



AI-Powered Imaging: New Horizons in Fighting Cancer

Corporate Update
March 2026

MEDIAN TECHNOLOGIES



ALMDT
EURONEXT
GROWTH

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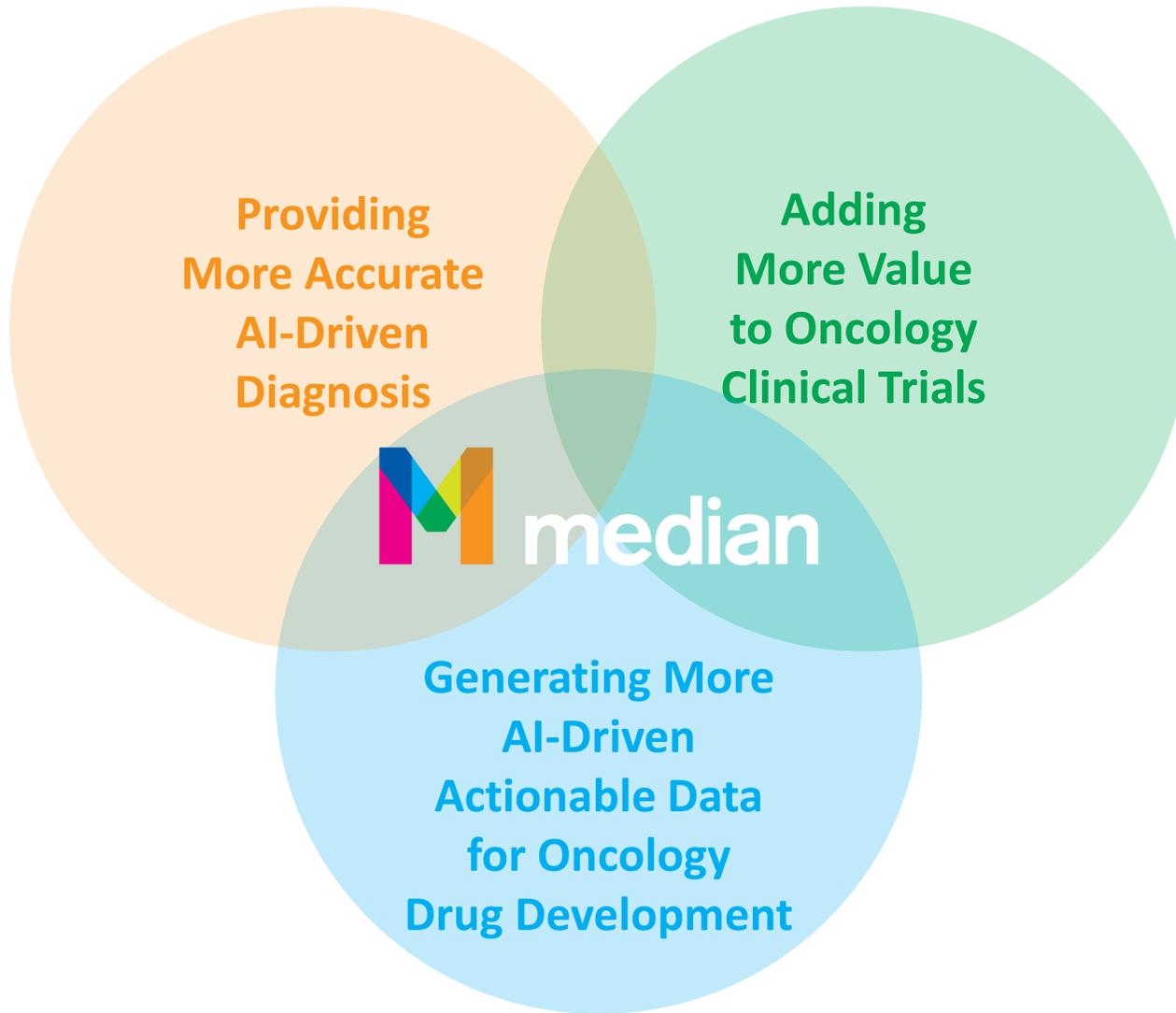
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These forward-looking statements can often be identified by the words "expects," "anticipates," "believes," "intends," "estimates," "plans" "can," "contemplate," "could," "is designed to," "may," "achieve," "potential," "objective," "target," "project," "strategy," "predict," "forecast," "opportunity," "goal," or the negative of these, and any other similar expressions are intended to identify forward-looking statements.

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**Providing Invaluable Clinical
Insights Where Others
Only See Images**

**Leveraging Proprietary AI,
Computer Vision, and Signal
Processing Technologies to
Develop Imaging Tests and
Services for Unmet Needs
in Oncology**

Proven & experienced management team



Fredrik Brag

Chief Executive Officer, Founder,
Board Member



Oran Muduroglu

Executive Chairman of the Board,
Named President of Median eyonis Inc.



