



FINANCE AVENUE | NOVEMBER 2024

# Reaching Patients Through Immunology Innovation

# Forward Looking Statements

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Est. 2008  
**argenx**



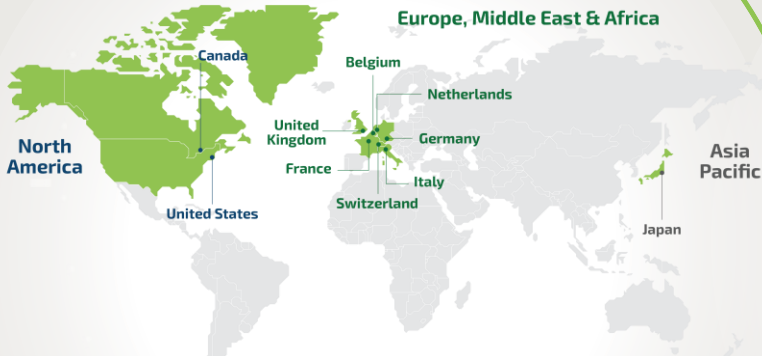
**Entrepreneurial** spirit –  
calculated risk based on data

**Immunology innovation** through  
model of **co-creation**

**Execution excellence**

# Building a Leading Immunology Company

## Global Footprint



**1,500+**

Global employees and consultants

## Rooted in Science

### Immunology Innovation Program

Translating immunology breakthroughs into first-in-class pipeline opportunities

### Wholly-Owned Pipeline

Efgartigimod	Empasiprubart	ARGX-119
FcRn Blocker	C2 Inhibitor	MuSK Agonist

### Key Validating Partnerships



## Reaching Patients

Global clinical programs ongoing in

**VYVGART<sup>®</sup>**  
(efgartigimod alfa-fcab)

**VYVGART<sup>®</sup> Hytrufo**  
(efgartigimod alfa and hyaluronidase-qvfc)

VYVGART first-approved FcRn blocker in US, Japan, China, Canada, EU, UK, Israel

**50,000**

Patients by 2030



Innovation has no  
meaning unless it  
reaches patients and  
provides real benefit

# Our Innovation Playbook

**Novel Disease  
Biology Insights**

**Foundational  
Immune  
Targets**

**Best-in-Field  
Antibody  
Engineering**

**First-in-Class  
Antibodies**

**Pipeline-in-  
a-Product  
Development**

**Differentiated  
Patient  
Outcomes**

# Our Innovation Horizons

ARGX-109  
(Anti-IL-6)

ARGX-213  
(Anti-FcRn)

ARGX-121  
(Anti-IgA)

ARGX-220

ARGX-119

CMS

ALS

Future  
Indications

PIPELINE

Empasiprubart

MMN  
Established PoC

DGF

DM

CIDP

Future  
Indications

VYVGART  
Opportunity

VYVGART®  
(efgartigimod alfa-fcab)  
Injection for Intravenous Use  
400 mg/20 mL vial

VYVGART® Hytrulo®  
(efgartigimod alfa and  
hyaluronidase-gvfc)  
Subcutaneous Injection  
180 mg/mL and 2000 U/mL vial

