GALDERMA

EST. 1981

Galderma:
The pure-play dermatology
category leader

Emil Ivanov

Head of Strategy, Investor Relations and ESG

NOVEMBER 2025



Disclaimer

THIS PRESENTATION AND ITS CONTENTS ARE NOT FOR RELEASE, PUBLICATION OR DISTRIBUTION, IN WHOLE OR IN PART, DIRECTLY OR INDIRECTLY, IN OR INTO OR FROM THE UNITED STATES OF AMERICA, CANADA, AUSTRALIA, JAPAN OR ANY JURISDICTION WHERE SUCH DISTRIBUTION IS UNLAWFUL.

This presentation (the "Presentation") has been prepared by Galderma Group AG ("we", "us") solely for informational purposes and has not been independently verified and no representation or warranty, express or implied, is made or given by or on behalf of us. We reserve the right to amend, update, revise, correct, complete or replace the Presentation at any time without notice, and we and the respective directors, officers, employees, advisors undertake no obligation to provide the recipients with access to any additional information. We are not obligated to update or correct the information set forth in this Presentation. Nothing in this Presentation is, or should be relied upon as, a promise or representation as to the future.

This Presentation must not be distributed, reproduced or used without our consent. No decision has been made on whether to proceed with the transaction contemplated in this Presentation. If such decision is made, any investment decision must be made exclusively on the basis of a prospectus that will supersede the information contained in this Presentation.

This Presentation does not constitute or form part of, and should not be construed as, an offer or invitation or inducement to subscribe for, underwrite or otherwise acquire, any securities of us, nor should it or any part of it form the basis of, or be relied on in connection with, any contract to purchase or subscribe for any securities of us, nor shall it or any part of it form the basis of, or be relied on in connection with, any contract or commitment whatsoever.

This document is (i) not a prospectus within the meaning of the Swiss Financial Service Act and (ii) not a prospectus under any other applicable laws, or offering document, and has not been reviewed or approved by any regulatory or supervisory authority. This Presentation is being made available to you solely for your information and background and is not to be used as a basis for an investment decision in our securities.

Certain statements in this Presentation are forward-looking statements. Forward-looking statements are statements that are not historical facts and may be identified by words such as "plans", "targets", "aims", "believes", "expects", "anticipates", intends", "estimates", "will", "may"," continues", "should" and similar expressions. Such statements are based on management's current intentions, expectations or beliefs and involve inherent risks, assumptions and uncertainties, including factors that could delay, divert or change any of them. By their nature, forward-looking statements involve a number of risks, uncertainties and assumptions that could cause actual results or events to differ materially from those expressed or implied by the forward-looking statements. These risks, uncertainties and assumptions could adversely affect the outcome and financial consequences of the plans and events described herein. Actual results may differ from those set forth in the forward-looking statements as a result of various factors (including, but not limited to, future global economic conditions, changed market conditions, intense competition in the markets in which we operate, costs of compliance with applicable laws, regulations and standards, diverse political, legal, economic and other conditions affecting our markets, and other factors beyond our control). Neither we nor any of our respective directors, officers, employees, advisors, or any other person is under any obligation to update or revise any forward-looking statements, whether as a result of new information, future events or otherwise.

This Presentation is intended to provide a general overview of our business and does not purport to be comprehensive. This Presentation is furnished solely for your information, should not be treated as giving investment advice and may not be printed or otherwise copied or distributed. The information contained in this Presentation is not to be viewed from nor for publication or distribution in nor taken or transmitted into the United States of America ("United States"), Australia, Canada or Japan and does not constitute an offer of securities for sale in any of these jurisdictions. The securities offered by us have not been, and will not be, registered under the U S Securities Act of 1933 as amended, or the securities laws of any state or other jurisdiction of the United States and such securities may not be offered or sold within the United States.

This Presentation does not constitute investment, legal, accounting, regulatory, taxation or other advice. By attending this Presentation and/or by accepting this Presentation and/or any related materials, you agree to be bound by the conditions and restrictions set out herein.

