FDA 510(k) Premarket Notification

CLEARED

K820391 ABUSCREEN RADIOIMMUNOASSAY FOR CANNABI

Mar 8, 1982
26 days to decision

K820391 · Product code: **LAT** · Toxicology
Source: https://www.510kdatabase.net/k820391/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Cannabinoid(s) (LAT)
Date received	Feb 10, 1982
Decision date	Mar 8, 1982
Days to decision	26 days
Third-party review	No

APPLICANT

Company	**Hoffmann-La Roche, Inc.**
Location	Mchenry, IL, US
Website	https://www.roche.com
510(k) history	49 submissions · 49 cleared · 1976-1985