

**K990131 PRO OSTEON IMPLANT 500R RESORBABLE BONE  
GRAFT SUBSTITUTE**Mar 2, 1999  
47 days to decisionK990131 · Product code: **MQV** · Orthopedic  
Source: <https://www.510kdatabase.net/k990131/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filler, Bone Void, Calcium Compound (MQV)
Date received	Jan 14, 1999
Decision date	Mar 2, 1999
Days to decision	47 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Interpore Cross Intl.</b>
Location	Irvine, CA, US
Contact	LYNN M RODARTI
510(k) history	39 submissions · 38 cleared · 1998-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k990131/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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