

K030105 ANESTHESIA MONITORMar 4, 2003
50 days to decisionK030105 · Product code: **MHX** · CardiovascularSource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k030105/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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