FDA 510(k) Premarket Notification

K120854 SYNTHES VA LCP ANKLE TRAUMA SYSTEM

CLEARED

Jun 18, 2012
89 days to decision

K120854 · Product code: **HRS** · Orthopedic
Source: https://www.510kdatabase.net/k120854/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Fixation, Bone (HRS)
Date received	Mar 21, 2012
Decision date	Jun 18, 2012
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k120854/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026