CHARGE: 505(a)—the article was a new drug which may not be introduced into interstate commerce since an application filed pursuant to law was not effective with respect to such drug.

DISPOSITION: 11-13-59. Default—destruction.

DRUG REQUIRING CERTIFICATE OR RELEASE, FOR WHICH NONE HAD BEEN ISSUED

DRUG FOR VETERINARY USE

6087. Entero-Sol Powder. (F.D.C. No. 43323. S. No. 71-708 P.)

QUANTITY: 10 cases, 12 btls. each, at Gainesville, Ga.

SHIPPED: 5-25-59, from Vineland, N.J., by Eastern Laboratories, Inc.

LABEL IN PART: (Btl.) "Rollins Entero-Sol Powder For the treatment of Enteritis, Air-Sac, Colds and Blue Comb in Chickens and Turkeys * * * Contents 156 Gms. Each Pound Contains: 24,000,000 units of Penicillin G. Potassium 24,000 mg. of Dihydrostreptomycin Base as sulphate, and 3,600 mg. of Menadione Sodium Bisulfite (Synthetic Vitamin K) In addition each pound contains: * * * Manufactured for Ben C. Rollins Company 602 Grove Street, Gainesville, Georgia."

LIBELED: 8-4-59, N. Dist. Ga.

CHARGE: 502(a)—when shipped, the label of the article contained false and misleading representations that the article was an adequate and effective treatment for colds in chickens and turkeys; and 502(l)—the article consisted in part of penicillin and dihydrostreptomycin and it was not from a batch with respect to which a certificate or release had been issued.

DISPOSITION: 9-23-59. Default—destruction.

DRUG IN VIOLATION OF PRESCRIPTION LABELING REQUIREMENTS

6088. Dextro-amphetamine sulfate tablets and amphetamine sulfate tablets. (F.D.C. No. 43391. S. Nos. 73-809/10 P.)

QUANTITY: 5 1,000-tablet btls. at Bay Saint Louis, Miss.

SHIPPED: Prior to 7-7-59, from outside the State of Mississippi.

RESULTS OF INVESTIGATION: The bottles were unlabeled. Analysis showed that 3 bottles contained *dextro-amphetamine sulfate tablets* and 2 bottles contained *amphetamine sulfate tablets.*

LIBELED: 7-13-59, S. Dist. Miss.

CHARGE: 502(b)—while held for sale, the article failed to bear (1) a label containing the name and place of business of the manufacturer, packer, or distributor, and (2) an accurate statement of the quantity of contents in terms of count; 502(e)(1)—the label failed to bear the common or usual name of the drug; 502(f)(1)—the label of the article failed to bear adequate directions for use and it was not exempt from such requirement since it was in possession of a person or persons not authorized to dispense a prescription drug; and 503(b)(4)—the label of the article failed to bear the statement "Caution—Federal law prohibits dispensing without prescription."

DISPOSITION: 8-18-59. Default—destruction.