

10th of November 2025

9M 2025 Financial Results & Business Update

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Agenda

Carlos Gallardo, Chairman & CEO

9M 2025 Highlights

Biologics Growth Drivers Update: Ilumetri® & Ebglyss®

Karl Ziegelbauer, CSO

Pipeline Updates

Jon Garay, CFO

Financial Review

Carlos Gallardo, Chairman & CEO

Closing Remarks

9M 2025 Highlights





9M 2025

Solid operational and commercial execution in Europe

Sustaining strong growth momentum

Net Sales

€820.7 MM +12.8% YoY, driven by strong dermatology performance in Europe +24.5% YoY

EBITDA

€180.7 MM +27.1% YoY, growing in line with expectations

On track to meet 2025 Guidance Biologics growth and operational strength driving results

Leading EU products fueling performance

Ilumetri® (psoriasis)

Steady performance in 9M 2025. Net sales €170.9 MM +12.1% YoY

Ebglyss® (atopic dermatitis)

Strong 9M 2025 results as European markets scale after launching in key countries. 9M 2025 Net Sales €75.5 MM +3x YoY

Wynzora® (psoriasis)

Strong growth & capturing leading market share in key countries. Net sales €25.4 MM +32.3% YoY

Klisyri® (actinic keratosis)

Solid performance in Europe. Net sales €20.1 MM +22.6% YoY

Innovation pipeline status

Phase 1 & 2 studies

Anti-IL-1RAP mAb entered Phase 2 for Hidradenitis Suppurativa

Aim to commence the remaining proof-of-concept studies over the next 9–12 months

Efinaconazole (onychomycosis)

Approved in Germany in August 2025, following the EU decentralized procedure. Launch planning preparations underway



Biologics Growth Drivers Update

Ilumetri® & Ebglyss®

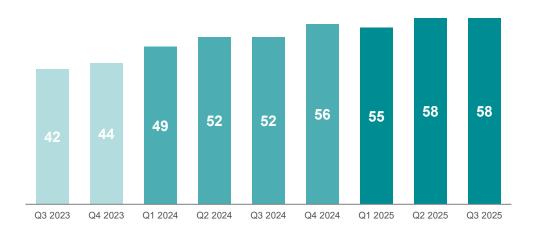




llumetri® highlights

POSITIVE study data strengthens Ilumetri's profile in the psoriasis space

Europe 9M 2025 Net Sales of €170.9 MM +12% YoY



* Source: IQVIA & ATU 2025

9M 2025 sales show steady double-digit growth YoY; on track to meet the >€300 MM peak target

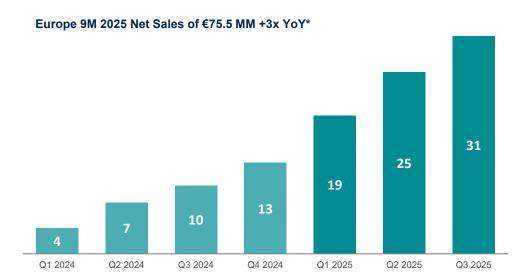
Anti-IL-23 remains the leading class in advanced psoriasis treatment*. The 200mg option adds dosing flexibility for llumetri's® patients, reinforcing its competitive profile

2-year POSITIVE study data, presented at EADV Paris 2025, underpins llumetri's® long-term value & real-world impact on patients' well-being



Ebglyss® highlights

Solid growth trajectory with key EU launches now completed



^{*}Growth figures may not correlate to published figures due to rounding

Strengthening our position in the AD segment: €75.5 MM sales in the 9M 2025 +3x YoY

Solid quarterly growth momentum as European markets scale after launch in most countries

Presented compelling real-world data at the 2025 EADV. Active in life cycle management & clinical collaboration to drive access & value

