FDA 510(k) Premarket Notification

K844214 COMPARTMENT SYNDROME PRESSURE MONITOR SYS

CLEARED

Apr 4, 1985
155 days to decision

K844214 · Product code: **LXC** · Orthopedic
Source: https://www.510kdatabase.net/k844214/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Pressure, Intracompartmental (LXC)
Date received	Oct 31, 1984
Decision date	Apr 4, 1985
Days to decision	155 days
Third-party review	No

APPLICANT

Company	**Stryker Corp.**
Location	Mchenry, IL, US
Contact	MARY C MASTENBROOK
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...