FDA 510(k) Premarket Notification

K833826 BTI BIOX III OXIMETER CRITICAL CAREUNI

CLEARED

Dec 30, 1983
65 days to decision

K833826 · Product code: **DQA** · Anesthesiology
Source: https://www.510kdatabase.net/k833826/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Oct 26, 1983
Decision date	Dec 30, 1983
Days to decision	65 days
Third-party review	No

APPLICANT

Company	**Biox Technology, Inc.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1982-1983