

**K883899 MANUAL PERCUSSOR CUPS**Dec 13, 1988  
89 days to decisionK883899 · Product code: **BYI** · AnesthesiologySource: <https://www.510kdatabase.net/k883899/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Percussor, Powered-electric (BYI)
Date received	Sep 15, 1988
Decision date	Dec 13, 1988
Days to decision	89 days
Third-party review	No

**APPLICANT**

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Company	<b>Medical Components, Inc.</b>
Location	Mchenry, IL, US
Contact	DUNCAN G JOHNSON
510(k) history	63 submissions · 55 cleared · 1980-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k883899/>; 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