

K242372 Mg OSTEOREVIVE™, Mg OSTEOCRETE™Oct 18, 2024
70 days to decisionK242372 · Product code: **MQV** · Orthopedic
Source: <https://www.510kdatabase.net/k242372/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Filler, Bone Void, Calcium Compound (MQV) |
| Date received | Aug 9, 2024 |
| Decision date | Oct 18, 2024 |
| Days to decision | 70 days |
| Third-party review | No |
| Combination product | No |
| PCCP authorized | No |
| Summary / Statement | Summary |

APPLICANT

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

REGULATORY CONSULTANT

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|-----------------|----------------------------------|
| Consulting firm | PaxMed International, LLC |
| Contact | Kevin Thomas |

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k242372/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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