FDA 510(k) Premarket Notification

K040980 ALLOMATRIX C PUTTY, ALLOMATRIX CUSTOM PUTTY, ALLOMATRIX DR PUTTY, IGNITE SCAFFOLD, MODELS 860C, 86XC, 86DR, 860D, 860T

CLEARED

Jul 14, 2004
90 days to decision

K040980 · Product code: **MQV** · Orthopedic
Source: https://www.510kdatabase.net/k040980/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filler, Bone Void, Calcium Compound (MQV)
Date received	Apr 15, 2004
Decision date	Jul 14, 2004
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k040980/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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