

TEXT OF PROPOSED REGULATIONS FEBRUARY 2012

TITLE 27, CALIFORNIA CODE OF REGULATIONS

Section 25305 Powers and Duties

... (a)(6) Review the scientific basis for proposed No Significant Risk Levels (NSRLs) and other regulations proposed for Sections 25701 through 25721 (No Significant Risk Levels).

... (b)(6) Review the scientific basis for proposed Maximum Allowable Dose Levels (MADLs) and other regulations proposed for Sections 25801 through 25821 (No Observable Effect Levels).

Section 25701 General

... (e)Whenever the lead agency proposes to formally adopt a regulation pursuant to Sections 25701 through 25721, such as a level of exposure to a listed carcinogen that shall be deemed to pose no significant risk of cancer, the lead agency shall provide to each member of the Carcinogen Identification Committee notice of the proposed action, the proposed change to the regulation, and a copy of the initial statement of reasons supporting the proposal for their review and comment. The Committee shall be given at least 45 days to comment. Any such comment by members of the Carcinogen Identification Committee shall become a part of the formal rulemaking record. Nothing in this section shall be construed to require members of the Carcinogen Identification Committee to submit any comment. This procedure complies with the peer review requirements of section 57004 of the California Health and Safety Code.

Section 25705 Specific Regulatory Levels Posing No Significant Risk

~~(b)(2) Whenever the lead agency proposes to formally adopt, pursuant to this subsection, a level which shall be deemed to pose no significant risk of cancer, assuming daily exposure at that level, the lead agency shall provide to each member of the Carcinogen Identification Committee notice of the proposed action, a copy of the proposed level, and a copy of the initial statement of reasons supporting the proposal. The close of the public comment period for any such proposal shall be scheduled by the lead agency so as to permit the Carcinogen Identification Committee the opportunity to review such proposal and provide comment to the lead agency. Any such comment by the Carcinogen Identification Committee shall become a part of the formal rulemaking file. Nothing in this subsection shall be construed to prevent members of the Carcinogen Identification Committee from providing comments individually on any such proposal, or to require the Carcinogen Identification Committee to submit any comment~~

Section 25801 General

...(f) Whenever the lead agency proposes to formally adopt a regulation pursuant to Sections 25801 through 25821, such as a maximum allowable dose level, the lead agency shall provide each member of the Developmental and Reproductive Toxicant Identification Committee notice of the proposed action, the proposed change to the regulation, and a copy of the initial statement of reasons supporting the proposal for their review and comment. The Committee shall be given at least 45 days to comment. Any such comment by members of the Developmental and Reproductive Toxicant Identification Committee shall become a part of the formal rulemaking record. Nothing in this section shall be construed to require the members of the Developmental and Reproductive Toxicant Identification Committee to submit any comments. This procedure complies with the peer review requirements of section 57004 of the California Health and Safety Code.