FDA 510(k) Premarket Notification

K822636 CLASS II DEVICE FOR I.C.P. MONITORING

CLEARED

Sep 14, 1982
14 days to decision

K822636 · Product code: **GWM** · Neurology
Source: https://www.510kdatabase.net/k822636/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Monitoring, Intracranial Pressure (GWM)
Date received	Aug 31, 1982
Decision date	Sep 14, 1982
Days to decision	14 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k822636/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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