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# Amended Safety Assessment of 5-Amino-6-Chloro-*o*-Cresol as Used in Cosmetics

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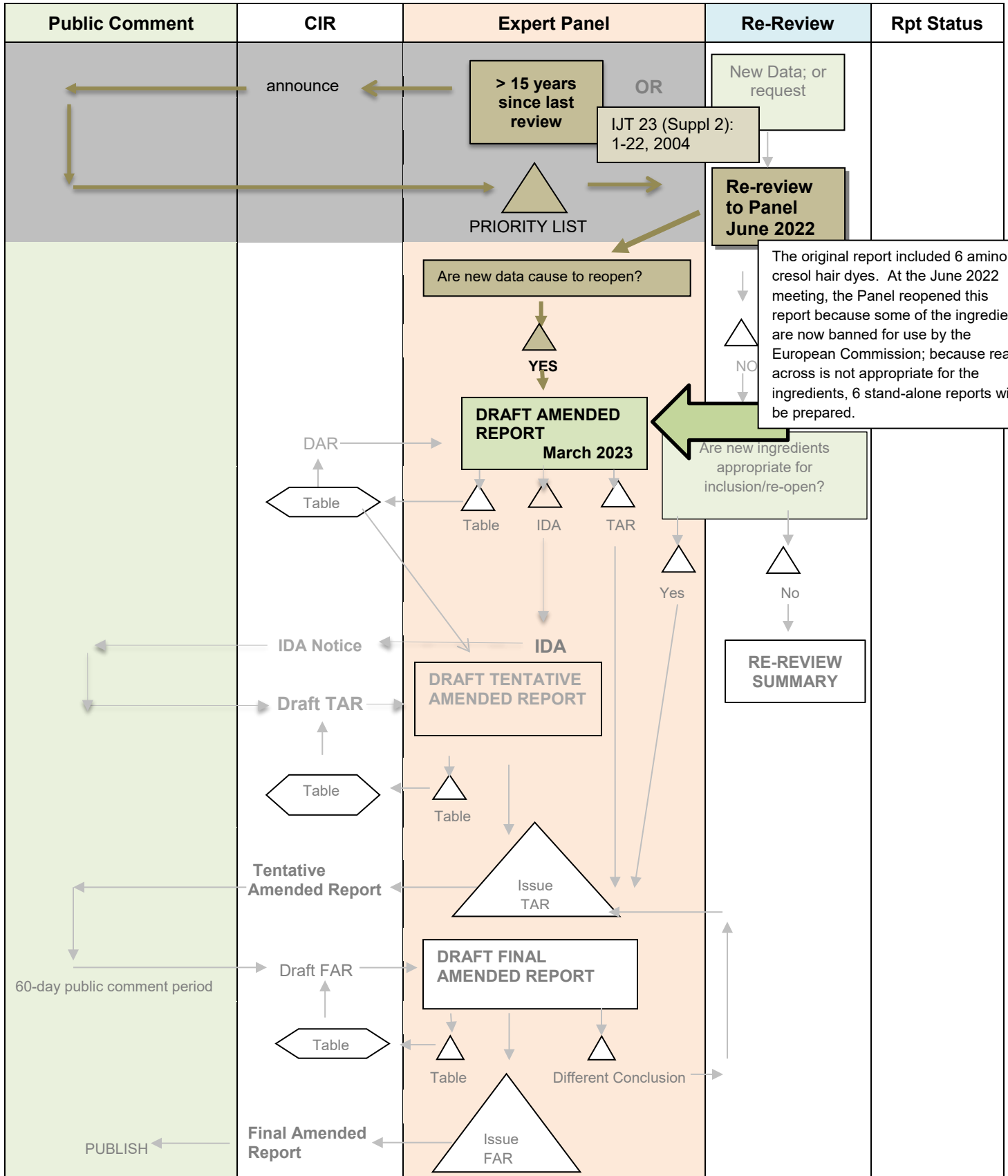
Status: Draft Amended Report for Panel Review  
Release Date: February 10, 2023  
Panel Meeting Date: March 6-7, 2023

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.; 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. The Cosmetic Ingredient Review (CIR) Executive Director is Bart Heldreth, Ph.D. This safety assessment was prepared by Christina Burnett, Senior Analyst/Writer, CIR.

# RE-REVIEW FLOW CHART

INGREDIENT/FAMILY 5-Amino-6-Chloro-*o*-Cresol

MEETING March 2023





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### Memorandum

To: Expert Panel for Cosmetic Ingredient Safety Members and Liaisons  
From: Christina L. Burnett, MSES, Senior Scientific Analyst/Writer, CIR  
Date: February 10, 2023  
Subject: Amended Safety Assessment of 5-Amino-6-Chloro-*o*-Cresol

The Panel previously reviewed the safety of 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, 5-Amino-4-Chloro-*o*-Cresol, 5-Amino-6-Chloro-*o*-Cresol, and 4-Chloro-2-Aminophenol in an assessment that was published in 2004 (identified as *originalreport\_5-Amino-6-Chloro-o-Cresol\_032023* in the pdf document). In June 2022, the Panel re-opened the safety assessment for these ingredients due to some of these hair dyes being banned for use in cosmetics by the European Commission. Because the Panel determined that data for these amino cresol hair dye ingredients could not be read-across, rather than including all 6 ingredients in one amended report, re-reviews of each hair dye included in that original 2004 report will now be presented as individual stand-alone reports. Accordingly, enclosed is the Draft Amended Report on the Safety Assessment of 5-Amino-6-Chloro-*o*-Cresol as Used in Cosmetics (*report\_5-Amino-6-Chloro-o-Cresol\_032023*). In the original report, the Panel concluded that 5-Amino-6-Chloro-*o*-Cresol is safe as used in oxidative and non-oxidative (semi-permanent) hair dyes.

According to 2022 VCRP survey data, 5-Amino-6-Chloro-*o*-Cresol has 27 reported uses in hair coloring products. The results of the concentration of use survey conducted by the Council in 2021 reported that 5-Amino-6-Chloro-*o*-Cresol is used at up to 0.24% in hair dyes and colors (*data\_5-Amino-6-Chloro-o-Cresol\_032023*). When the original safety assessment was published in 2004, 5-Amino-6-Chloro-*o*-Cresol was reported to have no uses, according to 1998 VCRP data. However, according to industry survey data submitted in 1996, 5-Amino-6-Chloro-*o*-Cresol was reported to be used at up to 2% in hair dyes and colors. As per the Panel's request at the December 2022 meeting, an updated use table format has been implemented. The frequency and concentration of use is presented both cumulatively by likely duration and exposure and individually by product category.

Since the June meeting, no new data have been submitted.

Additional supporting documents for this report package include a flow chart (*flow\_5-Amino-6-Chloro-o-Cresol\_032023*), report history (*history\_5-Amino-6-Chloro-o-Cresol\_032023*), a search strategy (*search\_5-Amino-6-Chloro-o-Cresol\_032023*), a data profile (*datapofile\_5-Amino-6-Chloro-o-Cresol\_032023*), transcripts from the meeting at which the re-review was discussed (*transcripts\_5-Amino-6-Chloro-o-Cresol\_032023*), and the minutes from all the meetings at which 5-Amino-6-Chloro-*o*-Cresol was discussed during the original review (*originalminutes\_5-Amino-6-Chloro-o-Cresol\_032023*).

If no further data are needed to reach a conclusion of safety, the Panel should formulate a Discussion and issue a Tentative Amended Report. However, if additional data are required, the Panel should be prepared to identify those needs and issue an Insufficient Data Announcement.

### **5-Amino-6-Chloro-*o*-Cresol History**

**2004**– The CIR’s Final Report on the Safety Assessment of 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, 5-Amino-4-Chloro-*o*-Cresol, 5-Amino-6-Chloro-*o*-Cresol, and 4-Chloro-2-Aminophenol in the *IJT* after the report was finalized by the Panel in 2000. Based on the available animal and clinical data available at that time, the Panel concluded that 5-Amino-6-Chloro-*o*-Cresol is safe as used in oxidative and non-oxidative (semi-permanent) hair dyes.

**June 2022** – Review of the available published literature since 2000 was conducted in accordance to CIR Procedures regarding re-review of ingredients after ~15 years. The Panel re-opened the safety assessment for this ingredient, due to it being banned for use in cosmetics by the European Commission.

**5-Amino-6-Chloro-*o*-Cresol Data Profile\* - March 2023 - 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 <sub>ow</sub>	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
<b>5-Amino-6-Chloro-<i>o</i>-Cresol</b>	OX		OX	X	OX	O	O	X	O	X		O	OX	OX					OX			OX					OX		

\* "X" indicates that new data were available in a category for the ingredient. "O" indicates data were reported in the original safety assessment.

**5-Amino-6-Chloro-o-Cresol**

Ingredient	CAS #	PubMed	FDA	HPVIS	NIOSH	NTIS	NTP	FEMA	EU	ECHA	ECETOC	SIDS	SCCS	AICIS	FAO	WHO	Web
5-Amino-6-Chloro-o-Cresol	110102-86-8	√	√	√	√	√	√	√	√	√	√	√	√	√	√	√	√

**Search Strategy (from 2002 on)****PubMed**

((“5-Amino-6-Chloro-o-Cresol”) OR (84540-50-1[EC/RN Number])) – 1 hit; 0 relevant

**ECHA**

No dossiers listed.

*Internet searches using trade names and other technical names. No 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/ingredientspackaginglabeling/gras/default.htm>
- SCOGS database: <http://www.fda.gov/food/ingredientspackaginglabeling/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](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](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/](http://www.who.int/biologicals/technical_report_series/en/)
- [www.google.com](http://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

**JUNE 2022 PANEL MEETING – RE-REVIEW CONSIDERATION (WITH SEVERAL OTHER HAIR DYES)**

**Belsito's Team Meeting – June 16, 2022**

**Dr. Belsito** - So hair dyes, this is going to take a... there is more than one hair dye here. I thought we were only going to look at one at a time. What's going on here? Reviewed as a group before.

**Monice Fiume (CIR)** - They were not, but being that this is the first time groups of reviews have been brought to you are the rereview documents. We were trying to figure out if there were ways to group hair dyes or preservatives or something like that together because they were similar types of functions. But I don't think this was the best example in retrospect.

**Dr. Belsito** - Yeah. So then let's go through this. So Orange 3 is now been banned in Europe, Acid Orange 3. I think we need to reopen it not only because of that, but in the use section, it says it's used and it's a new product now, which is a nail enamel.

And then the cresols are also, I think, problematic and need to be reopened. Some of them clearly seem to have carcinogenicity activity and I have a note here that seek Council comments and wave 3 of the cresols.

Will this be addressed in the rereview before publication? Let me get to wave 3.

**Dr. Snyder** - PCPC comments.

**Dr. Belsito** - Yeah, I'm just. So yeah.

**Christina Burnett (CIR)** - There was a typo in my memo that they pointed out at the very end. Two of the ingredients. Uh yeah, I think it's two or three of the ingredients that I said were on Annex Two are actually on Annex 3.

**Dr. Belsito** - Yeah. So this is starts on PDF page 8. I mean I think we need to open up the cresols as well.

**Dr. Liebler** - I agree.

**Dr. Belsito** - Because I think the amino position has significant effects on the toxicity. Then back to the next set of hair dyes. It was so it was just cresols and Acid Orange 3, right?

**Dr. Liebler** - There's one more.

**Dr. Belsito** - Oh yeah, the N,N Bis 2-Hydroxyethyl p-Phenylenediamine Sulfate. I didn't make any comment on that, so I don't think that I felt it needed to be reopened.

**Dr. Liebler** - Yeah, I thought it was a do not reopen unless Don identifies a rational rationale having to do with the EU, perhaps. Doesn't look like it, so do not reopen.

**Dr. Snyder** - Was the nitrosating issue in the original?

**Dr. Belsito** - It's. Yeah, the so the European Commission further advises this hair dye ingredient is a tertiary, meaning that is prone to nitrosation and should not be used in combination with nitrosating \*(inaudible) substances. I guess that is usually read in our discussion, not a conclusion. So you know in the rereview summary where we say we've decided not to reopen it, we can just point that out.

**Dr. Liebler** - Yeah.

**Dr. Belsito** - Yeah, I mean the you didn't ban at, they just issued caution when he you know with use.

**Dr. Liebler** - Right.

**Dr. Belsito** - And they limited the nitrosamine content should be less than 50 parts per billion.

When?

**Dr. Snyder** - They asked, well, they also said it was safe up to 2.5%, my notes say.

**Dr. Belsito** - Yeah. And what is the current use?

**Dr. Snyder** - I don't know.

**Christina Burnett (CIR)** - 1.3 is the maximum.

**Dr. Belsito** - To 1.3 right.

**Christina Burnett (CIR)** - Yes.

**Dr. Belsito** - Yeah. So it's well below what the EU restricted. So I don't think we need to reopen it. And then just in the discussion or in some point the document put about the \*(inaudible). But the cresols and the Acid Orange 3 I think unfortunately need to be reopened.

**Carol Eisenmann (PCPC)** - Hi I have one request for the cresols that they could be in the same report but all the data on each cresols be kept together because I think read across as we've said before on these materials is not appropriate. This was done before you started looking at each hair dye individually.

**Dr. Belsito** - Yeah. I agree, Carol.

**Monice Fiume (CIR)** - I'm sorry. Carol, can you please repeat what you are, clarify what you said?

**Carol Eisenmann (PCPC)** - That'd be nice. For all the so you, you can have them in the same report, but like all the data be in for one ingredient be together. So you can see what's the data on that ingredient rather than you know sometimes you're having a paragraph that has all the you summarize all the data, the acute tox data on all of the ingredients in one paragraph make it, you know, separate out. In other words, you the acute chronic reproductive development for one ingredient and then go to the next one and go through the order. In other words, it's going to be like several separate reports. In one report, rather than or make them separate reports because they should not read across isn't appropriate for them.

**Dr. Liebler** - So.

Yeah, it might be tricky to do that. I mean, one thing that could be done is the endpoint data summary tables could be organized by ingredient.

**Dr. Belsito** - Right.

**Christina Burnett (CIR)** - We can do that.

**Dr. Liebler** - And you could, like I don't mean to dismiss your suggestion, Carol entirely. But the, best way to get an eagle eye view of the data would be those summary tables and that should be, I agree that should be organized by ingredient.

**Dr. Belsito** - Exactly.

**Dr. Liebler** - And then you know, whatever Christina. Uh, you know, can come up with in terms of sort of organizing the various tox endpoints in the report text by ingredients to the maximum extent that's possible. I agree that's desirable.

**Dr. Belsito** - OK. Any other comments on this? Cresols are going to be fun.

**Christina Burnett (CIR)** - Wait until you see the other two I'm working on.

**Dr. Belsito** - Oh Lord.

**Christina Burnett (CIR)** - Sorry.

#### Cohen's Team Meeting – June 16, 2022

Minutes not captured.

#### Full Panel Meeting – June 17, 2022

**Dr. Bergfeld** - OK, we're off to the next set of items, which is other items called Hair Dyes. Doctor Belsito.

**Dr. Belsito** - OK, so this is not a rereview of one hair dye, that's it's a rereview of several. So we have Acid Orange 3, we have NN, Bis 2 hydroxyethyl parafenylenediamine sulfate. And then we have the cresols and the amino phenols. And we felt that among this group. We need to reopen Acid Orange 3. We need to reopen the cresol aminophenol group, but we did not need to reopen the NN, Bis 2 hydroxyethyl parafenylenediamine sulfate.

**Dr. Bergfeld** - Is there a second on the?

**Dr. Cohen** - 2nd.

**Dr. Bergfeld** - Any further discussion regarding which ones will be reopening?

**Dr. Belsito** - Uh, yeah. So the only discussion really is whether we do the cresol aminophenol group as a whole group, because the actually the positioning of the amino group on the cresol may have significant result in significant differences in the toxicology of the material. It was suggested that by our panel that they all be included in the same report. But that particularly would be presenting the data on toxicity etcetera that instead of as we typically would do like acute oral, you know subchronic chronic that we do that for each. So we do 6-amino-m-cresol and then we go through the various oral studies for that. Then we do 4 amino cresol and do all the tox studies for that so. It will be much clearer in our minds what we have for each of the different materials in this group, because I suspect that we may find that some are safe and some are insufficient. Maybe some should be banned, I don't know, but.

**Dr. Bergfeld** - I mean. I want to ask Bart about your recommendation.

**Dr. Bart Heldreth** - I think that sounds perfect. I think that sounds perfectly fine. You know we need to look at all of these one way or another, and it certainly makes complete sense to me to pull these out and make it very clear that they're separate and that there's really no chance for read across between them and that they're individuals. I think that makes perfect sense.



**Dr. Bergfeld** - OK, how about the orange dye? Anyone want to make a comment on that one? That one is going to be reopened at least this.

**Dr. Belsito** - Yeah. There's new data and it's just been banned by the EU. So I think we need to look at it.

**Dr. Bergfeld** - OK.

**Dr. Cohen** - Yeah, done. We had a lot of deliberation over this and it seems like there's also a paucity of data that may result in from the ban that results in the ban and we had gone back and forth whether this might not be reopened and put into a rereview summary, but I think we came around several times to your team's conclusion.

**Dr. Bergfeld** - Well, we are then voting on the reopening of the acid orange and we're not reopening the Bis, but also reopen the creosol. Is that right?

**Dr. Belsito** - Correct.

**Dr. Klaassen** - Correct.

**Dr. Bergfeld** - OK, I'm going to call the question then all those opposing. Abstaining. I assume it's unanimous that we're moving forward with reopening of two groups here.

### DECEMBER 2-3, 1998 PANEL MEETING

Dr. Belsito recalled that the following informal data requests on this group of ingredients were issued at the September 10, 1998 Team meeting:

- (1) Concentration of use
- (2) Physical and chemical properties
- (3) Method of manufacture
- (4) Impurities data, especially regarding the presence of *m*-cresol
- (5) UV absorption data; if absorption occurs in the UVA or UVB range, photosensitization data may be needed
- (6) Types of hair dye products (semi-permanent or oxidative) and the rate of reaction (bioavailability)
- (7) Metabolism data; if metabolism is not similar to that of 4-Amino-2-Hydroxytoluene and *p*-, *m*-, and *o*-Aminophenol (ingredients already reviewed by CIR), the following data are needed:
  - a. 28-day dermal toxicity data with histopathology
  - b. dermal reproductive toxicity data
  - c. two genotoxicity studies, one using a mammalian system; if positive, a 2-year dermal carcinogenicity study performed using NTP methods.

Dr. Belsito noted that because these ingredients are used in hair dyes and because hair dyes are exempt from sensitization and photosensitization testing as long as the requirement of testing prior to use appears on the label, his Team determined that item 5 above could be deleted. Dr. Belsito said that item 5 should be deleted, because, even if the UV absorption data were positive, the Panel would not have the authority to ask for photosensitization data.

Dr. Schroeter agreed with the revised list of data requests (item 5 deleted).

Dr. Bailey said that he is unsure of how the legal and regulatory status of an ingredient impacts the CIR review process. He said that if there are data that relate to safety, regardless of whether the FDA has legal authority to act, these data should still be of concern to the Panel.

Dr. Belsito said that even if the ingredients were found to be photosensitizers, this would not be a reason for saying that they are unsafe for use in hair dyes, because hair dyes carry a warning about possible sensitization and the need to test prior to use.

Dr. Bailey said that photosensitization is not necessarily being referred to in this case, but, more so, contact sensitization.

Dr. McEwen said that the CIR Procedures do not preclude the Panel from requesting any data that are needed. He said that the Panel needs to determine whether the patch test requirement on the product label sufficiently addresses the Panel's concern about photosensitization, not from a theoretical standpoint, but from the use standpoint of hair dyes.

Concerning the list of data needs included at the beginning of this section, Dr. Belsito said that items 1-4 and 6-7 should be requested for all of the ingredients included in the review. He also reiterated that if the metabolism of these ingredients is not similar, then additional data (e.g. 28-day dermal toxicity data) will be needed.

Dr. McEwen asked if the Panel could use the information on skin penetration from Dr. Walters' presentation to do some modeling on these ingredients to determine if 28-day dermal toxicity data would be needed. In other words, he wanted to know if the Panel would agree to review skin penetration modeling data before requesting 28-day dermal toxicity data.

Dr. Andersen said that after reviewing skin penetration modeling data, the Panel has the option of issuing an Insufficient Data Announcement if these data are not found to be sufficient.

Dr. Belsito said that Dr. Walters presented models that were based on absorption against a barrier of the stratum corneum and data indicating that the forehead is a very absorptive surface, more so than other areas of the body. Dr. Belsito also noted that the follicular shunting mechanism (which is discounted by the models, because, in general, it is not a major area of absorption) would be much more important for a hair dye. Dr. Belsito said that if the skin modeling results indicated a high extent of ingredient absorption, then the 28-day dermal toxicity data would be needed. However, he said that if the results indicated low absorption, he would still want to know what the results would be in a mouse or human, both of which have many hair follicles. He concluded that the computer-generated model would not be useful to him in the present safety assessment.

Someone in the audience commented that the models were generated on specific chemical compounds with similar structures, and that it is possible that the Panel will need absorption data on all four hair dyes included in the safety assessment in order to generate the model.

Dr. Klaassen said that having heard the presentation on skin absorption, he would like for the Panel to include the octanol/water partition coefficient in its request for data on chemical and physical properties. He said that this is the most important chemical parameter that the Panel could have on any ingredient.

Dr. Bailey urged the Panel to be very cautious and be sure to ask certain questions before compounds (especially aromatic amines) are grouped for review in a single report and, potentially, data on one ingredient are wrongfully extrapolated to others.

Dr. Andersen said that the effort by CIR to maximize the benefit from the effort of each review may lead to the creation of as large a family of ingredients as is reasonable. He noted that during reviews by the Panel, any Panel member has an opportunity to recommend the exclusion any ingredient(s) that should not be included in the review.

Dr. Bailey recommended that for ingredients that are reviewed as groups, a table should be created (as part of the report) that indicates which tests have been done on which ingredients.

Dr. Bergfeld said that it was brought to her attention by Dr. Belsito and others that there was a recent hair dye study (4,000 individuals) showing some safety parameters that should be incorporated into CIR's data bank and, perhaps, should be made available for use in the present safety assessment.

Dr. Bailey said that another hair dye study by the American Cancer Society will be published soon. He said that this is a follow-up study to one that was done a few years ago.

Based on the preceding discussion, the following data are needed for completion of the safety assessment on 6-Amino-m-Cresol, 6-Amino-o-Cresol, 4-Amino-m-Cresol, 5-Amino-4-Chloro-o-Cresol, 5-Amino-6-Chloro-o-Cresol, and 4-Chloro-2-Aminophenol (data needed on all ingredients):

- (1) Concentration of use
- (2) Physical and chemical properties
- (3) Method of manufacture
- (4) Impurities data, especially regarding the presence of m-cresol
- (5) Types of hair dye products (semi-permanent or oxidative) and the rate of reaction (bioavailability)
- (6) Metabolism data; if metabolism is not similar to that of 4-Amino-2-Hydroxytoluene and *p*-, *m*-, and *o*-Aminophenol (ingredients already reviewed by CIR), the following data are needed:
  - a. 28-day dermal toxicity data with histopathology
  - b. dermal reproductive toxicity data
  - c. two genotoxicity studies, one using a mammalian system; if positive, a 2-year dermal carcinogenicity study performed using NTP methods.

Note: The Panel responded to a suggestion that skin penetration modeling might help resolve some of the questions by noting that such an approach probably would not be useful for products that are used on the hair follicle rich scalp and could also contact the skin of the forehead.

An Insufficient Data Announcement containing the preceding data requests will be issued.

#### **JUNE 14-15, 1999 PANEL MEETING**

Dr. Belsito recalled that an insufficient data announcement with the following data requests was issued at the December 2-3, 1998 Panel meeting.

- (1) Concentration of use
- (2) Physical and chemical properties
- (3) Method of manufacture
- (4) Impurities data, especially regarding the presence of *m*-cresol
- (5) Types of hair dye products (semi-permanent or oxidative) and the rate of reaction (bioavailability)
- (6) Metabolism data; if metabolism is not similar to that of 4-Amino-2-Hydroxytoluene and *p*-, *m*-, and *o*-Aminophenol (ingredients already reviewed by CIR), the following data are needed:
  - a. 28-day dermal toxicity data with histopathology
  - b. dermal reproductive toxicity data
  - c. two genotoxicity studies, one using a mammalian system; if positive, a 2-year dermal carcinogenicity study performed using NTP methods.

Note: The Panel responded to a suggestion that skin penetration modeling might help resolve some of the questions by noting that such an approach probably would not be useful for products that are used on the hair follicle rich scalp and could also contact the skin of the forehead.

