

**K802205 ECG/RESPIRATION STIMULATOR**Oct 3, 1980  
22 days to decisionK802205 · Product code: **BZQ** · AnesthesiologySource: <https://www.510kdatabase.net/k802205/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Breathing Frequency (BZQ)
Date received	Sep 11, 1980
Decision date	Oct 3, 1980
Days to decision	22 days
Third-party review	No

**APPLICANT**

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Company	<b>Healthdyne, Inc.</b>
Location	Mchenry, IL, US
510(k) history	35 submissions · 35 cleared · 1978-1995

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

Device record: <https://www.510kdatabase.net/k802205/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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