FDA 510(k) Premarket Notification

K953607 TITANIUM LIMITED CONTACT DYNAMIC HIP SCREW PLATE (TILC-DHS)

CLEARED

Mar 8, 1996
219 days to decision

K953607 · Product code: **JDO** · Orthopedic
Source: https://www.510kdatabase.net/k953607/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Fixation, Proximal Femoral, Implant (JDO)
Date received	Aug 2, 1995
Decision date	Mar 8, 1996
Days to decision	219 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/k953607/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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