Galderma as a 'self-care' category leader in dermatology

Dermatological science & strong consumer heritage in dermatology



Only scaled company fully dedicated to dermatology spanning 3 of the most attractive segments in Dermatology



Global integrated commercial platform with presence in over 90 countries



Consumer-centric business with digitally-enabled execution

1947 year of invention of Cetaphil

770+
clinical trials
since 2019

130K+
Training participants
per year



Pharmacies/ Physician Proprietary
DTC¹ E-commerce Retailers Drugstores practices training events

Science-based endorsements & digital/ tech-enabled tools

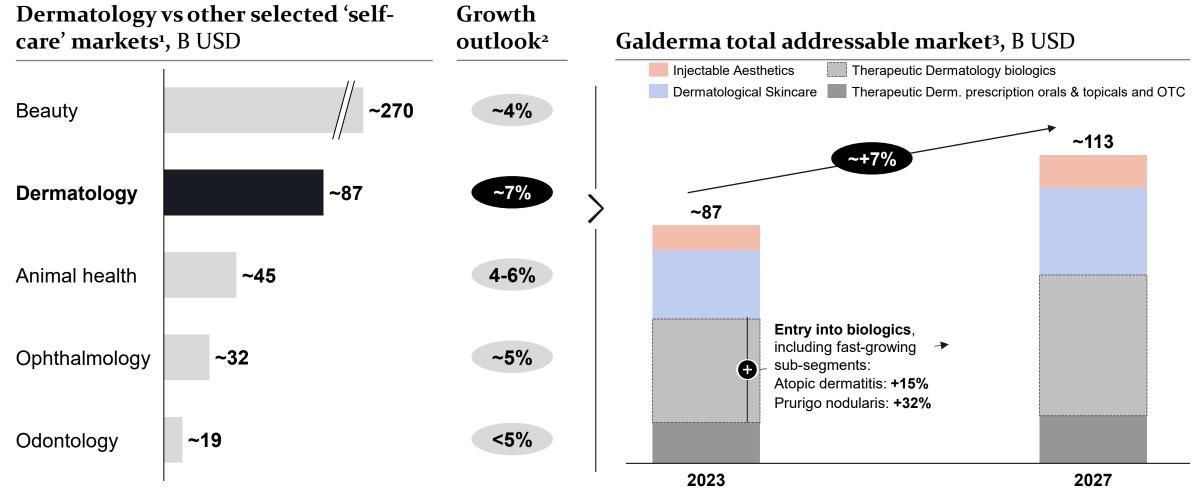




Loyalty programs & consumer-focused services

Consumer-driven decision making: ~90% of Galderma sales: self-pay

Fully focused on the fastest growing 'self-care' market and on attractive, high-growth segments, boosted by biologic entry



^{1.} Beauty – L'Oréal Annual Report (2022), Animal Health – Zoetis Investor Day Presentation (2023), Ophthalmology – Alcon Capital Markets Day (2023), Odontology – Straumann H12023 Earnings Presentation (2023) and Straumann Annual Report (2020) | 2. Mid-term growth outlook, may vary by sources – beauty market growth-outlook based on last 10 years average growth rate of L'Oréal beauty market | 3. Based on Galderma analyses using for Injectable Aesthetics: Medical Insights, The Global Aesthetic Market Study (Jan. 2024), Clarivate, EY Aesthetic Market Analysis – Dermal Fillers and Evaluate Pharma (Jan. 2023); for Dermatological Skincare: All numbers at Retail Selling Price, Internal Galderma database (TABS 2023), Nicholas Hall DB6 database and Euromonitor Beauty and Personal Care 2023 edition; for Therapeutic Dermatology: Numbers for prescription orals and topicals at Manufacturer Level Price, numbers for biologics at Public Price, IQVIA Analytics Link Disease Module (using moving annual total numbers as of Q3 2023 and gross to net ratio of 20%), Evaluate Pharma, Nicholas Hall DB6, Clarivate and Euromonitor Beauty and Personal Care 2023 edition, includes biologics and other molecules covering all modalities and modes of administration for atopic dermatitis (AD), prurigo nodularis (PN) and psoriasis (PSO)

