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### Agenda

- Opening Declaration
- Report on number of voting rights
- Audit Committee Audit Report
- Business Report
- Matters to be Resolved
- 06 Q&A
- ov Vote
- Closing Declaration



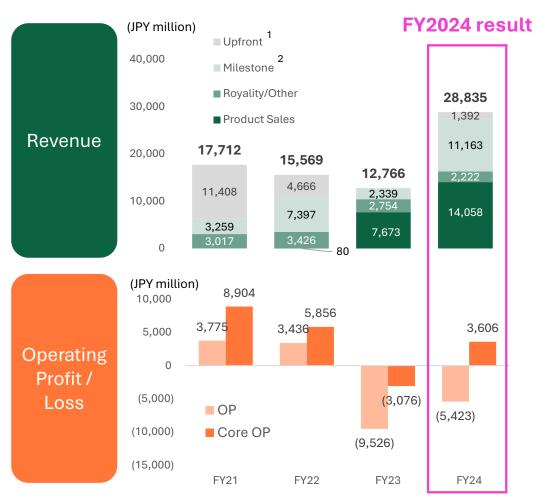
開会宣言 Opening Declaration







# NPJ/NPK product sales and cost base fully included in FY2024. Milestones increased due to progress of partnered programs.



JPY28,835m (+126% | JPY12,766m in FY2023)
Revenue

- PIVLAZ® sales grew significantly from JPY6.1bn to JPY12.7bn due to market penetration and full-year sales contribution (vs. 5 months in FY2023)
- Milestones increased from JPY2.3bn to JPY11.2bn due to the progress of partnered programs

#### **Core Operating Profit of JPY3,606m**

(Core Operating Loss of JPY3,076 in FY2023)

- R&D and SG&A expenses were lower than estimation as the integration progressed well
- Operating Loss of JPY5,423m
   (Operating Loss of JPY9,526m in FY2023)

## Cash/Cash equivalents/Fix deposit: JPY36.2bn (JPY49.0bn in FY2023)

 Purchased QUVIVIQ<sup>™</sup> API (equivalent to 1+ year of supply) to ensure the stable supply of QUVIVIQ<sup>™</sup> in advance



Profit

Cash

<sup>&</sup>lt;sup>1</sup> Upfront fee revenue recognised at deal inception

<sup>&</sup>lt;sup>2</sup> Milestone revenue recognised at milestone event + deferred revenue releases

# Priority objectives for FY2025

- <u>JPY 17 billion+</u> Net product sales (PIVLAZ® plus QUVIVIQ®)
- O2 Acquire/in-license <u>at least one</u> late-stage medicine for Japan/APAC (ex-China)

- Execute <u>at least one</u> new major partnership, and initiate <u>at least one</u> new in-house Ph.2 study
- 104 Investment in systems and applications for efficiency and scalability
- O5 Positive operating profit under IFRS (if GPR52 option is exercised)



#### 1

#### Full year product sales guidance

Target 13.0 - 14.0 Bn JPY (PIVLAZ) as net sales, and 4.0 - 5.0 Bn JPY (QUVIVIQ) as royalty and supply







13.0 - 14.0 Bn JPY

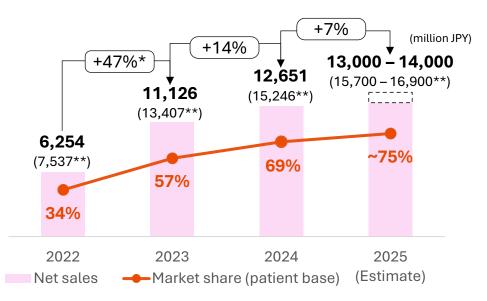
(NHI Sales:15.7 – 16.9Bn JPY)

