Gainesville, Fla.—Gainesville Municipal Airport; NDB Runway 28, Original; Established.
Jackson, Miss.—Hawkins Field; NDB-A, Amdt. 11; Revised.
Jackson, Miss.—Allen C. Thompson Field; NDB Runway 15L, Amdt. 9; Revised.
Macon, Ga.—Lewis B. Wilson Airport; NDB Runway 5, Amdt. 15; Revised.
Pensacola, Fla.—Pensacola Municipal (Hagler) Airport; NDB Runway 16, Amdt. 15; Revised.
Pensacola, Fla.—Pensacola Municipal (Hagler) Airport; NDB Runway 34, Amdt. 8; Revised.

5. Section 97.29 is amended by establishing, revising, or canceling the following ILS SIAPs, effective May 6, 1971.

Birmingham, Ala.—Municipal Airport; ILS Runway 5, Amdt. 26; Revised.
Daytona Beach, Fla.—Daytona Beach Regional Airport; ILS Runway 6L, Amdt. 14; Revised.
Jackson, Miss.—Allen C. Thompson Field; ILS Runway 15L, Amdt. 8; Revised.
Las Vegas, Nev.—McCarran International Airport; ILS Runway 25, Amdt. 2; Revised.
Macon, Ga.—Lewis B. Wilson Airport; ILS Runway 5, Amdt. 16; Revised.
Pensacola, Fla.—Pensacola Municipal (Hagler) Airport; ILS Runway 16, Amdt. 2; Revised.

6. Section 97.31 is amended by establishing, revising, or canceling the following Radar SIAPs, effective May 6, 1971.

Jackson, Miss.—Allen C. Thompson Field; Radar-1, Amdt. 7; Revised.
Las Vegas, Nev.—McCarran International Airport; Radar-1, Amdt. 4; Revised.
Macon, Ga.—Lewis B. Wilson Airport; Radar-1, Amdt. 8; Revised.

(Secs. 307, 313, 601, 1110, Federal Aviation Act of 1958; 49 U.S.C. 1438, 1354, 1421, 1510, sec. 6(c), Department of Transportation Act, 49 U.S.C. 1655(c) and 5 U.S.C. 552(a)(1))

Issued in Washington, D.C., on March 31, 1971.

JAMES F. RUDOLPH,
Director,
Flight Standards Service.

NOTE: Incorporation by reference provisions in §§ 97.10 and 97.20 approved by the Director of the FEDERAL REGISTER on May 12, 1969 (35 F.R. 5610).

[FR Doc.71-4915 Filed 4-9-71;8:45 am]

Title 21—FOOD AND DRUGS

Chapter I—Food and Drug Administration, Department of Health, Education, and Welfare

SUBCHAPTER A—GENERAL

PART 1—REGULATIONS FOR THE ENFORCEMENT OF THE FEDERAL FOOD, DRUG, AND COSMETIC ACT AND THE FAIR PACKAGING AND LABELING ACT

Animal Feed Labeling; Collective Names for Feed Ingredients

In the FEDERAL REGISTER of July 30, 1970 (35 F.R. 12212), the Commissioner of Food and Drugs proposed that Part 1 be amended to permit the use of collective names for feed ingredients in livestock feed labeling in lieu of each ingredient being specified by its common or usual name. The proposal was based on the fact that day-to-day variations in availability and cost of feed ingredients make it impractical to require the labeling of each formulation to bear the common or usual name of each such ingredient in accordance with section 403(i)(2) of the Federal Food, Drug, and Cosmetic Act.

In response to the proposal, 45 comments were received. Most of the 37 endorsing the proposal suggested changes which principally are as follows:

1. The collective names should conform to those adopted by the Association of American Feed Control Officials. This reasonable suggestion is adopted to achieve uniformity in labeling requirements.

2. The term "livestock" should be changed to "all animals other than man" or "livestock and poultry." For clarity the latter term is adopted. Since it is not intended that the collective terms apply to pet foods, the former suggested term is rejected.

3. The collective terms should apply to feeds that contain one or more of the ingredients from any category rather than two or more as proposed. This reasonable suggestion is consistent with the proposal's purpose and is adopted.

4. Certain additional feed constituents should be added to those in the proposal. The constituents in question have since been adopted by the Association of American Feed Control Officials and are therefore included in this order.