Dr. Belsito noted that, of the data requests listed, current concentration of use data and impurities data (only on 4-amino-*m*-cresol) were received from the cosmetics industry. He also stated that the CIR report contains a good amount of genotoxicity data on some, but not all, of the ingredients and that there is no information indicating how these chemicals are metabolized. Thus, his Team concluded that the current report is insufficient for arriving at a conclusion on the safety of these ingredients in cosmetics.

Dr. Belsito said that if the Panel continues to need data on chemical and physical properties, including the octanol/water partition coefficient, then impurities data (especially, regarding the presence of *m*-cresol and other organic molecules and heavy metals - modification of item 4 above) are needed. He noted that the impurities data are needed on all ingredients except 4-amino-*m*-cresol (data already received on this ingredient). Dr. Belsito added that the Panel still needs items 5 and 6 from the list of data needs, and that item 6c should refer to genotoxicity studies on 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol.

Dr. Schroeter said that his Team requested that 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol also be added to item 6c.

Dr. Shank said that a mammalian mutagenicity assay is needed on 4-Chloro-2-Aminophenol and that both mammalian and bacterial mutagenicity assays are needed on 6-Amino-*o*-Cresol.

The Panel voted unanimously in favor of issuing a Tentative Report with an insufficient data conclusion on 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, 5-Amino-4-Chloro-*o*-Cresol, 5-Amino-6-Chloro-*o*-Cresol, and 4-Chloro-2-Aminophenol. The data needed in order for the Panel to complete its safety assessment of this group of ingredients are listed in the report discussion as follows:

- (1) Physical and chemical properties, including the octanol/water partition coefficient
- (2) Impurities data, for all except 4-Amino-*m*-Cresol, especially regarding the presence of heavy metals, *m*-cresol, and other organic molecules
- (3) Types of hair dye products (semi-permanent or oxidative) in which these ingredients are used and the rate of reaction (bioavailability) in the hair dye product
- (4) Metabolism data; if metabolism is not similar to that of 4-Amino-2-Hydroxytoluene and *p*-, *m*-, and *o*-Aminophenol (ingredients already reviewed by CIR), the following data are needed:
  - a. 28-day dermal toxicity data with histopathology
  - b. dermal reproductive and developmental toxicity data
  - c. for 5-Amino-6-Chloro-*o*-Cresol, 5-Amino-4-Chloro-*o*-Cresol, and 6-Amino-*o*-Cresol, two genotoxicity studies, one using a mammalian system; for 4-Chloro-2-aminophenol, one genotoxicity study in a mammalian system; if any of these tests for any ingredient are positive, a 2-year dermal carcinogenicity study performed using NTP methods may be needed.

#### **DECEMBER 20-21, 1999 PANEL MEETING**

Because a significant amount of data was received one week before the Panel meeting, the Panel voted in favor of tabling any further discussion on this group of ingredients until the February 14-15, 2000 Panel meeting.

#### **FEBRUARY 14-15, 2000 PANEL MEETING**

Dr. Belsito noted that the report on this group of ingredients was tabled at the December 20-21, 1999 Panel meeting because of the large data submissions on 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol that were received. He also noted that additional data on 6-Amino-*m*-Cresol and 4-Amino-*m*-Cresol were received prior to today's meeting. Some of the information received indicates that these two dyes could be used in oxidative hair dyes. However, information indicating whether or not they are used in nonoxidative or semipermanent hair dyes was not received.

After reviewing all of the data on the safety of these ingredients, Dr. Belsito's Team concluded that all six are safe as used in oxidative hair dyes and that the following ingredients are safe as used in nonoxidative hair dyes: 6-Amino-*m*-Cresol, 4-Amino-*m*-Cresol, 5-Amino-4-Chloro-*o*-Cresol, and 5-Amino-6-Chloro-*o*-Cresol. The Belsito Team also concluded that the available data are insufficient for determining the safety of 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol in nonoxidative hair dyes, and that the data needs that were included in the Tentative Report (issued at June 14-15, 1999 Panel meeting) are applicable to these two ingredients.

Dr. Andersen noted that the Belsito Team's conclusion differs significantly from the conclusion that was issued in the Tentative Report (i.e., insufficient data conclusion on all six ingredients). Thus, if the proposed conclusion is approved, the Panel should issue a Revised Tentative Report.

It was moved and seconded that 6-Amino-*m*-Cresol, 4-Amino-*m*-Cresol, 5-Amino-4-Chloro-*o*-Cresol, and 5-Amino-6-Chloro-*o*-Cresol are safe as used in oxidative and non-oxidative hair dyes, that 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol are safe as used in oxidative hair dyes, and that the available data are insufficient for supporting the safety of 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol in non-oxidative hair dyes. The data that are needed in order for the Panel to complete the safety assessment of these two ingredients are listed in the discussion section of the report as follows:

- (1) Physical and chemical properties, including the octanol/water partition coefficient
- (2) Impurities data, especially regarding the presence of *m*-cresol, other organic molecules, and heavy metals
- (3) Metabolism data; if the metabolism is not similar to that of 4-Amino-2-Hydroxytoluene and/or *p*-, *m*-, and *o*-Aminophenol (ingredients already reviewed by CIR), the following data are needed:
  - (a) 28-day dermal toxicity data with histopathology
  - (b) dermal reproductive toxicity data
  - (c) an *in vitro* genotoxicity study for 6-Amino-*o*-Cresol, and a genotoxicity study in a mammalian system for 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol (if any of these data are positive, a two-year dermal carcinogenicity study performed using NTP methods may be needed)

The Panel voted unanimously in favor of issuing a Revised Tentative Report with the conclusions stated in the preceding paragraph.

#### **SEPTEMBER 11-12, 2000 PANEL MEETING**

Dr. Belsito recalled that at the February 14-15, 2000 Panel meeting, the Panel concluded that the available data support the safety of 6-Amino-*m*-Cresol, 4-Amino-*m*-Cresol, 5-Amino-4-Chloro-*o*-Cresol, and 5-Amino-6-Chloro-*o*-Cresol as used in oxidative and non-oxidative semipermanent hair dyes, and that the available data also support the safety of 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol as used in oxidative hair dyes. The Panel also concluded that the available data are insufficient to support the safety of 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol in nonoxidative semipermanent hair dyes. The issuance of a Revised Tentative Report with these conclusions was unanimously approved. Dr. Belsito noted that no data submissions in response to the insufficient data conclusion have been received.

The Panel voted unanimously in favor of issuing a Final Report on this group of ingredients with the following conclusion: The CIR Expert Panel concludes that the available data support the safety of 6-Amino-*m*-Cresol, 4-Amino-*m*-Cresol, 5-Amino-4-Chloro-*o*-Cresol, and 5-Amino-6-Chloro-*o*-Cresol as used in oxidative and non-oxidative (semi-permanent) hair dyes. The available data also support the safety of 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol for use in oxidative hair dyes, but are insufficient to support the safety of 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol in non-oxidative (semi-permanent) hair dyes. The data that are needed in order for the Panel to complete its safety assessment of 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol are listed in the discussion section of the report as follows:

- (1) Physical and chemical properties for all ingredients, including the octanol/water partition coefficient
- (2) Impurities data, especially regarding the presence of *m*-cresol, other organic molecules, and heavy metals for all ingredients except 4-Amino-*m*-Cresol
- (3) Metabolism data; if metabolism is not similar to that of 4-amino-2-hydroxytoluene and/or *p*-, *m*-, and *o*-aminophenol (ingredients already reviewed by CIR), the following data may be needed:
  - a. 28-day dermal toxicity with histopathology
  - b. dermal reproductive toxicity data
  - c. an *in vitro* genotoxicity study for 6-Amino-*o*-Cresol and one genotoxicity study in a mammalian system for 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol; if positive, a 2-year dermal carcinogenicity study using National Toxicology Program methods may be needed.

Dr. Belsito recommended that the last paragraph in the report discussion, which includes the data needs stated above, be reworded to clarify that the data needs listed refer to the data that are needed in order for the Panel to assess the safety of 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol for use in non-oxidative hair dyes.

# Amended Safety Assessment of 5-Amino-6-Chloro-*o*-Cresol as Used in Cosmetics

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Status: Draft Amended Report for Panel Review  
Release Date: February 10, 2023  
Panel Meeting Date: March 6-7, 2023

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.; 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. The Cosmetic Ingredient Review (CIR) Executive Director is Bart Heldreth, Ph.D. This safety assessment was prepared by Christina Burnett, Senior Analyst/Writer, CIR.

**ABBREVIATIONS**

CIR	Cosmetic Ingredient Review
CO <sub>2</sub>	carbon dioxide
Council	Personal Care Products Council
CPSC	Consumer Product Safety Commission
<i>Dictionary; wINCI</i>	web-based <i>International Cosmetic Ingredient Dictionary and Handbook</i>
DMSO	dimethyl sulfoxide
EC <sub>3</sub>	estimated concentrations for a SI of 3
FDA	Food and Drug Administration
HPLC	high-performance liquid chromatography
LLNA	local lymph node assay
LOAEL	lowest-observable-adverse-effect-level
NMR	nuclear magnetic resonance
NOAEL	no-observable-adverse-effect level
OECD	Organisation for Economic Co-operation and Development
Panel	Expert Panel for Cosmetic Ingredient Safety
SCCS	Scientific Committee on Consumer Safety
SED	systemic exposure dose
SI	stimulation indices
TG	test guideline
US	United States
VCRP	Voluntary Cosmetic Registration Program

## INTRODUCTION

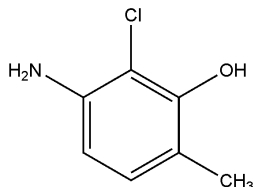
5-Amino-6-Chloro-*o*-Cresol, which according to the web-based *International Cosmetic Ingredient Dictionary and Handbook* (wINCI; *Dictionary*) is reported to function in cosmetics as a hair colorant,<sup>1</sup> was previously reviewed by the Expert Panel for Cosmetic Ingredient Safety (Panel) as part of a safety assessment of 6 amino-cresol hair dye ingredients that was published in 2004.<sup>2</sup> At that time, the Panel concluded that “the available data ... support the safety of 5-Amino-6-Chloro-*o*-Cresol... for use in oxidative and nonoxidative (semi-permanent) hair dyes.” In accordance with its Procedures, the Panel evaluates the conclusions of previously-issued reports approximately every 15 years, and it has been at least 15 years since this assessment has been issued. In June 2022, the Panel determined that this safety assessment should be re-opened for re-evaluation due to several of the other amino-cresol hair dye ingredients that were included in the original 2004 report being banned for use in cosmetics by the European Commission.<sup>3</sup> However, because the Panel determined that data for these amino-cresol hair dye ingredients could not be read-across, rather than including all 6 ingredients in one amended report, re-reviews of each hair dye will now be presented as individual stand-alone reports.

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 Cosmetic Ingredient Review (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. Excerpts from the summaries of the previous report on 5-Amino-6-Chloro-*o*-Cresol are disseminated throughout the text of this re-review document, as appropriate, and are identified by *italicized text*. (These data are not included in the tables or the Summary.)

## CHEMISTRY

### Definition and Structure

According to the *Dictionary*, 5-Amino-6-Chloro-*o*-Cresol (CAS No. 84540-50-1) is the organic compound that conforms to formula in Figure 1.<sup>1</sup> However, the use of regiochemical terms such as “ortho-” (i.e., the “-*o*-” in 5-Amino-6-Chloro-*o*-Cresol) is vague and inappropriate when an aromatic system such as a benzene ring has more than 2 substituents.



**Figure 1.** 5-Amino-6-Chloro-*o*-Cresol

5-Amino-6-Chloro-*o*-Cresol is a precursor in oxidative hair dye systems.<sup>4</sup> It reacts with primary intermediates to form the final hair-reactive dye. The reaction can be accelerated by addition of an oxidizing agent (e.g., hydrogen peroxide), but can also be achieved by air oxidation.

### Chemical Properties

Chemical properties for 5-Amino-6-Chloro-*o*-Cresol are summarized in Table 1. 5-Amino-6-Chloro-*o*-Cresol is soluble in water and has a symmetrical absorption peak below 300 nm, which falls off sharply above 300 nm.<sup>2</sup>

### Method of Manufacture

Method of manufacturing data for 5-Amino-6-Chloro-*o*-Cresol were not included in the original report and were not found in the updated literature search, and unpublished data were not submitted.

### Composition and Impurities

*A high-performance liquid chromatography (HPLC) analysis of 5-Amino-6-Chloro-*o*-Cresol yielded 94.19% of the ingredient in one peak.<sup>2</sup> Near the major peak were small peaks for 5-amino-4-chloro-2-methylphenol (2.76%) and *p*-amino-*o*-cresol (1.99%). The only other significant peak (0.83%) was identified as a dichloro derivative.*

The purity of 5-Amino-6-Chloro-*o*-Cresol is reported to be > 98% (w/w; free base) through nuclear magnetic resonance (NMR) spectroscopy and > 98% (area) through HPLC.<sup>4</sup> Solvent content as water is < 0.5% (w/w). 4-Amino-2-hydroxy-toluene (up to 6%) and 5-amino-4-chloro-*o*-cresol (up to 2.5%) are reported impurities. 5-Amino-4-chloro-*o*-cresol hydrochloride is a possible impurity.



## USE

### **Cosmetic**

The safety of the cosmetic ingredient 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 this ingredient 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, 5-Amino-6-Chloro-*o*-Cresol has 27 reported uses in hair coloring products (Table 2).<sup>5</sup> The results of the concentration of use survey conducted by the Council in 2021 (provided in 2022) reported that the maximum concentration of use of 5-Amino-6-Chloro-*o*-Cresol is 0.24% in hair dyes and colors.<sup>6</sup> When the original safety assessment was published in 2004, 5-Amino-6-Chloro-*o*-Cresol was reported to have no uses (according to VCRP data acquired from the FDA in 1998).<sup>2</sup> However, according to industry survey data submitted in 1996, 5-Amino-6-Chloro-*o*-Cresol was reported to be used at up to 2.0% in hair dyes and colors.

This ingredient is considered a coal tar hair dye for which regulations require caution statements and instructions regarding patch tests in order to be exempt from certain adulteration and color additive provisions of the US Federal Food, Drug, and Cosmetic Act. In order to be exempt, the following caution statement must be displayed on all coal tar hair dye products:

Caution - this product contains ingredients which may cause skin irritation on certain individuals and a preliminary test according to accompanying directions should be made. This product must not be used for dyeing the eyelashes or eyebrows; to do so may cause blindness.

Product labels shall also bear patch test instructions for determining whether the product causes skin irritation. However, whether or not patch testing prior to use is appropriate is not universally agreed upon. The Panel recommends that an open patch test be applied and evaluated by the beautician and/or consumer for sensitization 48 h after application of the test material and prior to the use of a hair dye formulation. Conversely, a report in Europe suggests that self-testing has severe limitations, and may even cause morbidity in consumers.<sup>7,8</sup> Hair dye products marketed and sold in the US, though, must follow the labeling requirements established by the Food, Drug, and Cosmetic Act.

In the European Union, 5-Amino-6-Chloro-*o*-Cresol is categorized in Annex III, the list of substances which cosmetic products must not contain except subject to the restrictions laid down.<sup>3</sup> It is limited to 0.5% in non-oxidative hair dyes; no concentration limit was described for oxidative hair dyes. The Scientific Committee on Consumer Safety (SCCS) concluded that "because of the low margin of safety for the use in both oxidative and non-oxidative hair dye formulations, ... use of 5-Amino-6-Chloro-*o*-Cresol as a hair dye ingredient up to a final on-head concentration of 2.0% under oxidative and non-oxidative conditions poses a risk to the health of the consumer."<sup>4</sup>

## TOXICOKINETIC STUDIES

### **Dermal Absorption/Penetration**

#### In Vitro

The dermal absorption/percutaneous penetration potential of [<sup>14</sup>C]labelled 5-Amino-6-Chloro-*o*-Cresol (> 99% pure) through dermatomed pig skin was determined for a cream formulation containing 4% of the test material.<sup>4</sup> The material was mixed with a developer and was tested with and without hydrogen peroxide. The study was performed in accordance with Organisation for Economic Co-operation and Development (OECD) test guideline (TG) 428. The concentration of 5-Amino-6-Chloro-*o*-Cresol in the final application formulation was 2.1%. Using Franz diffusion cells, 20 mg/cm<sup>2</sup> (nominal dose of test substance = 0.42 mg/cm<sup>2</sup>; skin discs = 1.0 cm<sup>2</sup>) was applied for 30 min. Application of the test material was terminated by gently rinsing with 0.01% Tween 80 solution and water. The formulation was analyzed in 2 experiments with 8 replicates per experiment for adsorbed, absorbed, and penetrated amount of the test material. The receptor fluid (Dulbecco' phosphate buffered saline) was analyzed at defined intervals for up to 48 h post application.

The amount of 5-Amino-6-Chloro-*o*-Cresol systemically available from a standard cream formulation mixed with a developer with hydrogen peroxide was found to be 30.21 ± 9.78 µg/cm<sup>2</sup> (range 18.82 - 49.21 µg/cm<sup>2</sup>) or 5.98 ± 1.45% (range 4.24 - 8.22%) of the applied dose. For the formulation tested without hydrogen peroxide, the systemically available amount was 53.8 ± 6.38 µg/cm<sup>2</sup> (range 45.47 - 62.1 µg/cm<sup>2</sup>) or 11.78 ± 1.26% (range 9.91 - 13.73%) of the applied dose. The SCCP noted the mean values + 2 standard deviations were used for calculating the margin of safety.<sup>4</sup>

**Animal**

*[<sup>14</sup>C]5-Amino-6-Chloro-*o*-Cresol hydrochloride (1.14%) in formulation was readily absorbed (93.2%) in a skin penetration study in female rats.<sup>2</sup> Radioactivity was excreted in urine (87.7%) and feces (2.22%). Only 0.48% was found in the carcass. The recovery rate of <sup>14</sup>C from the urine samples was 115% of the applied dose. No detectable radiolabel was found in the expired carbon dioxide (CO<sub>2</sub>). In a similar study, the test formulation was mixed 1:1 with 3% hydrogen peroxide developer solution prior to application. The skin penetration was only 0.116%.*

**Absorption, Distribution, Metabolism, Elimination (ADME) Studies****In Vivo**

*A metabolism study of [<sup>14</sup>C]5-Amino-6-Chloro-*o*-Cresol (0.25% in water) in female rats via a single subcutaneous injection found that excretion was mainly via urine (88.5%), of which most (88.1%) was eliminated in the first 24 h.<sup>2</sup> Only 3.97% was excreted in feces, 0.674% was found in the carcass, and 0.04% remained at the injection site skin. No detectable radioactivity was found in expired CO<sub>2</sub>. In an oral metabolism study in male rats, [<sup>14</sup>C] 5-Amino-6-Chloro-*o*-Cresol (1.7% in water) was mainly excreted via urine (90.93%), and mostly (90%) in the first 24 h. There was 6% in the gastrointestinal tract and 0.58% in the remaining carcass. No radiolabel was detected in expired CO<sub>2</sub>. In a distribution study, male rats received a single oral dose of [<sup>14</sup>C]5-Amino-6-Chloro-*o*-Cresol (1.7% in water). After 6 h, radioactivity was in the stomach, intestine, or colon content, and in the cecum. After 24 and 48 h, only residual radioactivity was found in the colon, cecum, and kidneys. After 96 h, excretion was nearly complete and only a small amount of label appeared (in bone). Within the first 24 h, 91% of the radioactivity was excreted via urine.*

**TOXICOLOGICAL STUDIES****Acute Toxicity Studies**

*The LD<sub>50</sub> was 1360 mg/kg in a gavage study of male rats that received 5-Amino-6-Chloro-*o*-Cresol hydrochloride at doses of 501, 1000, 1250, 1580, and 1990 mg/kg.<sup>2</sup> Observations included apathy, staggering, rapid breathing, dyspnea (at later stages), and yellow-orange discoloration of the urine.*

**Oral**

In an acute oral study, 10 male CF1 mice received 501, 631, 794, 1000, 1250, 1580, or 1900 mg/kg bw 5-Amino-6-Chloro-*o*-Cresol dissolved in distilled water by gavage.<sup>9</sup> Clinical symptoms observed in all dose groups included apathy, enhanced breathing, abdominal position, cramps, and uttering of tone. The LD<sub>50</sub> of the test material was determined to be 1200 mg/kg bw.

**Subchronic Toxicity Studies****Oral**

*In a 13-wk study, male and female rats received daily dose of 50 mg/kg 5-Amino-6-Chloro-*o*-Cresol hydrochloride with tragacanth (1%) by gavage.<sup>2</sup> No clinical observations, biochemical alterations, or pathological findings were indicative of systemic toxicity. The no-observable-adverse-effect-level (NOAEL) was 50 mg/kg.*

In a 90-d gavage study, groups of 12 male and 12 female Wistar Crl: (WI) BR rats received 0, 100, 300, or 600 mg/kg bw 5-Amino-6-Chloro-*o*-Cresol (> 99.6% pure) in propylene glycol.<sup>4</sup> The study was performed in accordance with OECD TG 408. A recovery group of 5 animals of each sex was also utilized and observed up to 28 d after treatment. Clinical signs of toxicity were monitored daily and feed consumption and body weight gains were measured weekly. Ophthalmologic examinations, hearing tests, and motor tests were also conducted. Blood samples were taken to study hematology, clotting potential, and clinical biochemistry. At necropsy, organ weights were recorded and histopathology of organs was performed.

Three deaths occurred during the study, one in the low-dose group, one in the high-dose group, and one in the recovery control group. All dose groups had a brown discoloration of the urine, and animals in the medium- and high-dose groups were observed with salivation from week 3 of treatment onwards. Histopathology revealed centrilobular hypertrophy of the liver in all groups in a dose-related manner that occurred primarily in males. This finding persisted in all high-dose males through the end of the recovery phase. Other microscopic findings were only related to the high-dose males, including cortical tubular basophilia in the kidneys and limiting ridge hyperplasia of the forestomach with squamous hyperplasia of the main stomach. In the high-dose group (gender not specified), increased relative and absolute liver weights were also observed, along with increased bilirubin, potassium levels, reduced urea levels, increased cholesterol levels, and increased alanine aminotransferase activity. Statistically significant deviations in hematological parameters in the high-dose group included reduced blood cell counts in males, reduced platelet counts in males, increased mean corpuscular volume in males and females, increased corpuscular hemoglobin level in females, and reduced corpuscular hemoglobin concentration in males and females. In the low-dose group, increased hematocrit values and reduced corpuscular hemoglobin concentration were observed in males, and increased mean corpuscular volume was observed in both males and females. Females in all dose groups were observed to have reduced corpuscular hemoglobin concentrations. Some of these effects persisted through the recovery period. The NOAEL was established at 100 mg/kg/d; however, the SCCS noted that this value should be considered the lowest-observable-adverse-effect-level (LOAEL).<sup>4</sup>

## **DEVELOPMENTAL AND REPRODUCTIVE TOXICITY STUDIES**

*No developmental toxicity was associated with treatment with 5-Amino-6-Chloro-o-Cresol hydrochloride in water in rats exposed on days 6 to 15 of pregnancy.<sup>2</sup> The rats received 0, 30, 90, or 270 mg/kg/d of the test material via oral gavage. The dams were killed on day 21 of gestation and the fetuses removed for examination. The only maternal effects were slight reduction in feed consumption and reduced body weight gain in the highest dose group. The NOAEL was considered to be 90 mg/kg/day.*

Development and reproductive toxicity studies were not found in the updated literature search, and unpublished data were not submitted.

