

§ 720.1

40 CFR Ch. I (7-1-03 Edition)

720.3 Definitions.

Subpart B—Applicability

720.22 Persons who must report.
720.25 Determining whether a chemical substance is on the Inventory.
720.30 Chemicals not subject to notification requirements.
720.36 Exemption for research and development.
720.38 Exemptions for test marketing.

Subpart C—Notice Form

720.40 General.
720.45 Information that must be included in the notice form.
720.50 Submission of test data and other data concerning the health and environmental effects of a substance.
720.57 Imports.

Subpart D—Disposition of Notices

720.60 General.
720.62 Notice that notification is not required.
720.65 Acknowledgment of receipt of a notice; errors in the notice; incomplete submissions; false and misleading statements.
720.70 Notice in the FEDERAL REGISTER.
720.75 Notice review period.
720.78 Recordkeeping.

Subpart E—Confidentiality and Public Access to Information

720.80 General provisions.
720.85 Chemical identity.
720.87 Categories or proposed categories of uses of a new chemical substance.
720.90 Data from health and safety studies.
720.95 Public file.

Subpart F—Commencement of Manufacture or Import

720.102 Notice of commencement of manufacture or import.

Subpart G—Compliance and Inspections

720.120 Compliance.
720.122 Inspections.

AUTHORITY: 15 U.S.C. 2604, 2607, and 2613.

SOURCE: 48 FR 21742, May 13, 1983, unless otherwise noted.

Subpart A—General Provisions

§ 720.1 Scope.

This part establishes procedures for the reporting of new chemical substances by manufacturers and import-

ers under section 5 of the Toxic Substances Control Act, 15 U.S.C. 2604. This part applies to microorganisms only to the extent provided by part 725 of this chapter. The rule defines the persons and chemical substances subject to the reporting requirements, prescribes the contents of section 5 notices, and establishes procedures for submitting notices. The rule also establishes EPA policy regarding claims of confidentiality for, and public disclosure of, various categories of information submitted in connection with section 5 notices.

[48 FR 21742, May 13, 1983, as amended at 58 FR 34204, June 23, 1993; 62 FR 17932, April 11, 1997]

§ 720.3 Definitions.

(a)(1) For the purposes of this part, the terms *cosmetic*, *device*, *drug*, *food*, and *food additive* have the meanings contained in the Federal Food, Drug, and Cosmetic Act, 21 U.S.C. 321 *et seq.*, and the regulations issued under it. In addition, the term “food” includes poultry and poultry products, as defined in the Poultry Products Inspection Act, 21 U.S.C. 453 *et seq.*; meats and meat food products, as defined in the Federal Meat Inspection Act, 21 U.S.C. 60 *et seq.*; and eggs and egg products, as defined in the Egg Products Inspection Act, 21 U.S.C. 1033 *et seq.*

(2) The term *pesticide* has the meaning contained in the Federal Insecticide, Fungicide, and Rodenticide Act, 7 U.S.C. 136 *et seq.* and the regulations issued under it.

(3) The terms *byproduct material*, *source material*, and *special nuclear material* have the meanings contained in the Atomic Energy Act of 1954, 42 U.S.C. 2014 *et seq.* and the regulations issued under it.

(b) *Act* means the Toxic Substances Control Act, 15 U.S.C. 2601 *et seq.*

(c) *Article* means a manufactured item (1) which is formed to a specific shape or design during manufacture, (2) which has end use function(s) dependent in whole or in part upon its shape or design during end use, and (3) which has either no change of chemical composition during its end use or only those changes of composition which have no commercial purpose separate from that of the article and that may