

**K945572 CODMAN(R) RIDIG AND STEERABLE ENDOSCOPE  
HOLDER, MODEL 83-1352**Feb 3, 1995  
81 days to decisionK945572 · Product code: **GWG** · Neurology  
Source: <https://www.510kdatabase.net/k945572/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscope, Neurological (GWG)
Date received	Nov 14, 1994
Decision date	Feb 3, 1995
Days to decision	81 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Johnson &amp; Johnson Professionals, Inc.</b>
Location	Raynham, MA, US
Contact	PATRICIA BANKERT
Website	<a href="https://www.jnj.com">https://www.jnj.com</a>
510(k) history	206 submissions · 184 cleared · 1976-2000

Johnson & Johnson Professionals, Inc. is a medical device company based in Raynham, Massachusetts. The company specializes in surgical and orthopedic devices. The company has received FDA 510(k) clearances from total submissions between 1976 and 2000. Orthopedic devices and neurosurgical instruments represent core product categories. Notable cleared devices include hip and elbow prostheses, programmable valve systems, and aneurysm clips. The company is inactive and represents a historical regulatory record with no submissions in more than two decades. Explore the complete...

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Device record: <https://www.510kdatabase.net/k945572/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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