FDA 510(k) Premarket Notification

K883749 PROBE(TM) MODELS 700 AND 1000

CLEARED

Nov 10, 1988
70 days to decision

K883749 · Product code: **JQW** · Chemistry
Source: https://www.510kdatabase.net/k883749/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Station, Pipetting And Diluting, For Clinical Use (JQW)
Date received	Sep 1, 1988
Decision date	Nov 10, 1988
Days to decision	70 days
Third-party review	No

APPLICANT

Company	**Packard Instrument Co., Inc.**
Location	Downers Grove, IL, US
Contact	GEORGE L RUSSELL
510(k) history	4 submissions · 4 cleared · 1988-1992

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k883749/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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