



Veritas In Silico

This document has been translated from the Japanese original for reference purposes only. In case of discrepancy, the Japanese original shall prevail.

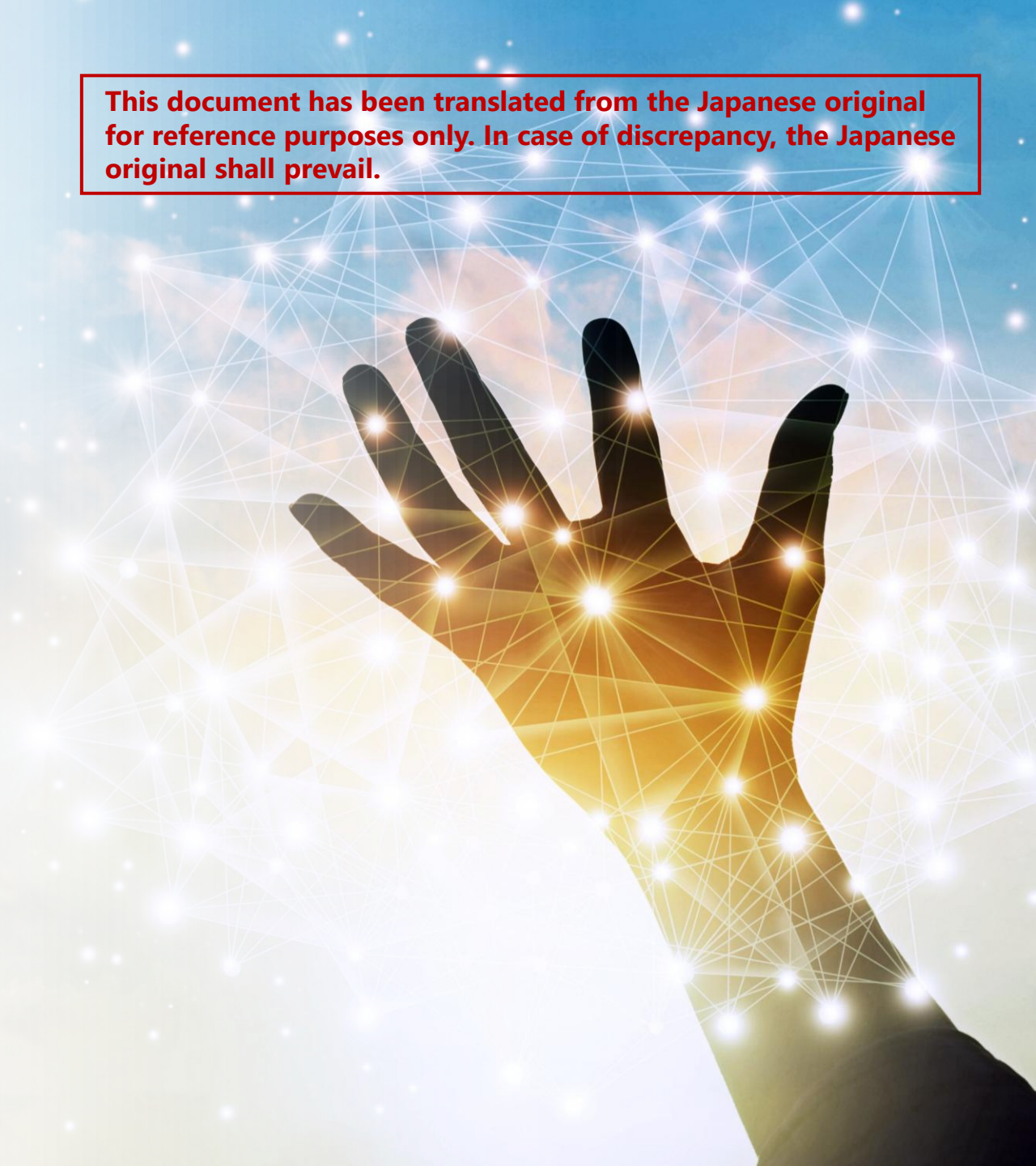
Fiscal Year Ended December 31, 2025

Financial Results

Veritas In Silico Inc.

Ticker code: 130A

February 12, 2026



Disclaimer

This document has been prepared solely to provide information on Veritas In Silico Inc. (the "Company"), and does not constitute an offer to sell or a solicitation of an offer to buy any securities in Japan, the United States or any other jurisdiction. The Company's securities may not be offered or sold in Japan, the United States or any other jurisdiction without registration or a filing, or an exemption therefrom, under the applicable laws and regulations.

In preparing this material, the Company has relied upon and assumed the truth, accuracy and completeness of the information available to it at the time of its preparation, and with respect to forward-looking information, external data and the like, the Company makes no representation or warranty as to their truth, accuracy or completeness. The information contained herein is subject to change without prior notice.

Statements made in this document with respect to future business activities and performance are forward-looking statements. Forward-looking statements include, but are not limited to, expressions such as "aim," "forecast," "assume," "believe," "continue," "attempt," "estimate," "anticipate," "measures," "intend," "plan," "may," "plan," "potential," "probability," "project," "risk," "pursue," "should," "strive," "target," "scheduled," or other similar expressions that describe future business activities, performance, events or circumstances. Forward-looking statements are based on the judgments of the Company's management in light of the information available to it at the time this material was prepared and involve a number of risks and uncertainties. Accordingly, such forward-looking statements are subject to various risks and uncertainties that could cause actual future business activities and results to differ materially from those expressed or implied by the forward-looking statements. Therefore, you are cautioned not to place undue reliance on forward-looking statements.

Highlights for FY2025

Platform Business

- Joint drug discovery research progressed, with a milestone achieved in May with Shionogi.
 - US patent for ibVIS® core technology was granted in July, securing IP in Japan, the EU, and the US.
-

Pipeline Business

- 1st In-house pipelines (AKI* prevention drug): ASO disease treatment project progressed, Patent filed in Dec., KPI achieved.
 - June: Signed joint research agreement with Mitsubishi Gas Chemical for nucleic acid drugs.
-

Business revenue for FY2025 was 91 million yen, and Business expenses were 487 million yen, resulting in the Net loss of 425 million yen.

- Business Performance essentially in line with the forecast (as disclosed on Oct. 14),
 - An Impairment loss of 31million yen incurred as an Extraordinary loss.
 - R&D investment was front-loaded in order to generate 1st In-house pipelines and new technology development.
-

Business revenue for FY2026 is estimated at 113 million yen, and an Ordinary loss at 564 million yen, resulting in a Net loss of 567 million yen.

- Anticipate upfront expenditures, including R&D investments for 2nd In-house pipelines generation, relocation of the research institute, and capital expenditures for facility enhancement.

Background to Our Strategy: Initiatives aligned with Industry Trends

In the pharmaceutical industry, major pharma companies are increasingly prioritizing the in-licensing of drug candidates and scaling back in-house drug discovery. Pharma companies in Japan and Europe, which had been cautious about this shift, are now rapidly following suit, and opportunities for platform agreements with big pharma are globally declining.

