FDA 510(k) Premarket Notification

K813440 PHILLY BOLT

CLEARED

Dec 18, 1981
10 days to decision

K813440 · 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