

K881235 Q-PLEX 1Mar 31, 1989
374 days to decisionK881235 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k881235/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 20, 2026