

K093935 SMITH & NEPHEW FOOTPRINT RCX PK SUTURE ANCHORApr 5, 2010
104 days to decisionK093935 · Product code: **MBI** · Orthopedic
Source: <https://www.510kdatabase.net/k093935/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fastener, Fixation, Nondegradable, Soft Tissue (MBI)
Date received	Dec 22, 2009
Decision date	Apr 5, 2010
Days to decision	104 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Smith & Nephew, Inc.
Location	McHenry, IL, US
Contact	CHRISTINA FLORES
Website	http://www.smith-nephew.com/
510(k) history	530 submissions · 517 cleared · 1980-2026

Smith & Nephew, Inc. is a medical technology company focused on repair, regeneration, and replacement of soft and hard tissues. The company operates with a manufacturing facility in McHenry, US. Smith & Nephew has established a significant regulatory track record with the FDA. The company has received FDA 510(k) clearances from total submissions since 1980. Orthopedic devices represent the dominant category, accounting for 71% of submissions. The company remains active, with the latest clearance in 2025. Recent cleared devices reflect a strong focus on orthopedic surgical...

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Device record: <https://www.510kdatabase.net/k093935/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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