## **GENOTOXICITY STUDIES**

*In an Ames test, 5-Amino-6-Chloro-o-Cresol hydrochloride was evaluated for mutagenicity using concentrations of 4 to 2500 µg/plate, with and without metabolic activation.<sup>2</sup> 5-Amino-6-Chloro-o-Cresol hydrochloride was mutagenic with metabolic activation in two Salmonella typhimurium strains. No increases in the number of mutations were observed in cell mutation test at the HGRPT locus in Chinese hamster lung fibroblast V79 cells exposed to 5-Amino-6-Chloro-o-Cresol hydrochloride dissolved in ethanol at up to 300 µg/ml, with or without metabolic activation. 5-Amino-6-Chloro-o-Cresol hydrochloride at concentrations at up to 1100 µg/ml did not induce chromosome aberration in Chinese hamster lung fibroblast V79 cells, with or without metabolic activation. No indications of a dose-related increase in unscheduled DNA synthesis were observed in rat liver hepatocytes exposed to 5-Amino-6-Chloro-o-Cresol hydrochloride at up to 2000 µg/ml.*

*In an vivo micronucleus test for chromosome mutations, 1200 mg/kg of 5-Amino-6-Chloro-o-Cresol hydrochloride dissolved in water was administered to male and female mice via gavage.<sup>2</sup> Bone marrow was extracted for analysis of 1000 polychromic erythrocytes per animal. The ratio of chromatic/polychromatic erythrocytes was slightly increased, suggesting some toxicity to the bone marrow, but the investigators concluded that 5-Amino-6-Chloro-o-Cresol hydrochloride was not mutagenic in this assay.*

In vitro and in vivo genotoxicity studies on 5-Amino-6-Chloro-o-Cresol summarized here are detailed in Table 3. 5-Amino-6-Chloro-o-Cresol (99.6% pure) was not mutagenic in an Ames test at up to 5000 µg/plate, with and without S9 metabolic activation.<sup>4</sup> 5-Amino-6-Chloro-o-Cresol (99.6% pure) was not genotoxic in an L5178 mouse lymphoma cells assay at the *tk* locus at up to 1000 µg/ml without metabolic activation or at up to 18.9 µg/ml with metabolic activation. In another gene mutation test, 5-Amino-6-Chloro-o-Cresol (80 - 90% pure) was not genotoxic in Chinese hamster lung fibroblast V79 cells at the HPRT locus, with or without metabolic activation.<sup>9</sup> In vivo testing found that 5-Amino-6-Chloro-o-Cresol (99.9% pure) did not induce micronuclei in mice that received a single intraperitoneal dose of up to 400 mg/kg bw.<sup>4</sup>

## **CARCINOGENICITY STUDIES**

Carcinogenicity studies were not included in the original report and were not found in the updated literature search, and unpublished data were not submitted.

## **DERMAL IRRITATION AND SENSITIZATION STUDIES**

### **Irritation**

#### **Animal**

*No signs of primary skin irritation were observed in male hairless mice exposed to 5-Amino-6-Chloro-o-Cresol at 10% in an aqueous formulation following twice daily applications for 5 consecutive days.<sup>2</sup> This formulation was also not a dermal irritant in male rabbits following a 2 h application with an occlusive patch or in an open application on the same test site every 30 s for a total of 60 applications.*

The irritation potential of undiluted 5-Amino-6-Chloro-o-Cresol (99.6% pure) was assessed in 3 male New Zealand albino rabbits in accordance with OECD TG 404.<sup>4</sup> The test material was applied as a single dose (0.5 g) moistened with 0.7 ml purified water to shaved, intact skin. Semi-occlusive patches (2 cm x 3 cm) were applied and left in place for 4 h. The skin was examined for erythema, eschar formation, and edema at 1, 24, 48, and 72 h after the patches were removed. No reactions were observed. It was concluded that 5-Amino-6-Chloro-o-Cresol was not irritating or corrosive.

### **Sensitization**

#### **Animal**

*In a guinea pig maximization study, induction was done with an injection of 5.0% aqueous solution of 5-Amino-6-Chloro-o-Cresol and two injections of 5.0% aqueous 5-Amino-6-Chloro-o-Cresol diluted 1:1 with Freund's complete adjuvant.<sup>2</sup> A topical induction was done 1 wk later with a 5% cream of 5-Amino-6-Chloro-o-Cresol in petroleum jelly under an occlusive patch for 48 h. The challenge was done with a 25% cream of the test substance applied under an occlusive patch. One quarter of the test animals had slight erythema 24 h after challenge, but no effects were evident after 48 h. It was concluded that 5-Amino-6-Chloro-o-Cresol is not a sensitizer.*

A local lymph node assay (LLNA) was performed using 5-Amino-6-Chloro-*o*-Cresol (99.6% pure) in accordance with OECD TG 429.<sup>4</sup> Female CBA mice were divided into groups of 6 and received 0, 5, 25, or 50% (2 groups) of the test material in ethanol:water (7:3, v/v) on the ear surface (25 µl) once daily for 3 consecutive days.  $\alpha$ -Hexylcinnamaldehyde was used as the positive control. Five days after the first topical application, all animals were injected intravenously with [<sup>3</sup>H]methyl thymidine and the proliferation of lymphocytes in the draining lymph nodes was measured. The stimulation indices (SI) were calculated to be 1.0, 0.9, 4.0, and 1.0 for the 5%, 25%, the initial 50%, and the additional 50% dose groups, respectively. The estimated concentrations for an SI of 3 (EC<sub>3</sub>) was not calculated as there was no clear evidence the test material could elicit an SI  $\geq$  3. The controls yielded expected results. It was concluded that 5-Amino-6-Chloro-*o*-Cresol was not sensitizing when tested at up to 50% in mice.

## **OCULAR IRRITATION STUDIES**

### **Animal**

In an ocular irritation study in accordance with OECD TG 405, 3 male New Zealand rabbits received approximately 0.1 ml (38.3 mg) undiluted 5-Amino-6-Chloro-*o*-Cresol (99.6% pure) in the conjunctival sac of one eye.<sup>4</sup> The eye was rinsed with warm tap water after 24 h. The untreated eye served as the control. Ocular reactions were recorded at 1, 24, 48, and 72 h and 7 d after instillation. Slight redness of the conjunctiva was reported in 3 rabbits within 6 h of instillation, which resolved in all animals by 24 h. Exudate was observed in all rabbits 1 h after instillation, in 3 rabbits after 6 h, and in 1 rabbit until 24 h. Opacity (maximum grade 1) and epithelial damage (maximum 55% of the corneal area) of the cornea were observed. The corneal injury resolved with 24 h in one animal and within 7 d in the other animals. Iridial irritation grade 1 was observed in one animal only 24 h after instillation. The irritation of the conjunctivae consisted of redness and chemosis, which resolved within 7 d in two animals and within 14 d in the third. It was concluded that 5-Amino-6-Chloro-*o*-Cresol was an ocular irritant under the conditions of the experiment.

## **MARGIN OF SAFETY**

The SCCS calculated the margin of safety for 5-Amino-6-Chloro-*o*-Cresol under oxidative conditions to be 68.<sup>4</sup> This calculation is based on the adjusted LOAEL of 33 mg/kg bw/d (from a 90-d oral rat study with an adjustment factor of 3) and a systemic exposure dose (SED) of 0.482 mg/kg bw (skin area surface of 580 cm<sup>2</sup> x absorption through skin of 49.77 µg/cm<sup>2</sup> x 0.001 (unit conversion)/typical human bw of 60 kg). The margin of safety for 5-Amino-6-Chloro-*o*-Cresol under non-oxidative conditions was calculated to be 51. In this calculation, the SED value of 0.643 mg/kg bw was used (the absorption through skin was 66.6 µg/cm<sup>2</sup>).

## **HAIR DYE EPIDEMIOLOGY**

Hair dyes may be broadly grouped into oxidative (permanent) and direct (temporary or semi-permanent) dyes. The oxidative dyes consist of precursors mixed with developers to produce color, while direct hair dyes consist of preformed colors. 5-Amino-6-Chloro-*o*-Cresol is reported to be used in direct and oxidative hair dye formulations. While the safety of individual hair dye ingredients is not addressed in epidemiology studies that seek to determine links, if any, between hair dye use and disease, such studies do provide broad information. The Panel determined that the available hair dye epidemiology data do not provide sufficient evidence for a causal relationship between personal hair dye use and cancer. A detailed summary of the available hair dye epidemiology data is available at <https://www.cir-safety.org/cir-findings>.

## **SUMMARY**

5-Amino-6-Chloro-*o*-Cresol is reported to function in cosmetics as a hair colorant. 5-Amino-6-Chloro-*o*-Cresol was previously reviewed by the Panel in a safety assessment of 6 amino-cresol hair dye ingredients that was published in 2004. At that time, the Panel concluded that 5-Amino-6-Chloro-*o*-Cresol was safe for use in oxidative and non-oxidative hair dyes. In accordance with its Procedures, the Panel evaluates the conclusions of previously-issued reports approximately every 15 years, and it has been at least 15 years since this assessment has been issued. In June 2022, the Panel determined that this safety assessment should be re-opened for re-evaluation due to several of the other amino-cresol hair dye ingredients that were included in the original 2004 report being banned for use in cosmetics by the European Commission.

According to 2022 VCRP survey data, 5-Amino-6-Chloro-*o*-Cresol has 27 reported uses in hair coloring products. The results of the concentration of use survey conducted by the Council in 2021 reported that the maximum concentration of use of 5-Amino-6-Chloro-*o*-Cresol is 0.24% in hair dyes and colors. When the original safety assessment was published in 2004, 5-Amino-6-Chloro-*o*-Cresol was reported to have no uses, according to 1998 VCRP data. However, according to industry survey data submitted in 1996, 5-Amino-6-Chloro-*o*-Cresol was reported to be used at up to 2% in hair dyes and colors.

Under European regulations for cosmetic ingredients, 5-Amino-6-Chloro-*o*-Cresol is categorized in Annex III, the list of substances which cosmetic products must not contain except subject to the restrictions laid down. It is limited to 0.5% in non-oxidative hair dyes; no concentration limit was described for oxidative hair dyes. The SCCS concluded that “because of the low margin of safety for the use in both oxidative and non-oxidative hair dye formulations, ... use of 5-Amino-6-Chloro-

*o*-Cresol as a hair dye ingredient up to a final on-head concentration of 2.0% under oxidative and non-oxidative conditions poses a risk to the health of the consumer.”

In a dermal absorption/percutaneous penetration study, a cream formulation containing [<sup>14</sup>C]5-Amino-6-Chloro-*o*-Cresol (> 99% pure) with a developer was applied to excised pig skin, with and without hydrogen peroxide, with the final concentration of the radiolabel reported as 2.1%. The amount of 5-Amino-6-Chloro-*o*-Cresol systemically available from a standard cream formulation mixed with a developer with hydrogen peroxide was found to be  $30.21 \pm 9.78 \mu\text{g}/\text{cm}^2$  (range 18.82 - 49.21  $\mu\text{g}/\text{cm}^2$ ) or  $5.98 \pm 1.45\%$  (range 4.24 - 8.22%) of the applied dose. For the formulation tested without hydrogen peroxide, the systemically available amount was  $53.8 \pm 6.38 \mu\text{g}/\text{cm}^2$  (range 45.47 - 62.1  $\mu\text{g}/\text{cm}^2$ ) or  $11.78 \pm 1.26\%$  (range 9.91 - 13.73%) of the applied dose.

In an acute oral study in male mice, the LD<sub>50</sub> for 5-Amino-6-Chloro-*o*-Cresol was 1200 mg/kg bw. The NOAEL in a 90-d oral study in rats was 100 mg/kg/d. Adverse effects included centrilobular hypertrophy of the liver observed at all dose levels in a dose-dependent manner that persisted through the recovery period. High-dose male group also had cortical tubular basophilia in the kidneys, limiting ridge hyperplasia of the forestomach with squamous hyperplasia of the main stomach, and increased relative and absolute liver weights. A reduced mean corpuscular hemoglobin concentration was observed in all dose groups of female rats.

5-Amino-6-Chloro-*o*-Cresol (99.6% pure) was not mutagenic in an Ames test at up to 5000  $\mu\text{g}/\text{plate}$ , with and without S9 metabolic activation. 5-Amino-6-Chloro-*o*-Cresol (99.6% pure) was not genotoxic in a L5178 mouse lymphoma cell assay at the *tk* locus at up to 1000  $\mu\text{g}/\text{ml}$  without metabolic activation or at up to 18.9  $\mu\text{g}/\text{ml}$  with metabolic activation. In another gene mutation test, 5-Amino-6-Chloro-*o*-Cresol (80-90% pure) was not genotoxic in Chinese hamster lung fibroblast V79 cells at the HPRT locus, with or without metabolic activation. In vivo testing found that 5-Amino-6-Chloro-*o*-Cresol (99.9% pure) did not induce micronuclei in mice that received a single intraperitoneal dose of up to 400 mg/kg bw.

Undiluted 5-Amino-6-Chloro-*o*-Cresol was not irritating or corrosive in a dermal study in rabbits. This ingredient was also not sensitizing in mice in an LLNA when tested at up to 50%. Undiluted 5-Amino-6-Chloro-*o*-Cresol was an ocular irritant in rabbit eyes.

A margin of safety for 5-Amino-6-Chloro-*o*-Cresol under oxidative conditions was calculated to be 68. This calculation is based on the adjusted LOAEL of 33 mg/kg bw/d (from a 90-d oral rat study with an adjustment factor of 3) and an SED of 0.482 mg/kg bw. The margin of safety for 5-Amino-6-Chloro-*o*-Cresol under non-oxidative conditions was calculated to be 51. In this calculation, the SED value of 0.643 mg/kg bw was used.

The Panel determined that the available hair dye epidemiology data do not provide sufficient evidence for a causal relationship between personal hair dye use and cancer.

Method of manufacturing data and carcinogenicity studies on 5-Amino-6-Chloro-*o*-Cresol were not included in the original report and were not found in the updated literature search, and unpublished data were not submitted.

## **PREVIOUS DISCUSSION**

*The Panel recognizes that irritation and sensitization data on [several cresol hair dyes] are absent from this report.<sup>2</sup> However, the hair dyes containing the ingredients included in this report, as coal tar hair dye products, are exempt from the principal adulteration provision and from the color additive provisions in sections 601 and 706 of the Federal Food, Drug, and Cosmetic Act of 1938 when the label bears a caution statement and patch test instructions for determining whether the product causes skin irritation. The Expert Panel expects that following this procedure will identify individuals who would have an irritation/sensitization reaction and allow them to avoid significant exposures.*

*The information available on the use of 5-Amino-6-Chloro-*o*-Cresol ... in hair dye formulations indicate that [this ingredient is] reacted with a developer and is not available for absorption into the skin of the scalp. [This compound], when tested alone, [is a] moderate skin sensitizer, but when combined with the developer, [this ingredient is] not a sensitizer in animal tests. In addition, no toxicologically significant impurities are present with [this ingredient].*

*This information, coupled with the available animal test data, support the safety of 5-Amino-6-Chloro-*o*-Cresol ... for use in oxidative hair dyes. Were 5-Amino-6-Chloro-*o*-Cresol... to have application in non-oxidative (semi-permanent) hair dyes, there is concern about the potential for skin sensitization because [this ingredient is a] moderate sensitizer. Because individuals would be pretested to determine if they would develop skin sensitization and because there is an absence of any significant systemic toxic effects in animal tests, the Panel believes that [this ingredient] could be used safely in semi-permanent hair dyes. Even though there is currently no use of [this ingredient] as semi-permanent hair dye, the Panel believes it useful to conclude that [it] could be used safely...<sup>2</sup>*

**DISCUSSION**

To be determined.

**CONCLUSION**

To be determined.

**TABLES****Table 1. Chemical properties**

Property	Value	Reference
<b>5-Amino-6-Chloro-o-Cresol</b>		
Physical Form	Beige crystals	2
Formula Weight (g/mol)	194.07 (hydrochloride)	2
Molecular Weight (g/mol)	157.6 (free base)	4
Melting Point (°C)	144-183	2
	82-86	4
Water Solubility (g/l @ 20 °C)	< 10	4
Other Solubility (g/l @ 20 °C)	ethanol: < 100 dimethyl sulfoxide (DMSO): > 100	4
log P <sub>ow</sub>	1.644 (experimental) 1.44 (estimated)	4

**Table 2. Updated and historical frequency (2022; 1998) and concentration (2021; 1996) of use according to duration and exposure and by product category**

	# of Uses		Max Conc of Use (%)	
	<b>5-Amino-6-Chloro-o-Cresol</b>			
	2022 <sup>5</sup>	1998 <sup>2</sup>	2021 <sup>6</sup>	1996 <sup>2</sup>
<b>Totals</b>	<b>27</b>	<b>NR</b>	<b>0.007 – 0.24</b>	<b>2</b>
<b>summarized by likely duration and exposure*</b>				
<b>Duration of Use</b>				
Leave-On	NR	NR	NR	NR
Rinse-Off	27	NR	0.007 – 0.24	2
Diluted for (Bath) Use	NR	NR	NR	NR
<b>Exposure Type</b>				
Eye Area	NR	NR	NR	NR
Incidental Ingestion	NR	NR	NR	NR
Incidental Inhalation-Spray	NR	NR	NR	NR
Incidental Inhalation-Powder	NR	NR	NR	NR
Dermal Contact	NR	NR	NR	NR
Deodorant (underarm)	NR	NR	NR	NR
Hair - Non-Coloring	NR	NR	NR	NR
Hair-Coloring	27	NR	0.007 – 0.24	2
Nail	NR	NR	NR	NR
Mucous Membrane	NR	NR	NR	NR
Baby Products	NR	NR	NR	NR
<b>as reported by product category</b>				
<b>Hair Coloring Preparations</b>				
Hair Dyes and Colors (all types requiring caution statements and patch tests)	26	NR	0.24	2
Hair Bleaches	1	NR	0.007	NR

NR – not reported

\*likely duration and exposure is derived based on product category (see Use Categorization <https://www.cir-safety.org/cir-findings>)

**Table 3. Genotoxicity studies**

Ingredient	Concentration/Dose	Vehicle	Test System	Procedure	Results	Reference
<b>IN VITRO</b>						
5-Amino-6-Chloro- <i>o</i> -Cresol; 99.6% pure	3 - 5000 µg/plate	DMSO	<i>Salmonella typhimurium</i> strains TA98, TA100, TA102, TA1535, TA1537	Bacterial reverse mutation test in accordance with OECD TG 471; with and without S9 metabolic activation	Not mutagenic; no biologically relevant increase in revertant colonies observed in any strain, with or without metabolic activation	4
5-Amino-6-Chloro- <i>o</i> -Cresol; 99.6% pure	Test 1: up to 1000 µg/ml without metabolic activation and up to 18.9 µg/ml with metabolic activation Test 2: up to 400 µg/ml without metabolic activation and up to 18 µg/ml with metabolic activation	DMSO	L5178 mouse lymphoma cells	Mammalian cell gene mutation test at the <i>tk</i> locus in accordance with OECD TG 476; with and without metabolic activation	Not genotoxic; no reproducible increase in mutant frequency observed in both tests without metabolic activation; a dose-related increase in mutant frequency observed only in test 1 with metabolic activation, but increases were not reproducible in test 2 and the effect was considered not biologically relevant	4
5-Amino-6-Chloro- <i>o</i> -Cresol; 80 - 90% pure	Up to 350 µg/ml without metabolic activation and up to 3000 µg/ml with metabolic activation	DMSO	Chinese hamster fibroblast V79 cells	Mammalian cell gene mutation test at the HPRT locus in accordance with OECD TG 476; with and without metabolic activation	Not genotoxic; no biologically relevant significant increase in mutant colony numbers, with or without metabolic activation; negative and positive controls according to guidelines	9
<b>IN VIVO</b>						
5-Amino-6-Chloro- <i>o</i> -Cresol; 99.9% pure	100, 200, and 400 mg/kg bw	DMSO (30%)	5 NMRI mice per sex	Mammalian erythrocyte micronucleus test in accordance with OECD TG 474; single intrapertitoneal dose; groups of animals killed at 24 or 48 h post-treatment; appropriate negative and positive controls used	Not genotoxic; test material did not induce micronuclei. No mortalities observed; clinical signs included reduced spontaneous activity, ruffled fur, eyelid closure, and abdominal position	4



## **REFERENCES**

1. Nikitakis J, Kowcz A. Web-Based International Cosmetic Ingredient Dictionary and Handbook. <http://webdictionary.personalcarecouncil.org/jsp/Home.jsp>. Washington, DC: Personal Care Products Council. Last Updated 2023. Accessed 01/13/2023.
2. Andersen FA (ed.). Final report on the safety assessment of 6-Amino-m-Cresol, 6-Amino-o-Cresol, 4-Amino-m-Cresol, 5-Amino-4-Chloro-o-Cresol, 5-Amino-6-Chloro-o-Cresol, and 4-Chloro-2-Aminophenol. *Int J Toxicol*. 2004;23(Suppl. 2):1-22.
3. European Commission. Cosing database; following Cosmetic Regulation (EC) No. 1223/2009. <http://ec.europa.eu/growth/tools-databases/cosing/> Last updated 2022. Accessed: 01/11/2023.
4. Scientific Committee on Consumer Safety (SCCS). Opinion on 5-Amino-6-chloro-o-cresol: COLIPA No. A94. 2009. SCCS/1225/09. [http://ec.europa.eu/health/scientific\\_committees/consumer\\_safety/docs/sccs\\_o\\_005.pdf](http://ec.europa.eu/health/scientific_committees/consumer_safety/docs/sccs_o_005.pdf). Accessed 01/13/2023.
5. 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.
6. Personal Care Products Council. 2022. Concentration of Use by FDA Product Category: Amino Cresol Ingredients. Unpublished data submitted by the Personal Care Products Council on January 12, 2022.
7. Thyssen JP, Sosted H, Uter W, et al. Self-testing for contact sensitization to hair dyes - scientific considerations and clinical concerns of an industry-led screening programme. *Contact Dermatitis*. 2012;66(6):300.
8. Goossens A. Self-testing for contact sensitization to hair dyes. *Contact Dermatitis*. 2012;66(6):299.
9. Scientific Committee on Cosmetic and Non-Food Products (SCCNFP). Opinion of the Scientific Committee on Cosmetic Products and Non-Food Products intended for consumers concerning 2-Chloro-6-Methyl-3-Aminophenol Hydrochloride. COLIPA No. A94. 2003. SCCNFP/073/03. [http://ec.europa.eu/health/ph\\_risk/committees/sccp/documents/out233\\_en.pdf](http://ec.europa.eu/health/ph_risk/committees/sccp/documents/out233_en.pdf). Accessed 01/23/2023.

# Final Report on the Safety Assessment of 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, 5-Amino-4-Chloro-*o*-Cresol, 5-Amino-6-Chloro-*o*-Cresol, and 4-Chloro-2-Aminophenol<sup>1</sup>

Each of these ingredients function as hair colorants. 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol are identified as oxidative hair dyes, that is, they are combined with an oxidizing agent before being applied to the hair. 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, and 5-Amino-4-Chloro-*o*-Cresol are used in oxidative hair dyes, but it is not known if they are also used in nonoxidative (semipermanent) hair dyes. No toxicologically significant impurities are present with these two ingredients. To supplement the safety test data on these ingredients, available data on related ingredients (4-amino-2-hydroxytoluene and *p*-, *m*-, and *o*-aminophenol) previously found safe as used by the Cosmetic Ingredient Review (CIR) Expert Panel were summarized. 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol do not absorb significant ultraviolet radiation in the UVB region and none in the UVA region, although 4-Amino-*m*-Cresol had a symmetrical UV absorption peak at 300 nm. Percutaneous penetration of 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol alone was significant, but when combined with oxidative developer, skin absorption was extremely low. Both of these dyes are excreted rapidly via the urine. Repeated exposure of animal skin to 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol failed to produce any cumulative irritation and single exposures up to 10% were not irritating to animal skin. 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol combined with oxidizer were not sensitizers in guinea pig maximization tests. Ocular irritation resulted from exposure of animals to undiluted 5-Amino-4-Chloro-*o*-Cresol, but not to a 5% solution. Only minor irritation was observed with 5% 5-Amino-6-Chloro-*o*-Cresol. Subchronic toxicity testing in animals using 5-Amino-4-Chloro-*o*-Cresol, 5-Amino-6-Chloro-*o*-Cresol, and 4-Amino-*m*-Cresol did not yield any adverse reactions. 6-Amino-*m*-Cresol and 4-Amino-*m*-Cresol were generally not mutagenic in *in vitro* and *in vivo* tests. Exposure to 5-Amino-4-Chloro-*o*-Cresol, 5-Amino-6-Chloro-*o*-Cresol, 6-Amino-*m*-Cresol and 4-Amino-*m*-Cresol from cosmetics were several orders of magnitude below developmental toxicity no-observed-adverse-effect levels (NOAELs). Although irritation data on several ingredients are absent, products containing these ingredients must

include a caution statement and patch test instructions for determining whether the product causes skin irritation. The Expert Panel expects that following this procedure would identify individuals who would have an adverse reaction and allow them to avoid significant exposures. These compounds, when tested alone, are moderate skin sensitizers, but when combined with the developer, these ingredients are not sensitizers in animal tests. This information, coupled with the available animal test data, supports the safety of these ingredients in oxidative hair dyes. In the absence of systemic toxicity data, however, the available data are insufficient to support the safety of 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol in semipermanent hair dyes. The types of data required for these two ingredients for this use include (1) physical and chemical properties, including the octanol/water partition coefficient; (2) impurities data, especially regarding the presence of *m*-cresol, other organic molecules, and heavy metals; (3) data demonstrating that the metabolism is similar to that of 4-amino-2-hydroxytoluene and/or *p*-, *m*-, and *o*-aminophenol, or 28-day dermal toxicity with histopathology, dermal reproductive toxicity data, and an *in vitro* genotoxicity study for 6-Amino-*o*-Cresol and one genotoxicity study in a mammalian system; if positive, a 2-year dermal carcinogenicity study using National Toxicology Program methods may be needed.

