

K932948 MEDTRONIC MODEL 3625 SCREENER SYSTEMFeb 3, 1994
231 days to decisionK932948 · Product code: **GZB** · Neurology
Source: <https://www.510kdatabase.net/k932948/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Spinal-cord, Implanted (pain Relief) (GZB)
Date received	Jun 17, 1993
Decision date	Feb 3, 1994
Days to decision	231 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medtronic Vascular
Location	Walker, MI, US
Contact	DAVID MUELLER
510(k) history	475 submissions · 453 cleared · 1977-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k932948/> 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 May 14, 2026