Leading positions in dermatology worldwide

Injectable Aesthetics

Dermatological Skincare

Therapeutic Dermatology

Dysport. Alluzience

relfydess™

#2 global neuromodulator

Spearheading next generation neuromodulation





#2 global filler

The world's most diverse range of HA fillers & skinboosters



#1 global biostimulator

The first & original collagen stimulator with PLLA-SCA1



9/10 U.S. derms

recommend

Over 75 years of heritage dedicated to sensitive skin



#1 peri-procedure skincare

Preferred U.S. physician-dispensed peri-procedure brand



#1 across² acne, rosacea, & actinic keratosis

Leading portfolio in Therapeutic Derm. orals & topicals



1St

IL-31RA³ in the market

Differentiated in prurigo nodularis & atopic dermatitis













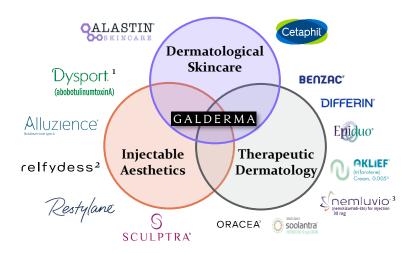




Uniquely positioned with its integrated dermatology platform

Broadest portfolio with leading science & innovation

Broadest dermatology portfolio of clinically-proven flagship brands to meet consumers' & patients' needs:



Leading science & innovation driving differentiation & long-term sustainable growth

Global scale with omni-channel execution excellence

Global commercial presence with notable headroom for high growth through continued penetration in fast-growing geographies

Scaled omni-channel strategy covering the whole spectrum:



Market-leading education & services

Broad education, training, and medical awareness activities

>225,000 healthcare participants reached⁴

GAIN >10,000 events per year

Differentiated value-adding platforms

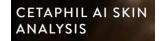
ASPIRE_{GALDERMA REWARDS} >4.3 M consumers in the U.S. loyalty program









