

K813440 PHILLY BOLTDec 18, 1981
10 days to decisionK813440 · Product code: **GWM** · Neurology
Source: <https://www.510kdatabase.net/k813440/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Monitoring, Intracranial Pressure (GWM)
Date received	Dec 8, 1981
Decision date	Dec 18, 1981
Days to decision	10 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k813440/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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