

**K180573 Responsive Arthroscopy Interference Screw System**Nov 23, 2018  
263 days to decisionK180573 · 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	<b>Responsive Arthroscopy, LLC</b>
Location	Minneapolis, MN, US
Contact	Doug Kohrs
510(k) history	11 submissions · 11 cleared · 2018-2025

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k180573/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 26, 2026