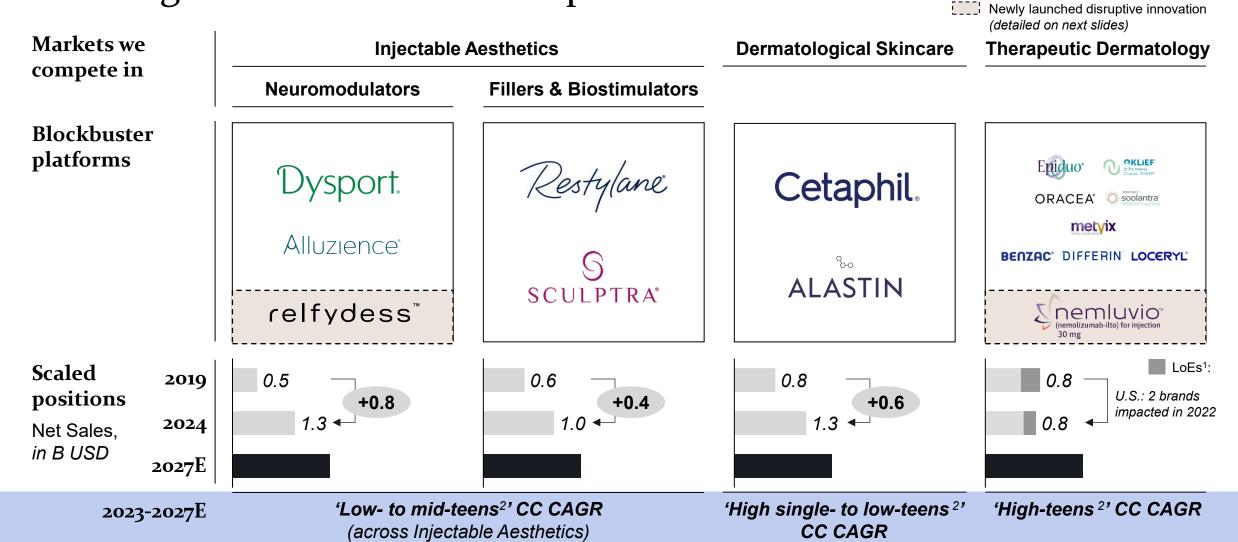




^{1.} Marketed under the brand name of Azzalure for aesthetic use in the European region and Dysport in the rest of the world for aesthetic indications – applies throughout the document | 2. relabotulinumtoxinA, previously referred to as QM-1114 |

^{3.} nemolizumab-ilto | 4. Single contact through medical education or awareness activities – one healthcare professional can attend more than one training

Mid-term growth: Attractive outlook for each blockbuster platform – adding a fourth multi-billion platform with Nemluvio



^{1.} LoE: Loss of Exclusivity | 2. 'Teens' defined as numbers greater than 10% and lower than 20%

Relfydess®: Next-generation neuromodulator with blockbuster potential

Launched for frown lines & crow's feet in first International markets

starting with Germany and Spain in Q4 2024, with a progressive roll-out planned in 18 countries approved¹

Highly differentiated neuromodulator



Sustained results

Up to 75% of patients maintaining improvements through 6 months



Fast onset

Up to 39% of patients seeing effects from Day 1



Designed for aesthetic use

Introducing a ready-to-use liquid neuromodulator² optimized for simple volumetric dosing



Neuromodulators: Deepening and expanding portfolio reach

relfydess

17 markets launched

New approvals in Q3: Hong Kong, New Zealand, & UAE – just in time for AMWC¹ Dubai

Neuromodulators

H490 bps Market share gains in Europe over 12 months where Relfydess is launched

Double-digit solution of Pouble-digit growth Service S



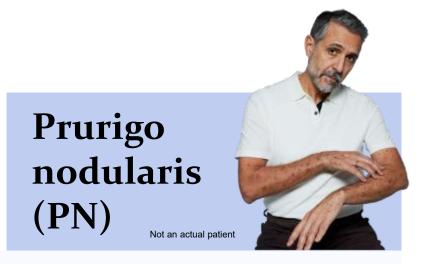






^{1.} AMWC: Aesthetic & anti-aging Medicine World Congress | 2. Commercialized as Azzalure in Europe

Nemluvio: Entering attractive markets, prurigo nodularis and atopic dermatitis, characterized by persistent itch



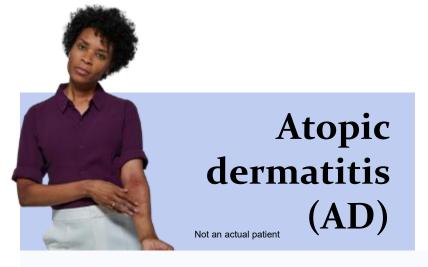
~2 B USD mid-term global market size, +32% CAGR¹

88% rate itch as the worst symptom²



Lower prevalence

Estimated 200--300~K U.S. patient population



 $\begin{array}{c|c} \sim & 20 \; B \; USD \; \text{mid-term} \\ \text{global market size, +15\% CAGR}^1 & \text{are seeking} \end{array}$

87% are seeking freedom from itch³



Higher prevalence

Estimated >20 M U.S. patient population

High unmet patient needs, with few approved biologics and patients still inadequately treated

1. 2023-2027 CAGR | 2. Rodriguez D et al JAMA Dermatol 2023: Patient Perspectives on Living With Severe Prurigo Nodularis | 3. Augustin M 2022 Real-World Treatment Patterns and Treatment Benefits among Adult Patients with Atopic Dermatitis: Results from the Atopic Dermatitis Patient Satisfaction and Unmet Need Survey

