

K822864 INTRACRANIAL PRESSURE MONITORING DEVICENov 16, 1982
50 days to decisionK822864 · Product code: **GWM** · Neurology
Source: <https://www.510kdatabase.net/k822864/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Monitoring, Intracranial Pressure (GWM)
Date received	Sep 27, 1982
Decision date	Nov 16, 1982
Days to decision	50 days
Third-party review	No

APPLICANT

Company	Medex, Inc.
Location	Mchenry, IL, US
510(k) history	48 submissions · 46 cleared · 1977-2005

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k822864/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 29, 2026