

The information described in §201.66(c) of this chapter shall be printed in accordance with the following specifications.

(1) The labeling shall meet the requirements of §201.66(c) of this chapter except that the information in §201.66(c)(3) of this chapter may be omitted, and the information in §201.66(c)(5) and (c)(6) of this chapter may be presented as follows:

(i) The words “Contains salicylate.” may be omitted from the warning in §335.50(c)(2)(i)(B).

(ii) The subheading “When using this product” in §335.50(c)(2)(iv) may be omitted.

(iii) The words “continue to” may be omitted from the directions in §335.50(d)(3).

(2) The labeling shall be printed in accordance with the requirements of §201.66(d) of this chapter except that any requirements related to §201.66(c)(3) of this chapter and the bullet in the warning in §335.50(c)(1)(i) may be omitted.

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PART 336—ANTIEMETIC DRUG PRODUCTS FOR OVER-THE-COUNTER HUMAN USE

Subpart A—General Provisions

Sec.

336.1 Scope.

336.3 Definition.

Subpart B—Active Ingredients

336.10 Antiemetic active ingredients.

Subpart C—Labeling

336.50 Labeling of antiemetic drug products.

336.80 Professional labeling.

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SOURCE: 52 FR 15892, Apr. 30, 1987, unless otherwise noted.

Subpart A—General Provisions

§336.1 Scope.

(a) An over-the-counter antiemetic drug product in a form suitable for oral administration is generally recognized as safe and effective and is not mis-

branded if it meets each of the conditions in this part and each of the general conditions established in §330.1.

(b) References in this part to regulatory sections of the Code of Federal Regulations are to chapter I of title 21 unless otherwise noted.

§336.3 Definition.

As used in this part:

Antiemetic. An agent that prevents or treats nausea and vomiting.

Subpart B—Active Ingredients

§336.10 Antiemetic active ingredients.

The active ingredient of the product consists of any of the following when used within the dosage limits established for each ingredient in §336.50(d):

- (a) Cyclizine hydrochloride.
- (b) Dimenhydrinate.
- (c) Diphenhydramine hydrochloride.
- (d) Meclizine hydrochloride.

Subpart C—Labeling

§336.50 Labeling of antiemetic drug products.

(a) *Statement of identity.* The labeling of the product contains the established name of the drug, if any, and identifies the product as an “antiemetic.”

(b) *Indications.* The labeling of the product states the following under the heading “Indications,” “For the prevention and treatment of the nausea, vomiting, or dizziness associated with motion sickness.” Other truthful and nonmisleading statements, describing only the indications for use that have been established and listed in this paragraph (b), may also be used, as provided in §330.1(c)(2), subject to the provisions of section 502 of the act relating to misbranding and the prohibition in section 301(d) of the act against the introduction or delivery for introduction into interstate commerce of unapproved new drugs in violation of section 505(a) of the act.

(c) *Warnings.* The labeling of the product contains the following warnings under the heading “Warnings:”

(1) *For products containing any ingredient identified in §336.10—(i) When labeled for use in adults and for those products that can be and are labeled for use in children under 12 years of age.* “Do not