GALDERMA

Nemluvio: Fast and safe itch relief with skin healing that lasts, underpinning guidance to above 2 B USD in peak sales



The first and only biologic to directly target IL-3IRA

NEMLUVIO blocks the signaling that drives itch, inflammation, skin barrier dysfunction, and fibrosis

Differentiated profile



FAST ITCH RELIEF as soon as Day 2



LASTING SKIN CLEARANCE through 1 year



FAVOURABLE SAFETY PROFILE to prescribe with confidence



Q4W¹ DOSING FROM THE START for added convenience

>2 B USD peak sales, beyond the mid-term period

PN: Well-positioned to be the **preferred treatment** choice (1L)

AD: Expected to be the 2nd largest product, especially for itch-dominant & refractory patients on IL-4/ IL-13 treatments

PN & AD launches









+ New indications²: Galderma is investigating (Phase 2) nemolizumab in Systemic Sclerosis and Chronic pruritus of unknown origin (CPUO) with potential for other dermatological indications

^{1.} Every 4 weeks, possibility of Q8W (every 8 weeks, just 6 doses/ year) after the first 16 weeks of NEMLUVIO treatment in AD | 2. Not included in the peak sales guidance

Nemluvio: U.S. on a strong launch trajectory

~37⁰/₀ in prurigo nodularis

~7.3% in atopic dermatitis

Majority

share of patients new to advanced therapies ("biologic naïve")

Percentage of patients covered through commercial plans² in both indications as 1st line biologic treatment

>8o%

Before nemolizumab







After 8 weeks on nemolizumab

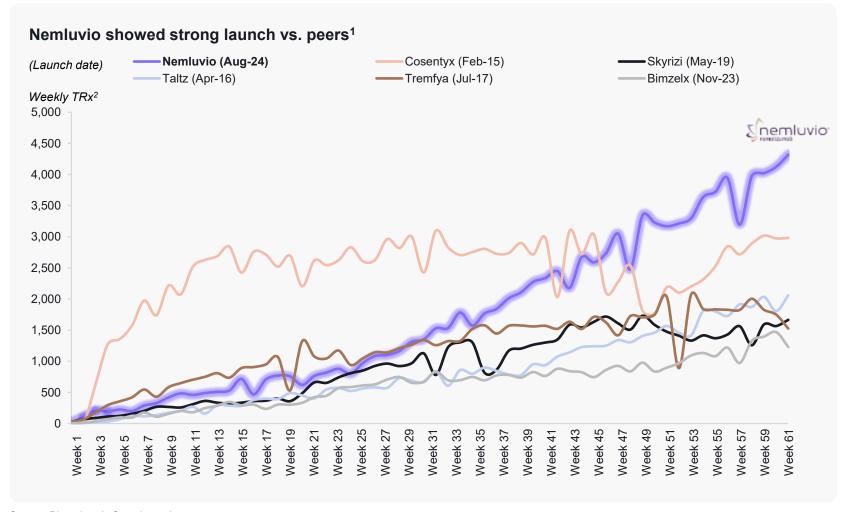






^{1.} NBRx: New-to-brand prescriptions; rolling 6 week average as of the week ending October 10, 2025 | 2. MMIT Commercial covered lives, as of October 10, 2025 | Source: IQVIA; LAAD; MMIT; Company estimates

Nemluvio: Launch exceeding forecasts, indicating strong uptake



Source: Bloomberg's Symphony data

Nemluvio offers a differentiated clinical profile, providing rapid and significant itch relief – onset as early as week 1 vs. week 4 for key competitors

The product's well-tolerated safety profile and convenient dosing schedule (Q4W, with Q8W for maintenance in atopic dermatitis) offer advantages over other biologics in the market

Nemluvio's launch continues gaining **significant traction**, with **TRx rising c. 5x** vs. beginning of 2025

