

**K770141 INSPIROMETER RESPIRATORY EXERCISER**Feb 1, 1977  
8 days to decisionK770141 · Product code: **BWF** · AnesthesiologySource: <https://www.510kdatabase.net/k770141/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Therapeutic (incentive) (BWF)
Date received	Jan 24, 1977
Decision date	Feb 1, 1977
Days to decision	8 days
Third-party review	No

**APPLICANT**

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Company	<b>Medi, Inc.</b>
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1977-1977

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

Device record: <https://www.510kdatabase.net/k770141/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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