## INTRODUCTION

This report reviews the safety of 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, 5-Amino-4-Chloro-*o*-Cresol, 5-Amino-6-Chloro-*o*-Cresol, and 4-Chloro-2-Aminophenol, all of which function as hair colorants (Pepe, Wenninger, and McEwen 2002).

Data from the Cosmetic Ingredient Review (CIR) reports on 4-amino-2-hydroxytoluene and *p*-, *m*-, and *o*-aminophenol, and relevant data on other structurally similar ingredients (including the hepatotoxicity of acetaminophen derivatives), are included in this review. Elder (1989) found 4-amino-2-hydroxytoluene and Elder (1988) found *p*-, *m*-, and *o*-aminophenol safe in the present practices of use and concentrations. For purposes of comparison with the ingredients reviewed in this safety assessment, 4-Amino-2-hydroxytoluene was used in hair dyes and tints at concentrations  $\leq 5\%$  and *p*-, *m*-, and *o*-aminophenol were

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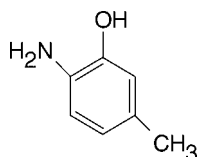
<sup>1</sup>Reviewed by the Cosmetic Ingredient Review Expert Panel. Monice Zondlo Fiume and Torill A. Yamarik prepared this report. Address correspondence to F. Alan Andersen, PhD, Director, Cosmetic Ingredient Review, 1101 17th Street, NW, Suite 310, Washington, DC 20036, USA.

used in hair tints and hair dyes and colors at concentrations of  $\leq 1\%$ ,  $\leq 5\%$ , and  $\leq 1\%$ , respectively.

## CHEMISTRY

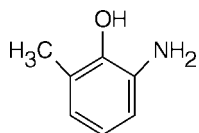
### Definition and Structure

6-Amino-*m*-Cresol (CAS no. 2835-98-5) is the substituted aromatic compound that conforms to the formula (Pepe, Wenninger, and McEwen 2002):



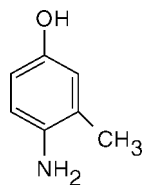
6-Amino-*m*-Cresol is also known as 4-Amino-3-Hydroxytoluene; 2-Amino-5-Methylphenol; Phenol, 2-Amino-5-Methyl-; 2-Hydroxy-4-Methylaniline (Pepe, Wenninger, and McEwen 2002); *m*-Cresol, 6-Amino; 6-Amino-3-Cresol; 6-Amino-3-Methylphenol; 2-Hydroxy-*p*-Toluidine; 5-Methyl-2-Aminophenol (Regulated Chemicals Listing 1998); 6-Amino-*meta*-Cresol; 4-Amino-3-Oxy-1-Methyl-Benzol; 4-Amino-3-Oxy-Toluol (Beilstein File of Organic Compounds 1998); and Toluene, 4-Amino-3-Hydroxy (CRC Handbook of Data on Organic Compounds 1998).

6-Amino-*o*-Cresol (CAS no. 17672-22-9) is the substituted aromatic compound that conforms to the formula (Pepe, Wenninger, and McEwen 2002):



6-Amino-*o*-Cresol is also known as 3-Amino-2-Hydroxytoluene; 2-Amino-6-Methylphenol; Phenol, 2-Amino-6-Methyl-; 6-Amino-2-Methylphenol; Phenol, 6-Amino-2-Methyl-; 2-Hydroxy-3-Methylaniline (Pepe, Wenninger, and McEwen 2002); *o*-Cresol, 6-Amino; 6-Methyl-2-Aminophenol (Regulated Chemicals Listing 1998); 3-Amino-2-Oxy-1-Methylbenzol; and 3-Amino-2-Oxy-Toluol (Beilstein File of Organic Compounds 1998).

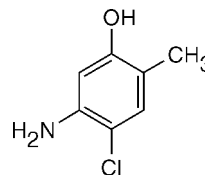
4-Amino-*m*-Cresol (CAS no. 2835-99-6) is the substituted aromatic compound that conforms to the formula (Pepe, Wenninger, and McEwen 2002):



4-Amino-*m*-Cresol is also known as 2-Amino-5-Hydroxytoluene; 4-Amino-3-Methylphenol; Phenol, 4-Amino-3-Methyl-;

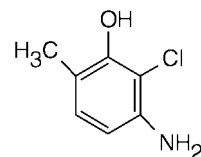
4-Hydroxy-*o*-Toluidine (Pepe, Wenninger, and McEwen 2002); 3-Methyl-4-Aminophenol (James Robinson Ltd. 1998); *p*-Amino-*m*-Cresol; *m*-Cresol, 4-Amino-; 4-Hydroxy-2-Methylaniline; *p*-Hydroxy-*o*-Toluidine; *m*-Methyl-*p*-Aminophenol; 3-Methyl-4-Aminophenol; 2-Methyl-4-Hydroxyaniline (Regulated Chemicals Listing 1998); 4-Amino-*meta*-Cresol; 6-Amino-3-Oxy-1-Methylbenzol; 6-Amino-3-Oxy-Toluol; *p*-Hydroxy-*o*-Toluidine; and Toluene, 2-Amino-5-Hydroxy (CRC Handbook of Data on Organic Compounds 1998).

5-Amino-4-Chloro-*o*-Cresol (CAS no. 110102-86-8) is an organic compound that conforms to the formula:



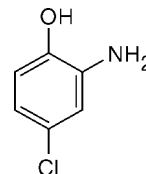
5-Amino-4-Chloro-*o*-Cresol is also known as 5-Amino-4-Chloro-2-Methylphenol; Phenol, 5-Amino-4-Chloro-2-Methyl- (Pepe, Wenninger, and McEwen 2002); and 2-Methyl-4-Chloro-5-Aminophenol (Henkel KGaA 1994).

5-Amino-6-Chloro-*o*-Cresol (CAS no. 84540-50-1) is an organic compound that conforms to the formula:



5-Amino-6-Chloro-*o*-Cresol is also known as 3-Amino-2-Chloro-6-Methylphenol; Phenol, 3-Amino-2-Chloro-6-Methyl- (Pepe, Wenninger, and McEwen 2002; Regulated Chemicals Listing 1998); 2-Chloro-3-Amino-6-Methylphenol; 2-Chloro-6-Methyl-3-Aminophenol; 3-Amino-2-Chloro-6-Methylphenol; 2-Methyl-5-Amino-6-Chlorophenol (Regulated Chemicals Listing 1998); 2-Hydroxy-3-Chloro-4-Aminotoluene; 2-Hydroxy-3-Chloro-4-Aminotoluol; and 5-Amino-6-Chloro-Benzol (Henkel KGaA 1996).

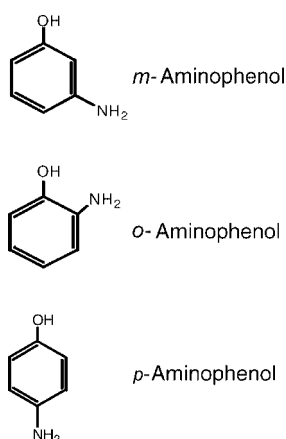
4-Chloro-2-Aminophenol (CAS no. 95-85-2) is the hair colorant that conforms to the formula:



4-Chloro-2-Aminophenol is also known as 2-Amino-4-Chlorophenol; Phenol, 2-Amino-4-Chloro-; 2-Hydroxy-5-Chloroaniline; CI 76525 (Pepe, Wenninger, and McEwen 2002; Regulated Chemicals Listing 1998); 5-Chloro-2-Hydroxyaniline; *o*-Amino-*p*-Chlorophenol; *p*-Chloro-*o*-Aminophenol; and C.I. Oxidation Base 18 (Regulated Chemicals Listing 1998).

**TABLE 1**Physical and chemical properties of 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol (Henkel KGaA 1994, 1996)

Property	Description	
	5-Amino-4-Chloro- <i>o</i> -Cresol	5-Amino-6-Chloro- <i>o</i> -Cresol
Form	Brown crystals	Beige crystals
Melting point	248°C (with decomposition)	144–183°C
Odor	None	None
Solubility	Soluble in water, propylene glycol, and triethanolamine	Soluble in water
Purity	97% (by HPLC)	>94% (by HPLC)
Molecular weight	157.59 (free base)	194.07 (hydrochloride)

**Structure of Related Ingredients**

The structures of *p*-, *m*-, and *o*-aminophenol are given above for comparison purposes. These ingredients were found safe in the present practices of use and concentrations (Elder 1988). Those use concentrations were  $\leq 1\%$ ,  $\leq 5\%$ , and  $\leq 1\%$  for *p*-, *m*-, and *o*-aminophenol, respectively, in hair tints and hair dyes and colors.

**Physical and Chemical Properties**

6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, and 4-Amino-*m*-Cresol all have a molecular weight of 123.07 and 4-Chloro-

2-Aminophenol has a molecular weight of 143.01 (Spectral Database Information System 1998). Other data on the physical and chemical properties of 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol were not available. 6-Amino-*m*-Cresol (purity grade not defined) is a solid at room temperature (Goel, Kansal, and Sharma 1979). 4-Amino-*m*-Cresol has a melting point of 176°C to 178°C, is soluble in water and organic solvents, and a 1% solution had a pH of 8.2 (James Robinson Ltd. 1998).

Physical and chemical properties of 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol are shown in Table 1.

The melting point for 6-Amino-*m*-Cresol is 163°C (CTFA 1999a). It is slightly soluble in water and soluble in many organic solvents. It is 99.9% pure as determined by elemental analysis. 6-Amino-*m*-Cresol is a crystalline powder with a beige to reddish-brown color. Upon exposure to air it becomes darker. The ultraviolet (UV) absorption data for 6-Amino-*m*-Cresol indicated absorption maxima at 210, 235, and 291 nm in ethanol. Physical and chemical properties of 6-Amino-*m*-Cresol are listed in Table 2.

The melting point for 4-Amino-*m*-Cresol is 178°C (CTFA 1999b). It is slightly soluble in water and is a crystalline powder with a reddish-brown color. It is 99.9% pure as determined by elemental analysis. When heated to decomposition it emits toxic fumes of NO. 4-Amino-*m*-Cresol is stable at normal conditions and hazardous polymerization will not occur. According to the classification of the European Directive on Classification of Hazardous Preparations, 90/492/EEC, 4-Amino-*m*-Cresol is not

**TABLE 2**Physical and chemical properties of 6-Amino-*m*-Cresol and 4-Amino-*m*-Cresol (CTFA 1999a, 1999b)

Property	Description	
	6-Amino- <i>m</i> -Cresol	4-Amino- <i>m</i> -Cresol
Form	Beige to reddish-brown crystals	Reddish-brown crystals
Melting point	163°C	178°C
Odor	Not available	Emits toxic fumes of NO when heated
Solubility	Slightly soluble in water, and many organic solvents	Slightly soluble in water
Purity	99.9% (by HPLC/GC)	99.9% (by HPLC/GC)
Molecular weight	123.16	123

a dangerous substance. The UV absorption data for 4-Amino-*m*-Cresol indicated absorption maxima at 206, 234, and 300 nm in ethanol. Physical and chemical properties of 4-Amino-*m*-Cresol are also listed in Table 2.

### Manufacture and Production

Published data on the manufacture and production of 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, 5-Amino-4-Chloro-*o*-Cresol, 5-Amino-6-Chloro-*o*-Cresol, or 4-Chloro-2-Aminophenol were not found.

### Analytical Methods

6-Amino-*m*-Cresol and 4-Amino-*m*-Cresol have each been separated using capillary electrophoresis and high-performance liquid chromatography (HPLC) utilizing crown ethers (Nishi et al. 1997). 4-Amino-*m*-Cresol has been determined using thin-layer chromatography, and identified in urine using HPLC (Son, Everett, and Fiala 1980).

### Ultraviolet Absorbance

Published data on the UV absorbance of 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol were not found. 6-Amino-*m*-Cresol has maximum absorption peaks at 210, 235, and 291 nm in ethanol (CTFA 1999a). 4-Amino-*m*-Cresol had a symmetrical absorption peak at 300 nm (James Robinson, Ltd. 1998) and maximum absorption peaks at 206, 234, and 300 nm in ethanol (CTFA 1999b).

5-Amino-4-Chloro-*o*-Cresol has a symmetrical absorption peak below 300 nm, which falls off sharply above 300 nm (Henkel KGaA 1994), and 5-Amino-6-Chloro-*o*-Cresol has a similar pattern with an even sharper fall off (Henkel KGaA 1996).

4-Amino-2-hydroxytoluene has a maximum UV absorbance at approximately 285 nm (Elder 1989).

### Impurities

Published data on the impurities of 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, or 4-Chloro-2-Aminophenol were not found.

The impurity limits for 4-Amino-*m*-Cresol specify >99.5% solid content, <1.0% sulfated ash, and <50 ppm iron, with assay of >98.0% (James Robinson Ltd. 1998). The typical analysis was >99.9% solid content, <0.5% sulfated ash, and <10 ppm iron, with assay of 98.5% to 99.5%. No *m*-cresol was detected by HPLC.

The specification of 97% purity for 5-Amino-4-Chloro-*o*-Cresol is supported by HPLC analysis; impurities include an early peak identified as 2-Methyl-5-Aminophenol (2%), and two unidentified peaks (1% combined), one of which was close to the peak of the ingredient and one that eluted later (Henkel KGaA 1994).

An HPLC analysis of 5-Amino-6-Chloro-*o*-Cresol yielded 94.19% of the ingredient in one peak. Near the major peak were

small peaks for 5-Amino-4-Chloro-2-Methylphenol (2.76%) and *p*-Amino-*o*-Cresol (1.99%). The only other significant peak (0.83%) was identified as a dichloro derivative (Henkel KGaA 1996).

### USE

#### Cosmetic

6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, 5-Amino-4-Chloro-*o*-Cresol, 5-Amino-6-Chloro-*o*-Cresol, and 4-Chloro-2-Aminophenol function as hair colorants (Pepe, Wenninger, and McEwen 2002).

5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol are specifically for use in oxidative hair dyes, with the former being used in combination with hydrogen peroxide (Henkel KGaA 1994, 1996).

The product formulation data submitted by the Food and Drug Administration (FDA) in 1998 stated that 6-Amino-*m*-Cresol was used in two hair dye and color formulations (FDA 1998). The other ingredients reviewed in this assessment were not reported to FDA as being used in 1998.

Concentration of use values are no longer reported to the FDA by the cosmetic industry (FDA 1992); the last reported concentration of use data available to CIR is from 1984 (FDA 1984). None of the ingredients reviewed in this report, however, were listed as being used in 1984.

Current information from industry indicated that 6-Amino-*m*-Cresol was used at a concentration of 2.4%, 6-Amino-*o*-Cresol was used at a concentration of 0.7%, and 4-Amino-*m*-Cresol was used at a concentration of 0.3% in all types of hair dye and colors (which require a caution statement and patch test) (CTFA 1999c).

In addition, 5-Amino-4-Chloro-*o*-Cresol is reported to be used in oxidation hair dye formulations at concentrations up to 2%, but because it is combined with hydrogen peroxide, the use concentration is only up to 1% (Henkel KGaA 1994). 5-Amino-6-Chloro-*o*-Cresol is also reported to be used in oxidative hair dyes formulations up to a final concentration of 2% (Henkel KGaA 1996).

Hair-coloring formulations are applied to or can come in contact with hair, skin (particularly at the scalp), eyes, and nails. Individuals dyeing their hair could use such formulations once every few weeks, whereas hairdressers could come in contact with products containing these ingredients several times a day. Under normal conditions of use, skin contact with hair dye is restricted to 30 min.

The hair dyes containing 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, 5-Amino-4-Chloro-*o*-Cresol, 5-Amino-6-Chloro-*o*-Cresol, and 4-Chloro-2-Aminophenol, as coal tar hair dye products, are exempt from the principal adulteration provision and from the color additive provisions in sections 601 and 706 of the *Federal Food, Drug, and Cosmetic Act* of 1938 when the label bears a caution statement and patch test instructions for determining whether the product causes skin

irritation. The following caution statement should be displayed conspicuously on the labels of coal tar hair dyes:

**Caution**—This product contains ingredients that may cause skin irritation on certain individuals, and a preliminary test according to accompanying directions should be made. This product must not be used for dyeing eyelashes or eyebrows; to do so may cause blindness.

The CIR Expert Panel has reviewed the cosmetic industry's current coal tar hair dye product labeling, which recommends that an open patch test be applied and evaluated by the beautician and/or consumer for sensitization 24 h after application of the test material and prior to the use of a hair dye formulation.

Because the recommendation on the industry's adopted labeling establishes a procedure for individual user safety testing, it is most important that the recommended procedure be consistent with current medical practice.

There is a consensus among dermatologists that screening patients for sensitization (allergic contact dermatitis) should be conducted by the procedures used by the North American Contact Dermatitis Group and the International Contact Dermatitis Group (North American Contact Dermatitis Group 1980; Eiermann et al. 1982; Adams et al. 1985). These procedures state that the test material should be applied at an acceptable concentration to the patient, covered with an appropriate occlusive patch, and evaluated for sensitization 48 and 72 h after application. The CIR Expert Panel has cited the results of studies conducted by both the North American Contact Dermatitis Group and the International Contact Dermatitis Group in its safety evaluation reports on cosmetic ingredients (Elder 1985).

During the August 26–27, 1991, public meeting of the CIR Expert Panel, all members agreed that the cosmetic industry should change its recommendation for the evaluation of the open patch test from 24 h to 48 h after application of the test material.

The industry was advised of this recommendation and asked to provide any compelling reasons why this recommendation should not be made by the Expert Panel and adopted by the cosmetic industry. No opposition to this recommendation was received. At the February 11, 1992, public meeting of the CIR Expert Panel, this policy statement was adopted.

6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, 5-Amino-4-Chloro-*o*-Cresol, 5-Amino-6-Chloro-*o*-Cresol, and 4-Chloro-2-Aminophenol do not appear in Annex II (list of substances which must not form part of the composition of cosmetic products) or Annex III (list of substances which cosmetic products must not contain except subject to the restrictions and conditions laid down) of the *Cosmetics Directive of the European Union* (European Union 1995).

### Noncosmetic

No uses for these ingredients other than in cosmetics were found.

## GENERAL BIOLOGY

### Absorption, Distribution, and Metabolism

#### *6-Amino-m-Cresol, 6-Amino-o-Cresol, and 4-Amino-m-Cresol*

Published data on the absorption, distribution, metabolism, and excretion of 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, or 4-Amino-*m*-Cresol were not found.

#### *5-Amino-4-Chloro-o-Cresol*

Skin absorption of radioactive ( $^{14}\text{C}$ ) 5-Amino-4-Chloro-*o*-Cresol was studied using six female Sprague-Dawley rats (mean weight 189.5 g). A formulation containing the ingredient, with *p*-toluenediamine sulfate, basic fatty acid emulsion, propylene glycol, water, and ammonia, was diluted 1:1 with water to make a final test ingredient concentration of 1.85%. This formulation (0.2 g) was applied to an intact, clipped area of skin (9 cm<sup>2</sup>) for 72 h under semioclusive conditions. The concentration of ingredient on the skin was 0.41 mg/cm<sup>2</sup>.

Feces and urine were monitored for 72 h, after which time the animals were sacrificed and adrenal glands, blood, brain, fat, bone, heart, kidneys, liver, lungs, muscle tissue, ovaries, spleen, thyroid glands, untreated skin, and the remaining carcass were analyzed. The mean skin absorption was 32.7%. 5-Amino-4-Chloro-*o*-Cresol was excreted via urine (92%) and feces (8%). The concentration in kidneys (0.003%) at 72 h was the greatest of any of the organ/tissue samples. The stratum corneum at the site of application, obtained by tape stripping, had 0.22% of the radioactivity (Henkel KGaA 1994).

A similar study was performed using the same strain of female rats of the same weight range except that the formulation was diluted 1:1 with a developer consisting of 6% hydrogen peroxide before application. After 30 min contact, the test material was rinsed off. Samples were taken as above. The skin absorption in this case was only 1.28%. Excretion via urine (91%) and feces (9%) accounted for all that was absorbed; the concentration in organs/tissues was at or near the detection limit of the  $^{14}\text{C}$ . The stratum corneum had 0.2% of the radioactivity and the dermis, likewise, had 0.2% (Henkel KGaA 1994).

In a third study, the metabolism of ingested 5-Amino-4-Chloro-*o*-Cresol Hydrochloride was investigated using six female Sprague-Dawley rats (mean weight 200 g). A 1.27% solution of  $^{14}\text{C}$  5-Amino-4-Chloro-*o*-Cresol Hydrochloride in a 1:1 propylene glycol/water solution was given by oral administration at a dose of 21.5 mg/kg. Feces, urine, organs, and tissues were examined as described above. 5-Amino-4-Chloro-*o*-Cresol Hydrochloride was readily absorbed in the intestine (91.7%). It was excreted via urine (94%) and feces (6%). The greatest concentration in the organ/tissue samples was 0.001% in the liver (Henkel KGaA 1994).

#### *5-Amino-6-Chloro-o-Cresol*

Skin penetration/absorption of radioactive ( $^{14}\text{C}$ ) hydrochloride was determined in a study using 12 female Wistar rats (mean weight 231 ± 7 g). Test animals were clipped and their skin

anesthetized with an i.m. injection of Ketanest<sup>®</sup> (12 ml/kg). In addition to the radioactive test ingredient, the formulation contained fatty alcohol, anionic surfactant, ammonium sulfate, water, and ammonia. The test article concentration was 1.14% and the pH was adjusted to 9.5. A dose of 20 mg/cm<sup>2</sup> was applied for 48 h without occlusive patches. Urine fractions were taken 0–8 h, 8–24 h, and 24–48 h. Feces were sampled daily. After 48 h, the animals were sacrificed and the skin and carcass assayed for radioactivity.

5-Amino-6-Chloro-*o*-Cresol hydrochloride was readily absorbed (93.2%). Radioactivity was excreted in urine (87.7%) and feces (2.22%). Only 0.48% was found in the carcass. The recovery rate of <sup>14</sup>C from the urine samples was 115% of the applied <sup>14</sup>C. An additional two animals were treated in the same manner, except that their expired CO<sub>2</sub> was monitored. No detectable <sup>14</sup>C was found in the expired CO<sub>2</sub> (Henkel KGaA 1996).

A similar study in six rats (mean body weight 217 ± 7 g) was conducted, except that the formulation was mixed 1:1 with 3% hydrogen peroxide developer solution prior to application. The test material was applied at a concentration of 15.3 mg/cm<sup>2</sup> and washed off after 30 min. Samples were collected as above. The skin penetration was only 0.116% (Henkel KGaA 1996).

The metabolism of radioactive (<sup>14</sup>C) 5-Amino-6-Chloro-*o*-Cresol was determined in five female Wistar rats (weight 254 to 270 g). A single subcutaneous (s.c.) injection of 1 g of a 5-Amino-6-Chloro-*o*-Cresol solution (0.25% in water) was given into the neck. Urine, expired CO<sub>2</sub>, and feces were collected over a period of 96 h. The animals were sacrificed and the skin and carcasses analyzed for residual radioactivity. Excretion was mainly via urine (88.5%) of which most (88.1%) was eliminated in the first 24 h. Only 3.97% was excreted in feces, and 0.674% was in the carcass and 0.04% in the injection site skin. No detectable radioactivity was found in expired CO<sub>2</sub> (Henkel KGaA 1996).

Metabolism was further studied using a single oral application of <sup>14</sup>C 5-Amino-6-Chloro-*o*-Cresol to 5 male Wistar rats (weight 321 to 336 g). Each animal received 49.4 mg/kg of the test article (1.7% in water) by gavage. Urine, expired CO<sub>2</sub>, and feces were collected as daily fractions for 96 h. The animals were sacrificed and the gastrointestinal tract and the remaining carcass were analyzed. Excretion was again mainly via urine (90.93%) and mostly (90%) in the first 24 h. There was 6% in the gastrointestinal tract and 0.58% in the remaining carcass. No <sup>14</sup>C was detected in expired CO<sub>2</sub> (Henkel KGaA 1996).

