FDA 510(k) Premarket Notification

K790920 INTRACRANIAL PRESSURE MONITORING

CLEARED

Aug 3, 1979
86 days to decision

K790920 · Product code: **GWM** · Neurology
Source: https://www.510kdatabase.net/k790920/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Monitoring, Intracranial Pressure (GWM)
Date received	May 9, 1979
Decision date	Aug 3, 1979
Days to decision	86 days
Third-party review	No

APPLICANT

Company	**Codman & Shurtleff, Inc.**
Location	Mchenry, IL, US
510(k) history	152 submissions · 151 cleared · 1976-2020