

**K033395 CODMAN DURAL GRAFT IMPLANT, MODELS 80-1471,  
80-1472, 80-1473, 80-1474, 80-1475, 80-1476, 80-1477, 80-1478,  
80-1480, 80-1**Feb 25, 2004  
125 days to decisionK033395 · Product code: **GXO** · Neurology  
Source: <https://www.510kdatabase.net/k033395/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Strip, Craniosynostosis, Preformed (GXO)
Date received	Oct 23, 2003
Decision date	Feb 25, 2004
Days to decision	125 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Codman &amp; Shurtleff, Inc.</b>
Location	Mchenry, IL, US
Contact	LIZ DOLAN
510(k) history	152 submissions · 151 cleared · 1976-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k033395/> 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 June 15, 2026