FDA 510(k) Premarket Notification

K072922 3S HEMI TOE

CLEARED

Nov 28, 2007
44 days to decision

K072922 · Product code: **KWD** · Orthopedic
Source: https://www.510kdatabase.net/k072922/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Toe, Hemi-, Phalangeal (KWD)
Date received	Oct 15, 2007
Decision date	Nov 28, 2007
Days to decision	44 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Trilliant Surgical, Ltd.**
Location	Round Rock, TX, US
Contact	JD WEBB
510(k) history	14 submissions · 14 cleared · 2007-2017