Misbranding, Section 502 (a), the label statement "This product is sterile and non-pyrogenic" was false and misleading as applied to an article contaminated with living micro-organisms and pyrogen.

DISPOSITION: September 13, 1948. Default decree of condemnation. The product was ordered delivered to the Food and Drug Administration, for testing purposes.

2561. Adulteration of sodium iodide and sodium salicylate. U. S. v. 67 Ampuls * * *. (F. D. C. No. 23967. Sample Nos. 79517-H, 14603-K.)

LIBEL FILED: On November 24, 1947, Northern District of Illinois.

ALLEGED SHIPMENT: On or about July 18, 1947, by Bristol Laboratories, Inc., from Syracuse, N. Y.

PRODUCT: 67 ampuls of *sodium iodide and sodium salicylate* at Chicago, Ill.

LABEL, IN PART: "20 cc. size ampuls Sodium Iodide and Sodium Salicylate Sterile Solution for Intravenous Use."

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

DISPOSITION: February 3, 1948. Default decree of condemnation and destruction.

2562. Adulteration of sodium salicylate and iodide with colchicine. U. S. v. 4 Cartons * * *. (F. D. C. No. 25415. Sample No. 46005-K.)

LIBEL FILED: August 26, 1948, Eastern District of Missouri.

ALLEGED SHIPMENT: On or about June 11, 1948, from Philadelphia, Pa.

PRODUCT: 4 cartons, each containing 12 20-cc ampuls, of *sodium salicylate and iodide with colchicine* at St. Louis, Mo.

NATURE OF CHARGE: Adulteration, Section 501 (b), the article purported to be 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. The article was adulterated while held for sale after shipment in interstate commerce.

DISPOSITION: January 6, 1949. Default decree of condemnation and destruction.

2563. Adulteration of vitamin B complex with distilled water. U. S. v. 92 Packages * * *. (F. D. C. No. 25631. Sample No. 25868-K.)

LIBEL FILED: September 11, 1948, District of Minnesota.

ALLEGED SHIPMENT: On or about August 4, 1948, by Hyland Laboratories, from Los Angeles, Calif.

PRODUCT: 92 packages of *vitamin B complex with distilled water* at Minneapolis, Minn.

LABEL, IN PART: "10cc. B-Complex dried * * * with sterile diluent containing * * * Distilled Water 10cc."

NATURE OF CHARGE: Adulteration, Section 501 (c), the diluent purported to be and was represented as "Water for Injection," a drug the name of which is recognized in the United States Pharmacopoeia, an official compendium, and