^{1.} Weekly TRx data of Nemluvio and its peers since launch; Week 61 as of week ending October 3rd, 2025; | 2. Total prescriptions

Record financial performance in Q3 2025

Q₃ YTD 2025 ACTUALS

3,737Net sales in M USD

Record net sales, ahead of expectations, with focused execution of both, the existing portfolio and new innovation; growth predominantly volume-based

NET SALES GROWTH

Q3 YTD: +15.0% Q3: +21.0% constant currency year-on-year growth

Widespread growth with Q3 acceleration
 across all product categories and geographies,
 especially Nemluvio, while Neuromodulators
 benefitted from favorable phasing

2025 FULL YEAR GUIDANCE **+17.0-17.7%** Net sales growth **23.1-23.6%** Core EBITDA margin

at constant currency Raising full-year guidance on net sales
(previously +12-14%) and Core EBITDA margin
(previously approximately 23%) on strong
growth trajectory, especially of Nemluvio

^{1.} Constant currency year-on-year growth means the annual growth rate of net sales, excluding the impact of exchange rates movements and excluding hyperinflation economies. The impact of changes in foreign exchange rates are excluded by translating all reported revenues during the 2 periods at average exchange rates in effect during the previous year – applies throughout the document

Continued strong net sales growth momentum across the portfolio

Q3 YTD 2025 constant currency year-on-year growth

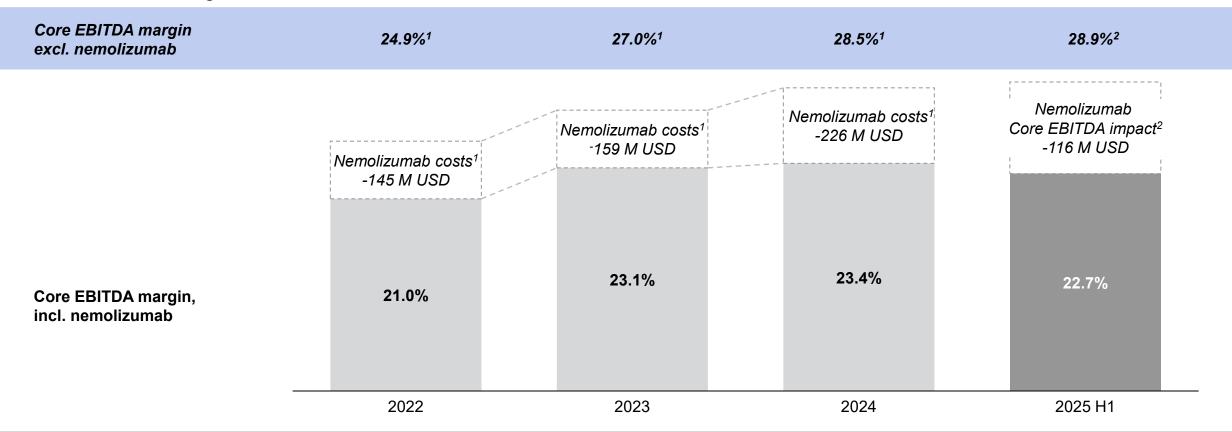
| +10.5% | | DERMATOLOGICAL SKINCARE +8.2% | THERAPEUTIC DERMATOLOGY +40.4% | GALDERMA +15.0% |
|---|-----------|-------------------------------|---|-------------------------|
| | | | | |
| Dysport. (abobotulinumtoxinA) Azzalure* | Restylane | Cetaphil. | (nemolizumab-ilto) for injection 30 mg | +13.2% International |
| relfydess Alluzience* | SCULPTRA® | ALASTIN | ORACEA° o sociantra metyix BENZAC° DIFFERIN' LOCERYL' | +17.5% u.s. |

