FDA 510(k) Premarket Notification

CLEARED

K071300 MODIFICATION TO SMITH & NEPHEW RF DENERVATION PROBES & RF CANNULAE

Jul 20, 2007
72 days to decision

K071300 · Product code: **GXI** · Neurology
Source: https://www.510kdatabase.net/k071300/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe, Radiofrequency Lesion (GXI)
Date received	May 9, 2007
Decision date	Jul 20, 2007
Days to decision	72 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Smith & Nephew, Inc.**
Location	Mchenry, IL, US
Contact	KATHY REDDIG
Website	http://www.smith-nephew.com/
510(k) history	530 submissions · 517 cleared · 1980-2026

Smith & Nephew, Inc. is a medical technology company focused on repair, regeneration, and replacement of soft and hard tissues. The company operates with a manufacturing facility in McHenry, US. Smith & Nephew has established a significant regulatory track record with the FDA. The company has received FDA 510(k) clearances from total submissions since 1980. Orthopedic devices represent the dominant category, accounting for 71% of submissions. The company remains active, with the latest clearance in 2025. Recent cleared devices reflect a strong focus on orthopedic surgical...