

K832355 CRASH CARTOct 28, 1983
102 days to decisionK832355 · Product code: **BZN** · Anesthesiology
Source: <https://www.510kdatabase.net/k832355/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cart, Emergency, Cardiopulmonary (excluding Equipment) (BZN)
Date received	Jul 18, 1983
Decision date	Oct 28, 1983
Days to decision	102 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k832355/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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