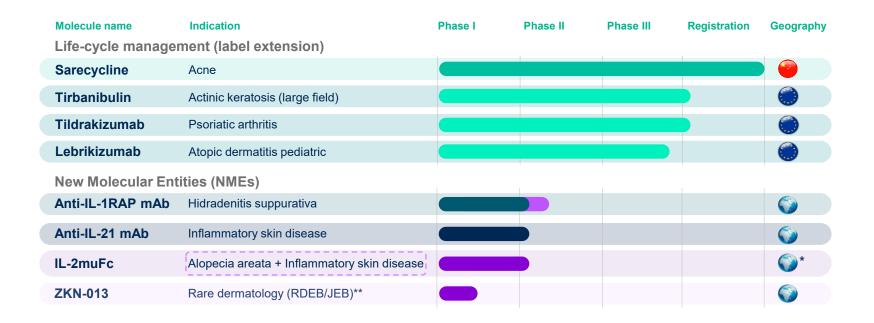


Pipeline Updates





A dynamic pipeline spanning early & late-stage programs





^{*} Worldwide ex-Greater China. ** RDEB / JEB - Recessive Dystrophic Epidermolysis Bullosa / Junctional Epidermolysis Bullosa

Financial Review





Consistent top-line growth & solid cash flow generation

9M 2025 Results

Highlights

Net Sales €820.7 MM +12.8% yearon-year, driven by strong momentum in European dermatology, on track with 2025 Guidance

Gross Margin moderating to **64.9%**, reflecting ongoing pressure linked to llumetri[®] royalties

EBITDA of €180.7 MM, +27.1% vs 9M 2024, driven primarily by incremental sales and lower SG&A weight in Q3 2025 YoY

R&D at €102.4 MM, 12.5% of Net Sales, in line with expectations

SG&A at €366.7 MM +6.2% vs 9M 2024, with lower growth in Q3 2025 YoY, expected to pick up in Q4

Net Debt of €18.6 MM:

Net Debt/EBITDA at 0.1x. Provides strategic flexibility to pursue potential inorganic growth opportunities

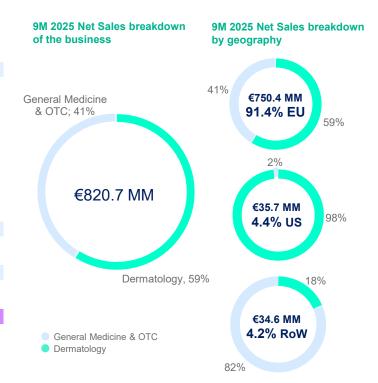


Net Sales Breakdown by Products

9M 2025 Results

Million€	9M 2025	9M 2024	% Chg YoY
Europe	750.4	647.2	15.9%
Dermatology	442.4	355.2	24.5%
General Medicine & OTC	308.0	292.0	5.5%
Ebastel franchise	44.3	45.6	(2.9%)
Crestor	31.0	32.9	(5.8%)
Sativex franchise	27.2	28.1	(3.2%)
Almax	26.8	25.0	7.2%
Parapres	15.0	15.3	(2.0%)
Eklira franchise	15.0	11.6	29.3%
Almogran franchise	13.0	12.6	3.2%
Others Europe*	135.7	120.9	12.2%
US	35.7	42.2	(15.4%)
Dermatology	35.1	41.3	(15.0%)
General Medicine	0.6	0.9	(33.3%)
RoW	34.6	38.2	(9.4%)
Dermatology	6.1	8.4	(27.4%)
General Medicine	28.5	29.8	(4.4%)
Net Sales	820.7	727.6	12.8%

^{*} Includes Algidol® divestment & Sekisan® out-licensing in Q1 2025, with a €12 MM upfront payment and net full year impact of approximately €15 MM vs 2024





