

K190956 SCS 17-01Jan 30, 2020
294 days to decisionK190956 · Product code: **MQV** · Orthopedic
Source: <https://www.510kdatabase.net/k190956/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filler, Bone Void, Calcium Compound (MQV)
Date received	Apr 11, 2019
Decision date	Jan 30, 2020
Days to decision	294 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Anika Therapeutics, Inc.
Location	Bedford, MA, US
Contact	Jeffrey Cyr
Website	http://www.anikatherapeutics.com/
510(k) history	9 submissions · 9 cleared · 2017-2025

Anika Therapeutics, Inc. is a global leader in hyaluronic acid-based orthopedic regenerative solutions and osteoarthritis pain management. The company develops advanced tissue repair, cartilage regeneration, and injectable bone substitute technologies with a manufacturing facility in Bedford, US. Anika has received FDA 510(k) clearances from total submissions since 2017. Orthopedic devices represent the dominant focus of the company's regulatory portfolio. The latest clearance was received in 2025, reflecting active ongoing innovation and market engagement. The company's ...