\$573M in gMG and CIDP  
revenue in Q3 2024



CIDP launched

ITP launched

5 registrational  
trials by YE 2024:  
oMG, snMG, TED,  
SjD, ITP-US

PFS MG, CIDP  
PDUFA April, 2025



# Delivering Innovation Across VYVGART Globally

## Driving impact with VYVGART

**MG** Rapid, deep, sustained responses achieved in patients

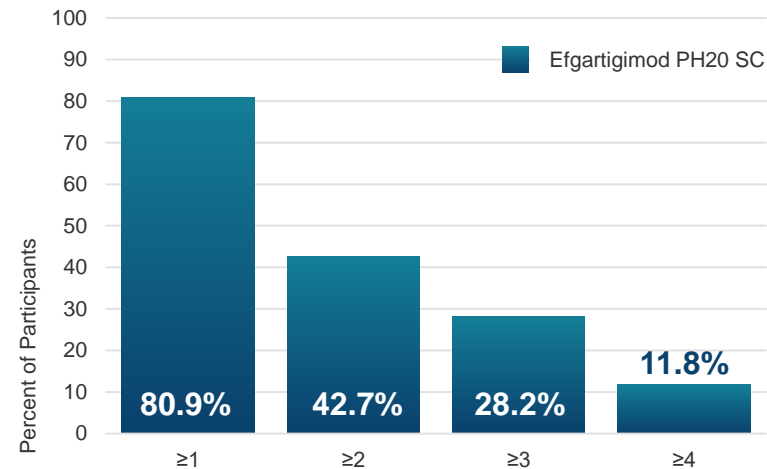
**54%**  
MSE<sup>1</sup>

**42%**  
TAPERED TO ≤ 5 MG  
STEROIDS<sup>2</sup>

**25%**  
TAPERED TO 0 MG  
STEROIDS<sup>2</sup>

**CIDP** Consistent responses and regain of function

### Functional Ability (aINCAT)



## Reaching Patients Globally with VYVGART Franchise

>10,000 patients on treatment<sup>1</sup>

VYVGART and VYVGART Hytrulo<sup>2</sup> approved across 3 continents within one calendar year

1. ADAPT/ADAPT+ combined real world and clinical data  
2. IQVIA Claims analysis: 1-year post efgartigimod initiation



# Driving Patient Growth with VYVGART

## PATIENT GROWTH



**60%**

Hytrulo patients new to VYVGART YTD<sup>2</sup>

Expanding VYVGART Hytrulo share

## EARLIER LINE PATIENTS

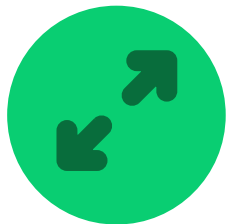


**~60%**

New VYVGART patients coming from orals<sup>1</sup>

US patients

## PRESCRIBER EXPANSION



**>3k**

Total VYVGART prescribers<sup>1</sup>

Breadth of prescribers

## PATIENT EXPERIENCE



**PFS**

PDUFA April 10th

Expanding patient reach

1. VYVGART and VYVGART Hytrulo  
2. VYVGART Hytrulo only

# We Aim to Address the Unseen Suffering in CIDP

A man with grey hair, wearing a dark quilted vest over a light-colored long-sleeved shirt and blue trousers, sits on a wooden bench in a garden. He is leaning on a black cane with both hands. The background shows a path leading through green foliage and a warm, orange-toned sky, suggesting a sunset or sunrise. The overall mood is contemplative and serene.

**≤20%**

of patients achieve remission on current SOC (CDAS=2)\*

**>50%**

of patients are dissatisfied with their symptom burden\*\*

**>88%**

of treated patients report residual neurological symptoms, including muscle weakness, sensory symptoms, pain, and fatigue \*\*\*

