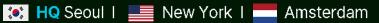
Conquer Cancer through Al



DISCLAIMER

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About Lunit Since 2013 | 1st Generation Medical Al Company | Listed in KOSDAQ since 2022

Global Investors

Backed by leading global investors

Raised \$122M while unlisted

Foreign investment =\$73M 60%

- 9% of global shareholders after IPO
- → Top 12% of KOSDAQ listed 1,800 companies in Korea SoftBank



Industry Leader

Best-in-class Al technology and products

- · Unprecedented "AA-AA" highest level in technology assessment among healthcare companies
- Best performance proven by direct comparison with global competitors in various studies
- · Highest market cap, among listed medical Al companies (2025.09)



Global Partners

Established partnerships with industry giants

Chosen by leading medical device companies

FUJIFILM PHILIPS

Chosen by leading bio companies

(A) GUARDANT

Multiple global pharmas

Commercial Success

Successfully scaling out revenue growth



Core Competence



Best-in-Class A.I.

Global leader in Al technology

- Top-tier proprietary Al technology
- 30+ papers presented in top AI conferences
- Large-scale medical data for R&D (N = +4 million cases)







+480 Publications in Major Peerreviewed Journals & Abstracts



THE LANCET JAMA Oncology





SCIENTIFIC REPORTS



AACR



RSNA

Clinical Infectious Diseases



XUSCAP

+100 Research Partnerships























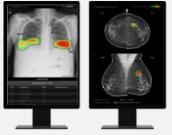
Conquer Cancer through Al Increase cancer survival through Al-powered cancer diagnosis and treatment

Cancer diagnosis support solutions

Lunit INSIGHT®

+50% 50% more patients can

be screened earlier



Flagship **Lunit INSIGHT CXR / MMG**







Chest CT



3D Mammo (DBT)

Al research platform

Cancer treatment support solutions

Lunit scope®

+50% 50% more patients can be eligible for immunotherapy



Flagship **Lunit SCOPE IO**





DNA/RNA

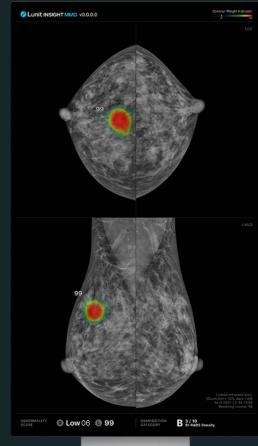


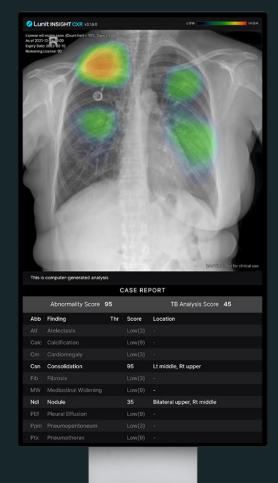
Tissue



Increase Survival of **Cancer Patients**

Lunit INSIGHT®













Product Roadmap | Beyond cancer risk prediction to autonomous Al

Current

Risk Assessment

Foundation Model

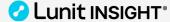
Autonomous Al

AI-CAD

Final clinical decision: physicians







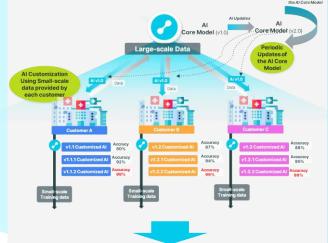
Al-based Risk Prediction

 Hybrid Al model combining EMR data with synthetic mammography (DBT) images



Al Customization

- Periodic updates of the Al core model
- Al customization using data provided from each site

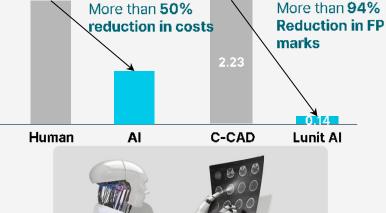


Improve Al accuracy across all customers

Autonomous Al in Cancer Screening

- Cost efficiency maximized through AI
- Near-perfect accuracy achievable through large-scale learning
- Final clinical decision: Al

Expected Performance with Autonomous AI





Lunit INSIGHT Value

Higher Reading Accuracy | Better performance than specialists, proven in major studies

Evidence Extensively validated through studies Studies/abstracts

THE LANCET Digital Health

JAMA Oncology

Radiology

Network Open.

