

K830230 PROGRAMMABLE VENTILATORFeb 9, 1983
12 days to decisionK830230 · Product code: **CBK** · Anesthesiology
Source: <https://www.510kdatabase.net/k830230/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Continuous, Facility Use (CBK)
Date received	Jan 28, 1983
Decision date	Feb 9, 1983
Days to decision	12 days
Third-party review	No

APPLICANT

Company	Bio-Med Devices, Inc.
Location	Mchenry, IL, US
510(k) history	14 submissions · 13 cleared · 1980-2022

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Device record: <https://www.510kdatabase.net/k830230/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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