FDA 510(k) Premarket Notification

K830319 AUTO PREP

CLEARED

Mar 29, 1983
57 days to decision

K830319 · Product code: **JQW** · Chemistry
Source: https://www.510kdatabase.net/k830319/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Station, Pipetting And Diluting, For Clinical Use (JQW)
Date received	Jan 31, 1983
Decision date	Mar 29, 1983
Days to decision	57 days
Third-party review	No

APPLICANT

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