

**K882604 PULMO-PATTY**Jan 4, 1989  
194 days to decisionK882604 · Product code: **BYI** · Anesthesiology  
Source: <https://www.510kdatabase.net/k882604/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Percussor, Powered-electric (BYI)
Date received	Jun 24, 1988
Decision date	Jan 4, 1989
Days to decision	194 days
Third-party review	No

**APPLICANT**

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Company	<b>Tower Stopper, Inc.</b>
Location	Hancock, MN, US
Contact	ARTHUR NOHL
510(k) history	1 submissions · 1 cleared · 1989-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k882604/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 27, 2026