industry. And, I agree with Don. I think using that expert interpretation, the risk of sensitization would be low and we have irritation data. So, I would reiterate the IDA for further information on that and see what we get back from the industry.

DR. LIEBLER: This report is in an early stage and we've got the industry trade group very motivated to work with us to help us with our data needs. So I think that's a good place to be for now. So, that'll help us as we go forward.

DR. BERGFELD: So, what is the Belsito group doing now? Dan's going with the suggested conclusion. Don? Paul?

DR. BELSITO: I'm fine.

DR. BERGFELD: Paul?

DR. SNYDER: I'm fine. It's early stage, we're fine.

DR. BERGFELD: Okay. So, we're considering that a second to the Cohen conclusion, correct? It's an IDA.

DR. COHEN: It's an IDA, but I'd like to just restate that the IDA did not include sensitization at max use.

DR. BERGFELD: Okay. Did you want to add that?

DR. COHEN: No, we had it originally and then I think Don's comments and his team were provocative enough to change that motion.

DR. BERGFELD: You can put it in the discussion, though, if need be.

DR. COHEN: Yeah.

DR. BERGFELD: Okay, any other discussion regarding this particular ingredient? Lisa?

MS. BURNETT: Just to clarify for me.

DR. BERGFELD: Okay, Christina?

MS. BURNETT: The data needs are clarification on the method of manufacturing for the three types. And clarification from industry as to what type might be used in cosmetics, and any composition and impurities that can be gathered from that.

DR. COHEN: Yes.

DR. BERGFELD: I think the concentration on crystalline silica is there, need to know.

DR. COHEN: During the conversation asked if there was any additional composition and impurities data that might be brought into the report, if it was available.

MS. BURNETT: Thank you.

DR. BERGFELD: Okay. All right, any other comments, or, Christina, do we need anything else for clarification?

MS. BURNETT: I believe I have the two points that are needed. Thank you.

DR. BERGFELD: Okay. All right, I'll call the question. All those opposed? Abstaining? Unanimous approval of an IDA with the stated needs.

MARCH 2022 PANEL MEETING – SECOND REVIEW/DRAFT TENTATIVE REPORT

Belsito's Team Meeting – March 7, 2022

Dr. Donald Belsito: OK, so diatomaceous earth. So is your astrological sign an Earth sign, Christina?

Christina Burnett (CIR): No, it's an air sign actually.

Dr. Donald Belsito: OK, At the September 2021 meeting, we issued an IDA. We wanted clarification on the types of diatomaceous earth that are used in cosmetic products, whether they're natural, calcined, or flux calcined, method of manufacturing, and composition or impurities. Data including crystalline silica content on the types of diatomaceous earth used. We've received information from a supplier providing information on diatomaceous earth used in cosmetics, soda ash, flux calcined. A method of manufacturing, composition or impurities...we received in vitro ocular. Information that clarified some of the previous safety data testing. We also got comments from the IDPA for the panel's consideration. The use tables have been updated. Now 2% rinse off products. As opposed to 62.2% and point out 1% in leave on products mainly now polish and enamels. So where are we with these documents? We also got wave twos and threes on this one.

Dr. Dan Liebler: Well, I think we last time we learned that, the overwhelming majority of the diatomaceous earth supplied is natural, not calcined or flux calcined. But then we got one outlier, apparently, which does produce a flux calcined containing material. But then again, we have the acute tox study in the dust aerosol study on PDF 32, which there are no abnormalities

observed after administering flux calcined, diatomaceous earth. So, I took that as pretty strong weight of evidence that even the highest reporting crystalline silica content, 4%, would not be expected to produce lung injury if inhaled.

Dr. Donald Belsito: So you thought the calcined and the flux were OK?

Dr. Dan Liebler: You know, we don't have extensive data, not on it. We only had a little bit, but I think that sort of allows you know for the fact that the majority of, the overwhelming majority of what's produced or included in cosmetics is natural and not calcined and flux calcined.

Dr. Donald Belsito: I'm not so sure, Dan. Look at the dermal irritation and sensitization studies. They were all done on flux calcined.

Dr. Dan Liebler: Right. So, we have two things. One is we have the reports of method of manufacture, which is overwhelming majority natural, not calcined and flux calcined. Then we have a tox study and we have your dermal irritation and sensitization that include calcined and flux calcined. So I don't know exactly how we word this in a discussion, but it indicates that even if there is calcined or flux calcined as a contaminant or a component of what's used, this provides pretty strong weight of evidence, for lack of concern.

Dr. Donald Belsito: Yeah, I mean, I just want to make sure because I understand what the ID, whatever PA or whatever their acronym is saying, but they then go back and note that there is one producer who is marketing flux calcined to the cosmetic industry and that producer may be supplying the vast bulk of diatomaceous earth that goes into cosmetic industry. I'm just curious as to why all of our skin and eye studies were done on flux calcined?

Dr. Dan Liebler: Yeah.

Dr. Donald Belsito: If that's not what's being used.

Dr. Dan Liebler: Yeah. You know, I think this goes back to our concern from the silica report, the way we sort of looked at it with the silicates, we kind of had some synthetic, amorphous was good, natural mined, crystalline bad and we have let that kind of seep into our way of looking at all the other silica derived ingredients. And you know, we just mentioned it with the zeolites and now the diatomaceous earth. And I think our assumption that the crystalline content of diatomaceous earth is toxicologically equivalent to the crystalline content of silicates. That assumption may not be valid, and in fact all the data we have suggested it's not.

Dr. Donald Belsito: Paul, Curt?

Dr. Paul Snyder: Well, I had put on Page 36 of the report, the draft discussion that last paragraph there, I mean. My take was that, you know, as pointed out by the Council that, you know, it's not a method of manufacture because it's mined, it's sourced. So, I said just quote the IDPA stating only natural diatomaceous earth should be sourced for cosmetic to knock the recommended use, calcined or flux calcined for the crystalline silicon purity. I mean it's again; I don't think there's any room for wiggle there, I mean. Crystalline silica is it? Impurity is not a good thing.

Dr. Donald Belsito: Or we have chronic infection.

Dr. Paul Snyder: So I took more of a hard stance like the IDPA.

Dr. Donald Belsito: We have chronic inhalation studies with flux calcined. Right. I mean.

Dr. Dan Liebler: PDF 32.

Dr. Donald Belsito: Thanks. We have.

Dr. Dan Liebler: Acute.

Dr. Paul Snyder: Yeah, we don't have it. We don't have it. We don't have any carc studies. We don't have any inhalation, chronic, no.

Christina Burnett (CIR): On PDF page, yeah 32, 28 day study. You have a two year study.

Dr. Paul Snyder: Yeah, that's what that's not very long study for four.

Christina Burnett (CIR): You have a 2.

Dr. Paul Snyder: Crystalline silica.

Christina Burnett (CIR): You have a two year study for a flux calcined. In rats.

Dr. Donald Belsito: And we have a guinea pig 1.5 years. Right.

Dr. Paul Snyder: Yeah. I just. I just was taking what the IDPA, I mean they're the experts and they're, if they're recommending not to use it due to crystalline silicon purity. But I guess if we're going to, we'll have to make that very clear in the discussion that it cannot contain any.

Dr. Dan Liebler: I mean it, the safe, you know, sort of the conservative approach would be to do it as Paul just suggested in the discussion and say that you know, that cosmetic ingredient, cosmetic products will all be formulated with natural diatomaceous earth, not calcined or flux, calcined, diatomaceous earth. And not try and make a safety conclusion of around flux calcined or calcined, based on the available data because, you know, it's consistent that it's... that there's not a hazard. But, you know, concerned, concerned experts could differ in terms of the level of risk.

Dr. Donald Belsito: Yeah, I mean, and if you look at those long term studies, it states, and I'm not sure I noted this, what this means... 100% pure flux, calcined, diatomaceous earth? So does that mean they checked it ahead of time to make sure it was free of silicates? Do you see what page 30? PDF 32?

Dr. Paul Snyder: Probably doesn't, because if you go under the impurities section under composition, impurities on page 31. It says, flux calcined diatomaceous earth is used in the finished products. That concentration below 10%. It has a respirable crystalline silica content of less than 1%.

Dr. Donald Belsito: Right. So, I mean we can say that we appreciated the negative long term inhalation studies with the calcined, with the flux calcined diatomaceous earth, but noted that these were 100% pure, which cannot be guaranteed in cosmetic material necessarily or something to that effect.

Christina Burnett (CIR): And I also want to note that the natural DE can still have crystalline silica. It has a range of .1 to 4%. We do have two hands raised. So if you want to move to that.

Dr. Dan Liebler: Yeah. Could shed some light.

Thomas Gremillion (CFA): I had a clarifying question. This is Thomas. You're talking about crystalline silica being an impurity in the flux calcine. It says the crystalline silica content of a calcined diatomaceous earth is point 1 to 4% cristobalite content of calcined flux products. 20- to 28, between 40 and 60 for flux calcined. Is cristobalite content the same as the crystalline silica content. Is it? It's 60% of some of the flux calcined products is crystalline silica. Or am I misunderstanding that?

Christina Burnett (CIR): Yeah, I think they measure, the, the crystalline silica in terms of the cristobalite.

Thomas Gremillion (CFA): So they're synonymous.

Christina Burnett (CIR): Not quite because crystalline silica can be different. I want to say species but just like you'd have different amorphous silicas, you can have different crystalline silica. In terms of rock form.

Thomas Gremillion (CFA): I guess is the 40 to 60% content. It, you know. That the Element that the material that that poses the elevated risk hazard like crystalline is somewhere crystalline silica or it's. I'm just trying to understand what level of impurity in the flux calcined product says. It's like crystalline silica.

Shripal Sharma: So perhaps I could clarify. Can you hear me?

Dr. Donald Belsito: Yes.

Dr. Dan Liebler: Sure.

Shripal Sharma: So my name is Shripal Sharma and I work for Emeris as Production Director. And we are the supplier of a natural diatomaceous earth. And calcined, flux calcined... Diatomaceous earth to the world, you know, we had plants all over the world. And so, what I'd like to clarify the difference between natural DE and flux calcined DE. Is that natural DE's more or less amorphous silica natural with some impurity of crystalline silica? Yeah, like you know, mention .1224%. When it comes to how it is manufactured. You know it's calcined in the presence of a fluxing agent, which is sodium carbonate. So sodium carbonate is used as a fluxing agent. And calcination, at a high temperature. That converts some of the natural DE into flux calcined DE. Those amorphous, crystalline, amorphous silica particles are converted into cristobalite. And therefore the cristobalite content depending on the- on the manufacturing conditions could be somewhere between 30 to 60 to 70%. Ah, that's where the high level of crystalline silica in the form of cristobalite increases in flux calcined DE products.

Dr. Dan Liebler: Can you? Yeah. As far as it goes. On one other piece of information would be helpful is how does cristobalite compared to crystalline silica like in quartz for example?

Shripal Sharma: Does that help? Yeah, in it, that's a good question. I think OSHA back in 2015, I believe. They come up, came up with this, this regulation on occupational exposure limit. And prior to 2015, there was a different occupation exposure limit for quartz and cristobalite. Cristobalite it was .05 and 4 quartz it was .1. At that time, I think back in 2016, if I recall correctly. They lowered the quartz content exposure limit to .05. With the rationale that from a scientific understanding standpoint, there is no difference in toxicity. Of cristobalite versus quartz. In other words, they are they behave the same way when it comes to their toxicity.

Dr. Dan Liebler: So if I'm, if I'm not mistaken, we have considerable concern about crystalline silica in the form of quartz. For inhalation tox. And if cristobalite is comparable based on what you're saying in terms of, the threshold level? Then we - and cristobalite is a major component of flux calcined DE. Then we have a problem with flux calcined DE in terms of toxicity hazard. Don't we?

Shripal Sharma: Well, it depends on it. It depends. You know how a product is consumed.

Dr. Dan Liebler: Yeah.

Shripal Sharma: Whether it is in a, in a, in a powder form, and how it is, how it is applied, and how long it is, it is exposed. So, for example in our plant, you know we have employees working there, you know, 8 to 12 hours a day. And you know, we have been in the business since the early 20s and 30s and a number of epidemiology studies have come out, you know, over the last 70-80 years. And the one thing I can tell you is that you know, even though our employees are exposed to both forms of DE natural and flux calcined. You know, we don't have any, any, you know, silicosis cases. And other plants...anywhere in the world, I would say. Ah, or the reason is that we have a very good, you know, just management program through engineering control measures and use of proper PPE. That's not to say that, you know, every operation is different, and so what we see in our operation doesn't mean that it's going to be, you know, translate it or transfer into a different scenario. But when it comes to cosmetic, if I may say. You know, it's a very small level of DE either fluxed calcined or natural DE that goes into cosmetics.

Dr. Curtis Klaassen: They keep.

Shripal Sharma: And when it comes to flux calcined DE, the majority of that, as far as we know, goes into the mask. The facial mask and that's applied in a paste form. So, although overtime, it's get dried naturally.

Dr. Dan Liebler: Yes.

Shripal Sharma: But when it comes to removing that face mask after it is dry, you know, we again, this is our understanding. We don't expect that, you know, a lot of that dried material is going to come out in the form of powder and cause an inhalation risk.

Dr. Dan Liebler: So I think that our problem with flux calcined and calcined is the fact that it is a form of crystalline silica and we're concerned about crystalline silica because, at least in the case of quartz, crystalline silica. There are data demonstrating inhalation toxicity and even if it's set, you know much higher concentrations that might be in cosmetic products. Our problem is to establish the safety under relatively conservative options, so... We don't have any data right now to suggest that flux calcined DE produces any toxic effects. Unless I'm overlooking something, but we still have this concern about crystalline forms of silica as associated with inhalation tox in some crystalline forms such as quartz. And so, what you're telling us is helpful, except for the part where you point out that quartz and cristobalite have similar safety thresholds.

Shripal Sharma: That's correct. And one thing, if I made add and hopefully it helps you, is that you know, natural DE is...that's what the industry is promoting for cosmetics right now. Ah, we're not promoting the flux calcined. And the reason we missed it, you know, during the June of last year, communication that you received from IDPA. Because that's what, you know, majority of the members, you know, communicate it to IDPA including our company. And later on we learned, you know, through our European colleagues, European sources that you know, calcined DE is more like a legacy use. Ah, and because of legacy use, you know, continuing to sell that, you know it'll be 10 and a lower quantity. But when it comes to promoting? Ah, and, you know, promoting any, any that sort for cosmetic it's only the natural DE that the industry continues to promote.

Dr. Dan Liebler: Yeah.

Shripal Sharma: Although flux calcined DE to a limited extent remains in the market.

Dr. Dan Liebler: Yeah, it's helpful to know what's being promoted because it points to the future. But for us, we basically have to go on what's available to cosmetic ingredient producers now and what are they buying and using. And so, I appreciate you being, you know, forthright with us to explain that that's still can be purchased and it still is sold, and it still may be incorporated into cosmetic ingredients, and that leaves the ball in our court as to how we deal with it in our report.

Dr. Donald Belsito: Also explains why most do they all of the sensitization and ocular data was done on the flux calcined because we just heard that, so it's used in the masks. It's legacy.

Shripal Sharma: Yeah, I don't know that one thing that I could perhaps share with you is that flux also, and DE, registered in under each in Europe. Because flux calcined DE is made with the fluxing agent sodium carbonate, and because of that using fluxing agent under each, it is considered a not a, not a natural product. It's a synthetic product. And when it comes to REACH requirement, any synthetic products should be registered under each and in order for the industry and our company specifically to register that back in 2010. I believe under each you know all those studies had to be included in REACH dossiers.

Dr. Dan Liebler: Thanks, Doctor Sharma, for chipping in with additional information for us. We appreciate it.

Dr. Donald Belsito: Yes.

Shripal Sharma: Thank you.

Dr. Donald Belsito: So where does this leave us, folks, with these documents?

Dr. Paul Snyder: It would seem to me... It would seem to me that they're safe as used. Except for when there's a potential for inhalation. And if there's potential for inhalation, then that we have to have discuss the crystalline silica content like we did before, right?

I mean, the mask doesn't bother me. Now based upon that, and then that's also why...like you said, Don. That's why they did all the sensitization irritation data because they're applying that wet to the skin and if there's no potential for irritation then the

calcined or flux calcined could be used safely that way because that's not a source for inhalation, but if there's and then we can kind of merge the IDPA stating that only natural to be sourced and should not contain crystalline silica impurity.

Jay Ansell (PCPC): And that's about where we net it out as well.

Dr. Paul Snyder: Yeah, I think that seems reasonable.

Dr. Donald Belsito: OK, I'm not sure that I followed all of your ins and outs there, Paul. So, you're saying that the flux calcined? Can be used as long as it's not respirable.

Dr. Paul Snyder: As long as you...there's no potential for incidental inhalation. And those products in which it is, formulated to be used where there's a potential for inhalation, only natural DE should be sourced for cosmetics and should not contain crystalline silica impurity.

Dr. Dan Liebler: Are you thinking of that in the in the discussion or the conclusion, Paul?

Dr. Paul Snyder: I think it would be in the conclusion actually. I mean, that's kind of where I was at before, but when I thought I did not..I better understand this after Doctor Sharma's explanation there. But I thought we shouldn't say that only natural DE should be sourced for cosmetics and not recommend the use of the calcined or flux calcined based on crystalline silicon purity.

For those masks, it would appear to be that either applying them or taking them off there is no risk for incidental inhalation.

Dr. Donald Belsito: Or we heard that even natural DE has a certain level of silica, right, Christina?

Christina Burnett (CIR): Correct.

Dr. Paul Snyder: Yeah, but we're...but we're saying we could...we're saying it shouldn't contain it. And then the flux calcined. He said it when they flux it, but a lot of the amorphous goes crystalline, and then they get an increase crystalline silica content. So that's a greater risk for increasing the crystalline silica content. So that should not be used in ones that are formulated where there could be incidental inhalation. That's what I wrote.

Dr. Donald Belsito: OK then.