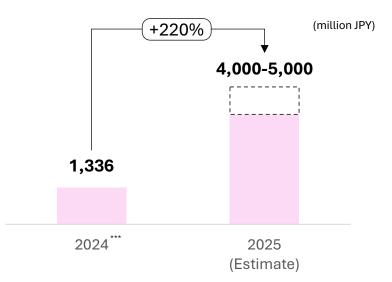












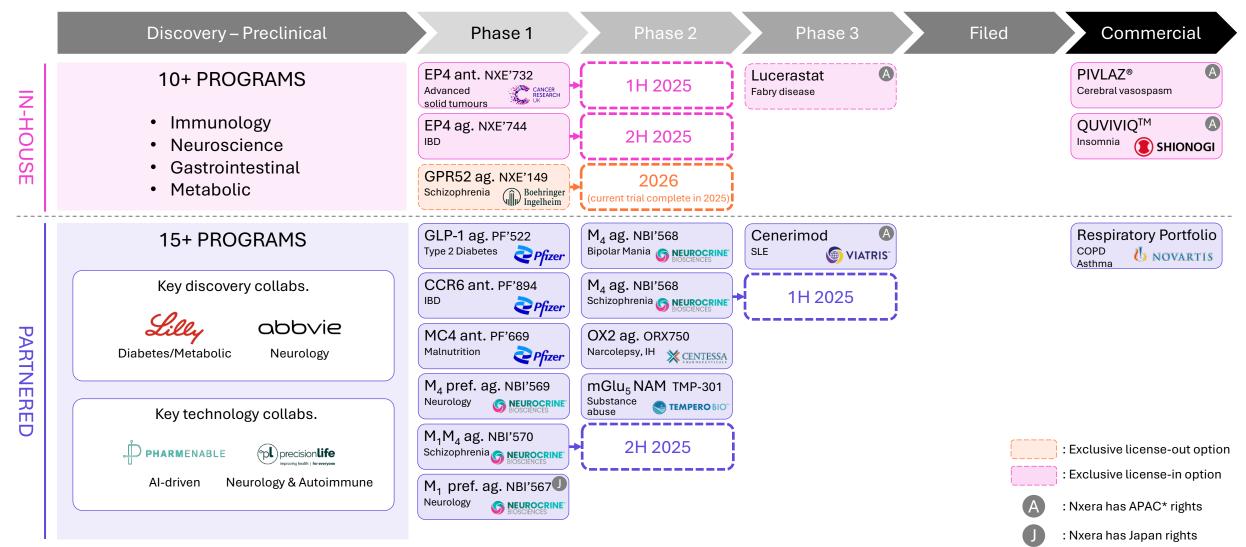
Source: MDV DPC hospital data



<sup>\*:</sup> Comparison of 2-4Q of 2022 and 2023, \*\* NHI sales, \*\*\* 2024 sales includes upfront, milestone, royalty and product supply while 2025 sales includes royalty and product supply

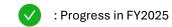
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#### Major Pipeline Overview (with future projection)





## Looking ahead to potential catalysts in 2025\*



	PROGRAM	PARTNER	TIMING	EVENT
<b>&gt;</b>	Cenerimod	100618 SVIATRIS"	Feb. 2025	Assignment of JAPAC rights
<b>V</b>	TMP-301 (mGlu5 NAM)	TEMPERO BIO"	Mar. 2025	Phase 2 study start in alcohol use disorder
	Lucerastat	idorsia	H1 2025	Exclusive opt-in decision
	NXE'732 (EP4 antagonist)	NXEIG CANCER RESEARCH UK	H1 2025	Phase 2a study start in Advancing Solid Tumours
	NBI'568 (M4 agonist)	NEUROCRINE BIOSCIENCES	H1 2025	Phase 3 study start in Schizophrenia
	NBI'568 (M4 agonist)	NEUROCRINE® BIOSCIENCES	H2 2025	Phase 2 study start in Bipolar Mania
	NBI'570 (M1/M4 agonist)	NEUROCRINE® BIOSCIENCES	H2 2025	Phase 2 study start in Schizophrenia
	NXE'744 (EP4 agonist)	NXera ~	H2 2025	Phase 2 study start in IBD
	NXE'149 (GPR52 ag)	NXEIO Boehringer Ingelheim	H2 2025	Phase 1b completion
	NXE'732 (EP4 antagonist)	NXEIO CANCER RESEARCH	H2 2025	Phase 1b topline data
	ORX750 (OX2 agonist)	CENTESSA	H2 2025	Phase 2a data across NT1, NT2, and IH
	Multiple discovery collaboration progress	abbvie <i>Lilly</i>	2025	Progression through discovery stage
	NBI'567 (M1 ago) / NBI'569 (M4 ago) / NBI'570 (M1/M4 ago)	NEUROCRINE® BIOSCIENCES	2025	Phase 1 data readout
<b>V</b>	QUVIVIQ™	Holling Bio-Pharma Corp.	Feb. 2025	Out licensing in Taiwan
	New global out-licenses		Anytime	Out licensing and/or discovery collabs
	New Japan / APAC in-licenses		Anytime	Acquire/in-license late-stage medicines
	QUVIVIQ™		Anytime	APAC out-licensing deals



