A Leading Specialty Vaccine Company

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September 2026

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This presentation presents information about investigational vaccine candidates that have not been approved for use and have not been determined by any regulatory authority to be safe or effective.

Management uses and presents IFRS results, as well as the non-IFRS measure of Adjusted EBITDA to evaluate and communicate its performance. While non-IFRS measures should not be construed as alternatives to IFRS measures, management believes non-IFRS measures are useful to further understand Valneva's current performance, performance trends, and financial condition. Adjusted EBITDA is a supplemental measure of performance used by investors and financial analysts. Management believes this measure provides additional analytical tools.

A Leading Specialty Vaccine Company: Focused on vaccines that make a difference

We **develop, manufacture, and commercialize** prophylactic vaccines for infectious diseases addressing unmet medical needs

Cash-Generating Specialty Vaccines Business:

Proprietary travel vaccines: IXIARO®, DUKORAL®, IXCHIQ®

Blockbuster Opportunity:

Lyme disease vaccine partnered with Pfizer showed strong efficacy in Phase 3; Regulatory submission validated by EMA; FDA submission planned with decisions expected within next 12 months

Deep Vaccine Expertise:

Integrated operations enhances know-how and accelerates development

Proven Development and Execution Track Record:

Three proprietary vaccines brought from the bench to people through multiple approvals

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Our Strategy as a Leading Vaccine Biotech Company

Contributing to a world in which no one dies or suffers from a vaccine preventable disease