Dr. Dan Liebler: I think that we can do kind of along the lines of what Paul says. In which we have...as long as it's safe for use, safe as used. As long as any flux calcined or calcined are used in products that don't have potential for inhalation, is that correct, Paul?

Dr. Paul Snyder: Yes.

Dr. Dan Liebler: And in the discussion we can clarify that a bit more.

Dr. Paul Snyder: It appears to be that the flux in the calcite not calcined flux or flux calcined are for of the masks specifically.

Dr. Dan Liebler: And I think we feel comfortable perhaps that the mask,s or at least our team, if Kurt agrees, that the masks don't present a potential for inhalation because of the way they are applied and taken off.

Dr. Donald Belsito: Yeah.

Dr. Paul Snyder: Yeah, that's what I think.

Dr. Dan Liebler: What do you think on that?

Dr. Curtis Klaassen: (inaudible)

Dr. Dan Liebler: OK.

Dr. Donald Belsito: So Paul, you're saying the conclusion would be safe as used as long as the flux or flux calcined DE has no potential for inhalation, is that correct?

Dr. Paul Snyder: Correct.

Dr. Donald Belsito: There's flux or flux calcined DE?

Dr. Paul Snyder: Well, no, it would be...It would be natural diatomaceous earth is safe for use? With the crystalline silica content to be what X or zero? And calcined or flux calcined? Diatomaceous earth is safe for use in products in which there is no potential for inhalation, incidental inhalation.

Dr. Donald Belsito: So natural DE for use of crystalline silica is what?

Dr. Paul Snyder: Whatever. What did we say before? Did we say 0 on when we did the crystalline silica? Before what? Where did we? Did we have a level? We had lots of...

Dr. Dan Liebler: We did.

Christina Burnett (CIR): No, it did not.

Dr. Dan Liebler: I think we, I think we get into trouble if we try and put a number on that Paul? If we try to restrict.

Dr. Paul Snyder: Well, how did we handle before?

Dr. Donald Belsito: We said for silicates, Paul, the expert panel concluded the following 24 silicate ingredients are safe in cosmetics in the present practices of use as described in this safety assessment, when formulated to be nonirritating, with the exception that the available data are insufficient for the use of the naturally sourced silicate ingredients in products that may be incidentally inhaled.

Dr. Paul Snyder: Well then, that's what we need to do. We have to be consistent because it's the same issue. Crystalline silica.

Dr. Dan Liebler: So we could just flip that. We could use the same construction. Like instead of...so we lead with panel concluded that natural DE is safe as used, except in products containing flux calcined and calcined DE.

Dr. Donald Belsito: I mean, but we know that natural DE can also have it. So why don't we just include them all - natural DE, flux, and flux calcined are safe for use?

Dr. Paul Snyder: Yeah.

Dr. Donald Belsito: When? Three of these are used in products. Yeah. Pose no risk of incidental inhalation.

Dr. Dan Liebler: Yeah, that would be the simplest. Just say all of them.

Dr. Paul Snyder: Yeah.

Dr. Dan Liebler: Are safe as used except under certain circumstances, so in uses which may have incidental inhalation.

Dr. Donald Belsito: But there are some inhalation uses now, Christina?

Christina Burnett (CIR): Yes. Powders. And we don't know the concentration on that. And we see.

Dr. Paul Snyder: Sprays.

Christina Burnett (CIR): That's right. I'm sorry I'm scrolling the wrong way.

Dr. Paul Snyder: No sprays and powders so...

Dr. Donald Belsito: Well, we...

Dr. Paul Snyder: So we definitely, we can't say those are safety issues because we don't know some concentration of use.

Christina Burnett (CIR): Well, we don't know...

Monice Fiume (CIR): Sprays.

Christina Burnett (CIR): We don't know specifically if they are used in sprays. That's those categories that where it is in a skin product, but we don't know if it's a liquid or a spray.

Dr. Paul Snyder: Wow. Yeah, I see it right? Yep.

Christina Burnett (CIR): So but we have in the VCRP 5 uses in like a face powder.

Dr. Donald Belsito: So, I mean, we know from the chronic inhalation studies on flux calcined DE that it can be made 100% pure, which presumably means free of silicates, right? I mean there are inhalation studies - long term or negative on those?

Dr. Dan Liebler: Right.

Dr. Donald Belsito: So natural DE, flux or flux calcined DE are safe in cosmetics in the present practices of use and concentration described in this safety assessment. When formulated to be silicate free. Or when used in products where the potential for incidental inhalation does not exist. Something like that.

Dr. Paul Snyder: Yeah.

Christina Burnett (CIR): So. I do want to point out in the conclusion of the silicates report. Where we have the...because we have that insufficiency on incidentally inhaled products. When we were in the discussion, we have the, you know, data needs. And the two data needs for the silicates was composition and impurities data, especially quantification as crystalline silica and concentration of use of naturally sourced silicate ingredients or negative repeated dose inhalation data on naturally sourced silicate ingredients. In the case of DE, have at least the second point.

Dr. Donald Belsito: Just a second. Could you go over that again, Christina? I didn't follow. I was typing...the second point being?

Christina Burnett (CIR): That you were asking for a negative repeated dose inhalation data on naturally sourced silicate ingredients, so you have...

Dr. Donald Belsito: Right.

Christina Burnett (CIR): Negative repeated inhalation data for diatomaceous earth.

Dr. Donald Belsito: Right, and it was flux calcined, but it was 100% pure. Which I took to mean silicate free.

Christina Burnett (CIR): Then...

Dr. Donald Belsito: So this is what I just wrote... And if we don't like it, we can change it. Natural DE, flux, or flux calcined DE are safe in cosmetics in the present practices of use and concentration described in the safety assessment when formulated to be silicate free or used in products without risk of incidental inhalation. I don't know what people think about that.

Dr. Dan Liebler: Silicate free parts is where I'm hung up. I mean there are...in composition impurities we do have notation that crystalline silica content of uncalcined is .1 to 4%.

Dr. Donald Belsito: Right, I understand. But if you go back to the chronic inhalation studies?

Dr. Dan Liebler: Yeah.

Dr. Donald Belsito: This is PDF.

Dr. Dan Liebler: 32.

Dr. Donald Belsito: Yeah. If you look at. The last paragraph it says.

Dr. Curtis Klaassen: I mean (inaudible) it shouldn't.

Dr. Donald Belsito: It says inhalation sources in, in inhalation studies and no NOAEC could be determined 28 day of 100% pure flux calcined diatomaceous earth. Which I assume meant silicate free, because then they use that same product, and they did these two year, 1.5 years studies that were all negative.

Dr. Dan Liebler: It's a different reference.

Dr. Paul Snyder: Yeah, but that was...that was the study that's set up the second study. So, they couldn't get, they didn't get an observed effect.

Dr. Donald Belsito: You're right.

Dr. Dan Liebler: No.

Dr. Paul Snyder: In a 28 day study. So then they did a two year study.

Dr. Dan Liebler: I see.

Dr. Paul Snyder: Yeah.

Dr. Dan Liebler: I mean, the thing is it says 100% pure. It is a (inaudible), I just have a hard time with that 100% pure where it says that there, but it's the composition impurities data that's provided to us shows that these, you know, these products have, depending on which product you're looking at, have anywhere from about .1 or less than 1% or .1 to 4% crystalline silicate contamination.

Dr. Paul Snyder: That's exactly data we don't have here, Dan. We don't know what the crystalline silica content of this test material was.

Dr. Dan Liebler: Yeah, right. Exactly that. Yeah, that's what I'm saying. It says 100% pure in that in that inhalation tox description. But I'm not sure what that means. I'm not sure that's the same criteria that are applied for the composition and impurities data that we have.

Dr. Paul Snyder: Yeah.

Dr. Dan Liebler: And I think this puts us in a position where we really can't define a limit, a limit, and we can't say none. Because that's, you know, that's practically non achievable so...

Shripal Sharma: If I may clarify one more point. Which is when you say 100% pure. What that normally means is that there is no blend of that product with the other forms. So, for example 100% pure flux calcined DE does not mean that it is crystalline silica free or it does not say what the content of the crystalline silica is. All it says is that that product was used alone by itself without any blend with the other forms of DE, such as calcined DE or natural DE.

Dr. Dan Liebler: Yep.

Dr. Donald Belsito: OK.

Dr. Dan Liebler: Thank you.

Shripal Sharma: You're welcome.

Dr. Donald Belsito: So where are we? Both natural, calcined, and flux calcined can contain silicates.

Dr. Curtis Klaassen: Correct.

Shripal Sharma: That...

Dr. Paul Snyder: Today there's insufficient evidence.

Dr. Curtis Klaassen: I think we are. I think we're going a little over protective on this silica. And the concern about, you know, lung cancer. Ah. You know there, you know, even though there's some silica in there. There's still a dose response, and in fact we're breathing this stuff every day. Ah. And when we have this study here. I mean, this is a pretty good inhalation study compared to what we usually see. And with any compound, I guess I'm not so concerned. With a small amount of silica that might be in these compounds, so that's where I'm at.

Shripal Sharma: And if I may add to that.

Dr. Curtis Klaassen: It's like...

Shripal Sharma: You know, 12 to 16% of Earth's crust is actually made of crystalline silica. And the beach that we enjoy every day is actually, it's all crystalline silica. So it depends. In our house, someone is exposed - the duration of their exposure, the level of that exposure to have inhalation in a health effect, if you will, again, you know we, we've been manufacturing all three forms of the natural, calcined, and flux calcined and (inaudible)...

Dr. Curtis Klaassen: Yeah.

Shripal Sharma: Now, 70-80 years and our employees are exposed to it, you know we handle it, you know, with care, making sure our employees have the proper PPE on them and the dust generation in the plant is minimized. Through our safety data sheet, you know, we communicate to our customer how best to protect them, you know, from any exposure to dust and crystalline silica in particular and respect the local laws, etc. So, you know, needless to say, the crystalline silica, (inaudible) pieces is not, you know, hazardous if, you know, you can manage how best to control the exposure. And reduce that exposure, the duration of it, but also the extent to which it is, someone is exposed.

Dr. Dan Liebler: So I tend to agree with Curt, but not for...there isn't much as...I appreciate Doctor Sharma's last comments, not for any of those reasons but more because our data in the report show exposures to calcined and flux calcined with no toxicity. And considering flux calcined and calcined have much higher levels of crystalline silica in the diatomaceous earth. Then do...natural. We can argue till the cows come home about how much diatomaceous earth...crystalline silica...may be present in the natural stuff.

Dr. Curtis Klaassen: Right. And.

Dr. Dan Liebler: So we've got no toxicity data to point to a problem with any of it. So that's why I...that's why I'm inclined to not put restrictions on these, because I don't think we have any data that supports it.

Dr. Paul Snyder: But I think it's pretty compelling that the IDPA sent us a letter saying it...only natural DE should be sourced for cosmetics and not recommended to use calcined or flux calcined due to crystalline silica purity.

Dr. Dan Liebler: Well, that is...it can be interpreted that way, Paul. I agree. And in fact, Doctor Sharma's comments about, you know, we're trying to move the marketing towards using natural, you know, be interpreted as sort of sending the same message. The problem is our data just don't necessarily support it because we don't have a hazard or we don't have a, you know evidence of a...of a toxicity in the data that we cited the report. So I agree with your point, Paul, but I just don't see how we can support a restriction in the conclusion based on the data that we have in the report.

Dr. Donald Belsito: Yeah. Hi, Bart.

Dr. Bart Heldreth (CIR): I just wanted to suggest an alternative, maybe meeting in the middle, having a very simple safe as used conclusion, but the detailed explanation in the discussion stating something to the effect that manufacturers should take care to formulate products which may be incidentally inhaled so not to contain significant levels of crystalline silicon to cause concern or something to that effect because, as Dan said, the data in our report did say safe.

Dr. Dan Liebler: I would support that approach.

Dr. Donald Belsito: So. OK, can someone read that conclusion?

Dr. Dan Liebler: Basically, safe as used for the conclusion.

Dr. Donald Belsito: So.

Dr. Dan Liebler: With no restriction and then the discussion talks about formulation. Manufacturers are advised to formulate to minimize the...the content of crystalline silica in products that may have risk of inhalation.

Dr. Donald Belsito: So that goes in the discussion.

Dr. Dan Liebler: Correct. Yeah, I think you're reporting on this tomorrow, Don.

Dr. Donald Belsito: Right. It's crashing... We have a chronic inhalation and some discussion. We're just going to add...try to limit the amount of silicates is that where we said?

Dr. Dan Liebler: Crystalline silica and in products that might have incident...incidental inhalation.

Dr. Donald Belsito: Well.

Dr. Dan Liebler: We have another hand up from a Richard Brown.

Richard Brown (Guest): This is Richard Brown. I've just wanted to mention that there's one critical factor here that has not yet been discussed, and I think should be included in your recommendation, and that is that in order for crystalline silica, inhaled crystalline silica to be an issue, it has to be of respirable size. There was a comment made earlier about couple of comments about crystalline silica being 12 to 16% of the earth crust. Then the beach sand that we all enjoy when we get to the ocean being significantly composed of crystalline silica. The issue there is it's not of respirable size, so there is no inherent risk regardless of the nature of the crystalline silica. Another aspect of this that is important is that there's a significant body of published peer reviewed information that says...crystalline silica in order to be an issue, a significant issue, has to have an abundance of reactive moieties, reactive sites on its surface. Which primarily occurs through industrial processes that grind, crush, or break crystalline silica particles. And that these reactive sites are significantly (inaudible) when environmental contaminant, or other materials, come in contact with these surfaces. When crystalline silica is a component in a blend used for cosmetics, for example, those surfaces get covered with materials that will result in quenching of these sites. They also typically cause respirable size crystalline silica particles. Some do become parts of agglomerates that exceed respirable size so they're no longer a problem. So I would recommend that those things be considered in whatever recommendation you come up. Thank you.

Dr. Dan Liebler: Thanks. We've talked about those issues a lot on this panel and when we reviewed lots of potentially inhaled cosmetic ingredients and we have our inhalation resource documents that discuss those issues. We agree that they are relevant. Sounds like we got to plan here and on.

Dr. Donald Belsito: Well, I mean the last I got is that the conclusion is that they're safe in cosmetics, in the present practices of use and concentration and the discussion simply says limit crystalline silica products that may be inhaled.

Dr. Dan Liebler: Right. I guess the question is, does our team support that?

Dr. Paul Snyder: Well, I guess my question to Bart is, is this inconsistent with the silica silicates report where we said they don't...were insufficient for incidental inhalation because it's the same, it's the same impurity we're discussing.

Dr. Bart Heldreth (CIR): The main difference I see here is what Dan pointed out is we have inhalation studies using the flux calcined version of the ingredient that's supposed to contain the most silica, and there's no safety concern. To me that that makes this report a little different than the silicates report. We couldn't say that those ones in the silicate report were safe as use because we didn't have the data to support the inhalation tox.

Dr. Dan Liebler: I think one of the big unknowns, Paul, is that the chemical identity of any crystalline silica that may contaminate diatomaceous earth compared to silicates. And they may not be the same. Particularly in terms of the hazard that they present for pulmonary tox.

Dr. Paul Snyder: OK.

Dr. Donald Belsito: Anything else? I mean, are we going with the conclusion and discussion that I just stated or where are we? Paul, you look like you're not quite happy.

Dr. Paul Snyder: No, no, I'm fine with it. I just want to make sure that we don't end up with a...with 2 reports that are dealing with the same impurity but have different conclusions, and so I think we just need to be clear in the discussion as to why we're concluding that and I think that we can do that. I think we can craft that that we do have some data. And so I'm, I'm still not 100% convinced that we have everything we need. I mean, do we have particle size distribution data?

Christina Burnett (CIR): On PDF page 30. You have particle size distribution for flux calcined diatomaceous earth – coarse, medium, and fine grade materials at less than 90, like the percentage distribution at 90 micrometers and a 100 - or ten less than. 10. Sorry.

Dr. Paul Snyder: Less than 10. Yeah, it's pretty. Pretty high, less than 10, 60%.

Christina Burnett (CIR): The full table is on... there's a table that describes the full amount on, PDF page 37.

Dr. Donald Belsito: Yeah. And then of course, we don't know the particle size in the inhalation studies.

Christina Burnett (CIR): You do on the guinea pig one.

Dr. Donald Belsito: What was that, Christina?

Christina Burnett (CIR): You had the particle size distribution on the guinea pig inhalation study.

Dr. Donald Belsito: What was that distribution?

Christina Burnett (CIR): .Point four five micrometers to greater than 10.

Dr. Donald Belsito: It is small.

Dr. Dan Liebler: Yep.

Dr. Curtis Klaassen: Yeah, it is.

Dr. Paul Snyder: Yeah, we don't, we don't know, but that's just the range. We don't know what the, what, how many, how much, what percent was less than 10 microns. It's just that there's a range to .45 microns to greater than 10 microns, which is very informative. Well, we could discuss tomorrow.

Dr. Donald Belsito: Yeah, I think we need to move on. It's 2:45 and we still have some difficult ones. So, what we're recommending is safe as used and limit silicates and other products that could be inhaled in the discussion.

Dr. Dan Liebler: Yep.

Dr. Donald Belsito: OK. Very good. We'll see what the Cohen team comes up with.

Cohen's Team Meeting – March 7, 2022

Dr. David Cohen: OK. Well, hold on, let's move on. Alright, so the next one is diatomaceous earth. Just pulling it up. So. Right at the September meeting, we issued an IDA. We wanted clarification for the types of diatomaceous earth, whether natural, calcined, or flux calcined, method of manufacturing, composition and impurities. Oh, in the...in our first load of information we were told that only natural DE, not calcined DE was used in cosmetics, although in the second wave, a letter shared with us suggested that a company has sold flux calcined DE to one cosmetic manufacturer. And we got a letter from the International Diatomaceous Earth Producers Association clarifying a number of issues that I thought their responses were reasonable (inaudible) they indicated it's often in an aqueous formulation, rinsing off the issue with flaking and mudmasks was discussed. And I'm wondering what the group's thoughts are on silicates here as well. So. Ron?