Jean-Christophe Montigny

Chief Financial Officer



Thomas Bonnefont

Chief Operating Officer &
Chief Commercial Officer of Median eyonis



Nicolas Dano

Chief Operating Officer &
Chief Commercial Officer of Median iCRO

MANAGEMENT AND BOARD EXTENSIVE PRIOR EXPERIENCE



HealthCenter
/ Focus Imaging



BOARD OF DIRECTORS

Oran Muduroglu

Fredrik Brag

Tim Haines

Kapil Dhingra

Oern Stuge

Ben McDonald

Didric Cederholm





Helping Save the
Lives of Patients
Through
Innovation



Global Business – People Everywhere Need Better Answers to Cancer

France

Headquarters

U.S. + China

Subsidiaries

190+

Employees

€23.5M

2025 Revenue

Strong Scientific and Commercial Validation

7K+

Site Network

150+

Publications &
Conference Papers

340+

Clinical Studies

95+

Biopharma
Clients

2 of 3

Top Oncology
Pharmas are Clients

Leading Imaging Solutions



End-to-End CADe/CADx
Software As Medical Devices



Imaging Services for
Oncology Clinical Trials



End-to-End CADe/CADx Software as Medical Devices (SaMDs)

Enabling earlier detection & characterization of cancer with unprecedented performance – lung cancer as initial market

-  Unprecedented accuracy
-  Helps save patients' lives
-  Reduces unnecessary procedures & healthcare spend
-  Optimizes radiologist's time with simple workflows



Imaging Services for Oncology Clinical Trials

Empowering biopharma to leverage AI, ML, data mining, and radiomics in clinical development strategy to:

-  Select better patients
-  Predict responses to therapy
-  Accurately monitor disease progression
-  Rapidly assess safety and efficacy

Imaging Solutions Make an Impact Throughout the Entire Patient Journey

Strong momentum from recent business catalysts



**High Commercial
Launch Readiness in Lung Cancer**

- Achieved FDA 510(k) clearance & filed CE marking submission
- Named Oran Muduroglu as President of Median eyonis Inc. to lead the U.S. launch and scale-up, building on strong foundation of preparation
- Signed non-exclusive distribution agreement with Tempus
- Achieved best-in-class clinical results in pivotal studies
- Expanded KOL network across radiology, pulmonology & thoracic oncology

iCRO

**Continued Business Momentum
& Improved Financial Profile**

- Achieved record backlog – driven by U.S. demand
- Cash flow positive in 2025⁽¹⁾
- Expanded traction with top-tier pharmas

Company also completed up to €61.4M refinancing⁽²⁾ in summer 2025 to significantly strengthen financial position

[1] Unaudited, iCRO segment.

[2] €23.9M in capital increase and remainder in European Investment Bank (EIB) financing of €37.5M of which €18.5M remains undrawn.



AI imaging is revolutionizing cancer care

Median's next generation AI Software as Medical Devices (SaMD) can diagnose early-stage cancer when it can be cured

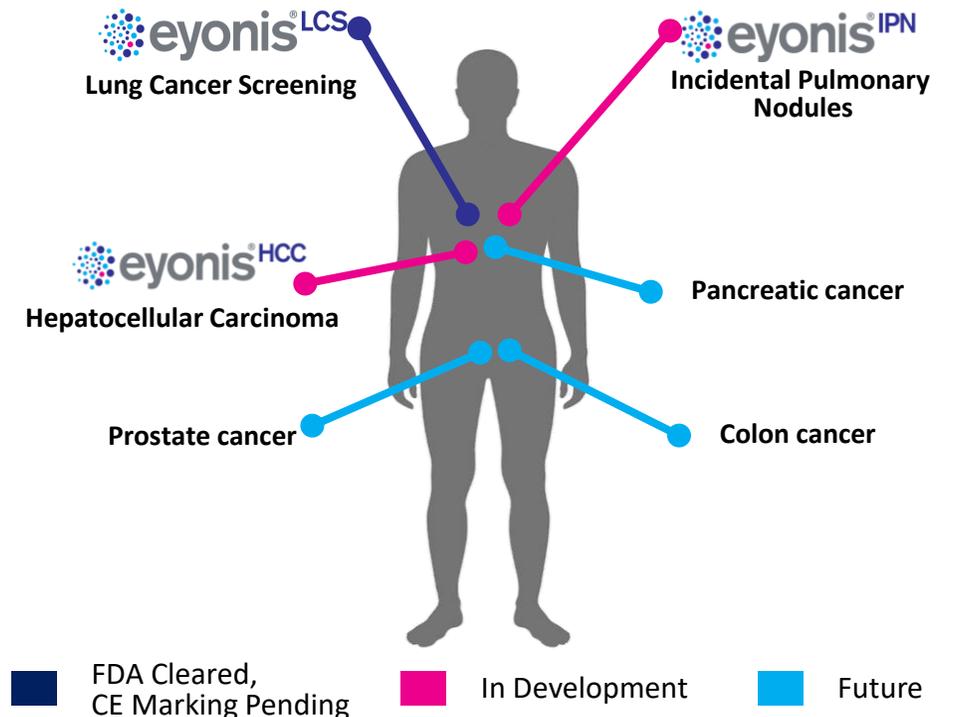


Catching the Unseen – Supporting Radiologists and Clinicians Worldwide in Screening, Early Diagnosis & Treatment of Cancer

End-to-End, AI/ML-Based CADe/CADx SaMDs

- Enabling earlier and more accurate cancer diagnosis from medical images
- Helping clinicians deliver better outcomes while reducing unnecessary procedures and lowering healthcare cost
- Detection, localization, and characterization of lesions into probably benign, suspicious, and very suspicious
- Unprecedented sensitivity and specificity versus standard of care and liquid biopsy

Widely Applicable Foundational Technology – First Commercial Opportunity in Lung Cancer Screening (LCS)



Why target lung cancer screening as eyonis' beachhead market?

