

**K253147 FIBERGRAFT™ BG Putty GPS Bone Graft Substitute  
 FIBERGRAFT™ BG Putty Bone Graft Substitute FIBERGRAFT™  
 AERIDYAN™ Matrix Bone Graft Substitute FIBERGRAFT™ BG  
 Matrix Bone Graft Substitute**

Oct 22, 2025  
 27 days to decision

K253147 · Product code: **MQV** · Orthopedic  
 Source: <https://www.510kdatabase.net/k253147/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Filler, Bone Void, Calcium Compound (MQV)
Date received	Sep 25, 2025
Decision date	Oct 22, 2025
Days to decision	27 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Prosidyan, Inc.</b>
Location	Philedelphia, PA, US
Contact	Tamala Wampler
510(k) history	12 submissions · 12 cleared · 2014-2025

**REGULATORY CONSULTANT**

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Consulting firm	<b>Mcra, LLC</b>
Contact	Dr. Mehdi Kazemzadeh-Narbat

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k253147/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).

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