UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
WASHINGTON, D.C. 20549

FORM 6-K

REPORT OF FOREIGN PRIVATE ISSUER
PURSUANT TO RULE 13a-16 OR 15d-16
UNDER THE SECURITIES EXCHANGE ACT OF 1934

For the Month of March 2023

Commission File Number: 001-38097

ARGENX SE

(Translation of registrant's name into English)

Laarderhoogtweg 25
1101 EB Amsterdam, the Netherlands
(Address of principal executive offices)

Indicate by check mark whether the registrant files or will file annual reports under cover of Form 20-F or Form 40-F.

Form 20-F ☒ Form 40-F ☐

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1): ☐

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7): ☐

argenx SE

On March 2, 2023, argenx SE (the "Company") issued press releases, copies of which are attached hereto as Exhibit 99.1 and 99.2, and an investor presentation, a copy of which is attached hereto as Exhibit 99.3. Each of Exhibit 99.1, 99.2 and 99.3 is incorporated by reference herein.

The information contained in this Current Report on Form 6-K, including Exhibit 99.1 and Exhibit 99.2, and an investor presentation, a copy of which is filed hereto as Exhibit 99.3, each of which is incorporated by reference into the Company's Registration Statements on Forms F-3 (File No. 333-258251) and S-8 (File Nos. 333-225375 and 333-258253).

EXHIBITS

Exhibit	Description
99.1	Press Release dated March 2, 2023
99.2	Press Release dated March 2, 2023
99.3	Investor Presentation dated March 2, 2023

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

ARGENX SE

Date: March 2, 2022

By: /s/ Hemamalini (Malini) Moorthy
Hemamalini (Malini) Moorthy
General Counsel

Exhibit 99.1

argenx Announces Planned Transition of Chief Operating Officer

- Karen Massey appointed as Chief Operating Officer, effective March 13, 2023
- Keith Woods to retire and serve as Advisor on argenx Board of Directors

March 2, 2023

Amsterdam, the Netherlands — argenx (Euronext & Nasdaq: ARGX), a global immunology company committed to improving the lives of people suffering from severe autoimmune diseases, today announced that Karen Massey will join argenx as Chief Operating Officer, effective March 13, 2023. Ms. Massey will succeed Keith Woods, who will remain at argenx for a transition period to support the launch of subcutaneous efgartigimod, after which he will retire and serve as a strategic advisor on the Commercial Committee of the argenx Board of Directors.

"Karen brings extensive commercial launch and leadership experience to argenx, specifically within neurology and autoimmune disease. Most recently, her leadership accelerated the post-launch performance of Ocrevus in multiple sclerosis, a dynamic market that is often compared to myasthenia gravis. Karen has a genuine passion for innovation and an ability to inspire large, diverse and global teams to deliver results by creating a sense of shared purpose and community," commented Tim Van Hauwermeiren, Chief Executive Officer at argenx. "Karen's appointment is the result of careful succession planning and a thorough process to find the right leader at this important time in our company history. I am confident that she will make a significant impact at argenx as we work together to bring innovative medicines to patients."

"I would like to extend a warm thank you to my friend and colleague, Keith Woods, who joined the company in 2018 with a commitment to build a global infrastructure ahead of our first commercial launch. He accomplished this and so much more – leading us through a very successful first launch-year while always inspiring our colleagues to keep the patient at the center of everything we do. I wish him well in his much-deserved next phase of life and look forward to his ongoing contributions as an advisor on our Board," continued Mr. Van Hauwermeiren.

Ms. Massey has over 20 years' experience in the pharmaceutical and biotechnology industry with an impressive breadth of experience across commercial, product development, corporate strategy & innovation. She joins argenx from Genentech (Roche Group), where she most recently served as Senior Vice President of Product Development and Global Clinical Operations. In this capacity, she led a global organization of more than 2,000 people across 24 countries and was accountable for operationalizing over 300 clinical trials within oncology, neuroscience, infectious diseases, immunology and ophthalmology. During her 9-year tenure at Genentech, Ms. Massey held various commercial leadership roles across marketing and business operations, including as the Vice President of the Multiple Sclerosis and Neuromyelitis Optica business, leading the team to accelerate the post-launch performance of Ocrevus. Ms. Massey started her biopharmaceutical career in marketing at Pfizer and returned there, after two years as a management consultant at Bain, to take on leadership positions in corporate strategy, sales and as a commercial lead in Latin America. Ms. Massey holds a Bachelor of Economics from the University of Sydney and a Master of Business Administration from the New York University Stern School of Business.

“argenx has demonstrated incredible growth transitioning from a research and development organization to an integrated, immunology company. This success has been built on a commitment to creating patient value through innovation. I am very excited by the opportunity to continue to build on that foundation as we look to realize the full potential of VYVGART and the pipeline candidates within our commercial franchises,” commented Ms. Massey.

