

**K234013 Mg OSTEONJECT™**Feb 5, 2024  
48 days to decisionK234013 · Product code: **MQV** · Orthopedic  
Source: <https://www.510kdatabase.net/k234013/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filler, Bone Void, Calcium Compound (MQV)
Date received	Dec 19, 2023
Decision date	Feb 5, 2024
Days to decision	48 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	Mg OSTEOREVIVE™; Mg OSTEOCRETE™

**APPLICANT**

---

Company	<b>Bone Solutions, Inc.</b>
Location	Santa Barbara, CA, US
Contact	Drew Diaz
510(k) history	8 submissions · 8 cleared · 2009-2025

**REGULATORY CONSULTANT**

---

Consulting firm	<b>PaxMed International, LLC</b>
Contact	Kevin Thomas

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA [accessdata.fda.gov](https://accessdata.fda.gov)

---

510k Database - [www.510kdatabase.net](https://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k234013/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 13, 2026