Right responses expressed reservations on the grounds that the proposed amendments would permit the use of labeling that might not be as informative as feed purchasers would desire. Currently, a statement of actual quantities of each feed constituent is not required. Accordingly, a complete evaluation of the nutritional value of mixed feeds is generally not possible, even with the names of each feed constituent.

Having considered the comments received and other relevant material, the Commissioner concludes that the proposed amendments, with changes, should be adopted as set forth below.

Therefore, pursuant to provisions of the act (secs. 403(i), 701(a), 52 Stat. 1048, 1055; 21 U.S.C. 343, 371(a)) and under authority delegated to the Commissioner (21 CFR 2.120), Part 1 is amended by revising § 1.10(a) and by adding a new § 1.16, as follows:

§ 1.10 Food; labeling; designation of ingredients.

(a) The name of an ingredient (except a spice, flavoring, or coloring which is an ingredient of a food other than one sold as a spice, flavoring, or coloring) required by section 403(i)(2) of the act to be on the label of a food shall be a specific name and not a collective name; however, this provision does not apply if the food is a livestock or poultry feed within the meaning of section 201(x) of the act and meets the requirements for the use of collective names for certain feed ingredients as prescribed by § 1.16. If an ingredient (which itself contains two or more ingredients) conforms to a definition and standard of identity prescribed by regulations under section 401 of the act, such ingredient may be designated on the label of such food by the name specified in the definition and standard, supplemented, in case such regulations require the naming of optional ingredients present in such ingredient, by a statement showing the optional ingredients which are present in such ingredient.

* * * * *

§ 1.16 Animal feed labeling; collective names for feed ingredients.

(a) An animal feed shall be exempt from the requirements of section 403(i)(2) of the act with respect to its label bearing the common or usual names of the animal feed ingredients listed in paragraph (b) of this section under the following prescribed conditions:

(1) The animal feed is intended solely for livestock and poultry.

(2) The label of the animal feed bears the collective name(s) prescribed in paragraph (b) of this section in lieu of the corresponding common or usual names of the individual feed ingredients contained therein.

(3) The label of the animal feed otherwise conforms to the requirements of section 403(i)(2) of the act.

(4) The ingredients of any feed listed in paragraph (b) of this section neither contain nor are food additives as defined in section 201(s) of the act unless provided for by and in conformity with applicable regulations established pursuant to section 409 of the act.

(b) Each collective name referred to in this paragraph may be used for the purpose of labeling where one or more of the ingredients listed for that collective name are present. The animal feed ingredients listed under each of the collective names are the products defined by the Association of American Feed Control Officials. The collective names are as follows:

(1) "Animal protein products" include one or more of the following: Animal products, marine products, and milk products.

(2) "Forage products" include one or more of the following: Alfalfa meals, entire plant meals, hays, and stem meals.

(3) "Grain products" include one or more of the following: Barley, grain sorghums, maize (corn), oats, rice, rye, and wheat.

(4) "Plant protein products" include one or more of the following: Algae meals, coconut meals (copra), cottonseed meals, guar meal, linseed meals, peanut meals, safflower meals, soybean meals, sunflower meals, and yeasts.

(5) "Processed grain byproducts" include one or more of the following: Brans, brewers dried grains, distillers grains, distillers solubles, flours, germ meals, gluten feeds, gluten meals, grits, groats, hominy feeds, malt sprouts, middlings, pearled, polishings, shorts, and wheat mill run.

(6) "Roughage products" include one or more of the following: Cobs, hulls, husks, pulps, and straws.

Effective date. This order shall become effective 30 days after its date of publication in the FEDERAL REGISTER.

(Secs. 403(i), 701(a), 52 Stat. 1048, 1055; 21 U.S.C. 343, 371(a))

Dated: April 1, 1971.

SAM D. FINE,
Associate Commissioner for Compliance.

[FR Doc.71-5010 Filed 4-9-71;8:46 am]

PART 8—COLOR ADDITIVES

Subpart C—Listing of Color Additives for Food Use Subject to Certification

Subpart E—Listing of Color Additives for Drug Use Subject to Certification

FD&C RED NO. 40

The Commissioner of Food and Drugs, on the basis of a petition submitted by Allied Chemical Corp., Specialty Chemicals Division, c/o Hazleton Laboratories, Inc., Post Office Box 30, Falls Church, Va. 22046, and other relevant material, finds that FD&C Red No. 40 (identified below) is safe for use in food and drugs under the conditions prescribed in this order and that certification is necessary for the protection of the public health. In the petition and the notice of filing thereof which was published June 27, 1970 (35 F.R. 10529), the additive's name was "Allura Red AC"; however, the petitioner has requested that the additive be known as "FD&C Red No. 40."

