

**K251556 Device 300423 Granules**Jul 17, 2025  
57 days to decisionK251556 · Product code: **MQV** · Orthopedic  
Source: <https://www.510kdatabase.net/k251556/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filler, Bone Void, Calcium Compound (MQV)
Date received	May 21, 2025
Decision date	Jul 17, 2025
Days to decision	57 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Geistlich Pharma AG</b>
Location	Washington, DC, US
Contact	Jan Van der Ploeg
510(k) history	26 submissions · 26 cleared · 2011-2026

**REGULATORY CONSULTANT**

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Consulting firm	<b>Gxpartners, LLC</b>
Contact	Gretchen Upton

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k251556/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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