FDA 510(k) Premarket Notification

K831278 LIQUI-SERA ANTICONVULSANTS CONTROLS

CLEARED

May 27, 1983
38 days to decision

K831278 · Product code: **DIF** · Toxicology
Source: https://www.510kdatabase.net/k831278/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Apr 19, 1983
Decision date	May 27, 1983
Days to decision	38 days
Third-party review	No

APPLICANT

Company	**Biodiagnostic Intl**
Location	Mchenry, IL, US
510(k) history	11 submissions · 11 cleared · 1983-2004

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k831278/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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