

medincell.

Shift to Growth

September 2026

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This presentation and the Fiscal Year 2024-25 results conference (French and English sessions) contain forward-looking statements, including statements regarding Company's expectations for (i) the timing, progress and outcome of its clinical trials; (ii) the clinical benefits and competitive positioning of its product candidates; (iii) the ability of its products to obtain regulatory approvals, commence commercial production and achieve market penetration and sales; (iv) its future product portfolio; (v) its future partnering arrangements; (vi) its future capital needs, capital expenditure plans and ability to obtain funding; and (vii) prospective financial matters regarding our business. Although the Company believes that its expectations are based on reasonable assumptions, any statements other than statements of historical facts that may be contained in this This presentation and the Fiscal Year 2024-25 results conference (French and English sessions) relating to future events are forward-looking statements and subject to change without notice, factors beyond the Company's control and the Company's financial capabilities.

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Any forward-looking statements made by or on behalf of the Company speak only as of the date they are made. Except as required by law, the Company does not undertake any obligation to publicly update these forward-looking statements or to update the reasons why actual results could differ materially from those anticipated by the forward-looking statements, including in the event that new information becomes available. The Company's update of one or more forward-looking statements does not imply that the Company will make any further updates to such forward-looking statements or other forward-looking statements. Readers are cautioned not to place undue reliance on these forward-looking statements.

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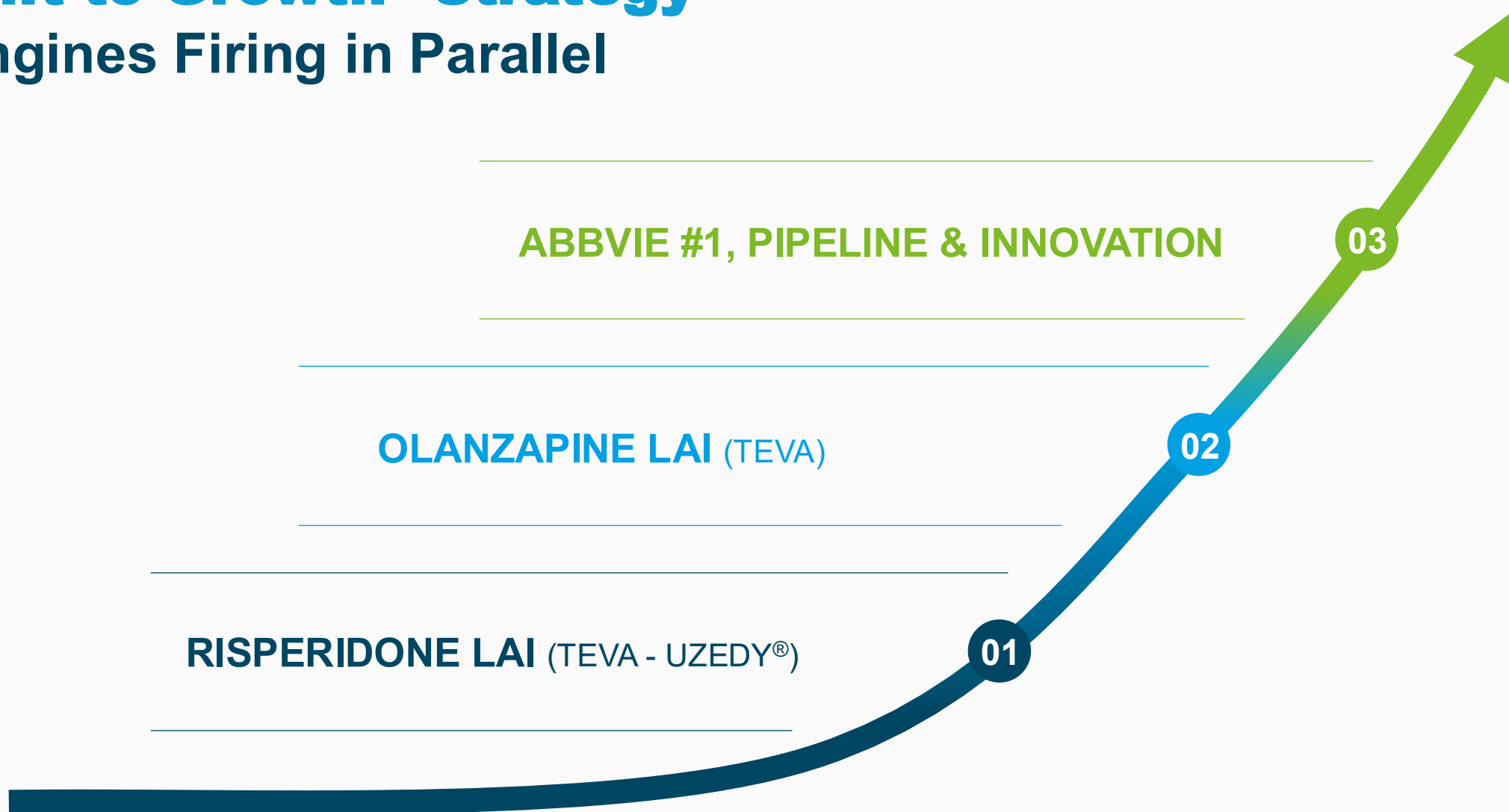
UZEDY® is a trademark of Teva Pharmaceuticals.

