

K240157 InVision™ 1.5 Surgical TheatreFeb 15, 2024
27 days to decisionK240157 · Product code: **LNH** · Radiology
Source: <https://www.510kdatabase.net/k240157/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Jan 19, 2024
Decision date	Feb 15, 2024
Days to decision	27 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Deerfield Imaging, Inc.
Location	Minnetonka, MN, US
Contact	Liz Ashworth
510(k) history	2 submissions · 2 cleared · 2021-2024

REGULATORY CONSULTANT

Consulting firm	Regulatory Technology Services, LLC
Contact	Prithul Bom

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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