#1

Cancer Killer Worldwide⁽¹⁾

18.6%

of All Lung Cancer Patients
Survive 5 years⁽²⁾

14.5M

U.S. Eligible Population⁽³⁾

19%

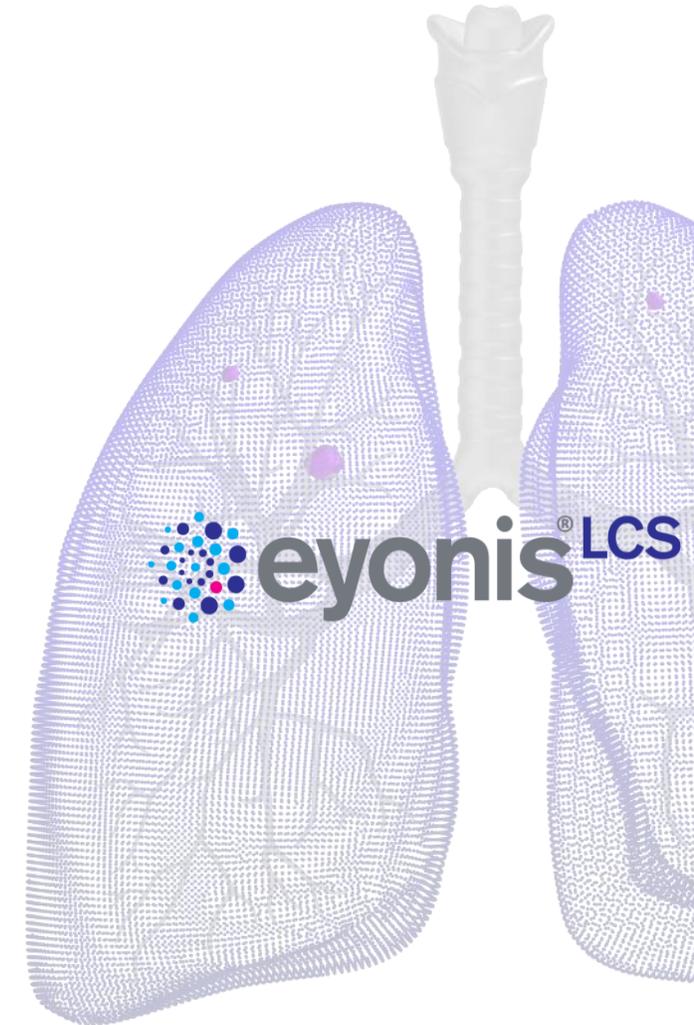
Of All Cancer Deaths:
1.8M Deaths Globally⁽¹⁾

80%

of Patients Diagnosed at
an Early Stage via LDCT
Screening Survive
20 Years⁽²⁾

\$10B

Estimated U.S. TAM⁽⁴⁾



[1] Cancer Tomorrow, IARC, Global Cancer Observatory 2022 – WHO.

[2] Mount Sinai - Lung Cancer Screening Dramatically Increases Long-term Survival Rate.

[3] USPSTF – U.S. Preventive Services Task Force 2021.

[4] Based on a \$650 per procedure reimbursement opportunity & 14.5M estimated population size.

The only FDA approved AI-based SaMD solution for LCS



The Historical Diagnostic Paradigm for LCS is Inefficient

- ✗ No end-to-end solutions to support radiologists
- ✗ LCS is done by radiologist alone or aided by detection (high false positive rate)
- ✗ No detection & characterization SaMD currently approved in the U.S. for LCS
- ✗ 2/3 false positives without eyonis LCS AI remain a key barrier to scaling LCS



- ✓ Only end-to-end solution
- ✓ Enhancing screening accuracy and efficiency

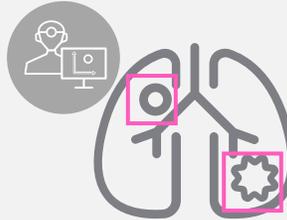
Radiologist Alone



Manual detection & diagnosis

Tedious, time consuming & error prone

Radiologist with Computer-Assisted Detection (CADe) SaMD only

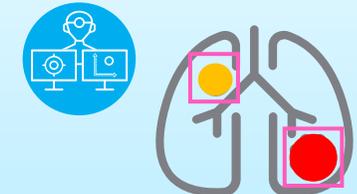


□ Detection

Manual diagnosis – one nodule at a time

High rates of false positives

Radiologist with End-to-End Computer-Assisted Detection & Diagnosis (CADe/CADx)



