INDIVIOR®

Indivior, Powering Recovery, Renewing Hope.

Investor Presentation

January 8, 2026

IMPORTANT CAUTIONARY STATEMENT REGARDING FORWARD-LOOKING STATEMENTS

This presentation contains certain statements that are forward-looking. Forward-looking statements include, among other things, express and implied statements regarding: the Company's financial guidance for both 2025 and 2026, including total net revenue, SUBLOCADE® net revenue, Non-GAAP gross margin, Non-GAAP operating expenses, Non-GAAP SG&A, R&D expenses, and Adjusted EBITDA; expected future operating expense savings; expected future increases in dispensed units and cash flow; our expectation that we can grow and accelerate SUBLOCADE net revenue, generate immediate accretion from profitability and cash flow growth exceeding revenue growth, and leverage strengthened financial profile to acquire next growth drivers; potential future patents that might be awarded; expectations of increased LAI usage; our product development pipeline and potential future products, the timing of clinical trials, expectations regarding regulatory approval of such product candidates, the timing of such approvals, and the timing of commercial launch of such products or product candidates, and eventual annual revenues of such future products; and other statements containing the words "believe," "anticipate," "plan," "expect," "intend," "estimate," "forecast," "strategy," "target," "guidance," "outlook," "potential," "project," "priority," "may," "will," "should," "would," "could," "can," the negatives thereof, and variations thereon and similar expressions.

By their nature, forward-looking statements involve risks and uncertainties as they relate to events or circumstances that may or may not occur in the future. Actual results may differ materially from those expressed or implied in these forward-looking statements due to a number of factors, including: lower than expected future sales of our products; greater than expected impacts from competition; unanticipated costs including the effects of potential tariffs and potential retaliatory tariffs; whether we are able to identify efficiencies and fund additional investments that we expect to generate increased revenue, and the timing of such actions; market acceptance of long-acting injectables; and the results of pending and future clinical trials, and the decisions of relevant regulators. For additional information about some of the risks and important factors that could affect our future results and financial condition, see "Risk Factors" in our Annual Report on Form 10-K filed March 3, 2025, in our Quarterly Reports on Forms 10-Q filed May 1, 2025, July 31, 2025, and October 30, 2025, our other filings with the U.S. Securities and Exchange Commission.

Forward-looking statements speak only as of the date that they are made and should be regarded solely as our current plans, estimates and beliefs. Except as required by law, we do not undertake and specifically decline any obligation to update, republish or revise forward-looking statements to reflect future events or circumstances or to reflect the occurrences of unanticipated events.

LONGSTANDING LEADERSHIP IN THE TREATMENT OF OPIOID USE DISORDER

20+

Years of leadership in OUD treatment

Long history of helping people **achieve long-term recovery from opioid use disorder (OUD)** through accessible, science-driven care

465K+

Patients treated

SUBLOCADE® is a durable growth driver and is the **#1 prescribed, first-in-class,** monthly subcutaneous long-acting injectable (LAI) medication for the treatment of moderate to severe OUD

$1.2B

Revenue expected in 2025[1]

Strong financial position and poised to **accelerate SUBLOCADE** and grow adjusted EBITDA and cash flow at a faster rate

1. Based on financial guidance ranges provided by Indivior in its press release on Form 8-K filed with the SEC on October 30, 2025.

EXECUTING THE INDIVIOR ACTION AGENDA AND ENTERING 2026 AS A FOCUSED, SIMPLIFIED AND STRONGER INDIVIOR

Sharpened focus on highest growth opportunity – U.S. SUBLOCADE

New operating model in place to drive significant bottom-line growth and cash flow generation

Improved financial profile and strength enables capital allocation optionality

THE INDIVIOR ACTION AGENDA

Phase III – Breakout (H2'26 – Beyond)

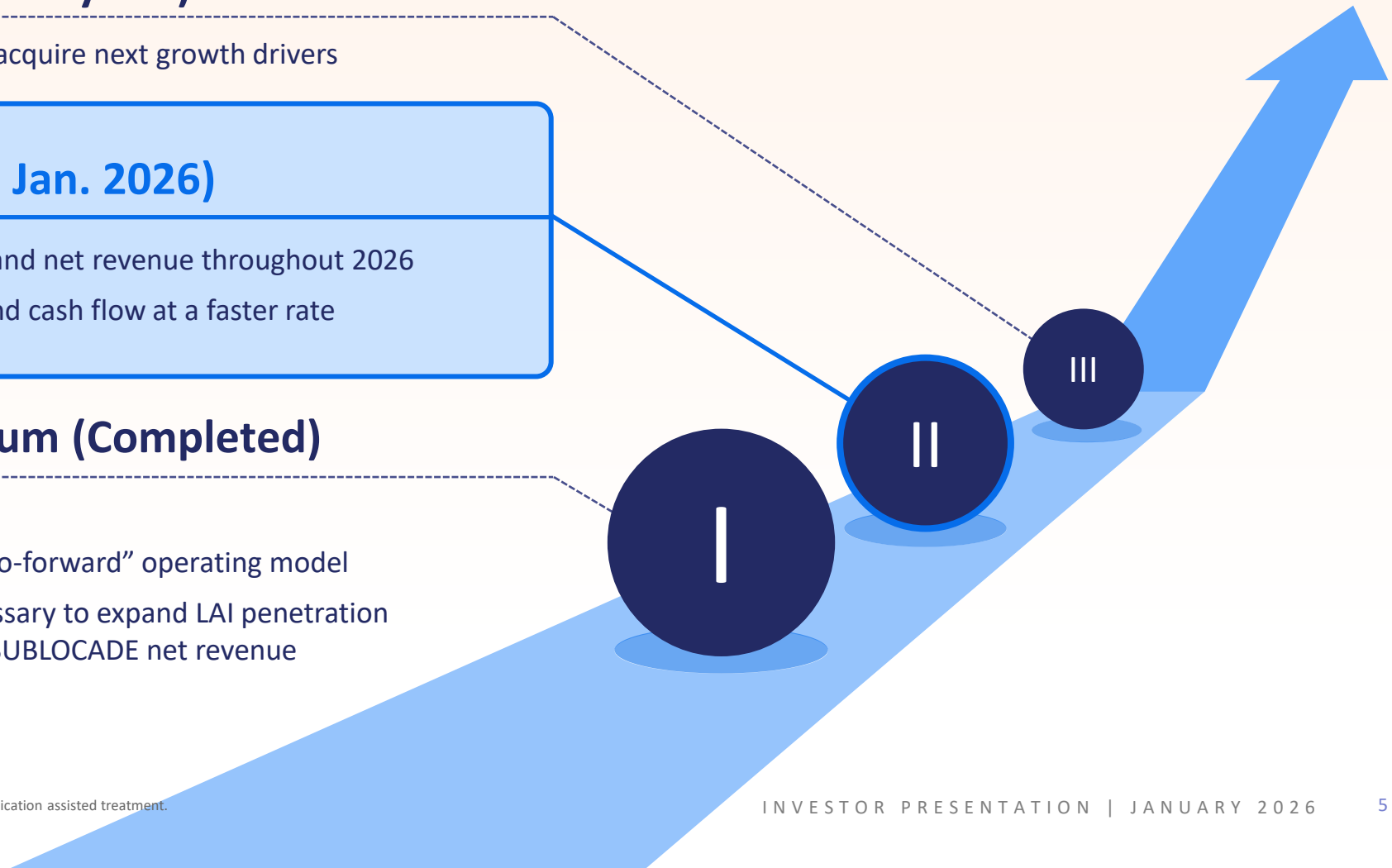
- Leverage strengthened financial profile to acquire next growth drivers

