FDA 510(k) Premarket Notification

K073329 SYGNAL DBM

CLEARED

Feb 28, 2008
93 days to decision

K073329 · Product code: **MBP** · Orthopedic
Source: https://www.510kdatabase.net/k073329/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filler, Bone Void, Osteoinduction (w/o Human Growth Factor) (MBP)
Date received	Nov 27, 2007
Decision date	Feb 28, 2008
Days to decision	93 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Musculoskeletal Transplant Foundation**
Location	Edison, NJ, US
Contact	NANCY BENNEWITZ
510(k) history	24 submissions · 24 cleared · 2004-2019