

**K892222 9.5F PERCU DOUBLE LUMEN INTRA-AORTIC
BALLOON**Sep 14, 1990
529 days to decisionK892222 · Product code: **DSP** · Cardiovascular
Source: <https://www.510kdatabase.net/k892222/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Balloon, Intra-aortic And Control (DSP)
Date received	Apr 3, 1989
Decision date	Sep 14, 1990
Days to decision	529 days
Third-party review	No

APPLICANT

Company	Kontron Instruments, Inc.
Location	Mchenry, IL, US
Contact	DAVID CROMWICK
510(k) history	57 submissions · 57 cleared · 1981-1993

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k892222/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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