FDA 510(k) Premarket Notification

K840999 PHILLY INFANT BOLT

CLEARED

Apr 4, 1984
26 days to decision

K840999 · Product code: **GWM** · Neurology
Source: https://www.510kdatabase.net/k840999/

SUBMISSION DETAILS

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

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