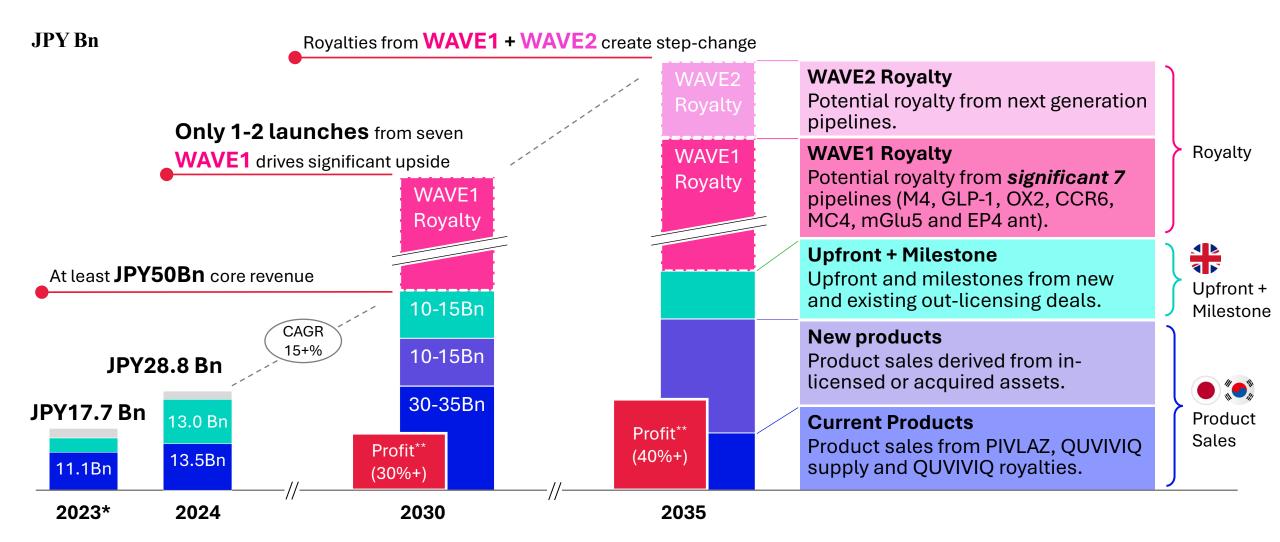
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#### Our Wave 1 and Wave 2 programs are positioned across fast growing areas of healthcare

