FDA 510(k) Premarket Notification

K062260 OSTEOBOOST-BMA

CLEARED

Oct 26, 2006
83 days to decision

K062260 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k062260/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Aug 4, 2006
Decision date	Oct 26, 2006
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Orthos Limited**
Location	Bristol, North Somerset, GB
Contact	ALAN RORKE
510(k) history	1 submissions · 1 cleared · 2006-2006