**The next two years will be the most
transformative of Medincell's history**

Medincell - June 2025

“Shift to Growth” Strategy

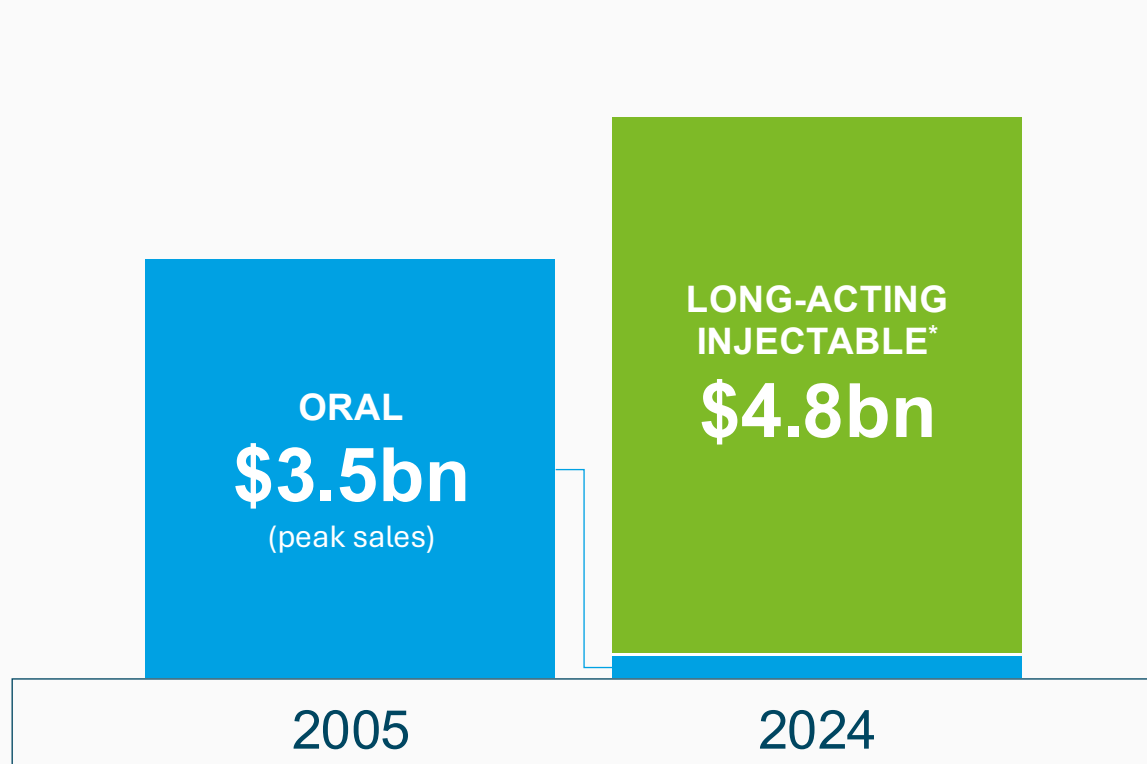
3 Engines Firing in Parallel



Schizophrenia Strategy

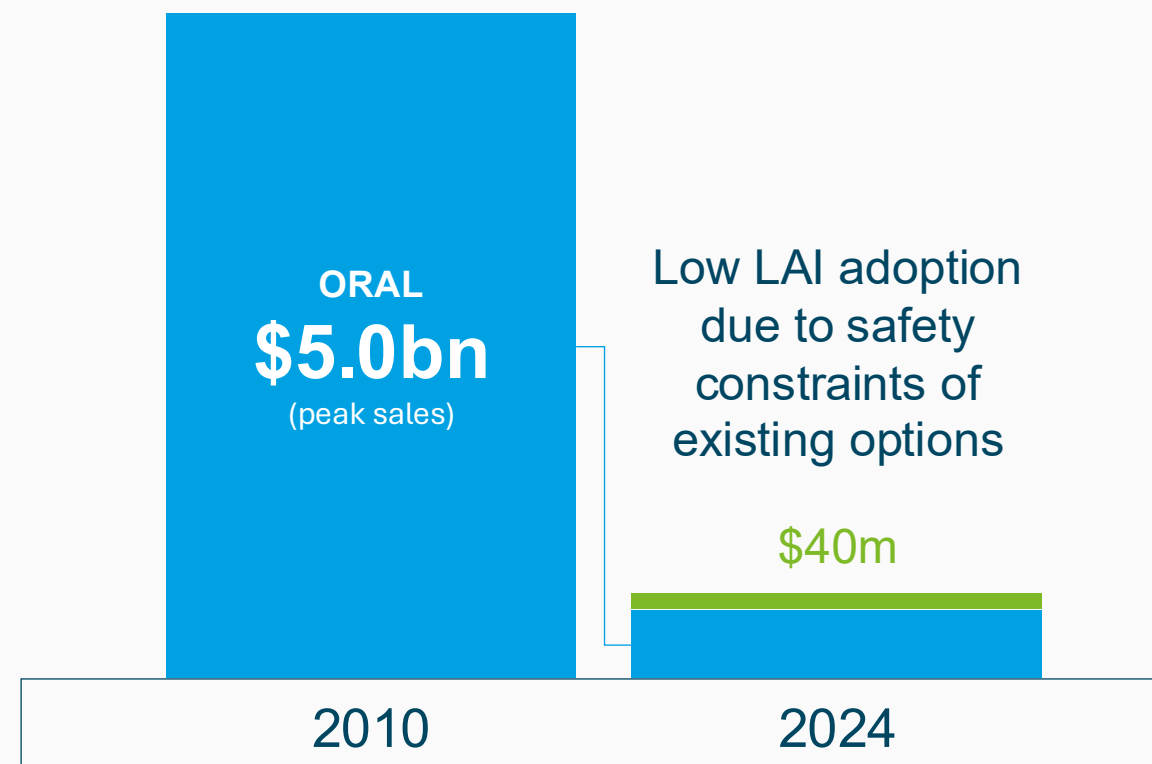
RISPERIDONE

Established LAI segment



OLANZAPINE

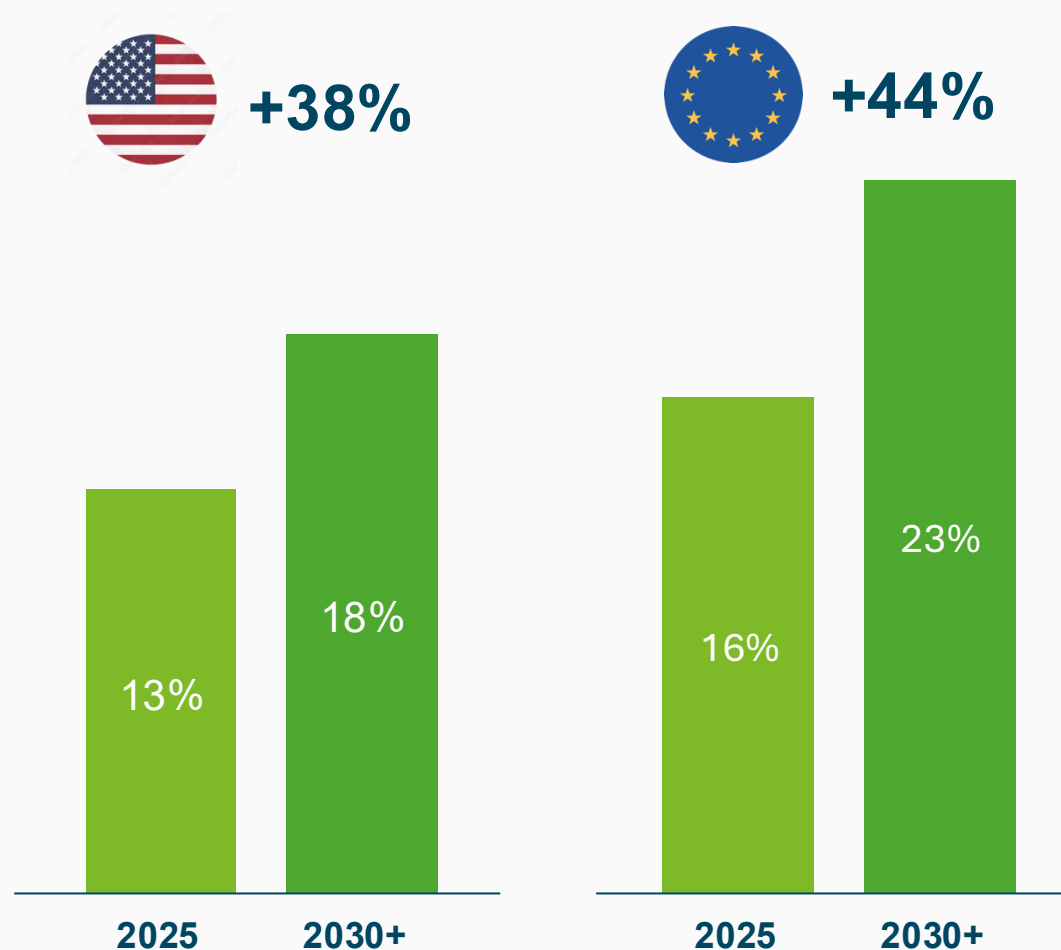
Significant untapped LAI opportunity



Very Strong Momentum in Schizophrenia LAI

LAI penetration in total treated schizophrenia population, in %

Source: LAI penetration based on IQVIA sales in Month of Therapy volume, with MIDAS (sales) dataset for EU and NDA Trx dataset for US / Teva's Innovation Strategy Day, May 2025



Teva is an Ideal Partner

**UZEDY® and Olanzapine LAI are key assets in Teva's
Pivot to Growth strategy**

**Teva's psychiatry sales force has shown very strong recent
commercial performance in the psychiatry field**

Worldwide reach

RISPERIDONE LAI (UZEDY®)

Monthly and every 2 months subcutaneous risperidone

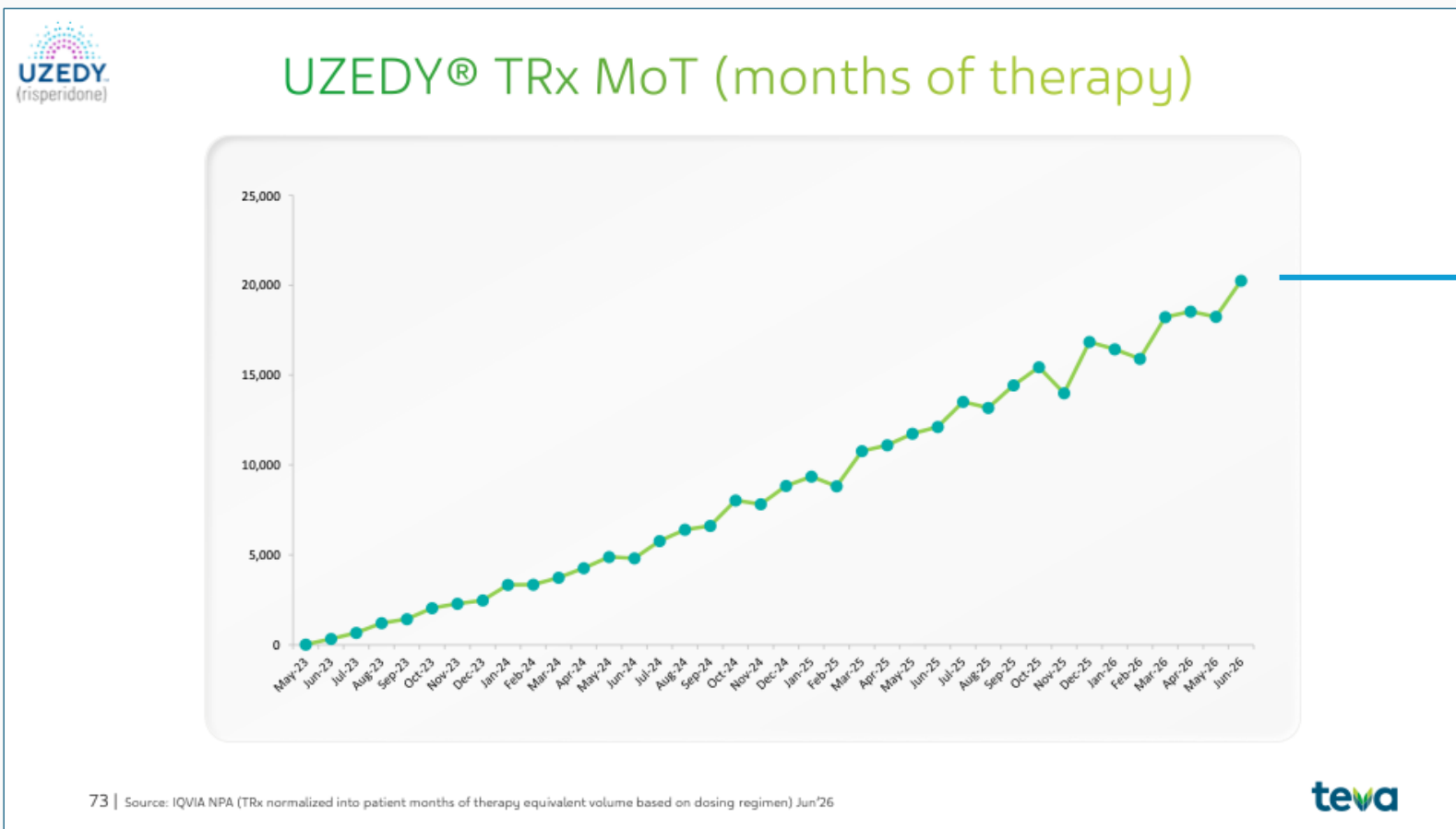
Approved for the maintenance treatment of Schizophrenia in adults

- 2023 United States
- 2025 South Korea (유제디), Canada (LONGAVO®)

Approved for the maintenance treatment of bipolar I disorder in adults (one-month dosing)

