

**K782145 SPIROMETER, MINI POSITIVE**Jan 31, 1979  
35 days to decisionK782145 · Product code: **BZK** · Anesthesiology  
Source: <https://www.510kdatabase.net/k782145/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Monitoring (w/wo Alarm) (BZK)
Date received	Dec 27, 1978
Decision date	Jan 31, 1979
Days to decision	35 days
Third-party review	No

**APPLICANT**

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Company	<b>Boehringer Laboratories</b>
Location	Mchenry, IL, US
Website	<a href="http://www.boehringerlabs.com">http://www.boehringerlabs.com</a>
510(k) history	38 submissions · 38 cleared · 1976-2024

Boehringer Laboratories is a family-owned American medical technology company headquartered in Phoenixville, Pennsylvania, with operations in McHenry, US. The company specializes in respiratory therapy and minimally invasive surgical devices. Boehringer Laboratories has maintained a strong FDA 510(k) regulatory record since 1976. The company has received FDA 510(k) clearances from total submissions, with no denied submissions. Recent clearances span 2024, demonstrating continued active development. The company's portfolio focuses primarily on anesthesiology devices, inclu...

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

Device record: <https://www.510kdatabase.net/k782145/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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