FDA 510(k) Premarket Notification

K830564 FEB TEST

CLEARED

Mar 29, 1983
34 days to decision

K830564 · Product code: **CFM** · Chemistry
Source: https://www.510kdatabase.net/k830564/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bathophenanthroline, Colorimetry, Iron (non-heme) (CFM)
Date received	Feb 23, 1983
Decision date	Mar 29, 1983
Days to decision	34 days
Third-party review	No

APPLICANT

Company	**Wako Chemicals USA, Inc.**
Location	Mchenry, IL, US
510(k) history	124 submissions · 123 cleared · 1981-2011