About argenx

argenx is a global immunology company committed to improving the lives of people suffering from severe autoimmune diseases. Partnering with leading academic researchers through its Immunology Innovation Program (IIP), argenx aims to translate immunology breakthroughs into a world-class portfolio of novel antibody-based medicines. argenx developed and is commercializing the first-and-only approved neonatal Fc receptor (FcRn) blocker in the U.S., the EU and Japan. For more information, visit www.argenx.com and follow us on LinkedIn, Twitter, and Instagram.

For further information, please contact:

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Forward-looking Statements

The contents of this announcement include statements that are, or may be deemed to be, “forward-looking statements.” These forward-looking statements can be identified by the use of forward-looking terminology, including the terms “believes,” “hope,” “estimates,” “anticipates,” “expects,” “intends,” “may,” “will,” or “should” and include statements argenx makes concerning the impact of the transition of the chief operating officer; argenx’s ability to bring innovative medicines to patients and the potential of VYVGART and the pipeline candidates within argenx’s commercial franchise. By their nature, forward-looking statements involve risks and uncertainties, and readers are cautioned that any such forward-looking statements are not guarantees of future performance. argenx’s actual results may differ materially from those predicted by the forward-looking statements as a result of various important factors. A further list and description of these risks, uncertainties and other risks can be found in argenx’s U.S. Securities and Exchange Commission (SEC) filings and reports, including in argenx’s most recent annual report on Form 20-F filed with the SEC as well as subsequent filings and reports filed by argenx with the SEC. Given these uncertainties, the reader is advised not to place any undue reliance on such forward-looking statements. These forward-looking statements speak only as of the date of publication of this document. argenx undertakes no obligation publicly update or revise the information in this press release, including any forward-looking statements, except as may be required by law.

Exhibit 99.2

argenx Reports Full Year 2022 Financial Results and Provides Fourth Quarter Business Update

$173 million in fourth quarter and $401 million in full year 2022 VYVGART® (efgartigimod alfa-fcab) global net product sales

FDA review ongoing for SC efgartigimod BLA with PDUFA target action date of June 20, 2023; MAA filed in Japan with approval decision expected by first quarter of 2024

ADHERE topline results continue to be expected in second quarter of 2023

Karen Massey appointed as Chief Operating Officer as part of planned transition; Keith Woods to retire and serve as advisor on Board of Directors

Management to host conference call today at 2:30 pm CET (8:30 am ET)

March 2, 2023

Amsterdam, the Netherlands – argenx (Euronext & Nasdaq: ARGX), a global immunology company committed to improving the lives of people suffering from severe autoimmune diseases, today reported financial results for the full year 2022 and provided a fourth quarter business update.

In a separate press release, argenx also announced today the appointment of Karen Massey to Chief Operating Officer as part of a planned transition. Ms. Massey will succeed Keith Woods who will remain at the Company through the launch of subcutaneous (SC) efgartigimod after which he will retire and serve as a strategic advisor on the argenx Board of Directors.

"We began 2023 from a position of strength following the successful first year of our VYVGART launch where we were able to reach more than 3,000 gMG patients globally with our transformative therapy. Our focus for the year ahead is expansion; both through upcoming regulatory approvals and launches, and more broadly by reaching additional patient segments with the planned launch of SC efgartigimod and through our ongoing stakeholder engagement efforts," said Tim Van Hauwermeiren, Chief Executive Officer of argenx. "Looking ahead, we have the opportunity this year to showcase the innovation within our pipeline through multiple data catalysts, including three pivotal data readouts from efgartigimod and the first clinical efficacy data from ARGX-117, our next pipeline-in-a-product candidate. We are well-equipped to build on this significant momentum as we advance our mission to redefine the treatment of autoimmune disease."

FOURTH QUARTER 2022 AND RECENT BUSINESS UPDATE

VYVGART Expansion

VYVGART is the first-and-only approved neonatal Fc receptor (FcRn) blocker in the U.S., Japan and the EU. argenx is planning for multi-dimensional expansion to reach more patients with VYVGART through additional regulatory approvals for generalized myasthenia gravis (gMG), the launch of SC efgartigimod for gMG, and new autoimmune indications with the VYVGART regulatory submission for immune thrombocytopenia (ITP) in Japan.

