

K253927 LiveMedica Enterprise PACS

Feb 6, 2026
60 days to decision

K253927 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k253927/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Dec 8, 2025
Decision date	Feb 6, 2026
Days to decision	60 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Live Medica, LLC
Location	Raleigh, NC, US
Contact	Nick Ponder
Website	https://livemedica.com
510(k) history	1 submissions · 1 cleared · 2026-2026

Live Medica, LLC specializes in cloud-based medical imaging solutions, with a manufacturing facility in Raleigh, US. The company focuses on enterprise-level Picture Archiving and Communication Systems (PACS) technology for healthcare providers. Live Medica has received FDA 510(k) clearance from total submission. The company's regulatory activity centers on Radiology devices, with its most recent clearance in 2026, demonstrating active engagement in the medical device market. Explore the company's cleared device names, product codes, and clearance dates in the database to ...

REGULATORY CONSULTANT

Consulting firm	MEDlcept, Inc.
Contact	Danielle Short

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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Device record: <https://www.510kdatabase.net/k253927/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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