

**K991413 CODMAN ETHISORB DURA PATCH**Mar 13, 2000  
325 days to decisionK991413 · Product code: **GXQ** · Neurology  
Source: <https://www.510kdatabase.net/k991413/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dura Substitute (GXQ)
Date received	Apr 23, 1999
Decision date	Mar 13, 2000
Days to decision	325 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	DEANA BOUSHELL
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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