FDA 510(k) Premarket Notification

K832355 CRASH CART

CLEARED

Oct 28, 1983
102 days to decision

K832355 · 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