FDA 510(k) Premarket Notification

CLEARED

K063480 STRYKER RF INTRADISCAL ADAPTOR, MODEL 406-750

May 22, 2007
186 days to decision

K063480 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k063480/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 17, 2006
Decision date	May 22, 2007
Days to decision	186 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Stryker Corp.**
Location	Mchenry, IL, US
Contact	JEAN SHEPPARD
510(k) history	124 submissions · 121 cleared · 1976-2023

Stryker Corp. is an American multinational medical technology company headquartered in Portage, Michigan. The company develops and markets surgical equipment, implants, and patient safety technologies used globally across multiple medical specialties. Stryker has received FDA 510(k) clearances from total submissions since its first clearance in 1976. The company maintains active regulatory engagement, with its latest clearance in 2023. Its product portfolio spans orthopedic devices, neurosurgical implants, surgical instruments, and endoscopy systems, reflecting a broad pr...

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