FDA 510(k) Premarket Notification

K802149 TM-5 INTRACRANIAL PRESSURE MONITOR

CLEARED

Nov 12, 1980
65 days to decision

K802149 · 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