

K242280 Vitoss® BiModal Bioactive Bone Graft Substitute Foam Strip, Vitoss® BiModal Bioactive Bone Graft Substitute, Vitoss BBTrauma® Bioactive Bone Graft Substitute, Vitoss® BA2X Bioactive Bone Graft Substitute, Vitoss® Bioactive Bone Graft Substitute Pack, Vitoss® Bioactive Bone Graft Substitute, Vitoss® Foam Bone Graft Substitute, Vitoss® Bone Graft Substitute Filled Canister, Vitoss® Bone Graft Substitute, Vitoss® Bone Graft Substitute - Synthetic Cancellous Chips

Aug 29, 2024
27 days to decision

K242280 · Product code: **MQV** · Orthopedic
Source: <https://www.510kdatabase.net/k242280/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Filler, Bone Void, Calcium Compound (MQV)
Date received	Aug 2, 2024
Decision date	Aug 29, 2024
Days to decision	27 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Stryker Spine
Location	Allendale, NJ, US
Contact	Fiorella Lopardo
510(k) history	74 submissions · 73 cleared · 2004-2026

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k242280/> Data sourced from FDA 510(k) public records (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. - Generated May 14, 2026