- Generated global net VYVGART revenues of $173 million in the fourth quarter of 2022 and $401 million in the full year of 2022
- Additional VYVGART regulatory approvals and commercial launches expected in 2023
    - Approval decision expected in Canada in third quarter of 2023 and in China and Israel by end of 2023
    - gMG launch in France, United Kingdom and Italy expected by end of 2023 following review of respective reimbursement dossiers
- Regulatory reviews of SC efgartigimod for gMG ongoing in the U.S., EU and Japan
    - Prescription Drug User Fee Act (PDUFA) target action date extended to June 20, 2023 following notification from the U.S. Food and Drug Administration (FDA) in January 2023
    - Marketing authorization application (MAA) filed in Japan in first quarter of 2023 with approval decision expected by first quarter of 2024
    - MAA review underway by European Medicines Agency with approval decision expected in fourth quarter of 2023
- Submission of MAA in Japan for VYVGART for the treatment of ITP expected mid-2023

Efgartigimod Research and Development

argenx aims to solidify its FcRn leadership by expanding the scope of IgG-mediated autoimmune diseases in development and further demonstrating the potential of FcRn blockade in ongoing clinical trials. By the end of 2023, efgartigimod is expected to be approved, in regulatory review or in development in 13 severe autoimmune diseases.

Multiple data readouts expected from ongoing efgartigimod trials in 2023 and 2024:

- ADHERE: Topline data in chronic inflammatory demyelinating polyneuropathy (CIDP) expected in second quarter of 2023
- ADDRESS: Topline data in pemphigus expected in second half of 2023
- ADVANCE-SC: Topline data from SC trial in ITP expected in second half of 2023
- BALLAD and ALKIVIA: Interim data in bullous pemphigoid expected in first half of 2024 and in myositis in second half of 2024
- ALPHA and RHO proof-of-concept (POC) trials through IQVIA collaboration: Topline data from ALPHA in post-COVID-19 postural orthostatic tachycardia syndrome (PC-POTS) expected in fourth quarter of 2023 and from RHO in primary Sjogren's syndrome expected in 2024
- POC trials underway in membranous nephropathy and lupus nephritis through Zai Lab collaboration

Clinical trials of efgartigimod in additional autoimmune indications to start this year:

- Registrational trial to start in thyroid eye disease in fourth quarter of 2023
- POC trials to start in ANCA-associated vasculitis (ANCA) and antibody mediated rejection (AMR) in kidney transplant in fourth quarter of 2023

Pipeline Progress

argenx is advancing a robust portfolio of innovative clinical programs, including ARGX-117 (C2 inhibitor) and ARGX-119 (muscle-specific kinase (MuSK) agonist). Both programs have the potential to be first-in-class opportunities for multiple severe autoimmune indications.

- ARDA: Interim data from POC trial of ARGX-117 in multifocal motor neuropathy expected mid-2023
- POC trial of ARGX-117 for prevention of delayed graft function after kidney transplantation expected to start in second half of 2023 following regulatory discussions
- Dermatomyositis selected as third autoimmune indication for development of ARGX-117
- Initiated Phase 1 dose-escalation trial of ARGX-119 in healthy volunteers; subsequent Phase 1b trial to assess early signal detection in patients

Continued investment in Immunology Innovation Program (IIP) to broaden autoimmune pipeline for sustained value creation opportunities

- argenx continues to invest in its discovery engine, the Immunology Innovation Program, to foster a robust innovation ecosystem and drive early-stage pipeline growth. argenx expects to nominate one new development candidate in 2023.

Steve Krognes appointed as non-executive director and Chair of the Audit and Compliance Committee of the Company's Board of Directors

FOURTH QUARTER AND FULL YEAR 2022 FINANCIAL RESULTS

ARGENX SE
UNAUDITED CONDENSED CONSOLIDATED STATEMENT OF PROFIT OR LOSS

	Three Months Ended December 31,		Twelve Months Ended December 31,	
(in thousands of $ except for shares and EPS)	**2022**	**2021**	**2022**	**2021**
Product net sales	$ 173,396	$ —	$ 400,720	$ —
Collaboration revenue	764	26,022	10,026	497,277
Other operating income	7,956	7,662	34,520	42,141
Total operating income	**182,116**	**33,684**	**445,267**	539,418
Cost of sales	(12,786)	—	(29,431)	—
Research and development expenses	(147,798)	(167,173)	(663,366)	(580,520)
Selling, general and administrative expenses	(135,287)	(97,422)	(472,132)	(307,644)
Loss from investment in joint venture	(677)	—	(677)	—
Total operating expenses	**(296,548)**	**(264,596)**	**(1,165,607)**	**(888,164)**
Operating loss	**$ (114,432)**	**$ (230,912)**	**$ (720,341)**	**$ (348,746)**
Financial income	13,925	1,199	27,665	3,633
Financial expense	(990)	(1,104)	(3,906)	(4,578)
Exchange gains/(losses)	60,259	(14,063)	(32,732)	(50,053)
Loss for the period before taxes	**$ (41,238)**	**$ (244,880)**	**$ (729,314)**	**$ (399,743)**
Income tax (expense)/benefit	$ 2,625	$ 7,062	$ 19,720	$ (8,522)
Loss for the period	**$ (38,613)**	**$ (237,818)**	**$ (709,594)**	**$ (408,265)**
Loss for the period attributable to:				
Owners of the parent	(38,613)	(237,818)	(709,594)	(408,265)
Weighted average number of shares outstanding	55,364,124	51,538,191	54,381,371	51,075,827
Basic loss per share (in $)	(0.70)	(4.61)	(13.05)	(7.99)
Diluted loss per share (in $)	(0.70)	(4.61)	(13.05)	(7.99)
Net increase/(decrease) in cash, cash equivalents and current financial assets compared to year-end 2021 and 2020			$ (144,180)	$ 340,276
Cash, cash equivalents and current financial assets at the end of the period			$ 2,192,548	$ 2,336,728