Platform Business: Expanding opportunities while aligning with industry trends

01

- Advance ongoing joint drug discovery to establish a success case in small-molecule drug discovery against mRNA targets
 - Demonstrate a technical breakthrough to shift the industry trends
- Conclude agreements with highly motivated for drug discovery biotech companies in Japan and overseas, and advance projects with strong value potential
- Continue pursuing contracts with major pharmaceutical companies to realize projects that generate early revenue

02

Expand in-house pipeline: Create drug candidates aligned with pharmaceutical company needs

- For pipeline 1, leverage DDS to fundamentally reduce clinical trial cost and duration, aiming to enhance future value and enable early monetization
- For pipeline2, create in FY2026

03

Enhance business stability via diversification

- Capitalize on the applicability of our AI drug discovery platform aibVIS to expand into agrochemicals
- Consider revenue generation by licensing out the DDS



Contents

- 1 Corporate Overview**
- 2 Business Highlights**
- 3 Financial Highlights**
- 4 FY2026 Forecast**



1 Corporate Overview

Corporate information

A Biotechnology Company Specializing in AI drug discovery, Led by a small, Highly Skilled Team

Name	Veritas In Silico Inc. (VIS)
Established	November 17, 2016
Main office	1-11-1 Nishigotanda, Shinagawa-ku, Tokyo 141-0031, Japan
Research facilities	Shinkawasaki Research Institute: Kanagawa, Japan Niigata Research Institute: Niigata, Japan
CEO	Shingo Nakamura
Employees	19 (As of December 31, 2025)
Business description	Creating mRNA-targeted small molecule drugs with pharmaceutical companies as partner using our proprietary drug discovery platform, aibVIS , as well as developing its pipelines from 2025.



Shinkawasaki Research Institute
(At the Kawasaki Business Incubation Center)

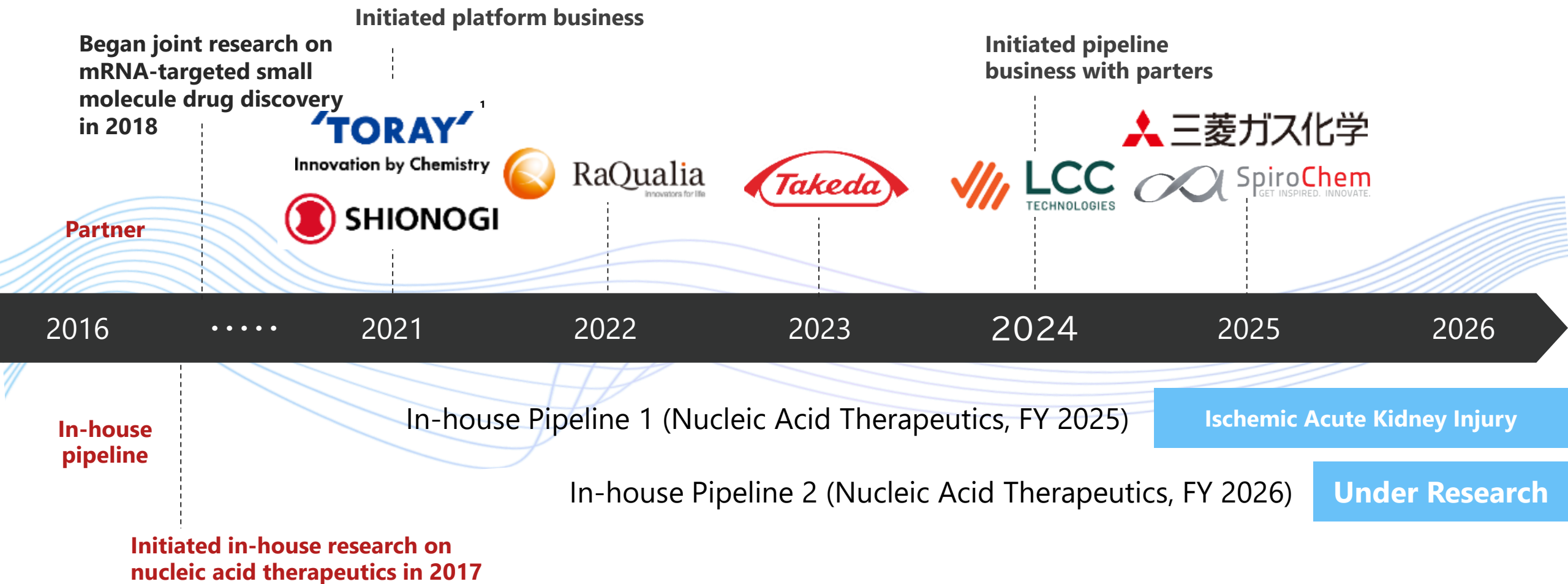


Niigata Research Institute
(At the Niigata University of Pharmacy and Medical and Life Sciences)

History

Balanced Approach: Partner Drug Discovery & In-house Pipeline

In addition to our joint drug discovery targeting mRNA with small-molecule therapeutics, we are forming partnerships with multiple companies to create pipelines. Furthermore, from FY2025 onward, we will drive the creation of one new in-house pipeline per year.



Our Business Portfolio Leveraging the Application Capabilities of **aibVIS**

Leveraging the high versatility and scalability of our proprietary AI drug discovery platform, **aibVIS**, to strategically expand multiple business opportunities.

Platform Business

Small-molecule pharmaceuticals targeting mRNA

A solution for diseases lacking effective therapies under conventional protein-targeted small-molecule discovery, and for needs met only by high-cost treatments

In-house pipeline

Nucleic acid drugs

aibVIS enables rapid creation of nucleic acid drugs, offering new treatment possibilities for rare diseases

Platform Business

Small-molecule agrochemicals targeting mRNA

aibVIS can be applied not only to human medicines but also agrochemicals, enabling agrochemicals with minimal impact on the environment and humans

Out-licensing

Drug delivery system (DDS)

Effective drug delivery can fundamentally solve challenges associated with nucleic acid drugs

Business diversification
driven by

aibVIS

:strengthening
the conventional AI drug
discovery platform with
specialized AI.

Our Business Portfolio Leveraging the Application Capabilities of aibVIS

10 years from the start of research until the product is launched on the market¹



Revenue structure that secures stable revenue from the early stages of drug discovery through joint research



Revenue structure that prioritizes creating in-house pipelines and out-licensing them to obtain large upfront payments and milestones