**>42K**

treated CIDP patients in US & ROW argenx markets (ex-China)\*\*\*\*

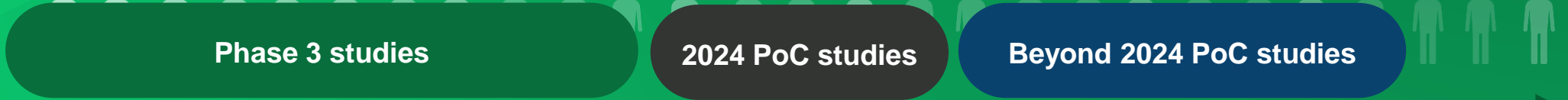
\*Gorson KC, et al. 2010

\*\* Mendoza M, et al. 2023

\*\*\*Bunschoten C et al. 2019

\*\*\*\* argenx market research

# This is Just the Beginning



REMAINING READOUTS IN 2024

**Myositis**  
**Bullous Pemphigoid**

**MG**  
Launched 2022  
**80K**

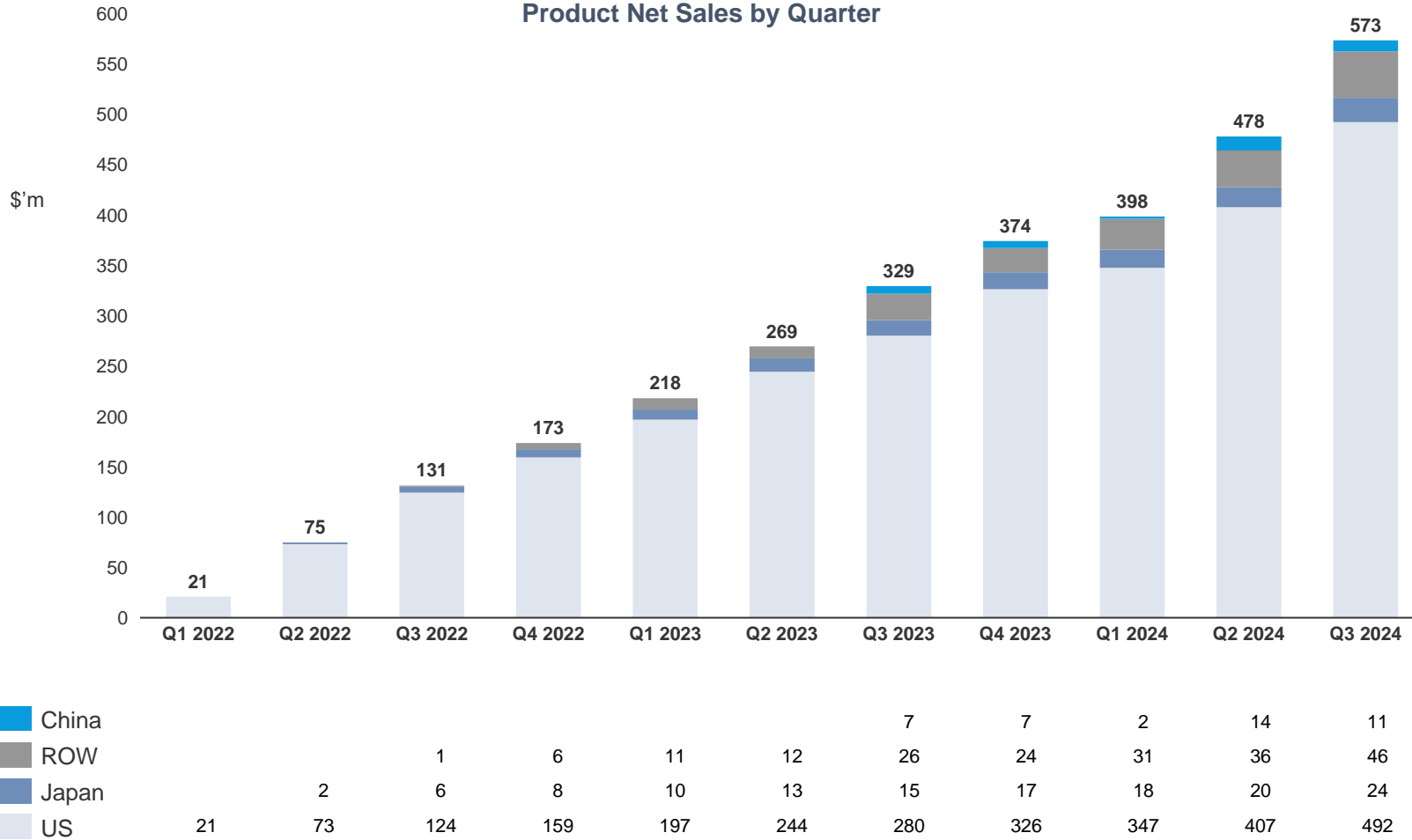
**CIDP**  
Approved June 21, 2024  
**42K**

**ITP**  
Approved in Japan | March 26, 2024  
**17K**

● On track for 2024 Ph3 start

# Product Net Sales: Q3 of \$573 million

Product Net Sales by Quarter



Q3 2024: growth of 74% vs Q3 2023

(in millions of \$)	Q3 2024	Q3 2023	Growth % *
US	492	280	76%
Japan	24	15	49%
ROW	46	26	76%
China supply	11	7	44%
<b>Total</b>	<b>573</b>	<b>329</b>	<b>74%</b>

Q3 2024: growth of 20% vs Q2 2024

(in millions of \$)	Q3 2024	Q2 2024	QoQ % Growth *
US	492	407	21%
Japan	24	20	20%
ROW	46	36	28%
China supply	11	14	(21)%
<b>Total</b>	<b>573</b>	<b>478</b>	<b>20%</b>
<b>Total excluding China</b>	<b>562</b>	<b>464</b>	<b>21%</b>