RSNA°

REPORTS

Clinical Infectious Diseases

European Radiology



ВМС Pulmonary Medicine

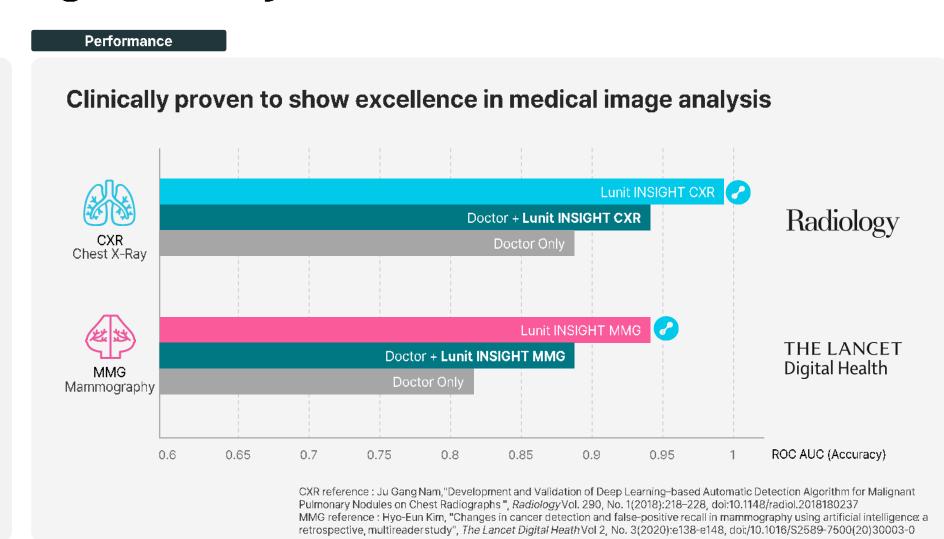
AJR

ACTA RADIOLOGICA

ITD

THE LANCET Regional Health

EUROPEAN RESPIRATORY journal



*Randomized Controlled Trial

Real World Evidence | Building solid evidence through prospective studies

Study Method

What is prospective study? Study designed

Retrospective

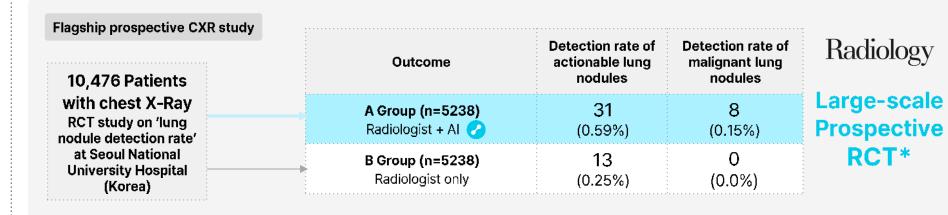
Prospective

Outcome is measured after the baseline state of the subjects is determined and controlled intervention applied.



Higher possibility to receive reimbursement if backed by prospective studies

Lunit INSIGHT CXR



Lunit INSIGHT MMG

: Research method in which subjects are randomly divided into control and experimental groups for comparison

Flagship prospective MMG study	Outcome	Detection rate of cancer per 1,000 women	Recall rate per 1,000 women	THE LANCET Digital Health
55,579 women screened for	1 Radiologist + Lunit INSIGHT MMG 🕜	4.3	28	
breast cancer Study on 'Double Reading' by Karolinska Institutet (Sweden)	2 Radiologists	4.1	29.3	Large-scale Prospective Study
msututet (sweden)	Lunit INSIGHT MMG 🕜	4.1	15.5	Study

Current Commercialization Status

Usage

Number of paying sites worldwide 1

Portion of revenue from

Overseas cumulative through Q3 2025

Lunit Users among Top 10 Hospitals in Korea

Chest X-Ray exams in Korea² Analyzed by Lunit INSIGHT CXR 2024

Lunit INSIGHT MMG Users among 47 Large Hospitals in Korea 2025

57%

20%

75%

Partner Testimonials

First Al company collaboration for our x-ray business

— General Manager K**



"This is actually out first Al company collaboration for our x-ray business. The offering will be available on all of GE digital, fixed, and mobile, and even fluoroscopy. Leveraging the long-term established expertise, that GE has on x-ray equipment in combination with Lunit's technology will help to keep up with the customer needs, care deeply about patients and diagnosing physicians or radiologists."

Felt very high customer satisfaction

— Global Marketing Manager M★★



"In Japan especially, many of the elder people cannot move from home to the clinic to take x-ray. Our x-ray with Lunit Al can visit to take x-ray immediately at home, so it is a significant difference. After we used Lunit Al in the market, I didn't have any complaints. From Fujifilm's standpoint, whatever Al lunit made has a very high quality, so we would want to implement it in our x-ray system or PACS system. Customers have already started to use Al and expect more support."

Lunit has both strong clinical record and national scalability

— Global Software Product Manager P★★



10

"Lunit has a strong clinical validation track record and is registered in Europe and also more than 19 countries, making it an excellent partner for developing solutions that bring our customers intelligent and definitive answers to their daily needs. We believe that the integration of Lunit's Al in Agfa's advanced medical imaging workstation will help facilitate more effective clinical workflows in various hospital situations."



Excludes demo and research use; only commercial sale

²⁾ Source: KOSTAT.go.kr Appx. 47M Chest X-ray exams performed annually in Korea

11

Key Success Factor: Proven to Be Best-in-Class

Accuracy



JAMA Oncology Salim, et al. JAMA Oncol. 2020 Aug 27.



IF 24,7991)

Comparing 3 Commercial Mammography Al

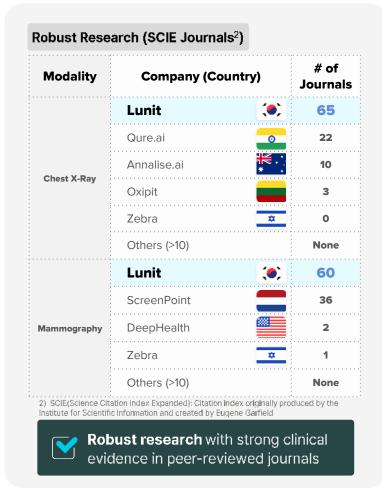
Retrospective analysis of 8 years of mammography screening Compared Al screening performance with data from 739 breast cancerdiagnosed women and 112,924 healthy women

	Lunit	Company A	Company B
AI	✓ Lunit INSIGHT MMG°	Algorithm A	Algorithm B
Sensitivity	81.9%	67.0%	67.4%
Accuracy (ROC AUC)	95.6%	92.2%	92.0%

1) Impact factor, Top 7 among Oncology Journals

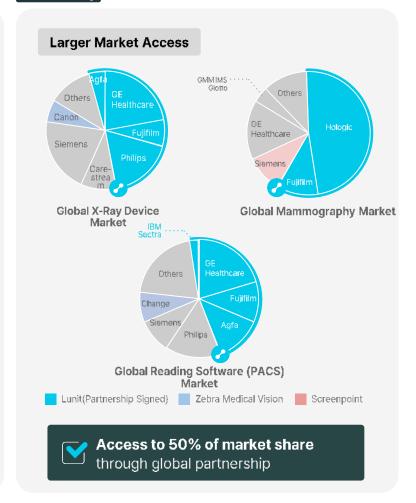


Research



Partnership

Company Overview



Lunit INSIGHT Strategy

Next Step | Becoming global standard of care by B2G expansion and successful reimbursement

