

CHARGE: 502(a)—the name "Multizyme," by which the drug was designated while held for sale, was misleading since it suggested the presence of multiple enzymes in the article as the valuable factor needed in a food supplement for human nutrition, whereas such was not the fact; and, in addition, the leaflets and order cards accompanying the article, while held for sale, contained false and misleading representations that the article would keep the bloodlines of the body clean and so protect one from high blood pressure, and that it would correct the suffering caused by the lack of balance of forces within the body cells; 502(b) (1) and (2)—when shipped, the article failed to bear a label containing the name and place of business of the manufacturer, packer, or distributor, and an accurate statement of the quantity of contents; 502(e) (2)—when shipped, the article failed to bear a label containing the common or usual name of each active ingredient contained therein; and 502(f) (1)—when shipped, the labeling of the article failed to bear adequate directions for use for the purposes for which it was intended.

DISPOSITION: 12-23-57. Consent—claimed by Lee V. Schneider, Seattle, Wash., and relabeled.

DRUGS AND DEVICES ACTIONABLE BECAUSE OF DEVIATION FROM OFFICIAL OR OWN STANDARDS*

5449. Posterior pituitary injection. (F.D.C. No. 40200. S. No. 54-805 M.)

QUANTITY: 99 1-cc. vials at Lynwood, Wash.

SHIPPED: 3-19-57, from Philadelphia, Pa., by Lustgarten Laboratories, Inc.

LABEL IN PART: "1 cc Amp. Pituitary Extract 10 U. S. P. Units (Obstetrical)
* * * Control No. 020477 * * * Vitamix Corporation Philadelphia, Pa."

RESULTS OF INVESTIGATION: Analysis showed that the potency of the article was less than 0.062 U. S. P. posterior pituitary units per cubic centimeter.

LIBELED: 5-16-57, W. Dist. Wash.

CHARGE: 501(b)—the strength of the article, when shipped, differed from the standard for "Posterior Pituitary Injection" set forth in the United States Pharmacopeia; and 502(a)—the label statement "1 cc Amp. Pituitary Extract 10 U. S. P. Units" was false and misleading as applied to a product, the potency of which is less than 10 U. S. P. posterior pituitary units per cubic centimeter.

DISPOSITION: 7-23-57. Default—destruction.

5450. Progesterone-estrogen. (F.D.C. No. 40367. S. No. 65-086 M.)

QUANTITY: 31 vials in a carton at Columbus, Ohio.

SHIPPED: 4-12-57, from Sarasota, Fla., by Stilco Laboratories.

LABEL IN PART: (Ctn.) "Stilco Laboratories * * * Sarasota, Florida,
35 x 10 cc — Control #956 — Progesterone-Estrogen In Sesame Oil — Intramuscular — Per CC Progesterone USP 25 MGM — Estrogens (95-98% Estrone) 25,000 I. U."

RESULTS OF INVESTIGATION: Analysis showed that the article was an oil solution containing 21.6 percent of the declared amount of estrogens.

LIBELED: 7-2-57, S. Dist. Ohio.

*See also Nos. 5443, 5446.