DETAILS OF THE FINANCIAL RESULTS

Total operating income for the fourth quarter and year-to-date in 2022 was $182.1 million and $445.3 million, respectively, compared to $33.7 million and $539.4 million for the same periods in 2021, and consists of:

- **Product net sales** from the sales of VYVGART for the three months ended December 31, 2022 were $173.4 million. The product net sales in the twelve months ended December 31, 2022 were $400.7 million. No product net sales were recognized during the same period in 2021.
- **Collaboration revenue** for the fourth quarter and year-to-date in 2022 was $0.8 million and $10 million, respectively, compared to $26 million and $497.3 million for the same periods in 2021. The collaboration revenue during the twelve months ended December 31, 2022 primarily relates to the recognition of milestone revenue following the option exercise by LEO Pharma to enter into a commercial license for ARGX-112. The collaboration revenue for the twelve months ended December 31, 2021 was primarily attributable to the recognition of the transaction price as a consequence of the termination of the collaboration agreement with Janssen, resulting in the recognition of $315.1 million, and the recognition of $177.5 million in collaboration revenue related to the strategic collaboration with Zai Lab, including the development milestone which was triggered by the FDA approval of VYVGART.
- **Other operating income** for the fourth quarter and year-to-date in 2022 was $7.9 million, and $34.5 million, respectively, compared to $7.7 million and $42.1 million for the same periods in 2021. During the twelve months ended December 31, 2022, and December 31, 2021, the fair value of the argenx profit share in AgomAb Therapeutics NV increased by $4.3 million and $11.2 million respectively. The increase is a result of the extension of a Series B financing round by AgomAb for which argenx maintains a profit share in exchange for granting the license for the use of HGF-mimetic antibodies from the SIMPLE Antibody™ platform.

Total operating expenses for the fourth quarter and year-to-date in 2022 were $296.5 million and $1,165.6 million, respectively, compared to $264.6 million and $888.2 million for the same periods in 2021, and consists of:

- **Cost of sales** for the fourth quarter and year-to-date in 2022 was $12.8 million and $29.4 million, respectively. The cost of sales was recognized with respect to the sale of VYVGART during 2022. There was no cost of sales recognized in the comparable prior year periods.
- **Research and development expenses** for the fourth quarter and year-to-date in 2022 were $147.8 million and $663.4 million, respectively, compared to $167.2 million and $580.5 million for the same periods in 2021. The research and development expenses mainly relate to external research and development expenses and personnel expenses incurred in the clinical development of efgartigimod in various indications and the expansion of other clinical and preclinical pipeline candidates. The increase in research and development expense during 2022 is mainly driven by the recognition of the priority review voucher submitted with the BLA filing for SC efgartigimod for the treatment of gMG, which resulted in an expense of $99.1 million.
- **Selling, general and administrative expenses** for the fourth quarter and year-to-date in 2022 were $135.3 million and $472.1 million, respectively, compared to $97.4 million and $307.6 million for the same periods in 2021. The selling, general and administrative expenses mainly relate to professional and marketing fees linked to the commercialization of VYVGART in the U.S., Japan and the EU and personnel expenses.

- **Loss from investment in joint venture** for the fourth quarter and year-to-date in 2022 was $0.7 million. The loss recognized was argenx's share of losses in Oncoverity, Inc. There was no losses from investment in joint venture in the comparable prior year periods.

Financial income for the fourth quarter and year-to-date in 2022 were $13.9 million and $27.7 million respectively, compared to $1.2 million and $3.6 million for the same periods in 2021. The increase in financial income is mainly due to an increase in interest income on current financial assets and cash and cash equivalents attributable to higher interest rates.

Exchange gains/losses for the fourth quarter and year-to-date in 2022 were $60.3 million of gains and $32.7 million of losses, respectively, compared to $14.1 million and $50.1 million of exchange losses for the same periods in 2021. Exchange gains/losses are mainly attributable to unrealized exchange rate losses on the cash, cash equivalents and current financial assets position in Euro.

Income tax for the fourth quarter and year-to-date in 2022 was $2.6 million and $19.7 million of tax benefit, respectively, compared to $7.1 million of tax benefit and $8.5 million of tax expense for the same periods in 2021. Tax benefit for the three months ended December 31, 2022 consists of $12.1 million of income tax expense and $14.7 million of deferred tax income, compared to $1.2 million of income tax expense and $8.2 million of deferred tax income for the same period in 2021.

