

K991522 MICROTARGETING ELECTRODEAug 4, 2000
459 days to decisionK991522 · Product code: **GZL** · Neurology
Source: <https://www.510kdatabase.net/k991522/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Depth (GZL)
Date received	May 3, 1999
Decision date	Aug 4, 2000
Days to decision	459 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	FHC, Inc.
Location	Bowdoinham, ME, US
Contact	FREDERICK HAER
510(k) history	12 submissions · 12 cleared · 2000-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k991522/> 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 27, 2026