LABEL, IN PART: "10cc Ampul Sodium Thiosulfate * * * For Intravenous Use."

NATURE OF CHARGE: The article was adulterated while held for sale after shipment in interstate commerce under Section 501 (b), in that it purported to be and was represented as "Sodium Thiosulfate Ampuls," a drug the name of which is recognized in the National Formulary, an official compendium, and its quality and purity fell below the official standard since it was contaminated with undissolved material.

DISPOSITION: November 12, 1948. Default decree of forfeiture and destruction.

2614. Adulteration of sodium thiosulfate ampuls. U. S. v. 136 Boxes * * *. (F. D. C. No. 25629. Sample No. 43450–K.)

LIBEL FILED: September 15, 1948, Northern District of Illinois.

ALLEGED SHIPMENT: On or about April 20, 1948, by the Sherman Laboratories, from Detroit, Mich.

PRODUCT: 136 boxes, each containing 5 10-cc. ampuls, of *sodium thiosulfate* at Chicago, Ill.

NATURE OF CHARGE: Adulteration, Section 501 (b), the article purported to be and was represented as "Sodium Thiosulfate Ampuls," a drug the name of which is recognized in the National Formulary, an official compendium, and its quality and purity fell below the official standard since it was contaminated with viable micro-organisms and undissolved material.

DISPOSITION: March 28, 1949. Default decree of condemnation and destruction.

2615. Adulteration of sodium salicylate and sodium iodide with colchicine. U. S. v. 50 Vials * * *. (F. D. C. No. 25738. Sample No. 19572–K.)

LIBEL FILED: October 27, 1948, Eastern District of Tennessee.

ALLEGED SHIPMENT: On or about September 15, 1948, by the Direct Sales Co., Inc., from Buffalo, N. Y.

PRODUCT: 50 250-cc. vials of *sodium salicylate and sodium iodide with colchicine* at Chattanooga, Tenn.

NATURE OF CHARGE: Adulteration, Section 501 (b), the article purported and was represented as "Sodium Salicylate and Iodide with Colchicine Ampuls," a drug the name of which is recognized in the National Formulary, an official compendium, and its quality and purity fell below the official standard since it was contaminated with undissolved material.

DISPOSITION: On or about December 31, 1948. Default decree of condemnation and destruction.

2616. Adulteration of aminophylline. U. S. v. 375 Ampuls * * *. (F. D. C. No. 25702. Sample No. 15559–K.)

LIBEL FILED: October 12, 1948, Eastern District of Michigan.

ALLEGED SHIPMENT: On or about September 15, 1948, by the Direct Sales Co., Inc., from Buffalo, N. Y.

PRODUCT: 375 10-cc. size ampuls of *aminophylline* at Detroit, Mich.

LABEL, IN PART: "Each 10 cc represents Aminophylline U. S. P. 3¾ Grs. * * * For Intravenous Use."

NATURE OF CHARGE: Adulteration, Section 501 (b), the article purported to be and was represented as "Aminophylline Injection," a drug the name of which is recognized in the United States Pharmacopoeia, an official compendium, and its quality and purity fell below the official standard since it was contaminated with undissolved material.

DISPOSITION: November 12, 1948. Default decree of condemnation and destruction.

2617. Adulteration of calcium gluconate. U. S. v. 1,043 Ampuls * * *. (F. D. C. No. 25913. Sample Nos. 29360–K, 29361–K.)

LIBEL FILED: November 8, 1948, District of Colorado.

ALLEGED SHIPMENT: On or about March 2 and September 29, 1948, by the Carroll Dunham Smith Pharmacal Co., from Kansas City, Mo.

PRODUCT: 1,043 10-cc. ampuls of *calcium gluconate* at Colorado Springs, Colo.

NATURE OF CHARGE: Adulteration, Section 501 (b), the article purported to be and was represented as "Calcium Gluconate Injection," a drug the name of which is recognized in the United States Pharmacopoeia, an official compendium, and its quality and purity fell below the official standard since it was contaminated with undissolved material.

DISPOSITION: December 23, 1948. Default decree of condemnation and destruction.

2618. Adulteration of thiamine hydrochloride. U. S. v. 26 Vials * * *. (F. D. C. No. 25765. Sample No. 10781–K.)

LIBEL FILED: September 21, 1948, Eastern District of New York.

ALLEGED SHIPMENT: On or about April 21, 1948, from Newark, N. J.

PRODUCT: 26 30-cc. vials of *thiamine hydrochloride* at Brooklyn, N. Y.

LABEL, IN PART: "Sterile Multiple Dose Vial Thiamine Hydrochloride Vitamin B_1, * * * intramuscularly or intravenously."

NATURE OF CHARGE: Adulteration, Section 501 (b), the article purported to be and was represented as "Thiamine Hydrochloride Injection," a drug the name of which is recognized in the United States Pharmacopoeia, an official compendium, and its quality and purity fell below the official standard since it was contaminated with undissolved material. The article was adulterated while held for sale after shipment in interstate commerce.

DISPOSITION: December 15, 1948. Default decree of condemnation and destruction.

2619. Adulteration of solution of vitamin B complex. U. S. v. 17 Vials * * *. (F. D. C. No. 25752. Sample No. 30143–K).

LIBEL FILED: September 16, 1948, District of Arizona.

ALLEGED SHIPMENT: On or about July 26, 1948, by the American Bio-Chemical Corp., from Los Angeles, Calif.

PRODUCT: 17 30-cc. vials of *solution of vitamin B complex* at Phoenix, Ariz.

LABEL, IN PART: "Sterile Solution Some Factors of Vitamin B Complex."

NATURE OF CHARGE: Adulteration, Section 501 (c), the purity and quality of the article fell below that which it purported and was represented to possess, namely, "Sterile Solution Some Factors of Vitamin B Complex * * * For intravenous or intramuscular use," since the article contained undissolved