

K802137 CHEEK EPIDURAL SCREWOct 31, 1980
57 days to decisionK802137 · 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

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Device record: <https://www.510kdatabase.net/k802137/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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