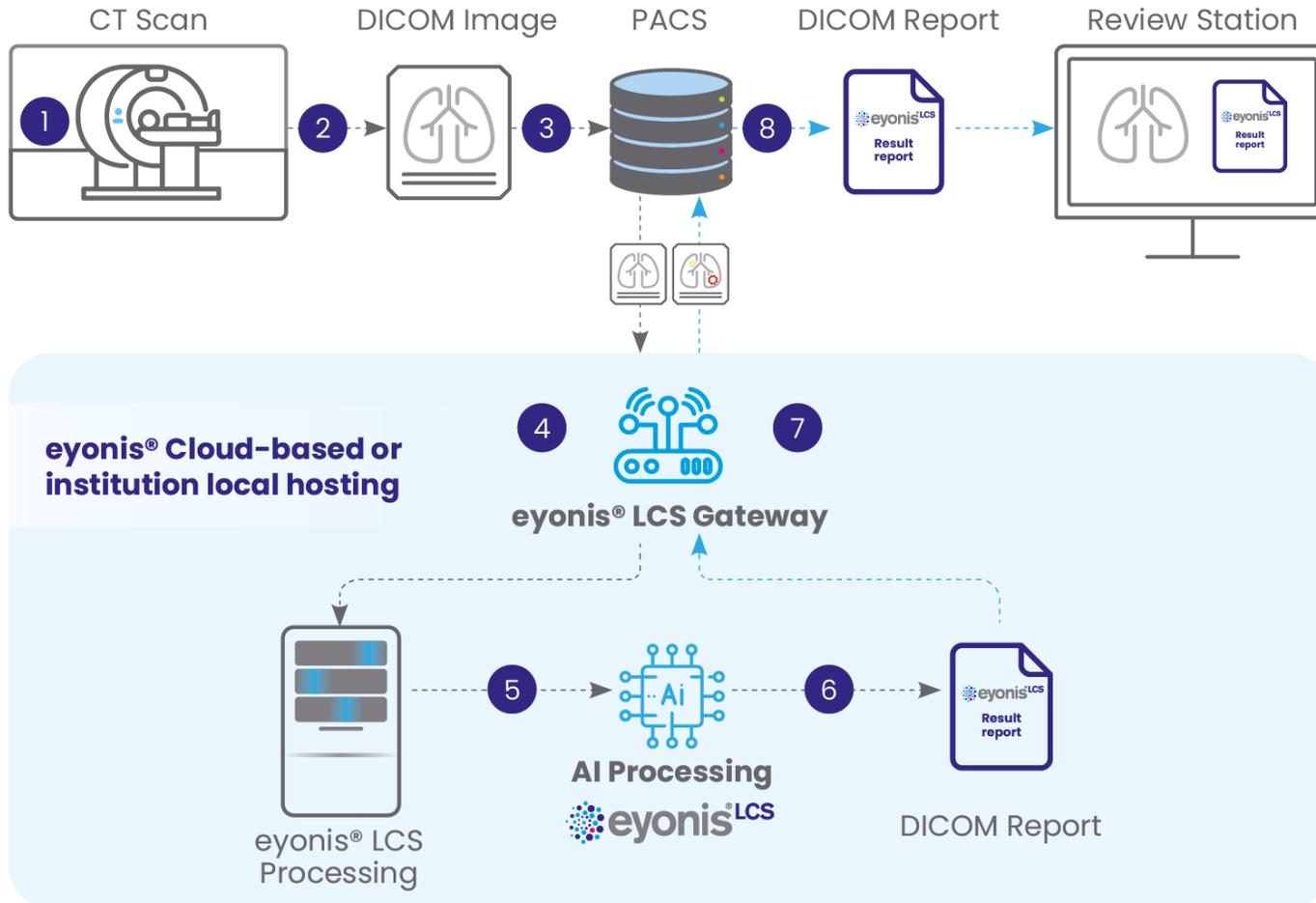
■ Probably Benign
■ Suspicious
■ Very suspicious

No Manual Intervention

Detection & diagnosis of all nodules from probably benign to very suspicious reduce false positives by up to 2/3

Seamlessly enabling scale-up of lung cancer screening

eyonis[®]LCS Workflow



Easy-to-use



Optimizes radiologist's time



Increases workflow standardization



Compatible with any CT-scan and any PACS system



No manual intervention



Greater accuracy & efficiency

**eyonis LCS
Manufacturer Values
(MV)**

**Performed to determine
eyonis LCS performance**

Evaluated on a U.S. cohort of 2,163 cases: 2,027 benign / 136 cancers, of which 66.18% were stage I

Negative Predictive Value (NPV): 99.9%

	Sensitivity	Specificity
	TP-positive Cases Detected (Malignant)	TN-Negative Cases Detected (Benign)
<p>Radiologist</p>	80.3%	76.4%
	93.3%	92.4%
	<p>66% False Negative Reduction</p>	<p>68% False Positive Reduction</p>

eyonis[®] LCS

- Significantly reduces false positives and false negatives (missed cancers)
- Reduces unnecessary invasive diagnostic procedures
- Allows high-confidence rule-out (screening default) for annual follow-up
- Supports high-confidence efficient rule-in (recall) pathways for at-risk patients expedited diagnostic work-up
- Saves lives with earlier lung cancer detection & characterization
- **Enables nationwide LCS at scale**

Strong platform validation

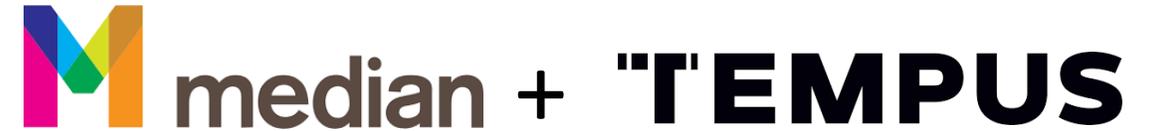
Leveraging Top U.S. & EU-based Academic Clinical Sites Involved in Pivotal Studies



23 eyonis Conference Papers, Oral Presentations, and Journal Publications since 2021



1st Distribution Agreement Signed with Tempus AI



- Integration of eyonis LCS into Tempus Pixel, an FDA-cleared, CE-marked AI-enabled solution that provides advanced analysis, tools, and automated reporting from radiology images
- Leverages Tempus' established position in oncology and AI-based precision medicine, and its strong network of healthcare providers, oncologists, and diagnostic centers
- Reinforces the clinical utility of the technology and facilitates adoption via an established enterprise workflow

Plans to broaden commercial adoption of eyonis LCS by targeting additional non-exclusive distribution agreements with top-tier imaging, cloud technology & Dx partners

eyonis LCS commercial launch and scale-up in the U.S.



