FDA 510(k) Premarket Notification

K760456 TABLE, HAND SURGI

CLEARED

Nov 2, 1976
77 days to decision

K760456 · Product code: **FWZ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k760456/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Operating Room Accessories Table Tray (FWZ)
Date received	Aug 17, 1976
Decision date	Nov 2, 1976
Days to decision	77 days
Third-party review	No

APPLICANT

Company	**Stryker Corp.**
Location	Mchenry, IL, US
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...