Net loss for the fourth quarter and year-to-date in 2022 was $38.6 million and $709.6 million, respectively, compared to net loss of $237.8 and $408.3 million for the same periods in 2021.

Cash, cash equivalents and current financial assets totaled $2.2 billion as of December 31, 2022, compared to $2.3 billion as of December 31, 2021. Net change in Cash and cash equivalents and current financial assets is primarily a result of the closing of a global offering of shares, which resulted in the receipt of $761.0 million in net proceeds in March 2022, offset by net cash flows used in operating activities.

FINANCIAL GUIDANCE

Based on current plans to fund anticipated operating expenses, working capital and capital expenditures, argenx expects to utilize up to $500 million cash in 2023.

EXPECTED 2023 FINANCIAL CALENDAR:

- May 4, 2023: Q1 2023 financial results and business update
- July 27, 2023: HY 2023 financial results and business update
- October 27, 2023: Q3 2023 financial results and business update

CONFERENCE CALL DETAILS

The full year 2022 results and fourth quarter business update will be discussed during a conference call and webcast presentation today at 2:30 pm CET/8:30 am ET. A webcast of the live call may be accessed on the Investors section of the argenx website at argenx.com/investors. A replay of the webcast will be available on the argenx website.

Dial-in numbers:

Please dial in 15 minutes prior to the live call.

Belgium	32 800 50 201
France	33 800 943355
Netherlands	31 20 795 1090
United Kingdom	44 800 358 0970
United States	1 888 415 4250
Japan	81 3 4578 9752
Switzerland	41 43 210 11 32

About argenx

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For further information, please contact:

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Investors:
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Forward-looking Statements

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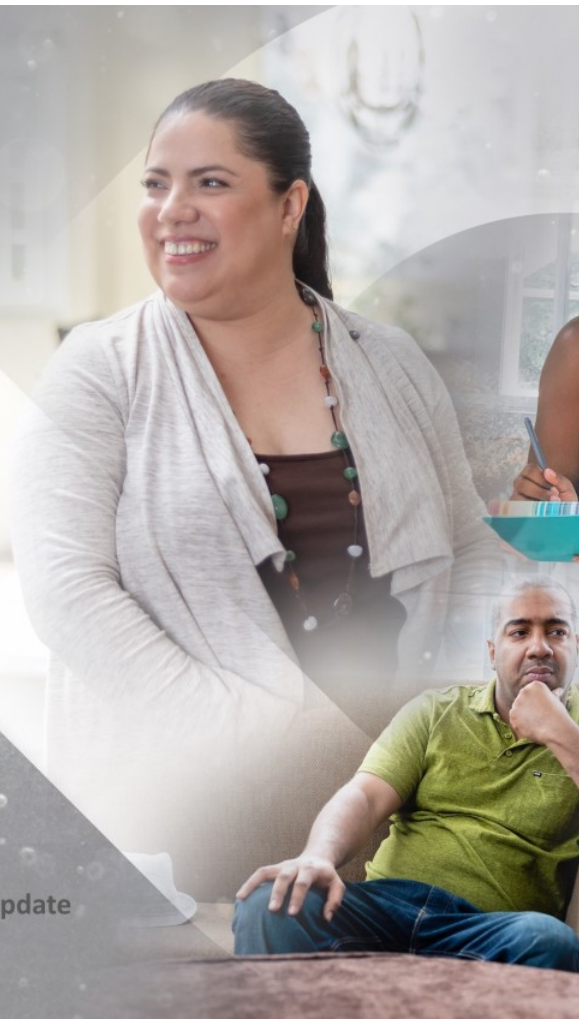
argenx

Together We Discover



Fourth Quarter and Full Year 2022 Financial Results and Business Update

March 2, 2023

Forward Looking Statements

This presentation has been prepared by argenx se ("argenx" or the "company") for informational purposes only and not for any other purpose. Nothing contai
presentation is, or should be construed as, a recommendation, promise or representation by the presenter or the company or any director, employee, agent, c
company. This presentation does not purport to be all-inclusive or to contain all of the information you may desire. Certain information contained in this prese
is based on studies, publications, surveys and other data obtained from third-party sources and our own internal estimates and research. While we believe th
studies, publications, surveys and other data to be reliable as of the date of this presentation, we have not independently verified, and make no representatio
adequacy, fairness, accuracy or completeness of, any information obtained from third-party sources. In addition, no independent source has evaluated the re
accuracy of our internal estimates or research and no reliance should be made on any information or statements made in this presentation relating to or base
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the date of publication of this document. argenx undertakes no obligation to publicly update or revise the information in this press release, including any forw
statements, except as may be required by law.