B to G

National Cancer Screening Programs

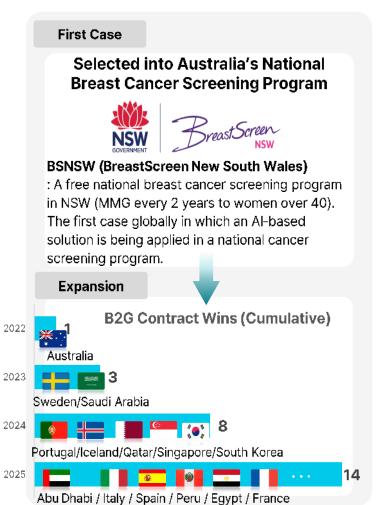
Reimbursement

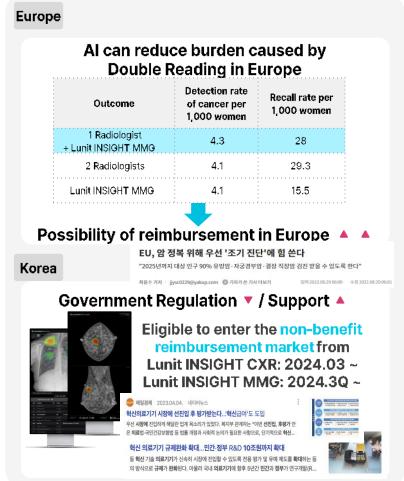
Active Reimbursement in Japan/Korea, **Expanding into Other Markets**

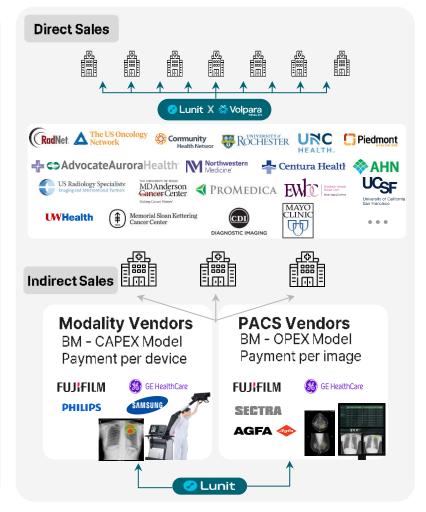


Company Overview

Expansion into Direct Sales

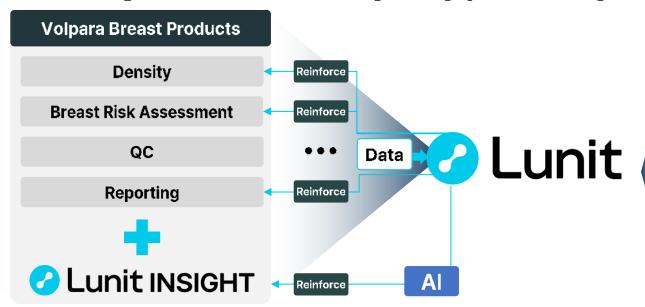






Lunit Acquires Volpara Health (May, 2024)





Future Direction

Lung cancer screening

XX cancer screening

Expand Data-Al flywheel model in all cancer screening











- Stable revenue (>\$30m in ARR in 2024)
- Large-scale data for training (imaging + clinical data)
- Distribution channel in US for Lunit INSIGHT
- KOL engagement in US for autonomous Al
- Enable business development for lung and other cancer screening product suites

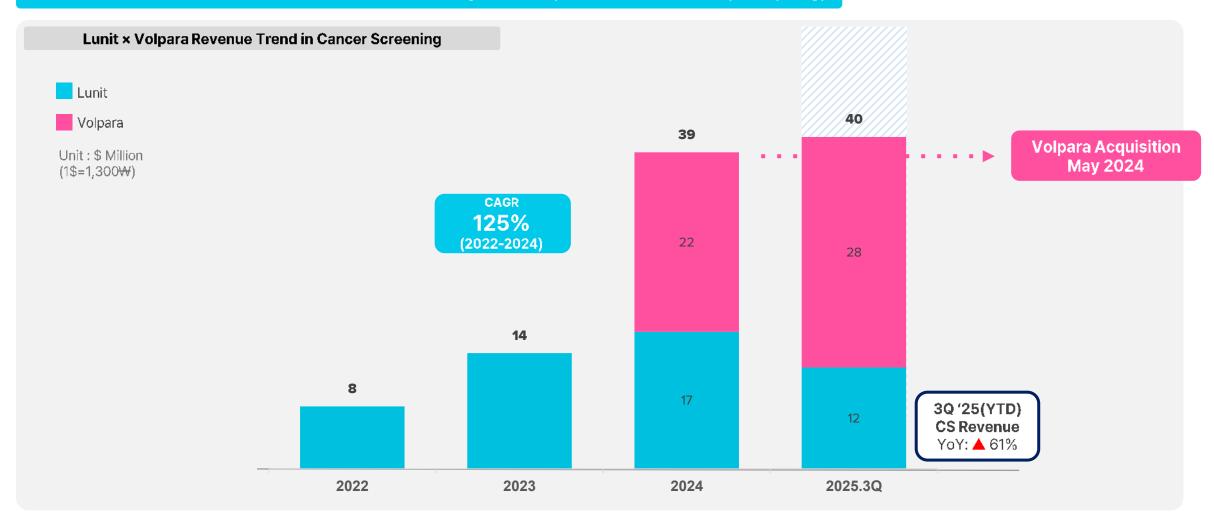


- Boost Al capabilities to strengthen all products
- Synergy with Lunit INSIGHT enables faster market penetration of Volpara products
- Distribution of Volpara products in non-US territories
- Enable development of lung cancer and other cancer related screening

Lunit SCOPE

Strong Revenue Growth Driven by Lunit x Volpara Synergy

Accelerated and Sustained Revenue Growth in Cancer Diagnostics Expected from Lunit x Volpara Synergy



Direct to Patient Business Model | SecondRead Al Gaining Strong Traction in US Market

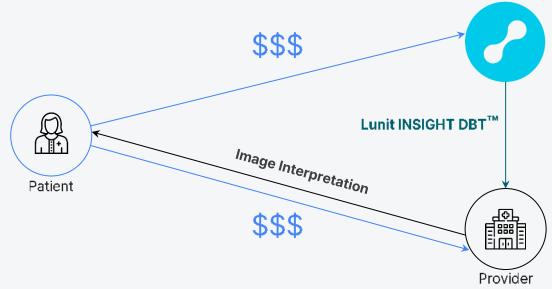
Traditional Business Model



New Business Model

Provider offers SecondReadAl™ direct to patients for an out-ofpocket fee at registration or point of care.