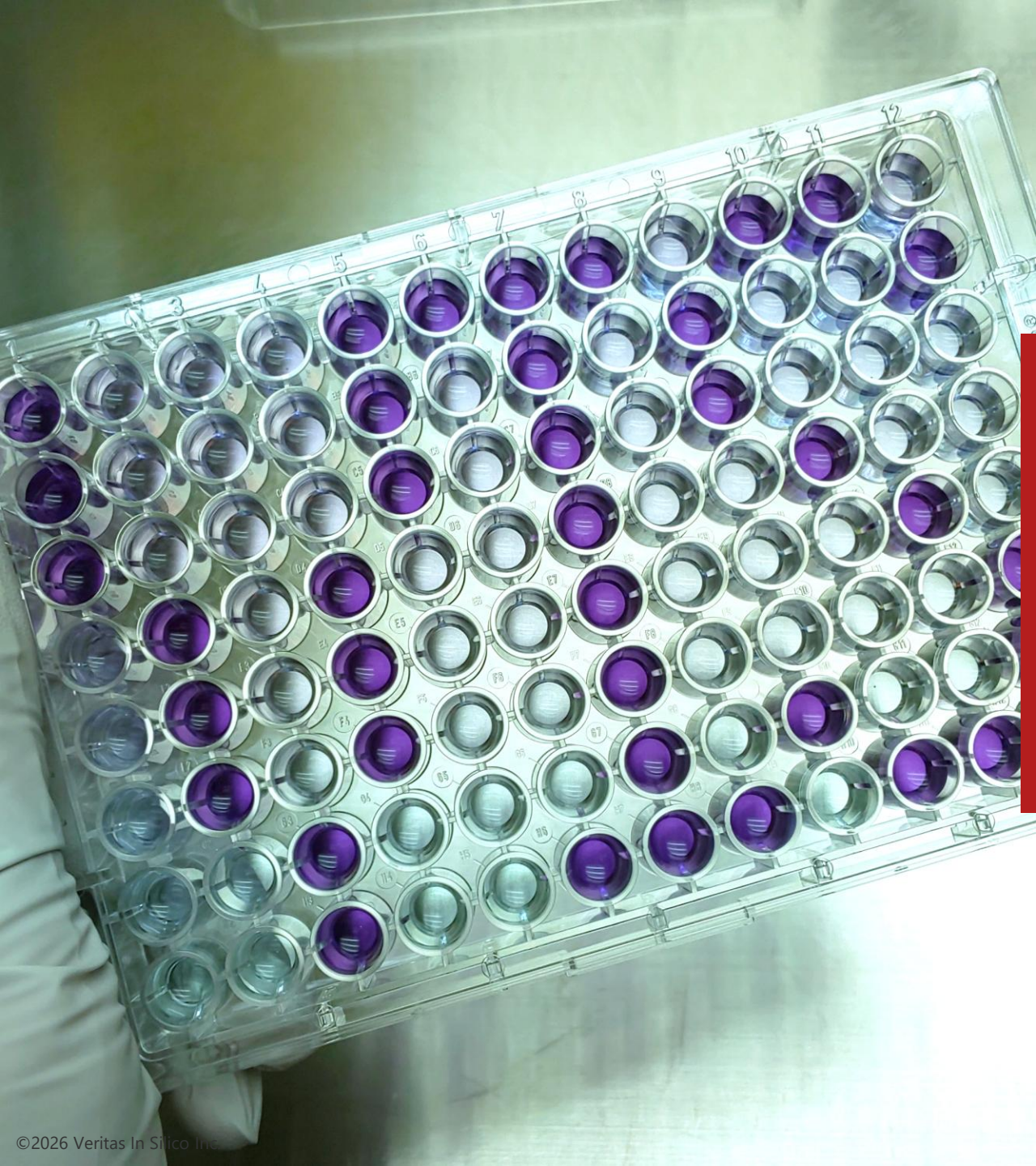
↑ Revenue Generation Point

¹ The average duration is based on the typical duration of drug discovery R&D, and the actual R&D durations may differ significantly from those given here.

² At present (as of Dec. 31, 2025), no partner has progressed to the development and sales step.

³ The use of aibVIS is limited to the duration of the joint drug discovery research between our company and pharmaceutical companies.

⁴ In principle, the initial joint drug discovery research agreement includes arrangements regarding development, manufacturing, and sales licenses.



2

Business Highlights

Business Topics for FY2025

Platform Business

Core Technology of Drug Discovery Platform Patent Granted in US in July 2025

Patent for the analysis of target mRNA substructures and their occurrence probabilities, the identification of substructures suitable for drug discovery targets, and screening utilizing these—the core technology of the AI drug discovery platform ibVIS®.

Achieved Milestone in Joint Drug Discovery Research with Shionogi

Through joint drug discovery research for mRNA-targeted small molecule drugs, achieving high-activity compounds exhibiting specific effects according to the stringent standards of Shionogi.

KPI item

Agreement for a Joint research with SpiroChem

With combining SpiroChem's world-class expertise in compounds with VIS's **ai**bVIS, with aim to establish design criteria that will contribute to future drug discovery.

Pipeline Business

KPI item

Generate 1st In-house pipelines (Applying a patent for a substance)

As 1st In-house pipelines, a nucleic acid medicine project targeting ischemic acute kidney injury induced after cardiovascular surgery is underway.

KPI item

Signed Joint research agreement with Mitsubishi Gas Chemical Co., Ltd.

Agreed on joint research for nucleic acid drug (ASO) discovery and establishment of manufacturing methods

Patent obtained for a drug delivery system used in nucleic acid drugs

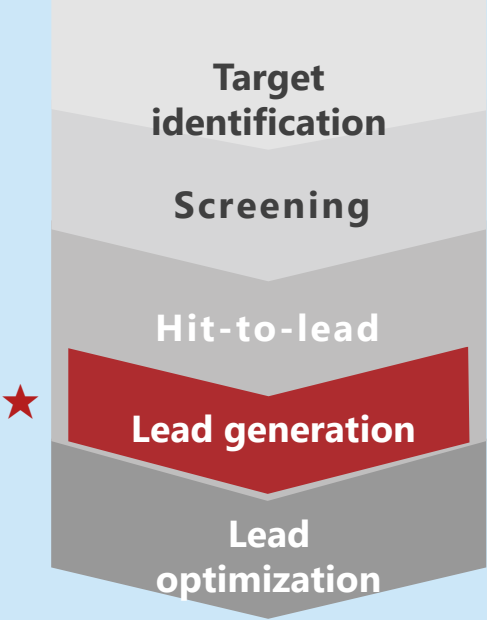
To fundamentally resolve the challenges facing nucleic acid therapeutics, we have acquired a patent for a drug delivery system (DDS). We aim for its early practical application.

Major Landmark in Platform & Pipeline Businesses Driving Future Revenue

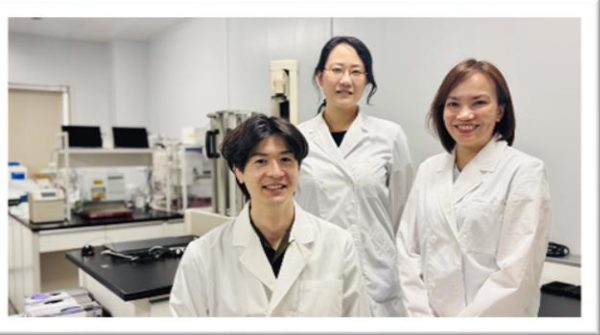
Platform business

Milestone achievement in joint research with Shionogi & Co., Ltd. on mRNA-targeted small molecule drug discovery

Stages of drug discovery



We successfully identified highly active compounds that exhibit specific effects on the targets difficult to achieve with protein targets, which meets the high standards of Shionogi.



Core members in this joint drug discovery research

Press release: [2025.05.23 Shionogi and Veritas In Silico Announce Milestone Achievement in Joint Drug Discovery Research of mRNA-targeted Small Molecules](#)

