

**K780718 SOF-WICKLAP PAD**Jun 28, 1978  
58 days to decisionK780718 · Product code: **GDY** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k780718/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Gauze/sponge, Internal, X-ray Detectable (GDY)
Date received	May 1, 1978
Decision date	Jun 28, 1978
Days to decision	58 days
Third-party review	No

**APPLICANT**

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Company	<b>Johnson &amp; Johnson Professionals, Inc.</b>
Location	Raynham, MA, US
Contact	Jean Grieve
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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