

Consumer Product Safety Commission

§ 1115.11

considered by staff in making the determination of whether and what type of corrective action may be required.

[71 FR 42030, July 25, 2006]

§ 1115.9 [Reserved]

§ 1115.10 Persons who must report and where to report.

(a) Every manufacturer (including importer), distributor, or retailer of a consumer product that has been distributed in commerce who obtains information that such consumer product contains a defect which could create a substantial risk of injury to the public shall immediately notify the Office of Compliance, Division of Corrective Actions, Consumer Product Safety Commission, Washington, DC 20207 (telephone: 301-504-0608), or such other persons as may be designated. Manufacturers (including importers), distributors, and retailers of consumer products subject to regulation by the Commission under provisions of the FFA, FHSA, PPPA, as well as consumer products subject to regulation under the CPSA and RSA, must comply with this requirement.

(b) Every manufacturer (including importer), distributor, or retailer of a consumer product that has been distributed in commerce who obtains information that such consumer product fails to comply with an applicable consumer product safety standard or ban issued under the CPSA shall immediately notify the Commission's Office of Compliance and Enforcement, Division of Corrective Actions or such other persons as may be designated. A subject firm need not report a failure to comply with a standard or regulation issued under the provisions of the RSA, FFA, FHSA, or PPPA unless it can be reasonably concluded that the failure to comply results in a defect which could create a substantial product hazard. (See paragraph (a) of this section.)

(c) Every manufacturer (including importer), distributor, and retailer of a consumer product that has been distributed in commerce who obtains information that such consumer product fails to comply with a voluntary consumer product safety standard upon which the Commission has relied under

section 9 of the CPSA, shall immediately notify the Commission's Office of Compliance and Enforcement, Division of Corrective Actions or such other persons as may be designated.

(d) Every manufacturer (including importer), distributor, and retailer of a consumer product that has been distributed in commerce who obtains information that such consumer product creates an unreasonable risk of serious injury or death shall immediately notify the Commission's Office of Compliance and Enforcement, Division of Corrective Actions or such other persons as may be designated. This obligation applies to manufacturers, distributors and retailers of consumer products subject to regulation by the Commission under the Flammable Fabrics Act, Federal Hazardous Substances Act, Poison Prevention Packaging Act, and Refrigerator Safety Act as well as products subject to regulation under the CPSA.

(e) A distributor or retailer of a consumer product (who is neither a manufacturer nor an importer of that product) is subject to the reporting requirements of section 15(b) of the CPSA but may satisfy them by following the procedure detailed in § 1115.13(b).

(f) A manufacturer (including an importer), distributor, or retailer need not inform the Commission under section 15(b) of the CPSA if that person has actual knowledge that the Commission has been adequately informed of the defect or failure to comply. (See section 15(b) of the CPSA.)

[43 FR 34998, Aug. 7, 1978, as amended at 57 FR 34229, Aug. 4, 1992; 62 FR 46667, Sept. 4, 1997]

§ 1115.11 Imputed knowledge.

(a) In evaluating whether or when a subject firm should have reported, the Commission will deem a subject firm to have obtained reportable information when the information has been received by an official or employee who may reasonably be expected to be capable of appreciating the significance of the information. (See § 1115.14(b).)

(b) In evaluating whether or when a subject firm should have reported, the Commission will deem a subject firm to know what a reasonable person acting in the circumstances in which the firm finds itself would know. Thus, the

§ 1115.12

16 CFR Ch. II (1–1–11 Edition)

subject firm shall be deemed to know what it would have known if it had exercised due care to ascertain the truth of complaints or other representations. This includes the knowledge a firm would have if it conducted a reasonably expeditious investigation in order to evaluate the reportability of a death or grievous bodily injury or other information. (See § 1115.14.)

§ 1115.12 Information which should be reported; evaluating substantial product hazard.

(a) *General.* Subject firms should not delay reporting in order to determine to a certainty the existence of a reportable noncompliance, defect or unreasonable risk. The obligation to report arises upon receipt of information from which one could reasonably conclude the existence of a reportable noncompliance, defect which could create a substantial product hazard, or unreasonable risk of serious injury or death. Thus, an obligation to report may arise when a subject firm received the first information regarding a potential hazard, noncompliance or risk. (See § 1115.14(c).) A subject firm in its report to the Commission need not admit, or may specifically deny, that the information it submits reasonably supports the conclusion that its consumer product is noncomplying, contains a defect which could create a substantial product hazard within the meaning of section 15(b) of the CPSA, or creates an unreasonable risk of serious injury or death. After receiving the report, the staff may conduct further investigation and will preliminarily determine whether the product reported upon presents a substantial product hazard. This determination can be based on information supplied by a subject firm or from any other source. If the matter is adjudicated, the Commission will ultimately make the decision as to substantial product hazard or will seek to have a court make the decision as to imminent product hazard.

(b) *Failure to comply.* A subject firm must report information indicating that a consumer product which it has distributed in commerce does not comply with an applicable consumer product safety standard or ban issued under the CPSA, or a voluntary consumer

product safety standard upon which the Commission has relied under section 9 of the CPSA.

(c) *Unreasonable risk of serious injury or death.* A subject firm must report when it obtains information indicating that a consumer product which it has distributed in commerce creates an unreasonable risk of serious injury or death.

(d) *Death or grievous bodily injury.* Information indicating that a noncompliance or a defect in a consumer product has caused, may have caused, or contributed to the causing, or could cause or contribute to the causing of a death or grievous bodily injury (e.g., mutilation, amputation/dismemberment, disfigurement, loss of important bodily functions, debilitating internal disorders, severe burns, severe electrical shocks, and injuries likely to require extended hospitalization) must be reported, unless the subject firm has investigated and determined that the information is not reportable.

(e) *Other information indicating a defect or noncompliance.* Even if there are no reports of a potential for or an actual death or grievous bodily injury, other information may indicate a reportable defect or noncompliance. In evaluating whether or when a subject firm should have reported, the Commission will deem a subject firm to know what a reasonable and prudent manufacturer (including an importer), distributor, or retailer would know. (See § 1115.11.)

(f) *Information which should be studied and evaluated.* Paragraphs (f)(1) through (7) of this section are examples of information which a subject firm should study and evaluate in order to determine whether it is obligated to report under section 15(b) of the CPSA. Such information may include information that a firm has obtained, or reasonably should have obtained in accordance with § 1115.11, about product use, experience, performance, design, or manufacture outside the United States that is relevant to products sold or distributed in the United States. All information should be evaluated to determine whether it suggests the existence of a noncompliance, a defect, or an unreasonable risk of serious injury or death: