FDA 510(k) Premarket Notification

CLEARED

K070052 MET-HEAD MODEL# MHDD-XX:CC DIAMETER, CC CANNULATION

Mar 28, 2007
83 days to decision

K070052 · Product code: **KWD** · Orthopedic
Source: https://www.510kdatabase.net/k070052/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Toe, Hemi-, Phalangeal (KWD)
Date received	Jan 4, 2007
Decision date	Mar 28, 2007
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vilex, Inc.**
Location	Pleasant Hills, PA, US
Contact	ABRAHAM LAVI
510(k) history	13 submissions · 12 cleared · 1998-2014