FDA 510(k) Premarket Notification

K925547 MAXSOFTWARE

CLEARED

Jan 26, 1993
84 days to decision

K925547 · Product code: **JQW** · Toxicology
Source: https://www.510kdatabase.net/k925547/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Station, Pipetting And Diluting, For Clinical Use (JQW)
Date received	Nov 3, 1992
Decision date	Jan 26, 1993
Days to decision	84 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

APPLICANT

Company	**Diagnostic Products Corp.**
Location	Mchenry, IL, US
Contact	KENNETH B ASARCH
510(k) history	321 submissions · 321 cleared · 1976-2006