Oran Muduroglu to lead U.S. Commercialization

 >3 decades of experience building, engineering, and scaling enterprise imaging and workflow platforms



Stentor/PHILIPS



 **Experience leading complex national deployments requiring:**

- Deep EMR and PACS interoperability
- Rigorous product development
- Disciplined enterprise sales
- Alignment with evolving reimbursement models

Key Pillars of the U.S. Commercial Strategy

Phased launch strategy for disciplined national, scalable expansion

- Detailed customer and payor mapping to prioritize regions with high lung-cancer screening volumes
- Targeting patients eligible for reimbursement
- Prospective health-economics studies initiating in 2026

Build-out of direct enterprise sales organization

- Expansion of robust U.S.-based commercial organization:
 - IT infrastructure and management
 - Clinical specialists
 - Demand generation direct sales / inside sales & IDTF
 - Collaboration with advocacy groups
 - U.S. marketing

Strategic non-exclusive distribution agreements

- Tempus partnership announced on February 12, 2026, covering U.S. and EU
- Plans to secure additional distribution collaborations with top-tier PACS, workflow and diagnostics partners

Existing reimbursement to support launch



- Existing CPT III codes for tissue characterization under CT: 0721T & 0722T
- CMS payment for 2 CPTIII codes assigned to New Tech APC 1508 - Level 8 (\$601 - \$700)

Total Medicare Hospital Payment is LDCT \$107 + eyonis LCS \$650⁽¹⁾

CPT Code	CPT Description	Existing LDCT Images	Hospital Outpatient Payment
71271 APC 5522	Computed tomography, thorax, low dose for lung cancer screening , without contrast material(s)	LDCT	\$107
0721T APC 1508	Quantitative CT tissue characterization , including interpretation and report, obtained <i>without</i> concurrent CT examination of any structure contained in previously acquired diagnostic imaging Do not report 0721T in conjunction with 71271	YES	\$650
0722T APC 1508	0721T but with concurrent CT examination is code 0722T Use 0722T in conjunction with 71271	NO	\$650 + \$107

These two procedures provide an LDCT and tissue characterization & are billed on separate days

The tissue characterization is “added-on” to the LDCT & are billed on the same day

Existing reimbursement creates potential opportunity for ~\$10B TAM – ~14.5M patients at \$650 reimbursement/test

Clear growth strategy designed to expand eyonis' reach

Increase Presence in U.S.



- Additional non-exclusive distribution agreements, IDTF & VA focus
- Leverage KOL partnerships to enable thought leadership across LCS programs
- Initiate health-economics studies to work towards broader, long-term insurance coverage

Drive Traction in Europe



- Achieve eyonis LCS CE marking (expected in Q2:2026)
- Leverage existing non-exclusive distribution agreement to cover Europe
- Focus on larger countries (Germany, France, Italy, Poland) and quick wins

Expand Portfolio



- Training eyonis unique proprietary CADe/CADx AI algorithm within our platform on additional cancer key indications
- Targets: Incidental Pulmonary Nodules, HCC (liver cancer), pancreatic cancer, colon cancer, and prostate cancer

iCRO

AI-powered imaging to revolutionize drug development

Our **central imaging services** provide global biopharma customers key data on patient response to therapies from phase I to III oncology studies

Our **Imaging Lab services** drive oncology drug development success with transformative AI insights

iCRO

Empowering Sponsors and CROs to Accelerate the Development and Delivery of Life-Saving Cancer Treatments

Global Leader in Central and AI Imaging Services for Oncology Clinical Trials

- Best-in-class, end-to-end image management for Phase I to Phase III clinical trials
- Imaging Lab, Median's iCRO enriched offering, drives drug development success with transformative

AI-powered insights



Select better patients



Predict response to therapy



Accurately monitor disease progression



Rapidly access safety and efficacy

Strong Financial Profile

€23.5M

FY2025 Revenue
Approx. 3% growth YoY

€76.6M

Backlog as of 12/31/25
Approx. 8% growth YoY
(17% constant forex)