The organ distribution of <sup>14</sup>C after a single oral dose of <sup>14</sup>C 5-Amino-6-Chloro-*o*-Cresol was studied in five male Wistar rats (mean weight 323 ± 9 g). A single dose of the test article (1.7% in water) was delivered by gavage. One rat was sacrificed at each of 1, 6, 24, 48, and 96 h after administration. Whole body autoradiography was used to detect the distribution of <sup>14</sup>C. Urine and feces were collected. One hour post administration the skin, kidneys, and the content of the intestine, liver, and especially the content of the stomach were collected for analysis. After 6 h, radioactivity was in the stomach, intestine, or colon content, and in the caecum. After 24 and 48 h, only residual radioactivity was

found in the colon, caecum, and kidneys. After 96 h, excretion was nearly complete and only a small amount of label appeared (in bone). Within the first 24 h, 91% of the radioactivity was excreted via urine (Henkel KGaA 1996).

#### *4-Amino-2-Hydroxytoluene and p-Aminophenol*

Elder (1989) reported the percutaneous absorption of radioactive 4-amino-2-hydroxytoluene in a hair dye applied to the dry hair of humans under normal use conditions. The total excretion of 4-amino-2-hydroxytoluene was 0.2% ± 0.1%. This is contrasted with the oral administration in humans of radioactive 4-amino-2-hydroxytoluene in which there was a 94% recovery of the radioactivity in the urine. Elder (1988) reported the percutaneous absorption of 4-amino-2-hydroxytoluene (nonradioactive) coupled with radioactive *p*-aminophenol. The resultant <sup>14</sup>C-indamine was determined in rats under the conditions of oxidative hair dyeing. As much as 11% of the radioactivity introduced as <sup>14</sup>C-*p*-aminophenol was detected in the excreta, viscera, and skin of rats (Elder 1988); the penetration of *p*-aminophenol was similar when not coupled with 4-amino-2-hydroxytoluene. The <sup>14</sup>C-indamine formed during the oxidation did not substantially penetrate the cutaneous barrier.

### Immunological Effects

#### *4-Chloro-2-Aminophenol*

The response of leukocytes from female guinea pigs treated with 4-Chloro-2-Aminophenol was evaluated using the leukocyte adherence inhibition (LAI) technique (Naniwa 1982). Both 4-Chloro-2-Aminophenol and *p*-aminophenol were conjugated with protein by similar condensation reactions. Significantly greater amounts of LAI were found for *p*-aminophenol–protein conjugates in the treated guinea pigs, indicating that 4-Chloro-2-Aminophenol–sensitized lymphocytes could not differentiate between 4-Chloro-2-Aminophenol–and *p*-aminophenol–protein conjugates. This suggested that cross-sensitization can occur with *p*-aminophenol.

### Nephrotoxicity

#### *4-Chloro-2-Aminophenol*

Renal cortical slices from male Fischer 344 rats were used in gluconeogenesis and lactate dehydrogenase (LDH) release studies (Hong et al. 1996). The tissue slices were incubated with 0.01 to 0.5 mM 4-Chloro-2-Aminophenol in dimethyl sulfoxide (DMSO), 4-amino-2-chlorophenol, or vehicle. Renal gluconeogenesis was inhibited by ≥0.01 mM 4-Chloro-2-Aminophenol and ≥0.05 mM 4-amino-2-chlorophenol. LDH leakage was increased at concentrations of ≥0.5 mM 4-Chloro-2-Aminophenol and ≥0.1 mM 4-amino-2-chlorophenol.

#### *p-Aminophenol*

Hong et al. (1996), in an introduction to their study of chloro amino phenols, characterized *p*-Aminophenol as an acute

nephrotoxicant and a mild hepatotoxicant; *o*-Aminophenol as not toxic to the kidney or liver; and neither 4-Amino-3-chlorophenol nor 2-amino-5-chlorophenol as marked nephrotoxicant(s).

### Hepatotoxicity

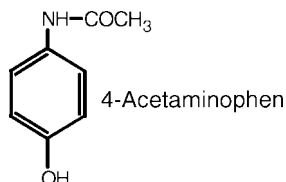
No data were available on ingredients in this safety assessment, but data on related ingredients are summarized below.

#### *p*-Aminophenol and *o*-Aminophenol

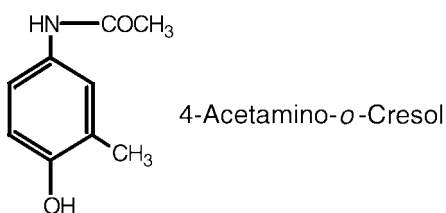
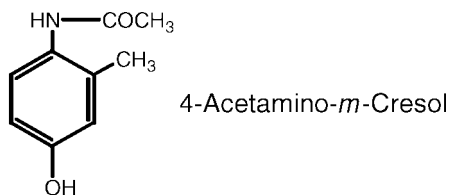
Elder (1988) reported that *p*-Aminophenol induces mild hepatotoxicity characterized by a twofold increase in serum transaminase levels, but that *o*-Aminophenol has no toxic effects on kidney or liver.

#### Acetaminophen

Acetaminophen, structure shown below, is somewhat similar to ingredients considered in this report and can be hepatotoxic in humans and experimental animals at large doses (Harvison, Forte, and Nelson 1986).



In a study to examine the role of mono-methylation in both the analgesic effect and hepatotoxicity of acetaminophen, Harvison, Forte, and Nelson (1986) prepared the following analogues that are structurally very similar to ingredients in this report:



Male Swiss-Webster mice (20 g) were injected intraperitoneally (i.p.) with either acetaminophen or the analogues shown above at various doses from 400 to 1000 mg/kg. Animals had been pretreated with either phenobarbital or cobaltous chloride and received a single i.p. dose of piperonyl butoxide 30 min before receiving the test substances. Animals were sacrificed and liver and kidney samples were taken and fixed in buffered formalin. Paraffin sections were prepared and stained with hematoxylin and eosin and examined for severity of necrosis.

The hepatotoxicity of 4-Acetamino-*o*-Cresol was comparable to that seen with acetaminophen, but 4-Acetamino-*m*-Cresol was less hepatotoxic. To the extent that these acetamino cresols are predictive of the hepatotoxicity of amino cresols, the results of these studies indicate that no greater hepatotoxicity would likely occur with the hair dye than is seen with acetaminophen, which isn't seen until g/kg doses are reached (Fethke, personal communication<sup>2</sup>).

### ANIMAL TOXICOLOGY

Published data on the toxicity of 6-Amino-*o*-Cresol in animals was not found.

#### Acute Intraperitoneal Toxicity

##### 4-Chloro-2-Aminophenol

Four male Fischer 344 rats per group were given a single i.p. injection of 0.4, 0.8, or 1.2 mmol/kg 4-Chloro-2-Aminophenol hydrochloride in 50% DMSO in distilled water, 0.4, 0.8, or 1.0 mmol/kg 4-amino-2-chlorophenol hydrochloride in distilled water, or vehicle (Hong et al. 1996). The animals were killed 48 h after dosing. 4-Chloro-2-Aminophenol had very few effects on renal function; no apparent morphological damage was observed at nonlethal doses of <0.8 mmol/kg. Changes in hepatic function or morphology were not observed. A dose of 1.2 mmol/kg 4-Chloro-2-Aminophenol killed 75% of the animals, but little evidence of nephrotoxicity was observed in the surviving animals. However, 4-amino-2-chlorophenol induced marked changes in renal function and morphology in a dose-dependent manner; no effect on hepatic function or hepatic morphology was observed.

#### Acute Dermal Toxicity

##### 4-Amino-2-Hydroxytoluene

In an acute dermal toxicity study, 4-amino-2-hydroxytoluene did not produce any systemic/dermal toxicity in rabbits at a dose of 5 g/kg (Elder 1989).

##### *p*-Aminophenol

The dermal LD<sub>50</sub> of *p*-aminophenol was >8 g/kg for rabbits (Elder 1988).

#### Acute Oral Toxicity

##### 5-Amino-4-Chloro-*o*-Cresol

Male and female Wistar rats (average body weight of 164 g for females and 183 g for males) were given 5-Amino-4-Chloro-*o*-Cresol hydrochloride by gavage at doses of 1184, 1539, and 2000 mg/kg. Observations included apathy, piloerection, cyanosis, tremor, crouch, diarrhea, semiclosed eyes, and impaired hearing. Gross observations included brightened coloration of the liver and kidneys, ulcerations in the glandular

<sup>2</sup>Available for review: Director, Cosmetic Ingredient Review, 1101 17th Street, N.W., Suite 310, Washington, DC 20036, USA.



stomach, hydrometra, brown-colored hydrocele in the intestine, and emphysema (in the one animal that died). For males, the LD<sub>50</sub> was between 1.54 and 2.0 g/kg and for females, the LD<sub>50</sub> was >2.0 g/kg (Henkel KGaA 1994).

#### *5-Amino-6-Chloro-o-Cresol*

Male albino TNO-Wistar rats (average body weight of 200 g) were given 5-Amino-6-Chloro-*o*-Cresol hydrochloride by gavage at doses of 501, 1000, 1250, 1580, and 1999 mg/kg. Observations included apathy, staggering, rapid breathing, dyspnea (at later stages), and yellow-orange discoloration of the urine. The LD<sub>50</sub> was 1.36 g/kg (Henkel KGaA 1996).

#### *4-Amino-m-Cresol*

Male CD-1 mice were dosed for 2 consecutive days (6 mice/group, route of administration not specified) with 1000, 1200, 1440, 1728, or 2074 mg/kg 4-Amino-*m*-Cresol. At 4 hours through day 2 of dosing, the following observations were observed: piloerection was observed in all groups; hypokinesia was observed in all but the low-dose group; ataxia occurred in the 1440- and 2074-mg/kg dose groups; and only mice in the 1200-mg/kg dose group had prostration. At least one mouse in all groups survived until day 14, but most mice died on day 1 or 2. The LD<sub>50</sub> value was calculated as 1000 mg/kg (Holmstroem 1980).

#### *6-Amino-m-Cresol*

Holmstroem (1980), using the same protocol described above, calculated the LD<sub>50</sub> of 6-Amino-*m*-Cresol as 1500 mg/kg.

In a pre-experiment toxicity study, Völkner and Heidemann (1991) dosed NMRI mice (2/sex/group) once with 500, 750, 1000, and 1500 mg/kg 6-Amino-*m*-Cresol in polyethylene glycol 400. Toxic reactions were observed in all groups and included reduction of spontaneous activity, eyelid closure, abdominal position, tremor, and death. One death occurred in each of the 750-, 1000-, and 1500-mg/kg groups by 6 h posttreatment. No deaths occurred in the 500-mg/kg group and the only toxic reaction observed in this group was reduction of spontaneous activity. Therefore, the 500-mg/kg group was estimated to be the maximum tolerated dose.

Leimbeck and Grötsch (1991) dosed two male and two female mice orally with 666 mg/kg 6-Amino-*m*-Cresol. In the first two hours all animals had tremor, anemia, and a slight to moderate reduction in activity. No animals died 72 h post application.

Fautz (1994) dosed two male rats once orally with 1200 mg/kg 6-Amino-*m*-Cresol in 1% carboxymethylcellulose. The rats had reduction of spontaneous activity, abdominal position, eyelid closure, and piloerection. In another experiment, two male rats each received a single oral dose of 1500 or 2000 mg/kg 6-Amino-*m*-Cresol in 1% carboxymethylcellulose, respectively. The animals in the 1500 mg/kg group had no toxic reactions except

brown-colored urine. One animal in the 2000-mg/kg group died 24 h after treatment. The 1500-mg/kg group was estimated to be the maximum tolerated dose.

#### *4-Amino-2-Hydroxytoluene*

Using rats, 10% to 20% 4-amino-2-hydroxytoluene was slightly toxic in three separate acute oral studies (Elder 1989).

#### *m-Aminophenol, o-Aminophenol, and p-Aminophenol*

The oral LD<sub>50</sub> values for rats of *p*-, *m*-, and *o*-aminophenol were 671–1270, 812–1660, and 1300 mg/kg, respectively (Elder 1988).

### Short-Term Oral Toxicity

#### *6-Amino-m-Cresol*

Male and female Wistar rats (15/sex/group) were dosed orally with 50, 250, and 500 mg/kg 6-Amino-*m*-Cresol daily for 4 weeks (Forschungs GmbH 1985). The control group was dosed with 1 ml/100 g body weight 0.5% carboxymethylcellulose (CMC). Prior to study initiation and after 4 weeks, 10 rats/sex/group had ophthalmological and reflex examinations (5/sex/group), hearing tests and blood tests.

No significant observations occurred in the 50-mg/kg group. The 250-mg/kg group had increased activity 10 min after dosing during the third and fourth week of treatment and increased, discolored urine excretion. Water consumption was also increased. Significant results included reduced erythrocyte counts in males (highly significant) and females; increased reticulocytes in females; decreased hemoglobin in males and a highly significant decrease in females; increased hematocrit in both sexes, but highly significant in males; decreased iron in females; increased hepatic weight in females; increased kidney weight in males and females; and increased spleen weights in both sexes, but highly significant in females.

The 500-mg/kg group had initial decreased activity during week 1 and later increased activity as in the previous group. Increased, discolored urine excretion was also observed. Borderline significant results were observed for decreased body weight gain and food consumption during weeks 1 and 2 in females. Highly significant results were reported for increased water consumption in both sexes at all phases of the study; decreased erythrocytes and hemoglobin and increased reticulocytes in both sexes; and decreased hematocrit in males and females, although females were within normal range. The mean corpuscular volume (MCV) and prothrombin time was significantly increased in females, but still in the normal range. Iron was significantly reduced in females. At necropsy, dark, discolored spleens were observed (sex not specified). Liver, kidney, and spleen weights were all increased in both sexes. No treatment related observations were observed at microscopic evaluation. The no-observed-adverse-effect level (NOAEL) for 6-Amino-*m*-Cresol was established at 50 mg/kg.

### Subchronic Dermal Toxicity

#### *m*-Aminophenol, *o*-Aminophenol, and *p*-Aminophenol

The dermal toxicity of hair dyes containing *m*-, *o*-, and/or *p*-aminophenol was determined using New Zealand white rabbits (Burnett et al. 1976). A dose of 1 ml/kg of oxidative hair dyes containing 0.7% *m*-aminophenol and 1.0% *p*-aminophenol, 0.7% *m*-aminophenol, 0.3% *o*-aminophenol, or 1.0% *N*-methyl-*p*-aminophenol sulfate mixed with an equal volume of 6% hydrogen peroxide or semipermanent hairdyes containing 0.09% and 0.2% *m*-aminophenol and *p*-aminophenol, respectively, or 0.02%, 0.04%, and 0.05% *m*-aminophenol, *p*-aminophenol, and *N*-methyl-*p*-aminophenol, respectively, were applied topically to the intact or abraded skin on the shaved backs of each animal twice weekly for 13 weeks, and no evidence of systemic toxicity was observed after application of the hairdyes.

### Subchronic Oral Toxicity

#### *5*-Amino-4-Chloro-*o*-Cresol

Male and female Sprague Dawley rats (males, 152 to 160 g; females, 128 to 135 g) were given 5-Amino-4-Chloro-*o*-Cresol hydrochloride by gavage daily, 5 days a week, for 90 days. Daily doses were 0, 20, 60, and 180 mg/kg. No clinical observations or pathological findings indicative of systemic toxicity were observed. Only minor deviations in a few biochemical and hematological parameters were noted. The NOAEL was established at the highest dose of 180 mg/kg (Henkel KGaA 1994).

#### *5*-Amino-6-Chloro-*o*-Cresol

Male and female Wistar rats (males, 102 to 149 g; females, 98 to 138 g) were given 5-Amino-6-Chloro-*o*-Cresol hydrochloride with tragacanth (1%) by gavage daily, 5 days a week, for 13 weeks. Daily doses were 50 mg/kg. No clinical observations, biochemical alterations, or pathological findings were indicative of systemic toxicity. The NOAEL was established at the highest dose of 50 mg/kg (Henkel KGaA 1996).

#### *4*-Amino-*m*-Cresol

Male and female Wistar rats were dosed orally with 15, 60, or 120 mg/kg 4-Amino-*m*-Cresol for 13 weeks (Forschungs GmbH 1984a). A control group was also included. The control group and the 120-mg/kg group had 25 rats/sex/group and the low- and mid-dose groups had 20 rats/sex/group. Prior to study initiation and again at 6 and 13 weeks, 5 rats/sex/group had ophthalmological, hearing, and reflex examinations. Blood samples were taken at the same time intervals on 20 rats/sex/group. Urinalyses were performed on 5 rats/sex/group.

No specific observations occurred in the 15-mg/kg group. The 60- and 120-mg/kg groups had dark, discolored urine due to compound discoloration in both sexes from treatment weeks 8 to 13. The 120-mg/kg group had significantly increased creatinine values in the female rats after 13 weeks of treatment, although the values were still within the normal range. The spleen weights were significant in female rats and increased in male rats. No

observations attributed to the test compound were found during microscopic evaluation. The NOAEL was established at the mid-dose, 60 mg/kg.

#### *4*-Amino-2-Hydroxytoluene

Elder (1989) reported that the administration of 4-amino-2-hydroxytoluene in the diet of rats at concentrations of  $\leq 3\%$  for 3 to 6 months caused reduction in body weight, a slight anemia, and sporadic microfollicular goiter. Feeding rats  $\leq 0.7\%$  *p*-aminophenol for 3 to 6 months resulted in decreased body weights and feed consumption, increased relative liver and kidney weights, and nephrosis. Feeding rats  $\leq 1\%$  *m*-aminophenol for 90 days resulted in decreased body weights and feed consumption, deposition of iron positive pigment in the spleen, liver, and kidneys, and increased thyroid gland activity.

### Acute Dermal Irritation

#### *6*-Amino-*m*-Cresol, *6*-Amino-*o*-Cresol, *4*-Amino-*m*-Cresol, and *4*-Chloro-2-Aminophenol

Published data on the dermal irritation potential of 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, or 4-Chloro-2-Aminophenol were not found.

#### *5*-Amino-4-Chloro-*o*-Cresol

The acute dermal toxicity of 5-Amino-4-Chloro-*o*-Cresol was determined using 3 adult female albino New Zealand white (SPF) rabbits. A 0.5-ml aliquot of 5-Amino-4-Chloro-*o*-Cresol was applied to intact, shaved skin on the dorsal back of each animal. A semioclusive patch was applied. After 4 h the patch was removed and the site rinsed. The skin was examined immediately after patch removal and then at 1, 24, 48, and 72 h thereafter. Only very slight erythema and edema were seen at 24 h, which disappeared at 48 and 72 h. Brown-yellow/yellow staining was seen at the application site. No information on systemic toxicity was provided (Henkel KGaA 1994).

The acute dermal irritation of 5-Amino-4-Chloro-*o*-Cresol was determined using six adult male albino New Zealand rabbits. A 0.5-ml aliquot of a 10% formulation (3 g of 5-Amino-4-Chloro-*o*-Cresol, 10 ml of distilled water, and 5 ml ammonium sulfate dissolved to a total volume of 30 ml in 96% ethanol) was applied to intact, shaved skin on the dorsal back of each animal. An occlusive patch was applied for 2 h. The skin was examined immediately after patch removal and then at 24 and 48 h. No signs of erythema, edema, or eschar formation were seen and the animals had no signs of systemic toxicity (Henkel KGaA 1994).

#### *5*-Amino-6-Chloro-*o*-Cresol

The acute dermal toxicity of 5-Amino-6-Chloro-*o*-Cresol was determined using six adult male albino New Zealand rabbits. A 10% aqueous formulation (3 g of 5-Amino-6-Chloro-*o*-Cresol, 10 ml distilled water, and 5 ml ammonium sulfate dissolved to a total volume of 30 ml in 96% ethanol) was applied to a shaved area (0.5 ml/10 cm<sup>2</sup>) on the dorsal back of each

animal. An occlusive patch was applied for 2 h. The skin was examined immediately after patch removal and then at 24 and 48 h. No signs of erythema, edema, or eschar formation were seen and the animals had no signs of systemic toxicity (Henkel KGaA 1996).

### Repeated Dermal Application

#### *5-Amino-4-Chloro-o-Cresol*

Five adult male hairless mice (hr/hr strain) were used to assess skin irritation associated with repeated application of a 10% dilution of 5-Amino-4-Chloro-*o*-Cresol hydrochloride, adjusted to pH 8 with ammonia. Applications (one or two drops only) were made to the same area of the back once a day for 5 working days and twice a day for 4 working days for a total of 9 consecutive working days. Animals were examined before each application and the responses scored. No primary skin irritation was observed (Henkel KGaA 1994).

#### *5-Amino-6-Chloro-o-Cresol*

Five adult male hairless mice (hr/hr strain) were used to assess skin irritation associated with repeated application of a 10% aqueous formulation (3 g of 5-Amino-6-Chloro-*o*-Cresol, 10 ml distilled water, and 5 ml ammonium sulfate dissolved to a total volume of 30 ml in 96% ethanol). One drop was applied to the same spot on the dorsal back, twice per day, for 5 consecutive days. No signs of primary skin irritation were observed (Henkel KGaA 1996).

Repeated application of 5-Amino-6-Chloro-*o*-Cresol to 6 adult male New Zealand rabbits was studied by Henkel KGaA (1996). One drop of a 10% aqueous formulation (3 g of 5-Amino-6-Chloro-*o*-Cresol, 10 ml distilled water, and 5 ml ammonium sulfate dissolved to a total volume of 30 ml in 96% ethanol) was applied to the same shaved area of the dorsal back every 30 s for a total of 60 applications. No signs of primary irritation were observed.

#### *4-Amino-2-Hydroxytoluene*

Elder (1989) reported that a concentration of 2.5%, 4-amino-2-hydroxytoluene was essentially nonirritating.

#### *m-Aminophenol, o-Aminophenol, and p-Aminophenol*

Elder (1988) reported that *p*- and *m*-Aminophenol were mildly irritating to rabbit skin; that *p*- and *o*-Aminophenol were both nonirritating when applied to intact and abraded rabbit skin under occlusive patches and to intact rabbit skin under semioclusive patches; and that *m*-Aminophenol, 3%, was not irritating when applied to the backs of rabbits.

### Sensitization

#### *6-Amino-m-Cresol, 6-Amino-o-Cresol, and 4-Amino-m-Cresol*

Published data on the sensitization potential of 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, or 4-Amino-*m*-Cresol were not found.

#### *4-Chloro-2-Aminophenol*

The sensitization potential of 4-Chloro-2-Aminophenol and cross-sensitization potential with *p*-aminophenol was determined using guinea pigs (Naniwa 1982). (4-Chloro-2-Aminophenol and *p*-aminophenol belong to the same amino derivative class and have common side chains on the benzoic ring.) Fifteen female guinea pigs were first injected with an emulsion of 200 mg of 4-Chloro-2-Aminophenol in 0.5 ml *N,N*-dimethylformamide and 0.5 ml Freund's complete adjuvant. At 2 or 3, 4, and 6 weeks after treatment, the animals were patch tested with 0.1%, 0.5%, and 1.0% 4-Chloro-2-Aminophenol in equal volumes of dioxan and acetone. The solutions, 0.05 ml, were applied to the shaved dorsal area of each animal, and the sites were not covered. The test sites were scored 24 h after application of 4-Chloro-2-Aminophenol. Following patch testing with 4-Chloro-2-Aminophenol, a 1.0% *p*-aminophenol solution was applied using the same procedure. Five animals that were not treated were patch tested with 4-Chloro-2-Aminophenol and *p*-aminophenol and served as a control group.

One test animal died by week 6 of the study (reason for death not stated.) At weeks 2 to 3, 1, 1, and 3 of the 15 test animals had reactions (weak or strong erythema) at the 0.1%, 0.5%, and 1.0% 4-Chloro-2-Aminophenol sites, respectively. During the fourth week of the study, 2, 8, and 13 animals had reactions at the 0.1%, 0.5%, and 1.0% 4-Chloro-2-Aminophenol sites, respectively. During the sixth week of the study, 2, 7, and 13 of the 14 remaining test animals had reactions at the 0.1%, 0.5%, and 1.0% 4-Chloro-2-Aminophenol sites, respectively. None of the test animals reacted to *p*-aminophenol and none of the control animals reacted to 4-Chloro-2-Aminophenol or *p*-aminophenol.

#### *5-Amino-4-Chloro-o-Cresol*

Henkel KGaA (1994) conducted a guinea pig maximization study of 5-Amino-4-Chloro-*o*-Cresol using 20 female Pirbright White animals. Fifteen animals were used to determine the minimum irritant and maximum nonirritant concentration. Induction was done with injection of 0.1 ml of a 0.25% aqueous solution of 5-Amino-4-Chloro-*o*-Cresol (adjusted to pH 8 with ammonia) as the minimum irritant concentration and two injections of 0.1 ml of a 0.5% aqueous solution of 5-Amino-4-Chloro-*o*-Cresol diluted 1:1 with Freund's complete adjuvant (FCA). Controls were treated with FCA and vehicle only. The second topical induction was done 1 week later with 1.0 ml of a 5% aqueous solution of 5-Amino-4-Chloro-*o*-Cresol under an occlusive patch for 48 h. The challenge was done 14 days after the second induction with 0.2 ml of a 2% aqueous solution of 5-Amino-4-Chloro-*o*-Cresol applied to the animals' flanks under an occlusive patch. Animals were examined at 24 and 48 h after removal of the patch.