Dr. Ron Shank: Oh. I thought the report was in good shape. So as far as the discussion is concerned, I would just make it very similar to the one in the silicates report. Which you've done very, very nicely. So just make the discussion here in harmony with the discussion in the silicates report. And then the conclusion would be that they are safe as used in the present practices of use and concentration when they are not expected to be incidentally inhaled. And when formulated to be non-irritating. There is inhalation data on page 32, several repeat dose inhalation studies on diatomaceous earth and those produced negative results. So, I would go with safe.

Dr. Wilma Bergfeld: Alright. Do you think that there needs to be a statement in the discussion regarding heavy metals? Silica. Those should be avoided. We have a heavy metal statement.

Dr. Ron Shank: Yeah, that's fine.

Dr. David Cohen: Yeah.

Dr. Wilma Bergfeld: And we had some mention of that in Wave 3.

Dr. Ron Shank: Just the boiler plate, right?

Dr. Wilma Bergfeld: Yeah.

Dr. Ron Shank: OK.

Dr. Thomas Slaga: Yeah.

Dr. Wilma Bergfeld: And you're referring to the inhalation because with silica, too, and there wasn't a product that could be inhaled.

Dr. Ron Shank: Yes.

Dr. Wilma Bergfeld: And the inhalation boilerplate, would that go in here to the discussion?

Dr. Ron Shank: We have negative data. So I don't think we need the boiler plate.

Dr. Wilma Bergfeld: OK.

Dr. David Cohen: Ron, can you just help me with non irritating?

Dr. Ron Shank: OK, so let's see where that came from? I guess that came from the ocular.

Dr. David Cohen: Just ocular? What page is he on?

Dr. Ron Shank: Well, let's see.

Dr. Bart Heldreth: PDF Page 34 has ocular irritation studies.

Dr. Ron Shank: Yeah.

Dr. David Cohen: Moderately irritating at 10% dilution. OK. Yeah. OK, well.

Dr. Thomas Slaga: So like you were suggesting to the conclusion, should have non irritating in it?

Dr. David Cohen: That's what Ron suggests... that, I didn't have that in mind.

Dr. Ron Shank: That's what I suggested, but...

Dr. Thomas Slaga: Ah.

Dr. Ron Shank: That's what I suggested. But if you don't want it, I'm not going to push it.

Dr. Thomas Slaga: Yeah, I didn't think we needed it.

Dr. Ron Shank: OK.

Dr. David Cohen: I didn't have it there either. That's why I asked you. Ron, what?

Dr. Ron Shank: OK, it was from the ocular so...

Dr. Thomas Slaga: But I mean we don't, we have it in the discussion.

Dr. David Cohen: OK. And so it's safe as used, with the exception when it's going to be incidentally inhaled. Is that it?

Dr. Ron Shank: I'm sorry, could you say that again?

Dr. David Cohen: So we're going to go, we're going to go out as safe as used. In the concentrations in this safety assessment, with the exception, with the exception that...

Dr. Ron Shank: Yes. Yes.

Dr. David Cohen: It is insufficient. What is it insufficient...if it is incidentally exposed. Incidentally inhaled?

Dr. Ron Shank: Well, we have inhalation data and it's negative. So I don't think we need to put that in the conclusion. I had said that at first, yes, but thinking about that more. Oh. Because we have the inhalation data and it's negative I don't think we have to put in the caveat, formulated to be not inhaled.

Dr. David Cohen: But, but what about the...what about the notes suggesting that flux calcined DE might wind up in cosmetics? Now, as I think wave two or three, there's just a note, it's, it's very incidental that they said yeah we sold flux calcined DE to a cosmetic manufacturer. That's all we have. And so that crystalline component just it is an overhang for me.

Christina Burnett (CIR): You do have safety test data on that product. The manufacturer had submitted that data before IDPA confirmed it. So some of the highlighted data in the report is on the flux calcined and I believe in the impurities section they listed their crystalline silica content.

Dr. Ron Shank: That was for a sensitization.

Dr. David Cohen: No, no composition and impurities. It's on I think page 31.

Christina Burnett (CIR): Yes, it's on page 31 in middle of, it's towards the end of the composition impurities. It's highlighted.

Dr. Ron Shank: Yes.

Christina Burnett (CIR): That says the supplier has reported that flux calcined diatomaceous earth is used in finished products, that concentrations below 10% and has a respirable crystalline silica content of less than 1% as cristobalite based on the size weighted relevance fine fraction method of (inaudible) analysis.

Dr. David Cohen: So are we being consistent with silicate document? If we're saying like, oh, less than 1% is OK. And then we were going through this whole discussion about undetectable or limits of detection or not we have no conclusion if it's respirable.

Dr. Wilma Bergfeld: I think you're right on. I think we have to be consistent and in that wave three, you do have mentioned of the suggestions from the committee that we there should be a suggestion to monitor heavy metals and crystalline silica.

Dr. David Cohen: So why don't we leave in the caveat? That it's insufficient for incidental inhalation? Unless I'm really asking for it for you, for your input guys. I got jammed up in these.

Dr. Ron Shank: OK. The inhalation data we have on diatomaceous earth was apparently done on non calcined material. These are used in hair dyes, nail polish and nail enamel, and face masks.

Dr. David Cohen: It's the face masks that that I have the issue with.

Dr. Ron Shank: OK, it's the same thing that if they do dry and flake, the flakes would be too large to be inhaled.

Dr. David Cohen: So. You're advocating for as safe as used.

Dr. Ron Shank: Yes. Yes.

Dr. David Cohen: And in the in the discussion...

Dr. Thomas Slaga: In theory.

Dr. David Cohen: Go ahead, Tom. Sorry.

Dr. Thomas Slaga: I said I agree with that. There is some inhalation data that Ron emphasized.

Dr. David Cohen: And we're dealing with the possible presence of crystalline silica simply in the discussion and what in the discussion are we saying?

Dr. Ron Shank: Well, if you're going to worry about the crystalline silica, then I would use the same discussion that's in the silicates report. They had, if it's present, its presence should be less than zero point 1% in the raw material.

Dr. David Cohen: I think our silicate conclusion was safe as used. We had formulated to be non-irritating for the silicates, with the exception that the available data are insufficient to make a determination of safety for the use of naturally sourced mined silicate ingredients and products that may be incidentally inhaled.

Christina Burnett (CIR): Correct. And I don't remember...in the discussion, I don't believe we discuss a limit on the crystalline silica because of the discussion of how it was difficult to limit and quantify, and the background ambient silica, and there's too many confounding factors.

Dr. David Cohen: Oh we went back and forth several meetings.

Dr. Ron Shank: That was taken out.

Christina Burnett (CIR): Yes.

Dr. David Cohen: Right. We don't have a, we don't have a concentration limit. We proposed in the...data are insufficient for incidentally exposed, incidentally inhaled silicates.

Christina Burnett (CIR): Correct.

Dr. David Cohen: Right. And by the way this, so that prosecution went on for a year about or so and we're coming right off of that one. On this one right and it just it, that's what, you know, got me wrapped around the axle was that there's discussion of crystalline silica. And, yeah, that some flux calcined DE can get into the pipeline of DE in cosmetics and how we how we going to address it?

Dr. Wilma Bergfeld: Definitely has been mentioned in the discussion, definitely asked to go there.

Dr. David Cohen: Yes, so.

Dr. Ron Shank: Well, it should be the same as in the silicate report.

Dr. David Cohen: OK, we, we, we.

Dr. Ron Shank: And in my copy of the silicates report, the concentration limit of less than .1 percent is still there.

Christina Burnett (CIR): In the discussion?

Dr. Ron Shank: Both. Both the discussion and the conclusion?

Dr. David Cohen: So.

Christina Burnett (CIR): That I don't think...that was the December copy.

Dr. David Cohen: Now wasn't the final, final, so I mean if the team...

Dr. Ron Shank: OK.

Dr. David Cohen: So if the Team is in agreement we can go out with a silicate like conclusion. And the Belsito team is presenting this one.

Dr. Ron Shank: Yes.

Dr. David Cohen: And if they come out as safe as used, we can bring up there are concerns about the crystalline silicates. Using face masks, and I understand the size of the flaking, and we can have that discussion.

Dr. Ron Shank: OK.

Dr. David Cohen: How does that that sound?

Dr. Thomas Slaga: Sounds good.

Dr. David Cohen: Reasonable.

Dr. Thomas Slaga: Yeah. Yep.

Christina Burnett (CIR): Would you like me to read the current paragraph in the silicates report that discusses that?

Dr. David Cohen: Sure.

Christina Burnett (CIR): So silicates used in cosmetics or solid inorganic oxides comprising in part silicon dioxide that can be derived from naturally occurring minerals or can be produced synthetically. The panel considered a method of manufacture of these ingredients i.e. whether synthetic or mined to be of significant importance to safety as synthetically derived ingredients are expected to have controlled silica crystalline material formation. The panel is of the understanding that only naturally sourced silica contains crystalline silica is a known cause of significant lung disease, including cancer. Thus, the available data are insufficient for determining safety for a formulations containing naturally sourced silicate. When the potential for incidental inhalation exists, the additional data needed to determine safety of naturally sourced ingredients i.e. potentially containing crystalline silica in cosmetics that may be incidentally inhaled are composition and impurities data, especially quantification of crystalline silica, and concentration of use of naturally sourced silicate ingredients or negative repeated dose inhalation data on naturally sourced silicate ingredients.

Dr. Ron Shank: OK. Thank you.

Dr. David Cohen: Does that change any of your thoughts?

Dr. Ron Shank: No, just so long as...

Dr. Thomas Slaga: No.

Dr. Ron Shank: And as it agrees with the silicates report.

Dr. Wilma Bergfeld: But it would agree if you put in the discussion regarding the fact that there were studies here allowing the amount of silica that is in this product to go forward as safe something on that order. Would it not? If you repeat a little bit about the crystalline silica? And then say that there were some long term pulmonary studies.

Dr. Ron Shank: Well, the inhalation studies done on diatomaceous earth didn't say anything about crystalline silica.

Dr. David Cohen: They were non calcined, right?

Dr. Ron Shank: And the inhalation studies were negative. What?

Christina Burnett (CIR): Right.

Dr. David Cohen: They were non calcined?

Dr. Ron Shank: OK.

Dr. David Cohen: No, no, I'm asking. They inhalation studies were done on non calcined DE.

Dr. Ron Shank: Oh. Let me find it.

Christina Burnett (CIR): So I will note that none of the diatomaceous earth products are synthetically produced.

Dr. Wilma Bergfeld: Amorphous (inaudible) says.

Dr. David Cohen: Right. Yeah, it's by its definition, they're naturally sourced, right?

Christina Burnett (CIR): Right. So. Right, right. They're processed and that's where the crystalline silica is produced is through the heating.

Dr. Bart Heldreth: Yeah, some of the studies for inhalation, both in the acute and the repeated, those are on flux calcined on PDF page 32.

Dr. Wilma Bergfeld: Yeah.

Dr. Ron Shank: Right.

Dr. David Cohen: Yeah, 100% pure flux calcined DE. So with that data make us not have, incidental inhalation warning on that? I mean, if we know this potential crystalline silica in it, and you have a study that says, yeah, we studied this and that was it. I mean, that's kind of what Christina had read in the, in the provision of the silicate document. Then maybe we don't need it.

Dr. Ron Shank: I agree. I don't think we need it. So now it's just safe as used.

Dr. Thomas Slaga: Yeah.

Dr. David Cohen: It's a long road to that.

Dr. Thomas Slaga: Yeah.

Christina Burnett (CIR): OK.

Dr. David Cohen: I'm making it.

Christina Burnett (CIR): So what is the discussion then? If it's not going to be...

Dr. Wilma Bergfeld: Could I just ask, can I ask a question about those are called pulmonary studies, not inhalation studies. And they were injections. The first one, is IV injection. And the second one is intratracheal study. Is that is that different than inhalation?

Dr. Ron Shank: Very.

Dr. Wilma Bergfeld: Yeah, I thought so.

Dr. Ron Shank: But on page 32.

Christina Burnett (CIR): You're. Yeah. And that's under the pulmonary response for the Other relevant studies. I think.

Dr. Wilma Bergfeld: Yeah.

Christina Burnett (CIR): So the other repeated dose studies are on page at the bottom of page 32. And there is a table.

Dr. David Cohen: OK. I'm so we will go out as safe as used, but in the discussion, we should mention that crystalline silica may be present particularly if it's, if it's in through certain processing. And that effort should be made to limit that.

Dr. Ron Shank: Well, we have the inhalation studies done with flux calcined diatomaceous earth, so for more than one. And they were negative. These were two years studies. Well, two years and 1 1/2 year.

Dr. David Cohen: Yeah, it took to me like it's a quandary. That's all because you have the data we asked for in the silicate paper. Yet we know that if this was a different situation where we had crystalline silica in something, we would want to limit inhalation, right? It's difficult, this one.

Dr. Ron Shank: It is.

Dr. David Cohen: But I think where? Christina, I don't know if you are satisfied with our discussion?

Dr. Ron Shank: No. She isn't.

Christina Burnett (CIR): No. I can make you flush it out more tomorrow once, once you combined with Dr. Belsito.

Dr. Bart Heldreth: You want to have something perhaps very similar to the discussion for silica, but then follow up with, however,

Dr. David Cohen: Yeah.

Dr. Bart Heldreth: After this data that fill that insufficiency it like I think the last sentence or two of the discussion that Christina read said, you know we are insufficient for these inhalation studies. Well here we have them.

Christina Burnett (CIR): Yes.

Dr. Ron Shank: I like that. I have the wrong copy of the silicate reports so.

Dr. David Cohen: OK. Alright so Ron and Tom, you'll be on standby tomorrow if we if we heat up on the discussion.

Dr. Ron Shank: Yeah. Will be there.

Christina Burnett (CIR): OK.

Dr. Wilma Bergfeld: I did just mentioned in the clinical studies, which are not particularly valid, but in the preparation of the DE there is crystalline silica.

Dr. David Cohen: Yes. Yeah, I'm not sure anyone is going to argue the presence or absence of the potential for crystalline silica. It...it's the issue of how we are clear products we think might have crystalline silica in it that may be incidentally inhaled. Right? And then Ron's point is, yeah, what we asked for inhalation studies in those circumstances. And not only did we get them and they were negative, they use calcined DE to do those studies. So we...

Dr. Thomas Slaga: Right.

Dr. David Cohen: We got exactly what we asked for. And you know where I, I guess we're just still contemplating that but ah it's I think it's just it will be discussed enough ultimately.

Dr. Wilma Bergfeld: Agreed.

Full Panel Meeting – March 8, 2022

Dr. Don Belsito: Yeah. So diatomaceous earth. After the September 2021 meeting, we issued an IDA. We wanted clarification on the types of diatomaceous earth used in cosmetic products. That is whether they're natural, calcined, or flux calcined... manufactured types of diatomaceous earth used in cosmetic products, composition and impurities data including crystalline silica content on the types of diatomaceous earth. Or it could be used in cosmetics, we got a lot of data and we got a lot of comments come in. Wave two and wave three. Some good discussion yesterday from the manufacturing group, and based upon all of that, we also noted that there were two year inhalation studies on flux calcined. We were told that the manufacturers group is recommending only the natural be used, but we also noted that all of the dermal studies and the (inaudible) studies used flux calcined, but again we had very good long term inhalation data on them. So with all that in the long winded background, my team felt that natural diatomaceous earth, flux, or flux calcined diatomaceous earth are safe in cosmetics in the present practices of use and concentration as described in this safety assessment.

Dr. Wilma Bergfeld: That's a motion.

Dr. Don Belsito: That's our motion.

Dr. Wilma Bergfeld: It's not a second, David.

Dr. David Cohen: Yes, it's a second.

Dr. Wilma Bergfeld: Any comments or?

Dr. David Cohen: Yeah, we spent a lot of time going through the same things you did, Don. And the issue of the presence of crystalline silica came up. We wanted to harmonize with our previous silica report. But you mentioned all of that respiratory tox data that we had, which was good. And it seemed like the overwhelming use is natural DE and there was that one note that flux calcined was sold by one distributor to a cosmetic manufacturer. Yeah, that's we, we came to the same conclusion and we thought the IDPA had a lot of great points. And sort of assuaged many of our concerns, including the discussion of the masks. So Don, anything for the comments for further discussion about? Using natural DE over calcined or flux calcined or the pulmonary data was enough for you to just clear everything.

Dr. Don Belsito: Yeah, I mean we, I mean the pulmonary data was all on the inhalation data was all on flux calcined and, you know, between that and the concentration of use, you know, we just felt that these were safe as used. You know, we couldn't come up with a reason to say that they were not safe. We had the data as opposed to the silicates and all of the others, the clays. We're going to be talking about later. We have the inhalation data here.

Dr. David Cohen: I know, I know.

Dr. Don Belsito: You know we have the data that we need. And so that obviously needs to go in the discussion. I think it's a good point that they should also go on the discussion that the manufacturing group really recommends the use of only natural. You know, really our data needs were all met here. I mean, we got a lot of information.

Dr. David Cohen: We have nothing further.

Dr. Wilma Bergfeld: And now, do you think that that natural description of why they recommend only the natural should go in the discussion as well? I mean, just to say that.

Dr. Dan Liebler: They're not saying why.

Dr. Wilma Bergfeld: Then maybe you shouldn't say that.

Dr. Dan Liebler: We're just recommending it.

Dr. Don Belsito: Right.

Dr. Wilma Bergfeld: Right.

Dr. David Cohen: Does it influence crystalline silica formation when it's calcium calcined?

Dr. Wilma Bergfeld: The testing work (inaudible).

Dr. Don Belsito: Well, yeah, the heating supposedly increases the risk of forming crystalline silica. That's the issue. But even the natural DE has it, so I can add. I mean, it's listed as, you know, part of the composition.