Therefore, pursuant to provisions of the Federal Food, Drug, and Cosmetic Act (sec. 706(b), (c)(1), (d), 74 Stat. 399-403; 21 U.S.C. 376(b), (c)(1), (d)) and under authority delegated to the Commissioner (21 CFR 2.120): *It is ordered,* That Part 8 be amended by adding § 8.244 to Subpart C and § 8.4104 to Subpart E, as follows:

§ 8.244 FD&C Red No. 40.

(a) *Identity.* (1) The color additive FD&C Red No. 40 is principally the disodium salt of 6-hydroxy-5-[(2-methoxy-5-methyl-4- sulfophenyl)azo]- 2- naphthalenesulfonic acid.

(2) Color additive mixtures for food use (including dietary supplements) made with FD&C Red No. 40 may contain only those diluents that are suitable and that are listed in Subpart D of this part as safe for use in color additive mixtures for coloring foods.

(b) *Specifications.* FD&C Red No. 40 shall conform to the following specifications and shall be free from impurities other than those named to the extent that such other impurities may be avoided by good manufacturing practice:

Sum of volatile matter (at 135° C.) and chlorides and sulfates (calculated as sodium salts), not more than 14.0 percent.
Water-insoluble matter, not more than 0.2 percent.
Higher sulfonated subsidiary colors (as sodium salts), not more than 1.0 percent.
Lower sulfonated subsidiary colors (as sodium salts), not more than 1.0 percent.
Disodium salt of 6-hydroxy-5-[(2-methoxy-5-methyl-4-sulfophenyl)azo]-8-(2-methoxy-5-methyl-4-sulfophenoxy)-2-naphthalenesulfonic acid, not more than 1.0 percent.
Sodium salt of 6-hydroxy-2-naphthalenesulfonic acid (Schaeffer's salt), not more than 0.3 percent.
4-Amino-5-methoxy-o-toluenesulfonic acid, not more than 0.2 percent.
Disodium salt of 6,6'-oxybis(2-naphthalenesulfonic acid), not more than 1.0 percent.
Lead (as Pb), not more than 10 parts per million.
Arsenic (as As), not more than 1 part per million.
Total color, not less than 85.0 percent.

(c) *Uses and restrictions.* FD&C Red No. 40 may be safely used for coloring foods (including dietary supplements) generally in amounts consistent with good manufacturing practice except that it may not be used to color foods for which standards of identity have been promulgated under section 401 of the act unless added color is authorized by such standards.

(d) *Labeling.* The label of the color additive and any mixtures prepared therefrom intended solely or in part for coloring purposes shall conform to the requirements of § 8.32.

(e) *Certification.* All batches of FD&C Red No. 40 shall be certified in accordance with regulations in Subpart A of this part.

§ 8.4104 FD&C Red No. 40.

(a) *Identity and specifications.* (1) The color additive FD&C Red No. 40 shall conform in identity and specifications to the requirements of § 8.244 (a)(1) and (b).

(2) Color additive mixtures for drug use made with FD&C Red No. 40 may contain only those diluents that are suitable and that are listed in Subpart F of this part as safe for use in color additive mixtures for coloring drugs.

(b) *Uses and restrictions.* FD&C Red No. 40 may be safely used in coloring drugs, subject to the restrictions on use of color additives in § 8.101, in amounts consistent with good manufacturing practice.

(c) *Labeling.* The label of the color additive and any mixtures prepared therefrom intended solely or in part for coloring purposes shall conform to the requirements of § 8.32.

(d) *Certification.* All batches of FD&C Red No. 40 shall be certified in accordance with regulations in Subpart A of this part.

Any person who will be adversely affected by the foregoing order may at any time within 30 days after its date of publication in the FEDERAL REGISTER file with the Hearing Clerk, Department of Health, Education, and Welfare, Room 6-62, 5600 Fishers Lane, Rockville, Md. 20852, written objections thereto. Objections shall show wherein the person filing will be adversely affected by the order and specify with particularity the provisions of the order deemed objectionable and the grounds for the objections. If a hearing is requested, the objections must state the issues for the hearing and such objections must be supported by grounds legally sufficient to justify the relief sought. Objections may be accompanied by a memorandum or brief in support thereof. All documents shall be filed in six copies.

