

**K893232 MODEL V420 DIRECT PRESSURE MONITOR & 427
WAVEFORM**Jul 14, 1989
77 days to decisionK893232 · Product code: **DSK** · Cardiovascular
Source: <https://www.510kdatabase.net/k893232/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Blood-pressure (DSK)
Date received	Apr 28, 1989
Decision date	Jul 14, 1989
Days to decision	77 days
Third-party review	No

APPLICANT

Company	Camino Laboratories, Inc.
Location	Anaheim, CA, US
Contact	HAZEL SHUMWAY
510(k) history	18 submissions · 18 cleared · 1982-1995

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k893232/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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