Phase II – Accelerate (Began Jan. 2026)

- Accelerate U.S. SUBLOCADE dispense unit and net revenue throughout 2026
- Immediately accelerate adjusted EBITDA and cash flow at a faster rate

Phase I – Generate Momentum (Completed)

- ✓ Grow U.S. SUBLOCADE net revenue
- ✓ Simplify the organization and establish "go-forward" operating model
- ✓ Determine actions and investments necessary to expand LAI penetration in U.S. BMAT category to accelerate U.S. SUBLOCADE net revenue

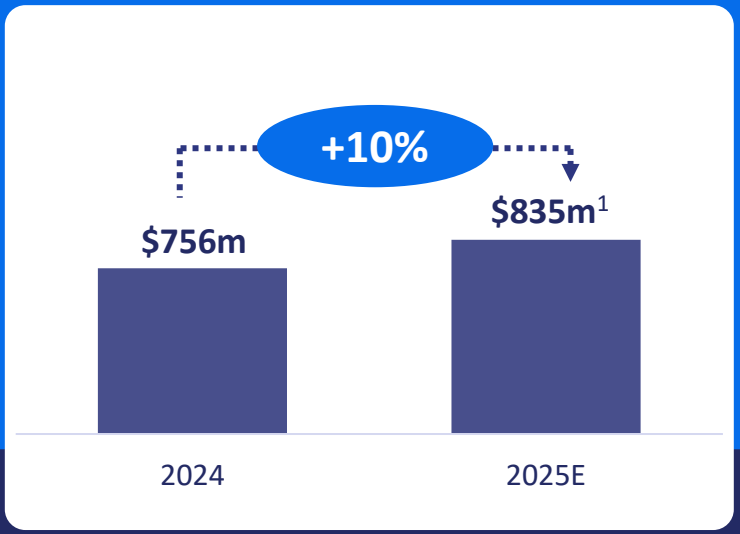


LAI: long-acting injectable. BMAT: buprenorphine medication assisted treatment.

COMPLETED PHASE I – GENERATE MOMENTUM

1

Grew SUBLOCADE in the U.S.



SUBLOCADE Net Revenue Growth

2

Simplified the organization and established "go-forward" operating model

- Completed LSE delisting
- Consolidated operating footprint
- Restructured R&D and Medical Affairs organizations
- Discontinued sales and marketing support of OPVEE®
- Optimized the Rest of World business
- Eliminated legacy DOJ obligation
- Received shareholder approval of U.S. redomicile

At least $150m in annual expense savings expected in 2026

3

Determined actions and investments necessary to expand LAI penetration in U.S. BMAT category to accelerate U.S. SUBLOCADE net revenue

Launched new DTC campaign in October 2025

Omnichannel patient activation initiative

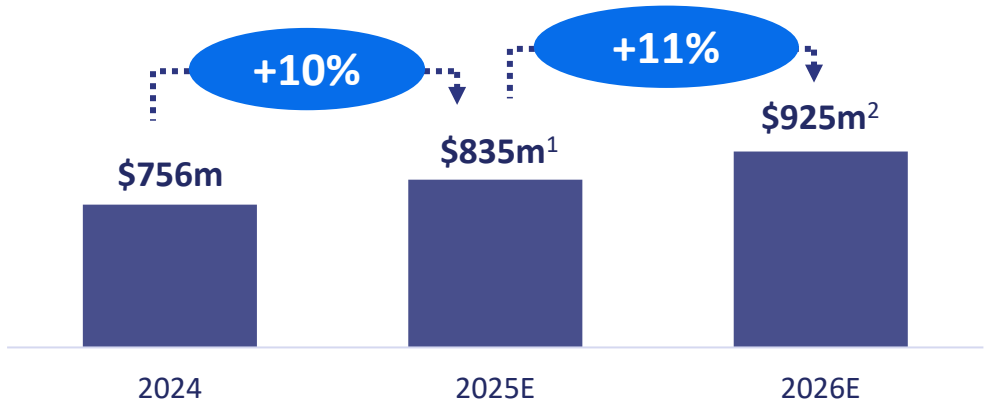
1. Based on financial guidance ranges provided by Indivior in its press release on Form 8-K filed with the SEC on October 30, 2025. LSE: London Stock Exchange. DOJ: Department of Justice. DTC: direct-to-consumer.

