DISPOSITION: 2–16–62. Certain pieces of the accompanying labeling having been seized and no capsules of the article having been found available for seizure, and The DePree Co. having consented to the entry of a decree, judgment of condemnation was entered and the labeling was ordered destroyed.

7047. Mineral Life capsules. (F.D.C. No. 45300. S. No. 23–318 R.)

QUANTITY: 25 20-capsule boxes at Oklahoma City, Okla.

SHIPPED: 10–8–60 and 10–10–60, from Logan, Utah, by Nutritional Progress Scientific Co.

LABEL IN PART: "MINERAL LIFE

Active Ingredients	Major ingredient Not More Than
Silicon	
Iron	0.5 to 2.0%
Calcium	0.2 to 1.5%
Magnesium	0.2 to 1.5%
Vanadium	0.5 to 1.5%
Titanium	0.3 to 1.0%
Copper	0.3 to 2.0%
Potassium	0.02 to 0.1%
Strontium	0.005 to 0.05%
Lead, Zirconium, Cobalt, Manganese	0.01 to 0.1%
Chromium, Molybdenum	0.0005 to 0.005%
Nickel	0.0001 to 0.001%

Dosage: 4 yrs. to 12 yrs.–1 capsule; 12 years and up–up to 5 capsules. This is a natural mineral. Not compounded by man. * * *

Nutritional Progress Scientific Co., 599 West Center, Logan, Utah."

ACCOMPANYING LABELING: Leaflets entitled "Mineralife And What It Means To You" and "Eject Death From The Temple"; and order blanks entitled "Mineral Life."

LIBELED: 12–30–60, W. Dist. Okla.

CHARGE: 502(a)—when shipped and while held for sale, the article's accompanying labeling contained false and misleading representations that the article was adequate and effective as a treatment for and preventive of cancer, cancer of the face, disease, illness, inefficient absorption of food, inefficient elimination, circulatory system coated and clogged with calcium and other sludge deposits, premature aging, starving body cells, toxic secretions, excessive moisture, fats and dead cell tissues, heart trouble, heart flutter and laborious pounding and breathlessness after strenuous exercise, sleeplessness, loss of appetite, shortness of breath following Asiatic flu, extreme tiredness; arthritis with soreness, weakness, stiffness, and throbbing; tumor, pain, severe stomach pain and failure to eat and sleep following flu, stomach trouble, colon trouble, ulcers, rheumatoid arthritis, pancreatic and stomach injuries, constipation, nervousness, weakness, bursitis, neuralgia, neuritis, heart palpitations, overweight condition, and anemic condition; and to eliminate any condition or substance which causes disease of the body; regulate body functions; control water balance; maintain acid-base equilibrium and promote utilization of foodstuffs; promote sound teeth and give rigidity and relative prominence to skeletal tissues; maintain elasticity and irritability of muscles and nerves; promote good health; become a relaxed, calmer person; promote strong, vibrant blood, and for pep; and 502(e)(2)—the article was fabricated from two or more ingredients and its label failed to bear the common or usual name of each active ingredient.

DISPOSITION: On 2–17–61, the shipper filed a claim, an answer, and a notice of a motion to remove the action to the District of Colorado. On 5–22–61, the action was transferred to the District of Colorado. On 9–13–61, the claimant, without admitting the allegations of the libel, consented to a decree of condemnation and destruction; an order adjudging the article misbranded and ordering its destruction was entered.

7048. B_{12} injection. (F.D.C. No. 46532. S. No. 18–681 T.)

QUANTITY: 593 10-cc. vials at San Antonio, Tex., in possession of Knight Pharmacal Co.

SHIPPED: 6–23–61 and 8–23–61, from Philadelphia, Pa.

LABEL IN PART: "10 cc Multiple-Dose Vial Crystalline Vitamin B_{12} * * * Distributed by Knight Pharmacal Co. San Antonio, Texas."

ACCOMPANYING LABELING: Carton insert, reading in part, "Vitamin B_{12} Injection."

LIBELED: 11–16–61, W. Dist. Tex.

CHARGE: 502(a)—while held for sale, the accompanying labeling of the article contained false and misleading representations that the article was adequate and effective for treatment of neurological conditions, trigeminal neuralgia, multiple sclerosis, and neuro-anemic syndromes.

DISPOSITION: 12–20–61. Default—delivery to a public, charitable hospital.

7049. Lecitabs (lecithin tablets). (F.D.C. No. 46968. S. No. 34–677 T.)

QUANTITY: 56 90-tablet btls. and 13 180-tablet btls. at Minneapolis, Minn.

SHIPPED: Between 6–23–61 and 10–4–61, from Chicago, Ill., by National Lecithin, Inc.

LABEL IN PART: "National Lecitabs Lecithin Tablets A Natural Food Product Highly concentrated extra rich, Soya Lecithin formula of 95% oil free Phosphatides. Ingredients: Soya Lechithin, in a base of non-fat, dry milk solids and soy protein. Natural flavoring added. Sole Distributors: National Lecithin, Inc. Chicago 26, Ill. * * * a dietary supplement of natural lipotropic factors. * * * a rich, natural source of Lecithin, Cephalin, Choline and Inositol Phosphatides * * * rich in both linoleic and linolenic acids."

LIBELED: 2–20–62, Dist. Minn.

CHARGE: 502(a)—when shipped, the label contained false and misleading representations that the article was adequate and effective to promote utilization of fat and to lower blood cholesterol.

The libel alleged also that the article was misbranded under the provisions of the law applicable to foods, as reported in notices of judgment on foods.

DISPOSITION: 4–6–62. Default—destruction.

7050. Alfalfa tablets. (F.D.C. No. 46463. S. No. 81–971 R.)

QUANTITY: 1 drum, containing approximately 29,000 tablets, at Forest, Miss., in possession of Pasco Products. Inc.

SHIPPED: 6–27–61, from North Kansas City, Mo.

LABEL IN PART: "Alfalfa Tablets, Red."

ACCOMPANYING LABELING: Folder entitled "Alfa-Lite In Liquid or Tablet"; and package label reading in part "100 Tablets Alfa-Life Concentrated Alfalfa Extract 100 Mg. in each tablet. For the treatment of joint pains and stiffness resulting from arthritic and rheumatic like conditions * * * Distrib-