FDA 510(k) Premarket Notification

K081342 REPROCESSED EXTERNAL FIXATION ACCESSORIES

CLEARED

Aug 13, 2008
92 days to decision

K081342 · Product code: **KTT** · Orthopedic
Source: https://www.510kdatabase.net/k081342/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Nail/blade/plate Combination, Multiple Component (KTT)
Date received	May 13, 2008
Decision date	Aug 13, 2008
Days to decision	92 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Sterilmed, Inc.**
Location	Plymouth, MN, US
Contact	JOSHUA CLARIN
510(k) history	64 submissions · 64 cleared · 2001-2024