

K032406 STRYKER RF ELECTRODES AND CANNULAEApr 1, 2004
241 days to decisionK032406 · Product code: **GXI** · Neurology
Source: <https://www.510kdatabase.net/k032406/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe, Radiofrequency Lesion (GXI)
Date received	Aug 4, 2003
Decision date	Apr 1, 2004
Days to decision	241 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Stryker Instruments
Location	Kalamazoo, MI, US
Contact	NICOLE PETTY
510(k) history	73 submissions · 73 cleared · 1994-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k032406/> 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 28, 2026