Pipeline business

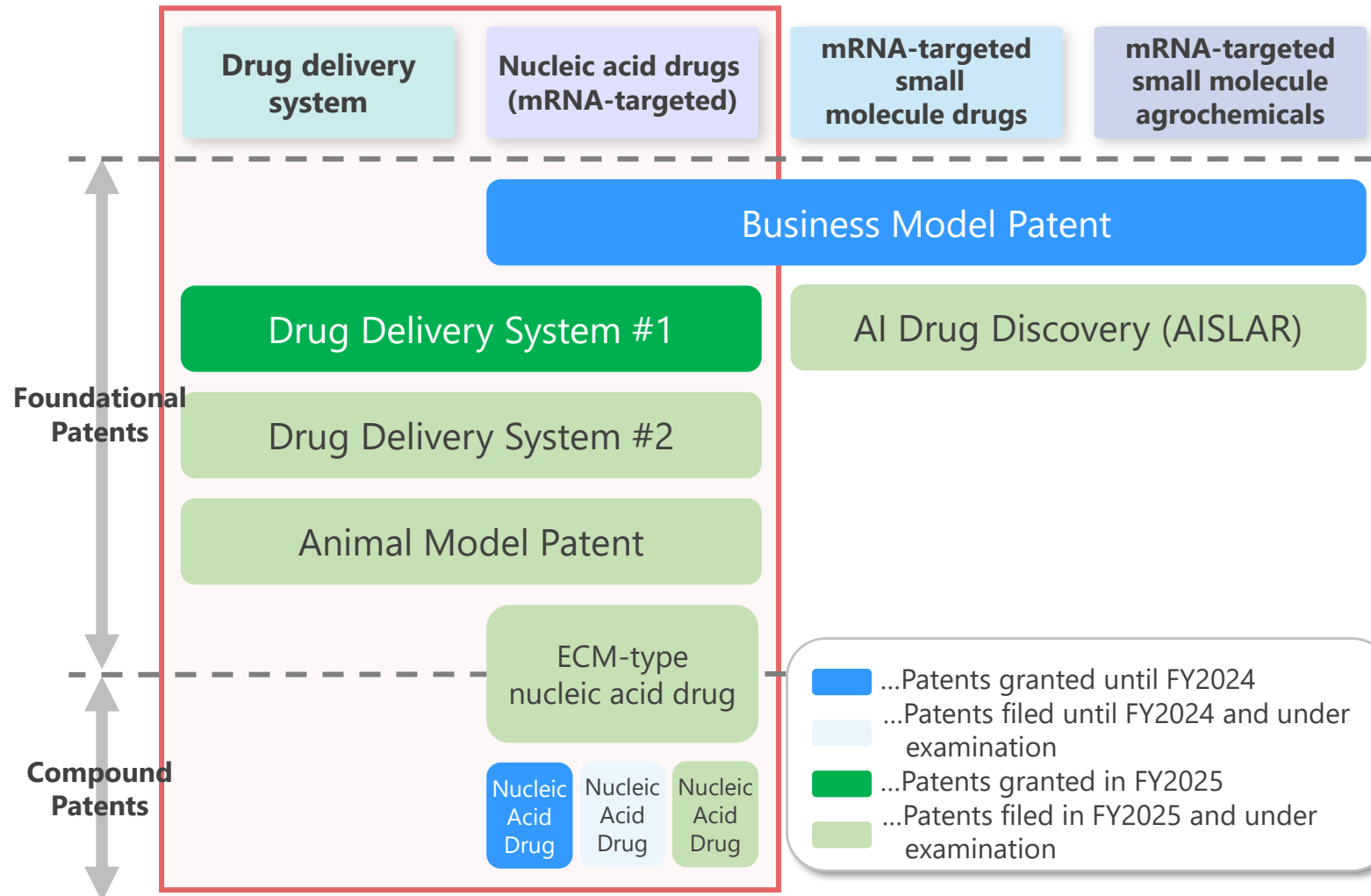
Joint research with Mitsubishi Gas Chemical, Inc. on nucleic acid drug discovery and manufacturing (First step for QbD).

Objective	Discovery of RNA-targeted nucleic acid drugs (ASOs) and establishment of manufacturing methods
Joint research term	3 years
Tech base	Utilize Veritas In Silico's drug discovery platform, aibVIS
Roles	<u>Veritas In Silico</u> : Acquisition of ASO compounds <u>Mitsubishi Gas Chemical</u> : Establishment of manufacturing methods
Key features	Incorporate QbD (Quality by Design) from the early stages of drug discovery research for rapidly moving to clinical trials

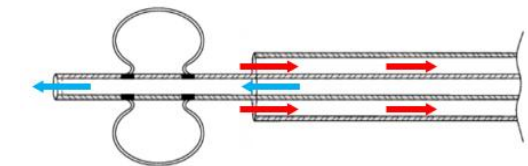
Release: [2025.07.03 Mitsubishi Gas Chemical and Veritas In Silico Announce Joint Research Agreement for Drug Discovery and Establishment of Manufacturing Methods for Nucleic Acid Drugs](#)

Positioning of our Drug Delivery System

By realizing new therapeutic methods utilizing the drug discovery system "Perfusio", we aim to expand our in-house pipeline value and revenue opportunities. Ultimately, by delivering drugs exclusively to target organs and avoiding systemic exposure, our goal is to drastically shorten clinical trials and reduce costs.



- Accelerating Pipeline Development: possibility of rapid realization of cost-effective, low-dose preventive drugs for ischemic acute kidney injury (AKI)
- As the approach leverages combinations of existing medical devices, it is well positioned for accelerated translation to the clinical setting
- Full-scale practical implementation and commercialization from 2026
- Out-licensing to pharmaceutical companies is within scope



Insert an occlusion balloon catheter through a standard catheter.

Utilizing Perfusio for Proprietary drug development is Our Core Strategy

For high-cost, highly potent medicines such as nucleic acid therapeutics, an appropriate drug delivery system is required to ensure precise delivery to the target organ. We will not only utilize Perfusio in-house but also aim to monetize it through out-licensing.

1

Establishing the operational structure

- Together with our partner medical device manufacturer, we will establish manufacturing and sales capabilities for the new catheter “Perfusio” as a general medical device usable across the entire body

2

Generalization

- Drive broader adoption of Perfusio-based drug delivery system
 - Actively promote external dissemination of results by Dr. Ota, who is conducting joint research with VIS
 - Permit clinical use for physician procedures/surgeries that use existing drugs approved for systemic administration with Perfusio at clinical setting because clinical trials are not required
 - **At clinical settings, permit uses that may implicate organ-perfusion patents, and adopt a policy of not charging licensing fees**

3

Use within VIS

- Create therapeutic drugs designed from the outset to be used with Perfusio
 - For in-house pipeline, aim to realize a new organ-perfusion therapy using Perfusio with novel therapeutics
 - Pursue a drug-approval pathway to drug approval that obviates clinical trials, use only on microdose toxicity testing

