FDA 510(k) Premarket Notification

K900130 MODIFIED SEISMOCARDIOGRAPH

CLEARED

Mar 28, 1990
82 days to decision

K900130 · Product code: **DSB** · Cardiovascular
Source: https://www.510kdatabase.net/k900130/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plethysmograph, Impedance (DSB)
Date received	Jan 5, 1990
Decision date	Mar 28, 1990
Days to decision	82 days
Third-party review	No

APPLICANT

Company	**Seismed Instruments, Inc.**
Location	Minneapolis, MN, US
Contact	SANDRA J GARLOUGH
510(k) history	3 submissions · 3 cleared · 1989-1992

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k900130/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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