

K823272 VENOUS RESERVOIR BAG #BMR-1400Dec 30, 1982
58 days to decisionK823272 · Product code: **DTN** · CardiovascularSource: <https://www.510kdatabase.net/k823272/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Reservoir, Blood, Cardiopulmonary Bypass (DTN)
Date received	Nov 2, 1982
Decision date	Dec 30, 1982
Days to decision	58 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...