FDA 510(k) Premarket Notification

K894470 STRYKER TRAUMA TRAY

CLEARED

Aug 18, 1989
31 days to decision

K894470 · Product code: **KYZ** · General Hospital
Source: https://www.510kdatabase.net/k894470/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Irrigating (non Dental) (KYZ)
Date received	Jul 18, 1989
Decision date	Aug 18, 1989
Days to decision	31 days
Third-party review	No

APPLICANT

Company	**Stryker Corp.**
Location	Mchenry, IL, US
Contact	SALMAN, PH.D.
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...