After the first and second inductions, all animals had typical reactions to FCA. Almost 50% of the test animals (9/19; no explanation provided for the fate of the 20th animal) had slight erythema 24 h after challenge, but only 5 animals had

this minimal effect after 48 h. It was concluded that 5-Amino-4-Chloro-*o*-Cresol is a moderate sensitizer in the maximization test.

Henkel KGaA (1994) performed a second maximization study using a hair dye formulation containing *p*-toluidine diamine and 5-Amino-4-Chloro-*o*-Cresol hydrochloride. The hair dye formulation was diluted 1:1 with 6% hydrogen peroxide before use in the experiment. As in the previous study, 15 female Pirbright White guinea pigs were used to determine irritant concentrations and 20 animals were included in the maximization test. Intradermal induction was done with injection of 0.1 ml of a 0.1% aqueous solution of the hair dye/oxidizer combination and two injections of a 0.2% solution diluted 1:1 with FCA. Controls were treated only with FCA and vehicle. The second, topical induction was done 1 week later with 1.0 ml of the test substance (hair dye/oxidizer combination) under an occlusive patch for 48 h. The challenge was done 14 days after the second induction using 0.2 ml of a 2.5% aqueous solution of the test material on the flank under an occlusive patch for 24 hours.

After the inductions, animals had typical reactions to FCA. None of the animals exposed to the test substance had any reactions. As found in a hair dye formulation mixed with an oxidizer, 5-Amino-4-Chloro-*o*-Cresol was a non-sensitizer in the maximization test.

Henkel KGaA (1994) conducted a third maximization test with a second hair dye formulation containing 2,4,5,6-tetra-amino-pyrimidine and 5-Amino-4-Chloro-*o*-Cresol. The hair dye formulation was diluted 1:1 with 6% hydrogen peroxide as an oxidizer before use in the experiment. As above, 15 female Pirbright White guinea pigs were used to determine irritant concentrations and 20 animals were included in the maximization test. Intradermal induction was done with injection of 0.1 ml of a 0.1% aqueous solution of the hair dye/oxidizer combination and two injections of a 0.2% solution diluted 1:1 with FCA. Controls were treated only with FCA and vehicle. The second, topical induction was done 1 week later with 1.0 ml of a 20% aqueous solution of the test substance (hair dye/oxidizer combination) under an occlusive patch for 48 h. The challenge was done 14 days after the second induction using 0.2 ml of a 2.5% aqueous solution of the test material on the flank under occlusive patches for 24 hours.

After the inductions, animals had typical reactions to FCA. None of the animals exposed to the test substance had any reactions. As found in this second hair dye formulation mixed with an oxidizer, 5-Amino-4-Chloro-*o*-Cresol hydrochloride was a nonsensitizer in the maximization test (Henkel KGaA, 1994).

Henkel KGaA (1994) also performed a Buehler method sensitization test using Dunkin-Hartley guinea pigs. Four animals were used to determine minimum irritant and maximum nonirritant concentrations and 20 animals were used in the sensitization test proper. Topical induction was done on the left body side on days 1, 8, and 15 with 0.5 ml of an ethanolic paste consisting of 5-Amino-4-Chloro-*o*-Cresol in ethanol (63% *w/w*) under occlusive patches for 6 h. Control animals were dosed with ethanol

only. The challenge was done 14 days later by exposing the animals' flanks to 0.5 ml of the paste for 6 h under occlusive patches. Animals were examined 24 and 48 h after patch removal.

Neither test animals nor controls had reactions on challenge, so 5-Amino-4-Chloro-*o*-Cresol was not considered to be a sensitizer in this test (Henkel KGaA 1994).

#### *5-Amino-6-Chloro-o-Cresol*

Henkel KGaA (1996) conducted a guinea pig maximization study of 5-Amino-6-Chloro-*o*-Cresol hydrochloride using 20 female Pirbright White animals. Induction was done with injection of 0.1 ml of a 5.0% aqueous solution of 5-Amino-6-Chloro-*o*-Cresol and two injections of 0.1 ml of a 5.0% aqueous solution of 5-Amino-6-Chloro-*o*-Cresol diluted 1:1 with FCA. Controls were treated with FCA and vehicle only. The second topical induction was done 1 week later with 1.0 ml of a 5% cream of 5-Amino-6-Chloro-*o*-Cresol in petroleum jelly under an occlusive patch for 48 h. The challenge was done 14 days after the second induction with a 25% cream of the test substance applied to the animals' flanks under an occlusive patch. Animals were examined at 24 and 48 h after removal of the patch.

After the first and second inductions, all animals had typical reactions to FCA. One quarter of the test animals had slight erythema 24 h after challenge, but no effects were evident after 48 h. It was concluded that 5-Amino-6-Chloro-*o*-Cresol is not a sensitizer in the maximization test (Henkel KGaA 1996).

Using guinea pigs, 4-amino-2-hydroxytoluene was a mild sensitizer in a maximization test and a very weak sensitizer in a test using an open epicutaneous method (Elder 1989). Application to guinea pigs of 0.1% to 2% *p*-aminophenol in petrolatum under occlusive patches resulted in a concentration-dependent incidence of sensitization, with 3 of 10 animals sensitized with 0.1% and 9 of 10 animals sensitized at 2% *p*-aminophenol (Elder 1988). *p*-Aminophenol, 3% in deionized water, was not a sensitizer in guinea pigs. In an open epicutaneous test using guinea pigs, 3% *p*-aminophenol produced weak reactions in 4 of 20 animals and 3% *m*-aminophenol was not a sensitizer. In a maximization test, moderately strong cross-reactions to *o*-aminophenol application were observed in some guinea pigs previously sensitized with *p*-phenylenediamine.

#### **Photosensitization**

Published data on the photosensitization potential of ingredients reviewed in this safety assessment were not found.

#### *4-Amino-2-Hydroxytoluene*

Elder (1989) reported that 4-Amino-2-hydroxytoluene, with induction and challenge concentrations of 5% and 10%, respectively, was not a photosensitizer when evaluated using guinea pigs.

*m-Aminophenol, o-Aminophenol, and p-Aminophenol*

Elder (1988) reported that *p*-Aminophenol and *m*-aminophenol, both with induction and challenge concentrations of 10% and 5%, respectively, were not photosensitizers, but they did induce a contact hypersensitivity reaction.

**Ocular Irritation***6-Amino-m-Cresol, 6-Amino-o-Cresol, 4-Amino-m-Cresol, and 4-Chloro-2-Aminophenol*

Published data on the ocular irritation potential of 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, or 4-Chloro-2-Aminophenol were not found.

*5-Amino-4-Chloro-o-Cresol*

A volume of 0.1 ml of 5% aqueous 5-Amino-4-Chloro-*o*-Cresol hydrochloride was instilled into the conjunctival sac of six male albino New Zealand rabbits; no rinsing was done. Eye irritation reactions were scored 2, 6, 24, and 48 h after exposure. No effects on the cornea or the iris, and only slight conjunctival erythema and edema up to 24 h were observed (Henkel KGaA 1994).

*5-Amino-6-Chloro-o-Cresol*

A quantity of 51 mg of 5-Amino-6-Chloro-*o*-Cresol hydrochloride was instilled into the conjunctival sac of the right eye of one female albino New Zealand rabbit; none of the eyes were rinsed. Ocular irritation reactions were scored 1, 24, 48, and 72 h after exposure. Instillation of the undiluted ingredient produced immediate severe ocular irritation, and additional study was terminated. Corneal opacity, injection of the iris, and irritation of the conjunctivae persisted throughout the duration of the study. Undiluted 5-Amino-6-Chloro-*o*-Cresol hydrochloride was considered a severe ocular irritant (Henkel KGaA 1996).

In a second study, a volume of 0.1 ml of 5% aqueous 5-Amino-6-Chloro-*o*-Cresol hydrochloride was instilled into the conjunctival sac of four male albino New Zealand rabbits; none of the eyes were rinsed. Ocular irritation reactions were scored 1, 6, 24, and 48 h after exposure. No effects on the cornea or the iris, and only slight conjunctival erythema up to 6 h were observed. Exudation was observed after 1 h in all four animals, in three animals at 6 h, and in one animal at 24 h; the effect was not seen at 48 h. The researchers considered 5% 5-Amino-6-Chloro-*o*-Cresol hydrochloride to be very slightly irritating (Henkel KGaA 1994).

*4-Amino-2-Hydroxytoluene, m-Aminophenol, o-Aminophenol, and p-Aminophenol*

At a concentration of 2.5%, 4-amino-2-hydroxytoluene (Elder 1989), *p*-aminophenol, and *m*-aminophenol (Elder 1988) were essentially nonirritating to rabbit eyes. In Draize tests, *p*-aminophenol (powder form) was not an eye irritant and

*o*-Aminophenol did not irritate the cornea or iris and produced a cumulative conjunctival irritation score of 3.3/20.

**REPRODUCTIVE AND DEVELOPMENTAL TOXICITY**

Published data on the reproductive and developmental toxicity of 6-Amino-*o*-Cresol or 4-Chloro-2-Aminophenol were not found.

**Dermal***m-Aminophenol, o-Aminophenol, and p-Aminophenol*

The teratogenic potential of hair dyes containing *m*-, *o*-, and/or *p*-aminophenol were determined using rats (Burnett et al. 1976). A dose of 2 ml/kg of oxidative hair dyes containing 0.7% *m*-aminophenol and 1.0% *p*-aminophenol, 0.7% *m*-aminophenol, 0.3% *o*-aminophenol, or 1.0% *N*-methyl-*p*-aminophenol sulfate mixed with an equal volume of 6% hydrogen peroxide or semipermanent hair dyes containing 0.09% and 0.2% *m*-aminophenol and *p*-aminophenol, respectively, or 0.02%, 0.04%, and 0.05% *m*-aminophenol, *p*-aminophenol, and *N*-methyl-*p*-aminophenol sulfate, respectively, were applied topically to the animals on days 1, 4, 7, 10, 13, 16, and 19 of gestation. The hair dyes were not teratogenic or embryotoxic.

Burnett and Goldenthal (1988) conducted a two-generation reproduction study using rats. Twice weekly, 0.5 ml of oxidative hair dye formulations containing 0.7% *m*-aminophenol and 1.0% *p*-aminophenol, 0.7% *m*-aminophenol, 0.3% *o*-aminophenol, or 1.0% *N*-methyl-*p*-aminophenol sulfate mixed with an equal volume of 6% hydrogen peroxide was applied to a shaved area of the back of each animal. Successive applications were made to adjacent areas to minimize dermal irritation. When the rats were 100 days old, they were mated to produce an F<sub>1a</sub> generation that was eventually used in a carcinogenicity study. The F<sub>0</sub> generation was reduced and re-mated to produce an F<sub>1b</sub> generation. Rats from the F<sub>1b</sub> litters were mated after 100 days to produce F<sub>2a</sub> and F<sub>2b</sub> litters. Male and female F<sub>2</sub> parents were selected and mated to produce an F<sub>3</sub> generation. However, a viral infection resulted in poor reproductive performance for all groups, including controls, invalidating the results. Dermal irritation consisting of intermittent mild dermatitis was noted during the treatment period in each generation. The topical application of oxidative hair dye formulations did not have an adverse effect on reproductive performance or on the health and survival of the developing fetus and postnatal animals.

**Oral***6-Amino-m-Cresol*

Female Sprague-Dawley rats were dosed orally with 5, 50, or 200 mg/kg 6-Amino-*m*-Cresol from days 6 to 15 of gestation (Hazleton Laboratories 1982). A control (distilled water) and positive control (vitamin A, 15 mg/kg) were also included. The control, positive-control, and 5- and 50-mg/kg groups had

23 animals per group, whereas 26 animals were used in the high-dose group. Rats were killed on day 19 of gestation.

No mortalities were attributed to treatment effects. No clinical changes were observed in any group. Body weight gain of all treated groups was comparable to the control group. No significant changes were observed at necropsy. No effect on pregnancy incidence was observed in the treated groups. The mean number of corpora lutea and the mean number of implantations per dam (preimplantation loss) were comparable to control groups. Postimplantation loss was not affected by 6-Amino-*m*-Cresol and postimplantation loss was lowest in the 200-mg/kg group. The number and sex of the fetuses and the litter and mean fetal weights in the treatment groups were comparable to the control group. Fetal defects, visceral and skeletal variations were the same as the control group. No malformations occurred in the treated groups. The positive control group had marked teratogenic effects: the majority of fetuses had exencephaly. 6-Amino-*m*-Cresol did not elicit embryotoxicity, embryoletality, or teratogenicity.

#### *5-Amino-4-Chloro-o-Cresol*

Pregnant Wistar/HAN rats (190 to 238 g) were dosed with 5-Amino-4-Chloro-*o*-Cresol hydrochloride in water (10 ml/kg) daily by gavage on days 6 to 15 of pregnancy (period of major organogenesis in the fetus). Four groups of 25 animals each received doses of 0, 20, 100, or 500 mg/kg/day of 5-Amino-4-Chloro-*o*-Cresol hydrochloride. Maternal mortality and body weight gain were recorded. The dams were killed on day 21 of gestation and the fetuses removed for examination. The number of alive and dead fetuses, fetal weight, sex, site of implantation in the uterus, early and late resorptions, and number of corpora lutea were determined. Half of the fetuses were selected at random and examined for visceral and brain abnormalities. The remaining fetuses were examined for abnormalities after staining with alizarin.

The only maternal effect seen was a brown discoloration of the urine. At examination of the fetuses, no developmental toxicity was associated with treatment with 5-Amino-4-Chloro-*o*-Cresol hydrochloride (Henkel KGaA 1994).

#### *5-Amino-6-Chloro-o-Cresol*

Pregnant Wistar/HAN rats (186 to 234 g) were exposed to 5-Amino-6-Chloro-*o*-Cresol hydrochloride in water daily by gavage on days 6 to 15 of pregnancy (period of major organogenesis in the fetus). Four groups of 25 animals each received doses of 0, 30, 90, or 270 mg/kg/day of 5-Amino-6-Chloro-*o*-Cresol hydrochloride. Maternal mortality and body weight gain were recorded. The dams were killed on day 21 of gestation and the fetuses removed for examination. The number of alive and dead fetuses, fetal weight, sex, site of implantation in the uterus, early and late resorptions, and number of corpora lutea were determined. Half of the fetuses were selected at random and examined for visceral and brain abnormalities. The remaining fetuses were examined for abnormalities after staining with alizarin.

The only maternal effects were slight reduction in feed consumption and reduced body weight gain in the highest dose group. The NOAEL was considered to be 90 mg/kg/day. No developmental toxicity was associated with treatment with 5-Amino-6-Chloro-*o*-Cresol hydrochloride (Henkel KGaA 1994).

#### *4-Amino-m-Cresol*

Female rats (strain BOR:WISW-SPF TNO) were dosed orally with 10, 40, or 80 mg/kg 4-Amino-*m*-Cresol from days 5 to 15 of gestation (Forschungs GmbH 1984b). A control group was included. Positive proof of sperm in the vaginal smear was considered day 0 of gestation. Each group consisted of 24 animals. Dams were killed on day 20 of gestation.

No abnormal clinical observations were found during the study and no mortalities occurred. Body weight gain and food consumption had no significant intergroup differences. No abnormalities were observed at gross necropsy. No significant differences were observed between groups in mean number of fetuses per dam, left-right intrauterine distribution, sex ratio, birth position, weight, death of fetuses and live birth index, number of resorptions, resorption indices, implantations, postimplantation loss index, corpora lutea and placenta, gravid uteri, and uteri weights. External and skeletal examination of fetuses revealed no malformations. Visceral examination included one fetus in the 40-mg/kg group with hydrocephaly and two fetuses in the 80-mg/kg group with minor visceral anomalies (increased renal pelvic cavitation). The malformation index for all groups was 0, except the 40-mg/kg group, which had a malformation index of 0.56%. The NOAEL was established at the high dose, 80 mg/kg.

#### *4-Amino-2-Hydroxytoluene*

Oral administration of  $\leq 3\%$  4-amino-2-hydroxytoluene produced maternal toxicity but was not teratogenic (Elder 1989).

#### *m-Aminophenol and p-Aminophenol*

Oral administration of 250 mg/kg *p*-aminophenol resulted in reduced maternal body weight gains and teratogenicity in offspring (external, skeletal, and visceral malformations) in a study using rats (Elder 1988). Chronic feeding of 0.7% *p*-aminophenol in the diet of rats produced embryotoxicity mediated by maternal toxicity. Chronic feeding of  $\leq 1\%$  *m*-aminophenol to rats resulted in maternal toxicity during gestation, but teratogenic effects were not observed. Oral administration of 100 to 200 mg/kg *p*-aminophenol to gravid hamsters did not produce teratogenic effects.

#### **Parenteral**

##### *m-Aminophenol, o-Aminophenol, and p-Aminophenol*

Elder (1988) reported that intravenous and i.p. administration of 100 to 200 mg/kg *p*-aminophenol induced fetal malformations; i.p. administration of *o*-aminophenol to hamsters resulted in teratogenic effects; but that no conclusive evidence was found for *m*-aminophenol using i.p. administration.

## GENOTOXICITY

### In Vitro

#### 6-Amino-*m*-Cresol

The mutagenic potential of 6-Amino-*m*-Cresol was evaluated in an Ames test using *Salmonella typhimurium* strains TA1535, TA1537, TA1538, TA98, and TA100 (Noser 1979a). Concentrations of 30 to 1000  $\mu\text{g}$  6-Amino-*m*-Cresol, alone and with equal amounts of 6% hydrogen peroxide, were tested with and without metabolic activation. Negative and positive controls were used. 6-Amino-*m*-Cresol was slightly mutagenic towards *S. typhimurium* TA100 with and without metabolic activation. It was not mutagenic towards the other strains.

*Saccharomyces cerevisiae* diploid D7 cell cultures were exposed to 0.1 ml of 6-Amino-*m*-Cresol in DMSO at concentrations of 0.6, 3.0, and 15.0  $\mu\text{g}/\text{ml}$  with and without metabolic activation (Bootman 1984a). Negative (DMSO) and positive (ethyl methanesulphonate) controls were used. 6-Amino-*m*-Cresol was highly toxic to the yeast cells, but it did not induce increases in the frequency of revertant or aberrant colonies with or without metabolic activation.

Mouse lymphoma L5178Y cells were treated for 2 h with 400  $\mu\text{l}$  of 12.5 to 200  $\mu\text{g}/\text{ml}$  6-Amino-*m*-Cresol in DMSO with and without metabolic activation (Martin 1983). DMSO was used as the negative control and benzopyrene with metabolic activation and 4-nitroquinoline-1-oxide without metabolic activation were used as the positive controls. All microtitre plates were incubated for 2 weeks, after which wells with viable clones were counted. Cell viability was measured by adding ouabain and 6-thioguanine to cell suspensions 48 h and 7 days after treatment, respectively. 6-Amino-*m*-Cresol did induce an increase in mutation to both ouabain and 6-thioguanine resistance in the presence of metabolic activation; however, the increase was not considered significant with or without metabolic activation.

The clastogenic potential of 6-Amino-*m*-Cresol hemisulfate was determined using cultured male human peripheral lymphocytes (Bootman 1984b). Cell cultures were incubated for 24 h with 25  $\mu\text{l}$  of the test compound dissolved in DMSO at concentrations of 0.6, 3.0, and 15.0  $\mu\text{g}/\text{ml}$  with and without metabolic activation. DMSO was used as the negative control and cyclophosphamide with metabolic activation was used as the positive control. 6-Amino-*m*-Cresol hemisulfate did not significantly increase the number of aberrations as compared to controls.

#### 4-Amino-*m*-Cresol

The mutagenic potential of 4-Amino-*m*-Cresol was evaluated in an Ames test using *S. typhimurium* strains TA1535, TA1537, TA1538, TA98, and TA100 (Noser 1979b). Concentrations of 15 to 600  $\mu\text{g}/\text{plate}$  4-Amino-*m*-Cresol, alone and with equal amounts of 6% hydrogen peroxide, were tested with and without metabolic activation. Negative and positive controls were used. 4-Amino-*m*-Cresol was not mutagenic with or without metabolic activation.

In an unscheduled DNA synthesis (UDS) assay, male rat primary hepatocytes were incubated with 1.0, 3.33, 10.0, 33.33, or 100.0  $\mu\text{g}/\text{ml}$  4-Amino-*m*-Cresol in DMSO (Miltenburger 1986). Negative controls were untreated or incubated with solvent and positive controls were incubated with 7,12-dimethylbenz(a)anthracene. 4-Amino-*m*-Cresol did not induce UDS in rat hepatocytes.

#### 4-Chloro-2-Aminophenol

The mutagenic potential of 4-Chloro-2-Aminophenol in DMSO was determined in a preincubation assay (Zeiger et al. 1988). Concentrations of 10 to 1500  $\mu\text{g}/\text{plate}$  were tested using *S. typhimurium* strains TA100, TA1535, TA97, and TA98 with and without metabolic activation. 4-Chloro-2-Aminophenol was weakly mutagenic.

#### 5-Amino-4-Chloro-*o*-Cresol

The mutagenic potential of 5-Amino-4-Chloro-*o*-Cresol hydrochloride was evaluated in an Ames test using *S. typhimurium* strains TA1535, TA1537, TA1538, TA98, and TA100 (Henkel KGaA 1994). Concentrations of 4 to 2500  $\mu\text{g}/\text{plate}$  with the 5-Amino-4-Chloro-*o*-Cresol hydrochloride dissolved in water and 75 to 1200  $\mu\text{g}/\text{plate}$  with the 5-Amino-4-Chloro-*o*-Cresol (the free base) dissolved in DMSO were tested with and without metabolic activation by Aroclor 1254-induced rat liver enzymes. Positive controls were used as follows: Sodium azide for TA 100 and TA 1535; 9-aminoacridine for TA 1537; 4-nitro-*o*-phenylenediamine for TA 98 and TA 1538; and 2-aminoanthracene for all strains. Toxic effects were noted at the greatest concentration tested (2500  $\mu\text{g}/\text{plate}$ ). Table 3 has a summary of the results of this study. On the basis of these data, the investigators concluded that the free base was mutagenic with metabolic activation.

V79 Chinese hamster lung cells were used to examine the mutagenicity of 5-Amino-4-Chloro-*o*-Cresol hydrochloride. Mutations to 6-thioguanine resistance at the *HGRPT* locus with

**TABLE 3**  
5-Amino-4-Chloro-*o*-Cresol Ames test results (Henkel KGaA 1994)

Strain	With metabolic activation		Without metabolic activation	
	Hydrochloride in water	Free base in DMSO	Hydrochloride in water	Free base in DMSO
TA 98	Neg	Weak pos	Neg	Neg
TA 100	Weak pos	Pos	Neg	Neg
TA 1535	Neg	Neg	Neg	Neg
TA 1537	Neg	Weak pos	Neg	Neg
TA 1538	Neg	Pos	Neg	Neg

Neg, negative; Pos, positive.

**TABLE 4**  
5-Amino-6-Chloro-*o*-Cresol Ames test results (Henkel KGaA 1996)

Strain	With metabolic activation		Without metabolic activation
	Phenobarbital	Aroclor 1254	
TA 98	Neg	Pos	Neg
TA 100	Neg	Pos	Neg
TA 1535	Neg	Neg	Neg
TA 1537	Neg	Neg	Neg
TA 1538	Neg	Pos	Neg

Neg, negative; Pos, Positive.

and without metabolic activation were measured. 5-Amino-4-Chloro-*o*-Cresol hydrochloride dissolved in ethanol at 6 to 60  $\mu\text{g/ml}$  without metabolic activation and 55 to 550  $\mu\text{g/ml}$  with metabolic activation (Aroclor 1254-induced rat liver enzyme fraction) were used. Ethyl methanesulfonate (EMS) and dimethylbenz[*a*]anthracene (DMBA) served as positive controls. At no concentration or metabolic activation status were any increases seen in the number of mutations (Henkel KGaA 1994).

#### 5-Amino-6-Chloro-*o*-Cresol

The mutagenic potential of 5-Amino-6-Chloro-*o*-Cresol hydrochloride was evaluated in an Ames test using *S. typhimurium* strains TA1535, TA1537, TA1538, TA98, and TA100 (Henkel KGaA 1996). Concentrations of 4 to 2500  $\mu\text{g/plate}$  with the 5-Amino-6-Chloro-*o*-Cresol hydrochloride was tested with and without metabolic activation by Aroclor 1254 or phenobarbital induced rat liver enzymes. Positive controls were used as follows: Sodium azide for TA 100 and TA 1535; 9-aminoacridine for the other strains. Table 4 presents the results of this study. On the basis of these data, the investigators concluded that 5-Amino-6-Chloro-*o*-Cresol hydrochloride was mutagenic with metabolic activation.

