

**K840999 PHILLY INFANT BOLT**Apr 4, 1984  
26 days to decisionK840999 · Product code: **GWM** · Neurology  
Source: <https://www.510kdatabase.net/k840999/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Monitoring, Intracranial Pressure (GWM)
Date received	Mar 9, 1984
Decision date	Apr 4, 1984
Days to decision	26 days
Third-party review	No

**APPLICANT**

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Company	<b>Philadelphia Medical Specialties</b>
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1981-1984

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k840999/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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