FDA 510(k) Premarket Notification

K981939 AUTOCORR PLUS

CLEARED

Aug 20, 1998
79 days to decision

K981939 · Product code: **DQA** · Anesthesiology
Source: https://www.510kdatabase.net/k981939/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Jun 2, 1998
Decision date	Aug 20, 1998
Days to decision	79 days
Third-party review	No
Summary / Statement	Summary
Other names	3404 OXIMETER/ECG MONITOR

APPLICANT

Company	**Bci Intl., Inc.**
Location	Waukesha, WI, US
Contact	DONALD J ALEXANDER
510(k) history	20 submissions · 20 cleared · 1994-1999