FDA 510(k) Premarket Notification

K830494 CONTOX DRUG SCREEN CONTROL

CLEARED

Mar 11, 1983
23 days to decision

K830494 · Product code: **DIF** · Toxicology
Source: https://www.510kdatabase.net/k830494/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Feb 16, 1983
Decision date	Mar 11, 1983
Days to decision	23 days
Third-party review	No

APPLICANT

Company	**Kaulson Laboratories, Inc.**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1983-1983