FDA 510(k) Premarket Notification

CLEARED

K911724 ACA(R) URINE DRUGS OF ABUSE CONTROL, MODIFICATION

May 22, 1991
35 days to decision

K911724 · Product code: **DIF** · Toxicology
Source: https://www.510kdatabase.net/k911724/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Apr 17, 1991
Decision date	May 22, 1991
Days to decision	35 days
Third-party review	No

APPLICANT

Company	**E.I. Dupont DE Nemours & Co., Inc.**
Location	Mchenry, IL, US
Contact	CHRISTOPHER BENTSEN
510(k) history	253 submissions · 252 cleared · 1976-1996