FDA 510(k) Premarket Notification

K031399 INTERGRO DBM

CLEARED

Feb 18, 2005
655 days to decision

K031399 · Product code: **MQV** · Orthopedic
Source: https://www.510kdatabase.net/k031399/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filler, Bone Void, Calcium Compound (MQV)
Date received	May 5, 2003
Decision date	Feb 18, 2005
Days to decision	655 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Interpore Cross Intl.**
Location	Irvine, CA, US
Contact	MARK LOAR
510(k) history	39 submissions · 38 cleared · 1998-2005