(ii) In addition to the information identified in paragraph (b)(2)(i) of this section, the labeling of the product may contain the following statement: “Clears up most athlete’s foot infection and with daily use helps keep it from coming back.”

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

(1) For products containing any ingredient identified in §330.210. (i) “Do not use on children under 2 years of age unless directed by a doctor.”

(2) For products labeled according to paragraph (b)(1) of this section for the treatment of athlete’s foot and ringworm. “If irritation occurs or if there is no improvement within 4 weeks, discontinue use and consult a doctor.”

(3) For products labeled according to paragraph (b)(1) of this section for the treatment of jock itch. “If irritation occurs or if there is no improvement within 2 weeks, discontinue use and consult a doctor.”

(4) For products labeled according to paragraph (b)(2) of this section for the prevention of athlete’s foot. “If irritation occurs, discontinue use and consult a doctor.”

(5) For products containing the ingredient identified in §333.210(a) labeled according to paragraph (b)(1) of this section. The following statements must appear in boldface type as the first warnings under the “Warnings” heading. (i) “Do not use on children under 2 years of age.” (This warning is to be used in place of the warning in paragraph (c)(1)(i) of this section.)

(2) For products labeled according to paragraph (b)(2) of this section for the prevention of athlete’s foot. “To prevent athlete’s foot,” (select one of the following: “clean” or “wash”) “the feet and dry thoroughly. Apply” (the word “spray” may be used to replace the word “apply” for aerosol products) “a thin layer of the product to the feet once or twice daily (morning and/or night). Supervise children in the use of this product. Pay special attention to spaces between the toes; wear well-fitting, ventilated shoes, and change shoes and socks at least once daily.”

(e) The word “physician” may be substituted for the word “doctor” in any of the labeling statements in this section.

[58 FR 49898, Sept. 23, 1993, as amended at 65 FR 52305, Aug. 29, 2000]

§ 333.280 Professional labeling.

The labeling provided to health professionals (but not to the general public) may contain the following additional indication:

(a) For products containing haloprogin or miconazole nitrate identified in §333.210 (a) and (c). “For the treatment of superficial skin infections caused by yeast (Candida albicans).”

(b) [Reserved]

Subpart D—Topical Acne Drug Products

SOURCE: 56 FR 41019, Aug. 16, 1991, unless otherwise noted.

§ 333.301 Scope.

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