

K250381 Deepsight NeedleVue LC1 Ultrasound SystemAug 1, 2025
171 days to decisionK250381 · Product code: IYO · Radiology
Source: <https://www.510kdatabase.net/k250381/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Feb 11, 2025
Decision date	Aug 1, 2025
Days to decision	171 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	DeepSight Technology, Inc.
Location	Santa Clara, CA, US
Contact	David Surber
510(k) history	1 submissions · 1 cleared · 2025-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k250381/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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