Entering 2023 in a Position of Strength

Driving **innovation** mission with an **entrepreneurial** spirit and commitment to **a strong culture**

Antibody Engineering Platform

8 Programs

demonstrated human proof-of-concept

Global VYVGART l

$401M

in product rever in first year

17

Autoimmune Di

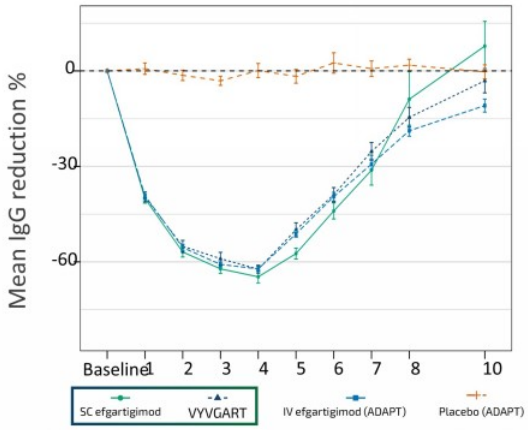
To be under evaluation efgartigimod and AR

argenx Today

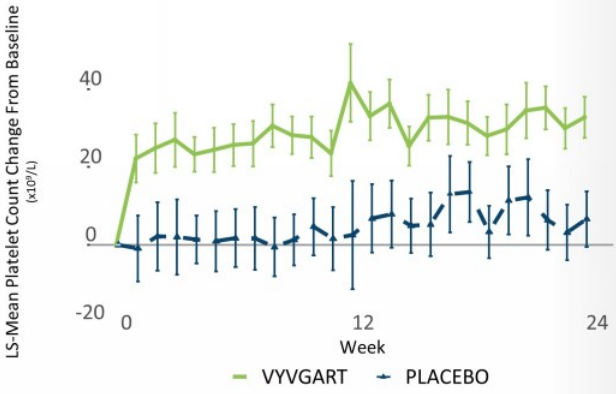
2022: Strengthened Efgartigimod Data Story

SC Noninferiority to IV

IgG reduction (%) in all ADAPT-SC and ADAPT participants



Clear Clinical Benefit in ITP



Broadened Sa

>1,300 clinical

Cyclic and ch

Cumulative
>1,000 pa

TEAEs consi
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Solidifying FcRn Leadership with Deep Repertoire of Preclinical and Translational Data

frontiers in Immunology

SOCIETY FOR INVESTIGATIVE DERMATOLOGY SID 1937

MDPI cells

ACR American College of Rheumatology

The Journal of Immunolo

2022: Broadened Scope of Efgartigimod Safety Database

Scope of Safety Database

>1,300 clinical study subjects

Cumulative exposure of **>1,000** patient years

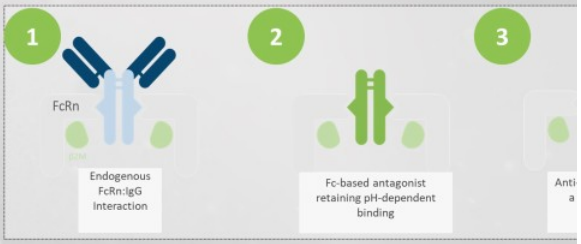
Different dosing regimens (up to 19 cycles of 4 weekly doses; up to 2 years of weekly dosing)

TEAEs consistent across **>4 indications;** typically mild to moderate

Low dis
rates d
effects
studies

Molecular design yields unique interaction with FcRn and differentiated safety profile