- 2025 United States

UZEDY® Continues Strong Commercial Momentum



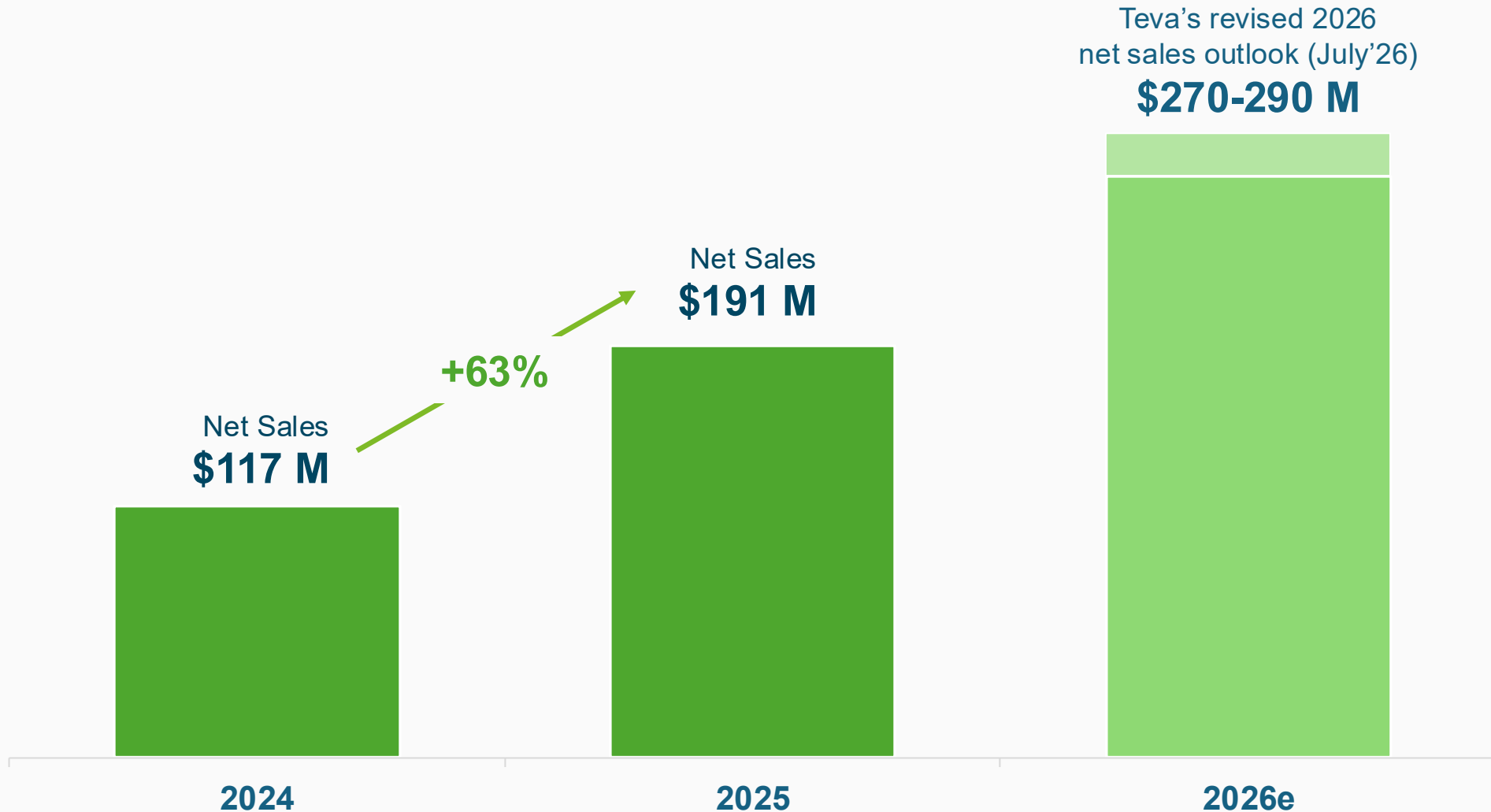
~20,000
Monthly Months of
Therapy in June 2026

+63%
YoY Growth in
Prescriptions

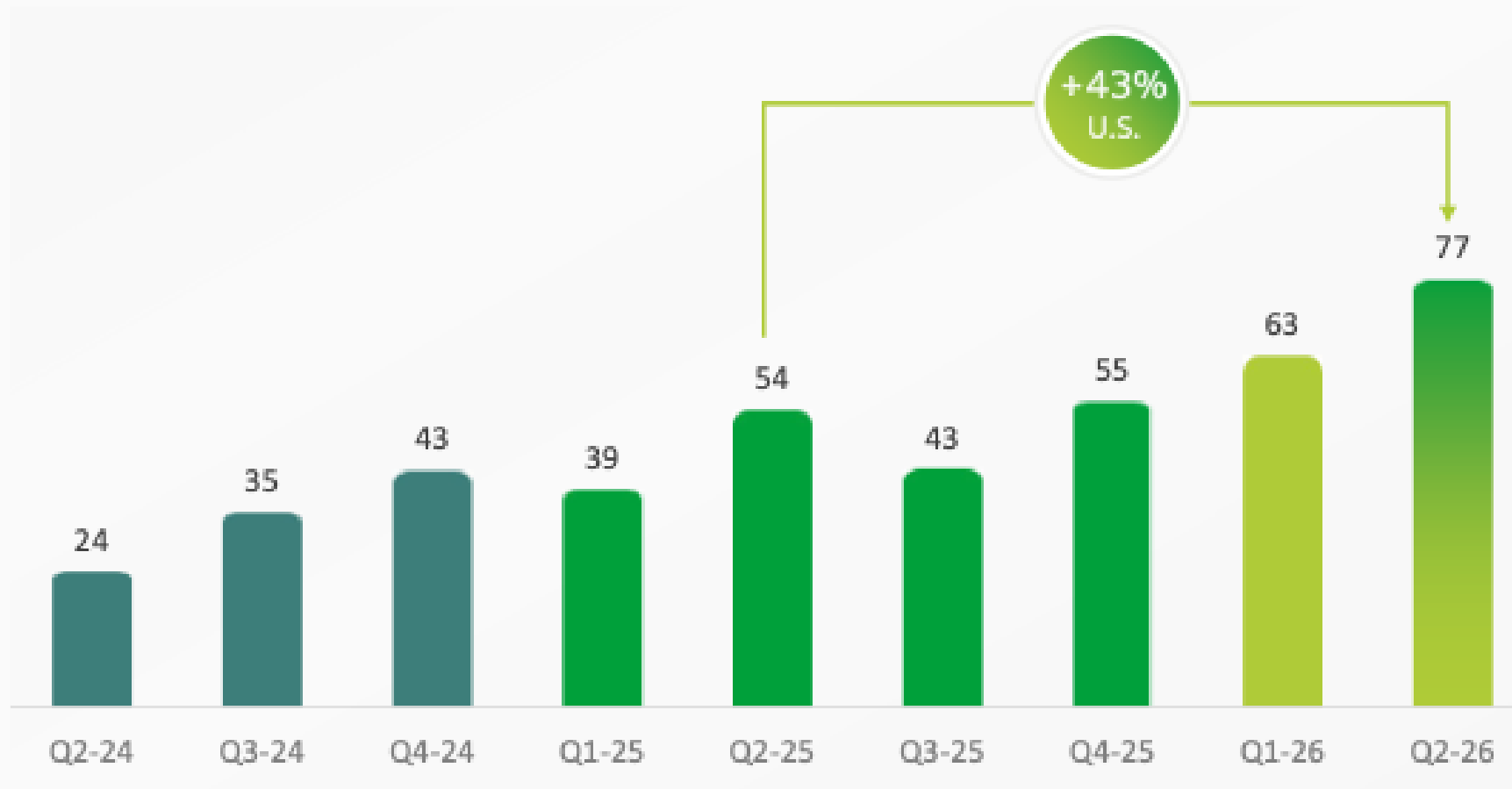
Source: IQVIA NPA data through June 2026, according to Teva. Prescription volume is normalized into months of therapy based on dosing regimen. Q2 2026 prescriptions increased 63% year-over-year.

Extract from Teva's Q2'26 Results Presentation - July 29, 2026

UZEDY® - 2025 Net Sales Growth: +63%



UZEDY® - Q2 2026 Net Sales Growth: +43%



Extract from Teva's Q2'26 Results Presentation - July 29, 2026

UZEDY® - Analysts' Peak Sales Forecasts¹

Analysts' Consensus: \$1.2B

Low: \$0.6B - High: \$2.2B

Revenue potential for Medincell



Royalties	Mid- to high-single digit on all net sales
Milestones	Commercial: \$105 M



¹ Based on most recent information received from analysts: Evercore, Truist, HC Wainwright, Leerinks, Jefferies, Stifel, Oddo BHF, Peter Degroof, Portzamparc, Kepler Cheuvreux, TP-Icap. Any opinions, estimates, or forecasts regarding Medincell's performance made by analysts are theirs alone and do not represent opinions, forecasts, or predictions of Medincell or its management.

