FDA 510(k) Premarket Notification

K802672 CARDIOACTIVE DRUGS CONTROL

CLEARED

Nov 24, 1980
28 days to decision

K802672 · Product code: **DIF** · Toxicology
Source: https://www.510kdatabase.net/k802672/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Oct 27, 1980
Decision date	Nov 24, 1980
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Fisher Scientific Co., LLC**
Location	Mchenry, IL, US
510(k) history	89 submissions · 89 cleared · 1976-1993