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§ 346.52 - Labeling of permitted combinations of anorectal active ingredients.

(a) Statement of identity. For a combination drug product that has an established name, the labeling of the product states the established name of the combination drug product, followed by the statement of identity established in §346.50(a). For a combination drug product that does not have an established name, the labeling of the product states the statement of identity established in §346.50(a).

(b) Indications. The labeling of the product states, under the heading “Indications,” the indication(s) for each ingredient in the combination, as established in the indications sections of this subpart.

(c) Warnings. The labeling of the product states, under the heading “Warnings,” the warning(s) for each ingredient in the combination, as established in the warnings sections of this subpart.

(d) Directions. The labeling of the product states, under the heading “Directions,” directions that conform to the directions established for each ingredient in the directions sections of this subpart. When the time intervals or age limitations for administration of the individual ingredients differ, the directions for the combination product may not exceed any maximum dosage limits established for the individual ingredients in the applicable OTC drug monograph.

PART 347—SKIN PROTECTANT DRUG PRODUCTS FOR OVER-THE-COUNTER HUMAN USE

Subpart A—General Provisions

§ 347.1 Scope.

(a) An over-the-counter skin protectant drug product in a form suitable for topical administration is generally recognized as safe and effective and is not misbranded if it meets each condition in this part and each general condition established in §330.1 of this chapter.

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

Subpart B—Active Ingredients

§ 347.10 Skin protectant active ingredients.

§ 347.12 Astringent active ingredients.

§ 347.20 Permitted combinations of active ingredients.

Subpart C—Labeling

§ 347.50 Labeling of skin protectant drug products.

§ 347.52 Labeling of astringent drug products.

§ 347.60 Labeling of permitted combinations of active ingredients.


SOURCE: 58 FR 54462, Oct. 21, 1993, unless otherwise noted.

Subpart A—General Provisions

§ 347.1 Scope.

(a) An over-the-counter skin protectant drug product in a form suitable for topical administration is generally recognized as safe and effective and is not misbranded if it meets each condition in this part and each general condition established in §330.1 of this chapter.

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