Dr. Dan Liebler: So this is this illustrates one of the hazards of our use of terminology regarding crystalline silica. If you imagine for a moment that we had not gotten the silicates report until sometime later this year. In other words, if we had never done the silicates report, we probably would have a very different view of crystalline versus amorphous forms. And each of

these materials, diatomaceous earth, clays, silicates... They're broadly similar chemically, but there are differences between crystalline silica is not all crystalline silica are alike so that cristobalite apparently is not the same as quartz, even though they're crystalline silica. They fall into two the same category. But, you know, quartz, you've got, tox data that suggests inhalation is a significant health hazard, whereas with the flux calcined, which contains a higher percentage of crystalline diatomaceous earth. It's definitely safe, so you know, I think that we get spooked by the word crystalline in the context of any silicate type mineral ingredient, and we just need to be on guard against that. If we've got data, then we can decide, if we don't have data, we can't only be informed by being spooked.

Dr. Don Belsito: Doctor Sharma has his hand up.

Sharma Shripal: Yeah. Good morning one so...

Dr. Wilma Bergfeld: Perfect.

Sharma Shripal: Well, it's a couple of responses here. One is to answer the question about, you know, not all crystalline silica or similar from their health hazard standpoint, you know, and I mentioned that yesterday as well. During Dr. Belsito's meeting. You know, back in 2016, OSHA came out with what they call crystalline silica rulemaking.

Dr. Curtis Klaassen: (inaudible).

Sharma Shripal: And in the rulemaking, OSHA concluded that there is no difference in toxicity of quartz, which is a natural (inaudible) process of a...of DE. So OSHA concluded and since there is no difference in toxicity or health hazard between the two forms of crystalline silica. One is a natural quartz and the other is cristobalite produced during calculation prior to 2016. Cristobalite and quartz had different occupational exposure limits by set by OSHA. And for quartz it was .1 milligram per cubic meter. And for cristobalite it was .05. So cristobalite had had half the (inaudible) of quartz, but then they harmonize in 2016 to be the same based on the scientific evidence that there's no health hazard differences between the two forms of crystalline silica.

Dr. Wilma Bergfeld: Is that the clarification that we need, Don and David?

Dr. David Cohen: I think it was a nice summary.

Dr. Don Belsito: Yeah.

Dr. Wilma Bergfeld: OK, Thank you very much.

Sharma Shripal: Thank you.

Dr. Wilma Bergfeld: Sure.

Dr. David Cohen: Dan, one quick question. You made an interesting comment like about the timing of the silicate report. How do you think this? Adjudication of DE would have been different.

Dr. Dan Liebler: Well, I think that we might have, we might have, I think we might have come into this with, you know, sort of less prior suspicion directed it at the at the flux calcined. But you know in this, in this report we have data. And in the silicates report, we had no data.

Dr. David Cohen: Yeah.

Dr. Dan Liebler: And then we've got upcoming reports where, you know, we don't necessarily have the data, I think I've just noticed that, that the silicates report kind of spooked us, and we just need to be careful not to let that drive our thinking as much as the data.

Dr. Don Belsito: Thomas, you have your hand up.

Thomas Gremillion (CFA): Thanks, Dr. Belsito. I just wanted to check my understanding. So there's not. Then how I sent that is sufficient (inaudible) some kind of ignore this tension between flux calcined and the natural. I want to check my understanding on that because the percentage of crystalline was so high on the flux calcined product, so it says little surprised when it.

Dr. Don Belsito: Yeah, I mean it, there were, you know, studies from 1.5 to two years and different species and they...and exposed to much higher amounts than you would get from using cosmetic products and they were all clean.

Thomas Gremillion (CFA): OK. Thank you.

Dr. Wilma Bergfeld: They were needing a second on this proposed conclusion, David.

Dr. David Cohen: Guys, second.

Dr. Wilma Bergfeld: And Don, would you repeat your conclusion?

Dr. Don Belsito: I've already moved on. I mean, safe as used essentially.

Dr. Wilma Bergfeld: OK, OK.

Dr. David Cohen: Yeah, that's what he came in.

Dr. Wilma Bergfeld: And I think that we have all the comments that we need. How about with comments for the discussion?

Dr. Don Belsito: Christina Burnett has her hand up.

Dr. Wilma Bergfeld: Christina.

Christina Burnett (CIR): Yeah. So I just wanted to clarify some of the discussion points I wrote down between the teams yesterday. I have that there should be a note about the ocular irritation.

Dr. Wilma Bergfeld: Go ahead.

Christina Burnett (CIR): Do you still want that in discussion?

Dr. David Cohen: Hello.

Christina Burnett (CIR): It was. It was only one study that showed it.

Dr. Don Belsito: Must have come from the current team. I don't have a note on that, Christina.

Christina Burnett (CIR): Yes.

Dr. David Cohen: Ron to you. I think you might have mentioned it yesterday.

Dr. Ron Shank: It was just should it be mentioned?

Dr. David Cohen: I have to go back to the report.

Dr. Ron Shank: First came up. Should we have when formulated to be non irritating? Having the eye in mind. But as we discussed it, we decided that that was not necessary. Flux calcined diatomaceous earth was an ocular irritant, but at very high concentrations.

Dr. David Cohen: Yeah.

Dr. Don Belsito: Yeah. And the purity was not reported on that.

Dr. Ron Shank: Right.

Dr. David Cohen: That's what I have in my notes from yesterday. OK.

Dr. Ron Shank: So we decided we didn't need it to discuss it.

Dr. Don Belsito: Right exposure was 99.1% and the purity of that material was not (inaudible).

Christina Burnett (CIR): OK so...

Dr. Don Belsito: Work specified rather.

Christina Burnett (CIR): So leave that out at the discussion.

Dr. Ron Shank: Yes.

Dr. Wilma Bergfeld: Sounds like it.

Christina Burnett (CIR): I have inclusion of the heavy metals boilerplate.

Dr. Don Belsito: Yes, of course.

Christina Burnett (CIR): OK.

Dr. Don Belsito: Respiratory boiler plate.

Christina Burnett (CIR): And the inhalation, OK and...

Dr. Don Belsito: You don't have like...fine, yeah.

Dr. Ron Shank: Yes.

Christina Burnett (CIR): OK. And then there's language about using face masks not being a concern? Because the flakes are too big to be respired?

Dr. David Cohen: Flakes are large and they are removed with water.

Christina Burnett (CIR): OK. And then what was discussed just a few minutes ago?

Dr. Wilma Bergfeld: Uh huh.

Dr. Don Belsito: Yeah.

Dr. Wilma Bergfeld: And I understand that Ron had asked to have the percentage as respired removed from the inhalation boilerplate.

Dr. Ron Shank: Yes, thank you.

Dr. Wilma Bergfeld: I don't know.

Dr. Don Belsito: Not the boiler plate.

Dr. David Cohen: Here (inaudible).

Dr. Wilma Bergfeld: I'm... is there a resource document or comment?

Dr. Don Belsito: (inaudible).

Dr. Wilma Bergfeld: Yeah. OK, I don't think we'd called the question on this. All those opposed to this conclusion of safe.

Dr. Dan Liebler: Yes.

Dr. Wilma Bergfeld: Stating approved

Safety Assessment of Diatomaceous Earth as Used in Cosmetics

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The Expert Panel for Cosmetic Ingredient Safety members are: Chair, Wilma F. Bergfeld, M.D., F.A.C.P.; Donald V. Belsito, M.D.; David E. Cohen, M.D.; Curtis D. Klaassen, Ph.D.; Daniel C. Liebler, Ph.D.; Allan E. Rettie, Ph.D.; David Ross, Ph.D.; Thomas J. Slaga, Ph.D.; Paul W. Snyder, D.V.M., Ph.D.; and Susan C. Tilton, Ph.D. Previous Panel members involved in this assessment: Lisa A. Peterson, Ph.D. and Ronald C. Shank, Ph.D. The Cosmetic Ingredient Review (CIR) Executive Director is Bart Heldreth, Ph.D. This safety assessment was prepared by Christina L. Burnett, Senior Scientific Analyst/Writer, CIR.

ABBREVIATIONS

BAL	bronchoalveolar lavage
CIR	Cosmetic Ingredient Review
CHO	Chinese hamster ovary
Council	Personal Care Products Council
CPSC	Consumer Product Safety Commission
DART	developmental and reproductive toxicity
<i>Dictionary</i>	<i>International Cosmetic Ingredient Dictionary and Handbook</i>
ECHA	European Chemicals Agency
FDA	Food and Drug Administration
GRAS	generally recognized as safe
HET-CAM	chorioallantoic membrane of a fertilized hen's egg
HRIPT	human repeated insult patch test
IARC	International Agency for Research on Cancer
IDLH	immediately dangerous to life or health
ILO	Intentional Labor Office
LLNA	local lymph node assay
mppcf	million particles per cubic foot
NIOSH	National Institute for Occupational Safety and Health
NMRD	non-malignant respiratory disease
NOAEC	no-observable-adverse-effect-concentration
NR	not reported/none reported
OECD	Organization for Economic Co-operation and Development
OSHA	Occupational Safety and Health Administration
Panel	Expert Panel for Cosmetic Ingredient Safety
PEL	permissible exposure limit
ppm	parts per million
REL	recommended exposure limit
SCOGS	Select Committee on GRAS Substances
SHE	Syrian hamster embryos
SI	stimulation index
SMR	standardized mortality ratio
SWeRF	size-weighted relevant fine fraction
TG	test guideline
TWA	time weighted average
UICC	Union for International Cancer Control
US	United States
VCRP	Voluntary Cosmetic Registration Program

ABSTRACT

The Expert Panel for Cosmetic Ingredient Safety (Panel) assessed the safety of Diatomaceous Earth as used in cosmetic formulations. It is reported to function as an abrasive, absorbent, anticaking agent, bulking agent, and opacifying agent in cosmetic products. The Panel reviewed all relevant data, and concluded that Diatomaceous Earth is safe in cosmetics in the present practices of use and concentration described in this safety assessment.

INTRODUCTION

This assessment reviews the safety of Diatomaceous Earth as used in cosmetic formulations. Diatomaceous Earth is reported to function as an abrasive, absorbent, anticaking agent, bulking agent, and opacifying agent in cosmetics, according to the web-based *International Cosmetic Ingredient Dictionary and Handbook* (wINCI; *Dictionary*).¹

The Expert Panel for Cosmetic Ingredient Safety (Panel) has reviewed related ingredients. In a report that was finalized in 2019, the Panel concluded that synthetically-manufactured amorphous silica and hydrated silica are safe in the present practices of use and concentration when formulated to be non-irritating.² Diatomaceous Earth is considered a natural amorphous form of silica. Synthetically-manufactured amorphous silica and hydrated silica are neither part of this safety assessment, nor are data from that report included in this assessment; however, the report on these ingredients are available on the Cosmetic Ingredient Review (CIR) website (<https://www.cir-safety.org/ingredients>).

This safety assessment includes relevant published and unpublished data that are available for each endpoint that is evaluated. Published data are identified by conducting an exhaustive search of the world's literature. A listing of the search engines and websites that are used and the sources that are typically explored, as well as the endpoints that the Panel typically evaluates, is provided on the CIR website (<https://www.cir-safety.org/supplementaldoc/preliminary-search-engines-and-websites>; <https://www.cir-safety.org/supplementaldoc/cir-report-format-outline>). Unpublished data are provided by the cosmetics industry, as well as by other interested parties.

Some chemical and toxicological data on Diatomaceous Earth included in this safety assessment were obtained from assessments by the International Agency for Research on Cancer (IARC)³ and the Agency for Toxic Substances and Disease Registry (ATSDR),⁴ as well as from robust summaries of data submitted to the European Chemical Agency (ECHA; listed as Kieselguhr)⁵ by companies as part of the REACH chemical registration process. These data summaries are available on the IARC, ATSDR, and ECHA websites, respectively, and when deemed appropriate, information from the summaries has been included in this report.

CHEMISTRY

Definition

Diatomaceous Earth (CAS No.61790-53-2 or 68855-54-9) is defined by the *Dictionary* as a mineral material consisting chiefly of the siliceous frustules and fragments of various species of diatoms, which may or may not be calcined.¹ [A frustule is the cell wall of a diatom]. Natural calcined and uncalcined forms are associated with the CAS No. 61790-53-2, and the flux-calcined form is associated with the CAS No. 68855-54-9.^{3,6} The “calcined” form is processed Diatomaceous Earth that is heated to 800 - 1000 °C to eliminate organic and carbonaceous material.⁷ The “flux-calcined” form is Diatomaceous Earth that is heated with the addition of sodium carbonate as a fluxing agent that results in a coarser material. Diatomaceous Earth is considered a natural amorphous form of silica.^{3,8}

Diatomaceous Earth is a polymorph of silica, or silicon dioxide.^{3,4} Silica may exist in amorphous or crystalline structures. While both forms are made up of silicon-oxygen tetrahedra, crystalline silica is determined by a regular, repeating arrangement of the silicon and oxygen tetrahedra, while the arrangement of bonds in amorphous silica is highly disordered and randomly linked. Silica can be sourced naturally as a mineral, biogenically through diatoms, or it can be synthetically produced. Natural and biogenic forms of amorphous silica include opal, Diatomaceous Earth, silicates and volcanic glass; while natural forms of crystalline silica include quartz, cristobalite, flint, and sandstone.

Chemical Properties

Available chemical properties for Diatomaceous Earth are provided in Table 1. Particle size distributions for Diatomaceous Earth (flux-calcined) for coarse, medium, and fine-grade materials were 59.5%, 81.6%, and 99.6% less than 90 µm, respectively, and 4.56%, 14.7%, and 58.7% less than 10 µm, respectively (Table 2).⁵ Diatomaceous Earth has an infinite variety of shapes, due to its origins in the living matter (diatoms) from which it formed.³

Method of Manufacture

Diatomaceous Earth is obtained by strip mining, commonly from the western portion of the United States (US).⁹ Diatomaceous Earth is also mined in western Canada, France, Denmark, Spain, Iceland, Romania, the Czech Republic, Algeria, Kenya, Morocco, Japan, South Korea, China, Australia, New Zealand, Mexico, Peru, Argentina, Costa Rica, Chile,

Brazil, Colombia, and Peru.¹⁰ Following extraction from a mine, the raw material is crushed, dried, ground, purified and alimmented.⁷ The resulting material may be used as-is (natural or milled product), or can be further process by heating (800 - 1000 °C) in one of two ways to produce a “calcined” product or a “flux-calcined” product.^{7,10} After heating, the material is then cooled and further ground before packaging. In commercial products, a large proportion of the amorphous silica in Diatomaceous Earth is converted into a crystalline form (cristobalite, up to 40% to 60%) during thermal processing.^{3,10}

The International Diatomite Producers Association and a supplier have reported that natural and flux-calcined Diatomaceous Earth are used in the formulation of cosmetic products.^{11,12} The flux-calcined material is produced through the following steps: harvesting, calcination, milling, sieving, quality control, packaging, and quality control.¹¹

Composition and Impurities

The composition of Diatomaceous Earth varies depending on where it is mined and how it is processed.¹³ Silica content in Diatomaceous Earth can vary between 68% to 96%.^{3,10,13-16} Other components may include aluminum (III) oxide (~4 - 7%), iron (III) oxide (~1 - 4%), titanium (IV) oxide; ions of calcium, magnesium, sodium, and potassium; and phosphates.^{3,10,13,15,16} Many elements are present in trace amounts, and co-deposited and secondary minerals can include clays, quartz, gypsum, mica, calcite, feldspars, salt, pyrite, sulfur, manganese nodules, and phosphates.¹⁰ Diatomaceous Earth usually contains 0.1% to 4% quartz.³ Chert and volcanic ash can be abundant constituents of the sediment, and common biogenic constituents include the siliceous remains of sponges, silicoflagellates, radiolaria, carbonized fossil leaves, and fossilized fish bones.¹⁰ Chemical and mineral impurities can affect the properties of the final Diatomaceous Earth product, including pH, solubles present, density, and abrasiveness: commercial uses can be adversely affected unless contaminants can be removed or made insoluble through processing.

Crystalline silica content of Diatomaceous Earth is dependent on the degree of exposure to high temperatures and pressures; surface chemistry of an individual Diatomaceous Earth sample may vary, depending upon production method and degree of hydration.⁴ The crystalline silica content of uncalcined Diatomaceous Earth is 0.1% to 4.0%. Cristobalite content of straight-calcined flux products is between 10% to 20%, and between 40% to 60% in flux-calcined products.^{3,17}

A supplier has reported that a product containing 100% Diatomaceous Earth has < 1% respirable crystalline silica.¹⁸ Another product containing 9-11% Diatomaceous Earth was reported to have < 0.11% respirable crystalline silica. This product also contained 57% - 61% *Lithothamnion calcareum* powder, 29% - 31% mannitol, and 0.7% - 1.5% zinc sulfate.¹⁹ This supplier has reported that flux-calcined Diatomaceous Earth is used in the finished products at concentrations below 10% and has a respirable crystalline silica content of < 1% (cristobalite) based on the size-weighted relevant fine fraction (SWeRF) method of analysis.¹¹

According to international standards for food additives, Diatomaceous Earth should not contain more than 10 mg/kg arsenic or lead.⁶

USE

Cosmetic

The safety of the cosmetic ingredients addressed in this assessment is evaluated based on data received from the US Food and Drug Administration (FDA) and the cosmetics industry on the expected use of these ingredients in cosmetics, and does not cover their use in airbrush delivery systems. Data are submitted by the cosmetic industry via the FDA’s Voluntary Cosmetic Registration Program (VCRP) database (frequency of use) and in response to a survey conducted by the Personal Care Products Council (Council) (maximum use concentrations). The data are provided by cosmetic product categories, based on 21CFR Part 720. For most cosmetic product categories, 21CFR Part 720 does not indicate type of application and, therefore, airbrush application is not considered. Airbrush delivery systems are within the purview of the US Consumer Product Safety Commission (CPSC), while ingredients, as used in airbrush delivery systems, are within the jurisdiction of the FDA. Airbrush delivery system use for cosmetic application has not been evaluated by the CPSC, nor has the use of cosmetic ingredients in airbrush technology been evaluated by the FDA. Moreover, no consumer habits and practices data or particle size data are publicly available to evaluate the exposure associated with this use type, thereby preempting the ability to evaluate risk or safety.