GALDERMA

Improvement in underlying profitability while investing in nemolizumab

Core EBITDA margin evolution

ILLUSTRATIVE ONLY - BAR SIZE NOT AT SCALE



Core EBITDA Margin 2023 – 2027E

+300 – 500bps Core EBITDA margin expansion (vs. 2023) by 2027E

Incl. nemolizumab

majority of which delivered in 2026 and 2027

16

^{1.} Nemolizumab costs include external R&D, Medical and Regulatory, Sales and Marketing, and Distribution | 2. Core EBITDA margin excluding nemolizumab Core EBITDA impact, i.e., gross profit and estimates on the allocation of external R&D, Medical and Regulatory, Sales and Marketing, and Distribution OpEx costs

GALDERMA

Currently effective U.S. tariffs factored in, while ramping-up U.S. investments

Current effective U.S. tariff treatment by product (as of October 22, 2025)

| Injectable Aesthetics | Dysport | ✓ Exempt : classified as 'pharmaceutical'; produced in the U.K. | | |
|----------------------------|--------------|--|--|--|
| | Restylane | × Non-exempt: 15% tariff on import value; produced in the EU | | |
| | Sculptra | × Non-exempt: 15% tariff on import value; produced in the EU | | |
| Dermatological Skincare | Cetaphil | ✓ Exempt : USMCA¹ compliant production in Canada | | |
| | Alastin | ✓ Exempt : produced in the U.S. | | |
| Therapeutic Dermatology | Nemluvio | ✓ Exempt : classified as 'pharmaceutical': manufacturing process is completed in the U.S. | | |
| | Differin | ✓ Exempt : USMCA¹ compliant production in Canada | | |
| | Other brands | ✓ Exempt : classified as 'pharmaceutical', predominantly in USMCA¹ compliant production in Canada | | |

Galderma continuing to expand its U.S. operations

Strong U.S. presence, including new U.S. headquarters, Miami

Significant healthcare professional education & training activities

Ramp-up of local manufacturing: Commitment to >650 M USD in U.S. production spend via contract manufacturing partners for Nemluvio (final assembly & packaging) and Alastin & select Cetaphil SKUs

Additional technology transfers initiated (via partners), including double-sourcing Relfydess U.S. capacity

^{1.} USMCA: United States-Mexico-Canada Agreement

Capital allocation priorities focused on organic growth and deleveraging

| | 2024 Actuals | 2025E | Mid-term |
|--------------------------------------|--------------|--|---|
| Non-core adjustments ¹ | 93 M USD | ~ 60 M USD | |
| Effective tax rate ² | 25.5% | 23 - 25% | ~ 20% |
| Core CAPEX | 3.3% | ~3% of Net sales | Low to mid-single digit as % of Net Sales |
| Leverage | 2.3x | | Targeting <2x for the mid-term |
| Net financial expenses ³ | 328 M USD | ~ 200 - 210 M USD | |
| Milestone and earnout payments | 176 M USD | 23 M USD | |
| Dividends ⁴ | ~17% | ◆ Ordinary dividend payout target of up to 20% → | |

^{1.} Includes assumptions for other income and expenses related to tangible asset impairments, ongoing litigation and onerous items, restructuring charges and others, excluding M&A fees | 2. On reported profit before tax |

^{3.} Includes interest income and interest expense, excluding FX impact | 4. Of reported net income based on prior year results, subject to Board and AGM approval

Continued strong trajectory, with focused execution of the existing portfolio and new innovation

Record net sales of 3,737 M USD for the first 9 months 0f 2025, growing +15.0% at constant currency, predominantly driven by volume

Widespread growth in Q3 2025 with acceleration across all product categories and geographies – especially Nemluvio in the U.S., while Injectable Aesthetics Neuromodulators benefited from some favorable phasing

Currently effective U.S. tariffs are factored in 2025 guidance, while Galderma continues to expand its U.S. operations

Raising full-year guidance on net sales to +17.0-17.7% year-on-year growth at constant currency (previously +12-14%), and on Core EBITDA margin to 23.1-23.6% at constant currency (previously approximately 23%), on a strong growth trajectory, especially of Nemluvio



GALDERMA

EST. 1981