ENTERED PHASE II – ACCELERATE – ON JANUARY 1, 2026

1

Accelerate U.S. SUBLOCADE

Total SUBLOCADE Net Revenue

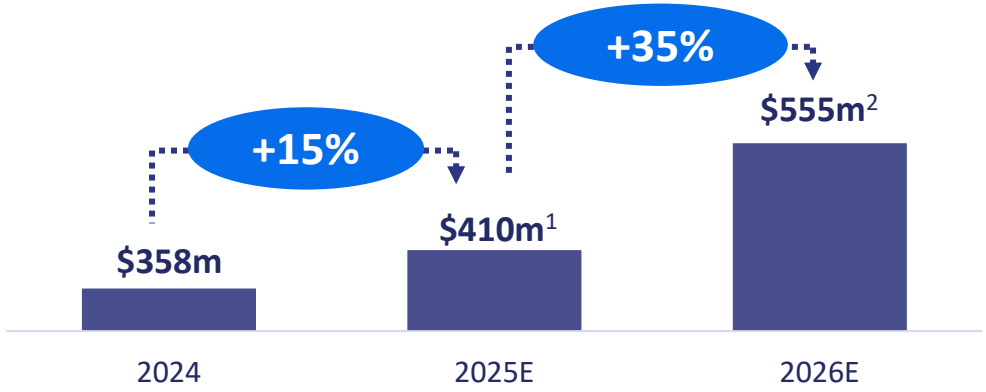


Expect to accelerate SUBLOCADE dispense unit growth from **~7%** in 2025 to the **mid-teens** in 2026

2

Immediately accelerate adjusted EBITDA and cash generation at a faster rate than revenue

Adjusted EBITDA[3]



~$300m in cash flow from operations expected in 2026[4]

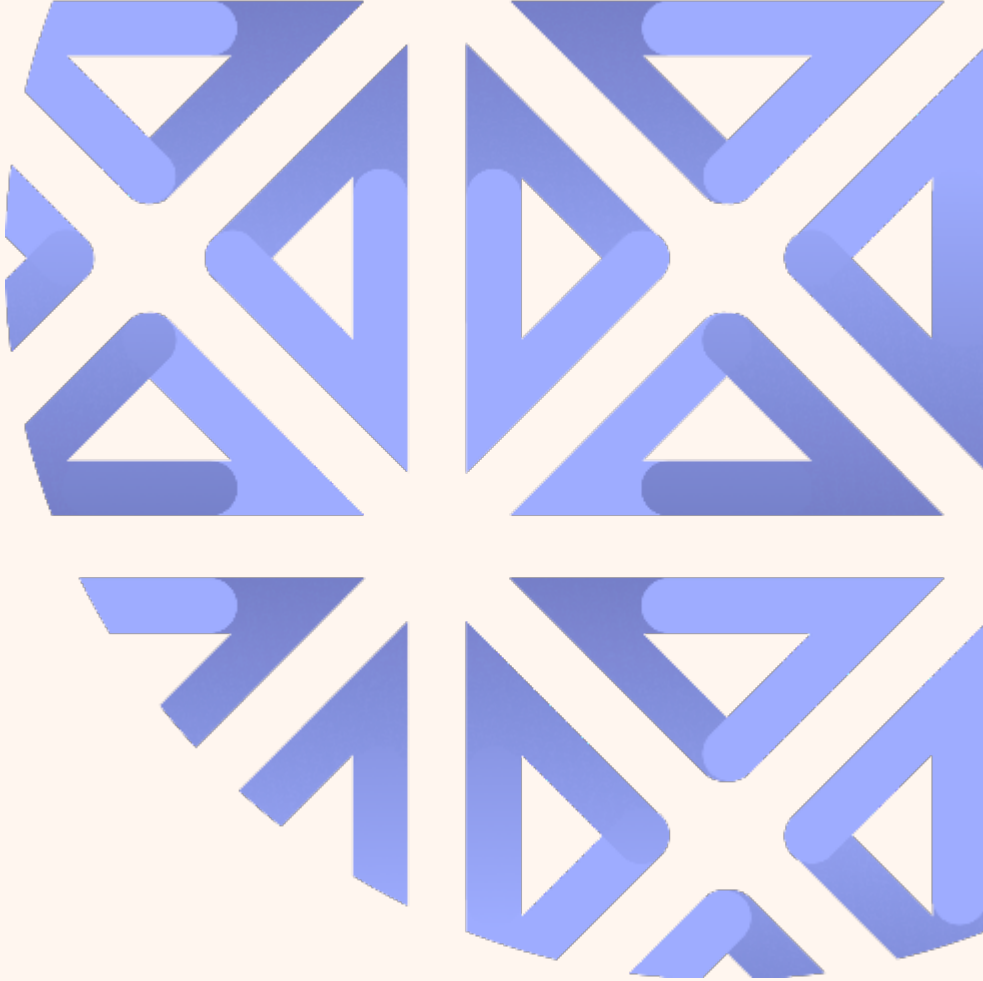
1. Based on financial guidance ranges provided by Indivior in its press release on Form 8-K filed with the SEC on October 30, 2025. 2. Based on financial guidance ranges provided by Indivior in its press release on Form 8-K filed with the SEC on January 8, 2026. 3. Adjusted EBITDA is a non-GAAP financial measure. See Non-GAAP Financial Measures in the Appendix for reconciliation. For non-GAAP guidance items, the Company has relied upon the exception in item 10(e)(1)(i)(B) of Regulation S-K to exclude such reconciliations, as the reconciliations of these non-GAAP guidance metrics to their corresponding GAAP equivalents are not available without unreasonable effort; See slides 8 for details. 4. Excludes cash flows from investing and financing activities.