According to 2022 VCRP survey data, Diatomaceous Earth is used in a total of 135 formulations (Table 3).²⁰ Of these reported uses, the majority are in leave-on products, with over a third of the uses (50) reported to be in nail products. Twenty-five uses are reported to be in rinse-off paste masks (mud packs). While uses were reported in a number of categories in the VCRP, the results of the concentration of use survey conducted by the Council in 2021 reported uses for Diatomaceous Earth in only 3 categories: at 0.001% in hair dyes and colors, up to 0.01% in nail polish and enamel, and at 2% in rinse-off products (paste masks).²¹

Diatomaceous Earth may be used in products that can come into contact with the eyes or mucous membranes; for example, it is reported to be used in eye shadow, eye lotion, bath soaps and detergents, and other personal cleanliness products (concentrations not reported).^{20,21} It is also reported to be used in products which maybe incidentally ingested, such as lipsticks and dentifrices (concentrations not reported). Additionally, Diatomaceous Earth is reported to be used in face

powders (concentration not reported), and could possibly be inhaled. In practice, as stated in the Panel's respiratory exposure resource document (<https://www.cir-safety.org/cir-findings>), most droplets/particles incidentally inhaled from cosmetics would be deposited in the nasopharyngeal and tracheobronchial regions and would not be respirable (i.e., they would not enter the lungs) to any appreciable amount. Conservative estimates of inhalation exposures to respirable particles during the use of loose powder cosmetic products are 400-fold to 1000-fold less than protective regulatory and guidance limits for inert airborne respirable particles in the workplace.

Although products containing some of these ingredients may be marketed for use with airbrush delivery systems, this information is not available from the VCRP or the Council survey. Without information regarding the frequency and concentrations of use of these ingredients, and without consumer habits and practices data or particle size data related to this use technology, the data are insufficient to evaluate the exposure resulting from cosmetics applied via airbrush delivery systems.

Diatomaceous Earth is not restricted from use in any way under the rules governing cosmetic products in the European Union.²²

Non-Cosmetic Use

Diatomaceous Earth has uses in food and beverages, including anticaking material in foodstuffs and clarifier in wine and beer.²³ In 1979, the Select Committee on GRAS (generally recognized as safe) Substances (SCOGS) opined that Diatomaceous Earth is GRAS as a filtering aid in such food and beverages as apple cider, beer, beet and cane sugar, vinegar, and wine in natural, calcined, or flux-calcined forms.²⁴ Diatomaceous Earth is also GRAS as a substance migrating to food from paper and paperboard products (21CFR§182.90). Diatomaceous Earth is approved as an indirect food additive with use as a polymer (21CFR§177.2410), as a component of paper and paperboard (21CFR176.170), and as a colorant for polymers (21CFR§178.3297). It is an approved food additive in animal feed with the restrictions that it cannot contain more than 15 ppm lead, 20 ppm arsenic, and 600 ppm fluorine (21CFR§573.340).

The use of Diatomaceous Earth as a drug carrier is being investigated.^{25,26} Diatomaceous Earth is an approved inactive ingredient in approved drug products, including capsules and tablets taken orally and in topical soaps.²⁷

Diatomaceous Earth is used in refractory and insulation bricks, filtration media, fertilizers, abrasives, insulation materials, lubricants, paints, rubbers, absorbents, bulking agents, and as carriers for catalysts.^{9,10,17,23} It is also widely used in pesticide formulations.^{10,14,17,23,28,29}

TOXICOKINETIC STUDIES

No toxicokinetic studies were discovered in the published literature, and no unpublished data were submitted. However, it should be noted that in a 13-wk study in which rats were given up to 5% Diatomaceous Earth via dietary pellets, residual silica values in the organs of the 5% dose group were comparable with the controls.³⁰ Repeated dose inhalation studies with 100% pure flux-calcined Diatomaceous Earth noted that the test material was detected in the alveoli when administered in high doses. In a study of nearly 2 years in guinea pigs exposed to uncalcined Diatomaceous Earth, total silica content per lung increased linearly.

TOXICOLOGICAL STUDIES

Acute Toxicity Studies

Oral

In an oral study performed in accordance with Organization for Economic Co-operation and Development (OECD) test guideline (TG) 401, female Wistar rats received a single dose of 300 mg/kg (1 rat) or 2000 mg/kg (5 rats) flux-calcined Diatomaceous Earth in arachis oil by gavage.⁵ The purity of the test material was not stated. Clinical observations were made at 0.5, 1, 2, and 4 h post-dosing, and then daily for 14 d. Morbidity and mortality were checked twice daily and body weights were recorded on days 0, 7, and 14. No mortalities were observed at either dose level. No signs of systemic toxicity were observed at 300 mg/kg; however, at 2000 mg/kg, clinical signs of toxicity included hunched posture in all animals and ataxia in one animal. All animals had expected body weights gains, and no abnormalities were observed at necropsy. The LD₅₀ for Diatomaceous Earth in this study was greater than 2000 mg/kg.

Inhalation

In a dust aerosol study performed in accordance with OECD TG 403, 5 male and 5 female Wistar rats received 2.7 mg/l flux-calcined Diatomaceous Earth (100%; target particle size 1 to 4 µm).⁵ The rats were exposed to the test material nose-only for 24 h. Clinical observations were made during exposure, immediately after exposure, and 1 h after exposure, and then once daily for 14 d. Body weights were recorded on test days 1 (before exposure), 2, 4, 8, and 15 (before necropsy). No mortalities were observed. Clinical signs of toxicity included moderately-ruffled fur in all animals on test day 1 that persisted until day 2, and slight nose scabbing on day 1 in all animals. Marginal to slight body weight loss was noted in all males and 4 females on day 1 and 2 but returned to expected gains thereafter. No abnormalities were observed at necropsy. The LC₅₀ for Diatomaceous Earth in this study was greater than 2.7 mg/l.

Short-Term, Subchronic, and Chronic Dose Toxicity Studies

Repeated dose oral and inhalation studies summarized here are described in Table 4. In 13-wk dietary studies, rats that received up to 5% natural or flux-calcined Diatomaceous Earth did not exhibit effects outside of increased body weight gains starting at 3% natural Diatomaceous Earth in one study.^{5,30}

In inhalation studies, a no-observable-adverse-effect-concentration (NOAEC) could not be determined in a 28-d inhalation rat study of 100% pure flux-calcined Diatomaceous Earth (particle size range 1 to 3 µm) tested at 0, 0.044, 0.207, or 0.7 mg/l.⁵ Effects in the high dose group included increased spleen, adrenal, and liver weights and slight and transient effect on body weight gains. In a 2-yr rat inhalation study of a flux-calcined Diatomaceous Earth at up to 5 million particles per cubic foot (mppcf) per day plus 50 mppcf for 1 h three times per week (5 + 50 mppcf), no fibrosis was observed.¹⁵ Perivascular and peribronchiolar localization of dust-laden macrophages were observed in both the 2 and 5 mppcf dose groups, and nodular lesions and reactions of the nodes were greater in the 5 mppcf dose group. A similar study of the same test material in guinea pigs also found no fibrosis after 1.5 yr, and a light increase in intra-alveolar macrophages with peribronchiolar localization in the 5 mppcf group. In another guinea pig study of unheated and heated Diatomaceous Earth (particle size range ~0.45 µm to > 10 µm), no fibrosis was noted during observations made at 2-3 mo intervals until study end at 2 yr.³¹ No fibrosis was observed in mongrel dogs exposed to up to 5 mppcf flux-calcined Diatomaceous Earth for up to 2.5 yr.¹⁵

DEVELOPMENTAL AND REPRODUCTIVE TOXICITY (DART) STUDIES

No DART studies were discovered in the published literature, and no unpublished data were submitted.

GENOTOXICITY STUDIES

In vitro genotoxicity studies summarized here are described in Table 5. Diatomaceous Earth (100% pure flux-calcined) was not mutagenic in an Ames test (up to 5000 µg/plate) or a mouse lymphoma cell gene mutation test (up to 40 µg/ml), and was not clastogenic in a human lymphocyte chromosome aberration test (up to 40 µg/ml).⁵ Abnormal cell proliferation, colony-forming efficiency, and nuclei formation was observed in Chinese hamster ovary (CHO) cells in assays with uncalcined and flux-calcined Diatomaceous Earth (1.3 µm and 2.1 µm, respectively; concentrations tested not reported).¹⁶ In studies with Syrian hamster embryo (SHE) cells treated with high temperature calcined and flux-calcined Diatomaceous Earth, concentration-dependent increases in cell division aberrations and cell transformations were observed; the induction of transforming potency was correlated with the amount of hydroxyl radicals generated.³²⁻³⁴ Cell transformation was decreased or not observed in SHE cells exposed to uncalcined Diatomaceous Earth samples where the likelihood of radical generation was decreased or non-existent.

CARCINOGENICITY STUDIES

The International Agency for Research on Cancer (IARC) has determined that “there is *inadequate evidence* in experimental animals for the carcinogenicity of uncalcined Diatomaceous Earth.” Overall, amorphous silica is not classifiable as to its carcinogenicity to humans (Group 3).³

Oral

In a feeding study, a group of 30 weanling Sprague-Dawley rats (sex not reported) received 20 mg/d Diatomaceous Earth (particle size not reported) mixed with cottage cheese at a concentration of 5 mg/g cheese.³⁵ The rats also received commercial rat chow and filtered tap-water ad libitum. A control group of 27 rats was only fed commercial rat chow. The animals were observed for their life span (mean survival following the start of treatment for treated rats was 840 d, and for control rats was 690 d). Complete gross and microscopic thoracic and abdominal necropsies were performed on each animal upon expiration, with special attention given to the gastrointestinal tract. During the course of the study, 5 malignant tumors (1 salivary gland carcinoma, 1 skin carcinoma, 2 sarcomas of the uterus, and 1 peritoneal mesothelioma) and 13 benign tumors (9 mammary fibroadenomas, 1 adrenal pheochromocytoma, and 3 pancreatic adenomas) were observed in the treated animals. The control group had 3 carcinomas (1 each in the lung, forestomach and ovary) and 5 mammary fibroadenomas. The authors determined that the difference in tumor incidence between treated and control rats was not statistically significant ($0.25 < p < 0.5$, χ^2 -test).

Subcutaneous

A group of 36 female Marsh mice, 3-mo-old, received a subcutaneous injection of 20 mg Diatomaceous Earth (uncalcined, particle size, 3 - 9 µm, with some crystalline material of larger size) suspended as a 10% slurry in isotonic saline (volume unspecified).³ Another group of 36 female littermates received an injection of 0.2 ml saline only. The numbers of mice still alive at 19 mo were 19/36 in the treated group and 20/36 in the control group. The treated group showed an extensive reactive granulomatous and fibroplastic reaction at the site of injection, but no malignant tumors. No further details were available.

Intraperitoneal

In another study by the same researchers, a group of 29 female Marsh mice, 3-mo-old, received an intraperitoneal injection of 20 mg Diatomaceous Earth suspended as a 10% slurry in isotonic saline.³ A group of 32 female littermates received an injection of the same volume of saline only (volume unspecified). The numbers of mice still alive at 19 mo were 11/29 in the treated group and 19/32 in the control group. Lymphosarcomas at the injection site in the abdominal cavity were reported in 6/17 treated animals and 1/20 controls ($p = 0.02$; method of statistical analysis unspecified). No further details were available.

OTHER RELEVANT STUDIES

Pulmonary Response

The following summaries demonstrate the physiological changes to the pulmonary system when Diatomaceous Earth enters the lung. In an intratracheal study, groups of 6 male Sprague-Dawley rats received a single instillation of Diatomaceous Earth (90% amorphous silica; particle size $< 7 \mu\text{m}$) suspended in isotonic saline.³⁶ Rats that underwent bronchoalveolar lavage (BAL) examinations received 10 mg/animal, and rats that underwent lung biochemical examinations received 15 mg/animal. Determinations in the BAL and phospholipids in the lung tissue were determined after 15, 60, and 180 d and 90, 180, and 360 d, respectively. Acute/subacute inflammation was observed that gradually became moderate after 60 d. No further details provided.

In another intratracheal study, groups of Hartley-Duncan guinea pigs (sex not specified) received a single instillation of 25 mg flux-calcined Diatomaceous Earth (particles $\leq 3.0 \mu\text{m}$ in diameter; 72% silica and 28% calcium silicates) in 0.5 ml physiological saline.³⁷ A control group of 2 animals received 0.5 ml saline only. After 2 or 4 h, 1, 2, 3, 4, 5, 6, or 7 d, and 5, 6, or 15 mo, 2 animals/time period were killed and lungs were dissected. No signs of infection nor significant individual variation in response within time period were observed. Pronounced neutrophil invasion of the bronchioles was observed by 4 h post-exposure, which remained well developed through 1 d post-exposure. The number of macrophages and neutrophils in the alveoli increased through 1 d post-exposure and remained greater than control values through 7 d post-exposure. The number of macrophages, many of which contained Diatomaceous Earth, remained elevated for the duration of the experiment. Phagocytosis of the particles was mainly performed by the macrophages, with some participation by the neutrophils. Many of the reactive macrophages in the groups longer than 2-h post-exposure had various types of pathological alterations. Some particles were found in type I epithelial cells. Edematous changes were observed in some type I epithelial cells, and proliferation of type II epithelial cells was observed in some alveoli, especially near the respiratory bronchiole. Mild, diffuse fibrosis was observed starting at 6 mo post-exposure and persisted at 15 mo post-exposure.

Cytotoxicity

Natural and flux-calcined Diatomaceous Earth (average diameters $1.3 \mu\text{m}$ and $2.1 \mu\text{m}$, respectively), in addition to titanium dioxide, crocidolite, chrysotile, quartz, and cristobalite, were studied for cell proliferation with cultured Chinese hamster ovary (CHO) cells.¹⁶ Concentration ranges were not reported; however, crystalline silica content of the natural Diatomaceous Earth was 4% quartz, and of the flux-calcined Diatomaceous Earth was 40% cristobalite and 2% quartz. For the assay, 100,000 cells were seeded/dish and incubated for 1 d prior to exposure to test dust for 3 d (vehicle not reported). Cells were then harvested and counted.

The ranking of toxicity as measured by the inhibition of cell proliferation was chrysotile $>$ crocidolite $>$ natural Diatomaceous Earth $>$ flux-calcined Diatomaceous Earth $>$ quartz $>$ cristobalite $>$ titanium dioxide. The effective concentration-50% (EC_{50}) for natural Diatomaceous Earth was $3.6 \mu\text{g}/\text{cm}^2$, and for flux-calcined Diatomaceous Earth was $10.8 \mu\text{g}/\text{cm}^2$. Responses were concentration-dependent. The researchers found that the toxicity of the dusts did not correlate with crystalline silica content, surface area, composition, volume, particles/ cm^2 , or fibrous geometry; however, toxicity was closely associated with the number of particles/ cm^2 culture surface that had one dimension $> 7.5 \mu\text{m}$. The authors indicated that particle size impacted toxicity.¹⁶

DERMAL IRRITATION AND SENSITIZATION STUDIES

Dermal irritation, sensitization, and phototoxicity studies summarized here are described in Table 6. Diatomaceous Earth (flux-calcined, up to 100% pure) was considered non-corrosive and non-irritating in EpiSkin™ reconstituted human epidermis model tests.⁵ In acute skin tolerance patch tests, Diatomaceous Earth (flux-calcined) was not irritating in 10 healthy volunteers at 100% or in 11 volunteers with sensitive skin in a product at 9% - 11%.^{38,39} Diatomaceous Earth was not sensitizing in a local lymph node assay (LLNA) at up to 10%.⁵ A cosmetic product containing 9% - 11% Diatomaceous Earth (soda ash flux-calcined) was not sensitizing in a human repeated insult patch test (HRIPT) of 100 healthy subjects when tested at a 10% dilution, nor was it phototoxic in a human single-application study in 10 healthy female subjects when tested neat.^{40,41}

OCULAR IRRITATION STUDIES

In Vitro

The ocular irritation potential of Diatomaceous Earth (flux-calcined, purity not reported) was assessed in a SkinEthic™ reconstituted human corneal epithelium model test.⁵ The test material was used as supplied, and 30 mg was applied to the tissue cultures. Triplicate cultures were exposed for 10 min, and then examined after 3 h. Viability of the tissues following exposure to the test material was 99.1% and the qualitative evaluation of the tissue following exposure indicated it was viable. The positive and negative controls yielded expected results. Based on the results of the study, the test material was considered non-irritating.

In another in vitro study, the ocular irritation potential of a formulations containing 9% - 11% Diatomaceous Earth (soda ash flux-calcined) was assessed using the chorioallantoic membrane of a fertilized hen's egg (HET-CAM test).⁴² The material was tested at 2%, 5%, and 10% w/v dilutions in water. Approximately 0.3 ml of the sample was spread over membrane and rinsed with 5 ml of demineralized water after 20 s. The test material was non-irritating at the 2% and 5% dilution, but moderately irritating at the 10% dilution. The 10% dilution had low solubility and rapid sedimentation; however, the results were reproducible between eggs and were considered relevant.

Animal

The ocular irritation potential of Diatomaceous Earth (flux-calcined, purity not reported) was assessed in 2 New Zealand White rabbits (sex not reported) in accordance with OECD TG 405.⁵ The undiluted test material was instilled at a volume of 0.1 ml in the right eye of the animals. The left eye was left untreated as a control. After instillation, the rabbits were observed for 72 h. No corneal effects were reported. Iridial inflammation was reported in one animal at 1 and 24 h post-instillation. Moderate conjunctival irritation was noted in both animals at 1 and 24 h post-instillation, and up to 48 h post-instillation in 1 animal. Both animals had recovered by 72 h post-instillation. The test material was considered to be non-irritating to the eye in this study.

