

**K823563 EEG & RESPIRATION MODULES POLYGRAPH**Jan 12, 1983  
41 days to decisionK823563 · Product code: **BZQ** · AnesthesiologySource: <https://www.510kdatabase.net/k823563/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k823563/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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