

**K931988 3P (PULMONARY PERCUSSIVE PACK)**Oct 28, 1994  
554 days to decisionK931988 · Product code: **BYI** · Anesthesiology  
Source: <https://www.510kdatabase.net/k931988/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Percussor, Powered-electric (BYI)
Date received	Apr 22, 1993
Decision date	Oct 28, 1994
Days to decision	554 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Delta Medical Manuf, Inc.</b>
Location	Raleigh, NC, US
Contact	DEN RAVELY
510(k) history	1 submissions · 1 cleared · 1994-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k931988/> 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 26, 2026