

K821363 TRANS-QMay 28, 1982
21 days to decisionK821363 · Product code: **GZJ** · Neurology
Source: <https://www.510kdatabase.net/k821363/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	May 7, 1982
Decision date	May 28, 1982
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Chesapeake Medical Assoc.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k821363/> 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 July 2, 2026