FDA 510(k) Premarket Notification

K840044 SDI PREPAX DILUKIT 29-1000

CLEARED

Mar 9, 1984
63 days to decision

K840044 · Product code: **JJO** · Toxicology
Source: https://www.510kdatabase.net/k840044/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Flame Emission Photometer For Clinical Use (JJO)
Date received	Jan 6, 1984
Decision date	Mar 9, 1984
Days to decision	63 days
Third-party review	No

APPLICANT

Company	**Scientific Distributors, Inc.**
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1984-1989