Dermatology Sales Breakdown

9M 2025 Results

Million€	9M 2025	9M 2024	% Chg YoY
Europe	442.4	355.2	24.5%
llumetri	170.9	152.5	12.1%
Ebglyss	75.5	20.4	n.m.
Ciclopoli franchise	38.7	38.0	1.8%
Decoderm franchise	29.1	27.8	4.7%
Wynzora	25.4	19.2	32.3%
Solaraze	17.6	17.7	(0.6%)
Skilarence	15.8	15.1	4.6%
Klisyri	14.8	12.2	21.3%
Others Europe	54.6	52.3	4.4%
US	35.1	41.3	(15.0%)
Seysara	14.4	16.8	(14.3%)
Klisyri	5.3	4.2	26.2%
Others US	15.4	20.3	(24.1%)
RoW	6.1	8.4	(27.4%)
Total Almirall Derma	483.6	404.9	19.4%





Income Statement

9M 2025 Results

Million€	9M 2025	9M 2024	% Chg YoY
Total Revenues	825.6	731.2	12.9%
Net Sales	820.7	727.6	12.8%
Other Income	4.9	3.6	36.1%
Cost of Goods	(287.8)	(256.3)	12.3%
Gross Profit	532.9	471.3	13.1%
% of sales	64.9%	64.8%	
R&D	(102.4)	(90.1)	13.7%
% of sales	(12.5%)	(12.4%)	
SG&A	(366.7)	(345.2)	6.2%
% of sales	(44.7%)	(47.4%)	
SG&A w/o Amort. & Dep.	(269.7)	(256.2)	5.3%
% of sales	(32.9%)	(35.2%)	
SG&A Amort. & Dep.	(97.0)	(89.0)	9.0%
Other Op. Exp	(1.3)	(1.1)	18.2%
EBIT	67.4	38.5	75.1%
% of sales	8.2%	5.3%	
Amort. & Dep.	113.3	103.7	9.3%
% of sales	13.8%	14.3%	
EBITDA	180.7	142.2	27.1%
% of sales	22.0%	19.5%	
Gains on sale of assets	-	(1.4)	(100.0%)
Other costs	(1.2)	(0.5)	140.0%
Restructuring costs	(2.4)	-	n.m.
Impairment reversals / (losses)	-	(1.7)	(100.0%)
Net financial income / (expenses)	1.8	(5.3)	n.m.
Exchange rate differences	(0.5)	(1.0)	(50.0%)
Profit before tax	65.1	28.6	127.6%
Corporate income tax	(26.0)	(21.4)	21.5%
Net Income	39.1	7.2	n.m.
Normalized Net Income	41.8	9.9	n.m.

9M 2025 **Net Sales** were driven by strong Dermatology growth in Europe, led by Ilumetri® & Ebglyss®

R&D in 9M 2025 primarily driven by early & midstage clinical trials

SG&A increased in 9M 2025, reflecting ongoing investments in recent & upcoming Ebglyss[®] launches and expected to pick up in Q4 2025

9M 2025 **EBITDA** bolstered by European sales momentum and lower SG&A weight in Q3 2025 YoY

Net financial result improved, mainly because of the gains from the Equity Swap valuation linked to recent share price appreciation



Balance Sheet

9M 2025 Results

Million€	Sep 2025	Dec 2024	Var € MM
Goodwill & Intangible assets	1,217.2	1,296.5	(79.3)
Property, plant & equipment	156.9	153.8	3.1
Financial assets	22.6	16.4	6.2
Other non current assets	183.4	188.9	(5.5)
Total Non Current Assets	1,580.1	1,655.6	(75.5)
Inventories	189.4	171.8	17.6
Accounts receivable	160.1	151.4	8.7
Other current assets	51.3	40.8	10.5
Cash & cash equivalents	376.6	377.1	(0.5)
Total Current Assets	777.4	741.1	36.3
Total Assets	2,357.5	2,396.7	(39.2)
Shareholders Equity	1,475.1	1,488.4	(13.3)
Financial debt	337.1	347.4	(10.3)
Non current liabilities	239.9	221.9	18.0
Current liabilities	305.4	339.0	(33.6)
Total Equity & Liabilities	2,357.5	2,396.7	(39.2)
Net Debt Position			
Financial debt	337.1	347.4	(10.3)
Pension plans	58.1	58.6	(0.5)
Cash and cash equivalents	(376.6)	(377.1)	0.5
Net Debt / (Cash)	18.6	28.9	(10.3)