2026 FINANCIAL GUIDANCE

	Guidance Range[1]	YoY Change[2]	Commentary
Total Net Revenue	**$1,125m - $1,195m**	**-3%**	U.S. SUBOXONE Film pressure; ROW optimization; PERSERIS® run-off; cessation of OPVEE® promotion
SUBLOCADE Net Revenue	**$905m - $945m**	**+11%**	Acceleration of U.S. SUBLOCADE dispense unit growth to **mid-teens**
Non-GAAP Operating Expenses[3]	**$430m - $450m**	**-26%**	At least **$150m** in operating expense savings
Adjusted EBITDA[3]	**$535m - $575m**	**+35%**	Margin expansion of **14 percentage pts. to 48%**

1. Based on financial data provided by Indivior in its press release on Form 8-K filed with the SEC on January 8, 2026. 2. Represents the midpoint of 2026 guidance ranges compared to the midpoint of 2025 guidance ranges provided by Indivior in its press release on Form 8-K filed with the SEC on October 30, 2025. 3. For non-GAAP guidance items, the Company has relied upon the exception in item 10(e)(1)(i)(B) of Regulation S-K to exclude such reconciliations, as the reconciliations of these non-GAAP guidance metrics to their corresponding GAAP equivalents are not available without unreasonable effort; See slides 24 to 26 for details.

SUBLOCADE®



SIGNIFICANT OPPORTUNITY TO INCREASE USE OF LAI BUPRENORPHINE MEDICATIONS IN THE TREATMENT OF OUD

8.9m[1]

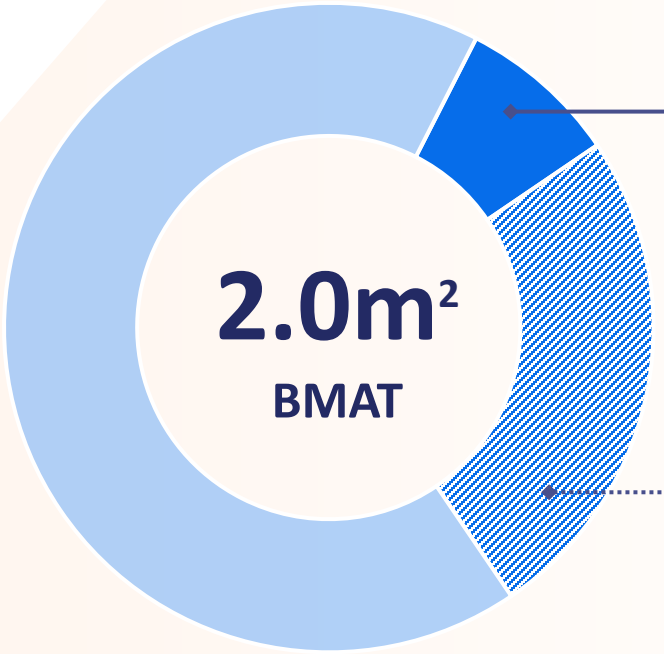
Misuse opioids in U.S.
(Total Addressable Market)

5.9m[1]

OUD diagnosed in U.S.
(Service Addressable Market)

2.0m[2]

Received Buprenorphine Medication Assisted Treatment (BMAT)



~8%

Current LAI usage in BMAT population allows for significant potential expansion

20-30%

HCPs expect LAI usage to increase to 20-30% of the BMAT population[3]

1. 2023 NSDUH Annual National Report (SAMSHA). 2. Symphony and Indivior analytics, patient treated over the last twelve months. 3. HCP Survey conducted Q3 2024. N=400 HCP and patients combined in qual and quant.

