FDA 510(k) Premarket Notification

K853303 LIFELINC LEADWIRE

CLEARED

Sep 18, 1985
43 days to decision

K853303 · Product code: **BZQ** · Anesthesiology
Source: https://www.510kdatabase.net/k853303/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Breathing Frequency (BZQ)
Date received	Aug 6, 1985
Decision date	Sep 18, 1985
Days to decision	43 days
Third-party review	No

APPLICANT

Company	**Tronomed, Inc.**
Location	Mchenry, IL, US
Contact	JIM LELTEN
510(k) history	12 submissions · 12 cleared · 1977-1996