FDA 510(k) Premarket Notification

K905056 PERCOR STAT-DL 9.5 FR W/OPT INSERT W/O INTRO SHEAT

CLEARED

Jun 11, 1991
215 days to decision

K905056 · Product code: **DSP** · Cardiovascular
Source: https://www.510kdatabase.net/k905056/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Balloon, Intra-aortic And Control (DSP)
Date received	Nov 8, 1990
Decision date	Jun 11, 1991
Days to decision	215 days
Third-party review	No

APPLICANT

Company	**Datascope Corp.**
Location	Mchenry, IL, US
Contact	ROBERT B SCHOCK
510(k) history	136 submissions · 135 cleared · 1976-2019

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k905056/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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