FDA 510(k) Premarket Notification

K180573 Responsive Arthroscopy Interference Screw System

CLEARED

Nov 23, 2018
263 days to decision

K180573 · Product code: **HWC** · Orthopedic
Source: https://www.510kdatabase.net/k180573/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screw, Fixation, Bone (HWC)
Date received	Mar 5, 2018
Decision date	Nov 23, 2018
Days to decision	263 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Responsive Arthroscopy, LLC**
Location	Minneapolis, MN, US
Contact	Doug Kohrs
510(k) history	11 submissions · 11 cleared · 2018-2025