FDA 510(k) Premarket Notification

K890887 ABUSCREEN FP FOR CANNABINOIDS

CLEARED

Mar 16, 1989
23 days to decision

K890887 · Product code: **LDJ** · Toxicology
Source: https://www.510kdatabase.net/k890887/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enzyme Immunoassay, Cannabinoids (LDJ)
Date received	Feb 21, 1989
Decision date	Mar 16, 1989
Days to decision	23 days
Third-party review	No
Combination product	No
PCCP authorized	No

APPLICANT

Company	**Roche Diagnostic Systems, Inc.**
Location	Mchenry, IL, US
Contact	ALEX WESOLOWSKI
510(k) history	296 submissions · 296 cleared · 1983-1999