SUBLOCADE: A DURABLE GROWTH ASSET WITH IP PROTECTION TO 2031-2038

ONCE-MONTHLY

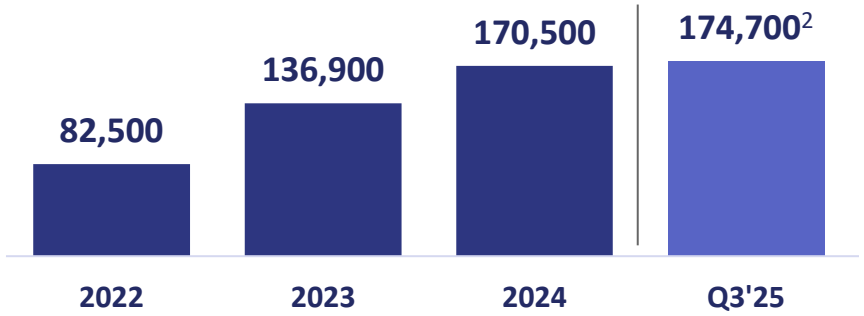
Sublocade®

(buprenorphine extended-release) injection for subcutaneous use Ⓒ

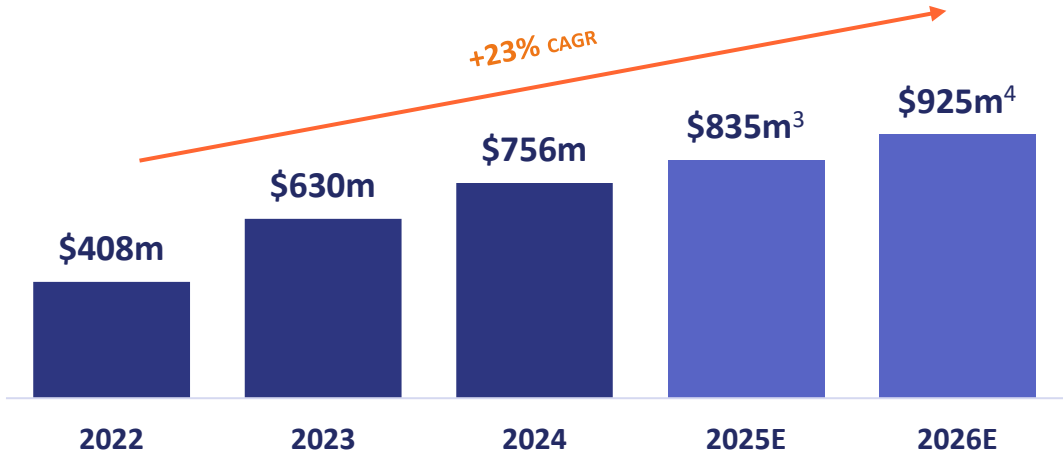
100mg·300mg

- **#1 prescribed LAI** in the U.S.
- **Over 465K** lives treated
- The **only once-monthly LAI with rapid initiation** on day 1
- **Significant IP** with 12 orange-book listed patents to 2031-2038[1]; pursuing 6 additional U.S. patent applications with potential expirations from 2035-2044

TTM SUBLOCADE PATIENTS



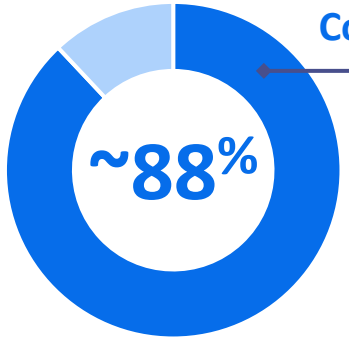
SUBLOCADE NET REVENUE



1. Patent expiring in 2038 is for 300mg/1.5ML dose - 11 other orange book patents expire 2031-2035. 2. Represents trailing twelve months of estimated patients in treatment. 3. Based on financial guidance ranges provided by Indivior in its press release on Form 8-K filed with the SEC on October 30, 2025. 4. Based on financial guidance ranges provided by Indivior in its press release on Form 8-K filed with the SEC on January 8, 2026.

STRONG FUNDAMENTALS POSITION SUBLOCADE FOR GROWTH

BROAD PAYOR ACCESS FOR SUBLOCADE



Coverage in Medicaid and Commercial

- Simple single prior authorization (PA)
- PA is label aligned
- 95% of people in the SUBLOCADE copay program pay $0

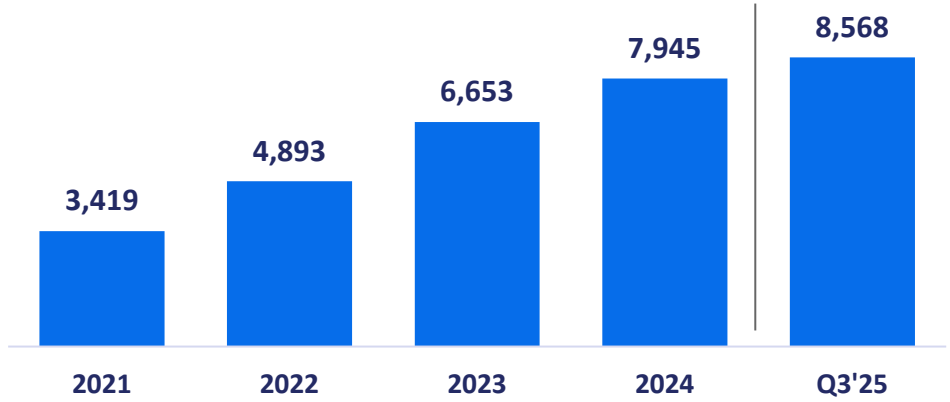
HIGH INTENT TO PRESCRIBE[1]

74% of HCPs consider SUBLOCADE to be appropriate for patients with severe OUD

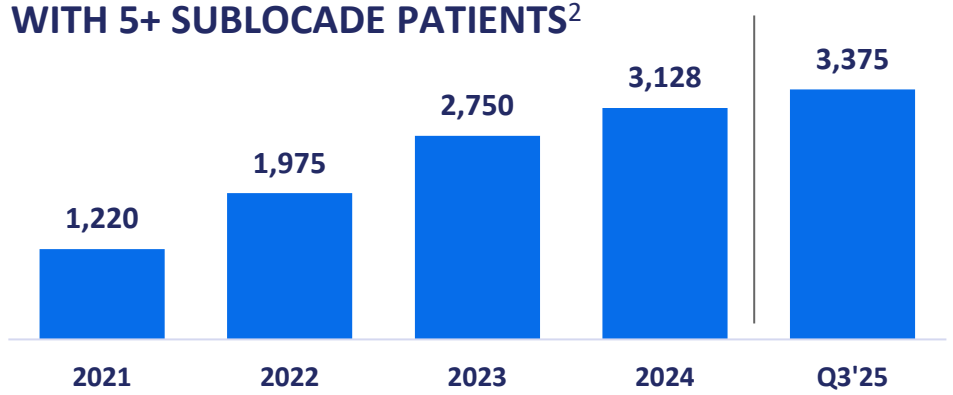
83% of HCPs consider SUBLOCADE to be appropriate for patients burdened by daily drug-taking

HCPs prescribing SUBLOCADE report that they will prescribe to **30%** more patients over the next 18 months

GROWING SUBLOCADE PRESCRIBER BASE[2]



PRESCRIBING DEPTH IMPROVING: HCPS WITH 5+ SUBLOCADE PATIENTS[2]



1. Indivior Internal Marketing – Q3 2025 Healthcare Practitioner ATU study. 2. Active count of prescribing HCPs; excludes delisted and Specialty HCPs. Represents trailing twelve months.