* EBITDA 12-month trailing

Goodwill & Intangible assets declined, primarily due to depreciation

Financial debt reflects the senior notes issued in September 2021, maturing in Q3 2026. The decline is mainly driven by EIB loan repayments

Strong liquidity & leverage, 0.1x Net Debt/EBITDA*



Cash Flow

9M 2025 Results

Million€	9M 2025	9M 2024
Profit Before Tax	65.1	28.6
Depreciation and amortization	113.3	103.7
Change in working capital	(16.1)	(11.5)
Other adjustments	16.2	11.3
CIT Cash Flow	(32.2)	(26.0)
Cash Flow from Operating Activities (I)	146.3	106.1
Interest Collections	5.1	5.9
Ordinary Capex	(55.5)	(45.9)
Investments	(59.6)	(96.0)
Divestments	5.9	12.1
Cash Flow from Investing Activities (II)	(104.1)	(123.9)
Interest Payment	(9.0)	(9.2)
Dividend Payment	(26.2)	(3.3)
Debt increase/(decrease) and Others	(14.6)	(14.4)
Other cash flows	7.1	(0.1)
Cash Flow from Financing Activities	(42.7)	(27.0)
Cash Flow generated during the period	(0.5)	(44.8)
Free Cash Flow (III) = (I) + (II)	42.2	(17.8)

Profit Before Tax improved significantly versus 9M 2024, in line with strong EBITDA growth

Working Capital increased, driven by higher receivables and elevated inventory levels for biologics

Investments include the Ilumetri® sales milestone, Anti-IL1-RAP mAb milestone, Wynzora® sales milestone and early-stage 2024 R&D milestones settled in H1 2025

Divestments include royalty inflows from the Covis agreement



Closing Remarks





Driving robust growth in biologics & progressing promising new pipeline opportunities to the next phase



Poised to lead in a rapidly expanding EU dermatology market



Building on a proven platform to drive sustainable growth



Turning strategy into results through disciplined execution





Appendices

Lebrikizumab

Life cycle management & clinical collaboration to drive access & value

	Indication	Title	Study	Objective
		ADlong	NCT05916365	Long-term safety up to 5 years
<u> </u>	Atopic	ADhope-1	NCT05990725	Charles to account of the still and a state of the
(almirall	Dermatitis	ADhope-2	NCT06526182	Study to assess 24-week effectiveness and safety
		ADTrust	NCT06815380	Observational study to assess impact on well-being and skin manifestations
		ADorable-1	NCT05559359	16-week efficacy & safety in pediatric patients
	Atopic ADorable-2 Dermatitis		NCT05735483	52-week long-term safety in pediatric patients
S:00.	Bomana	ADTouch	NCT06921759	Efficacy and safety in patients with atopic hand and foot dermatitis
any	PAR	PREPARED-1	NCT06339008	Efficacy & safety in adults with perennial allergic rhinitis
	CRSwNP	CONTRAST-NP	NCT06338995	Efficacy & safety in adults and adolescents with chronic rhinosinusitis and nasal polyps treated with intranasal corticosteroids



2025 EADV: Anti-IL1RAP mAb (LAD191)

Safety, pharmacokinetics & pharmacodynamics in adults with HS: results from part 3 of a Phase I study (link in image)

Safety, Pharmacokinetics, and Pharmacodynamics of LAD191, an IL-1RAP-Targeting Monoclonal Antibody, in Adults with Hidradenitis Suppurativa: Results from Part 3 of a Phase I Stude

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Submission Late Breaking Abstracts Deadline: August 21, 2025

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Maximum length: 4,000 characters (including spaces) - currently: 2,318 characters

³Almirall Ltd, Uxbridge, United Kingdom.

