FDA 510(k) Premarket Notification

K881235 Q-PLEX 1

CLEARED

Mar 31, 1989
374 days to decision

K881235 · Product code: **BZC** · Anesthesiology
Source: https://www.510kdatabase.net/k881235/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Calculator, Pulmonary Function Data (BZC)
Date received	Mar 22, 1988
Decision date	Mar 31, 1989
Days to decision	374 days
Third-party review	No

APPLICANT

Company	**Quinton, Inc.**
Location	Mchenry, IL, US
Contact	RON R DUCK
510(k) history	164 submissions · 160 cleared · 1976-2003