INITIATIVES TO ACCELERATE SUBLOCADE GROWTH

Improving Commercial Execution

- **Strengthen** field force messaging and productivity
- **Accelerate** growth with commercial patients
- **Drive** awareness of updated label and unique rapid initiation

Expanding Patient Awareness and Engagement

- **Increase** patient awareness of SUBLOCADE and LAI category
- **Launched** DTC Campaign ("Move Forward in Recovery") in October 2025

Unlocking Access Through Policy Leadership

- **Advance** state and federal policies that support durable access to increase long-term adoption of LAIs
- **Activate** advocates to accelerate access, reduce system barriers and increase awareness

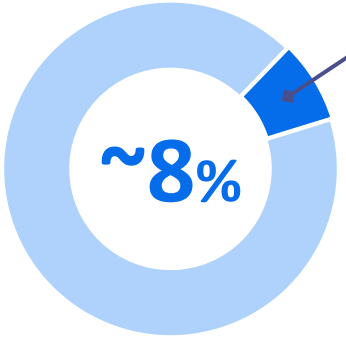
Committed to investing at sustained levels to expand LAI penetration in U.S. BMAT category to accelerate U.S. SUBLOCADE net revenue

SUBCLOCADE ON TRACK TO ACCELERATE IN 2026 WITH CONSUMER ACTIVATION EFFORTS

ACCELERATION IN NEW PATIENT STARTS

+25% Growth in new patient starts from November 2024 to November 2025

ADOPTION OF NEW PATIENTS RECEIVING ACCELERATED SECOND DOSE



Percent of new patients receiving accelerated dose **more than doubled** from August 2025 to October 2025

19% increase in new patients starting on accelerated dose in October 2025 vs. September 2025

POSITIVE EARLY INDICATORS OF DIRECT-TO-CONSUMER ACTIVATION

2x Increase in average daily search volume in October and November 2025 compared to January – September 2025

+80% Growth in FASTP Physician Locator usage vs. pre-National TV Launch

All Time High CRM engagement by patients November QTD

INDIVIOR

Source: Indivior internal analytics.
FASTP: Find a SUBLOCADE Treatment Provider. CRM: Customer Relationship Management.

Pipeline

OUD FOCUSED PIPELINE

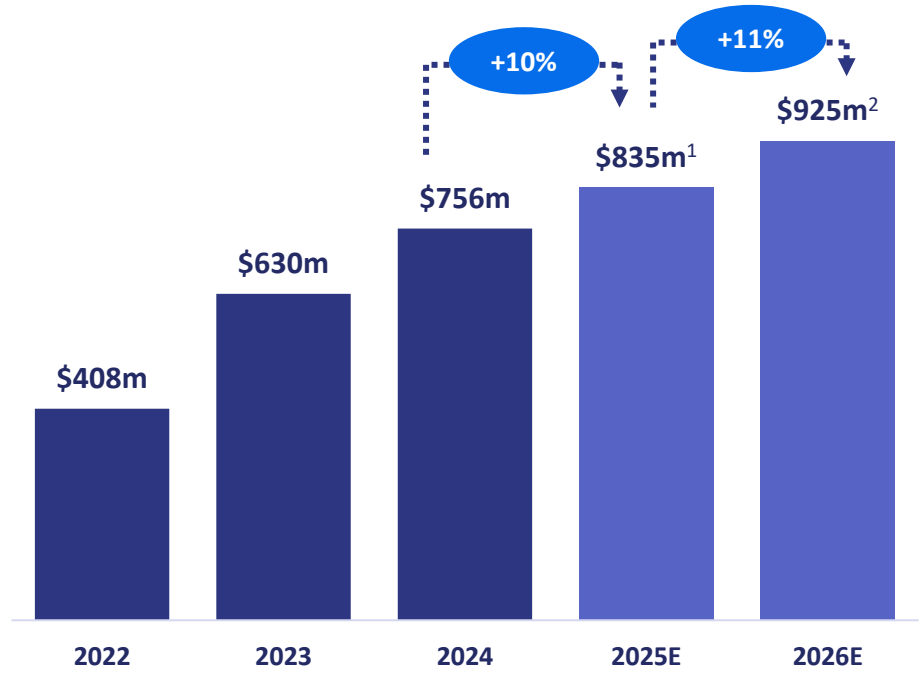
Trial	Patients & Population	Design	Primary Endpoints	Completion	Patent Protection
INDV-6001 3-month long-acting buprenorphine Phase II **NCT06576843**	**122 Patients** Moderate to severe OUD	Multiple dose Phase II PK study	Evaluate PK, safety and tolerability of INDV-6001 following multiple doses in participants with OUD	Last Patient Last Visit **Q4 2025**	2037-2043
INDV-2000 Selective Orexin-1 receptor antagonist (oral tablet) Phase II **NCT06384157**	**300 Patients** Moderate to severe OUD	Placebo or 3 dosing regimes of INDV-2000	Efficacy – Proportion (probability) of patients without treatment failure[1] by the end of week 12	Last Patient Last Visit **Q4 2025**	2035-2037

1. Treatment failure defined as either one of two criteria: (1) Urine Drug Screen positive for opioids, or fentanyl on 4 consecutive assessments while participants are on INDV-2000 or placebo alone, (2) Discontinued INDV-2000 or placebo prematurely.

