FDA 510(k) Premarket Notification

K960269 TITANIUM UNIVERSAL CLAMP

CLEARED

Apr 4, 1996
77 days to decision

K960269 · Product code: **LXT** · Orthopedic
Source: https://www.510kdatabase.net/k960269/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Nail/blade/plate Combination, Multiple Component, Metal Composite (LXT)
Date received	Jan 18, 1996
Decision date	Apr 4, 1996
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Synthes (Usa)**
Location	Mchenry, IL, US
Contact	ANGELA J SILVESTRI
510(k) history	411 submissions · 394 cleared · 1977-2015

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k960269/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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