Effective date. This order shall become effective 60 days after its date of publication in the FEDERAL REGISTER, except as to any provisions that may be stayed by the filing of proper objections. Notice of the filing of objections or lack thereof will be given by publication in the FEDERAL REGISTER.

(Sec. 706(b), (c)(1), (d), 74 Stat. 399-403; 21 U.S.C. 376(b), (c)(1), (d))

Dated: April 1, 1971.

SAM D. FINE,
Associate Commissioner for Compliance.

[FR Doc.71-5011 Filed 4-9-71;8:46 am]

PART 8—COLOR ADDITIVES

Subpart E—Listing of Color Additives for Drug Use Subject to Certification

FD&C BLUE NO. 2; CONFIRMATION OF EFFECTIVE DATE

In the matter of listing the color additive FD&C Blue No. 2 for use in or on nylon surgical sutures with certification required:

No objections were filed to the order in the above-identified matter published in the FEDERAL REGISTER of February 13, 1971 (36 F.R. 2967). Therefore, pursuant to provisions of the Federal Food, Drug, and Cosmetic Act (sec. 706 (b), (c)(1), (d), 74 Stat. 399-403; 21 U.S.C. 376 (b), (c)(1), (d)) and under authority delegated to the Commissioner of Food and Drugs (21 CFR 2.120), notice is given that the regulation established by the subject order (§ 8.4022) will become effective April 14, 1971.

Dated: April 1, 1971.

SAM D. FINE,
Associate Commissioner for Compliance.

[FR Doc.71-5012 Filed 4-9-71;8:46 am]

Title 41—PUBLIC CONTRACTS AND PROPERTY MANAGEMENT

Chapter 114—Department of the Interior

PART 114-26—PROCUREMENT SOURCES AND PROGRAMS

Subpart 114-26.7—Use of GSA Supply Sources by Grantees and Contractors

Pursuant to the authority of the Secretary of the Interior contained in 5 U.S.C. 301 (Supp. V, 1965-69) and Sec. 205(c), 63 Stat. 390; 40 U.S.C. 486(c), a

new Subpart 114–26.7 is added to Chapter 114, Title 41 of the Code of Federal Regulations, as set forth below.

This subpart shall become effective on the date of its publication in the FEDERAL REGISTER (4–10–71).

RICHARD R. HITE,
Deputy Assistant Secretary, of the Interior.

APRIL 5, 1971.

Sec.
114–26.700 Scope of subpart.
114–26.701 Applicability.
114–26.702 Policy on use of GSA supply sources.
114–26.704 Agency authorizations.
114–26.707 Acquisition of GSA stock.

AUTHORITY: The provisions of this Subpart 114–26.7 issued under 5 U.S.C. 301 (Supp. V, 1965–1969); sec. 205(c), 63 Stat. 390; 40 U.S.C. 486(c).

Subpart 114–26.7—Use of GSA Supply Sources by Grantees and Contractors

§ 114–26.700 Scope of subpart.

This subpart prescribes the Department's policy regarding the use of GSA supply sources by grantees and cost-reimbursement type contractors.

§ 114–26.701 Applicability.

The provisions of this subpart are applicable to all bureaus and offices of the Department.

§ 114–26.702 Policy on use of GSA supply sources.

(a) It is the policy of the Department that grantees shall be authorized to use GSA supply sources when it is in the best interest of the Government and not prohibited by law.

(b) It is the policy of the Department that, pursuant to the provisions of Federal Procurement Regulation Subpart 1–5.9, cost-reimbursement type contractors shall be authorized to use GSA supply sources when such use is in the best interest of the Government and is not prohibited by law.

§ 114–26.704 Agency authorizations.

(a) (1) In addition to other limitations and conditions, each authorization shall specify that the issuing office will not be liable for any obligations incurred by the grantee or contractor through the use of the authorization to use GSA supply sources.

(2) Authorizations shall also be limited to purchases within approved budgetary limitations, with a requirement for the prompt payment of bills upon receipt of billing.

(3) Each issuing office shall be responsible for monitoring the purchases made under each authorization, and for requiring the maintenance of purchase records needed for audit purposes.

(b) (1) Public Law 85–934 (72 Stat. 1793; 42 U.S.C. 1892) authorizes the vesting of title to equipment purchased with Federal funds pursuant to grants or contracts for the conduct of basic or applied scientific research at nonprofit institutions of higher education, or at nonprofit organizations whose primary purpose is the conduct of scientific research (205 DM 9).

