

K831759 INTRACARDIA SUCKERSAug 12, 1983
72 days to decisionK831759 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k831759/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Jun 1, 1983
Decision date	Aug 12, 1983
Days to decision	72 days
Third-party review	No

APPLICANT

Company	Research Industries Corp.
Location	Walker, MI, US
510(k) history	5 submissions · 5 cleared · 1983-1995

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Device record: <https://www.510kdatabase.net/k831759/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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