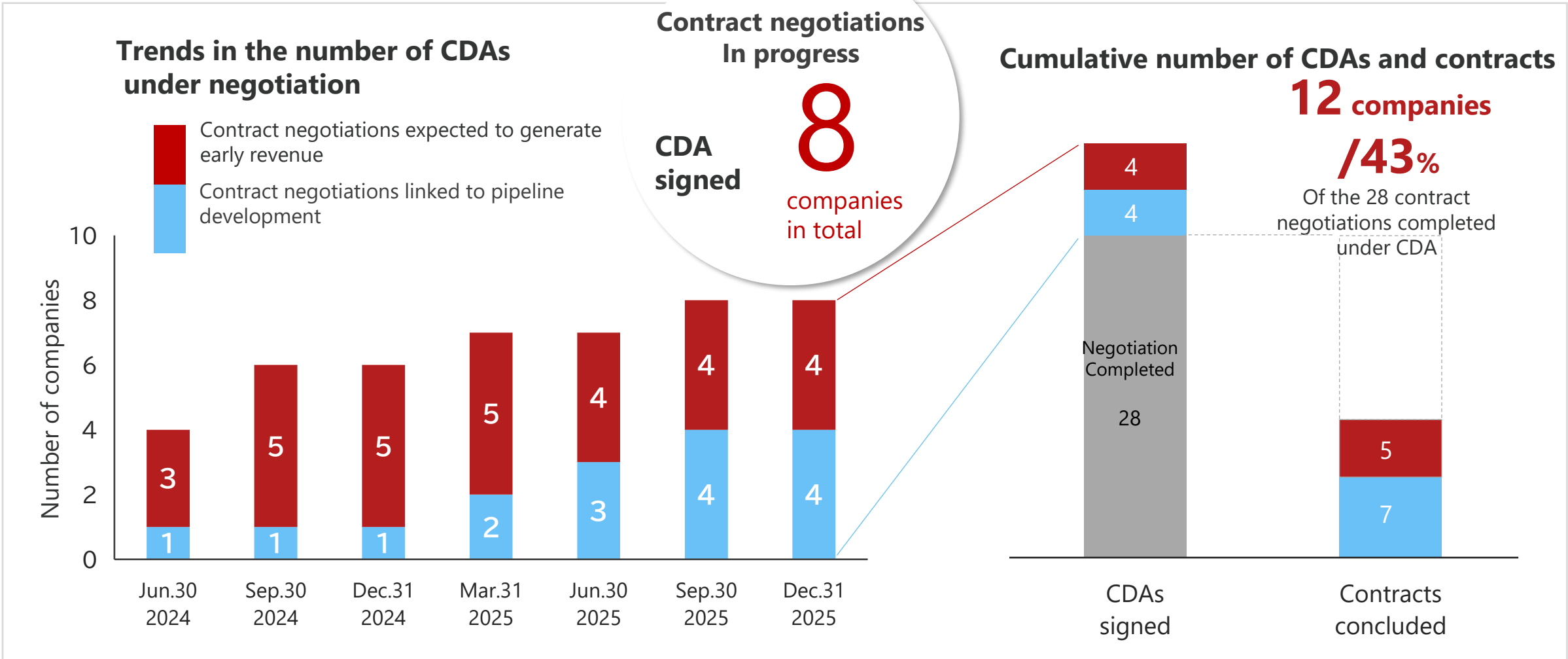
4

Driving additional revenue

- License out to pharmaceutical companies that wish to develop drugs using Perfusio, including for use during clinical trials

Secure CDAs and Target Early-Revenue Contracts

From 2026 onward, we will continue to secure an appropriate number of CDAs and aim to conclude two contracts per year, including at least one with early-revenue potential. Three out of eight progressing negotiations involve European companies. We will continue to pursue international expansion.



Progress toward Achieving KPIs for FY2025

Against an annual target of four new contracts, three were achieved. For our in-house pipeline creation, we filed a patent in December 2025, meeting the target.



New contracts
Four contracts within FY2025



LCC Technologies



Mitsubishi Gas Chemical



SpiroChem



In-house pipeline
One patent application within FY2025



Pipeline1

Amount of business revenue



(Note) As disclosed on October 14, 2025, regarding new contracts, one contract negotiation that had been delayed since fiscal year 2024 reached an impasse in terms of contract conditions and other points, making its completion within fiscal year 2025 difficult. As a result, the annual total of new contracts achieved was three. As of Dec. 31, 2025



3 Financial Highlights

Summary of FY2025 Business Results

Every Joint drug discovery research with our partners has been advanced, resulting in 91 million yen in Business revenue, including Milestone income and Research support funds.

Summary of P/L		(million of yen)		
	FY2024	FY2025	Change in amount	Change in rate
Business revenue	194	91	-103	53.2%
Business expenses	407	487	80	19.7%
Operating profit	-212	-396	-183	—
Non-operating Profit	-20	6	-17	—
Ordinary profit	-233	-390	-157	—
Net profit	-236	-425	-189	—

Breakdown

Milstone income: 10
Research support funds & Others: 81

R&D Expenses: 215
SGA Expenses: 272

Quarterly Performance Trends

Quarterly Performance (FY2023Q3~FY2025Q4) (million of yen)										
	FY2023 Q3	FY2023 Q4	FY2024 Q1	FY2024 Q2	FY2024 Q3	FY2024 Q4	FY2025 Q1	FY2025 Q2	FY2025 Q3	FY2025 Q4
Business revenue	29	81	32	83	49	29	24	19	22	25
Business expenses	80	83	97	85	104	120	105	124	124	134
Operating profit	-51	-2	-65	-1	-54	-91	-81	-105	-101	-108
Non-operating profit or loss	-1	0	-22	0	0	1	1	1	1	1
Ordinary profit	-53	-1	-87	-1	-54	-90	-79	-103	-100	-107
Net profit for the period	-53	-2	-87	-2	-55	-90	-79	-104	-101	-140

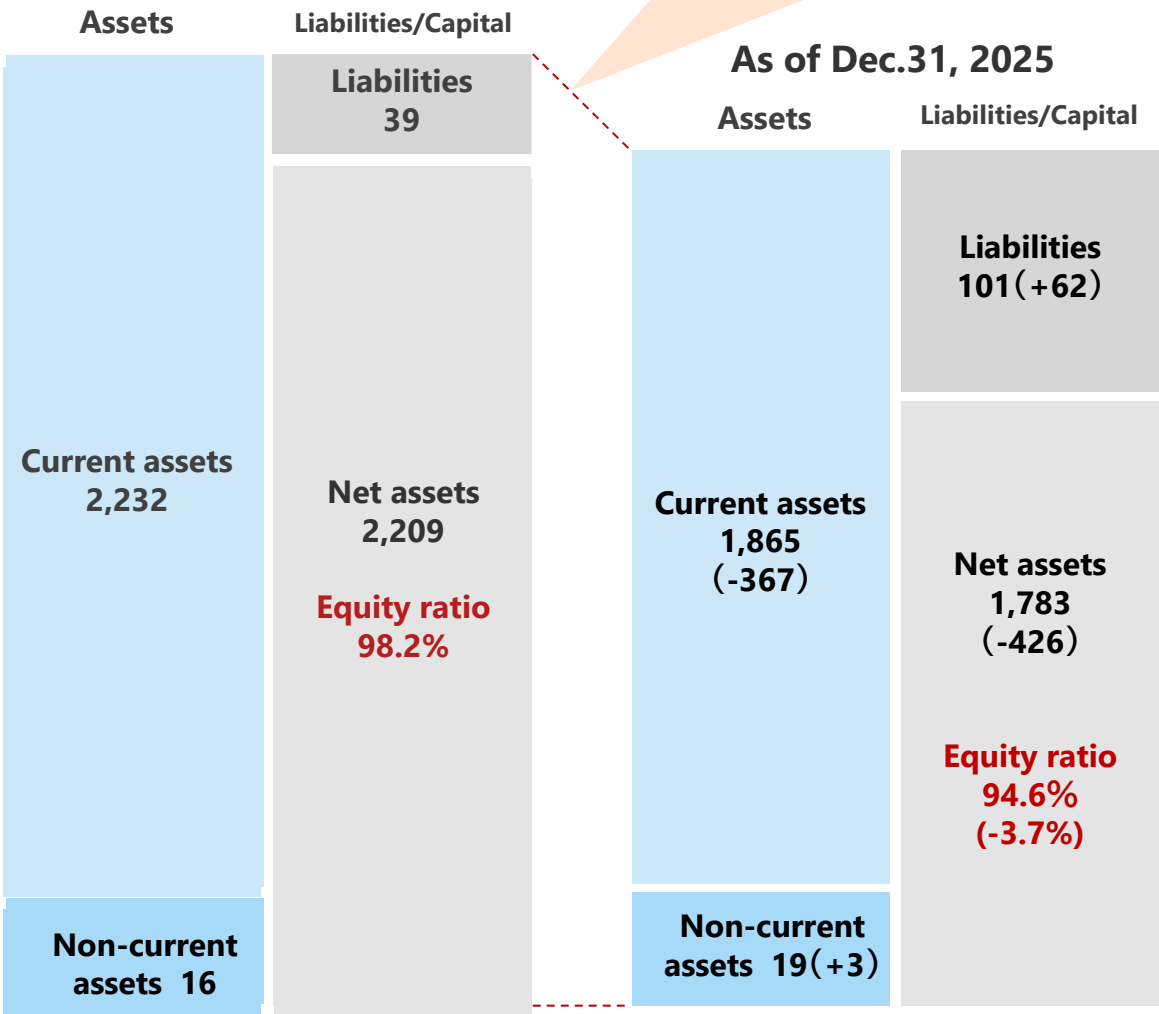
Financial Position

In Q2 2025, the Company performed a capital reduction: reducing Share capital from 77million to 10million yen, in order to enhance flexibility in our capital management.

