

**K071505 MODIFICATION TO NEUROSENSOR CEREBRAL  
BLOOD FLOW AND INTRACRANIAL PRESSURE MONITORING  
SYSTEM**Aug 6, 2007  
66 days to decisionK071505 · Product code: **GWM** · Neurology  
Source: <https://www.510kdatabase.net/k071505/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Monitoring, Intracranial Pressure (GWM)
Date received	Jun 1, 2007
Decision date	Aug 6, 2007
Days to decision	66 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Integra LifeSciences Corporation</b>
Location	Planisboro, NJ, US
Contact	JON CAPAROTTA
510(k) history	65 submissions · 65 cleared · 2004-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k071505/>; 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 May 15, 2026