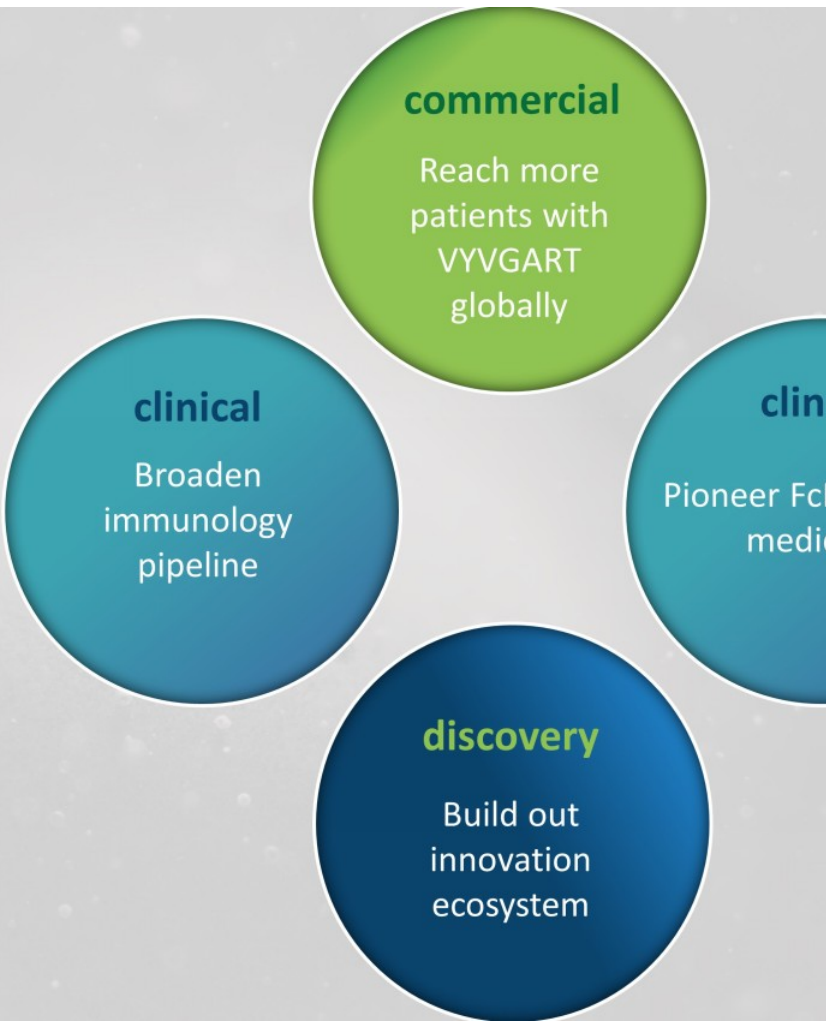
No reduction in albumin levels; no increase in lipid levels



Post-marketing data confirm positive benefit/risk profile as established in clinical trials

1. Ulrichts P, et al. *J Clin Invest*. 2018;128(10):4372-4386. doi: 10.1172/JCI97911.

2023:
Key Drivers of our Path to Profitability and ‘argenx 2025’



Pipeline Growth Driven By Immunology Innovation P

Internal Value Creation Strategy

FIRST IN CLASS | UNIQUE DESIGN | MULTIPLE INDICATIONS

Efgartigimod | ARGX-117 | ARGX-119

External Value Creation Strategy

ARGX-118 | Staten (ARGX-116) | AgoMAb (ARGX-114)

OncoVerity (Cusatuzumab) | ARGX-109 | LEO (ARGX-112)

Positioned for a Catalyst-Rich 2023

Commercial

- VYVGART gMG Approval in China
- VYVGART gMG Approval in Canada
- VYVGART gMG Launch in France, UK, Italy
- SC efgartigimod gMG Approval in US
- SC efgartigimod gMG Approval in EU
- SC efgartigimod gMG Submission in Japan
- VYVGART ITP Submission in Japan

Clinical

Efgartigimod

- ADHERE data in CIDP — **2Q 2023**
- ADDRESS data in Pemphigus — **2H 2023**
- ADVANCE (SC) data in ITP — **2H 2023**
- POC data in Post-COVID POTS — **4Q 2023**
- Initiate registrational trial in TED — **4Q 2023**
- Initiate POC studies in ANCA and AMR — **4Q 2023**

Additional pipeline

- ARGX-117: ARDA MMN interim results
- ARGX-117: Initiate DGF POC study
- ARGX-119: Initiate Phase 1 study

Quarterly Product Sales

2022 Vyvgart Net Sales

(in millions of $)	Q1	Q2	Q3	Q4	FY 22
US	21.2	73.2	124.1	159.1	377.6
Japan	-	1.5	6.0	8.3	15.8
Europe	-	-	0.6	5.1	5.7
Other*	-	0.1	0.6	0.9	1.6
Total	**21.2**	**74.8**	**131.3**	**173.4**	**400.7**

Full Year revenue of **$401M** in first year of launch

* The product net sales relate to sales made outside of US, Japan and Europe and relates to named patient sales made with the US label.

Fourth Quarter 2022 Financial Results

(in millions of $)	Three months ended December 31 2022	Three months ended December 31 2021	Twelve months ended December 31 2022	Twelve months ended December 31 2021
Product net sales	173.4	-	400.7	-
Collaboration revenue and other	8.7	33.7	44.6	539.4
Total operating income	**182.1**	**33.7**	**445.3**	**539.4**
Cost of sales	(12.8)	-	(29.4)	-
R&D expenses	(147.8)	(167.2)	(663.4)	(580.5)
SG&A expenses	(135.3)	(97.4)	(472.1)	(307.6)
Loss from investment in joint venture	(0.7)	-	(0.7)	-
Total operating expenses	**(296.5)**	**(264.6)**	**(1,165.6)**	**(888.2)**
Other (expenses) / income	75.8	(6.9)	10.7	(59.5)
(Loss) for the period	**(38.6)**	**(237.8)**	**(709.6)**	**(408.3)**

Cash of **$2.2B** at December 31 2022

Other income / (expenses) includes financial income / (expenses), exchange gains / (losses) and tax
Cash reflects cash, cash equivalents and current financial assets.

