

LABEL, IN PART: "Private Formula Control No. 04639 41,400 Special Formula Capsules #0 Clear Each Capsule contains: 2.0 Mg. Thiamin (2MDR) 2.0 Mg. Riboflavin (1MDR) * * * 9.0 Mg. Niacinamide."

NATURE OF CHARGE: Adulteration, Section 402 (b) (1), valuable constituents, thiamine, riboflavin, and niacinamide, had been in whole or in part omitted or abstracted therefrom.

Misbranding, Section 403 (a), the label statement "Each Capsule contains: 2.0 Mg. Thiamin (2MDR) 2.0 Mg. Riboflavin (1MDR) * * * 9.0 Mg. Niacinamide" was false and misleading as applied to the article, which contained less than these amounts of the stated substances.

DISPOSITION: July 10, 1951. National Drug Laboratories, Inc., claimant, having consented to the entry of a decree, judgment of condemnation was entered and the court ordered that the product be released under bond to be relabeled under the supervision of the Federal Security Agency.

17745. Adulteration and misbranding of Stur-Dee vitamin tablets. U. S. v. 128 Bottles * * *. (F. D. C. No. 30928. Sample No. 24442-L.)

LABEL FILED: April 17, 1951, Eastern District of New York.

ALLEGED SHIPMENT: On or about March 8, 1951, by the Midwest Chemical Development Corp., from Cleveland Ohio.

PRODUCT: 128 bottles each containing 55 Stur-Dee vitamin tablets at Brooklyn, N. Y.

LABEL, IN PART: (Bottle) "Stur-Dee Brand 55 Tablets Natural Vitamin A & D Tablets * * * Each tablet contains: Vitamin A 5,000 USP Units."

NATURE OF CHARGE: Adulteration, Section 402 (b) (1), a valuable constituent, Vitamin A, had been in whole or in part omitted or abstracted from the product.

Misbranding, Section 403 (a), the label statement "Each tablet contains: Vitamin A 5,000 USP Units" was false and misleading as applied to an article containing less than the declared amount of vitamin A.

DISPOSITION: June 22, 1951. Default decree of condemnation and destruction.

17746. Adulteration and misbranding of Patheba vitamin tablets. U. S. v. 4 Cases * * *. (F. D. C. No. 30497. Sample No. 94470-K.)

LABEL FILED: January 25, 1951, Southern District of Alabama.

ALLEGED SHIPMENT: On or about December 8, 1950, by the Brayten Pharmaceutical Co., from Chattanooga, Tenn.

PRODUCT: 4 cases, each containing 24 100-tablet bottles, of Patheba vitamin tablets at Mobile, Ala.

LABEL, IN PART: (Bottle) "Patheba Tabs * * * Each Tablet Contains * * * Thiamine Hydrochloride 2 Mg. * * * 13 Mg. Of Yeast Powder Containing 300 U. S. P. Units Vitamin D."

NATURE OF CHARGE: Adulteration, Section 402 (b) (1), valuable constituents, thiamine hydrochloride and vitamin D, had been in part omitted or abstracted from the product.

Misbranding, Section 403 (a), the label statement "Each Tablet Contains * * * Thiamine Hydrochloride 2 Mg. * * * 13 Mg. Of Yeast Powder Containing 300 U. S. P. Units Vitamin D" was false and misleading since the product contained less than 2 milligrams of thiamine hydrochloride and less than 300 U. S. P. units of vitamin D per tablet.