CLINICAL STUDIES

Case Report

A 51-yr-old male employed in the Diatomaceous Earth industry for 26 yr (20 yr in a mill, 6 yr in an office) was reported to have a history of a recurrent peptic ulcer, pleurisy, and bronchopneumonia, with frequent attacks of bronchitis.⁴³ The patient was a nonsmoker. An electrocardiogram indicated right ventricular hypertrophy. The patient had a 4-yr history of intermittent palpitation, severe exertional moderate paroxysmal dyspnea, and orthopnea. He also complained of wheezing and hoarseness, with productive cough, until a year and a half before presentation. Cough, but not dyspnea, was relieved by bronchodilator aerosols. At physical examination, no apparent distress or cyanosis were noted; however, slight clubbing of the fingers was observed. Rales were detected over most of the chest except in infraclavicular areas anteriorly. Resonance was diminished over the upper lung fields posteriorly, and on the left anteriorly. Chest films were interpreted as consistent with far-advanced coalescent pneumoconiosis. The patient died 5 yr after the chest films were made, reportedly due to heart failure from cor pulmonale.

OCCUPATIONAL EXPOSURE STUDIES

Occupational exposure studies are described in Table 7. Occupational exposure studies indicate a risk of pneumoconiosis in Diatomaceous Earth mine and mill workers, which can be mitigated with dust control measures and personal protective equipment.⁴⁴⁻⁴⁹ Studies were of quarry and mill workers in the western US and exposures were to raw, calcined, or flux-calcined Diatomaceous Earth.

OCCUPATIONAL EXPOSURE LIMITS

Occupational exposure to Diatomaceous Earth, and the quartz and amorphous silica dust it contains, can occur during mining, the calcination process, and through handling the calcined product in end-use industries as a filtration agent, mineral charge, refractory, abrasive, carrier, or adsorbent.³ The National Institute for Occupational Safety and Health (NIOSH) time weighted average (TWA) for recommended exposure limits (REL) for Diatomaceous Earth (also characterized as amorphous silica) is 6 mg/m³, and the Occupational Safety and Health Administration (OSHA) TWA permissible exposure limit (PEL) is 20 mppcf (80 mg/m³/100% silicon dioxide).^{50,51} The immediately dangerous to life or health (IDLH) value is 3000 mg/m³.⁵⁰

SUMMARY

Diatomaceous Earth is reported to function as an abrasive, absorbent, anticaking agent, bulking agent, and opacifying agent in cosmetics. The "calcined" form is processed Diatomaceous Earth that is heated to 800 - 1000 °C to eliminate organic and carbonaceous material. The "flux-calcined" form is Diatomaceous Earth that is heated with the addition of sodium carbonate as a fluxing agent that results in a coarser material). Diatomaceous Earth is considered a natural amorphous form of silica.

The composition of Diatomaceous Earth varies depending on where it is mined and how it is processed. Silica content in Diatomaceous Earth can vary between 83% to 96%. Crystalline silica content of Diatomaceous Earth is dependent on the degree of exposure to high temperatures and pressures; surface chemistry of an individual Diatomaceous Earth sample may vary, depending upon production method and degree of hydration. The crystalline silica content of uncalcined Diatomaceous Earth is 0.1% to 4.0%. Cristobalite content of straight-calcined flux products is between 10% to 20%, and between 40% to 60% in flux-calcined products.

According to 2022 VCRP survey data, Diatomaceous Earth is used in a total of 135 formulations. Of these reported uses, the majority are in leave-on products with over a third of the uses (50) reported to be in nail products. Twenty-five uses are reported to be in rinse-off paste masks (mud packs). The results of the concentration of use survey conducted by the Council in 2021 indicate that Diatomaceous Earth is used at 0.001% in hair dyes and colors, up to 0.01% in nail polish and enamel, and at 2% in rinse-off products (paste masks). Diatomaceous Earth is reported to be used in cosmetic powders, and could possibly be inhaled; for example, it is reported to be used in face powders (concentration not reported).

In a 90-d dietary study, male and female rats were fed a diet containing 5% Diatomaceous Earth. (Estimated intake ranged from about 12 g/kg bw/d at the start of the experiment to about 5 g/kg at the end of the experiment.) Residual silica values in the organs of treated rats were comparable with the controls.

In oral rat studies with flux-calcined Diatomaceous Earth, the LD₅₀ was greater than 2000 mg/kg. The LC₅₀ was greater than 2.7 mg/l in a 24 h inhalation rat study of flux-calcined Diatomaceous Earth.

In 13-wk dietary studies, rats that received up to 5% natural or flux calcined Diatomaceous Earth did not exhibit adverse effects outside of increased body weight gains in one study. An NOAEC could not be determined in a 28-d inhalation rat study of 100% pure flux-calcined Diatomaceous Earth (particle size range 1 to 3 µm) at up to 0.7 mg/l. In a 2-yr rat inhalation study of a flux-calcined Diatomaceous Earth at up to 5 mppcf, no fibrosis was observed. Perivascular and peribronchiolar localization of dust-laden macrophages were observed in the 2 and 5 mppcf dose groups and nodular lesions and reactions of the nodes was greater in the 5 mppcf dose group. A similar study of the same test material in guinea pigs also found no fibrosis after 1.5 yr and a light increase in intra-alveolar macrophages with peribronchiolar localization in the 5 mppcf group. In another guinea pig study of unheated and heated Diatomaceous Earth (particle size range ~0.45 µm to > 10 µm), no fibrosis was noted during observations made at 2-3 mo intervals until study end at 2 yr. No fibrosis was observed in mongrel dogs exposed to up to 5 mppcf flux-calcined Diatomaceous Earth for up to 2.5 yr.

Diatomaceous Earth (100% pure flux-calcined) was not mutagenic in an Ames test (up to 5000 µg/plate) or a mouse lymphoma cell gene mutation test (up to 40 µg/ml); and was not clastogenic in a human lymphocyte chromosome aberration test (up to 40 µg/ml). In studies with SHE cells, high temperature calcined and flux-calcined Diatomaceous Earth had increased cell division aberrations and cell transformations in a concentration-dependent manner; the induction of transforming potency was correlated with the amount of hydroxyl radicals generated. Cell transformation was decreased or not observed in SHE cells exposed to uncalcined Diatomaceous Earth samples where the likelihood of radical generation was decreased or non-existent.

IARC has determined that there is inadequate evidence in experimental animals for the carcinogenicity of uncalcined Diatomaceous Earth. In an oral feeding study in Sprague-Dawley rats that received 20 mg/d Diatomaceous Earth in cottage cheese, there was no statistically significant difference in cancer incidence between treated and control rats. A subcutaneous study in mice found uncalcined Diatomaceous Earth led to extensive reactive granulomatous and fibroplastic reactions at the injection site, but no malignant tumors were observed. The same research group performed an intraperitoneal study in mice and found lymphosarcomas at the injection site in the abdominal activity.

In an intratracheal rat study of Diatomaceous Earth that was 90% amorphous silica, acute/subacute inflammation was observed that gradually became moderate after 60 d. Guinea pigs that received a single 25 mg intratracheal instillation had mild, diffuse fibrosis observed starting 6 mo after exposure that persisted to 15 mo. Abnormal cell proliferation, colony-forming efficiency, and nuclei formation was observed in CHO cells in assays with uncalcined and flux-calcined Diatomaceous Earth (1.3 µm and 2.1 µm, respectively; concentrations tested not reported).

Diatomaceous Earth (flux-calcined, up to 100% pure) was considered non-corrosive and non-irritating in EpiSkin™ reconstituted human epidermis model tests. In acute skin tolerance patch tests, Diatomaceous Earth (flux-calcined) was not irritating in 10 healthy volunteers at 100% or in 11 volunteers with sensitive skin in a product at 9% - 11%. Diatomaceous Earth was not sensitizing in a LLNA at up to 10%. A cosmetic product containing 9% - 11% Diatomaceous Earth (soda ash flux-calcined) was not sensitizing in a HRIPT of 100 healthy subjects when tested at a 10% dilution, nor was it phototoxic in a human single application study in 10 healthy female subjects when tested neat.

In ocular studies, flux-calcined Diatomaceous Earth in a formulation at 9%-11% was non-irritating at 2% and 5% dilutions, but was moderately irritating at a 10% dilution. However, flux-calcined Diatomaceous Earth (tested neat) was not an ocular irritant in an in vitro reconstituted human corneal epithelium model test nor in a rabbit eye test.

A case report of a worker at a Diatomaceous Earth mill observed far-advanced coalescent pneumoconiosis. Occupational studies indicate a risk of pneumoconiosis in Diatomaceous Earth mine and mill workers, which can be mitigated with dust control measures and personal protective equipment. The TWA REL for Diatomaceous Earth set by

NIOSH is 6 mg/m³ and the TWA PEL set by OSHA is 20 mppcf (80 mg/m³/100% silicon dioxide). The IDLH value is 3000 mg/m³.

No DART studies were discovered in the published literature, and no unpublished data were submitted.

DISCUSSION

The Panel reviewed the safety of Diatomaceous Earth as used in cosmetic formulations. The available data are sufficient for determining safety, and the Panel concluded Diatomaceous Earth is safe in cosmetics in the present practices of use and concentration described in this safety assessment.

Diatomaceous Earth is naturally-occurring and is a polymorph of silica, or silicon dioxide. Regardless of whether Diatomaceous Earth is calcined or not, it can contain crystalline silica; crystalline silica is a known respiratory carcinogen.

Accordingly, the Panel discussed the issue of incidental inhalation exposure resulting from this ingredient. (Diatomaceous Earth is reported to be used in face powders (concentration not reported), and could possibly be inhaled.) The Panel noted that chronic inhalation studies of flux-calcined Diatomaceous Earth (which may comprise up to 60% crystalline silica) were negative for fibrosis or tumors in rats and guinea pigs. Additional data from acute and short-term inhalation studies suggest little potential for respiratory effects at relevant doses. Furthermore, droplets/particles deposited in the nasopharyngeal or tracheobronchial regions of the respiratory tract present no toxicological concerns based on the chemical and biological properties of these ingredients. Coupled with the small actual exposures in the breathing zone and the concentrations at which this ingredient is used (or expected to be used) in potentially inhaled products, the available information indicates that incidental inhalation would not be a significant route of exposure that might lead to local respiratory or systemic effects. (The Panel had an initial concern that face masks may flake during drying, and those flakes could incidentally be inhaled; this concern was mitigated once the Panel had a better understanding as to how those products work.) A detailed discussion and summary of the Panel's approach to evaluating incidental inhalation exposures to ingredients in cosmetic products is available at <https://www.cir-safety.org/cir-findings>.

The Panel's respiratory exposure resource document (see link above) notes that airbrush technology presents a potential safety concern, and that no data are available for consumer habits and practices thereof. As a result of deficiencies in these critical data needs, the safety of cosmetic ingredients applied by airbrush delivery systems cannot be assessed by the Panel. Therefore, the Panel has found the data insufficient to support the safe use of cosmetic ingredients applied via an airbrush delivery system.

The Panel expressed concern over the lack of DART studies for Diatomaceous Earth. However, the Panel noted that Diatomaceous Earth did not produce adverse effects in oral rodent studies, and is GRAS for uses in food and beverages. This information, coupled with noted lack of residual silica absorption in a 13-wk dietary study in rats, helped mitigate concern over the absence of DART data.

The Panel also expressed concern about the presence of heavy metals in Diatomaceous Earth. Although heavy metals may be present during mining, those should be readily avoidable/separable. Accordingly, the Panel stressed that the cosmetics industry should continue to use current good manufacturing processes (cGMPs) to limit impurities in cosmetic formulations.

CONCLUSION

The Expert Panel for Cosmetic Ingredient Safety concluded that Diatomaceous Earth is safe in cosmetics in the present practices of use and concentration described in this safety assessment.

TABLES**Table 1. Chemical properties**

Property	Value	Reference
Physical Form	Powder	5
Color	White or beige	5
	Calcined = pink to light brown or light yellow to light orange	6
	Flux-calcined = white to pink or light brown	6
Density/Specific Gravity (g/ml @ 20 °C)	2.36	5
Melting Point (°C)	1710	4
Boiling Point (°C)	2230	4
Water Solubility (mg/l @ 20 °C & pH 3)	3.7	5

Table 2. Particle size distributions for flux-calcined Diatomaceous Earth⁵

Diameter of particles (µm)	Volume % less than		
	Fine-Grade	Medium-Grade	Coarse-Grade
1	3.81	1.89	0.68
1.5	6.81	3.09	1.18
2	9.63	3.89	1.56
3	15.7	5.12	2.09
4	22.3	6.28	2.49
6	35.3	8.76	3.18
10	58.7	14.7	4.56
20	90.3	32.2	9.59
28	95.9	43.7	15.1
40	98.4	56.4	24.4
50	99.1	64.1	32.1
75	99.5	76.7	50
90	99.6	81.6	59.5
250	99.996	97	96.2
600	100	99.95	99.98

Table 3. Frequency (2022)²⁰ and concentration (2021)²¹ of use according to duration and exposure

	# of Uses	Max Conc of Use (%)
Totals*	135	0.001-2
<i>Duration of Use</i>		
Leave-On	92	0.0049-0.01
Rinse-Off	43	0.001-2
Diluted for (Bath) Use	NR	NR
<i>Exposure Type</i>		
Eye Area	2	NR
Incidental Ingestion	17	NR
Incidental Inhalation-Spray	9 ^a ; 8 ^b	NR
Incidental Inhalation-Powder	5; 9 ^a	NR
Dermal Contact	67	2
Deodorant (underarm)	3 ^b	NR
Hair - Non-Coloring	1	NR
Hair-Coloring	NR	0.001
Nail	50	0.0049-0.01
Mucous Membrane	20	NR
Baby Products	NR	NR

*Because this ingredient may be used in cosmetics with multiple exposure types, the sum of all exposure types may not equal the sum of total uses.

^a Not specified whether a spray or a powder, but it is possible the use can be as a spray or a powder, therefore the information is captured in both categories

^b It is possible these products are sprays, but it is not specified whether the reported uses are sprays.

NR – not reported

Table 4. Repeated dose toxicity studies of Diatomaceous Earth

Test Material Dose/Concentration	Animals/Group	Study Duration	Vehicle	Protocol	Results	Reference
ORAL						
0%, 1%, 3%, or 5% Diatomaceous Earth of freshwater origin, particle size range was 0.46 µm to 640 µm, with 90% smaller than 100 µm and 55% smaller than 12 µm	Groups of 15 male and 15 female Wistar rats	13-wk study	Dietary pellets	Body weights recorded weekly; at study end animals were killed and necropsied; livers, kidneys, and spleens of rats fed test material at 5% were analyzed for residual silica	Body weights of the 5% dose group were greater than the controls through the course of the study, with the maximum weight differential occurring at week 6; body weight gains in the 3% dose group were similar to those in the 5% group; body weight gains in the 1% dose group were similar to controls; histologic examination of organs of the 5% dose group were comparable to controls; residual silica values in the organs of the 5% dose group were comparable with the controls	³⁰
1% and 5% natural Diatomaceous Earth and 5% flux-calcined Diatomaceous Earth as feed; 5% natural mixture contained 4.8% silica, 0.44% quartz, and no cristobalite; 1% natural mixture contained 1.2% silica, 0.24% quartz, and 0.26% cristobalite; 5% flux calcined mixture contained 5.1% silica, 0.43% quartz, and 1.70% cristobalite	Groups of 20 male and 20 female Sprague-Dawley rats	13-wk study	Dietary pellets	Study performed in accordance with OECD TG 408; control animals received plain diet;	No clinical signs of toxicity or mortalities observed; no effects observed in body weight; feed consumption, ophthalmological findings, hematological findings, clinical biochemistry findings, or urinalysis findings; no treatment-related effects were observed at necropsy	⁵
INHALATION						
100% pure flux-calcined; 0, 0.018, 0.58, or 1.57 mg/l; target particle size range was 1 to 3 µm	5 male and 5 female Wistar rats/dose group	5-d range finding study	None described	Nose-only aerosol inhalation study; 6 h/exposure performed	No clinical signs of toxicity or mortalities observed; reduced feed consumption was observed in the high dose group; mean body weight loss was recorded in both male and female animals in the high dose group and a statistically-significant reduced body weight gain was observed in male rats in the high dose group only when compared with controls; dose-dependent alveolar histiocytosis was observed in all dose groups; alveolitis was observed in one male in the mid-dose group and in all animals in the high-dose group as well as increased absolute and relative lung weights in the mid- and high-dose groups; microgranulomas were found in one male and female in the mid-dose group and in all animals in the high-dose group; the test material was observed in the alveoli in most of the high-dose group animals; a no-observable-effect-concentration (NOEC) could not be determined	⁵

