"What Kind of Games Are Being Played?", editorial from California Pharmacist, page 9, November 1973	Page
"Some Manufacturers Disclose Sources of Supply" article from California	10167
"Manufacturer Disclosures Pert II" article from Colifornia Pharmanist	. 10168
Letter dated February 25, 1974, to Senator Gaylord Nelson, chairman Subcommittee on Monopoly, from Dr. Edward G. Feldmann, associate executive director for scientific affairs. American Pharmaceutical	. 10171
Letter dated February 22, 1974, to Dr. Edward G. Feldmann, associate executive director for scientific affairs, American Pharmaceutical Association, from Senator Gaylord Nelson, chairman, Subcommittee or	10184
Monopoly  Letter dated February 25, 1974, to Senator Gaylord Nelson, chairman, Subcommittee on Monopoly, from Dr. Edward G. Feldmann, associate executive director for scientific affairs. American Pharmaceutical	10205
Association  Letter dated February 27, 1974, to Senator Gaylord Nelson, chairman, Subcommittee on Monopoly, from Dr. Edward G. Feldmann, associate executive director for scientific affairs, American Pharmaceutical	10206
Association Statements appearing in Pharmaceutical Manufacturers Association publications in connection with their claim regarding responsibility for	10211
95 percent of the U.S. drug supply  Membership list of the National Association of Pharmaceutical Manuface	10224
turers as of January 1974.  Letter dated February 26, 1974, to Senator Gaylord Nelson, chairman, Subcommittee on Monopoly, from Joseph Barrows, chairman of the board National Association, of Pharmacettical Maria	10248
Subcommittee on Monopoly, from Joseph Barrows, chairman of the board, National Association of Pharmaceutical Manufacturers	10255
1973, for the period January 1 through December 31, 1974	10459
APPENDIX	•
Exhibits provided by the U.S. General Accounting Office:  Prepared statement of Hon. Elmer B. Staats, Comptroller General of the United States	10497
"Problems In Obtaining And Enforcing Compliance With Good Manufacturing Practices For Drugs," a report submitted to the U.S. Congress by the Comptroller General of the United States R.	
164031(2), March 29, 1973  "How to Improve the Procurement and Supply of Drugs in the Federal Government," a report submitted to the U.S. Congress by the Comptroller General of the United States, B-164031(2), December	10520
6, 1973Exhibits provided by the Food and Drug Administration:	10577
Prepared statement of Hon Alexander M Schmidt M D. Commis-	10638
Orige compliance information letter, dated February 28, 1973, from the Office of Compliance. Bureau of Drugs. Food and Drug Administra-	
Drug compliance information letter, dated May 7, 1973, from the	10657
omce of Compliance, Bureau of Drugs, Food and Drug Administration, to manufacturers, repackers, and relabelers of drug products	10660
Office of Compliance, Bureau of Drugs, Food and Drug Administra-	•
tion, to manufacturers, repackers, and relabelers of drug products	10000
Drug compliance information letter, dated October 5, 1973, from the	10003
Orug compliance information letter, dated October 5, 1973, from the Office of Compliance, Bureau of Drugs, Food and Drug Administration, to manufacturers, repackers, and relabelers of drug products	
Office of Compliance, Bureau of Drugs, Food and Drug Administration, to manufacturers, repackers, and relabelers of drug products  Drug compliance information letter, dated January 23, 1974, from the Office of Compliance, Bureau of Drugs, Food and Drug Administration, to manufacturers, repackers, and relabelers of drug products	10666
Office of Compliance, Bureau of Drugs, Food and Drug Administration, to manufacturers, repackers, and relabelers of drug products  Drug compliance information letter, dated January 23, 1974, from the Office of Compliance. Bureau of Drugs, Food and Drug Administra-	10666 10669