FDA 510(k) Premarket Notification

K190283 PROSTIM Injectable Inductive Graft

CLEARED

Mar 14, 2019
31 days to decision

K190283 · Product code: **MQV** · Orthopedic
Source: https://www.510kdatabase.net/k190283/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Filler, Bone Void, Calcium Compound (MQV)
Date received	Feb 11, 2019
Decision date	Mar 14, 2019
Days to decision	31 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Wrightmedicaltechnologyinc**
Location	Arlington, TN, US
Contact	Rachel Roberts
510(k) history	302 submissions · 291 cleared · 1993-2023