FDA 510(k) Premarket Notification

K173224 SPIN-SWI

CLEARED

Feb 23, 2018
143 days to decision

K173224 · Product code: **LNH** · Radiology
Source: https://www.510kdatabase.net/k173224/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Oct 3, 2017
Decision date	Feb 23, 2018
Days to decision	143 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Spintech, Inc.**
Location	Minneapolis, MN, US
Contact	Kay Fuller
510(k) history	10 submissions · 10 cleared · 1992-2022

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k173224/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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