FDA 510(k) Premarket Notification

K810235 ACETAMINOPHEN/SALICYLATE CONTROL

CLEARED

Feb 10, 1981
13 days to decision

K810235 · Product code: **DIF** · Toxicology
Source: https://www.510kdatabase.net/k810235/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Jan 28, 1981
Decision date	Feb 10, 1981
Days to decision	13 days
Third-party review	No

APPLICANT

Company	**Clin Tech Diagnostics Corp.**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1980-1982