FDA 510(k) Premarket Notification

K091606 VERTAPLEX HV (SINGLE PACK AND DUAL PACK)

CLEARED

Oct 23, 2009
142 days to decision

K091606 · Product code: **NDN** · Orthopedic
Source: https://www.510kdatabase.net/k091606/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cement, Bone, Vertebroplasty (NDN)
Date received	Jun 3, 2009
Decision date	Oct 23, 2009
Days to decision	142 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Stryker Corp.**
Location	Mchenry, IL, US
Contact	BECKY E DITTY
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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