Cash Flow Positive⁽¹⁾

Strong traction with leading biopharma

Global Footprint:
Across U.S., Europe, and Asia

95+
Biopharmaceutical Clients

7K+
Site Network

Preferred Imaging Services Provider to
2 of the Top 3 Pharma in Oncology

6
Successful FDA Inspections

Successful FDA Inspection in China in 2025

27
Successful Chinese NMPA Inspections

55
Phase I trials

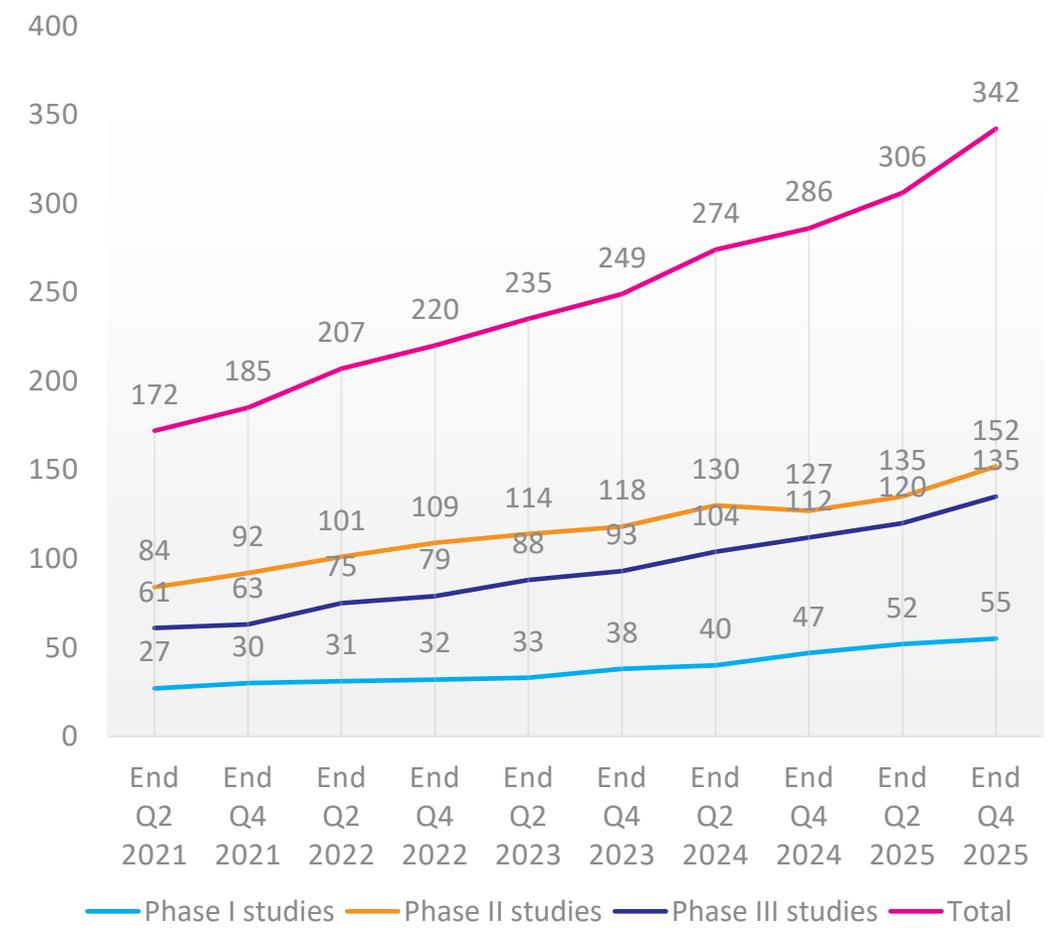
342 Oncology Studies

152
Phase II trials

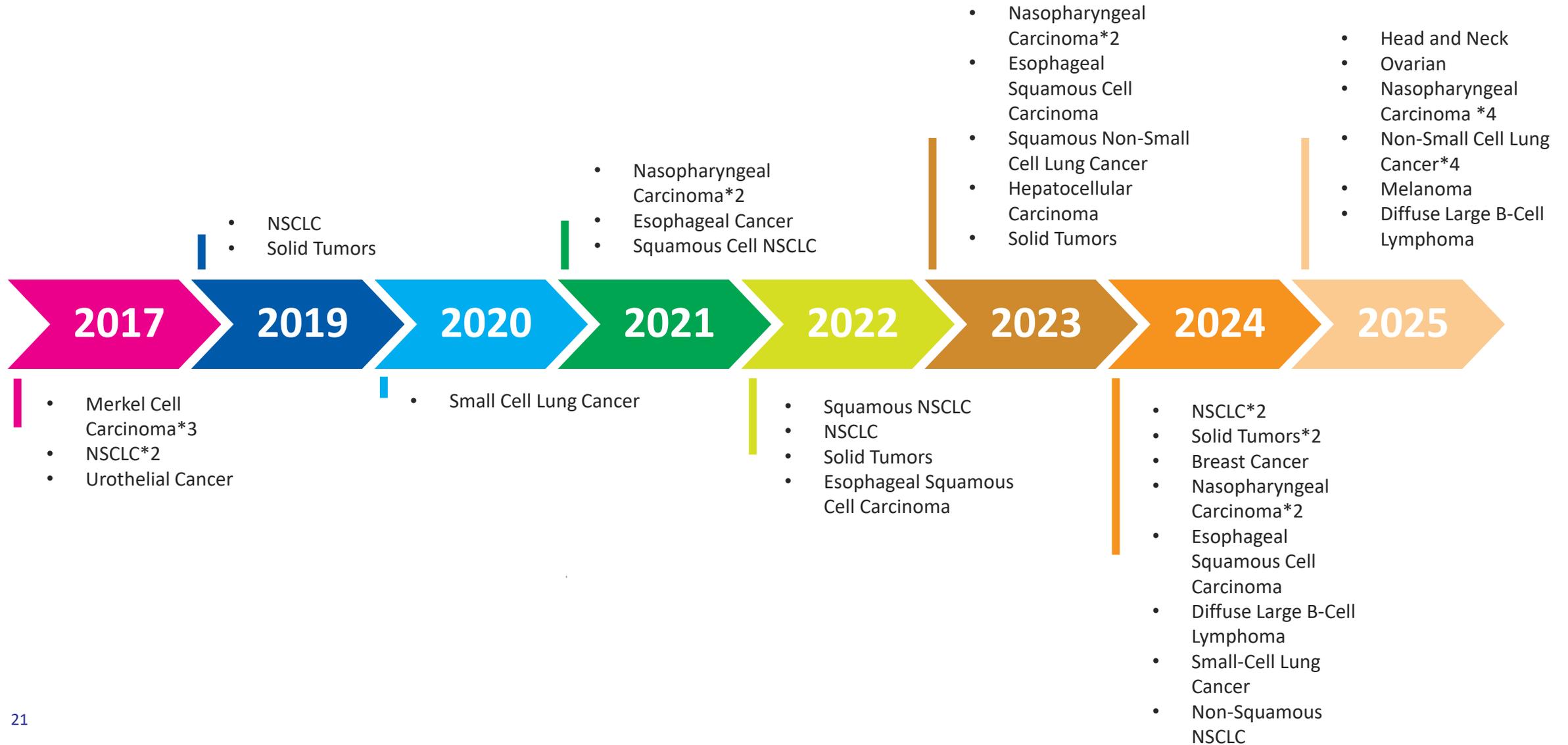
135
Phase III trials

All figures as of December 31, 2025.
Cumulative contracted and less than 12-month awarded studies, since the beginning of the iCRO activity, and until December 31, 2025.

