

**K896357 SAFE AID DEVICE IMPOTENCE DEVICE**Apr 25, 1990  
170 days to decisionK896357 · Product code: **LKY** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k896357/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, External Penile Rigidity (LKY)
Date received	Nov 6, 1989
Decision date	Apr 25, 1990
Days to decision	170 days
Third-party review	No

**APPLICANT**

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Company	<b>Dr. John P. March</b>
Location	Sacramento, CA, US
Contact	JOHN P MARCH
510(k) history	1 submissions · 1 cleared · 1990-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896357/>; 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 July 2, 2026