

K863042 SIRECUST 888/888R NON-INVASIVE BLOOD PRESS MONITORJan 14, 1987
155 days to decisionK863042 · Product code: **DSK** · Cardiovascular
Source: <https://www.510kdatabase.net/k863042/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Blood-pressure (DSK)
Date received	Aug 12, 1986
Decision date	Jan 14, 1987
Days to decision	155 days
Third-party review	No

APPLICANT

Company	Siemens Medical Solutions USA, Inc.
Location	Hoffman Estates, IL, US
Contact	ANDY LEVY
510(k) history	778 submissions · 778 cleared · 1980-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k863042/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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