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to the public and when it retracts certain information it has released.

(1) Generally, section 6(b)(1) requires the Commission to provide manufacturers or private labelers with advance notice and opportunity to comment on information the Commission proposes to release, if the public can readily ascertain the identity of the firm from the information. Section 6(b)(1) also requires the Commission to take reasonable steps to assure that the information is accurate and that disclosure is fair in the circumstances and reasonably related to effectuating the purposes of the Acts administered by the Commission. Disclosure of information may not occur in fewer than 30 days after notice to the manufacturer or private labeler unless the Commission finds the public health and safety requires a lesser period of notice. Exceptions to these requirements are established in section 6(b)(4). Additional limitations on the disclosure of information reported to the Commission under section 15(b) of the CPSA are established in section 6(b)(5).

(2) Section 6(b)(2) requires the Commission to provide further notice to manufacturers or private labelers where the Commission proposes to disclose product-specific information the firms have claimed to be inaccurate.

(3) Section 6(b)(3) authorizes manufacturers and private labelers to bring lawsuits against the Commission to prevent disclosure of product-specific information after the firms have received the notice specified.

(c) *Internal clearance procedures.* Section 6(b)(6) requires the Commission to establish internal clearance procedures for Commission initiated disclosures of information that reflect on the safety of a consumer product or class of products, even if the information is not product specific. This rule does not address section 6(b)(6) because the Commission has internal clearance procedures in its directives system. (Directive 1450.2 “Clearance Procedures for Commission Staff to Use in Providing Information to the Public.” April 27, 1983.

§ 1101.2 Scope.

Section 6(b) and these rules apply to information concerning products sub-

ject to the CPSA (15 U.S.C. 2051–2085), and to the four other acts the Commission administers (transferred acts). These transferred acts are the Flammable Fabrics Act, 15 U.S.C. 1191–1204 (FFA); the Poison Prevention Packaging Act of 1970, 15 U.S.C. 1471–1476 (PPPA); the Federal Hazardous Substances Act, 15 U.S.C. 1261–1276 (FHSA); and the Refrigerator Safety Act, 15 U.S.C. 1211–1214 (RSA). See section 6(b)(1) of the CPSA, 15 U.S.C. 2055(d)(1).

Subpart B—Information Subject to Notice and Analysis Provisions of Section 6(b)(1)

§ 1101.11 General application of provisions of section 6(b)(1).

(a) *Information subject to section 6(b)(1).* To be subject to the notice and analysis provisions of section 6(b)(1), information must meet all the following criteria:

(1) The information must pertain to a specific product which is either designated or described in a manner which permits its identity to be ascertained readily by the public.

(2) The information must be obtained, generated or received by the Commission as an entity or by individual members, employees, agents, contractors or representatives of the Commission acting in their official capacities.

(3) The Commission or its members, employees, agents or representatives must propose to disclose the information to the public (see § 1101.12).

(4) The manner in which the product is designated or described in the information must permit the public to ascertain readily the identity of the manufacturer or private labeler. [See § 1101.13.]

(b) *Information not subject to section 6(b)(1).* The requirements of section 6(b)(1) do not apply to:

(1) Information described in the exclusions contained in section 6(b)(4) of the CPSA (see subpart E of this rule).

(2) Information the Commission is required by law to make publicly available. This information includes, for example, Commission notifications to foreign governments regarding certain products to be exported, as required by section 18(b) of the CPSA, 15 U.S.C.

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2068(b); section 14(d) of the FHSA, 15 U.S.C. 1273(d); and section 15(c) of the FFA, 15 U.S.C. 1202(c). (See the Commission's Export Policy Statement, 16 CFR part 1017.)

(3) Information required to be disclosed to the President and Congress pursuant to section 27(j) of the CPSA, 15 U.S.C. 2076(j).

(4) Press releases issued by firms.

(5) Information filed or presented in administrative proceedings or litigation to which the Commission is a party and which is not expressly subject to the section 6(b)(4) exceptions.

§ 1101.12 Commission must disclose information to the public.

Public. For the purposes of section 6(b)(1), the public includes any person except:

(a) Members, employees, agents, representatives and contractors of the Commission, in their official capacity.

(b) State officials who are commissioned officers under section 29(a)(2) of the CPSA, 15 U.S.C. 2078(a)(2), to the extent that the Commission furnishes them information necessary for them to perform their duties under that section. Such officials may not release to the public copies of such information unless the Commission has complied with section 6(b) or the information falls within an exception to section 6(b).

(c) Members of a Commission Chronic Hazard Advisory Panel established under section 28 of the CPSA (15 U.S.C. 2077). However, disclosures of information by such a Panel are subject to section 6(b).

(d) The persons or firms to whom the information to be disclosed pertains, or their legal representatives.

(e) The persons or firms who provided the information to the Commission, or their legal representatives.

(f) Other Federal agencies or state or local governments to whom accident and investigation reports are provided pursuant to section 29(e) of the CPSA (15 U.S.C. 2078(e)). However, as required by that section, employees of Federal agencies or state or local governments may not release to the public copies of any accident or investigation report made under the CPSA by an officer, employee or agent of the Commission

unless CPSC has complied with the applicable requirements of section 6(b).

(g) The Chairman or ranking minority member of a committee or subcommittee of Congress acting pursuant to committee business and having jurisdiction over the matter which is the subject of the information requested.

§ 1101.13 Public ability to ascertain readily identity of manufacturer or private labeler.

The advance notice and analysis provisions of section 6(b)(1) apply only when a reasonable person receiving the information in the form in which it is to be disclosed and lacking specialized expertise can readily ascertain from the information itself the identity of the manufacturer or private labeler of a particular product. The Commission will provide the advance notice and opportunity to comment if there is a question whether the public could readily ascertain the identity of a manufacturer or private labeler.

Subpart C—Procedure for Providing Notice and Opportunity To Comment Under Section 6(b)(1)

§ 1101.21 Form of notice and opportunity to comment.

(a) *Notice may be oral or written.* The Commission will generally provide to manufacturers or private labelers written notice and opportunity to comment on information subject to section 6(b)(1). However, when the Commission makes a public health and safety finding pursuant to section 6(b)(1) of the CPSA, the Commission may determine that it is necessary to provide the notice and opportunity to comment orally, either in person or by telephone.

(b) *Content of notice.* The Commission will provide the manufacturer or private labeler with:

(1) Either the actual text of the information to be disclosed or, if appropriate, a summary of the information.

(2) A general description of the manner in which the Commission will disclose the information, including any other relevant information the Commission intends to include with the disclosure. If the Commission advises that the form of disclosure will be by press