**MARKET SIZE (2030) WAVE1 (Potential Launch by 2030) WAVE2 (Potential Launch by 2035)** CENTESSA PHARMACEUTICALS CENTESSA MEUROCRINE®
BIOSCIENCES NEUROCRINE® BIOSCIENCES NXeLCI ✓ TEMPERO BIO Neurology mGlu5 NAM Prec Ox2 agonists M4 pref. agonist **GPR52** agonist M4 agonist Ox2 agonist Substance Use Schizophrenia Neuropsych-related Schizophrenia Narcolepsy \$120bn+ M1 pref. agonist Disorders sleep disorders M4 agonist Cognitive & psychosis abbvie Bipolar Mania -related disorders Disc Multiple targets M1/M4 agonist Neurology Schizophrenia **Pfizer** Metabolic **GLP-1** agonist \$150bn+ Multiple targets T2D / Obesity T2D/Obesity and MC4 antagonist Others Malnutrition Immunology **Pfizer** NXeLa ✓ \$300bn+ **CCR6** antagonist EP4 antagonist + PD-L1 EP4 agonist IBD Immune-oncology for IBD Advanced Solid Tumors Multi billion USD milestones and royalties JPY250bn (max total royalty potential at peak)



#### Our 2030 vision is to build a high growth, highly profitable Japanese biopharma



Note: \* Revenue values are proforma the acquisition of Idorsia Pharmaceuticals Japan and Korea and reflect annual product sales of PIVLAZ in 2023.

\*\* WAVE1 and WAVE2 royalty is not included.





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#### Proposal No.1

Reduction of the Amount of Legal Capital Surplus and Disposition of Surplus

#### 1. Content of Reduction of the Amount of Legal Capital Surplus

In accordance with the provisions of Article 448, Paragraph 1 of the Companies Act, the amount of legal capital surplus will be reduced in its entirely and the same amount is transferred to other capital surplus.

(1) Amount of legal capital surplus to be reduced
Amount of legal capital surplus:

JPY35,288,890,082

(2) Effective date of reduction of legal capital surplus April 18, 2025 (planned)

#### 2. Content of Disposition of Surplus

In accordance with the provisions of Article 452 of the Companies Act, subject to the reduction in the amount of legal capital surplus taking effect, losses will be dealt with by transferring JPY 14,620,719,168, which is part of other capital surplus arising from the reduction in the amount of legal capital surplus, to retained earnings brought forward.

(1) Item and amount of surplus to be reduced
Amount of other capital surplus:

JPY14,620,719,168

(2) Item and amount of surplus to be increased
Amount of retained earnings brought forward: JPY14,620,719,168

(3) Effective date of reduction of disposition of surplus April 18, 2025 (planned)



## Proposal No.2

Partial amendment to the Articles of Incorporation

Current Articles of Incorporation	Proposed Amendments
CHAPTER 3 GENERAL MEETING OF SHAREHOLDERS	CHAPTER 3 GENERAL MEETING OF SHAREHOLDERS
(Convocation)	(Convocation)
Article 12	Article 12
The Ordinary General Meeting of Shareholders of the Company shall be convened within three months after the end of each business year and an Extraordinary General Meeting of Shareholders shall be convened whenever necessary.	(Same as current provision)
(Newly established)	2. The General Meeting of Shareholders of the Company may be held as shareholders' meetings without a location.



Proposal No.3
Election of Eight (8) Directors

1	Christopher Cargill	Director, Representative Executive Officer, President & CEO Member of the Compensation Committee	RE-APPOINTED		
2	David Roblin	External Director, Chair of the Compensation Committee, Member of the Nomination Committee	RE-APPOINTED	EXTERNAL	INDEPENDENT
3	Noriaki Nagai	External Director, Member of the Nomination Committee, Member of the Audit Committee	RE-APPOINTED	EXTERNAL	INDEPENDENT
4	Rolf Soderstrom	External Director, Chair of the Audit Committee, Member of the Compensation Committee	RE-APPOINTED	EXTERNAL	INDEPENDENT
5	Miwa Seki	External Director, Chair of the Nomination Committee, Member of the Audit Committee	RE-APPOINTED	EXTERNAL	INDEPENDENT
6	Eiko Tomita	External Director Member of the Audit Committee	RE-APPOINTED	EXTERNAL	INDEPENDENT
7	Naoko Shimura	_	NEWLY APPOINTED	EXTERNAL	
8	Nicola Rabson	-	NEWLY APPOINTED	EXTERNAL	INDEPENDENT

Note: Current positions and responsibilities at the Company are presented



# 質疑応答 Q&A









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