FDA 510(k) Premarket Notification

CLEARED

K072118 STRYKER VERTAPLEX RADIOPAQUE BONE CEMENT

Dec 7, 2007
128 days to decision

K072118 · Product code: **NDN** · Orthopedic
Source: https://www.510kdatabase.net/k072118/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cement, Bone, Vertebroplasty (NDN)
Date received	Aug 1, 2007
Decision date	Dec 7, 2007
Days to decision	128 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Stryker Corp.**
Location	Mchenry, IL, US
Contact	JEAN SHEPPARD
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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