UZEDY®: Medincell's Technology Enables Best-in-Class Features

Subcutaneous vs intramuscular

No loading dose or oral supplementation*

Ready-to-use

Same effect regardless of location of injection

*Therapeutic levels are reached within 6 to 24 hours after administration



UZEDY® Shows Strong Real-World Benefits

UZEDY® vs second-generation oral antipsychotics (SGOAs) – May 2025

RELAPSE RATE: -42%



INPATIENT RATE: -47%



MEAN TIME TO RELAPSE: +54%



MEAN LENGTH OF HOSPITAL STAY: -50%



MEAN ALL-CAUSE HCRU COSTS: -29%



Source: Poster presented by Teva Pharmaceuticals at the Annual Psych Congress Elevate, May 28–31, 2025, Las Vegas, NV: Treatment Patterns and Healthcare Resource Utilization Among Patients Receiving the Long-Acting Injectable Antipsychotic TV-46000 Versus Second-Generation Oral Antipsychotics. HCRU: healthcare resource utilization

Multiple Patents Granted on UZEDY®

RISPERIDONE LAI (UZEDY®)

US patent protection

2042

OLANZAPINE LAI

Once-monthly subcutaneous injection of olanzapine

December 9, 2025

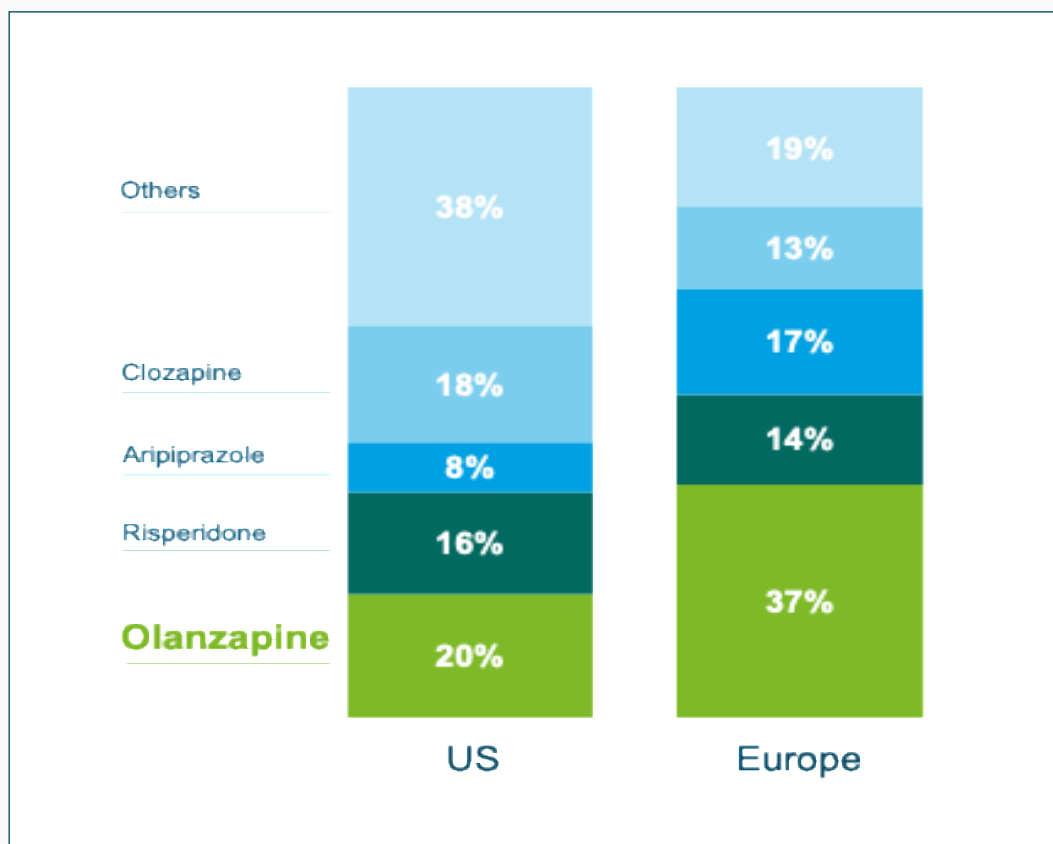
Submission to U.S. FDA for the maintenance treatment of Schizophrenia in adults

> *FDA approval anticipated in Q4 2026*

April 28, 2025

European submission to for the maintenance treatment of Schizophrenia in adults

Strong Unmet Need and Large Patient Opportunity for Subcutaneous Olanzapine LAI



→ One of the most widely used oral antipsychotic in schizophrenia

Used for most severe and refractory patients

No valid competitor

Olanzapine LAI - Analysts' Peak Sales Forecasts¹

Analysts' Consensus: \$2.0B

Low: \$1.5B - High: \$2.9B



EVERCORE

LEERINK PARTNERS

TRUIST 

Jefferies

 HCW
H.C. WAINWRIGHT & CO.

STIFEL



 ODDO BHF

 TP ICAP

 Degroof
Petercam

 Kepler
Cheuvreux

 PORTZAMPARC
BNP PARIBAS GROUP

¹ Based on most recent information received from analysts: Evercore, Truist, HC Wainwright, Leerinks, Jefferies, Stifel, Oddo BHF, Peter Degroof, Portzamparc, Kepler Cheuvreux, TP-Icap. Any opinions, estimates, or forecasts regarding Medincell's performance made by analysts are theirs alone and do not represent opinions, forecasts, or predictions of Medincell or its management.

Revenue potential for Medincell

Royalties	Mid- to high-single digit on all net sales
Milestones	US Approval: \$4 M EU Approval: \$3 M Commercial: \$105 M

Subcutaneous Olanzapine LAI was Designed to Resolve Eli Lilly's PDSS-Linked Market Failure

Risk of PDSS* associated with Eli Lilly product's

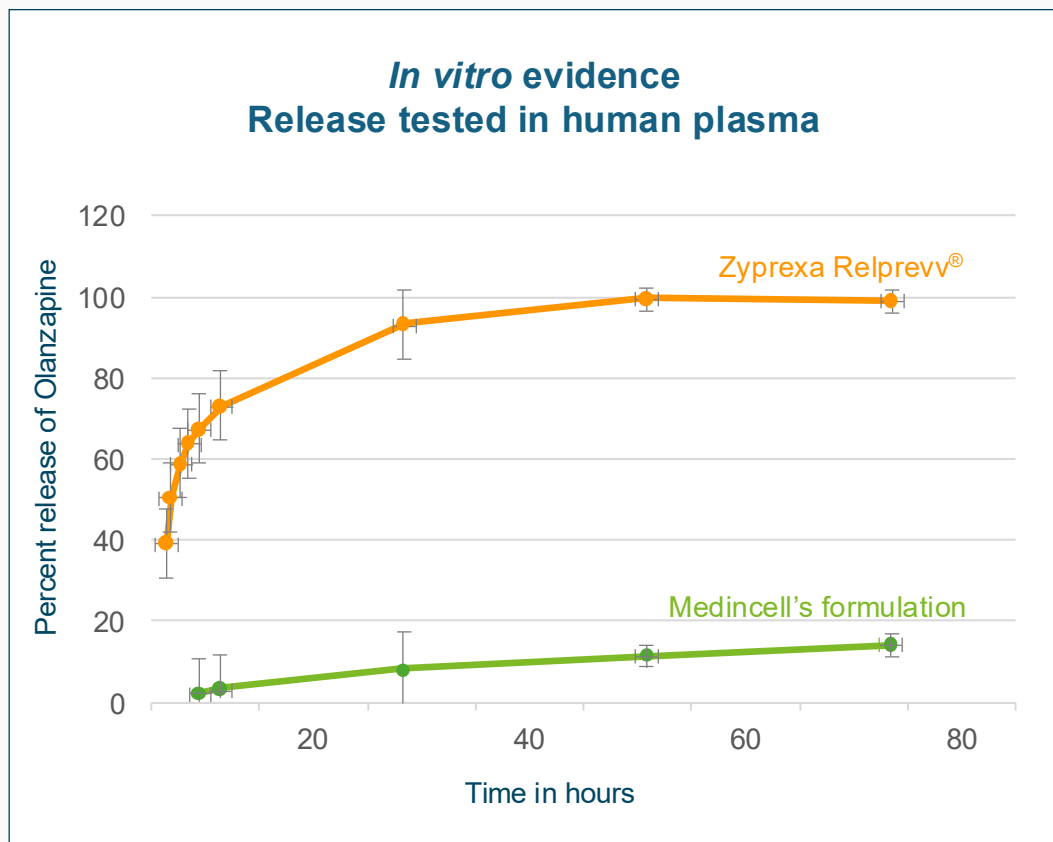
- Overdose-like reaction risk at each injection (< 0.1%)
- Caused by unintended intravascular burst exposure during injection

Regulatory restrictions on use

- Mandatory 3-hour monitoring on site after each injection
- Restricted distribution (REMS)

* Post-Injection Delirium and Sedation Syndrome

Data Support a Favorable Safety Profile, Including No Observed PDSS During Clinical Development



Existing long-acting olanzapine
constrained by REMS and 3-hour
PDSS monitoring

Subcutaneous Olanzapine LAI has
shown no observed PDSS across
4,000 injections during clinical studies