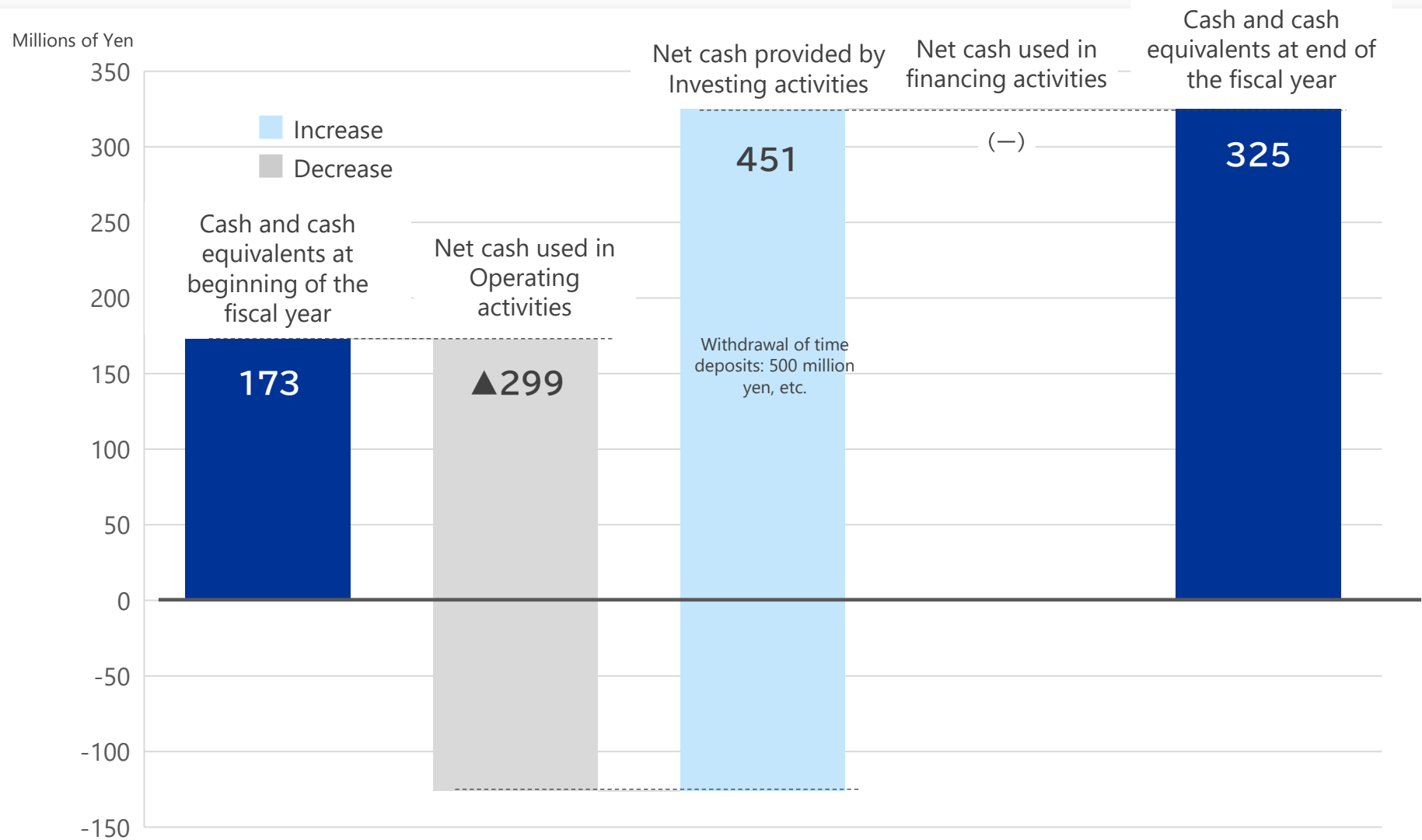
Balance Sheets (millions of yen)		
	As of Dec. 31, 2024	As of Dec. 31, 2025
Cash and deposits	2,173	1,825
Total current assets	2,232	1,865
Property, plant and equipment	14	0
Total non-current assets	16	19
Total assets	2,248	1,884
Total liabilities	39	101
Share capital	77	10
Total net assets	2,209	1,783
Total liabilities and net assets	2,248	1,884

Balance Sheets Transition (millions of yen)

As of Dec.31, 2024



Cash Flows for FY2025





FY2026 Forecast

4

FY2026 Annual Forecasts

Summary of forecasted P/L (million of yen)

	FY2025	FY2026 Forecast	Change amount	Change rate
Business revenue	91	113	22	24.2%
Business expenses	487	682	194	39.9%
Operating profit	△396	△569	△172	—
Non-operating Profit	6	4	△1	—
Ordinary profit	△390	△564	△173	—
Net profit	△425	△567	△141	—

