RECALL NUMBER	PRODUCT	LOT(S)	MFR. & RECALLER	REASON	TYPE	CLASS	OTC or Rx
D-399-4	Digoxin 0.25 mg. Tablet	1	Heather Drug Co. Inc. Cherry Hill, N.J.	Product failed USP Dissolution test and requirements set forth in the FR 1-22-74	FDA Init	II	Rx
D-401-4	Digoxin 0.25 mg. Tablets	9	Heather Drug Co. Cherry Hill, N.J.	Product fails USP Dissolution test & requirements set forth in the FR dated 1-22-74	FDA Init.	II	Rx
D-402-4	Sodium Phenobarbital Inj. USP 130 mg (2 gr) per cc.	1	Wyeth Labs Inc. Marietta, Pa. recaller: Wyeth Labs Inc. Radnor, Pa.	Mislabeled product. one or more units of Sodium Phenobarbital Inj. labeled as Sparine 50 mg. per cc.	Vol.	II	Rx
D-410-4	Haemolyte Dialysis Concentrate #7 (Special Formula)	1	Nutrilite Products Inc. Buena Park, Ca. recalling through its subsidiary Tera Pharmaceuticals Inc. Buena Park, Ca.	Distributed without analysis after reworking.	Vol.	II	Rx
D-414-4	Alcohol Absolute	1	Recalled: Abbott Labs. N. Chicago, Ill.	Product Exceeds the USP limit for non-volatile substances in ethyl alcohol.	Vol.	II	Rx
D-418-4	Kelfin (Sodium cephalothin)	1	Eli Lilly & Co. Indianapolis, Ind.	Mislabeling - a bulk package labeled as Loridine actually contained Keflin.	Vol.	II	Rx
D-419-4	Loridine (Cephaloridine)	1	Eli Lilly & Co. Indianapolis, Ind.	Mislabeling. A bulk package labeled as Keflin actually contained Loridine.	Vol.	II	Rx
D-429-4	Digoxin Tablets USP 0.25 mg.	1	Cord Labs. Inc. Detroit, Michigan	Product failed USP Dissolution test and requirements set forth in the FR 1-22-74	FDA Init.	II	Rx
D-430-4	Radiocaps - 131. Sodium Iodide I-131 Capsules packed 20/ caps/carton	1	Abbott Labs. N. Chicago, Ill.	Erroneous labeling re: expiration date	Vol.	II	Rx