FDA 510(k) Premarket Notification

K944752 ORTHOMET RESURFACING FEMORAL COMPONENT

CLEARED

Dec 28, 1994
93 days to decision

K944752 · Product code: **KXA** · Orthopedic
Source: https://www.510kdatabase.net/k944752/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Femoral, Resurfacing (KXA)
Date received	Sep 26, 1994
Decision date	Dec 28, 1994
Days to decision	93 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Orthomet, Inc.**
Location	Plymouth, MN, US
Contact	DAVID A CANNISTRACI
510(k) history	60 submissions · 41 cleared · 1986-1995