Breakdown and Comparison to FY2025

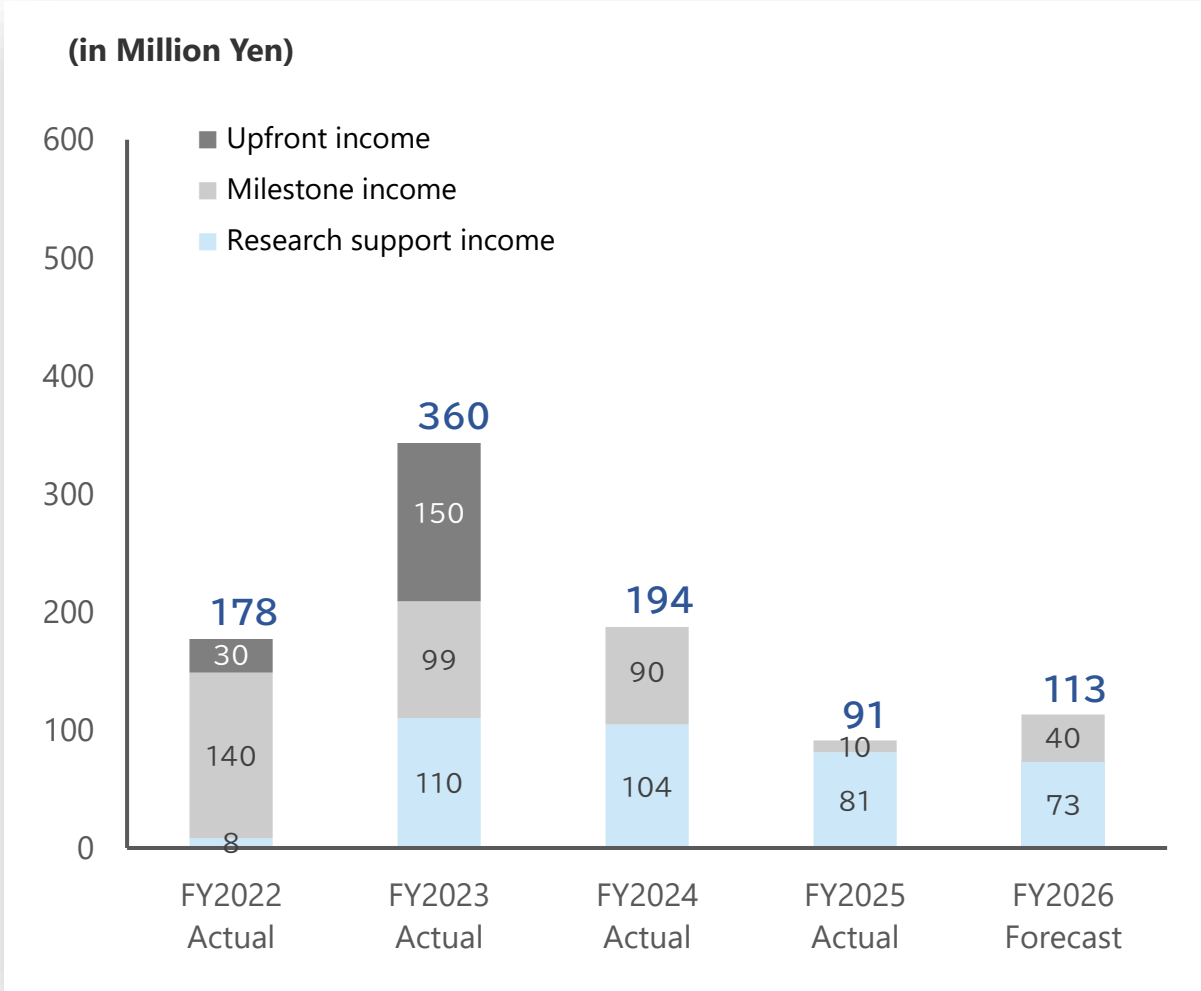
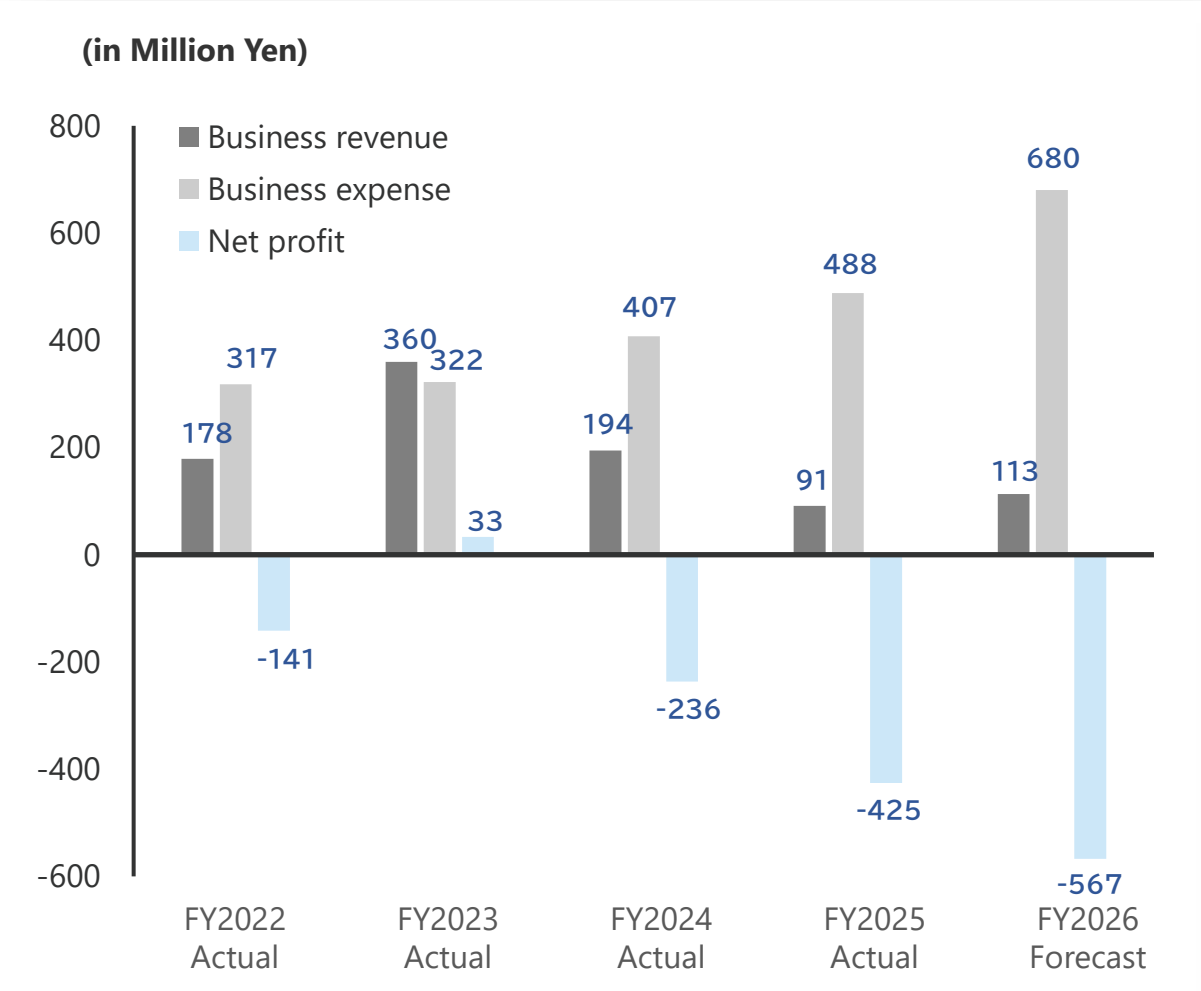
Milestone income: 40
Research support funds & Others: 73

Increase +30
Decrease -8

R&D Expenses: 380
SGA Expenses: 301

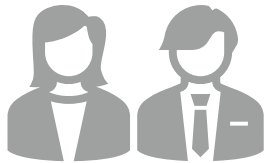
Increase +165
Increase +29

Trends of Business Performance and Forecast of FY2026



Status of Growth Investments Funded by Initial Public Offering

To build the internal foundation for our path toward becoming a specialty pharma, we will advance growth investments in human capital, R&D, facility expansion and renewal, and marketing.



Human Capital

Toward transformation into a specialty pharma

- Scaling up specialized workforce
- Establishing a governance structure
- Improving working condition via base-up

Invested 90
Planned 340 /430million yen

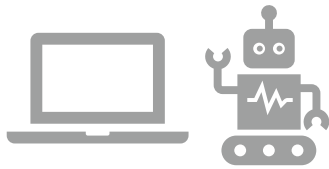


R&D

Creation of in-house pipelines

- In-house pipeline creation research
- Expanding IP
- Commercializing drug delivery system

Invested 40
Planned 350 /390million yen



Facility Expansion and Renewal

Expansion of facilities and computing capacity to strengthen in-house research

- Expending and relocating laboratory
- Expanding and renewing research facilities
- Strengthening computing capacity and AI
- DX of research processes

Invested 30
Planned 10 /40million yen



Marketing

Expanding drug discovery partnerships overseas

- Expand overseas business development activities
- Generate research data directly related to business development
- Presentations at international academic conferences

Invested 10
Planned 50 /60million yen

*As of Dec 31st, 2025 *Planned investments from 2026 onward (Million yen)

Key Initiatives for FY2026

To achieve growth as to be a specialty pharma, the following key initiatives are focused on in 2026:

Measure 1

Secure two new contracts

Secure two new contracts as KPI

Develop business with an eye to European companies and Pesticide business, in adoption to the changes in business environments.

Measure 4

Commercialization of Drug Delivery Systems

Advance commercialization of "Perfusio"

Secure partnerships with medical device manufacturers to accelerate commercialization.

Measure 2

Generate 2nd In-house Pipelines

Progress research to generate 2nd In-house pipelines

Aiming to generate drug candidate substances with high future value and secure patent applications.

Measure 5

Relocate Shin-Kawasaki Institute

Relocate Shin-Kawasaki Institute

To accommodate increasingly sophisticated research and improve working environment, the institute expanded and relocated to a new site in March 2026.

