FDA 510(k) Premarket Notification

K953777 SYNTHES WIRE MOUNT

CLEARED

Mar 8, 1996
207 days to decision

K953777 · Product code: **JDQ** · Orthopedic
Source: https://www.510kdatabase.net/k953777/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cerclage, Fixation (JDQ)
Date received	Aug 14, 1995
Decision date	Mar 8, 1996
Days to decision	207 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/k953777/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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