

**K812039 SPIROMETER 210/211**Sep 25, 1981  
66 days to decisionK812039 · Product code: **BZG** · Anesthesiology  
Source: <https://www.510kdatabase.net/k812039/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Diagnostic (BZG)
Date received	Jul 21, 1981
Decision date	Sep 25, 1981
Days to decision	66 days
Third-party review	No

**APPLICANT**

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Company	<b>Medix Medical Electronics (U.S.A.), Inc.</b>
Location	Mchenry, IL, US
510(k) history	14 submissions · 14 cleared · 1981-1986

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Device record: <https://www.510kdatabase.net/k812039/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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