Measure 3

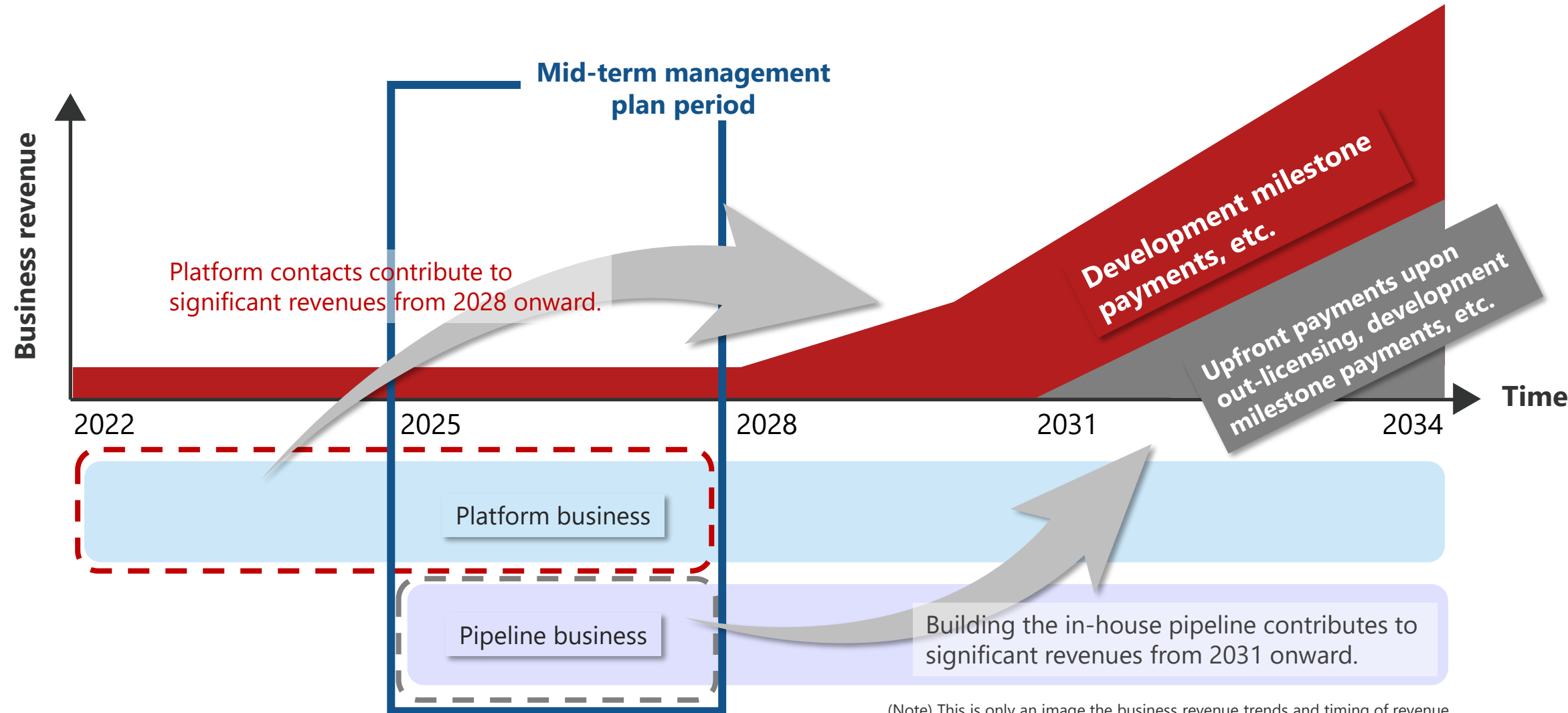
Conduct non-clinical trials of 1st In-house pipelines

Advance nonclinical trials (animal studies) in 1st in-house pipelines

Utilizing "Perfusio" with aim to significantly reduce the costs of non-clinical and clinical trials.

Leveraging Revenue Reinvestment to Maximize Future Value

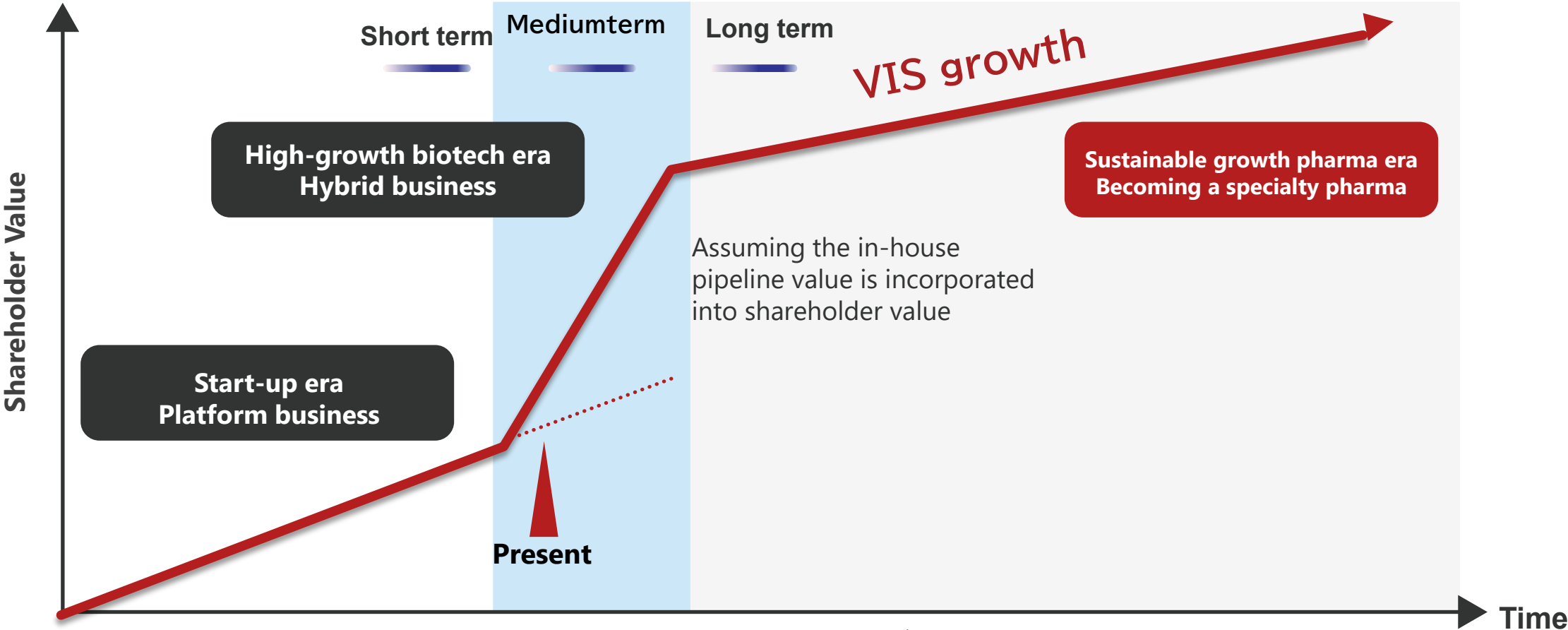
During the mid-term plan, we invest in in-house pipeline development to drive future growth, ensuring past and current investments lead to substantial milestone revenues.



(Note) This is only an image the business revenue trends and timing of revenue achievement, and does not guarantee the actual business revenue trends or timing.

Grow up to be a Specialty Pharma Company

The Company is going to make transition from a high-growth hybrid model to a specialty pharma with R&D and commercial capabilities.



(*This is only an image of the growth curve we aim to achieve and does not indicate the actual trends of shareholder value..



Mission

Build a society where every patient can look forward to a brighter future through the realization of mRNA-targeted drugs

Achieving breakthroughs in drug discovery, We aim to grow into a specialty pharma.

Aim to provide the best treatment for any disease



Veritas In Silico