FDA 510(k) Premarket Notification

K870835 ADULT APNEA MONITOR (20), MONITOR INTERFACE (30)

CLEARED

Mar 27, 1987
24 days to decision

K870835 · 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