

**K790920 INTRACRANIAL PRESSURE MONITORING**Aug 3, 1979  
86 days to decisionK790920 · Product code: **GWM** · Neurology  
Source: <https://www.510kdatabase.net/k790920/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Monitoring, Intracranial Pressure (GWM)
Date received	May 9, 1979
Decision date	Aug 3, 1979
Days to decision	86 days
Third-party review	No

**APPLICANT**

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Company	<b>Codman &amp; Shurtleff, Inc.</b>
Location	Mchenry, IL, US
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/k790920/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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