

K031399 INTERGRO DBMFeb 18, 2005
655 days to decisionK031399 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k031399/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 20, 2026