Introduction: LAD191 is a monoclonal antibody targeting the interleukin-1 receptor accessory protein (IL-1RAP). Parts 1 and 2 of this first-in-human (FiH) study evaluated single and multiple ascending doses of LAD191, respectively, in healthy volunteers (HV) and have demonstrated a favorable safety and tolerability profile, along with dose-proportional pharmacokinetics (PK). This Part 3 analysis aims to evaluate the safety, tolerability, PK, immunogenicity and pharmacodynamics of LAD191 among patients with hidradenitis suppurativa (HS).

Materials and Methods: This was a Phase I, randomized, 3-part, placebo-controlled study (NCT06488209). Part 3 included adult patients with HS (Hurley stage II or III). Patients were randomized to receive LAD191 or placebo, administered by subcutaneous injection once weekly for up to six doses.

Results: Five patients were randomized, of which three received LAD191 (mean age: 30.0 years; all females; Hurley stage I/IIII: 2/1 patients) and two received placeb (mean age: 31.0 years; all females; Hurley stage I/IIII: 2/10 patients). The mean International HS Severity Score (I/HS4) at baseline was 30.3 in the LAD191 arm and 13.5 in the placebo arm, with a higher number of severe patients (IHS4 ≥ 11) in the LAD191 arm compared to the placebo arm (3 vs. 1 patient). No treatment-emergent adverse events (IEAEs) were reported in the LAD191 arm vs. three TEAEs in the placebo arm (two moderate worsenings of HS and a mild acute pharyligits). No serious TEAEs or TEAEs leading to discontinuation were reported. Transient decreases in neutrophils count were observed in patients receiving LAD191, with spontaneous recovery noted in all cases. A trend toward lower LAD191 exposure (-25%) was observed in HS patients compared to what was previously observed in HV. The halt-life of LAD191 in HS patients was 17 days and no anti-drug artibodies were detected. An improvement in HS lesion count was observed following LAD191 treatment, accompanied by reductions in serum inflammatory biomarkers, such as IL-6 and lipocalin-2.

Conclusion: LAD191 was well tolerated and demonstrated a favorable safety and PK profile in patients with HS. Moreover, LAD191 has led to downstream cytokine reduction and early signs of clinical improvement in HS, supporting further clinical development.

Income Statement CER

9M 2025 Results

EURO	CER	Sep 2025
CZK	25.08	24.83
DKK	7.46	7.46
PLN	4.31	4.24
USD	1.08	1.11
CHF	0.96	0.94
GBP	0.85	0.85
NOK	11.59	11.71
SEK	11.41	11.10