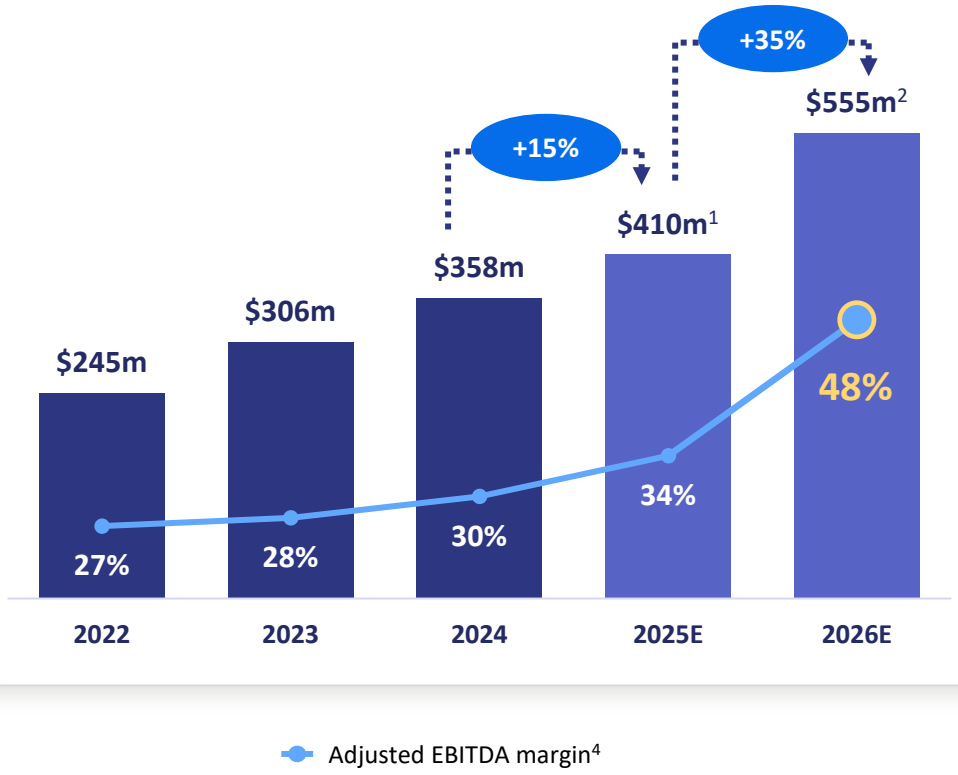
Financials

EXECUTION AGAINST THE INDIVIOR ACTION AGENDA DRIVES STRONG FINANCIAL PERFORMANCE

GROWING SUBLOCADE NET REVENUE



EXPANDING ADJUSTED EBITDA[3]



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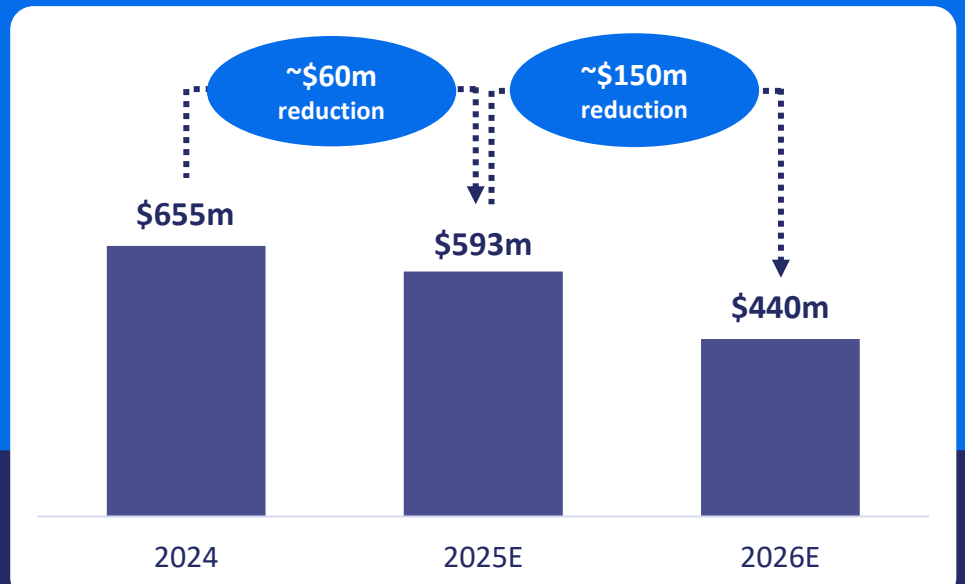
BOTTOM-LINE EXPANSION DRIVEN BY SIMPLIFIED OPERATING MODEL

Simplification Actions to Generate Savings

Completed LSE delisting	**Consolidated** operating footprint
Restructured R&D and Medical Affairs organizations	**Discontinued** sales and marketing support of OPVEE
Optimized The Rest of World business	**Received** Shareholder approval of U.S. redomicile

Expect at Least $150m in Non-GAAP Operating Expense Savings in 2026[1,2]



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SIGNIFICANT CASH FLOWS AND STRONG BALANCE SHEET ENABLE CAPITAL ALLOCATION OPTIONALITY

$473m in cash and investments as of 9/30/25

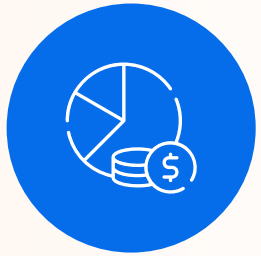
~$300m in cash flow from operations expected in 2026[1]

$295m Payment to DOJ on 11/20/25 eliminated legacy matter

0.8x leverage ratio[2]

DEBT MANAGEMENT

$350m term loan maturing in 2030 with **$50m** revolving credit facility



SHARE REPURCHASES

~$400m of share repurchases conducted since 2021 at average weighted price of **$14.60**

BUSINESS DEVELOPMENT

Earning our way to Phase III of Indivior Action Agenda – Breakout – to **acquire next commercial stage growth drivers**

1. Excludes cash flows from investing and financing activities. 2. Defined as Net Debt as of September 30, 2025, of $287m, divided by Adjusted EBITDA for September 2025 trailing 12 months of $361m; See Non-GAAP Financial Measures in the Appendix for reconciliation on slide 26.

