

**K011781 INTERCEPT-URETHRAL MICROCOIL**Aug 31, 2001  
85 days to decisionK011781 · Product code: **MOS** · Radiology  
Source: <https://www.510kdatabase.net/k011781/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Coil, Magnetic Resonance, Specialty (MOS)
Date received	Jun 7, 2001
Decision date	Aug 31, 2001
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Surgi-Vision, Inc.</b>
Location	Columbia, MD, US
Contact	NANCY E TAYLOR
510(k) history	9 submissions · 9 cleared · 1999-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k011781/> 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 17, 2026