V79 Chinese hamster lung cells were used to examine the mutagenicity of 5-Amino-6-Chloro-*o*-Cresol hydrochloride. Mutations to 6-thioguanine resistance at the *HGRPT* locus with and without metabolic activation were measured. 5-Amino-4-Chloro-*o*-Cresol hydrochloride dissolved in ethanol at 0, 35, 100, 200, and 300  $\mu\text{g/ml}$  without metabolic activation and 0, 25, 100, 200, and 300  $\mu\text{g/ml}$  with metabolic activation (Aroclor 1254-induced rat liver enzyme fraction) were used. EMS and DMBA served as positive controls. At concentrations  $\geq 50$   $\mu\text{g/ml}$ , the plating efficiency of the cells was slightly reduced. At no concentration or metabolic activation status were any increases seen in the number of mutations (Henkel KGaA 1996).

V79 Chinese hamster lung cells were used to examine the mutagenicity of 5-Amino-6-Chloro-*o*-Cresol hydrochloride at concentrations from 10 to 1100  $\mu\text{g/ml}$ . Chromosomes were prepared 7 (high dose), 18 (low, medium, and high dose), and 28

(high dose) h after the start of a 4-h treatment. Treatment was done with and without Aroclor 1254-induced rat liver enzymes. EMS was used as a positive control. Concentrations of 1000 and 3000  $\mu\text{g/ml}$  were toxic in range finding studies, with and without metabolic activation. Although no chromosome aberrations were seen at 7 h, chromosome aberrations were increased in all dose groups at 18 h and at 28 h. The authors concluded that 5-Amino-6-Chloro-*o*-Cresol hydrochloride does induce chromosome aberrations in the V79 line independent of metabolic activation (Henkel KGaA 1996).

Unscheduled DNA synthesis (a measure of DNA damage) was measured in rat liver hepatocytes exposed to 5-Amino-6-Chloro-*o*-Cresol hydrochloride at concentrations ranging from 6.67 to 2000  $\mu\text{g/ml}$ . Six cultures were used for each concentration and the experiments were repeated three times. Cells were incubated without the test compound for 1 h, at which time tritiated thymidine and the test substance were added and incubated a further 3 h. 2-Acetylaminofluorene (2-AAF) served as a positive control. Cells were washed, nuclei isolated, and the incorporated radioactivity was measured. Total DNA content was determined colorimetrically. No indications of a dose-related increase in unscheduled DNA synthesis were observed (Henkel KGaA 1996).

#### In Vivo

##### 6-Amino-*m*-Cresol

In a micronucleus test, male CD-1 mice (10 per group) were dosed orally with 30, 150, or 750 mg/kg 6-Amino-*m*-Cresol in 0.5% carboxymethylcellulose at a volume of 10 ml/kg once daily for 2 days (Holmstroem 1980). The mice were dosed during two separate studies 6 and 30 h before they were killed. The vehicle was used as a negative control and 100 mg/kg cyclophosphamide was used as a positive control. Body weights did not vary by more than 1 g during the study. 6-Amino-*m*-Cresol did not increase the frequency of micronuclei.

In another micronucleus test, groups of six male and female NMRI mice were orally dosed with 500 mg/kg 6-Amino-*m*-Cresol in polyethylene glycol 400 (Völkner and Heidemann 1991). Three negative and one positive control (cyclophosphamide) were dosed orally once at 10 ml/kg. Bone marrow smears for the treated groups and negative control were prepared 24, 48, and 72 h post treatment. Bone marrow smears for the positive control were prepared 24 h post treatment. 6-Amino-*m*-Cresol did not induce micronuclei.

Groups of five male and five female NMRI mice were dosed orally with 666 mg/kg 6-Amino-*m*-Cresol in carboxymethylcellulose in a third micronucleus test (Leimbeck and Grötsch 1991). One negative and one positive control (cyclophosphamide, 40 mg/kg) were used. Bone marrow smears were evaluated 24, 48, and 72 h post administration. Again, 6-Amino-*m*-Cresol did not induce micronuclei in bone marrow cells.

A chromosome aberration study was conducted using groups of five male and five female Chinese hamsters (King and



Harnasch 1991). The animals were dosed once orally with 3200 mg/kg 6-Amino-*m*-Cresol in 4% gum arabic, and slides were prepared 6, 24, and 48 h post treatment. One negative control group was dosed with 20 ml of 4% gum arabic per kg body weight and one positive control was dosed i.p. with 30 mg/kg cyclophosphamide. Preparations from the positive control group were made at 24 h. A cytotoxic effect was observed, which indicated a strongly decreased ratio of polychromatic and normochromatic erythrocytes in the bone marrow (55% reduction compared to control animals). 6-Amino-*m*-Cresol did not induce chromosome aberrations in Chinese hamster bone marrow cells.

A bromodeoxyuridine pellet was implanted subcutaneously into male CD rats, and 2 h later groups of five animals were given a single oral dose of 60, 192, or 600 mg/kg 6-Amino-*m*-Cresol hemisulfate in distilled water (McGregor 1985). A negative-control group was given vehicle and a positive-control group was dosed with 5 mg cyclophosphamide. The animals were injected with colchicine 20 h after implantation, and killed 2 h after injection. 6-Amino-*m*-Cresol hemisulfate did not cause sister chromatid exchanges (SCEs) in rat bone marrow chromosomes.

An unscheduled DNA synthesis assay was performed using groups of five male Wistar Hanlbm:WIST (SPF) rats (Fautz 1994). The animals were given a single oral dose of 6-Amino-*m*-Cresol in 0.5% aqueous carboxymethylcellulose at a volume of 10 ml/kg. For the 2 h treatment, a dose of 1500 mg/kg was given and for the 16 h treatment, doses of 150 and 1500 mg/kg were used. A negative control (carboxymethyl cellulose) and a positive control, 100 mg/kg 2-AAF, were used. One of the animals in the 1500-mg/kg dose group died within 16 h of treatment and the other animals in the group had signs of toxicity. Additionally, the hepatocyte viability of two animals out of the 1500-mg/kg group was decreased. 6-Amino-*m*-Cresol did not induce UDS.

#### *4-Amino-m-Cresol*

In another micronucleus test, groups of six male and six female NMRI mice were given a single oral dose of 100, 333, or 1000 mg/kg 4-Amino-*m*-Cresol in DMSO (Miltenburger and Völkner 1988). Vehicle was used as the negative control and cyclophosphamide was used as the positive control. Femoral bone marrow cells were prepared 24 h after dosing for all groups and 48 and 72 h after dosing for the high-dose and control groups. 4-Amino-*m*-Cresol did not induce micronuclei.

In a micronucleus test, CD-1 mice were dosed with 20, 100, or 500 mg/kg 4-Amino-*m*-Cresol (Holmstroem 1980). The mice were dosed during two separate studies 6 and 30 h before they were killed. The vehicle control was 0.5% carboxymethylcellulose. The positive control was cyclophosphamide, which induced a small but significant increase in micronucleus frequency. Body weights did not vary by more than 1 g during the study. 4-Amino-*m*-Cresol did not increase the frequency of micronuclei in polychromatic erythroblasts.

In an SCE assay, groups of  $\leq 25$  male Chinese hamsters were dosed orally with 100, 300, 1000, 1500, or 2000 mg/kg or i.p.

with 10, 30, 100, 300, or 400 mg/kg 4-Amino-*m*-Cresol hemisulfate in double distilled water (Bracher et al. 1984). Water was used as a negative control and 2-AAF was used as a positive control. Doses of 1500 and 2000 mg/kg p.o. and 400 mg/kg i.p. had cytotoxic effects, and a dose of 500 mg/kg i.p. was "partly lethal." 4-Amino-*m*-Cresol hemisulfate did not cause SCEs, regardless of administration.

A UDS assay was performed in which groups of five male Wistar rats were dosed with 4-Amino-*m*-Cresol in "aqua bidest" at a dose of 1000 mg/kg for the 4 h treatment and doses of 60 and 600 mg/kg for the 16 h treatment (Fautz and Völkner 1991). A negative control and a positive control (substances not specified) was used. 4-Amino-*m*-Cresol did not induce UDS.

Five male Wistar rats per group were dosed with 1000 mg/kg 4-Amino-*m*-Cresol and killed 4 hours post treatment and 60 and 600 mg/kg and killed 16 hours posttreatment (Fautz and Völkner 1991b). The negative-control group received DMSO/PEG 400 and the positive-control group received 2-AAF. The rats were killed at the designated times by liver perfusion. Three animals from each group were used in the UDS assay. Hepatocytes were cultured with  $^3\text{H}$ -radiolabeled thymidine ( $^3\text{HtdR}$ ) for 4 h. The hepatocytes were washed and incubated overnight prior to autoradiography. The nuclear and net grain counts of the treated groups were in the range of the corresponding controls, therefore a statistical evaluation was not performed. 4-Amino-*m*-Cresol did not induce DNA damage leading to repair synthesis in the hepatocytes of treated rats.

#### *5-Amino-4-Chloro-o-Cresol*

An in vivo micronucleus test for chromosome mutations was conducted using adult CFW1 mice (20–32 g). Seven male and seven female mice were used at each dose. The test substance was dissolved in water at doses of 50, 250, and 500 mg/kg of 5-Amino-4-Chloro-*o*-Cresol hydrochloride was administered once by gavage. Bone marrow extracted from the femurs was prepared 24, 48, and 72 h after dosing in the case of the highest dose group and at 24 h for the other two dose groups. Endoxan<sup>®</sup> was the positive control and the vehicle was the negative control. Analysis was done of 1000 polychromic erythrocytes per animal. No induced micronuclei were found at any dose. The investigators concluded that 5-Amino-4-Chloro-*o*-Cresol hydrochloride was not mutagenic in this assay (Henkel KGaA 1994).

#### *5-Amino-6-Chloro-o-Cresol Hydrochloride*

An in vivo micronucleus test for chromosome mutations was conducted using adult OF1 mice (28.7–37.8 g for males and 21.6–30.0 g for females). Five male and five female mice were used. The test substance was dissolved in water and administered once by gavage to a final dose of 1200 mg/kg of 5-Amino-6-Chloro-*o*-Cresol hydrochloride. Bone marrow extracted from the femurs was prepared 24, 48, and 72 h after dosing in the case of the highest dose group and at 24 h for the other two dose groups. Cyclophosphamide (10 mg/kg) was the positive control and the vehicle was the negative control. Analysis was done of

1000 polychromic erythrocytes per animal. The ratio of chromatic/polychromatic erythrocytes was slightly increased, suggesting some toxicity to the bone marrow, but the investigators concluded that 5-Amino-6-Chloro-*o*-Cresol hydrochloride was not mutagenic in this assay (Henkel KGaA 1994).

#### 4-Amino-2-Hydroxytoluene

In Ames tests, 4-amino-2-hydroxytoluene was not mutagenic using *S. typhimurium* strain TA1535 without and with metabolic activation; 4-amino-2-hydroxytoluene was not mutagenic in some studies using strains TA98 and TA100 without and with metabolic activation, but was mutagenic in one study towards strains TA98, TA97, and TA100 (Elder 1989). Negative results were obtained in a micronucleus assay and a dominant lethal study using 4-amino-2-hydroxytoluene. No significant effect on SCEs or increase in chromosomal aberrations was observed in human lymphocytes obtained from subjects that repeatedly dyed their hair with a formulation containing 4-amino-2-hydroxytoluene.

#### *p*-Aminophenol

Elder (1988) reported that *p*-Aminophenol was strongly mutagenic in an assay for SCEs (human peripheral blood lymphocytes,  $\leq 10^{-4}$  M), was mutagenic in a DNA synthesis inhibition assay (Epstein-Barr virus-transformed lymphoblastoid cells, 0.5 mM), three assays for DNA structural alterations (human lymphoblastoid cells, 0.05 to 0.5 mM; mouse bone marrow cells; plant cells), two erythrocyte micronucleus tests ( $\leq 2$  mmol/kg; 3%), and a sperm head abnormality test (200 to 400 mg/kg), was slightly mutagenic in an Ames assay without metabolic activation and one assay for SCEs, and was nonmutagenic in an Ames assay without and with metabolic activation ( $\leq 2$   $\mu$ mol/plate), an *Escherichia coli* genetic repair assay, two assays for SCEs (Chinese hamster bone marrow cells, 5 mg/kg; metaphase human fibroblasts, 5 to 50  $\mu$ M), one erythrocyte micronucleus test (0.5%), a thymidine kinase reversion assay (1% with metabolic activation), and a sperm head abnormality test (0.5 to 2.0 mmol/kg).

#### *m*-Aminophenol

Elder (1988) also reported that *m*-Aminophenol was mutagenic in an assay for DNA structural alterations (human lymphocytes); was slightly mutagenic in an assay for SCEs (human lymphocytes, 6.6  $\mu$ g/ml); and was nonmutagenic in an Ames assay ( $\leq 1$  mg/ml agar with metabolic activation), an *E. coli* genetic repair assay, a DNA synthesis inhibition assay (rat hepatocytes,  $\leq 500$  nmol/ml), an assay for DNA structural alterations (human lymphocytes, 6.6  $\mu$ g/ml), two SCE induction assays (Chinese hamster cells,  $0.5-2 \times 10^{-2}$  mM; Chinese hamster bone marrow cells, 5 mg/kg), two erythrocyte micronucleus tests (0.5–2 mmol/kg; 0.5%), a dominant lethal assay ( $\leq 1\%$ ), and a sperm head abnormality test (0.5 to 2 mmol/kg). Also, no significant effect on SCEs or increase in chromosomal aberrations was observed in human lymphocytes obtained from subjects that re-

peatedly dyed their hair with a formulation containing *p*- or *m*-aminophenol (Elder 1988)

#### *o*-Aminophenol

Elder (1988) reported that *o*-Aminophenol was mutagenic in one Ames assay (7 to 100  $\mu$ g/ml with metabolic activation), an *E. coli* genetic repair assay, three assays for SCE induction (human fibroblasts, 0.01 to 0.3 mM; Chinese hamster cells,  $0.5-2 \times 10^{-2}$  mM; human lymphocytes, 1.6 to 6.6  $\mu$ g/ml), an erythrocyte micronucleus test (0.5 to 2 mmol/kg), and a sperm head abnormality test (0.5 to 2 mmol/kg) and was nonmutagenic in two Ames assays (0.5 to 2.0  $\mu$ g/plate without and with metabolic activation; with metabolic activation), a DNA synthesis inhibition assay (rat hepatocytes,  $\leq 100$  nmol/ml), one SCE induction assay (Chinese hamsters, 5 mg/kg), and an assay for DNA structural alterations (implanted Ehrlich ascites tumor cells).

## CARCINOGENICITY

Published data on the carcinogenicity of the ingredients reviewed in this safety assessment were not found. Data from previous safety assessments of related ingredients are summarized.

#### *m*-Aminophenol, *o*-Aminophenol, and *p*-Aminophenol

The carcinogenic potential of an oxidative hair dye containing 0.5% and 1.5% *p*-amino-*o*-cresol and *p*-aminophenol, respectively, was determined using mice (Jacobs et al. 1984). A dose of 0.5 ml of the dye mixed with an equal volume of 6% hydrogen peroxide was applied to the skin of each mouse once weekly for 20 months. The oxidative dye was not carcinogenic.

The carcinogenic potential of hair dyes containing *m*-, *o*-, and/or *p*-aminophenol were determined using mice (Burnett et al. 1980). A dose of 0.05 ml of oxidative hair dyes containing 0.7% *m*-aminophenol and 1.0% *p*-aminophenol, 0.7% *m*-aminophenol, 0.3% *o*-aminophenol, or 1.0% *N*-methyl-*p*-aminophenol sulfate mixed with an equal volume of 6% hydrogen peroxide were applied once weekly for 21 months and 0.05 ml of semipermanent hair dyes containing 0.09% and 0.2% *m*-aminophenol and *p*-aminophenol, respectively, or 0.02%, 0.04%, and 0.05% *m*-aminophenol, *p*-aminophenol, and *N*-methyl-*p*-aminophenol sulfate, respectively, were applied once weekly for 23 month. The hair dyes were not carcinogenic, and toxicity was not observed.

Burnett and Goldenthal (1988) also conducted a study to determine the carcinogenic potential of oxidative hair dye formulations containing 0.7% *m*-aminophenol and 1.0% *p*-aminophenol, 0.7% *m*-aminophenol, 0.3% *o*-aminophenol, or 1.0% *N*-methyl-*p*-aminophenol sulfate using the F<sub>1a</sub> generation of rats from their reproduction study that was previously summarized. The formulations were mixed with equal volumes of 6% hydrogen peroxide and twice weekly a dose of 0.5 ml was applied topically to a shaved area of the back for approximately 2 years. Successive applications were made to adjacent areas to minimize dermal irritation.

The incidence of mammary gland adenomas was significantly increased for the female test animals as compared to the animals in one of three control groups; however, this value was not considered statistically different from the other two control groups. The incidence of pituitary adenomas significantly increased for female test animals as compared to all three control groups. The researchers noted that the "incidence of this tumor is known to be high and variable in untreated female Sprague-Dawley rats. The fact that no pituitary carcinomas occurred in this group suggests that the distribution of these tumors was not related to the experimental treatments." The oxidative hair dye formulations were not considered carcinogenic.

## CLINICAL ASSESSMENT OF SAFETY

### Irritation and Sensitization

Published data on the clinical irritation and sensitization potential of 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, 5-Amino-4-Chloro-*o*-Cresol, or 5-Amino-6-Chloro-*o*-Cresol were not found.

#### 4-Chloro-2-Aminophenol

Thirty-one factory workers were patch tested with 4-Chloro-2-Aminophenol, as well as with four other compounds (*p*-aminophenol, *p*-nitrophenol, *p*-dichloronitrobenzene, and 3'-chlorodiphenylamine-2-carboxylic acid) used or produced at the factory (Naniwa 1979). (4-Chloro-2-Aminophenol, *p*-aminophenol, and 3'-chlorodiphenylamine-2-carboxylic acid are amino derivatives of aromatic compounds and *p*-nitrophenol and *p*-dichloronitrobenzene are nitro derivatives of them.) Using adhesive plasters, 0.1%, 0.5%, and 1.0% 4-Chloro-2-Aminophenol (and the other four compounds) in petrolatum was applied to the back of each subject for 48 h. The tests sites were scored 20 min after removal of the patches. A challenge test was performed by dropping 0.1 ml of 0.1% dinitrochlorobenzene (DNCB) in acetone onto the flexural antibrachium of each person, and the reaction was evaluated 48 h after application. A group of five control subjects was tested in the same manner.

Of the 31 subjects tested, 7 had positive reactions to 0.1%, 0.5%, and 1.0% 4-Chloro-2-Aminophenol, 6 had positive reactions to 0.5% and 1.0%, 2 had positive reactions to 0.1% and 0.5%, 1 had a positive reaction to 1.0% only, and one had a positive reaction to 0.1% and 1.0%. Six of the seven subjects that reacted to all three concentrations of 4-Chloro-2-Aminophenol had been directly exposed to it on repeated occasions. Some cross-sensitization might have occurred between 4-Chloro-2-Aminophenol and *p*-aminophenol (four cases), *p*-nitrophenol (one case), *p*-dichloronitrobenzene (three cases), and 3'-chlorodiphenylamine-2-carboxylic acid (two cases). None of the test subjects had a cross-sensitization reaction with DNCB. None of the control subjects had a primary irritation reaction to any of the tested compounds.

#### 4-Amino-2-Hydroxytoluene

In modified Draize repeat-insult patch tests (RIPTs), two aqueous solutions containing 2.0% 4-amino-2-hydroxytoluene produced one (although not reconfirmed at challenge) and two significant cases of dermatitis using 23 and 31 subjects, respectively (Elder 1989). In two semioclusive (open) RIPTs with 3% *m*-aminophenol, slight irritation during induction and no sensitization reactions at challenge were observed in one study and some irritation and a low degree of sensitization in 2/99 subjects was observed in the other study.

## EPIDEMIOLOGY

Between 35% and 45% of American women dye their hair, often at monthly intervals, over a period of years (Cosmetic, Toiletory, and Fragrance Association [CTFA] 1993). This estimate is drawn from market research data on hair dye product use, generally from females aged 15 to 60.

Hair dyes may be broadly grouped into oxidative (permanent) and direct (semipermanent) hair dyes. The oxidative dyes consist of precursors mixed with developers to produce color, although direct hair dyes are a preformed color. The ingredients addressed in this safety assessment are oxidative hair dyes.

In 1993, an International Agency for Research on Cancer (IARC) working group evaluated 78 epidemiology literature citations and concluded that "personal use of hair colourants cannot be evaluated as to its carcinogenicity" and that "occupation as a hairdresser or barber entails exposures that are probably carcinogenic" (IARC 1993). The IARC report did not distinguish between personal use of oxidative/permanent versus direct hair dyes, or distinguish among the multiple chemical exposures in addition to hair dyes to which a hairdresser or barber might be exposed.

In 2003, an updated review of the available epidemiology literature was prepared (Helzlsouer, Rollison, and Pinney 2003). This review considered 83 literature citations available since the IARC review. The authors found that hair dye exposure assessment ranged from ever/never use to information on type, color, duration and frequency of use.

The authors found insufficient evidence to support a causal association between personal hair dye use and a variety of tumors and cancers. The review highlighted well-designed studies with an exposure assessment that included hair dye type, color, and frequency or duration of use, which found associations between personal hair dye use and development of bladder cancer, non-Hodgkin's lymphoma, and multiple myeloma. These findings, however, were not consistently observed across studies. The authors concluded that the available evidence is insufficient to conclude a causal association between personal hair dye use and bladder cancer, non-Hodgkin's lymphoma, and multiple myeloma. With respect to other cancers, including leukemia, breast cancer, or childhood cancers, and autoimmune disease or adverse developmental/reproductive effects, the

authors concluded that the evidence also did not demonstrate a causal association with hair dye use.

A case-control study (Gago-Dominguez et al. 2001, 2003), described in this 2003 review, did suggest a possible genetically susceptible subgroup, which detoxify arylamines to a lower degree than the general population. The study authors hypothesized that this subgroup may be at greater risk of bladder cancer from hair dye exposure. The review authors noted that these results were based on small sample sizes.

The 2003 review authors recommended the replication of studies to better understand the observed associations, but concluded that the available evidence is insufficient to conclude the association between personal hair dye use and the health outcomes discussed is causal.

In considering this information, the CIR Expert Panel agreed that the available epidemiology studies are insufficient to conclude there is a causal relationship between hair dye use and cancer and other end points described in the Helzlsouer, Rollison, and Pinney (2003) review.

The Panel stated that use of direct hair dyes, although not the focus in all investigations, appear to have little evidence of an association with adverse events as reported in epidemiological studies. However, direct hair dyes are a diverse group of chemicals and the determination of safety may hinge on other safety test data.

The Panel recognizes that hair dye epidemiological studies do not address the safety of individual hair dyes, but is concerned that studies have demonstrated an association between use of oxidative/permanent hair dyes and some cancer endpoints. The Panel, therefore, strongly supports the need to replicate these studies, along with further studies to examine the possibility of susceptible subpopulations. Additional studies examining bladder cancer, non-Hodgkin's lymphoma, and multiple myeloma and hair dye use are underway and it is the intent of the CIR Expert Panel to periodically review hair dye epidemiological studies and update this section.

## Occupational

### *4-Chloro-2-Aminophenol*

Blood samples were taken from 21 workers that handled 4-Chloro-2-Aminophenol (and other compounds) (Tomoda, Tomioka, and Minami 1989). Half-oxidized hemoglobins, such as  $(\alpha^{2+}\beta^{3+})_2$  and  $(\alpha^{3+}\beta^{2+})_2$ , and methemoglobin were significantly increased in circulating erythrocytes of some workers.

## Exposure Assessment

### *5-Amino-4-Chloro-o-Cresol*

Considering that 5-Amino-4-Chloro-*o*-Cresol hydrochloride is used in oxidative hair dye formulations up to a maximum concentration of 2%, Henkel KGaA (1994) assessed the risks that such exposure might pose. Dilution with an oxidant 1:1 reduces the available concentration to 1%. It was estimated that a maximum of 100 ml of this dyeing mixture would be applied monthly.

It was further noted that color development is completed within 30 min and the resulting oxidized hair dye is fixed at the hair cortex, with any excess rinsed off (80% to 90% of the dyeing mixture).

