LABEL, IN PART: "Sterile Solution Posterior Pituitary (obstetrical) 10 Units per cc. with Chlorobutanol (chloral Deriv.) 0.5%."

NATURE OF CHARGE: Adulteration, Section 501 (b), the article purported to be "Posterior Pituitary Injection," a drug, the name of which is recognized in the United States Pharmacopeia, an official compendium, and its strength differed from the official standard since the United States Pharmacopeia provides that each cubic centimeter of *posterior pituitary injection* possesses an activity equivalent to 10 U. S. P. posterior pituitary units, whereas each cubic centimeter of the article possessed an activity equivalent to less than 10 U. S. P. posterior pituitary units.

DISPOSITION: March 25, 1952. Default decree of condemnation. The court ordered that the product be turned over to the Federal Security Agency.

3752. Adulteration of conjugated estrogen tablets. U. S. v. 885 Bottles * * *. (F. D. C. No. 32587. Sample No. 21235–L.)

LIBEL FILED: January 11, 1952, Western District of Texas.

ALLEGED SHIPMENT: On or about January 29, 1951, from St. Louis, Mo.

PRODUCT: *Conjugated estrogen tablets.* 885 100-tablet bottles at San Antonio, Tex. Analysis showed that the product contained a total amount of estrogenic steroids calculated to be 0.60 mg. of sodium estrone sulfate per tablet.

LABEL, IN PART: "Each tablet contains water soluble conjugated estrogens, naturally occurring, expressed as sodium estrone sulfate 1.25 mg."

NATURE OF CHARGE: Adulteration, Section 501 (c), the strength of the article differed from that which it was represented to possess. The article was adulterated while held for sale after shipment in interstate commerce.

DISPOSITION: April 24, 1952. Default decree of condemnation and destruction.

3753. Adulteration and misbranding of Arvimin. U. S. v. 5 Cases * * *. (F. D. C. No. 33067. Sample No. 12091–L.)

LIBEL FILED: April 11, 1952, Southern District of Ohio.

ALLEGED SHIPMENT: On or about June 19, 1951, from Marion, Ohio.

PRODUCT: 5 cases, each containing 12 1-pound cans, of *Arvimin* at Cincinnati, Ohio. Analysis showed that the product contained approximately 16 percent of the declared amount of vitamin A and 50 percent of the declared amount of vitamin D_2.

LABEL, IN PART: "Arvimin Each 1 Lb. (453.6 Grams) Represents—Active Drug Ingredient—Sodium Arsanilate * * * 5.0 Grams Incorporated in a Nutritional Base Composed of * * * Vitamin A (U. S. P. Units) 300,000 Units – Vitamin D_2 (U. S. P. Units) 1,000,000 Units."

NATURE OF CHARGE: Adulteration, Section 501 (c), the strength of the article differed from that which it purported and was represented to possess.

Misbranding, Section 502 (a), the label statement "Each 1 Lb. * * * Represents * * * Vitamin A (U. S. P. Units) 300,000 Units – Vitamin D_2 (U. S. P. Units) 1,000,000 Units" was false and misleading as applied to the article, which contained less than those amounts of vitamins A and D_2.

The article was adulterated and misbranded while held for sale after shipment in interstate commerce.

DISPOSITION: May 23, 1952. Default decree of condemnation and destruction.