FDA 510(k) Premarket Notification

K813507 INFALERT

CLEARED

Jan 12, 1982
28 days to decision

K813507 · Product code: **BZQ** · Anesthesiology
Source: https://www.510kdatabase.net/k813507/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Breathing Frequency (BZQ)
Date received	Dec 15, 1981
Decision date	Jan 12, 1982
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Briox Technologies, Inc.**
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1979-1982