

**K000515 PRO OSTEON 200R RESORBABLE BONE GRAFT
SUBSTITUTE, MODELS 2RG051, 2RG101, 2RG151, 2RG201,
2RG301, 2RG050, 2RG100, 2RG150,**

Sep 15, 2000
212 days to decision

K000515 · Product code: **MQV** · Orthopedic
Source: <https://www.510kdatabase.net/k000515/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filler, Bone Void, Calcium Compound (MQV)
Date received	Feb 16, 2000
Decision date	Sep 15, 2000
Days to decision	212 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k000515/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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