

K830319 AUTO PREPMar 29, 1983
57 days to decisionK830319 · 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

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Device record: <https://www.510kdatabase.net/k830319/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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