FDA 510(k) Premarket Notification

K831065 SECTIONING AID KIT

CLEARED

Jun 8, 1983
65 days to decision

K831065 · Product code: **KEW** · Pathology
Source: https://www.510kdatabase.net/k831065/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Slides, Microscope (KEW)
Date received	Apr 4, 1983
Decision date	Jun 8, 1983
Days to decision	65 days
Third-party review	No

APPLICANT

Company	**Technicon Instruments Corp.**
Location	Mchenry, IL, US
510(k) history	157 submissions · 156 cleared · 1976-1991

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k831065/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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