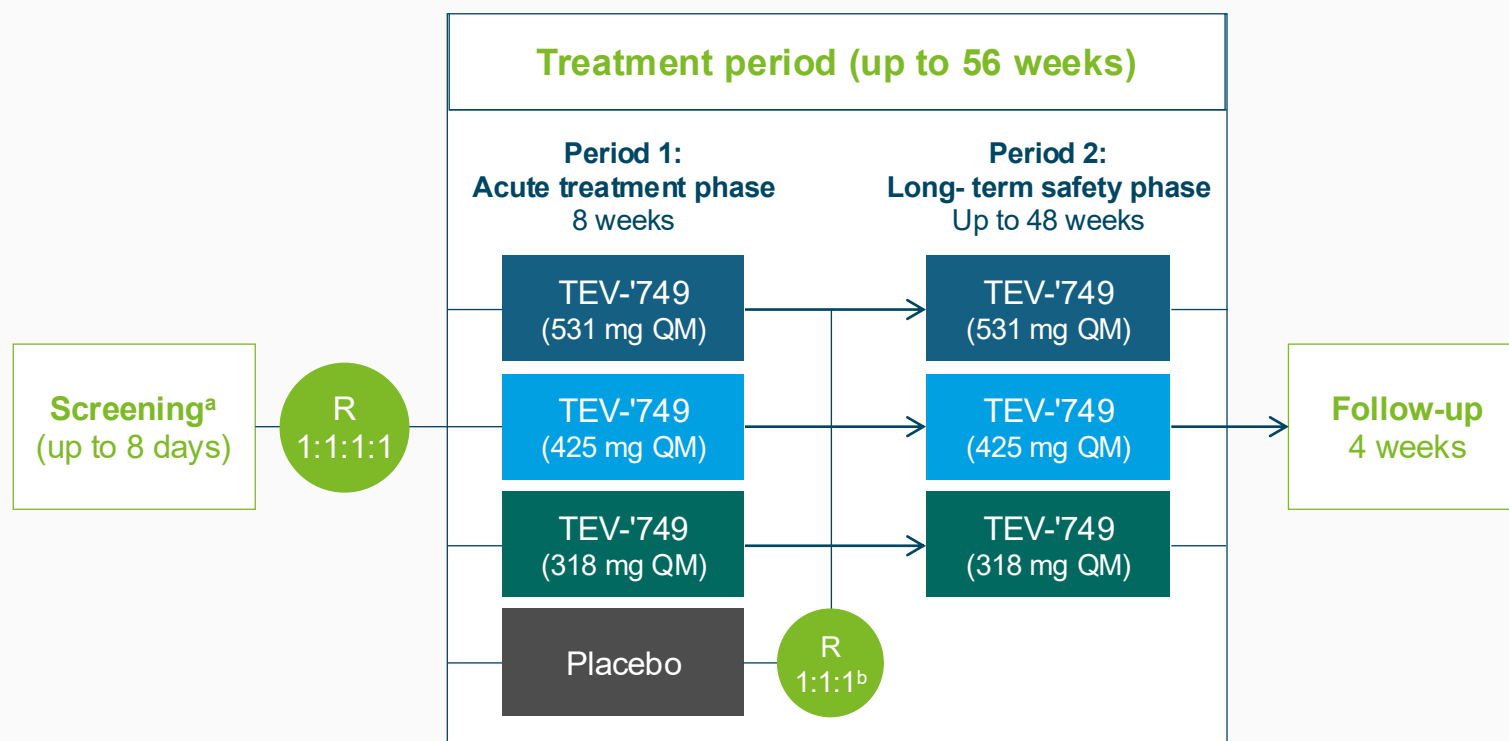
→ *In vitro* data support these findings

“No required monitoring for Medincell Olanzapine LAI is expected to drive Olanzapine LAI usage growth.”

Teva Q3 2025 Results presentation, Nov. 5, 2025

SOLARIS - Pivotal Phase 3

Phase 3 randomized, double-blind, placebo-controlled (Period 1), open-label long-term safety (Period 2) trial to evaluate efficacy, safety in adult patients with acute exacerbation of schizophrenia



Period 1 (8 weeks, n=675) aimed to assess the efficacy and safety, of TEV-'749 schizophrenia.

In Period 2 (n=423), Period 1 TEV-'749 participants retained their treatment, placebo patients were re-randomized 1:1:1 to TEV-'749 (318 mg, 425 mg, or 531 mg).

TEV-'749 doses were comparable to daily oral olanzapine doses of 10 mg, 15 mg, and 20 mg.

a. Participants entering the trial who had not previously received oral olanzapine within the last year received 2 oral doses of olanzapine for 2 consecutive days at the screening period to assess tolerability. The investigator verified the previous use, tolerability, and duration of olanzapine treatment to assure prior tolerability.
b. To maintain the blinding in Period 1, all participants were re-randomized between Periods 1 and 2; participants previously assigned to the active treatment groups retained their Period 1 treatment assignment (rerandomization was done to maintain blinding of Period 1, and de facto is a deterministic assignment and not randomization), and participants previously assigned to placebo were randomized to one of the active treatment groups in a 1:1:1 ratio. Participants were hospitalized for >28 days after receiving the first injection. QM, once monthly; R, randomization.
Correll CU, et al. Poster #97 presented at the 38th Psych Congress 2025; December 17-21, 2025; San Diego, CA, USA

SOLARIS - Positive Results

Demonstrated long-term clinical effectiveness in all groups

- **Improvements in PANSS Total Score**
 - **Sustained improvement in CGI-S and PSP scale scores**
-

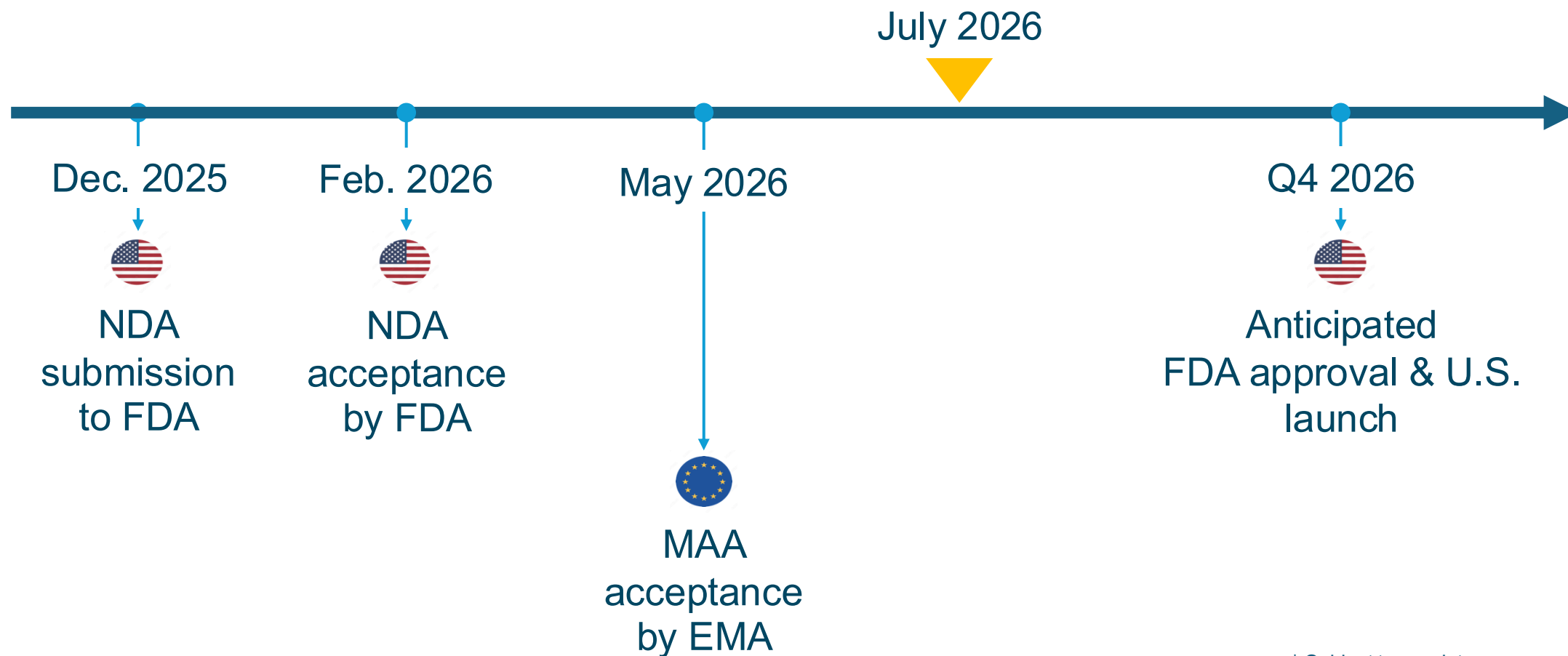
Long-term safety profile consistent with other olanzapine formulations

- **No new systemic safety signals**
- **Mild/moderate injection site reactions decreasing with continued dosing**
- **No PDSS Events**

Sources: Teva Pharmaceuticals – Poster presented at the 38th Psych Congress 2025; September 17-21, 2025; San Diego, CA, USA

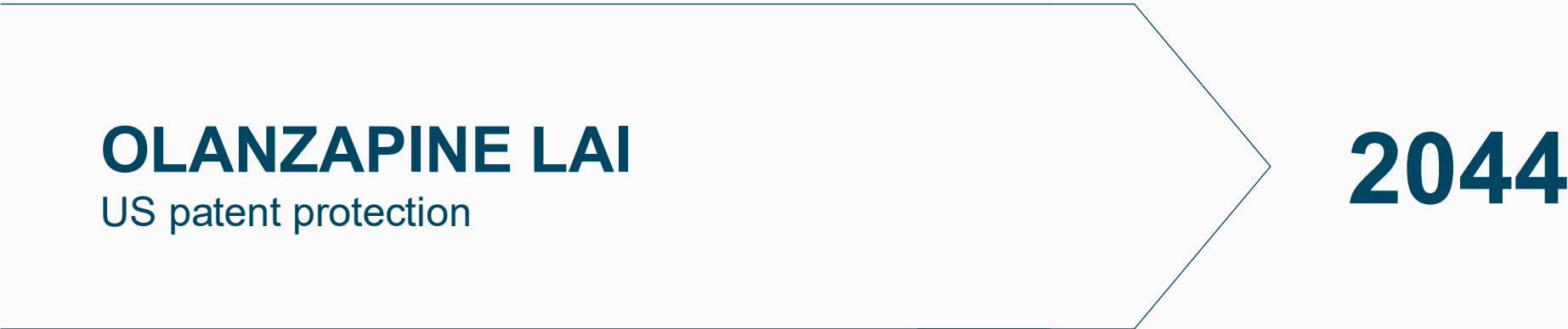
Final Countdown to Subcutaneous Olanzapine LAI

U.S. Launch Anticipated in Q4 2026*



* Subject to regulatory approval

Multiple Patents Granted on Olanzapine LAI



OLANZAPINE LAI

The diagram features a large, light blue chevron pointing to the right. Inside the chevron, the text 'OLANZAPINE LAI' is written in bold, dark blue capital letters, with 'US patent protection' in a smaller, dark blue font below it. To the right of the chevron's tip, the year '2044' is displayed in large, bold, dark blue numbers.

