

K802714 COBE/STOCKERT BATTERY MODULENov 26, 1980
26 days to decisionK802714 · Product code: **DWB** · CardiovascularSource: <https://www.510kdatabase.net/k802714/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Blood, Cardiopulmonary Bypass, Roller Type (DWB)
Date received	Oct 31, 1980
Decision date	Nov 26, 1980
Days to decision	26 days
Third-party review	No

APPLICANT

Company	Cobe Laboratories, Inc.
Location	Mchenry, IL, US
Website	https://www.gambro.com
510(k) history	77 submissions · 77 cleared · 1976-1993

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k802714/>; 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 May 26, 2026