FDA 510(k) Premarket Notification

K180090 Cure™ Lumbar Plate System

CLEARED

Apr 23, 2018
101 days to decision

K180090 · Product code: **KWQ** · Orthopedic
Source: https://www.510kdatabase.net/k180090/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Jan 12, 2018
Decision date	Apr 23, 2018
Days to decision	101 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Meditech Spine, LLC**
Location	Atlanta, GA, US
Contact	Bruce Dunaway
510(k) history	10 submissions · 10 cleared · 2015-2021