FDA 510(k) Premarket Notification

K033071 SYNTHES (USA) CANNULATED TITANIUM HUMERAL NAIL SYSTEM

CLEARED

Nov 5, 2003
37 days to decision

K033071 · Product code: **JDS** · Orthopedic
Source: https://www.510kdatabase.net/k033071/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nail, Fixation, Bone (JDS)
Date received	Sep 29, 2003
Decision date	Nov 5, 2003
Days to decision	37 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Synthes (Usa)**
Location	Mchenry, IL, US
Contact	LISA M BOYLE
510(k) history	411 submissions · 394 cleared · 1977-2015