CONSUMER PRODUCT SAFETY ACT

adopt a concurrent resolution, the matter after the resolving clause of which is as follow (with the blank spaces appropriately filled): “That the Congress disapproves the consumer product safety rule which was promulgated by the Consumer Product Safety Commission with respect to and which was transmitted to the Congress on and disapproves the rule for the following reasons: ”; or

(2) within the 60 calendar days of continuous session of the Congress which occur after the date of the promulgation of such rule, one House of the Congress adopts such concurrent resolution and transmits such resolution to the other House and such resolution is not disapproved by such other House within the 30 calendar days of continuous session of the Congress which occur after the date of such transmittal.

(c) Congressional inaction on, or rejection of, a concurrent resolution of disapproval under this section shall not be construed as an expression of approval of the rule involved, and shall not be construed to create any presumption of validity with respect to such rule.

(d) For purposes of this section—

(1) continuity of session is broken only by an adjournment of the Congress sine die; and

(2) the days on which either House is not in session because of an adjournment of more than 3 days to a day certain are excluded in the computation of the periods of continuous session of the Congress specified in subsection (b).

INFORMATION REPORTING


(a) If a particular model of a consumer product is the subject of at least 3 civil actions that have been filed in Federal or State court for death or grievous bodily injury which in each of the 24-month periods defined in subsection (b) result in either a final settlement involving the manufacturer or a court judgment in favor of the plaintiff, the manufacturer of such product shall, in accordance with subsection (c), report to the Commission each such civil action within 30 days after the final settlement or court judgment in the third of such civil actions, and, within 30 days after any subsequent settlement or judgment in that 24-month period, any other such action.

Unofficial compilation for convenience only.
(b) The 24-month periods referred to in subsection (a) are the 24-month period commencing on January 1, 1991, and subsequent 24-month periods beginning on January 1 of the calendar year that is two years following the beginning of the previous 24-month period.

(c)(1) The information required by subsection (a) to be reported to the Commission, with respect to each civil action described in subsection (a), shall include and in addition to any voluntary information provided under paragraph (2) shall be limited to the following:

(A) The name and address of the manufacturer.

(B) The model and model number or designation of the consumer product subject to the civil action.

(C) A statement as to whether the civil action alleged death or grievous bodily injury, and in the case of an allegation of grievous bodily injury, a statement of the category of such injury.

(D) A statement as to whether the civil action resulted in a final settlement or a judgment in favor of the plaintiff.

(E) In the case of a judgment in favor of the plaintiff, the number assigned, the name of the civil action, the number assigned the civil action, and the court in which the civil action was filed.

(2) A manufacturer furnishing the report required by paragraph (1) may include (A) a statement as to whether any judgment in favor of the plaintiff is under appeal or is expected to be appealed or (B) any other information which the manufacturer chooses to provide. A manufacturer reporting to the Commission under subsection (a) need not admit or may specifically deny that the information it submits reasonably supports the conclusion that its consumer product caused a death or grievous bodily injury.

(3) No statement of the amount paid by the manufacturer in a final settlement shall be required a part of the report furnished under subsection (a), nor shall such a statement of settlement amount be required under any other section of this Act.

(d) The reporting of a civil action described in subsection (a) by a manufacturer shall not constitute an admission of—

(1) an unreasonable risk of injury,

(2) a defect in the consumer product which was the subject of such action,

(3) a substantial product hazard,

(4) an imminent hazard,

(5) any other admission of liability under any statute or under any common law.

Unofficial compilation for convenience only.
(e) For purposes of this section:

(1) A grievous bodily injury includes any of the following categories of injury: mutilation, amputation, dismemberment, disfigurement, loss of important bodily functions, debilitating internal disorder, severe burn, severe electric shock, and injuries likely to require extended hospitalization.

(2) For purposes of this section, a particular model of a consumer product is one that is distinctive in functional design, construction, warnings or instructions related to safety, function, user population, or other characteristics which could affect the product’s safety related performance.