Consumer Product Safety Commission

(c) Section 6(b)(5) does not apply to information independently obtained or prepared by the Commission staff.

Subpart H—Delegation of Authority to Information Group

§1101.71 Delegation of authority.

(a) Delegation. Pursuant to section 27(b)(9) of the CPSA 15 U.S.C. 2076(b)(9) the Commission delegates to the General Counsel or his or her senior staff designees, the authority to render all decisions under this part concerning the release of information subject to section 6(b) when firms have furnished section 6(b) comment except as provided in paragraph (b). The Commission also delegates to the Secretary of the Commission, or his or her senior staff designee, authority to make all decisions under this part concerning the release of information under section 6(b) when firms have failed to furnish section 6(b) comment or have consented to disclosure except as provided in paragraph (b) of this section. The General Counsel shall have authority to establish an Information Group composed of the General Counsel and the Secretary of the Commission or their designees who shall be senior staff members.

(b) *Findings not deleted*. The Commission does not delegate its authority—

(1) To find, pursuant to section 6(b)(1)and §1101.23(b) of this part, that the public health and safety requires less than 15 days advance notice of proposed disclosures of information.

(2) To find, pursuant to section 6(b)(2) and §1101.25(b) of this part, that the public health and safety requires less than five (5) days advance notice of its intent to disclose information claimed to be inaccurate;

(3) To decide whether it should take reasonable steps to publish a retraction of information in accordance with section 6(b)(7) and §1101.52 of this part.

(c) Final agency action; Commission decision. A decision of the General Counsel or the Secretary or their designees shall be a final agency decision and shall not be appealable as of right to the Commission. However, the General Counsel or the Secretary may in his or her discretion refer an issue to the Commission for decision.

[48 FR 57430, Dec. 29, 1983, as amended at 73 FR 72335, Nov. 28, 2008]

PART 1102—PUBLICLY AVAILABLE CONSUMER PRODUCT SAFETY IN-FORMATION DATABASE

Subpart A—Background and Definitions

Sec.

1102.2 Purpose.

- 1102.4 Scope.
- 1102.6 Definitions.

Subpart B—Content Requirements

- 1102.10 Reports of harm.
- 1102.12 Manufacturer comments.
- 1102.14 Recall notices.
- 1102.16 Additional information.

Subpart C—Procedural Requirements

- 1102.20 Transmission of reports of harm to the identified manufacturer or private labeler.
- 1102.24 Designation of confidential information.
- 1102.26 Determination of materially inaccurate information.
- 1102.28 Publication of reports of harm.
- 1102.30 Publication of manufacturer comments.

Subpart D—Notice and Disclosure Requirements

1102.42 Disclaimers.

1102.44 Applicability of sections 6(a) and (b) of the CPSA.

AUTHORITY: 15 U.S.C. 2051, 2051 note, 2052, 2055, 2055a, 2065, 2068, 2070, 2071, 2072, 2076, 2078, 2080, 2087.

SOURCE: 75 FR 76867, Dec. 9, 2010, unless otherwise noted.

Subpart A—Background and Definitions

§1102.2 Purpose.

This part sets forth the Commission's interpretation, policy, and procedures with regard to the establishment and maintenance of a Publicly Available Consumer Product Safety Information Database (also referred to as the "Database") on the safety of consumer products and other products or substances regulated by the Commission.