FDA 510(k) Premarket Notification

K020739 BCI 3178 PEDIATRIC PULSE OXIMETRY SENSOR

CLEARED

Jun 3, 2002
89 days to decision

K020739 · Product code: **DQA** · Anesthesiology
Source: https://www.510kdatabase.net/k020739/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Mar 6, 2002
Decision date	Jun 3, 2002
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Bci, Inc.**
Location	Mchenry, IL, US
Contact	DONALD J ALEXANDER
510(k) history	13 submissions · 13 cleared · 1979-2003