

**K883806 ROCKET INTRAUTERINE SOUND**Oct 31, 1988  
54 days to decisionK883806 · Product code: **HHM** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k883806/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sound, Uterine (HHM)
Date received	Sep 7, 1988
Decision date	Oct 31, 1988
Days to decision	54 days
Third-party review	No

**APPLICANT**

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Company	<b>A &amp; A Medical, Inc.</b>
Location	Branford, CT, US
Contact	K BERNBERG
510(k) history	23 submissions · 23 cleared · 1988-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k883806/>; 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 June 29, 2026