Revenue per analysis is x10-20





15

SecondReadAl Expected Opt-in Rate > 50%

Powered by Lunit

Lunit Cancer Screening

Lunit INSIGHT Risk Assessment Product

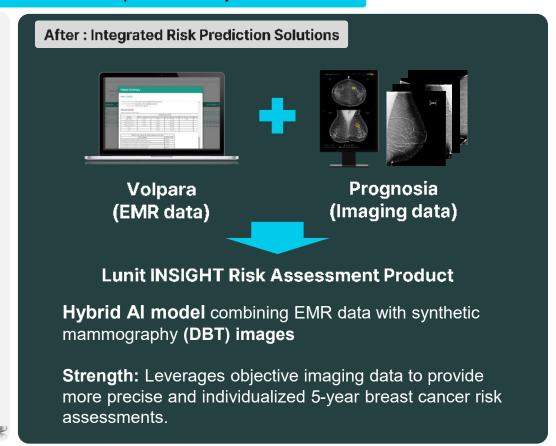
Establishing a new standard for early breast cancer detection and personalized preventive care

Before: Risk Assessment Solutions

Volpara 'Risk Pathways'

- A risk assessment software designed to identify and manage high-risk patients for breast, lung, colorectal, gastric, and pancreatic cancer
- Assesses risk based on **EMR data** such as family history, genetic information, lifestyle, etc
- Personalized prediction is **limited** due to the absence of imaging information





Product Development Timeline

2025.06 Completion of Prognosia acquisition and product integration

Late 2025 FDA submission

Q3 2026 **Product launch**

A New Standard for Personalized Screening

16

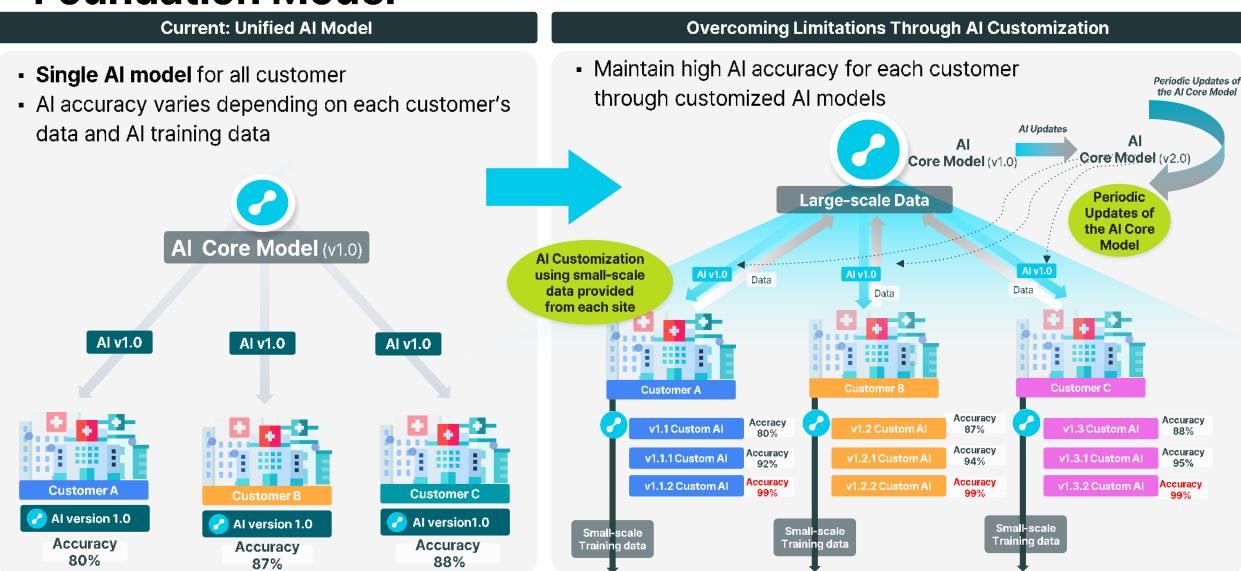
Prognosia Business Strengths

- •FDA (Breakthrough Device Designation): Recognized for technological and clinical value, enabling accelerated market entry
- •Optimized for U.S. market: Validated with U.S. SEER-based data and aligned with major guidelines (NCCN, USPSTF, ASCO)

