FDA 510(k) Premarket Notification

K800883 CARDIACS I TOXICOLOGY CONTROL

CLEARED

Apr 29, 1980
12 days to decision

K800883 · Product code: **DIF** · Toxicology
Source: https://www.510kdatabase.net/k800883/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Apr 17, 1980
Decision date	Apr 29, 1980
Days to decision	12 days
Third-party review	No

APPLICANT

Company	**Utak Laboratories, Inc.**
Location	Mchenry, IL, US
510(k) history	23 submissions · 23 cleared · 1980-1991