US patent protection

2044

AbbVie #1

API and indication not disclosed

April 2024

Strategic co-development and licensing agreement with AbbVie (up to 6 programs), \$35 million upfront

September 2024

Initiation of preclinical and supportive CMC work to advance AbbVie #1 into clinical development

Expected to Be Ready for Launch of Clinical Development in 2027

AbbVie #1 Expected to Be Ready for Launch of Clinical Development in 2027

All pre-IND activities conducted by Medincell

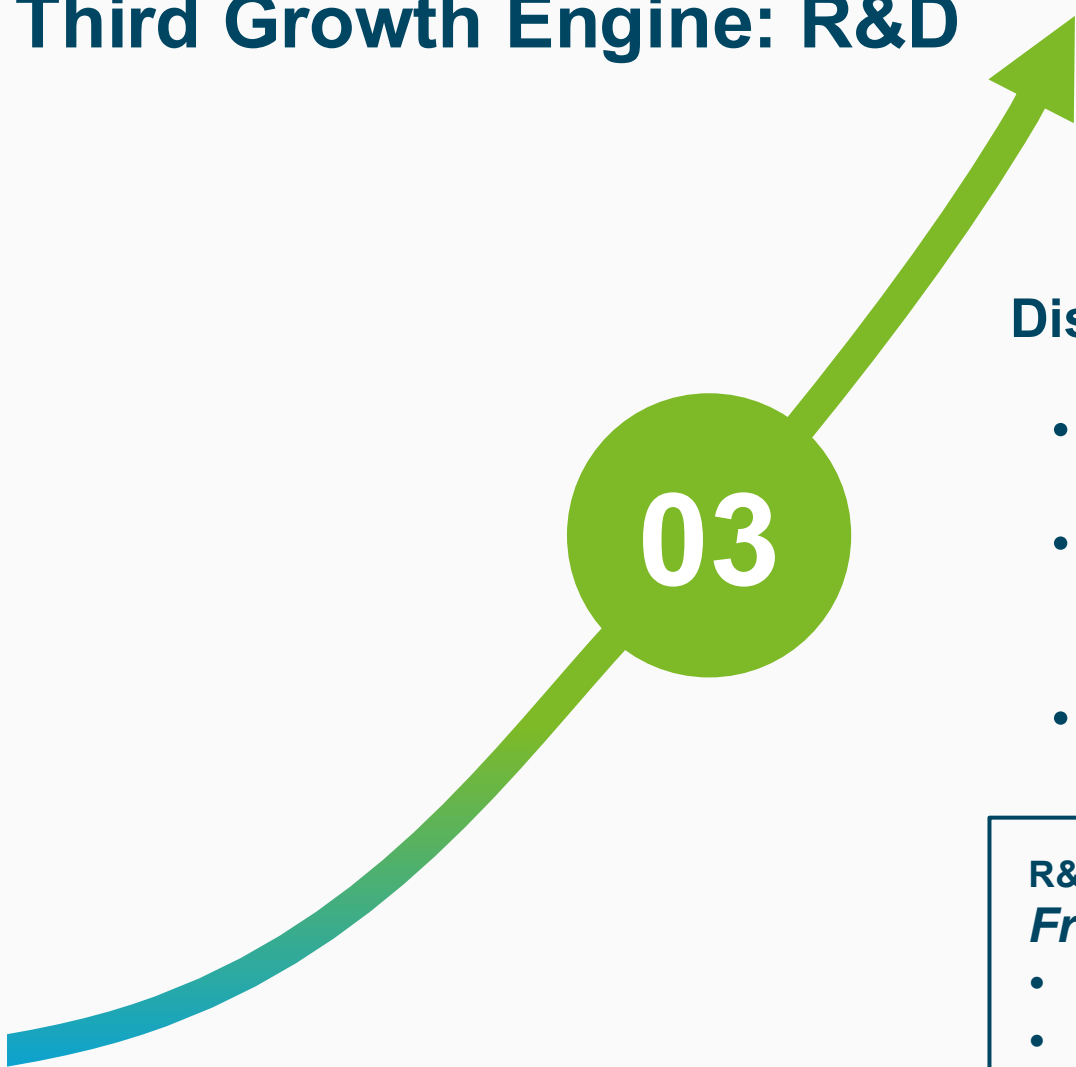
AbbVie to conduct all clinical development

**Mid-single- to low-double-digit royalties
\$315 million in potential milestones**

Third Growth Engine Pipeline & Innovation

“Shift to Growth” Strategy

The Third Growth Engine: R&D



03

Disciplined Capital Allocation to R&D

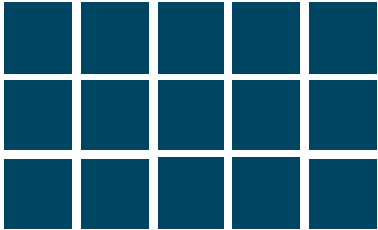
- **Broaden our Innovation Platform**
- **Expand our Portfolio with Blockbuster-Potential LAIs**
- **Extend our Network of Top-Tier Partners**

R&D Day – May 12, 2026

From Science to Long-Term Value Creation

- Replay & transcript, [click here](#)
- Presentation, [click here](#)

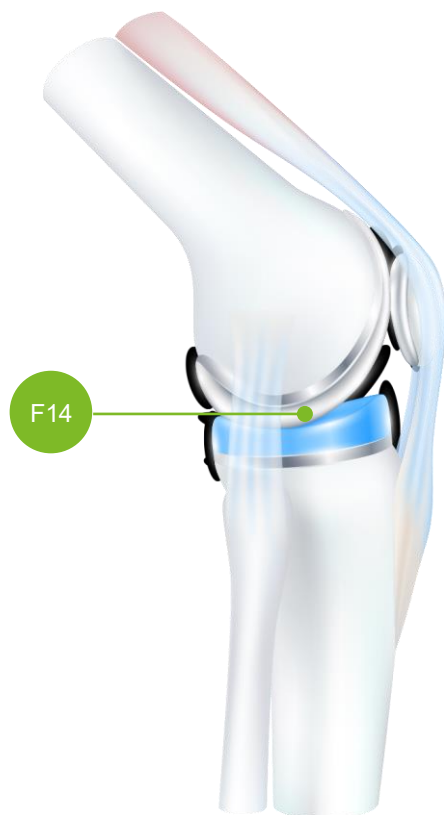
Advancing Innovation Across the Pipeline

FORMULATION	PRECLINICAL	CLINICAL PHASE 3	NDA	MARKET
~15 in-house and partnered programs 	AbbVie #1 Confidential API and indication 	Celecoxib Intraarticular (mdc-CWM) Postoperative pain 	Olanzapine LAI (mdc-TJK) Schizophrenia 	Risperidone LAI UZEDY® Schizophrenia / BP-I 

Global Health programs

Tuberculosis Macozinone 	Contraception (mdc-WWM) Progestin 6-Month Gates Foundation
	Malaria (mdc-STM) Ivermectin 3-Month Gates Foundation

mdc-CWM: Regulatory Path to Approval



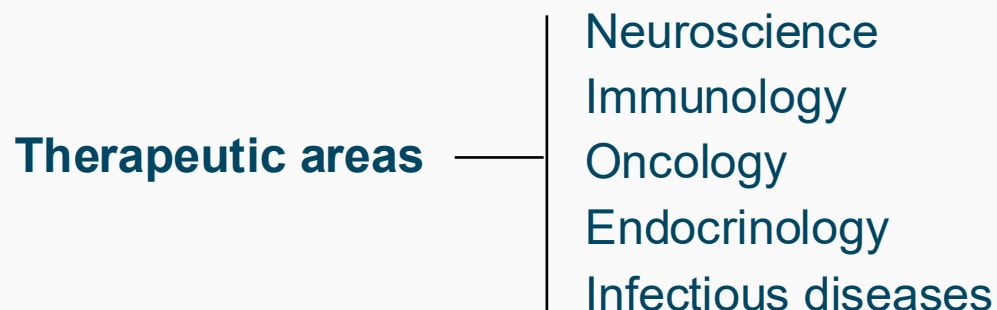
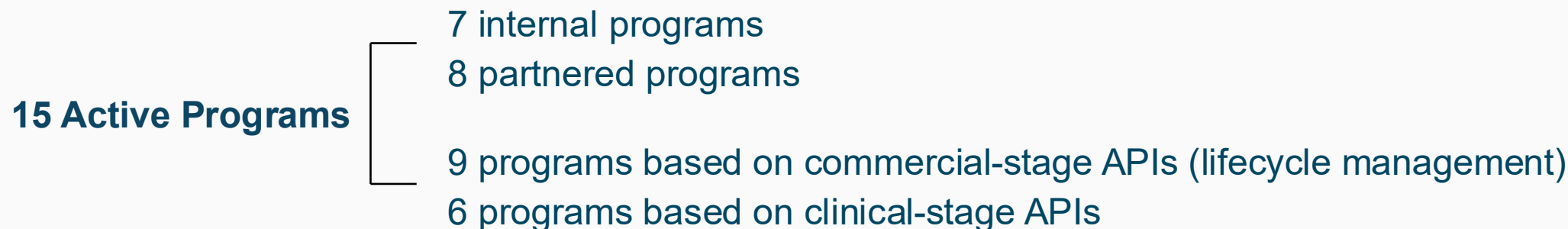
The first Phase 3 study missed the primary pain endpoint at Week 2, while showing positive results at Day 7 and on long-term functional outcomes