2022: Transformed into an Integrated Immunology Company

VYVGART launched in U.S., Japan & Germany

SUBMISSIONS IN **10+ COUNTRIES**

Built scalable global supply chain

LARGE SCALE CAPACITY TO **HANDLE DEMAND**

More than 3,000 patients on VYVGART®

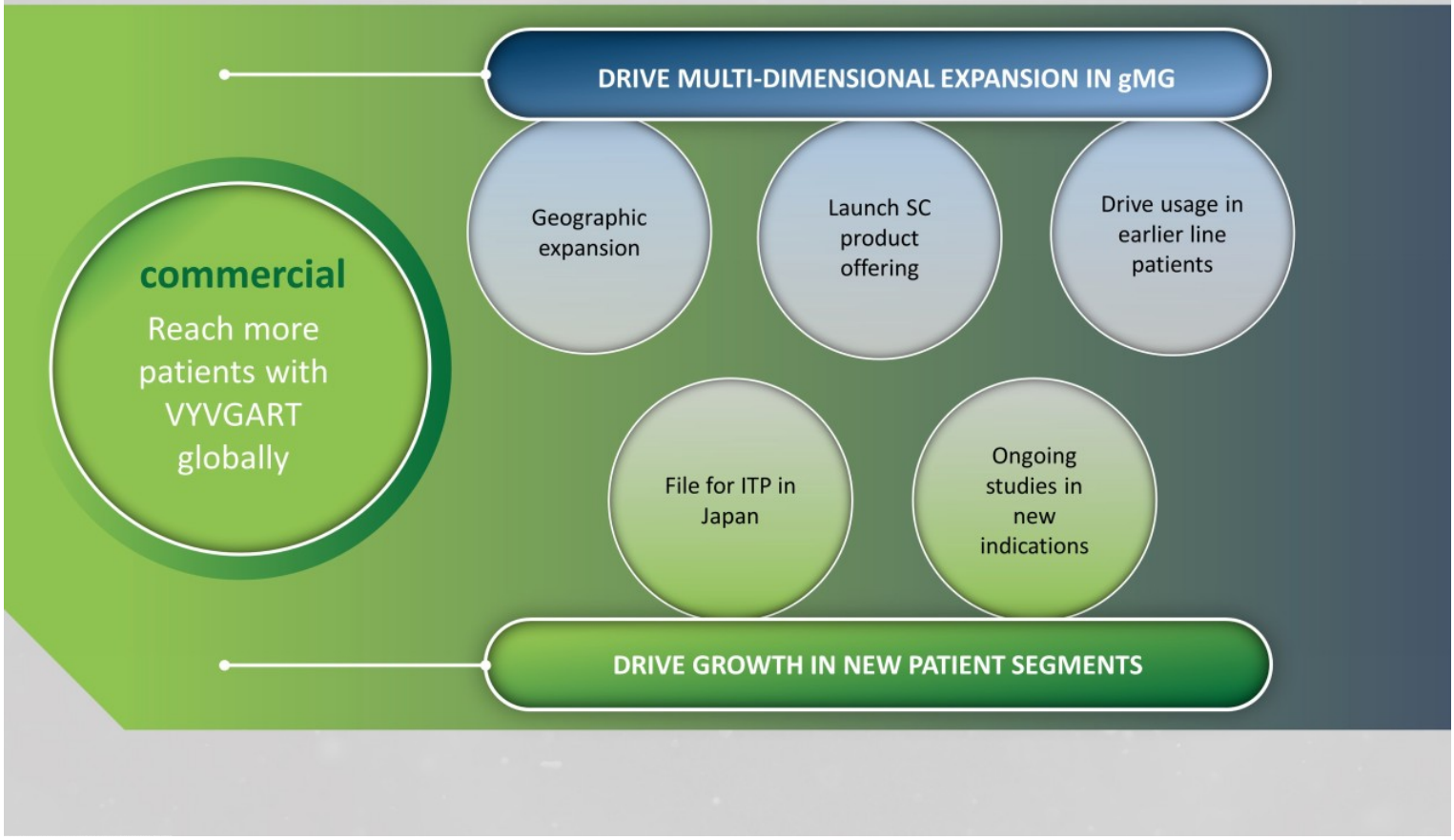
Rapid HC

90% of KI REACHE

Enabled a access thr based ag

90% US FAVO

Reach More Patients with VYVGART Globally




Our mission
continues...
Co-Creation
Empowerment
argenx
Innovation
Humility


...and gMG
is just the
beginning
Pioneer with
Our Science
Lead
Compass
our Pat
argenx 2025: A Leading, Sustai
Immunology Company
Drive Impact
Through
Innovation
Build
Compar
Want to
Fo