Million€	9M 2025 CER	9M 2025	Var	9M 2024	% Var CER YoY	% Chg YoY
Total Revenues	826.1	825.6	(0.5)	731.2	13.0%	12.9%
Net Sales	821.2	820.7	(0.5)	727.6	12.9%	12.8%
Other Income	4.9	4.9	_	3.6	36.1%	36.1%
Cost of Goods	(288.5)	(287.8)	0.7	(256.3)	12.6%	12.3%
Gross Profit	532.7	532.9	0.2	471.3	13.0%	13.1%
% of sales	64.9%	64.9%		64.8%		
R&D	(102.3)	(102.4)	(0.1)	(90.1)	13.5%	13.7%
% of sales	(12.5%)	(12.5%)		(12.4%)		
SG&A	(367.7)	(366.7)	1.0	(345.2)	6.5%	6.2%
% of sales	(44.8%)	(44.7%)		(47.4%)		
SG&A w/o Amort. & Dep.	(270.2)	(269.7)	0.5	(256.2)	5.5%	5.3%
% of sales	(32.9%)	(32.9%)		(35.2%)		
SG&A Amort. & Dep.	(97.5)	(97.0)	0.5	(89.0)	9.6%	9.0%
Other Op. Exp	(1.3)	(1.3)	-	(1.1)	18.2%	18.2%
EBIT	66.3	67.4	1.1	38.5	72.2%	75.1%
% of sales	8.1%	8.2%		5.3%		
Amort. & Dep.	113.8	113.3	(0.5)	103.7	9.7%	9.3%
% of sales	13.9%	13.8%		14.3%		
EBITDA	180.1	180.7	0.6	142.2	26.7%	27.1%
% of sales	21.9%	22.0%		19.5%		
Gains on sale of assets	-	-	-	(1.4)	(100.0%)	(100.0%)
Other costs	(1.1)	(1.2)	(0.1)	(0.5)	120.0%	140.0%
Restructuring costs	(2.4)	(2.4)	-	_	n.m.	n.m.
Impairment reversals / (losses)	-	-	_	(1.7)	(100.0%)	(100.0%)
Net financial income / (expenses)	1.8	1.8	-	(5.3)	n.m.	n.m.
Exchange rate differences	(0.5)	(0.5)	_	(1.0)	(50.0%)	(50.0%)
Profit before tax	64.1	65.1	1.0	28.6	124.1%	127.6%
Corporate income tax	(26.0)	(26.0)	-	(21.4)	21.5%	21.5%
Net Income	38.1	39.1	1.0	7.2	n.m.	n.m.
Normalized Net Income	40.7	41.8	1.1	9.9	n.m.	n.m.



Leading Product Net Sales

9M 2025 Results

Million€	9M 2025	9M 2024	% Chg YoY
llumetri [®]	170.9	152.5	12.1%
Ebglyss [®]	75.5	20.4	n.m.
Ebastel® franchise	54.7	57.5	(4.9%)
Ciclopoli® franchise	40.2	40.2	-
Almax [®]	31.5	31.6	(0.3%)
Crestor®	31.0	32.9	(5.8%)
Decoderm® franchise	29.4	28.2	4.3%
Sativex® franchise	27.2	28.1	(3.2%)
Wynzora [®]	25.4	19.2	32.3%
Klisyri [®]	20.1	16.4	22.6%
Rest of the products	314.8	300.6	4.7%
Net Sales	820.7	727.6	12.8%



Reconciliations with financial statements

Gross Margin & EBITDA

Million €	9M 2025	9M 2024
Net Sales ⁽¹⁾	820.7	727.6
Procurements ⁽¹⁾	(191.4)	(175.4)
Other manufacturing costs ⁽²⁾		
Staff costs	(31.5)	(29.1)
Amortization & Depreciation	(9.7)	(8.4)
Other operating costs	(18.5)	(18.2)
Royalties ⁽²⁾	(41.2)	(28.7)
Others ⁽²⁾	4.5	3.5
Gross Profit	532.9	471.3
As % of Revenues	64.9%	64.8%

Million €	9M 2025 9I	M 2024
Operating Profit	63.9	34.9
Directly traceable with annual accounts		
Amortization & Depreciation	113.3	103.7
Net gain (loss) on asset disposals	_	3.2
Non directly traceable with annual accounts		
Staff costs	2.4	-
Other gain / (Loss) from operating expense	es 1.1	0.4
EBITDA	180.7	142.2

⁽¹⁾ As per Annual Account Terminology.



⁽²⁾ Data included in the corresponding caption of the profit and loss account.

Reconciliations with audited financial statements

EBIT & Net Financial income/(expenses)

Million€	9M 2025	9M 2024
EBITDA	180.7	142.2
Amortization & Depreciation	(113.3)	(103.7)
EBIT	67.4	38.5
Financial income	5.5	5.5
Financial cost	(11.9)	(11.6)
Financial derivative	8.2	0.8
Net Financial income / (expenses)	1.8	(5.3)



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