Clear differentiation from approved and investigational therapies

FDA meeting confirmed clear guidance for a second Phase 3 design focused on TKR-naïve subgroup

AIC is actively preparing the next Phase 3 study

A Diversified Early-Stage Pipeline with Strong Value-Creation Potential*



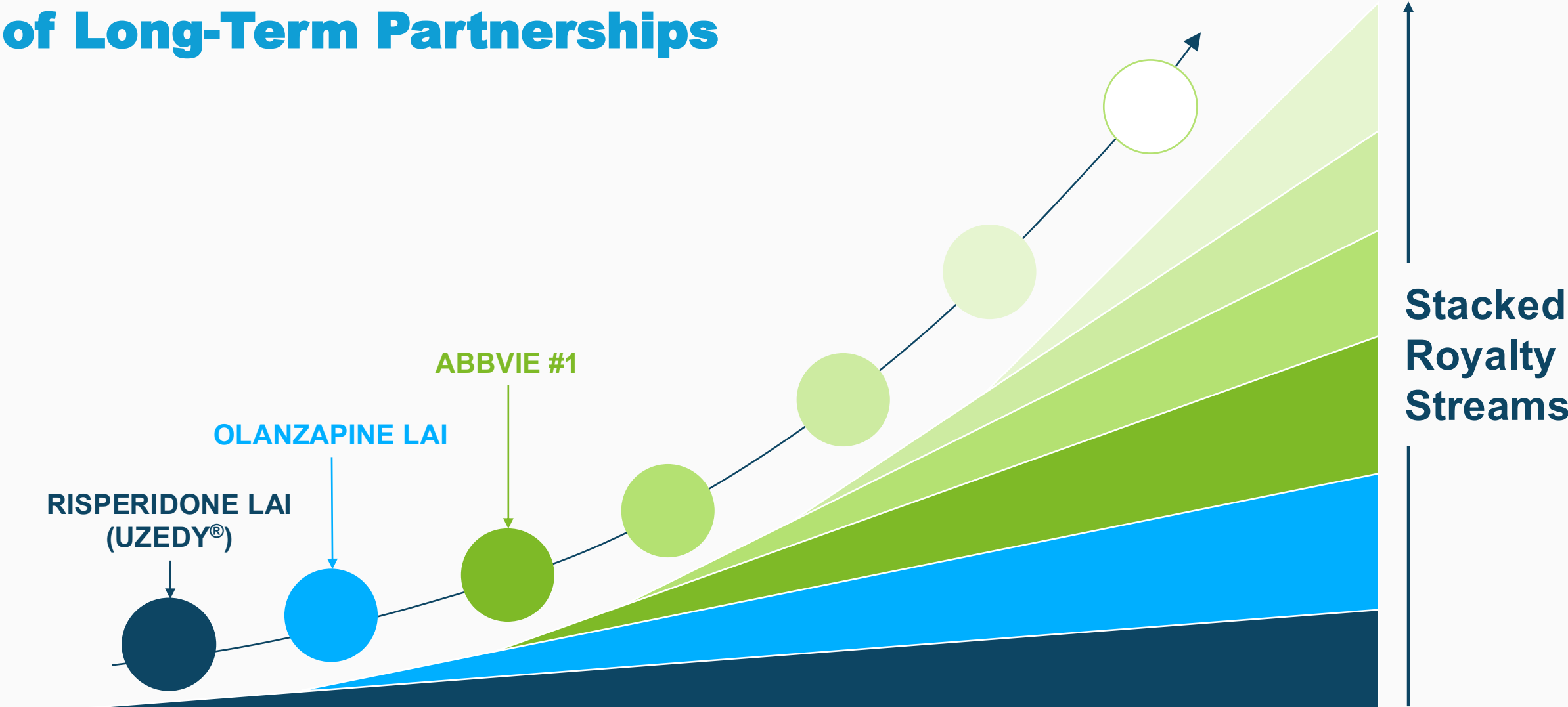
Several candidates with blockbuster potential

*As of May 12, 2026

Limited disclosure of early-stage pipeline to

- Proactively manage early technical and partner attrition
- Protect long-term competitive differentiation
- Respect confidentiality commitments

