

and this part require that the rates of assessment fixed for a particular marketing year shall be applicable to all assessable walnuts from the beginning of such year; and (2) the 1974-75 marketing year began August 1, 1974, and the rates of assessment hereinafter fixed will automatically apply to all such assessable walnuts beginning with that date.

The expenses and assessment rates are as follows:

§ 984.326 Expenses of the Walnut Marketing Board and rates of assessment for the 1974-75 marketing year.

(a) *Expenses.* Expenses in the amount of \$200,000 are reasonable and likely to be incurred by the Walnut Marketing Board during the marketing year beginning August 1, 1974, for its maintenance and functioning, and for such purposes as the Secretary may, pursuant to the provisions of this part, determine to be appropriate.

(b) *Rates of assessment.* The rates of assessment for said marketing year, payable by each handler in accordance with § 984.69, are fixed at 0.125 cent per pound for merchantable inshell walnuts and 0.30 cent per pound for merchantable shelled walnuts.

(Secs. 1-19, 48 Stat. 31, as amended; 7 U.S.C. 601-674)

Dated: October 24, 1974.

CHARLES R. BRADER,
Deputy Director, Fruit and
Vegetable Division.

[FR Doc.74-25219 Filed 10-29-74; 8:45 am]

Title 21—Food and Drugs

CHAPTER I—FOOD AND DRUG ADMINISTRATION, DEPARTMENT OF HEALTH, EDUCATION, AND WELFARE

SUBCHAPTER A—GENERAL

PART 8—COLOR ADDITIVES

Chromium-Cobalt-Aluminum Oxide Pigment; Confirmation of Effective Date

An order was published in the FEDERAL REGISTER of August 7, 1974 (39 FR 28421), listing chromium-cobalt-aluminum oxide pigment for safe use in coloring sutures (21 CFR 8.6010) and deleting it from the list of color additives in § 8.501(f).

Pursuant to provisions of the Federal Food, Drug, and Cosmetic Act (sec. 706 (b), (c), and (d), 74 Stat. 399-403; 21 U.S.C. 376 (b), (c), and (d)) and under authority delegated to the Commissioner of Food and Drugs (21 CFR 2.120), notice is given that no objections or requests for hearing were filed in response to the subject order. Accordingly, the amendments promulgated thereby will become effective October 7, 1974.

Dated: October 22, 1974.

SAM D. FINE,
Associate Commissioner
for Compliance.

[FR Doc.74-25158 Filed 10-29-74; 8:45 am]

PART 8—COLOR ADDITIVES

FD&C Red No. 40; Confirmation of Effective Date

An order was published in the FEDERAL REGISTER of August 6, 1974 (39 FR 28278), listing FD&C Red No. 40 as a safe color additive for use in dentifrices which are cosmetics subject to certification (21 CFR 8.7201).

Pursuant to provisions of the Federal Food, Drug, and Cosmetic Act (sec. 706 (b), (c), and (d), 74 Stat. 399-403; 21 U.S.C. 376 (b), (c), and (d)) and under authority delegated to the Commissioner of Food and Drugs (21 CFR 2.120), notice is given that no objections or requests for hearing were filed in response to the subject order. Accordingly, the amendment promulgated thereby will become effective October 7, 1974.

Dated: October 22, 1974.

SAM D. FINE,
Associate Commissioner
for Compliance.

[FR Doc.74-25155 Filed 10-29-74; 8:45 am]

SUBCHAPTER B—FOOD AND FOOD PRODUCTS

PART 19—CHEESES, PROCESSED CHEESES, CHEESE FOODS, CHEESE SPREADS, AND RELATED FOODS

Label Requirements for Cottage Cheese Products

In a notice published in the FEDERAL REGISTER of June 3, 1974 (39 FR 19501), the Commissioner proposed that Part 19 be amended by deleting the undesignated paragraph immediately following §§ 19.525(e) (3) and 19.530(e) (4) (21 CFR 19.525(e) (3) and 19.530(e) (4)).

The purpose of the proposal was to remove an inconsistency which exists between the standards for cottage cheese dry curd and cottage cheese under §§ 19.525 and 19.530 and the requirements of § 1.8d *Food labeling; information panel*, published in the FEDERAL REGISTER of March 14, 1973 (38 FR 6950). Since there is already in §§ 19.525(e) and 19.530(e) a cross-reference to the Part 1 requirements, deletion of the undesignated paragraph immediately following these paragraphs as proposed will remove the inconsistency.

No comments were received which responded to the issue raised in the proposal.

The only comment received, which was from a trade association, recommended that the term "dried milk" as used in § 19.530(e) (1) be revised to read "dry whole milk" to avoid confusion and to reaffirm the distinction between "dry whole milk" and "nonfat dry milk."

The subject raised by the one comment was not in response to the proposal; however it will be considered in response to the proposed revision of § 1.10 *Food; labeling; designation of ingredients* published in the FEDERAL REGISTER of June 14, 1974 (39 FR 20888).

In the absence of any adverse comments to the proposal and because the

aforementioned undesignated paragraphs are not consistent in every respect with the requirements of § 1.8d(c), the Commissioner concludes that the paragraphs should be deleted.

Therefore, pursuant to provisions of the Federal Food, Drug, and Cosmetic Act (secs. 401, 701, 52 Stat. 1046, 1055-1056, as amended by 70 Stat. 919 and 73 Stat. 948; 21 U.S.C. 341, 371) and under authority delegated to the Commissioner (21 CFR 2.120): It is ordered, That §§ 19.525(e) and 19.530(e) be revised to read as follows:

§ 19.525 Cottage cheese dry curd; identity; label statement of optional ingredients.

(e) The common or usual name of each of the ingredients used in the food shall be declared on the label as required by the applicable sections of Part 1 of this chapter, except that:

(1) Concentrated skim milk, nonfat dry milk, and reconstituted skim milk prepared by addition of water to concentrated skim milk or nonfat dry milk may be declared as "skim milk."

(2) Bacterial cultures may be declared by the word "cultured" followed by the name of the substrate, e.g., "made from cultured skim milk."

(3) Milk-clotting enzymes may be declared by the word "enzymes."

§ 19.530 Cottage cheese; identity; label statement of optional ingredients.

(e) The common or usual name of each of the ingredients used in the food shall be declared on the label as required by the applicable sections of Part 1 of this chapter, except that:

(1) Concentrated milk, dried milk, and reconstituted milk prepared by addition of water to concentrated milk or dried milk may be declared as "milk."

(2) Concentrated skim milk, nonfat dry milk, and reconstituted skim milk prepared by addition of water to concentrated skim milk or nonfat dry milk may be declared as "skim milk."

(3) Bacterial cultures may be declared by the word "cultured" followed by the name of the substrate, e.g., "made from cultured skim milk."

(4) Milk-clotting enzymes may be declared by the word "enzymes."

Any person who will be adversely affected by the foregoing order may at any time on or before November 20, 1974, file with the Hearing Clerk, Food and Drug Administration, Rm. 4-65, 5600 Fishers Lane, Rockville, MD 20852, written objections thereto. Objections shall show wherein the person filing will be adversely affected by the order, specify with particularity the provisions of the order deemed objectionable, and state the grounds for the objections. If a hearing is requested, the objections shall state the issues for the hearing, shall be supported by grounds factually and legally sufficient to justify the relief sought, and shall include a detailed description and