

**K980817 PRO OSTEON IMPLANT 500R RESORBABLE BONE
GRAFT SUBSTITUTE MODEL NUMBERS 5RG05, 5RG10, 5RG15,
5G15L, 5RG20, 5RG30, 5RG30L**

Sep 25, 1998
206 days to decision

K980817 · Product code: **MQV** · Orthopedic
Source: <https://www.510kdatabase.net/k980817/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filler, Bone Void, Calcium Compound (MQV)
Date received	Mar 3, 1998
Decision date	Sep 25, 1998
Days to decision	206 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Interpore Intl.
Location	Walker, MI, US
Contact	DAVID P BALDING
510(k) history	25 submissions · 25 cleared · 1984-1998

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k980817/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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