FDA 510(k) Premarket Notification

K830230 PROGRAMMABLE VENTILATOR

CLEARED

Feb 9, 1983
12 days to decision

K830230 · 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