FDA 510(k) Premarket Notification

K802137 CHEEK EPIDURAL SCREW

CLEARED

Oct 31, 1980
57 days to decision

K802137 · Product code: **GWM** · Neurology
Source: https://www.510kdatabase.net/k802137/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Monitoring, Intracranial Pressure (GWM)
Date received	Sep 4, 1980
Decision date	Oct 31, 1980
Days to decision	57 days
Third-party review	No

APPLICANT

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