

**K885154 INTERSEPT* CRYSTALLOID CARDIOPLEGIA DELIV.
SYSTEM**Feb 22, 1989
69 days to decisionK885154 · Product code: **KRJ** · Cardiovascular
Source: <https://www.510kdatabase.net/k885154/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filter, Prebypass, Cardiopulmonary Bypass (KRJ)
Date received	Dec 15, 1988
Decision date	Feb 22, 1989
Days to decision	69 days
Third-party review	No

APPLICANT

Company	Medtronic Blood Systems, Inc.
Location	Anaheim, CA, US
Contact	SHARON ROCKWELL
510(k) history	15 submissions · 15 cleared · 1988-1995

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k885154/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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