Lunit^e

17

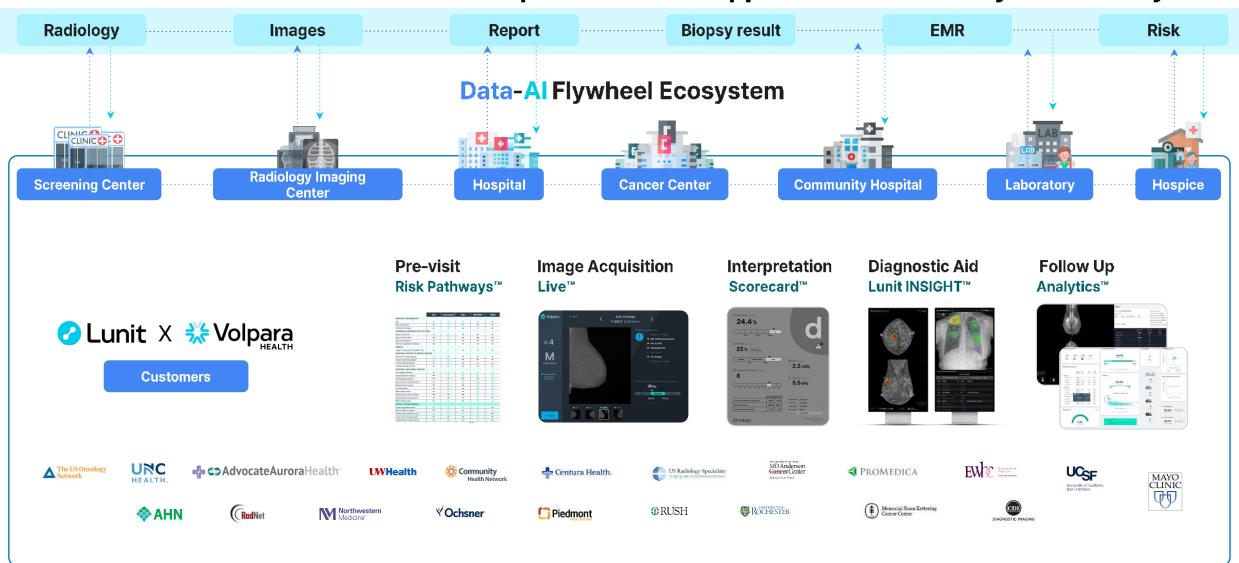
Foundation Model



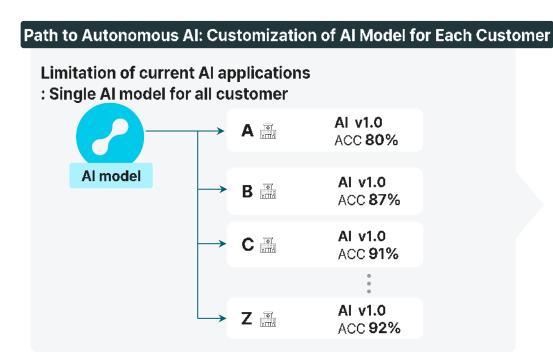
Lunit^e

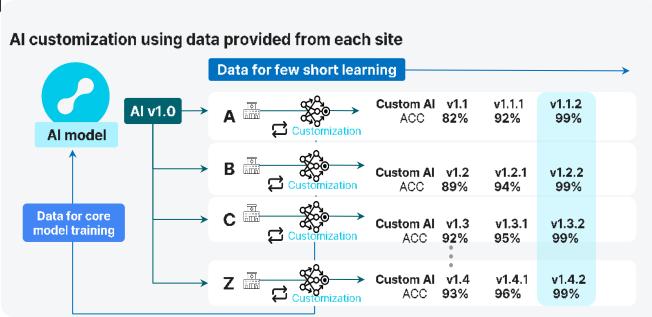
Lunit Cancer Screening

Customization Al for Autonomous Al Requires Platform Approach of Data-Al Flywheel Ecosystem



Our Future is Autonomous Al



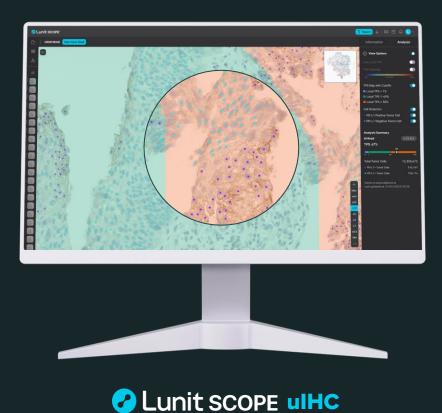


Expected Benefits of Autonomous AI Expected Performance with Autonomous AI More than 50% More than 94% reduction in costs reduction in FP marks 2.23 Iterative 24/7 No Increased Personalization Self-improvement **Efficiency Availability Human Errors** Human ΑI C-CAD Lunit Al

