FDA 510(k) Premarket Notification

K021932 SYNTHES 6.5 MM CANNULATED SCREW

CLEARED

Sep 6, 2002
86 days to decision

K021932 · Product code: **HWC** · Orthopedic
Source: https://www.510kdatabase.net/k021932/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screw, Fixation, Bone (HWC)
Date received	Jun 12, 2002
Decision date	Sep 6, 2002
Days to decision	86 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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