FDA 510(k) Premarket Notification

K991522 MICROTARGETING ELECTRODE

CLEARED

Aug 4, 2000
459 days to decision

K991522 · 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