

K802149 TM-5 INTRACRANIAL PRESSURE MONITORNov 12, 1980
65 days to decisionK802149 · Product code: **GWM** · Neurology
Source: <https://www.510kdatabase.net/k802149/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Monitoring, Intracranial Pressure (GWM)
Date received	Sep 8, 1980
Decision date	Nov 12, 1980
Days to decision	65 days
Third-party review	No

APPLICANT

Company	Nicolet Biomedical Instruments
Location	Mchenry, IL, US
510(k) history	26 submissions · 26 cleared · 1977-1988

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k802149/> 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