

K000852 RENEW NEUROSTIMULATION SYSTEMJan 19, 2001
310 days to decisionK000852 · Product code: **GZF** · Neurology
Source: <https://www.510kdatabase.net/k000852/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Peripheral Nerve, Implanted (pain Relief) (GZF)
Date received	Mar 15, 2000
Decision date	Jan 19, 2001
Days to decision	310 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Advanced Neuromodulation Systems
Location	Plano, TX, US
Contact	RASHMI MOZA
510(k) history	25 submissions · 25 cleared · 1999-2009

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k000852/> 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 June 22, 2026