Table 4. Repeated dose toxicity studies of Diatomaceous Earth

Test Material Dose/Concentration	Animals/Group	Study Duration	Vehicle	Protocol	Results	Reference
100% pure flux-calcined; 0, 0.044, 0.207, or 0.700 mg/l; target particle size range was 1 to 3 µm	20 male and 20 female Wistar rats/dose group	28-d study	Compressed air	Study performed in accordance with OECD TG 412; nose-only aerosol inhalation study; 6 h/exposure performed 5d/wk with a 9-wk recovery period	No clinical signs of toxicity or mortalities observed; a slight and transient effect on body weight gain occurred in the high dose group; dose-dependent increase in lung weights recorded at the end of treatment period that further increased at the end of the recovery period; lymph nodes were also increased in size at the end of the recovery period; increase in spleen, adrenal, and liver weights was observed in the high-dose group at the end of the recovery period; histiocytosis was observed in the alveoli with a dose-dependent increase in incidence and severity that progressed during the recovery period; test material was detected in the alveoli in the mid and high dose group animals at the end of the treatment period that persisted until the end of the recovery period; a NOAEC could not be determined	⁵
Flux-calcined Diatomaceous Earth (61% cristobalite); 0, 2, 5, 50, and 5+50 mppcf; mean particle size 0.7 µm	Male Wistar rats divided as follows in the 0, 2, 5, 50, and 5+50 mppcf dose groups: 47, 79, 82, 46, and 53 animals, respectively	2- yr study	None described	Rats exposed to test material in exposure chambers for 6 h/d, 5 d/wk for up to 2 yr except in the 50 mppcf (1 h, 3 times/wk) and the 5+50 mppcf (daily 5 mppcf exposure plus 50 mppcf 3 times/wk for 1 h each) dose groups; rats killed at 6 mo, 1 yr, 1.5 yr, and 2 yr; lungs, hilar lymph nodes, heart, liver, kidney, spleen, adrenal, small intestine (duodenum), and hepatic lymph nodes underwent histologic examination	Terminal body weights at 1 yr and 1.5 yr in treated groups were comparable to controls except for in the rats exposed to 5+50 mppcf, which were below the control and 5 mppcf group; tissues studied other than the lungs had no test material-related changes. At 6 mo, rats in 2 and 5 mppcf dose groups had scattered macrophages and occasional giant cell within alveolar spaces; there was no significant septal reaction; pulmonary hilar lymph nodes only slightly enlarged and contained small clusters of macrophages in medullary portions; 5+50 mppcf group had slightly enhanced cellular reaction, when compared to the 5 mppcf group, and macrophages were noted to accumulate around bronchioles. At 1 yr, an increased macrophagic infiltration of perivascular and peribronchiolar areas were observed in the 2 and 5 mppcf groups; reactions were dose dependent; in 5+50 mppcf, macrophagic cells accumulated in a nodular fashion and reticular condensation was evident in lung parenchyma and hilar nodes. At 1.5 yr, no definite parenchymal or lymph node fibrosis was observed. At 2 yr, perivascular and peribronchiolar localization of dust-laden macrophages was observed in the 2 and 5 mppcf dose groups; nodular lesions and reaction of the nodes was greater in the 5 mppcf dose group; no fibrosis evident.	¹⁵

Table 4. Repeated dose toxicity studies of Diatomaceous Earth

Test Material Dose/Concentration	Animals/Group	Study Duration	Vehicle	Protocol	Results	Reference
Flux-calcined Diatomaceous Earth (61% cristobalite); 0, 2, 5, 50, and 5+50 mppcf; mean particle size 0.7 µm	Male guinea pigs (strain not reported) divided as follows in the 0, 2, 5, 50, and 5+50 mppcf dose groups: 47, 57, 69, 20, and 20 animals, respectively	1.5-yr study	None described	Guinea pigs exposed to test material in exposure chambers for 6 h/d, 5 d/wk for up to 1.5 yr except in the 50 mppcf (1 h, 3 times/wk) and the 5+50 mppcf (50 mppcf for 3 d/wk plus daily 5 mppcf) dose groups; rats killed at 6 mo, 1 yr, and 1.5 yr; lungs, hilar lymph nodes, heart, liver, kidney, spleen, adrenal, small intestine (duodenum), and hepatic lymph nodes underwent histologic examination	<p>Terminal body weights at 1 yr and 1.5 yr were comparable to controls; tissues studied other than the lungs had no test material-related changes.</p> <p>At 6 mo, same as the findings for the rats above.</p> <p>At 1 yr, definite cellular reaction with large clusters of macrophages and multinucleated giant cells in alveolar spaces in the 5 mppcf group; macrophages observed to accumulate around bronchioles and alveolar ducts; hilar lymph nodes were markedly enlarged and medullary portions were packed with dust cells and interwoven reticulum fibers.</p> <p>At 1.5 yr, a slight increase in intra-alveolar macrophages with peribronchiolar localization was observed in the 5 mppcf group; alveolar septa were unaffected and no fibrosis evident</p>	¹⁵
Diatomaceous Earth at 171 mppcf (natural, unheated), cristobalite at 167 mppcf (from heat-treated Diatomaceous Earth), or sodium silicate; particle size range ~0.45 µm to > 10 µm	Albino guinea pigs (sex and number/group not reported)	21-24 mo study	None described	Guinea pigs were placed in separate cubical dust rooms (512 ft ³) for 24 h/d until killed for examination; dust was generated within the room for 7 to 8 h/d, 5.5 d/wk for 21-24 mo; control animals kept in ambient air; pairs of animals selected at random were killed at 2-3 mo intervals and lung tissues were collected and analyzed for total silica content and total ash	<p>In animals exposed to Diatomaceous Earth, fibrosis was only noted at 24 mo, and not at the same severity as in the cristobalite-exposed animals; in animals exposed to cristobalite, fibrosis first observed after 15 mo and was severe by 21 mo; no fibrosis observed in animals exposed to sodium silicate, but alveoli became heavily packed with phagocytic macrophages. Total silica content per lung increased linearly throughout at least 21 mo in each experiment, and total ash weight increased more rapidly than dust was accumulating. Cristobalite produced a greater increment in ash weight than Diatomaceous Earth and sodium silicate. Total amount of silica accumulated varied inversely with the degree of tissue damage occurring, even though atmospheric dust concentrations were comparable for the 3 silica types. Maximum total content of cristobalite reached only 68 mg/lung, while that of Diatomaceous Earth and sodium silicate was 120 mg/lung and 465/lung, respectively. Author noted that siliceous dust that produces cell damage may be cleared more effectively from the lung than innocuous dust.</p>	³¹

Table 4. Repeated dose toxicity studies of Diatomaceous Earth

Test Material Dose/Concentration	Animals/Group	Study Duration	Vehicle	Protocol	Results	Reference
Flux-calcined Diatomaceous Earth (61% cristobalite); 0, 2, and 5 mppcf; mean particle size 0.7 µm	Male mongrel dogs divided as follows in the 0, 2, and 5 mppcf dose groups: 8, 16, and 17 animals, respectively	2.5-yr study	None described	Dogs exposed to test material in exposure chambers for 6 h/d, 5 d/wk for up to 30 mo; an unreported number of dogs were killed at 6 mo, 1 yr, 1.5 yr, 2 yr, and 2.5 yr; one dog in the control and each dose group was killed 10 mo after cessation of exposure to examine recovery; lungs, hilar lymph nodes, heart, liver, kidney, spleen, adrenal, small intestine (duodenum), hepatic lymph nodes, and sections of the trachea, pancreas, and urinary bladder underwent histologic examination	<p>Terminal weights comparable or slightly greater than controls; no changes in hematology during the course of the study; tissues studied other than the lungs had no test material-related changes.</p> <p>At 6 mo, no reaction observed in the 2 mppcf group and minimal intra-alveolar macrophages observed in the 5 mppcf group; however, hilar nodes had greater macrophagic infiltration than rats and guinea pigs described above.</p> <p>At 1 yr, little to no changes observed.</p> <p>At 1.5 yr, clusters of dust cells in alveolar spaces adjacent to bronchioles observed in 5 mppcf group, with hilar lymph nodes enlarged and medulla replaced with hyalinized tissue.</p> <p>At 2 yr, slight perivascular and peribronchiolar localization of macrophages observed in 2 mppcf group that were definite nodules extending into bronchiolar lumina in the 5 mppcf group; hilar lymph nodes were enlarged and diffusely packed with macrophages; medulla had numerous nodules.</p> <p>At 2.5 yr, observations similar to those in the 2 yr group with no significant progression in reactions; no fibrosis evident.</p> <p>In the recovery animals, parenchymal and nodal changes did not increase compared to 2.5 yr group.</p>	¹⁵

Table 5. In vitro genotoxicity studies of Diatomaceous Earth

Concentration/Dose	Vehicle	Test System	Procedure	Results	Reference
0, 50, 150, 500, 1500, or 5000 µg/plate flux-calcined (100% pure)	polyethylene glycol 400	<i>Salmonella typhimurium</i> strains TA 1535, TA 1537, TA 98, and TA 100; <i>Escherichia coli</i> strain WP2 uvr A	Ames test in accordance with OECD TG 471, with and without metabolic activation	Not mutagenic	⁵
0, 2.5, 5, 10, 20, 30 or 40 µg/ml flux-calcined (100% pure)	R0 medium	Mouse lymphoma L5178Y cells	Mammalian cell gene mutation test in accordance with OECD TG 476, with and without S9 metabolic activation	Not mutagenic	⁵
0, 1.25, 2.5, 5, 10, 20, or 40 µg/ml flux-calcined (100% pure)	Minimal essential medium or dimethyl sulfoxide	Human lymphocytes	Mammalian chromosome aberration test in accordance with OECD TG 473, with and without S9 metabolic activation	Not clastogenic	⁵
Natural and flux-calcined Diatomaceous Earth (average diameters 1.3 µm and 2.1 µm, respectively) in addition to titanium dioxide, crocidolite, chrysotile, quartz, and cristobalite; concentration ranges not reported; crystalline silica content of the natural Diatomaceous Earth was 4% quartz and of the flux-calcined Diatomaceous Earth was 40% cristobalite and 2% quartz	Not reported	Cultured CHO cells	Colony-forming efficiency assays; 200 cells seeded/dish and the test dusts added 24 h later; cultures then incubated for 5 d before being fixed and number of colonies containing > 20 cells was determined for each dish.	Similar ranking of toxicity observed as in the cell proliferation assay described above; colony formation was not as inhibited as cell proliferation; results were concentration-dependent	¹⁶
Natural and flux-calcined Diatomaceous Earth as described above	Not reported	Cultured CHO cells	Abnormal nucleus induction assays; cultures prepared in the same manner as the above inhibition of cell proliferation assays, exposed for 2 d and then fixed; percentage of cells containing micronuclei and/or polynuclei was determined for each dish.	Similar qualitative, concentration-dependent results were observed as in the cell proliferation and colony-forming efficiency assays described above	¹⁶
Three different sourced uncalcined Diatomaceous Earth samples (96%-98% pure; 0.6% -1.4% iron impurities) and 2 calcined Diatomaceous Earth samples (~98% pure; 0.7% - 0.9% iron impurities); concentrations not well defined, but at least 3 concentrations per sample were tested starting at 2 µg/cm ² and were up to approximately 40 µg/cm ²	Suspended in sterile tridistilled water; culture medium without serum and complete medium	Syrian hamster embryo (SHE) cells	Cell transformation assay; without metabolic activation	Morphological transformation of the uncalcined and calcined Diatomaceous Earth samples occurred in a dose-dependent manner; authors concluded that samples with fractured surfaces and/or iron-active sites were able to generate reactive oxygen species-induced SHE cell transformation	³²
Uncalcined Diatomaceous Earth (100% amorphous), Diatomaceous Earth heated to 900°C (98.5% amorphous, 1% quartz, <0.5% cristobalite), Diatomaceous Earth heated to 1200°C (51% amorphous, 1% quartz, 48% cristobalite), a generically heated flux-calcined Diatomaceous Earth (53% amorphous, 47% cristobalite), and the generically heated flux-calcined Diatomaceous Earth (42% amorphous, 58% cristobalite) depleted of particles greater than 10 µm; concentrations tested for each material were 4.5, 9, and 18 µg/cm ² (also 36 µg/cm ² for generically heated Diatomaceous Earth)	Culture medium	SHE cells	Cell division aberration assay; without metabolic activation	A concentration-dependent increase in abnormal mitoses frequency was observed with all dusts tested, except uncalcined Diatomaceous Earth at 4.5 and 9 µg/cm ² ; Diatomaceous Earth heated to 900°C and 1200°C appeared “less active” than the uncalcined – the authors theorized this may be due to cytotoxic potential, which appeared “blunted” through heating	³³

Table 5. In vitro genotoxicity studies of Diatomaceous Earth

Concentration/Dose	Vehicle	Test System	Procedure	Results	Reference
Uncalcined Diatomaceous Earth (100% amorphous), Diatomaceous Earth heated to 900°C (98.5% amorphous, 1% quartz, <0.5% cristobalite), Diatomaceous Earth heated to 1200°C (51% amorphous, 1% quartz, 48% cristobalite), a generically heated flux-heated Diatomaceous Earth (53% amorphous, 47% cristobalite), and the generically heated flux-calcined Diatomaceous Earth (42% amorphous, 58% cristobalite) depleted of particles greater than 10 µm; concentrations tested for each material were between 1.9 and 30.4 µg/cm ² (up to 60.8 µg/cm ² for generically heated Diatomaceous Earth)	Culture medium	SHE cells	Cell transformation assay; without metabolic activation	Uncalcined Diatomaceous Earth did not induce morphological transformation while a concentration-dependent increase of the transformation frequency was induced by all other test materials; the heated samples exhibited a certain degree of transformation with the 1200°C heated sample greater than the 900°C (which was weakly active only above 15 µg/cm ²); transformation potential appears to be correlated with the ability to generate radicals	³³
Uncalcined Diatomaceous Earth with 0.03% iron impurities and uncalcined Diatomaceous Earth depleted of iron; concentrations started at 3.5 µg/cm ² and included up to 60 µg/cm ²	Not reported	SHE cells	Cell transformation assay, with and without antioxidants	Concentration-dependent increase in transformation frequency starting at 3.5 µg/cm ² was observed in samples with iron, transforming potency was 1.8-fold less in samples depleted of iron; in presence of antioxidants, transformation frequencies were significantly decreased; authors concluded iron may generate reactive oxygen species that increase transforming potency	³⁴
Uncalcined Diatomaceous Earth with 0.03% iron impurities and uncalcined Diatomaceous Earth depleted of iron; concentrations between 2.25 and 34 µg/cm ²	Not reported	SHE cells	Cell division aberration assay, with and without antioxidants	A significant concentration-dependent increase in frequency of abnormal mitoses was induced by sample with iron; mitotic spindle disturbances, mono- and multi-polar mitoses, and some chromosome lagging were most frequently observed; iron-depleted samples induced abnormal mitoses in a similar manner to the samples with iron; in presence of antioxidants, frequency of abnormal mitoses were significantly decreased	³⁴

Table 6. Dermal irritation and sensitization studies of Diatomaceous Earth

Test Article	Concentration/Dose	Test Population	Procedure	Results	Reference
IRRITATION					
IN VITRO STUDIES					
100% Diatomaceous Earth; flux-calcined	20 mg; undiluted	Reconstituted human epidermis samples	EpiSkin™ reconstituted human epidermis model test in accordance with OECD TG 431; duplicate tissues treated for 3, 60, and 240 min	Non-corrosive; relative mean viability after exposure to test material for 3, 60, and 240 min was 102.8%, 111.3%, and 114.1%, respectively; qualitative evaluation indicated tissue was viable at each time point following exposure to test material; positive and negative controls yielded expected results	5
Diatomaceous Earth; flux-calcined, purity not reported	Not reported	Reconstituted human epidermis samples	EpiSkin™ reconstituted human epidermis model test; tissues treated for 15 min before incubation for 42 h; no further details reported	Not irritating; relative mean viability after exposure to test material was 102.6%; qualitative evaluation indicated tissue was viable following exposure to the test material; positive and negative controls yielded expected results	5
HUMAN					
100% Diatomaceous Earth; flux-calcined	0.02 mg; undiluted	10 subjects	Acute skin tolerance test; 48-h single patch test using Finn Chambers; occluded; test material applied to external face of the arm	Not irritating	38
Product containing 9% - 11% Diatomaceous Earth (Diatomaceous Earth contained < 0.11% respirable crystalline silica); soda ash flux-calcined	Amount not reported; undiluted	11 subjects with sensitive skin	Acute 24-h skin tolerance patch test; occluded; no further details	Not irritating	39
SENSITIZATION					
ANIMAL					
Diatomaceous Earth; flux-calcined, purity not reported	0%, 2.5%, 5%, or 10% in propylene glycol; 25 µl	Groups of 4 female CBA mice	LLNA; animals received test material daily on dorsum of each ear lobe for 3 consecutive days; positive control group received 90% phenylacetaldehyde in a solution of propylene glycol (final concentration 2.5% v/v)	Not sensitizing; all treated animals survived treatment; no clinical signs of toxicity observed in any test groups; stimulation indices (SI) for 2.5%, 5%, and 10% dose groups were 1.13, 0.97, and 0.99, respectively; SI of positive control was 18.43	5
HUMAN					
Cosmetic formulation containing 0.9% - 1.1% Diatomaceous Earth (Diatomaceous Earth contained < 0.11% respirable crystalline silica); soda ash flux-calcined	25 µl; applied neat	100 healthy subjects with normal skin	HRIPT according to Marzulli-Maibach method; test material applied on back of subjects with Finn Chambers on Scanpor®; occluded; duplicate patches without test material applied to serve as control only during the induction phase; induction patches occurred 3 times a week for 3 wk and a 2-wk rest period occurred prior to the single challenge patch; patches were in place for 48 h	Not irritating and not sensitizing	40
PHOTOTOXICITY					
HUMAN					
Product containing 9% - 11% Diatomaceous Earth (Diatomaceous Earth contained < 0.11% respirable crystalline silica); soda ash flux-calcined	0.2 ml; undiluted	10 healthy female subjects	Phototoxicity study of single application of test material on each forearm; occluded for 24 h; one arm was irradiated with UV-A (4 F4OBL with fluorescent tubes; 320-400 nm), while the other arm served as control	Not phototoxic; no skin reactions observed on irradiated product site and control site without product; very slight transient erythema observed in 1 subject on non-irradiated product site	41

