FDA 510(k) Premarket Notification

K080029 MAKO SURGICAL CORP. PATELLOFEMORAL KNEE IMPLANT SYSTEM

CLEARED

May 16, 2008
133 days to decision

K080029 · Product code: **KRR** · Orthopedic
Source: https://www.510kdatabase.net/k080029/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Patello/femoral, Semi-constrained, Cemented, Metal/polymer (KRR)
Date received	Jan 4, 2008
Decision date	May 16, 2008
Days to decision	133 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Mako Surgical Corp.**
Location	Weston, FL, US
Contact	WILLIAM F TAPIA
Website	https://www.stryker.com
510(k) history	33 submissions · 33 cleared · 2005-2026

Mako Surgical Corp. is a medical device manufacturer based in Weston, US. Now part of Stryker, the brand continues to operate under the parent company with a focus on robotic-assisted surgical systems. The company maintains a strong FDA 510(k) regulatory record with FDA 510(k) clearances from total submissions. Orthopedic devices represent 88% of its submission portfolio. Mako's clearance history spans from 2005 to 2026, demonstrating sustained innovation in joint reconstruction and orthopedic surgery applications. Recent cleared devices include total knee and hip applica...

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k080029/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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