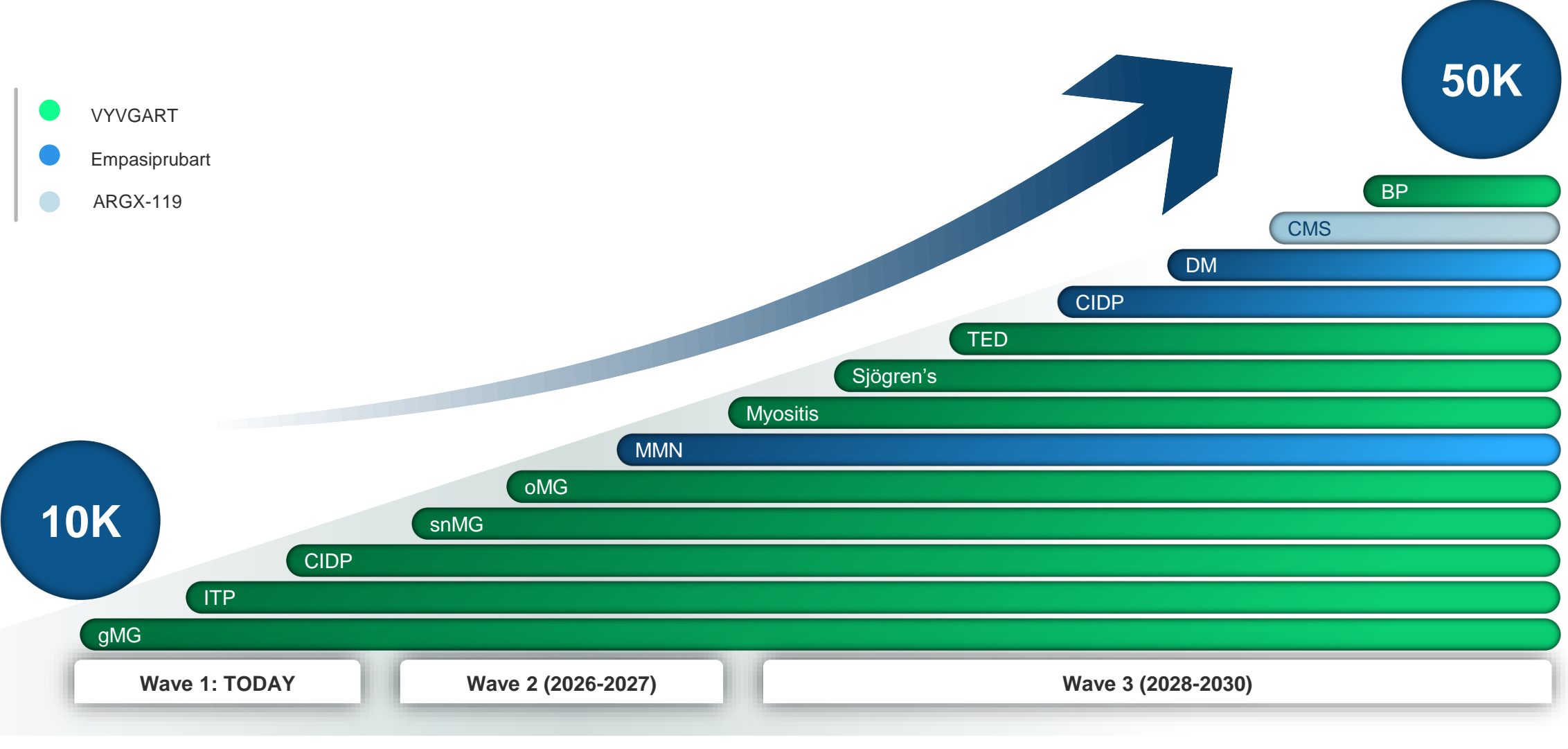
\*Net sales growth % excludes the impact of fx.



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# Opportunities Ahead Strengthen Our Leadership



# Our Innovation Horizons

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# Vision 2030

5

New Molecules  
in Phase 3

10

Labeled  
Indications

50k

Patients on  
Treatment

## COMMITMENT TO OUR TRANSFORMATION MISSION

Continuous Pipeline of  
Innovation

Leadership in FcRn

Disciplined Scaling