Table 7. Occupational exposure studies of Diatomaceous Earth

Diatomaceous Earth Composition	Study Population and Location	Time Frame Examined	Procedure/Parameters Measured/Limitations	Findings	Reference
Quarry dust was essentially amorphous silica with quartz content of crude Diatomaceous Earth being 2%; mill dust had high percentage of cristobalite	869 workers of 5 plants in California, Nevada, and Oregon	1953-1954	X-ray investigation	-9% of the workers had lung changes interpreted as pneumoconiosis and that an equal number had doubtful changes -prevalence of abnormal chest films especially high in employees in mills -exposure in quarries associated with a lower proportion of abnormal films; none of 25 employees who had worked there exclusively for over 5 yr had a positive film, but 40% showed doubtful linear nodular changes	44
Same as above	Follow-up study in 428 workers from one plant from the above study (state not specified); plant included a quarry and a mill	1974; including employees terminated between July 1, 1969 and July 1, 1974	X-ray investigation	-films interpreted as positive for pneumoconiosis (Union for International Cancer Control (UICC)/Cincinnati classification of 1/1) observed in 20 (4.7%) of the workers -another 6 films had a UICC/Cincinnati classification of 1/0 -of these 26, 14 were determined to have findings consistent with Diatomaceous Earth pneumoconiosis, and all but 2 of these 14 had been employed before 1953 -in 129 employees in the industry for 20 yr or more, 13 had positive films considered consistent with Diatomaceous Earth pneumoconiosis, of which 6 had negative films in 1953 -only 4 individuals had complicated or coalescent lesions: these workers had been mill workers employed 27- 46 yr -no massive coalescent lesions or distorting changes noted in the existing work force -researchers pointed out that this evidence agreed with earlier observations indicating that the risk of pneumoconiosis was relatively low in workers whose exposure was confined to crude Diatomaceous Earth, as compared with those exposed to calcined Diatomaceous Earth -researchers noted that strict occupational dust control measures and personal protective equipment led to the near elimination of new cases of Diatomaceous Earth pneumoconiosis	45
Raw material contained ~ 4% crystalline silica; calcined and fluxed-calcined material had 10-20% and 20-25% cristobalite, respectively	2570 white male Diatomaceous Earth mining and processing workers in California; at least 12 mo cumulative service	1942-1987	Mortality patterns analysis; mortality trends assessed in respect of an index of cumulative exposure to crystalline silica and crystalline silica index; workers with known potential occupational asbestos exposure excluded; cigarette smoking was a confounding factor	-all causes combined standardized mortality ratio (SMR) slightly increased when compared with rates among US white males (SMR 1.12; 628 observed) -increased risks from lung cancer (SMR 1.43; 59 observed) and non-malignant respiratory disease (NMRD; excluding infectious diseases and pneumonia; SMR 2.59, 56 observed) were main contributors to the observed excess -excess lung cancer also observed when rates were compared with local county rates instead of the US national rates -increasing gradients of risk detected for lung cancer and NMRD with both crystalline silica exposure indices -researchers stated smoking was not likely to account for all associations between dust exposure and lung cancer -prior to the 1950s, poor dust control measures likely largest contributors to lung cancer and NMRD; the absence of excess lung cancer in workers hired after 1960 and no deaths attributed to pneumoconiosis in workers hired after 1950 indicated exposure reductions were successful in reducing excess risks in workers	46
Same as above	2342 white male Diatomaceous Earth workers; a subset of the above California workers cohort (406 had been excluded due to potential inadequate exposure data or definitive asbestos exposure	1942-1987	Mortality patterns analysis as above; results not likely to be confounded by smoking or asbestos exposure	-mortality excesses detected for NMRD (SMR 2.01) and lung cancer (SMR 1.29) -mortality from NMRD rose sharply with cumulative exposure to respirable crystalline silica (mostly cristobalite), indicating a strong dose-response relationship for crystalline silica and NMRD mortality -while not as strong of a relationship, lung cancer results further support an etiologic role for crystalline silica	47

Table 7. Occupational exposure studies of Diatomaceous Earth

Diatomaceous Earth Composition	Study Population and Location	Time Frame Examined	Procedure/Parameters Measured/Limitations	Findings	Reference
Same as above	1809 white male Diatomaceous Earth workers; a subset of the above California workers cohort; workers had at least 1 yr of exposure to crystalline silica	1942-1987	X-ray investigation	-81 workers (4.5%) had opacities on chest radiographs -age-adjusted relative risk of opacities increased significantly with cumulative exposure to crystalline silica -risk of opacities for cumulative exposure to crystalline silica of 2.0 mg/m ³ -yr was 1.1% when average crystalline silica exposure was < 0.50 mg/m ³ , but was 3.7% when average crystalline silica exposure was > 0.50 mg/m ³	48
Same as above	759 white male Diatomaceous Earth workers; a subset of the above California workers cohort;	1942-1987	X-ray and spirometry investigation; chest radiographs interpreted by the International Labor Office (ILO) system; individual-based reconstructed exposure indices for total dust (largely Diatomaceous Earth) and cristobalite were used in performing regression analyses	-of 492 chest radiographs, 5% had ILO scores > 1/0 and 25% had score of 0/1 or higher -radiographic patterns were not typical of classic silicosis - regression analyses showed there was a relationship between both total cristobalite exposure and total dust exposure and the ILO score -differences observed in spirometric data according to radiographic ILO category, but the results were inconsistent and did not allow for determining if physiologic changes were associated with radiographic change or through confounding factors, such as smoking -researchers noted that recent exposure level may produce radiographic abnormalities, but a demonstrable physiologic effect may not be observed; this decrease in observed effects was noted to be due to modern dust control measures.	49

REFERENCES

1. Nikitakis J, Kowcz A. Web-Based International Cosmetic Ingredient Dictionary and Handbook (wINCI Dictionary). <http://webdictionary.personalcarecouncil.org/jsp/IngredientSearchPage.jsp>. Washington, D.C.: Personal Care Products Council. Accessed 08/09/2021.
2. Burnett CL, Bergfeld WF, Belsito DV, et al. Amended Safety Assessment of Synthetically-Manufactured Amorphous Silica and Hydrated Silica as Used in Cosmetics. Washington, DC. 2019. (Available from the Cosmetic Ingredient Review.: <https://www.cir-safety.org/ingredients>.)
3. International Agency for Research on Cancer. Silica, Some Silicates, Coal Dust and *para*-Aramid Fibrils. Lyon, France. 1997. <https://monographs.iarc.fr/wp-content/uploads/2018/06/mono68.pdf>.
4. Agency for Toxic Substances and Disease Registry (ATSDR). Toxicological Profile for Silica. 2019. <https://www.atsdr.cdc.gov/toxprofiles/tp211.pdf>. Accessed 02/18/2021.
5. European Chemicals Agency. Kieselguhr, soda ash flux-calcined. <https://echa.europa.eu/registration-dossier/-/registered-dossier/15119>. Accessed 10/16/2020.
6. Joint Food and Agriculture Organization of the United Nations (FAO)/World Health Organization (WHO) Expert Committee on Food Additives (JECFA). Diatomaceous Earth. Geneva. 2000. http://www.fao.org/fileadmin/user_upload/jecfa_additives/docs/Monograph1/Additive-151.pdf. Accessed 10/13/2020. (FNP 52 Add 8.)
7. Ghiazza M, Gazzano E, Bonelli B, et al. Formation of a vitreous phase at the surface of some commercial diatomaceous earth prevents the onset of oxidative stress effects. *Chem Res Toxicol*. 2009;22(1):136-145.
8. Haneke KE. Sodium Metasilicate, Anhydrous [6834-92-0], Sodium Metasilicate Pentahydrate [10213-79-3], and Sodium Metasilicate Nonahydrate [13517-24-3]: Review of Toxicological Literature. Research Triangle Park, NC: Integrated Laboratory Systems, Inc.;2002. https://ntp.niehs.nih.gov/ntp/htdocs/chem_background/exsumpdf/sodiummetasilicate_508.pdf. Accessed 10/13/2020.
9. Feigin DS. Misconceptions regarding the pathogenicity of silicas and silicates. *J Thorac Imaging*. 1989;4(1):68-80.
10. Breese ROY, Bodycomb FM. Diatomite. In: Kogel JE, Trivedi NC, Barker JM, Krukowski ST, eds. *Industrial Minerals & Rocks: Commodities, Markets, and Uses*. 7th ed. Littleton, CO: Society for Mining, Metallurgy, and Exploration, Inc.; 2006:433-450.
11. Seppic. 2021. Memo concerning type of Diatomaceous Earth used in cosmetic ingredients. Unpublished data submitted by the Personal Care Products Council on November 22, 2021.
12. Ellis M. 2021. Re: IDPA Comments on the Draft Safety Assessment of Diatomaceous Earth as Used in Cosmetics and Other Matters. Unpublished data submitted by the International Diatomite Producers Association on February 22, 2022.
13. Nattrass C, Horwell CJ, Damby DE, Kermanizadeh A, Brown DM, Stone V. The global variability of diatomaceous earth toxicity: a physicochemical and in vitro investigation. *J Occup Med Toxicol*. 2015;10:23.
14. Ulrichs C, Krause F, Rocks T, Goswami A, Mewis I. Electrostatic application of inert silica dust based insecticides onto plant surfaces. *Commun Agric Appl Biol Sci*. 2006;71(2):171-178.
15. Wagner WD, Fraser DA, Wright PG, Dobrogorski OJ, HE S. Experimental evaluation of the threshold limit of cristobalite - calcined diatomaceous earth. *Am Ind Hyg Assoc J*. 1968;29(3):211-221.
16. Hart GA, Hesterberg TW. In vitro toxicity of respirable-size particles of diatomaceous earth and crystalline silica compared with asbestos and titanium dioxide. *J Occup Environ Med*. 1998;40(1):29-42.
17. Seixas NS, Heyer NJ, Welp EAE, Checkoway H. Quantification of historical dust exposures in the diatomaceous earth industry. *Ann Occup Hyg* 1997;41(5):591-604.

18. Biotechmarine. 2021. Diatomaceous Earth Product Information. Unpublished data submitted by the Personal Care Products Council on June 25, 2021.
19. Biotech Marine. 2017. Statement Phycocorail Composition File (Lithothamnion Calcareum Powder and Diatomaceous Earth).
20. U.S. Food and Drug Administration Center for Food Safety & Applied Nutrition (CFSAN). Voluntary Cosmetic Registration Program - Frequency of Use of Cosmetic Ingredients. College Park, MD. 2022. (Obtained under the Freedom of Information Act from CFSAN; requested as "Frequency of Use Data" January 4, 2022; received January 11, 2022.)
21. Personal Care Products Council. 2021. Concentration of Use by FDA Product Category: Diatomaceous Earth. Unpublished data submitted by the Personal Care Products Council on December 16, 2021.
22. European Union. Regulation (EC) No. 1223/2009 of the European Parliament and of the Council of 30 November 2009 on Cosmetic Products. 2009. <http://eur-lex.europa.eu/LexUriServ/LexUriServ.do?uri=OJ:L:2009:342:0059:0209:en:PDF>. Accessed 11/9/2017.
23. Bunch TR, Bond C, Buhl K, Stone D. Diatomaceous Earth General Fact Sheet. <http://npic.orst.edu/factsheets/degen.html>. National Pesticide Information Center, Oregon State University Extension Services. Accessed 11/16/2020.
24. U.S. Food and Drug Administration Select Committee on GRAS Substances (SCOGS). Evaluation of the Health Aspects of Certain Silicates as Food Ingredients. 1979. SCOGS-61. Accessed 01/11/2021. (Available through National Technical Reports Library: PB301402.)
25. Janicijevic J, Krajisnik D, Calija B, et al. Inorganically modified diatomite as a potential prolonged-release drug carrier. *Mater Sci Eng C Mater Biol Appl*. 2014;42:412-420.
26. Delasoie J, Zobi F. Natural diatom biosilica as microshuttles in drug delivery systems. *Pharmaceutics*. 2019;11(10):537.
27. U.S. Food and Drug Administration Center for Drug Evaluation and Research. Inactive ingredient Search for Approved Drug Products. U.S. Food and Drug Administration. <https://www.accessdata.fda.gov/scripts/cder/iig/index.cfm> 2021. Accessed 02/19/2021.
28. Delgarm N, Ziaee M, McLaughlin A. Enhanced-efficacy Iranian diatomaceous earth for controlling two stored-product insect pests. *J Econ Entomol*. 2020;113(1):504-510.
29. Saeed N, Farooq M, Shakeel M, Ashraf M. Effectiveness of an improved form of insecticide-based diatomaceous earth against four stored grain pests on different grain commodities. *Environ Sci Pollut Res*. 2018;25(17):17012-17024.
30. Bertke EM. The effect of ingestion of diatomaceous earth in white rats: A subacute toxicity test. *Toxicol Appl Pharmacol*. 1964;6:284-291.
31. Pratt PC. Lung dust content and response in guinea pigs inhaling three forms of silica. *Arch Environ Health*. 1983;38(4):197-204.
32. Elias Z, Poirot O, Daniere MC, et al. Cytotoxic and transforming effects of silica particles with different surface properties in Syrian hamster embryo (SHE) cells. *Toxicol In Vitro*. 2000;14(5):409-422.
33. Elias Z, Poirot O, Fenoglio I, et al. Surface reactivity, cytotoxic, and morphological transforming effects of diatomaceous earth products in Syrian hamster embryo cells. *Toxicol Sci*. 2006;91(2):510-520.
34. Elias Z, Poirot O, Daniere MC, et al. Role of iron and surface free radical activity of silica in the induction of morphological transformation of Syrian hamster embryo cells. *Ann Occup Hyg*. 2002;46(Suppl 1):53-57.
35. Hilding AC, Hilding DA, Larson DM, Aufderheide AC. Biological effects of ingested amosite asbestos, taconite tailings, diatomaceous earth and Lake Superior water in rats. *Arch Environ Health*. 1981;36(6):298-303.

36. Adamis Z, Tatrai E, Honma K, Six E, Ungvary G. In vitro and in vivo tests for determination of the pathogenicity of quartz, diatomaceous earth, mordenite and clinoptilolite. *Ann Occup Hyg.* 2000;44(1):67-74.
37. Maeda H, Ford J, Williams MG, Dodson RF. An ultrastructural study of acute and long-term lung response to commercial diatomaceous earth. *J Comp Pathol.* 1986;96(3):307-317.
38. Bioethic. 2002. Study Summary: Study of acute skin tolerance of a cosmetic product after a 48-hour single patch-test (Micro Algues 80 - Diatomaceous Earth). Unpublished data submitted by the Personal Care Products Council on June 25, 2021.
39. Palmer Research. 2003. Study summary report: Determination of the irritation potential of a cosmetic product on human subjects: 24-hour single occlusive patch test (Phycocorail tested at 100%). Unpublished data submitted by the Personal Care Products Council on June 25, 2021.
40. Groupe DermScan. 2014. Assessment of the sensitizing potential of a cosmetic product: Final clinical security test under dermatological control (product LCA13049-13P379-1 was 10% Phycocorail which contains 9-11% Diatomaceous Earth). Unpublished data submitted by the Personal Care Products Council on June 25, 2021.
41. Palmer Research. 1995. Evaluation du potentiel phototoxique apres application et exposition uniques sur 10 volontaires (Phycocorail). (English translation using Google translate provided). Unpublished data submitted by the Personal Care Products Council on June 25, 2021.
42. Seppic. 2001. HET-CAM Test: Phycocorail (contains 57-61% Lithothamnion Calcareum Powder and 8-11% Diatomaceous Earth). Unpublished data submitted by the Personal Care Products Council on November 30, 2021.
43. Dutra FR. Diatomaceous earth pneumoconiosis. *Arch Environ Health.* 1965;11(5):613-619.
44. Cooper WC, Cralley LJ. Pneumoconiosis in diatomite mining and processing. Washington, DC: U.S. Department of Health, Education and Welfare;1958.
45. Cooper WC, Jacobson G. A 21-year radiographic follow-up of workers in the diatomite industry. *J Occup Med* 1977;19(8):563-566.
46. Checkoway H, Heyer NJ, Demers PA, Breslow NE. Mortality among workers in the diatomaceous earth industry. *Br J Ind Med.* 1993;50(7):586-597.
47. Checkoway H, Heyer NJ, Seixas NS, et al. Dose-response associations of silica with nonmalignant respiratory disease and lung cancer mortality in the diatomaceous earth industry. *Am J Epidemiol.* 1997;145(8):680-688.
48. Hughes JM, Weill H, Checkoway H, et al. Radiographic evidence of silicosis risk in the diatomaceous earth industry. *Am J Respir Crit Care Med.* 1998;158(3):807-814.
49. Harber P, Dahlgren J, Bunn W, Lockey J, Chase G. Radiographic and spirometric findings in diatomaceous earth workers. *J Occup Environ Med.* 1998;40(1):22-28.
50. National Institute for Occupational Safety and Health (NIOSH). NIOSH Pocket Guide to Chemical Hazards. Centers for Disease Control and Prevention (CDC). <https://www.cdc.gov/niosh/npg/npgd0552.html> 2019. Accessed 10/13/2020.
51. Occupational Safety and Health Administration (OSHA). Silica, Amorphous Including Natural Diatomaceous Earth. United States Department of Labor. <https://www.osha.gov/chemicaldata/613.2021>. Accessed 08/23/2022.

2022 FDA VCRP Raw Data

DIATOMACEOUS EARTH	03C	Eye Shadow	1
DIATOMACEOUS EARTH	03D	Eye Lotion	1
DIATOMACEOUS EARTH	05G	Tonics, Dressings, and Other Hair Grooming Aids	1
DIATOMACEOUS EARTH	07B	Face Powders	5
DIATOMACEOUS EARTH	07E	Lipstick	6
DIATOMACEOUS EARTH	08A	Basecoats and Undercoats	1
DIATOMACEOUS EARTH	08E	Nail Polish and Enamel	49
DIATOMACEOUS EARTH	09A	Dentifrices	11
DIATOMACEOUS EARTH	10A	Bath Soaps and Detergents	1
DIATOMACEOUS EARTH	10B	Deodorants (underarm)	3
DIATOMACEOUS EARTH	10E	Other Personal Cleanliness Products	2
DIATOMACEOUS EARTH	12A	Cleansing	4
DIATOMACEOUS EARTH	12C	Face and Neck (exc shave)	8
DIATOMACEOUS EARTH	12D	Body and Hand (exc shave)	1
DIATOMACEOUS EARTH	12F	Moisturizing	6
DIATOMACEOUS EARTH	12G	Night	1
DIATOMACEOUS EARTH	12H	Paste Masks (mud packs)	25
DIATOMACEOUS EARTH	12J	Other Skin Care Preps	9