(2) This authority to vest title to scientific research equipment shall be used to reduce the Government's cost and administrative burden of accounting, shipment, storage, disposition, and otherwise treating the equipment as Government property.

(3) Office of Management and Budget Circular No. A–101, dated January 9, 1971, recognizes the need for reducing such costs and burden while furthering the overall Government objective of strengthening the scientific capability of educational institutions. The policy set forth in OMB Circular No. A–101 shall be used to determine when title to equipment should be vested in such institutions. This policy applies to educational institutions only. Part 114–51 of this chapter establishes the Departmental policy for the management of equipment and supplies acquired under project grants.

§ 114–26.707 Acquisition of GSA stock.

(a) Grantees and contractors should obtain FEDSTRIP system orientation from the GSA Regional Supply Service Officer.

(b) Requisitions submitted to GSA under this system shall include a statement of authorization substantially as follows:

This order is placed pursuant to written authorization No. _____ dated _____ issued by _____ for (*contract number or grant*).

[FR Doc.71–5021 Filed 4–9–71;8:47 am]

Title 43—PUBLIC LANDS: INTERIOR

Chapter II—Bureau of Land Management, Department of the Interior

APPENDIX—PUBLIC LAND ORDERS

[Public Land Order 5037]

[Wyoming 27622]

WYOMING

Modification of Powersite Reserve No. 348 To Permit Grant of Right-of-Way

By virtue of the authority contained in section 24 of the Act of June 10, 1920, 41 Stat. 1075, as amended, 16 U.S.C. sec. 818 (1964), and pursuant to the determination of the Federal Power Commission in DA–165-Wyoming, it is ordered as follows:

The Executive Order of March 27, 1913, creating Powersite Reserve No. 348, is hereby modified to the extent necessary to permit the location of a right-of-way under section 2477, U.S. Revised Statutes, 43 U.S.C. sec. 932, by Park County, over the following described lands, as delineated on a map on file with the Bureau of Land Management in Wyoming 17877, for a maintenance of a public road, subject to the provisions of section 24 of the Federal Power Act, supra:

SIXTH PRINCIPAL MERIDIAN

T. 52 N., R. 104 W.,
Sec. 15, lots 28;
Sec. 16, lots 18, 22;
Sec. 19, lots 19, 22, 24.

The areas described aggregate 97.22 acres in Park County.

HARRISON LOESCH,
Assistant Secretary of the Interior.

APRIL 6, 1971.

[FR Doc.71–5018 Filed 4–9–71;8:47 am]

[Public Land Order 5038]

[Sacramento 3531]

CALIFORNIA

Withdrawal and Reservation of Lands in Connection with the Honey Lake Waterfowl Management Area

By virtue of the authority vested in the President and pursuant to Executive Order No. 10355 of May 26, 1952 (17 F.R. 4831), it is ordered as follows:

1. Subject to valid existing rights, the following described public lands, which are under the jurisdiction of the Secretary of the Interior, are hereby withdrawn from all forms of appropriation under the public land laws, including the mining laws (30 U.S.C., Ch. 2), but not from leasing under the mineral leasing laws, and reserved for management in cooperation with the State of California Department of Fish and Game in connection with the Honey Lake Waterfowl Management Area:

MOUNT DIABLO MERIDIAN

T. 28 N., R. 14 E.,
Sec. 9, NE¼SE¼, S½SE¼.
T. 28 N., R. 15 E.,
Sec. 7, lots 3, 4, 5, 6.

The areas described aggregate 191.46 acres in Lassen County.

2. Upon execution of a cooperative agreement with the Secretary of the Interior, or his delegate, the State of California is authorized to manage the withdrawn lands for the conservation of small game and waterfowl, and as a public hunting ground, consistent with Federal programs for the management of the lands.

3. The withdrawal made by this order does not alter the applicability of the public land laws governing the use of the lands under lease, license, or permit, or governing the disposal of their mineral or vegetative resources other than under the mining laws. However, leases, licenses, contracts, or permits will be issued only if the proposed use of the lands will not interfere with proper management of the Honey Lake Waterfowl Management Area.

HARRISON LOESCH,
Assistant Secretary of the Interior.

APRIL 6, 1971.

[FR Doc.71–5019 Filed 4–9–71;8:47 am]