Lunit scope®



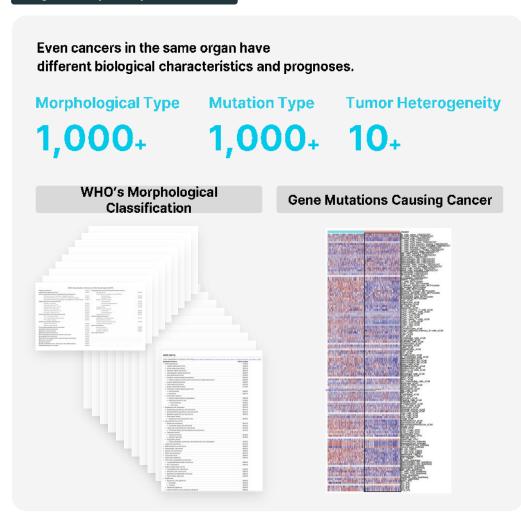




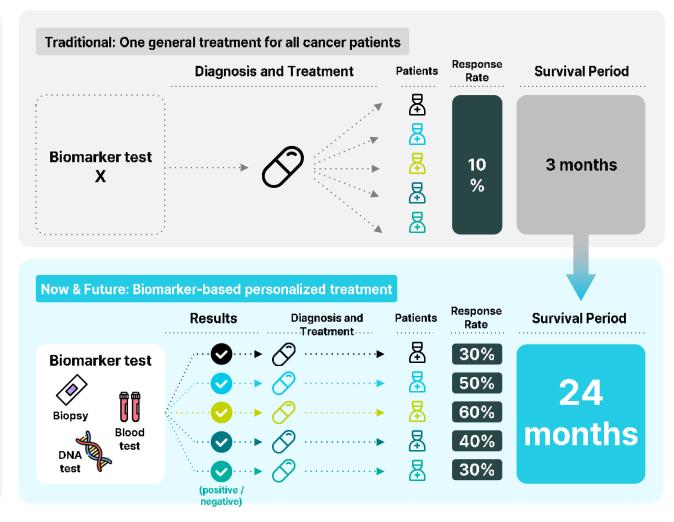


Personalized Approach Is Key to Cancer Treatment

High Complexity in Cancer

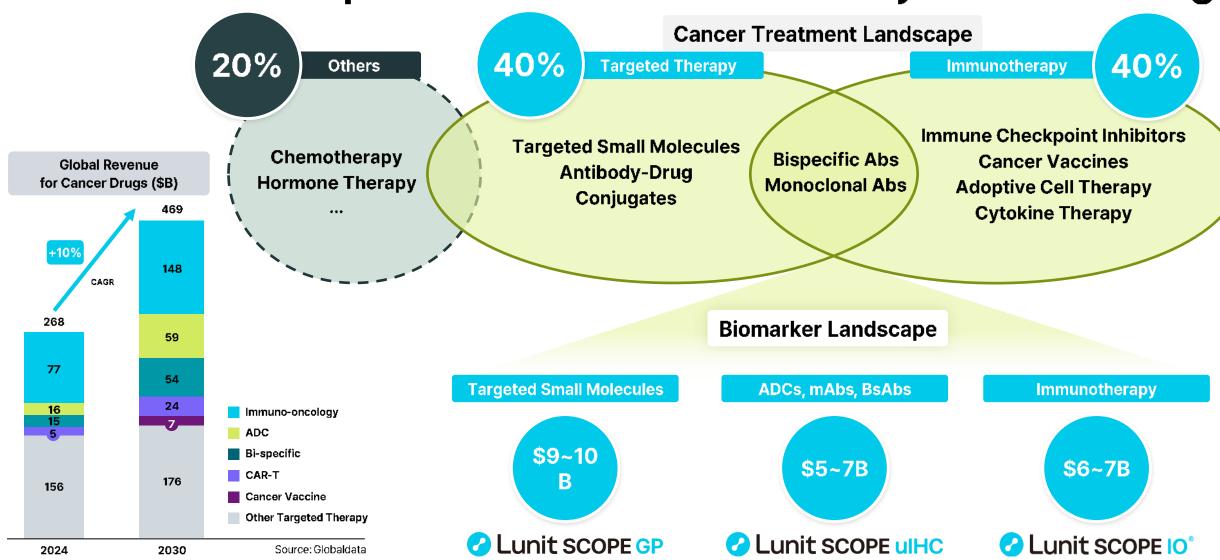


KEY: Personalized Treatment by Biomarkers



Lunit SCOPE Development

Lunit SCOPE: Al-powered Biomarkers for Major Cancer Drugs

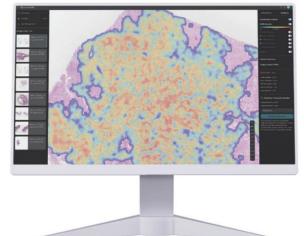


Lunit SCOPE Overview

Lunit SCOPE: Al-powered Biomarkers for Major Cancer Drugs













Purpose	Biomarkers for IO therapeutics	Biomarkers for ADCs therapeutics	Screening for multiple biomarkers
Key Features	ldentification of predictive tissue/cell features (H&E)	Quantification of protein expression level/pattern (IHC)	End-to-end prediction of specific genomic mutations
Potential Applications	CDx biomarkers for IO drugs Complimentary biomarkers	CDx biomarkers for ADC drugs	Prescreening genomic mutations Patient enrollment in clinical trials
Related Therapies	Immune Checkpoint Inhibitors	Targeted therapy - mostly ADCs	Targeted therapy - small molecules

Proven to Be Effective in Pharma Clinical Trials

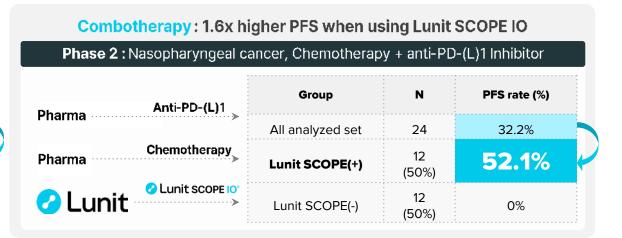
Monotherapy: 3x more responsive when using Lunit SCOPE IO Phase 1: "multiple cancer", anti-PD-(L)1 Inhibitor Ν ORR Group Anti-PD-(L)1 Pharma 32 All analyzed set 21.9% 62.5% Lunit SCOPE(+) Lunit SCOPE 10° (25%) Lunit 24 Lunit SCOPE(-) 8.3% (75%)

+20Global Big Pharma

Currently discussing research agreement/ collaboration with leading global pharmas

Combotherapy: 2x more responsive when using Lunit SCOPE IO				
Phase 1/2 : Colorectal cancer (MSS), TGF-beta Inhibitor + anti-PD-(L)1 Inhibitor				
	Group	N	ORR	
TGF-beta inhibit Pharma	or → All analyzed set	31	12.9%	K
Pharma Anti-PD-(L	.)1 Lunit SCOPE(+)	16 (51.6%)	25.0%	P
Lunit SCOPE	Lunit SCOPE(-)	15 (48.4%)	0%	

Monotherapy: 2.6x more responsive when using Lunit SCOPE IO Phase 2: Rectal cancer, Chemoradiotherapy + anti-PD-(L)1 Inhibitor Ν pCR Group Anti-PD-(L)1 38 28.9% Pharma All analyzed set 75.0% Lunit SCOPE(+) (21%)Lunit SCOPE 10° Lunit 30 16.7% Lunit SCOPE(-) (79%)



Key Clinical Studies for Lunit SCOPE



Lunit SCOPE uIHC

High Prediction of Response to Immunotherany in Lung Cancer

ininianomerapy in Lung Cance
[Regardless of PD-L1 status]
[Regardless of PD-L1 status]
ORR - 2.5 times higher
URR = 7 3 HHIES HIGHEL

