

**K020495 INTERCEPT ESOPHAGEAL INTERNAL MR COIL,
INTERCEPT URETHRAL INTERNAL MR COIL, AND INTERCEPT
VASCULAR INTERNAL MR COIL**

Apr 23, 2002
68 days to decision

K020495 · Product code: **MOS** · Radiology
Source: <https://www.510kdatabase.net/k020495/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Coil, Magnetic Resonance, Specialty (MOS)
Date received	Feb 14, 2002
Decision date	Apr 23, 2002
Days to decision	68 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k020495/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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