

K831422 BIOFLOWSep 12, 1983
132 days to decisionK831422 · Product code: **DSB** · Cardiovascular
Source: <https://www.510kdatabase.net/k831422/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plethysmograph, Impedance (DSB)
Date received	May 3, 1983
Decision date	Sep 12, 1983
Days to decision	132 days
Third-party review	No

APPLICANT

Company	Bomed Medical Mfr., Ltd.
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1981-1986

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k831422/> 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 July 1, 2026