

K820045 CARDIOVASCULAR DEVICESJan 29, 1982
22 days to decisionK820045 · Product code: **DTM** · Cardiovascular
Source: <https://www.510kdatabase.net/k820045/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filter, Blood, Cardiopulmonary Bypass, Arterial Line (DTM)
Date received	Jan 7, 1982
Decision date	Jan 29, 1982
Days to decision	22 days
Third-party review	No

APPLICANT

Company	Bentley Laboratories, Inc.
Location	Mchenry, IL, US
Website	https://www.bentleyinstruments.com
510(k) history	55 submissions · 55 cleared · 1976-1993

Bentley Laboratories, Inc. is located in McHenry, US. The company has a historical record of FDA 510(k) device clearances spanning from 1976 to 1993. Bentley Laboratories received FDA 510(k) clearances from total submissions. The company specialized primarily in Cardiovascular devices, which represented approximately 80% of its regulatory submissions. Notable cleared devices included blood cardioplegia heat exchangers, venous reservoir bags, membrane oxygenators, and central venous catheters used in cardiac surgery and perfusion applications. The company is inactive and s...

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k820045/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 29, 2026