Evolution of Oncology Studies Managed by Median vs Phases



Supported 40+ regulatory drug approvals across numerous indications



Leveraging the Transformative Power of Imaging AI for Drug Development

1

Collaborations

Establish AI imaging drug development collaborations with pharma groups

2

Geographical Strategy

EU/U.S.: be selected as preferred provider by big pharma groups, partner with CROs

China: grow market share

3

Operational Focus

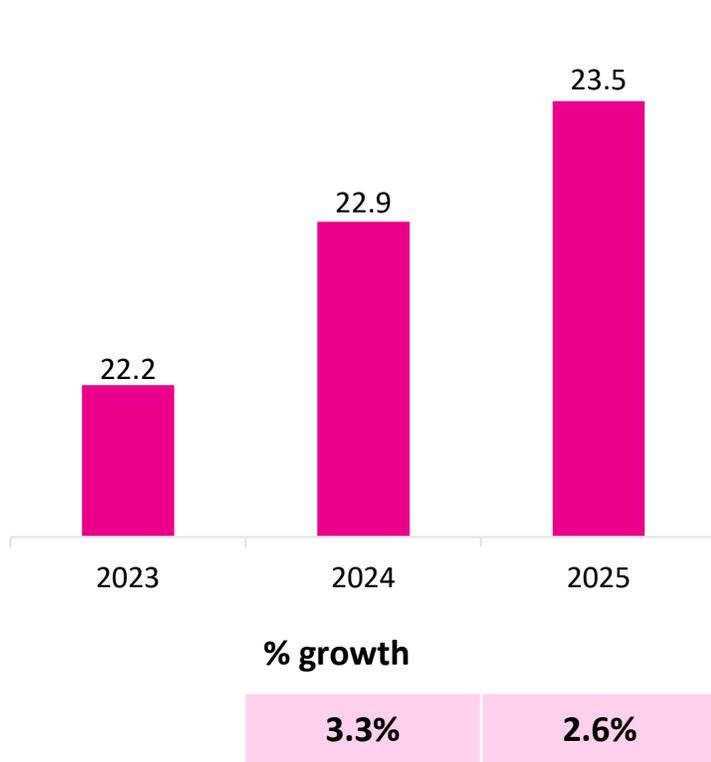
Further profitability improvement expected through operational optimization and automation

Summary Financials

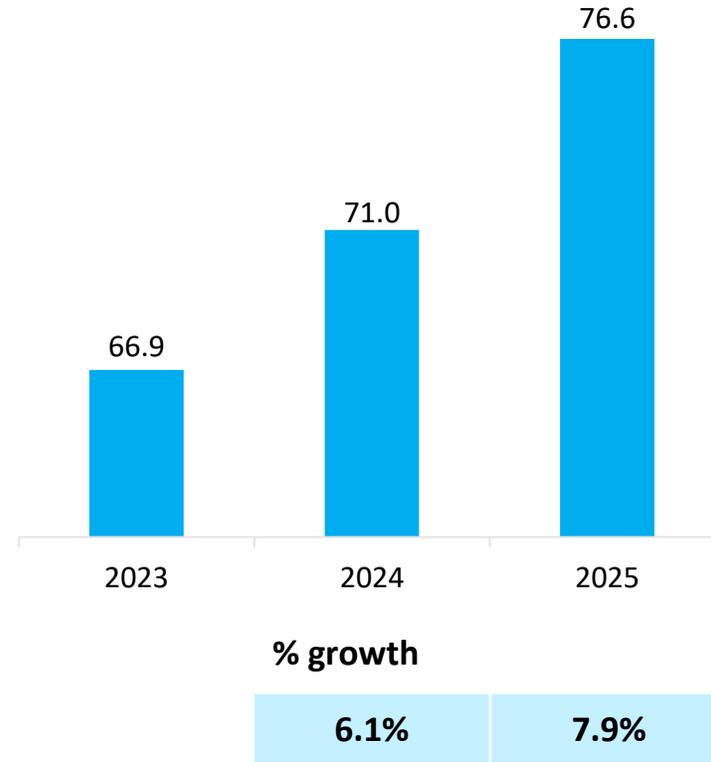
Continued growth while driving operational efficiency

