

K992591 CODMAN SINGLE AND DOUBLE LUMEN SKULL BOLT KITSOct 15, 1999
74 days to decisionK992591 · Product code: **GWM** · Neurology
Source: <https://www.510kdatabase.net/k992591/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Monitoring, Intracranial Pressure (GWM)
Date received	Aug 2, 1999
Decision date	Oct 15, 1999
Days to decision	74 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Codman & Shurtleff, Inc.
Location	Mchenry, IL, US
Contact	JAMES M FLAHERTY
510(k) history	152 submissions · 151 cleared · 1976-2020

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k992591/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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