FDA 510(k) Premarket Notification

K821034 SKATRON REPEATING DISPENSER

CLEARED

May 3, 1982
20 days to decision

K821034 · 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