FDA 510(k) Premarket Notification

K810355 ALT (UV) TEST

CLEARED

Feb 23, 1981
13 days to decision

K810355 · Product code: **CKA** · Chemistry
Source: https://www.510kdatabase.net/k810355/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nadh Oxidation/nad Reduction, Alt/sgpt (CKA)
Date received	Feb 10, 1981
Decision date	Feb 23, 1981
Days to decision	13 days
Third-party review	No

APPLICANT

Company	**Boehringer Mannheim Corp.**
Location	Mchenry, IL, US
510(k) history	340 submissions · 340 cleared · 1976-1999