FDA 510(k) Premarket Notification

K082002 PROGENIX PLUS

CLEARED

Nov 24, 2008
133 days to decision

K082002 · Product code: **MQV** · Orthopedic
Source: https://www.510kdatabase.net/k082002/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filler, Bone Void, Calcium Compound (MQV)
Date received	Jul 14, 2008
Decision date	Nov 24, 2008
Days to decision	133 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medtronic Sofamor Danek**
Location	Memphis, TN, US
Contact	MICHELLE OBENAUER
510(k) history	154 submissions · 147 cleared · 2002-2021