- iCRO business has continued to grow despite strong operational improvements – iCRO segment now cash flow positive⁽¹⁾
- eyonis LCS launch to drive further topline acceleration

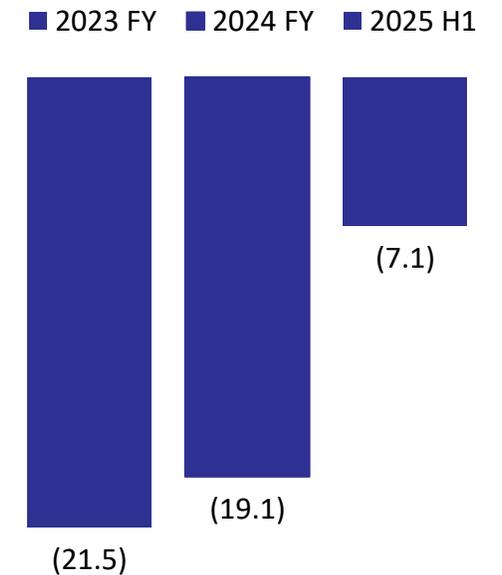
Revenue (€ Millions)



Backlog (€ Millions)



Cash Flow Before Financing Activities (€ Millions)



As of December 31, 2025, cash and cash equivalents at €18.2 million

Refinancing Overview

- Up to €61.4M refinancing completed in summer 2025
 - **August 1st**: €23.9M gross capital increase through ABSA issuance (new ordinary shares accompanied by a warrant at an exercise price of €2.39 per new ordinary share)
 - **July 11st**: Up to €37.5M EIB financing facility secured
 - **October 21st**: First €19.0M tranche drawn, following €20.7M repayment of the 2019 EIB loan tranche

Current cash runway extends through:

Q4:2026

Potential upside to cash position from 13.3M warrants:

€47.8M

In additional proceeds upon full exercise

Business Highlights

Several features drive business opportunity

1

More accurate, earlier AI-driven diagnosis through eyonis LCS for lung cancer is now FDA cleared – large market opportunity, exceptional data and well-formulated commercial strategy, including existing reimbursement

2

Providing invaluable clinical insights to medical images along the cancer patient journey, from drug development to patient care

3

Highly validated platform, with 150+ publications, 95+ biopharma customers and marquee partnership with Tempus AI to drive strong commercial traction of eyonis LCS

4

Global operations and customer base – providing better answers to cancer for patients worldwide

5

Highly experienced management team with track-record of successfully developing, commercializing and scaling life sciences solutions

Appendix



Demonstrated clinical strength through two pivotal studies

1- REALITY Standalone Study ClinicalTrials.gov ID: NCT06576232

Study design: Retrospective study, 12 months follow-up; 79.7% stage I, 4.7% stage II, 8.7% stage III, 6.9% stage IV

Sample size: Enriched cohort of 1,147 cases (342 cancer, 805 benign) enrolled in Europe & U.S.

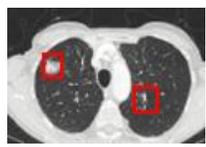
Objectives: Assess SaMD's performance in characterizing cancer positive & negative patients as well as detecting suspicious / malignant on LDCT images

Primary Endpoint: AUROC that measures eyonis LCS performance on patient-level data

eyonis^{LCS}
run
onto raw
cohort



Algorithm analysis



Truthed*
cohort

Compared to

Statistical Analysis

Comparison of truthers ground truth VS. Software as a Medical Device output

Standalone Study Result

2 - RELIVE MRMCC Study ClinicalTrials.gov ID: NCT06751576

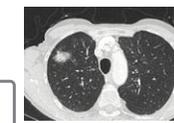
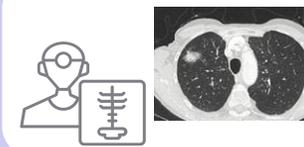
Study design: Two arm, randomized, controlled, blinded, paired-split-plot design study with 4 blocks and 16 readers

Sample size: Enriched cohort of 480 cases (160 cancers, 320 benign) and 16 readers enrolled in Europe & the U.S.

Objectives: Demonstrate that eyonis LCS improves clinician performance in analyzing LDCT lung screening scans

Primary Endpoint: Statistical radiologist performance superior with eyonis LCS report vs. without

Radiologist
without
eyonis^{LCS}



Report

Radiologist
with
eyonis^{LCS}

Compared to

Statistical Analysis

Compared reading with LCS report vs without

MRMC Trial Result



ALMDT
EURONEXT
GROWTH

Our Core Values

Leading innovation with purpose

Combine the spirit of innovation with our passion and conviction to help cure cancer and other debilitating diseases.

Committing to quality in all we do

Be dedicated to quality in everything we do. Quality begins with us and we are committed to it.

Supporting our customers in achieving their goals

Listen to the needs of our customers and help make their goals our goals through our innovation, imaging expertise, superior services, and quality solutions.

Putting the patient first

There is a person at the other end of the images we analyze who is counting on us to do everything we can to help make them healthier.

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