

K940130 STRYKER MODEL 6110 EMERGENCY/POLE STRETCHERMar 10, 1994
66 days to decisionK940130 · Product code: **FPP** · General Hospital
Source: <https://www.510kdatabase.net/k940130/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stretcher, Hand-carried (FPP)
Date received	Jan 3, 1994
Decision date	Mar 10, 1994
Days to decision	66 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Stryker Corp.
Location	Mchenry, IL, US
Contact	SCOTT GIAMBALVO
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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