

K230187 Terran NM-101Sep 29, 2023
249 days to decisionK230187 · Product code: **LNH** · Radiology
Source: <https://www.510kdatabase.net/k230187/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Jan 23, 2023
Decision date	Sep 29, 2023
Days to decision	249 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Terran Biosciences, Inc.
Location	New York, NY, US
Contact	Sam Clark
510(k) history	1 submissions · 1 cleared · 2023-2023

REGULATORY CONSULTANT

Consulting firm	Medical Device Regulatory Solutions, LLC
Contact	Kay Fuller

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

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