FDA 510(k) Premarket Notification

CLEARED

K150152 The Octane Straight Intervertebral Fusion Device, Ti Coated

May 11, 2015
108 days to decision

K150152 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k150152/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jan 23, 2015
Decision date	May 11, 2015
Days to decision	108 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Exactech, Inc.**
Location	Gainesville, FL, US
Contact	Dawn Davisson
Website	https://www.exac.com/
510(k) history	186 submissions · 174 cleared · 1986-2026

Exactech, Inc. operates with a manufacturing facility in Gainesville, US. The company does not offer direct sales or distribution in the United States. Product inquiries and safety concerns are handled through designated company contacts. Exactech has submitted FDA 510(k) applications, resulting in cleared devices. The company's regulatory activity spans from 1986 to 2026, demonstrating sustained engagement with FDA clearance processes. Orthopedic devices represent the dominant focus of the company's portfolio, accounting for approximately 99% of submissions. Recent FDA 5...

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k150152/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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