OKK - 2.5 tin	nes nignei
Patient Group	Overall Response Rate
Lunit SCOPE IO Positive	26.8%
unit SCOPE IO Negative	11.3%

[PD-L11~49% Group] ORR - 6 times higher

Patient Group	Overall Response Rate
Lunit SCOPE IO Positive	22.8%
Lunit SCOPE IO Negative	3.9%

Expanding Lunit SCOPE IO to All Cancer Types

[All cancer types] ORR - 1.7 times higher

Patient Group	Overall Response Rate
Lunit SCOPE IO Positive	26.3%
Lunit SCOPE IO Negative	15.8%

uIHC based AI Quantification for Various Antibodies Protein Expression Analysis

Staining (Antibodies)	No. of Negative TC / Positive TC	Cell detection performance (mF1)	TP\$ classification performance (Accuracy)
TROP2	1243 / 3289	78.9%	94.0%
MET	3493 / 1134	80.5%	96.0%
Claudin 18.2	2927 / 74	72.7%	94.0%
DLL3	2973 / 3	68.9%	100%
HER3	2768 / 138	60.9%	94.0%
FGFR2	2358 / 438	50.8%	81.0%
E-Cadherin	443 / 912	79.2%	90.0%

uIHC based AI Quantification **Shows Predictive Power in HER2**

Group	n	ORR, N (%)	PFS, months, median (95% CI)	HR (95% CI)	P
AI-H3-low	11	0	1.4 (1.3-NR)	Ref	
Al-H3- high	19	8 (42.1)	4.4 (4.0-12.0)	0.12 (0.04-0.38)	< .001

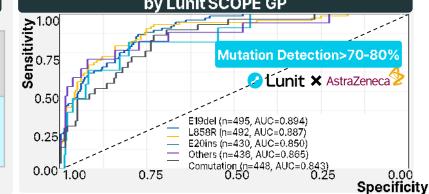
Survival Rate: 3x Gap

🕜 Lunit SCOPE GP

Prediction of Various Lung Cancer Driver Mutations by Lunit SCOPE GP

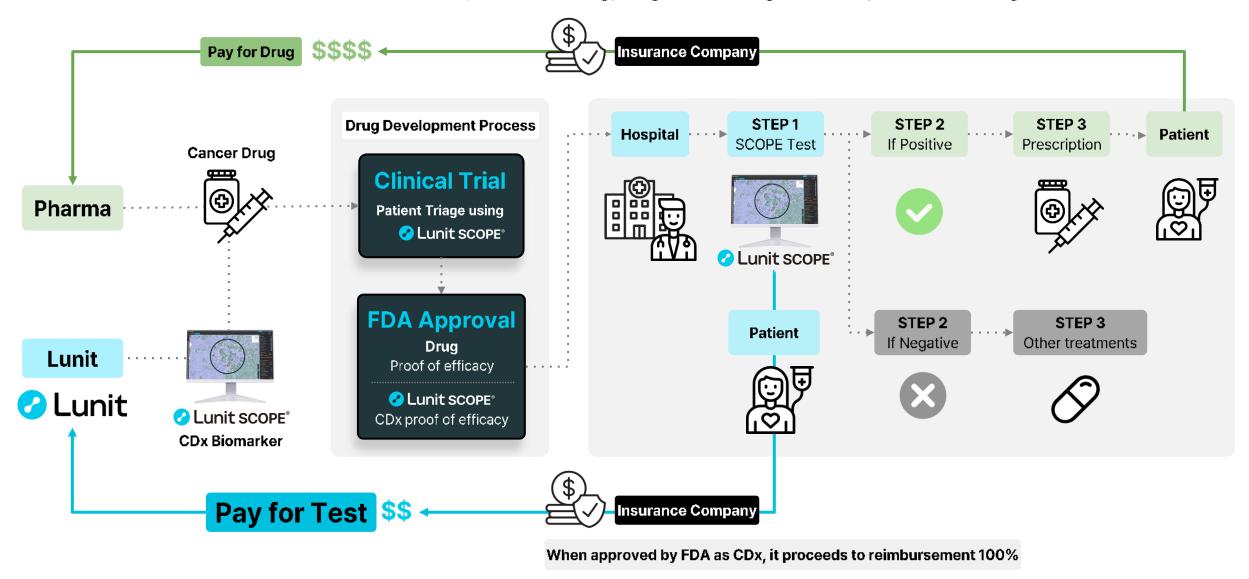
	Ex	External Validation set				
	N (%)	N (%) AUC Sensitivity		Specificity		
EGFR-mt	224 (28.3)	0.723	75.5%	52.6%		
KRAS-mt	130 (16.4)	0.721	86.2%	22.5%		
ALK-tr	46 (5.8)	0.738	39.1%	81.8%		
ROS1-tr	62 (7.8)	0.609	14.5%	70.6%		
<i>RET</i> -tr	11 (1.4)	0.683	18.2%	96.4%		
MET-ex	12 (1.5)	0.849	83.3%	74.4%		
All-WT	277 (35.0)	N/A	15.9%	99.2%		

EGFR Mutation Subtype Prediction by Lunit SCOPE GP



Lunit SCOPE BM

Business Model Paired with pharma's drug, to generate high sales by active testing before treatment



Commercialization Roadmap From Pharma Research to Companion Diagnostics (CDx)

Step 1: Biopharma Engagement

Step 2: Broader Use, RUO Service

Step 3: As a CTA

CDx Sales for Clinical Use

Pilot or Bake-off Study

Active Pharma

15+

Translational Research

Active Pharma



Clinical Trial Enrollment

Apply SCOPE in Clinical Trials

Screening patients with SCOPE for patient enrollment in clinical trials OR Analyzing existing clinical trial data with SCOPE

