FDA 510(k) Premarket Notification

K871816 STRYKERSCOPE TM/ORTHOSCOPE TM/STRYKER MINISCOPE TM

CLEARED

Aug 3, 1987
84 days to decision

K871816 · Product code: **EAY** · Dental
Source: https://www.510kdatabase.net/k871816/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Fiber Optic, Dental (EAY)
Date received	May 11, 1987
Decision date	Aug 3, 1987
Days to decision	84 days
Third-party review	No

APPLICANT

Company	**Stryker Corp.**
Location	Mchenry, IL, US
Contact	HARMON H WOODWORTH
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...