

K063117 MODIFICATION TO DUREPAIR DURA REGENERATION MATRIX

Nov 3, 2006
22 days to decision

K063117 · Product code: **GXQ** · Neurology
Source: <https://www.510kdatabase.net/k063117/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Dura Substitute (GXQ)
Date received	Oct 12, 2006
Decision date	Nov 3, 2006
Days to decision	22 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medtronic Neurosurgery
Location	Goleta, CA, US
Contact	JEFFERY HENDERSON
510(k) history	28 submissions · 27 cleared · 2001-2025

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k063117/>; 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