

**K881733 SHILEY FEMORAL CORONARY PERFUSION
CATHETER**Jul 6, 1988
75 days to decisionK881733 · Product code: **DWF** · Cardiovascular
Source: <https://www.510kdatabase.net/k881733/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Apr 22, 1988
Decision date	Jul 6, 1988
Days to decision	75 days
Third-party review	No

APPLICANT

Company	Shiley, Inc.
Location	Mchenry, IL, US
Contact	GANZ POBUDA
510(k) history	174 submissions · 174 cleared · 1976-1993

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k881733/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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