FDA 510(k) Premarket Notification

K812983 DSD

CLEARED

Nov 10, 1981
15 days to decision

K812983 · Product code: **JQW** · Chemistry
Source: https://www.510kdatabase.net/k812983/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Station, Pipetting And Diluting, For Clinical Use (JQW)
Date received	Oct 26, 1981
Decision date	Nov 10, 1981
Days to decision	15 days
Third-party review	No

APPLICANT

Company	**Pm America, Inc.**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1980-1983

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k812983/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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