Summary

DELIVERING ON STRATEGIC PRIORITIES TO ACCELERATE IN 2026

Make a **positive difference** in the lives of people living with OUD

Maximize the potential of the business

Create **long-term value** for shareholders

Appendix

FY 2024 NON-GAAP OPERATING EXPENSE RECONCILIATION

($ in mil.)	2024
Total Operating Expenses, net	**919**
Other operating expense (income), net	(4)
Acquired In-process R&D	(1)
Non-GAAP adjustments	(235)
Share based compesation	(24)
Non-GAAP operating expenses	**655**

Columns may not foot due to rounding.

FY 2022–2024 ADJUSTED EBITDA RECONCILIATIONS

($ in mil.)	2024	2023	2022
Net Income	**7**	**(126)**	**(42)**
Add Back:			
Interest Income	(23)	(43)	(19)
Interest Expense	41	35	27
Income Tax Expense / (Benefit)	13	(19)	(43)
Non-GAAP adjustments in Operations	280	265	297
Dep/Amort (excluding ROU Amort)	16	11	9
Share-Based Compensation Expense	24	21	16
Opiant Transaction		162	
Total Adjustments	**351**	**432**	**287**
Adjusted EBITDA	**358**	**306**	**245**
Net Revenue	1,188	1,093	901
Adjusted EBITDA Margin	30%	28%	27%

Columns may not foot due to rounding.

Q3 2025 TTM LEVERAGE RECONCILIATION

($ in mil.)	Q4 2024	Q1 2025	Q2 2025	Q3 2025
Net Debt[1]				**287**
Net income (loss)	**21**	**47**	**18**	**42**
Adjustments:				
Interest income	(5)	(4)	(6)	(6)
Interest expense	13	12	15	12
Income tax expense (benefit)	17	11	44	(5)
Depreciation/amortization (excluding ROU amortization)	6	3	3	2
Non-GAAP adjustments in operating income	17	3	6	67
Share-based compensation expense	6	6	8	6
Total Adjustments	**54**	**31**	**70**	**76**
Adjusted EBITDA	**75**	**78**	**88**	**120**
Adjusted Leverage				**0.8**

1. Net Debt represents $337m of the outstanding balance of the note purchase agreement less $50m of cash. Columns may not foot due to rounding.

SUBLOCADE® (buprenorphine extended-release) injection, for subcutaneous use (CIII)

INDICATION

SUBLOCADE is indicated for the treatment of moderate to severe opioid use disorder in patients who have initiated treatment with a single dose of a transmucosal buprenorphine product or who are already being treated with buprenorphine.

SUBLOCADE should be used as part of a complete treatment plan that includes counseling and psychosocial support.

HIGHLIGHTED SAFETY INFORMATION

WARNING: RISK OF SERIOUS HARM OR DEATH WITH INTRAVENOUS ADMINISTRATION; SUBLOCADE RISK EVALUATION AND MITIGATION STRATEGY
See full prescribing information for complete boxed warning.

- **Serious harm or death could result if administered intravenously.**
- **SUBLOCADE is only available through a restricted program called the SUBLOCADE REMS Program. Healthcare settings and pharmacies that order and dispense SUBLOCADE must be certified in this program and comply with the REMS requirements.**

CONTRAINDICATIONS

Hypersensitivity to buprenorphine or any other ingredients in SUBLOCADE.

WARNINGS AND PRECAUTIONS

Addiction, Abuse, and Misuse: SUBLOCADE contains buprenorphine, a Schedule III controlled substance that can be abused in a manner similar to other opioids. Monitor patients for conditions indicative of diversion or progression of opioid dependence and addictive behaviors.

Respiratory Depression: Life threatening respiratory depression and death have occurred in association with buprenorphine. Warn patients of the potential danger of self-administration of benzodiazepines or other CNS depressants while under treatment with SUBLOCADE.

Risk of Serious Injection Site Reactions: Likelihood of may increase with inadvertent intramuscular or intradermal administration. Evaluate and treat as appropriate. The most common injection site reactions are pain, erythema and pruritus with some involving abscess, ulceration and necrosis.

Neonatal Opioid Withdrawal Syndrome: Neonatal opioid withdrawal syndrome (NOWS) is an expected and treatable outcome of prolonged use of opioids during pregnancy.

Adrenal Insufficiency: If diagnosed, treat with physiologic replacement of corticosteroids, and wean patient off the opioid.

Risk of Opioid Withdrawal With Abrupt Discontinuation: If treatment with SUBLOCADE is discontinued, monitor patients for several months for withdrawal and treat appropriately.

Risk of Hepatitis, Hepatic Events: Monitor liver function tests prior to and during treatment.

Risk of Withdrawal in Patients Dependent on Full Agonist Opioids: Verify that patients have tolerated transmucosal buprenorphine before injecting SUBLOCADE.

Treatment of Emergent Acute Pain: Treat pain with a non-opioid analgesic whenever possible. If opioid therapy is required, monitor patients closely because higher doses may be required for analgesic effect.

ADVERSE REACTIONS

Adverse reactions commonly associated with SUBLOCADE (in ≥5% of subjects) were constipation, headache, nausea, injection site pruritus, vomiting, increased hepatic enzymes, fatigue, and injection site pain.

For more information about SUBLOCADE, the full Prescribing Information including BOXED WARNING, and Medication Guide, visit www.sublocade.com.