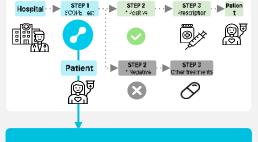


FDA Approval

Regulatory Approval as CDx

Drug Proof of efficacy

Lunit SCOPE® CDx proof of efficacy



Per Test Payment for Clinical Use

Research Revenue: Per Analysis Payment from Pharma Partners

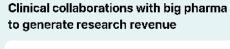
Global Top20 Pharma of 2024 based on oncology drug revenues

1	Merck	11	Astellas
2	BMS	12	Incyte
3	Roche	13	Takeda
4	J&J	14	Gliead
5	AZ	15	Sanofi
6	Novartis	16	BeiGene
7	Pfizer	17	Daiichi
8	Amgen	18	Bayer
9	Eli Lilly	19	GSK
10	Abbvie	20	Teva

We are engaged with



Lunit SCOPE Revenue Trend (RUO Sales)







Increase in US Labs that order PD-L1 tests



Lunit SCOPE Market

Cancer Biomarker Market

\$27B+



Strong Traction with Major Industry Vendors for Future Distribution Scalability

Pathology Labs

Top 5 Anatomic Pathology Labs Based on Revenue

- Labcorp
- Quest
- **Sonic** 3
- **Neogenomics**
- **Pathgroup**



Among Top5 Labs, We are engaged with - as well as other key CROs



Among Top5 antibody makers, We are engaged with





Staining Component

Antibody Manufacturer



Digitization Component

Scanner Manufacturer



Viewing Component

Path-PACS Manufacturer

















HAMAMATSU







Most pathology scanners are compatible with Lunit's Al

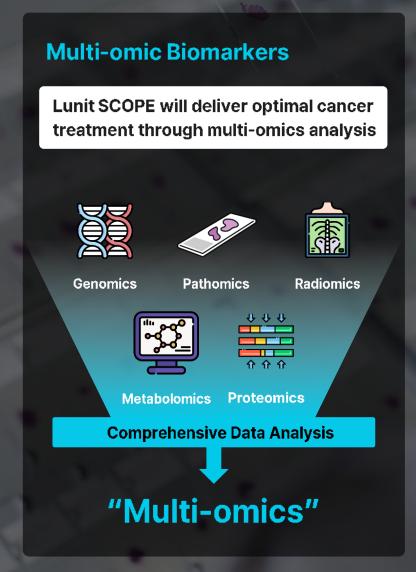


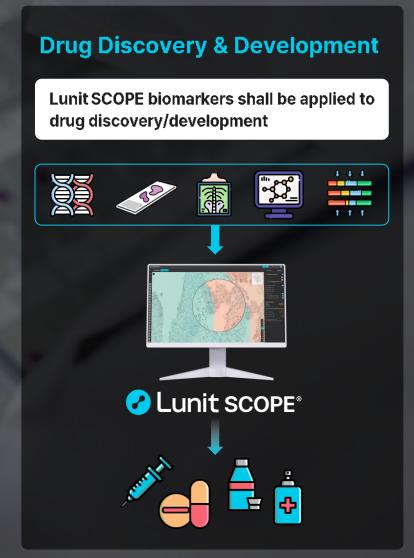
Example companies:



Next Steps

Cover All Types of Drugs Lunit SCOPE will be able to cover all types of oncology drugs Final Step **Hormonal** Immuno Therapy Therapy **Global Oncology** Drug Market 2029 Chemo **Targeted Therapy** Therapy **Second Step** Source: Maximize market research PVT. LTD

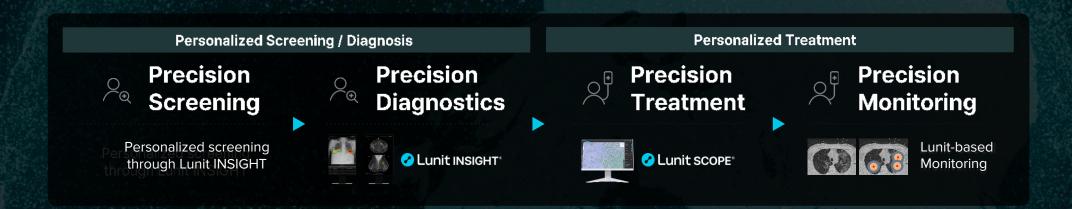




Conquer Cancer through Al-powered Precision Medicine

Having strong faith that a data-driven approach is the future,

Lunit will be the center of cancer care Globally become the new standard of care



THANK YOU



Appendix

Consolidated Financial Statements

Statement of Financial Position

Unit:KRW million

				Unit:KRW millio
	3Q 2025	2024	2023	2022
Current Asset	65,594	111,062	230,418	68,050
Non Current Asset	323,651	324,215	37,503	29,103
Total Assets	389,245	435,277	267,921	97,153
Current Liabilities	206,047	252,736	12,794	9,446
Non Current Liabilities	18,521	17,767	20,017	18,797
Total Liabilities	224,568	270,502	32,811	28,243
Capital Stock	14,602	14,471	14,351	6,114
Additional paid-in Capital	543,014	526,646	515,518	320,857
Others	157	1,495	181	142
Retained earnings	(393,096)	(377,837)	(294,941)	(258,202)
Total shareholders' equity	164,677	164,774	235,110	68,910

Income Statement	Unit:KRW million

	3Q 2025	2024	2023	2022
Operating Income	56,653	54,180	25,080	13,866
Operating Cost	120,120	121,833	67,297	64,517
Operating Income(Loss)	(63,467)	(67,654)	(42,217)	(50,651)
Financial Income	85,157	19,202	10,257	13,642
Financial Expenses	36,525	34,721	4,764	2,060
Other Income	200	107	80	43
Other Expenses	44	121	119	92
Income(Loss) before Tax	(14,678)	(83,187)	(36,764)	(39,118)
Tax Expenses(Income)	266	(758)	35	-
Net Income(Loss)	(14,943)	(82,428)	(36,798)	(39,118)

Source: DART, Company K-IFRS Audited