

K181076 Responsive Arthroscopy Wedge Push-In Suture Anchors

Jan 18, 2019
269 days to decision

K181076 · Product code: **MBI** · Orthopedic
Source: <https://www.510kdatabase.net/k181076/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fastener, Fixation, Nondegradable, Soft Tissue (MBI)
Date received	Apr 24, 2018
Decision date	Jan 18, 2019
Days to decision	269 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k181076/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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