FDA 510(k) Premarket Notification

K032406 STRYKER RF ELECTRODES AND CANNULAE

CLEARED

Apr 1, 2004
241 days to decision

K032406 · 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