

**K870835 ADULT APNEA MONITOR (20), MONITOR INTERFACE
(30)**Mar 27, 1987
24 days to decisionK870835 · Product code: **BZQ** · Anesthesiology
Source: <https://www.510kdatabase.net/k870835/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Breathing Frequency (BZQ)
Date received	Mar 3, 1987
Decision date	Mar 27, 1987
Days to decision	24 days
Third-party review	No

APPLICANT

Company	Applied Membrane Technology, Inc.
Location	Minnetonka, MN, US
Contact	STEPHEN P CONOVER
510(k) history	3 submissions · 3 cleared · 1985-1987

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k870835/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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