From the available percutaneous absorption data in rats (Henkel KGaA 1994), in which dilution with an oxidizer was done to produce a 1.85% hair dye solution and rinsing off after 30 min exposure was done, an intake of 5-Amino-4-Chloro-*o*-Cresol hydrochloride of 5.21  $\mu\text{g}/\text{cm}^2$  was determined. Assuming a scalp surface of 500  $\text{cm}^2$ , the total absorbed hair dye would be 2.6 mg. This quantity may be extrapolated to 2.8 mg if a hair dye solution at 2% were applied. Using this latter value and considering a 60-kg user, the dose is 47  $\mu\text{g}/\text{kg}$ . Comparing this dose with, for example, the 180-mg/kg dose reported to produce no observable effects in a 90-day oral toxicity study in rats, these investigators concluded a substantial safety factor was available for 5-Amino-4-Chloro-*o*-Cresol.

### *5-Amino-6-Chloro-o-Cresol*

Considering that 5-Amino-6-Chloro-*o*-Cresol hydrochloride is used in oxidative hair dye formulations up to a maximum concentration of 2%, Henkel KGaA (1996) assessed the risks that such exposure might pose. Dilution with an oxidant 1:1 reduces the available concentration to 1%. It was estimated that a maximum of 100 ml of this dyeing mixture would be applied monthly. It was further noted that color development is completed within 30 min and the resulting oxidized hair dye is fixed at the hair cortex, with any excess rinsed off (80 to 90% of the dyeing mixture).

From the available percutaneous absorption data in rats (Henkel KGaA 1996) in which dilution with an oxidizer was done to produce a 1.14% hair dye solution and rinsing off after 30 min exposure was done, only 0.116% of 5-Amino-6-Chloro-*o*-Cresol hydrochloride was absorbed. Assuming a scalp surface of 500  $\text{cm}^2$ , 100 ml of hair dye mixture applied, concentration of dye of 1.14%, and absorption of 0.116%, the total absorbed hair dye can be calculated to be only 8.87  $\mu\text{g}$ . This quantity may be extrapolated to 17.75  $\mu\text{g}$  if a hair dye solution at 2% were applied. Using this latter value and considering a 60-kg user, the dose is 0.3  $\mu\text{g}/\text{kg}$ . Comparing this dose with, for example, the 50-mg/kg dose that was reported to produce no observable effects in a 90-day oral toxicity study in rats, the investigators concluded that a substantial safety factor was available for 5-Amino-6-Chloro-*o*-Cresol.

## SUMMARY

6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, 5-Amino-4-Chloro-*o*-Cresol, 5-Amino-6-Chloro-*o*-Cresol, and 4-Chloro-2-Aminophenol function as hair colorants. 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol are identified as oxidative hair dyes, that is, they are combined with an oxidizing agent before being applied to the hair. Information is not available to determine if 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, and 5-Amino-4-Chloro-*o*-Cresol

are used only in oxidative hair dyes or have application as nonoxidative (commonly referred to as semipermanent) hair dyes.

In 1998, frequency of use data submitted by FDA indicated that 6-Amino-*m*-Cresol was used in two hair dye formulations. More recent data available from the industry indicate that 6-Amino-*m*-Cresol was used at 2.4%, 6-Amino-*o*-Cresol was used at 0.7%, and 4-Amino-*m*-Cresol was used at 0.3% in 1999. Recent data from industry also reports that 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol were used at a maximum concentration of 2% in oxidizing hair dyes, which is effectively reduced to 1% with the addition of oxidizing agents.

5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol do not absorb significant UV radiation in the UVB region and none in the UVA region, although 4-Amino-*m*-Cresol had a symmetrical UV absorption peak at 300 nm. Both 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol produce virtually a single peak in HPLC and no small peaks were identified as *m*-cresol. 4-Amino-*m*-Cresol did not contain *m*-cresol when analyzed using HPLC.

Percutaneous penetration of 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol alone was significant, but when combined with oxidative developer, the absorption was extremely low. Both of these dyes are excreted rapidly via the urine.

The hair dyes containing 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, 5-Amino-4-Chloro-*o*-Cresol, 5-Amino-6-Chloro-*o*-Cresol, and 4-Chloro-2-Aminophenol, as coal tar hair dye products, are exempt from the principal adulteration provision and from the color additive provisions in sections 601 and 706 of the *Federal Food, Drug, and Cosmetic Act* of 1938 when the label bears a caution statement and patch test instructions for determining whether the product causes skin irritation. The following caution statement should be displayed conspicuously on the labels of coal tar hair dyes:

**Caution**—This product contains ingredients that may cause skin irritation on certain individuals, and a preliminary test according to accompanying directions should be made. This product must not be used for dyeing eyelashes or eyebrows; to do so may cause blindness.

Repeated exposure of animal skin to 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol failed to produce any cumulative irritation and single exposures up to 10% were not irritating to animal skin.

The response of leukocytes from guinea pigs using the LAI technique suggested that cross-sensitization might occur between 4-Chloro-2-Aminophenol and *p*-aminophenol. However, in testing using guinea pigs in which induction was with 4-Chloro-2-Aminophenol and the animals were challenged first with 4-Chloro-2-Aminophenol and then *p*-aminophenol, animals reacted to 4-Chloro-2-Aminophenol but not *p*-amino phenol. In clinical testing using factory workers, some cross-sensitization was observed between 4-Chloro-2-Aminophenol and *p*-aminophenol, as well as *p*-nitrophenol, *p*-dichloronitrobenzene, and 3'-chlorodiphenylamine-2-carboxylic acid. Guinea pig maximization tests of 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-

Chloro-*o*-Cresol combined with oxidizer demonstrate no sensitization.

Ocular exposure of animals to undiluted 5-Amino-4-Chloro-*o*-Cresol was irritating, but exposure to a 5% solution produced no irritation. Only minor irritation was observed with 5% 5-Amino-6-Chloro-*o*-Cresol.

Subchronic toxicity testing in animals using 5-Amino-4-Chloro-*o*-Cresol, 5-Amino-6-Chloro-*o*-Cresol, and 4-Amino-*m*-Cresol did not yield any adverse reactions.

6-Amino-*m*-Cresol and 4-Amino-*m*-Cresol were generally negative in *in vitro* and *in vivo* mutagenicity tests. The only exception was 6-Amino-*m*-Cresol was slightly mutagenic in an Ames assay towards *S. typhimurium* strain TA100 with and without metabolic activation. 4-Chloro-2-Aminophenol was weakly mutagenic in a preincubation assay. 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol were positive in some Ames test strains, but were negative in the HGPRT test in mammalian cells. 5-Amino-4-Chloro-*o*-Cresol did not induce chromosome aberrations in mammalian cells, but 5-Amino-6-Chloro-*o*-Cresol induced chromosome aberrations in mammalian lung cells but not in bone marrow erythrocytes. Neither of these hair dyes induced unscheduled DNA synthesis.

5-Amino-4-Chloro-*o*-Cresol, 5-Amino-6-Chloro-*o*-Cresol, 6-Amino-*m*-Cresol and 4-Amino-*m*-Cresol were not developmental toxins.

An exposure assessment that compared likely exposure levels of 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol with adverse effects data found that exposure would be several orders of magnitude below NOAEL levels.

## DISCUSSION

The Expert Panel recognizes that irritation and sensitization data on 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, and 4-Chloro-2-Aminophenol are absent from this report. However, the hair dyes containing the ingredients included in this report, as coal tar hair dye products, are exempt from the principal adulteration provision and from the color additive provisions in sections 601 and 706 of the *Federal Food, Drug, and Cosmetic Act* of 1938 when the label bears a caution statement and patch test instructions for determining whether the product causes skin irritation. The Expert Panel expects that following this procedure will identify individuals who would have an irritation/sensitization reaction and allow them to avoid significant exposures.

The information available on the use of 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol in hair dye formulations indicate that these ingredients are reacted with a developer and are not available for absorption into the skin of the scalp. These compounds, when tested alone, are moderate skin sensitizers, but when combined with the developer, these ingredients are not sensitizers in animal tests. In addition, no toxicologically significant impurities are present with these two ingredients. This information, coupled with the available animal test data,

support the safety of 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol for use in oxidative hair dyes.

Were 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol to have application in nonoxidative (semipermanent) hair dyes, there is concern about the potential for skin sensitization because these ingredients are moderate sensitizers. Because individuals would be pretested to determine if they would develop skin sensitization and because there is an absence of any significant systemic toxic effects in animal tests, the Panel believes that these two ingredients could be used safely in semipermanent hair dyes. Even though there is currently no use of these ingredients as semipermanent hair dyes, the Panel believes it useful to conclude that they could be used safely.

Although 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol appear to be used only in oxidative hair dyes, it is not clear whether 6-Amino-*m*-Cresol, 4-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, and 4-Chloro-2-Aminophenol are used solely in oxidative hair dyes where they would be reacted with a developer and would not be available for absorption into the skin. Therefore, the Expert Panel has considered each ingredient separately for use in oxidative hair dyes and in semi-permanent hair dyes.

Because 6-Amino-*m*-Cresol, 4-Amino-*m*-Cresol, 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol would be chemically reacted with a developer in oxidative hair dyes, and because the available information has consistently shown that such reactions make the starting ingredient unavailable for skin absorption, the CIR Expert Panel believes these ingredients would present no safety concerns if used in oxidative hair dyes.

The use of 6-Amino-*m*-Cresol, 4-Amino-*m*-Cresol, 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol in semipermanent hair dyes, however, could lead to skin absorption that would raise the need to assess systemic toxicity.

Such data are available for 6-Amino-*m*-Cresol and 4-Amino-*m*-Cresol, i.e., there are no toxic impurities, the ingredients themselves are not significantly toxic when absorbed into the skin, and there is no reproductive or developmental toxicity or genotoxicity associated with exposure to them. Therefore, it is possible to conclude that 6-Amino-*m*-Cresol and 4-Amino-*m*-Cresol can also be used safely in semi-permanent hair dyes.

Such data are not available to assess the safety of 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol for use in semipermanent hair dyes. In this situation, where the ingredients would not be chemically reacted before they are absorbed into the skin, available data do not provide all the information needed. The types of data required for each ingredient include

1. Physical and chemical properties for all ingredients, including the octanol/water partition coefficient
2. Impurities data, especially regarding the presence of *m*-cresol, other organic molecules, and heavy metals
3. Metabolism data, if the metabolism is not similar to that of 4-amino-2-hydroxytoluene and/or *p*-, *m*-, and *o*-aminophenol

(ingredients already reviewed by CIR), the following data may be needed:

- a. 28-Day dermal toxicity with histopathology
- b. Dermal reproductive toxicity data
- c. An *in vitro* genotoxicity study for 6-Amino-*o*-Cresol and one genotoxicity study in a mammalian system for 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol; if positive, a 2-year dermal carcinogenicity study using National Toxicology Program methods may be needed.

## CONCLUSION

The CIR Expert Panel concludes that the available data support the safety of 6-Amino-*m*-Cresol, 4-Amino-*m*-Cresol, 5-Amino-4-Chloro-*o*-Cresol and 5-Amino-6-Chloro-*o*-Cresol as used in oxidative and nonoxidative (semipermanent) hair dyes. The available data also support the safety of 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol for use in oxidative hair dyes, but are insufficient to support the safety of 6-Amino-*o*-Cresol and 4-Chloro-2-Aminophenol in nonoxidative (semipermanent) hair dyes.

## REFERENCES

- Adams, R. M., H. I. Maibach, W. E. Clendenning, et al. 1985. A five-year study of cosmetic reactions. *J. Am. Acad. Dermatol.* 13:1062–1069.
- Beilstein File of Organic Compounds. 1998. 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, and 4-Amino-*m*-Cresol entries. *Beilstein database*. Columbus, OH: STN International.
- Bootman, J. 1984a. 1-Hydroxy-2-amino-5-methylbenzene hemisulphate: Assessment of its capacity to induce genetic damage in *Saccharomyces cerevisiae*. Unpublished data submitted by the Cosmetic, Toiletry, and Fragrance Association (CTFA). 18 pages.<sup>2</sup>
- Bootman, J. 1984b. *In vitro* assessment of the clastogenic activity of 1-hydroxy-2-amino-5-methylbenzene hemisulphate in cultured human peripheral lymphocytes. Unpublished data submitted by CTFA.
- Bracher, M. et al. 1984. Sister chromatid exchange assay on 4-amino-*m*-cresol using Chinese hamster ovary cells. Unpublished data submitted by CTFA. 19 pages.<sup>2</sup>
- Burnett, C. M., and E. I. Goldenthal. 1988. Multigeneration reproduction and carcinogenicity studies in Sprague-Dawley rats exposed topically to oxidative hair-colouring formulations containing *p*-phenylenediamine and other aromatic amines. *Food Chem. Toxicol.* 26:467–474.
- Burnett, C., E. I. Goldenthal, S. B. Harris, et al. 1976. Teratology and percutaneous toxicity studies on hair dyes. *J. Toxicol. Environ. Health* 1:1027–1040.
- Burnett, C., M. M. Jacobs, A. Seppala, and P. Shubik. 1980. Evaluation of the toxicity and carcinogenicity of hair dyes. *J. Toxicol. Environ. Health* 6:247–257.
- Cosmetic, Toiletry, and Fragrance Association (CTFA). 1993. Personal correspondence from GN McEwen.<sup>2</sup>
- CTFA. 1999a. Chemical information on 6-Amino-*m*-Cresol. Unpublished data submitted by CTFA. 29 pages.<sup>2</sup>
- CTFA. 1999b. Chemical description on 4-Amino-*m*-Cresol. Dated Dec 3. Unpublished data submitted by CTFA. 28 pages.<sup>2</sup>
- CTFA. 1999c. Concentration of use data on 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, and 4-Amino-*m*-Cresol. Dated Mar 11. Unpublished data submitted by CTFA.
- CRC Handbook of Data on Organic Compounds. 1998. 6-Amino-*m*-Cresol and 4-Amino-*m*-Cresol entries. *HODOC database*. Columbus, OH: STN International.

- Eiermann, H. J., W. Larsen, H. I. Maibach, et al. 1982. Prospective study of cosmetic reactions: 1977–1980. *J. Am. Acad. Dermatol.* 6:909–917.
- Elder, R. L., ed. 1985. Final report on the safety assessment of *p*-Phenylenediamine. *J. Am. Coll. Toxicol.* 4:203–266.
- Elder, R. L., ed. 1988. Final report on the safety assessment of *p*-, *m*- and *o*-Aminophenol. *J. Am. Coll. Toxicol.* 7:279–333.
- Elder, R. L., ed. 1989. Final report on the safety assessment of 4-Amino-2-Hydroxytoluene. *J. Am. Coll. Toxicol.* 8:569–587.
- European Union. 1995. *Cosmetics Directive of the European Union*. Updated version—Incorporating all amendments until August 1, 1995. Dir. 76/768/EEC, Annex IV.
- Fautz, R. 1994. *In vivo/in vitro* unscheduled DNA synthesis in rat hepatocytes with A75 (6-Amino-*m*-Cresol). Unpublished data submitted by CTFA. 30 pages.<sup>2</sup>
- Fautz, R., and W. Völkner. 1991a. *In vivo/in vitro* unscheduled DNA synthesis in rat hepatocytes with 4-amino-3-methylphenol in aqua bidest. Unpublished data submitted by CTFA. 1 page.<sup>2</sup>
- Fautz, R., and W. Völkner. 1991b. *In vivo/in vitro* unscheduled DNA synthesis in rat hepatocytes with 4-amino-3-methylphenol in DMSO/PEG 400 (1 + 9). Unpublished data submitted by CTFA. 30 pages.<sup>2</sup>
- Food and Drug Administration (FDA). 1984. Cosmetic product formulation and frequency of use data. *FDA database*. Washington, DC: FDA.
- FDA. 1992. Modification in voluntary filing of cosmetic product ingredient and cosmetic raw composition statements. Final rule. *Federal Register* 57:3128–3130.
- FDA. 1998. Frequency of use of cosmetic ingredients. *FDA database*. Washington, DC: FDA.
- Forschungs GmbH. 1984a. Three month toxicity study with “Oxyrot,” “1-hydroxy-3-methyl-4-aminobenzol-sulfate” [4-Amino-*m*-Cresol]. Unpublished data submitted by Forschung GmbH. 34 pages.<sup>2</sup>
- Forschungs GmbH. 1984b. Teratogenicity study of “Oxyrot,” “1-hydroxy-3-methyl-4-aminobenzyl-sulfate” [4-Amino-*m*-Cresol]. Unpublished data submitted by Forschungs GmbH. 55 pages.<sup>2</sup>
- Forschungs GmbH. 1985. Four week toxicity study with “Oxygelb,” “1-hydroxy-2-amino-5-methyl-benzyl” [6-Amino-*m*-Cresol]. Unpublished data submitted by Forschungs GmbH. 36 pages.<sup>2</sup>
- Gago-Dominguez, M., J. E. Castela, J. M. Yuan, M. C., Yu, and R. K. Ross. 2001. Use of permanent hair dyes and bladder-cancer risk. *Int. J. Cancer* 91:575–579.
- Gago-Dominguez, M., et al. 2003. Permanent hair dyes and bladder cancer: Risk modification by cytochrome P4501A2 and *N*-acetyltransferases 1 and 2. *Carcinogenesis* 24:483–489.
- Goel, R. K., K. P. Kansal, and S. N. Sharma. 1979. Infrared absorption spectra of 6-Amino-*m*-Cresol. *Indian J. Pure Appl. Phys.* 17:778–780.
- Harvison, P. J., A. J. Forte, and S. D. Nelson. 1986. Comparative toxicities and analgesic activities of three monomethylated analogues of acetaminophen. *J. Med. Chem.* 29:1737–1743.
- Hazleton Laboratories. 1982. 1-Hydroxy-2-amino-5-methylbenzene: Oral (gavage) teratology study in the rat. Unpublished data submitted by Hazleton Laboratories. 33 pages.<sup>2</sup>
- Helzlsouer, K., D. Rollison, and S. Pinney. 2003. Association between hair dye use and health outcomes: Review of the literature published since 1992. Unpublished data submitted by Clairol, Inc.
- Henkel KGaA. 1994. 5-Amino-4-chloro-2-methylphenol-hydrochloride: COLIPA-No. A117. Submission I. Toxicological data for 5-Amino-4-chloro-2-methylphenol-hydrochloride submitted by Henkel KGaA. 31 pages.<sup>2</sup>
- Henkel KGaA. 1996. 2-Chloro-6-methyl-3-aminophenol hydrochloride: COLIPA - No. A094. Submission I. Toxicological data for 2-Chloro-6-methyl-3-aminophenol hydrochloride submitted by Henkel KGaA. 31 pages.<sup>2</sup>
- Holmstroem, M. 1980. Mutagenicity micronucleus assay of 5 compounds and LD<sub>50</sub> assay. Unpublished data submitted by CTFA. 33 pages.<sup>2</sup>
- Hong, S. K., M. A. Valentovic, D. K. Anestis, et al. 1996. Nephrotoxicity of 4-amino-2-chlorophenol and 2-amino-4-chlorophenol in the Fischer 344 rat. *Toxicology* 110:47–58.
- International Agency for Research on Cancer (IARC). 1993. *IARC monographs on the evaluation of carcinogenic risks to humans*, Vol. 57, 43–118, 159–165. Lyon, France: IARC.
- Jacobs, M. M., C. M. Burnett, A. J. Penicnak, et al. 1984. Evaluation of the toxicity and carcinogenicity of hair dyes in Swiss mice. *Drug Chem. Toxicol.* 7:573–586.
- James Robinson Ltd. 1998. Specifications and UV absorption data. Unpublished data submitted by CTFA. 6 pages.<sup>2</sup>
- King, M. T., and D. Harnasch. 1991. Mutagenicity study of LGH 110583/3 (6-Amino-*m*-Cresol) with the chromosome aberration test in bone marrow cells of Chinese hamsters *in vivo*. Unpublished data submitted by CTFA. 14 pages.<sup>2</sup>
- Leimbeck, R., and W. Grötsch. 1991. *In vivo* micronucleus test of LGH 110583/3 (6-Amino-*m*-Cresol) in mice. Unpublished data submitted by CTFA. 25 pages.<sup>2</sup>
- Martin, C. N. 1983. Study to determine the ability of 1-hydroxy-2-amino-5-methylbenzohemisulphate to induce mutations at the Na<sup>+</sup>/K<sup>+</sup> ATPase and hypoxanthine/guanine-phosphoribosyl transferase loci in mouse lymphoma L5178Y cells using a fluctuation assay. Unpublished data submitted by CTFA. 15 pages.<sup>2</sup>
- McGregor, D. B. 1985. 1-Hydroxy-2-amino-5-methyl-benzenehemisulphate: Sister chromatid exchange in rat bone marrow. Unpublished data submitted by CTFA. 33 pages.<sup>2</sup>
- Miltenburger, H. G. 1986. Unscheduled DNA synthesis (UDS) *in vitro* of 4-Amino-*m*-Cresol. Unpublished data submitted by CTFA. 19 pages.<sup>2</sup>
- Miltenburger, H. G., and W. Völkner. 1988. Micronucleus test in bone marrow cells of the mouse with Cos 291 (4-Amino-*m*-Cresol). Unpublished data submitted by CTFA. 27 pages.<sup>2</sup>
- Naniwa, S. 1979. Industrial contact dermatitis due to nitro and amino derivatives. 1st report: Mass examinations of a factory. *J. Dermatol.* 6:59–63.
- Naniwa, S. 1982. Industrial contact dermatitis due to nitro and amino derivatives. 2nd report: Experimental study of cross sensitivities of amino derivatives. *J. Dermatol.* 9:367–373.
- Nishi, H., K. Nakamura, H. Nakai, and T. Sato. 1997. Separation of enantiomers and isomers of amino compounds by capillary electrophoresis and high-performance liquid chromatography utilizing crown ethers. *J. Chromatogr.* 757:225–235.
- North American Contact Dermatitis Group. 1980. *Patch testing in allergic contact dermatitis*. Evanston, IL: American Academy of Dermatology.
- Noser, F. 1979a. Ames test with 6-Amino-*m*-Cresol. Unpublished data submitted by CTFA. 10 pages.<sup>2</sup>
- Noser, F. 1979b. Ames test with 4-Amino-*m*-Cresol. Unpublished data submitted by CTFA. 10 pages.<sup>2</sup>
- Pepe, R. C., J. A. Wenninger, and G. N. McEwen, Jr., eds. 2002. *International cosmetic ingredient dictionary and handbook*, 9th ed., Vol. 1. Washington, DC: CTFA.
- Regulated Chemicals Listing. 1998. 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, 5-Amino-6-Chloro-*o*-Cresol, and 4-Chloro-2-Aminophenol entries. *Chemlist database*. Columbus, OH: STN International.
- Son, O. S., D. W. Everett, and E. S. Fiala. 1980. Metabolism of *o*-methyl-[<sup>14</sup>C]toluidine in the F344 rat. *Xenobiotica* 10:457–468.
- Spectral Database Information System. 1998. 6-Amino-*m*-Cresol, 6-Amino-*o*-Cresol, 4-Amino-*m*-Cresol, and 4-Chloro-2-Aminophenol entries. *Specinfo database*. Columbus, OH: STN International.
- Tomoda, A., K. Tomioka, and M. Minami. 1989. Increased concentrations of haemoglobin X and Y in the erythrocytes of workers in a chemical plant in Japan. *Br. J. Indus. Med.* 46:502–504.
- Völkner, W., and A. Heidemann. 1991. Micronucleus assay in bone marrow cells of the mouse with LGH 110583/3 (6-Amino-*m*-Cresol). Unpublished data submitted by CTFA. 27 pages.<sup>2</sup>
- Zeiger, E., B. Anderson, S. Haworth, T. Lawlor, and K. Mortelmans. 1988. *Salmonella* mutagenicity tests: IV. Results from the testing of 300 chemicals. *Environ. Mol. Mutagen.* 11(Suppl 12):1–158.

**Concentration of Use by FDA Product Category\***

6-Amino-m-Cresol

5-Amino-4-Chloro-o-Cresol

6-Amino-o-Cresol

5-Amino-6-Chloro-o-Cresol

4-Amino-m-Cresol

4-Chloro-2-Aminophenol

<b>Ingredient</b>	<b>Product Category</b>	<b>Maximum Concentration of Use</b>
6-Amino-m-Cresol	Hair dyes and colors	0.69%
4-Amino-m- Cresol	Hair dyes and colors	0.08-0.14%
5-Amino-6-Chloro-o-Cresol	Hair dyes and colors	0.24%
5-Amino-6-Chloro-o-Cresol	Hair bleaches	0.007%

\*Ingredients included in the title of the table but not found in the table were included in the concentration of use survey, but no uses were reported.

Information collected in 2021  
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