FDA 510(k) Premarket Notification

K823563 EEG & RESPIRATION MODULES POLYGRAPH

CLEARED

Jan 12, 1983
41 days to decision

K823563 · Product code: **BZQ** · Anesthesiology
Source: https://www.510kdatabase.net/k823563/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Breathing Frequency (BZQ)
Date received	Dec 2, 1982
Decision date	Jan 12, 1983
Days to decision	41 days
Third-party review	No

APPLICANT

Company	**Nihon Kohden America, Inc.**
Location	Foothill Ranch, CA, US
510(k) history	166 submissions · 163 cleared · 1979-2012