

K821034 SKATRON REPEATING DISPENSERMay 3, 1982
20 days to decisionK821034 · Product code: **JQW** · Chemistry
Source: <https://www.510kdatabase.net/k821034/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Station, Pipetting And Diluting, For Clinical Use (JQW)
Date received	Apr 13, 1982
Decision date	May 3, 1982
Days to decision	20 days
Third-party review	No

APPLICANT

Company	Skatron, Inc.
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1981-1982

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k821034/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 2, 2026