FDA 510(k) Premarket Notification

K030105 ANESTHESIA MONITOR

CLEARED

Mar 4, 2003
50 days to decision

K030105 · Product code: **MHX** · Cardiovascular
Source: https://www.510kdatabase.net/k030105/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Jan 13, 2003
Decision date	Mar 4, 2003
Days to decision	50 days
Third-party review	No
Summary / Statement	Summary
Other names	BEDSIDE MONITOR, SERIES BSM-5130A

APPLICANT

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