Compounding Value Through a String of Long-Term Partnerships



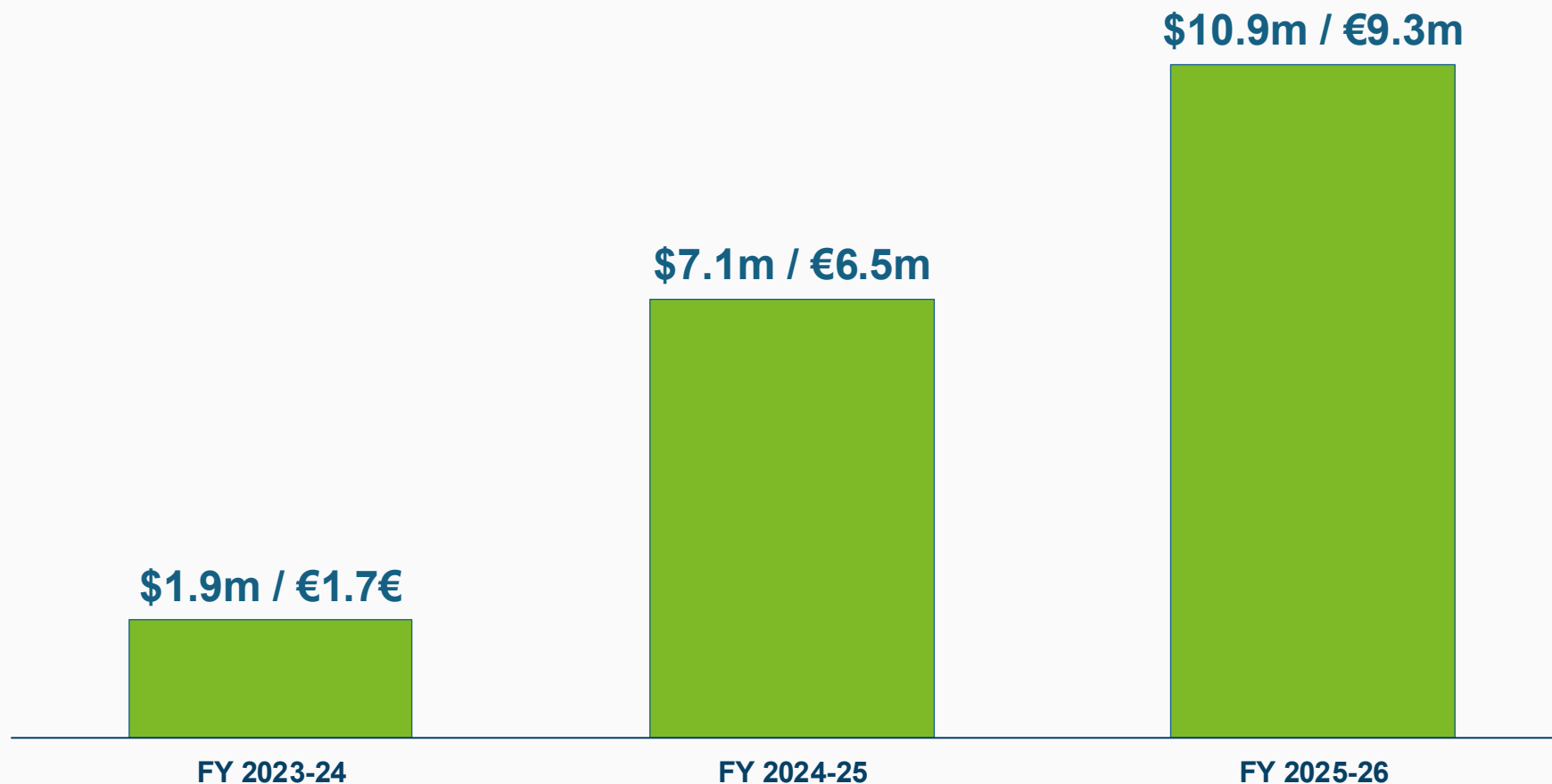
Financials

FY 2025-26

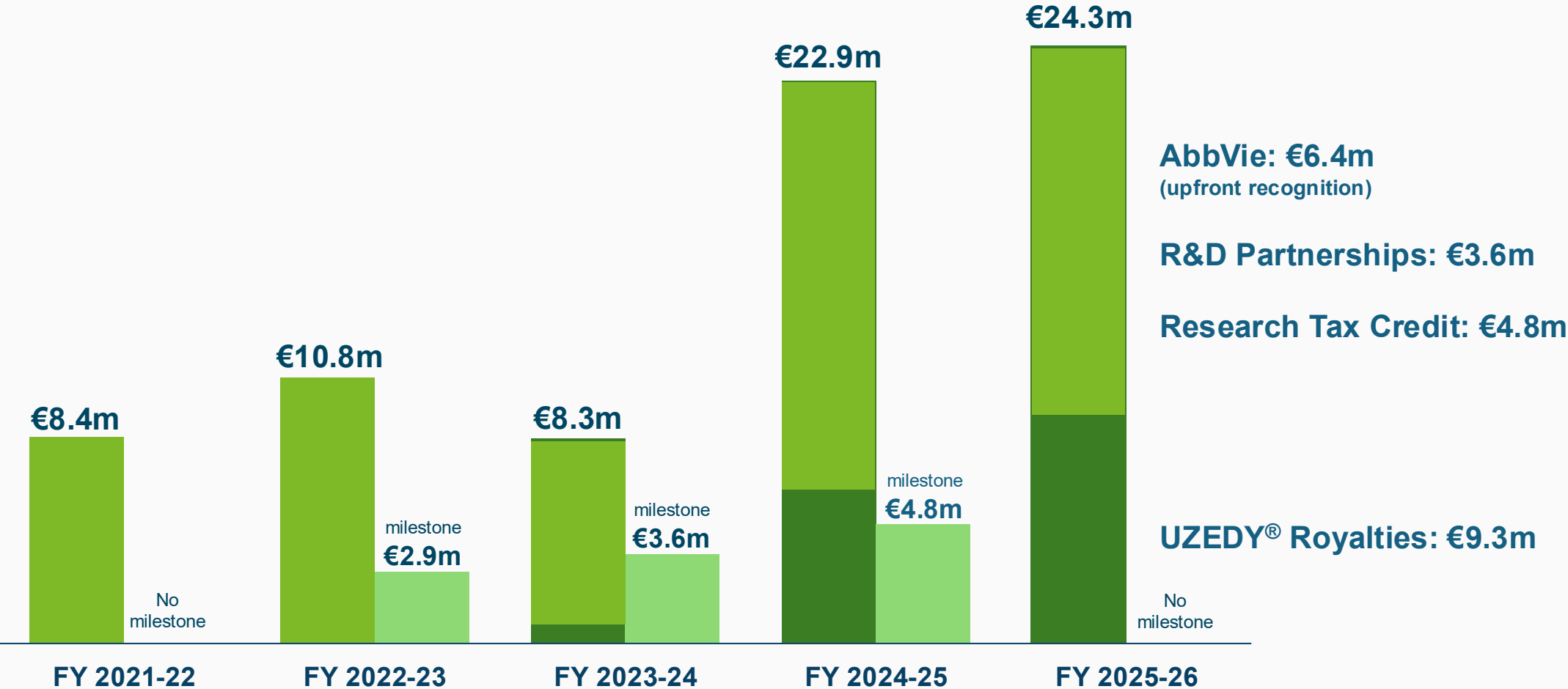
(ended March 31st, 2026)

UZEDY® Royalties

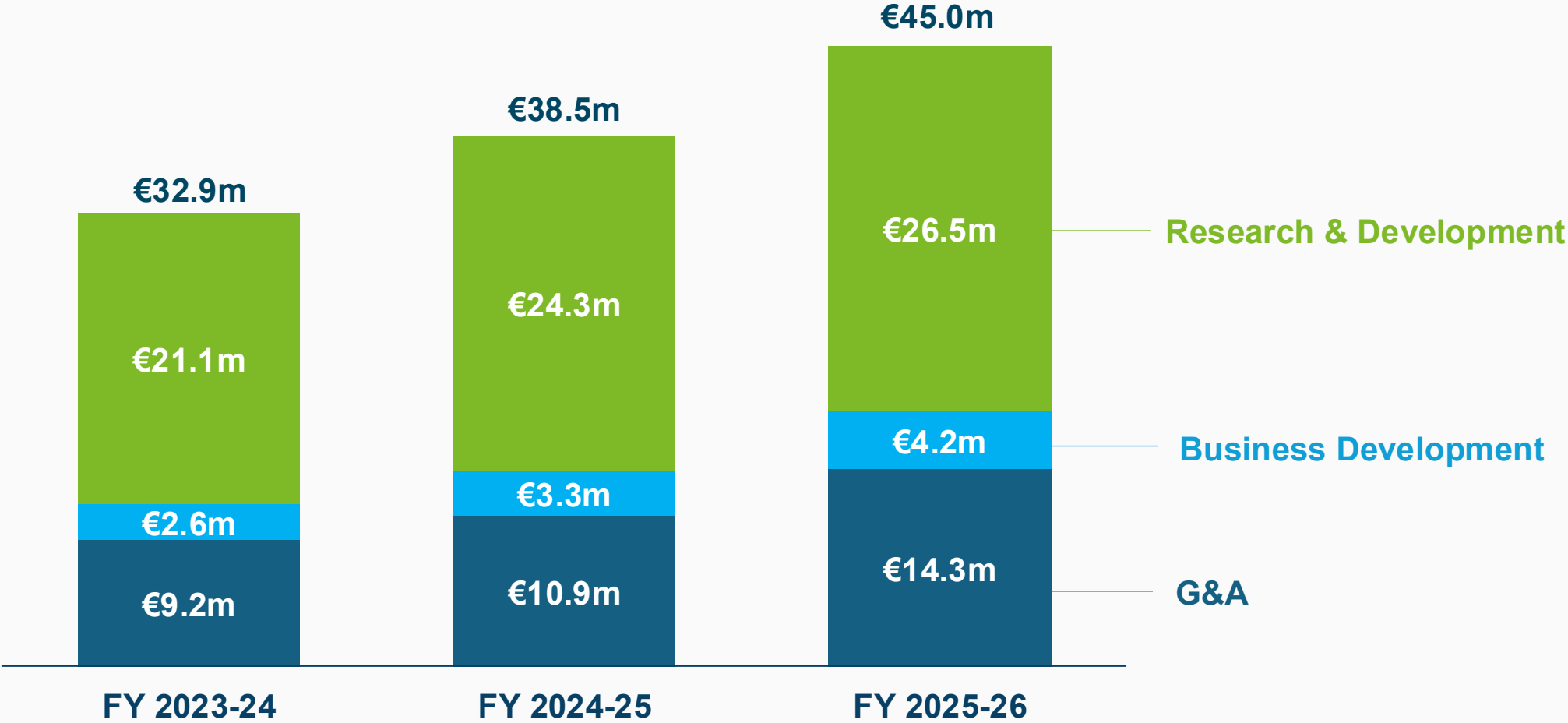
+54% in US\$, +42% in €



Investment Phase Continues While Royalties Scale



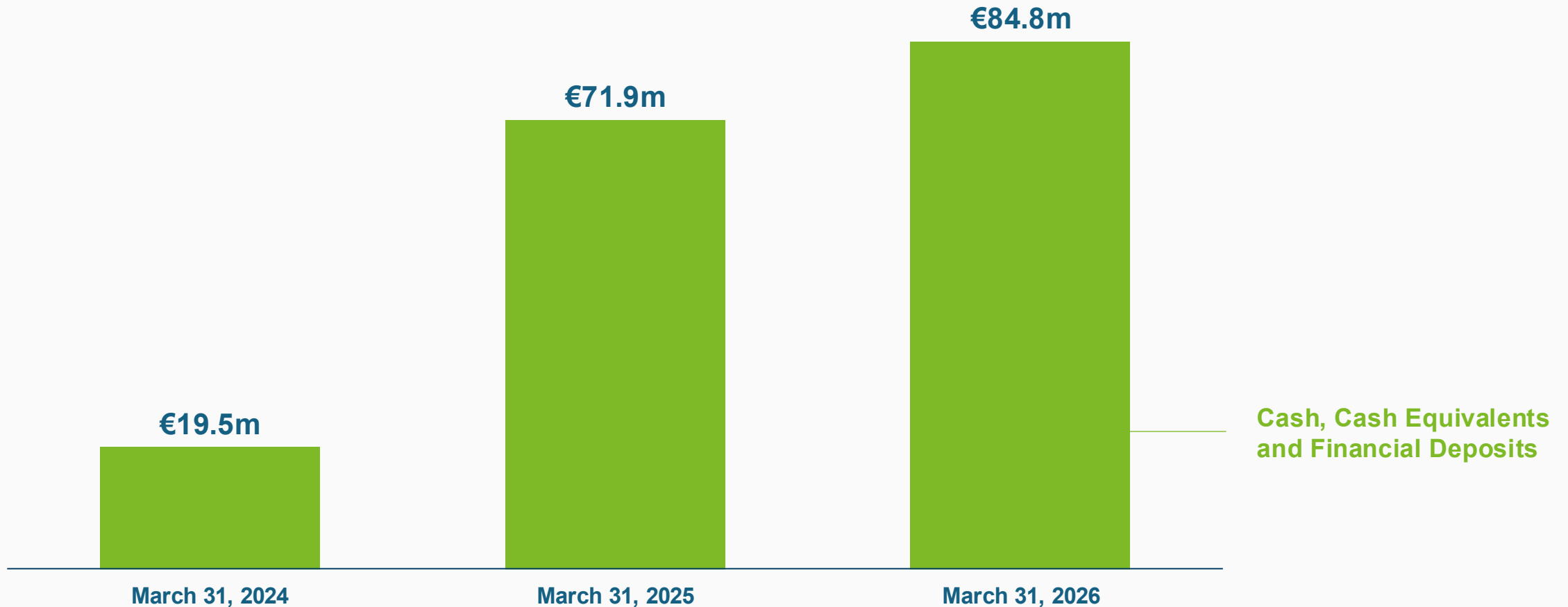
Expenses in Line with Scale-Up Strategy



P&L Driven by Growing Royalties and Program Phasing

Million €	FY2025-26	FY2024-25	Δ YoY
Total income	24.3	27.7	(3.4)
of which Milestones	0	4.8	n.a.
Operating expenses	(45.0)	(38.5)	(6.5)
Operating result	(20.8)	(10.8)	(10.0)
Financial result	(10.6)	(7.4)	(3.2)
Net result	(31.3)	(18.4)	(12.9)

Strengthened Balance Sheet Supports Continued Investment and Long-Term Value Creation



Path to Sustainable Growth



medincell.

Thank you