Drive Commercial Growth

Provide stability and generate cash to invest in innovative R&D

Maximize R&D Upside

Create value with proven R&D engine

Deliver on meaningful clinical milestones

Fully Leverage Integrated Biotech Model

Augment clinical pipeline to achieve scale with a balanced portfolio

Enhance manufacturing and commercial operations

Proprietary Commercial Portfolio of Differentiated Vaccines

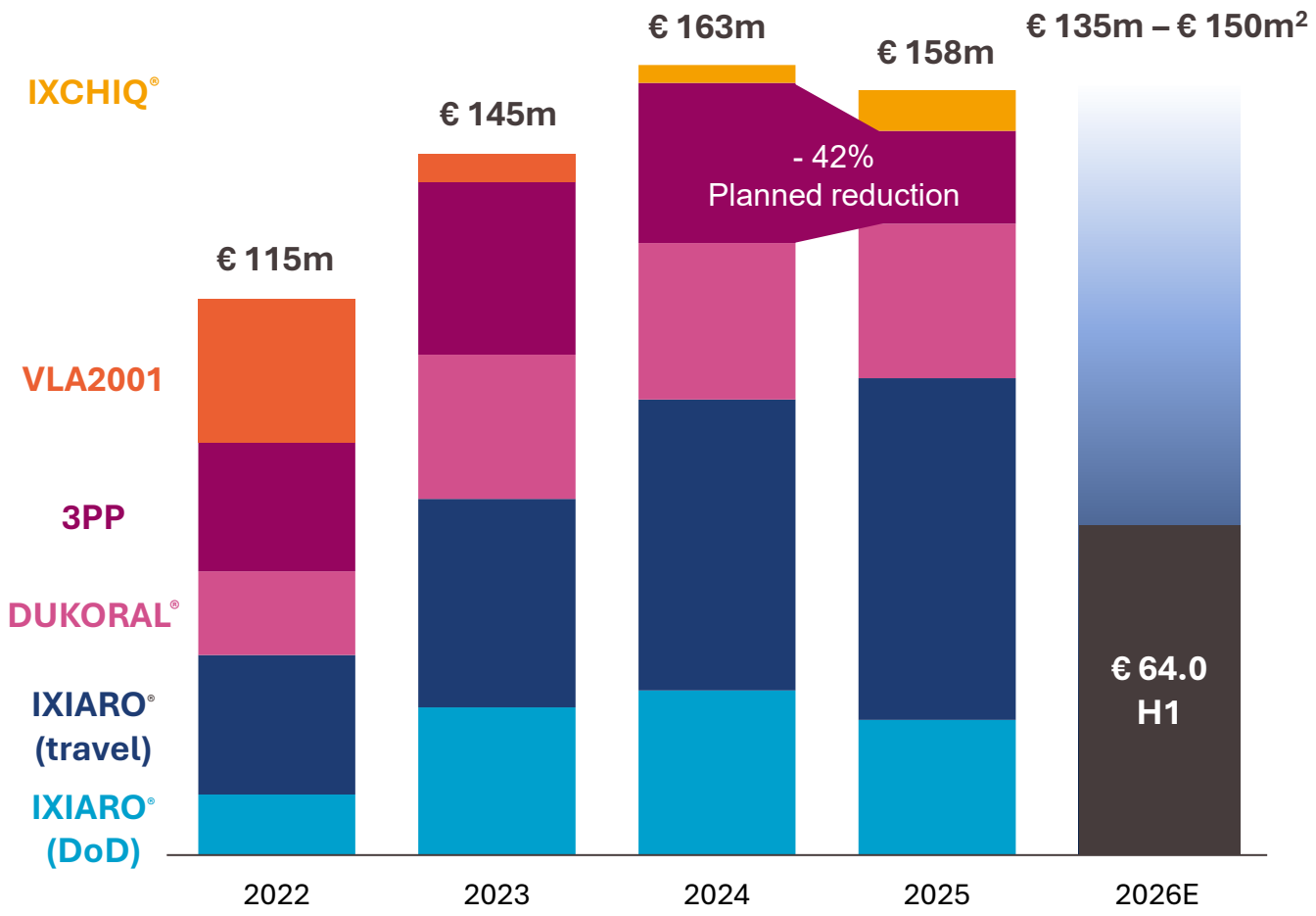
Differentiated vaccines

Brand	Indication	Differentiation	Key Markets
IXIARO®	Active immunization against **Japanese encephalitis** from 2 months of age	**Only** vaccine approved in U.S./Europe Requirement for U.S. military personnel deployed to parts of Asia	**Valneva direct markets**: US, CA, UK, FR, Nordics, BE, NL, AT **Key markets addressed by Partners**: DE, AU, IL
DUKORAL®	Active immunization against **Cholera and ETEC**[1] from 2 years of age	**Only** Cholera and LT-ETEC[1] vaccine approved in >30 countries	**Valneva direct markets**: CA, UK, FR, Nordics, AT **Key markets addressed by Partners**: DE, AU, IL, PL
IXCHIQ®	Active immunization against **chikungunya virus** in healthy individuals >12 years of age (Europe, Canada) and 18-59 years of age (UK, Brazil)[2]	**Strong and long-lasting** immunity across all age groups tested	**Valneva direct markets**: EU, CA, UK **Key markets addressed by Partners**: Brazil

1. ETEC indication in some markets only; 2. In August 2025, the U.S. FDA suspended the biologics license application (BLA) for IXCHIQ due to serious safety concerns. In January 2026, Valneva voluntarily withdrew the BLA and the associated investigational new drug application (IND) for IXCHIQ in the United States.

Vaccine Sales Fuel Strategic Investments into R&D

Focus on proprietary vaccines expected to drive revenue quality, margin expansion with strategic reduction in third-party product (3PP) sales[1]



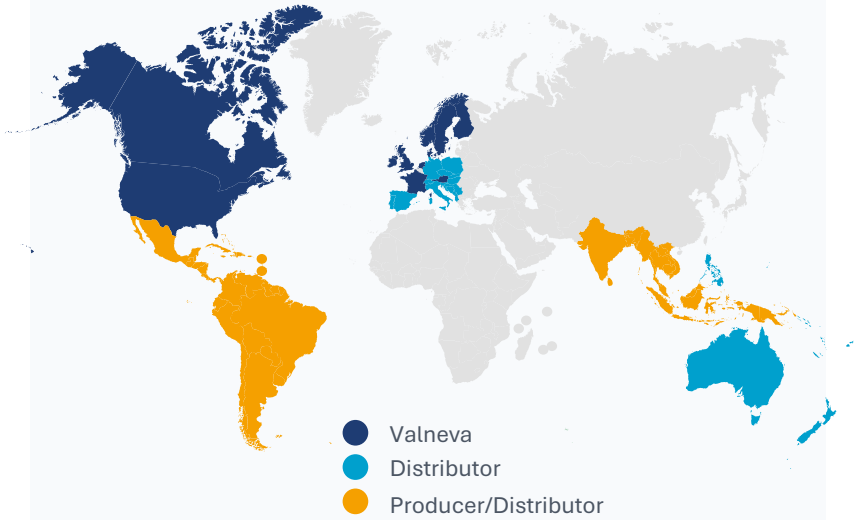
Lean and Efficient Commercial Infrastructure

Highly experienced teams

Deep vaccine expertise

Footprint extended through partners

Commercial Footprint

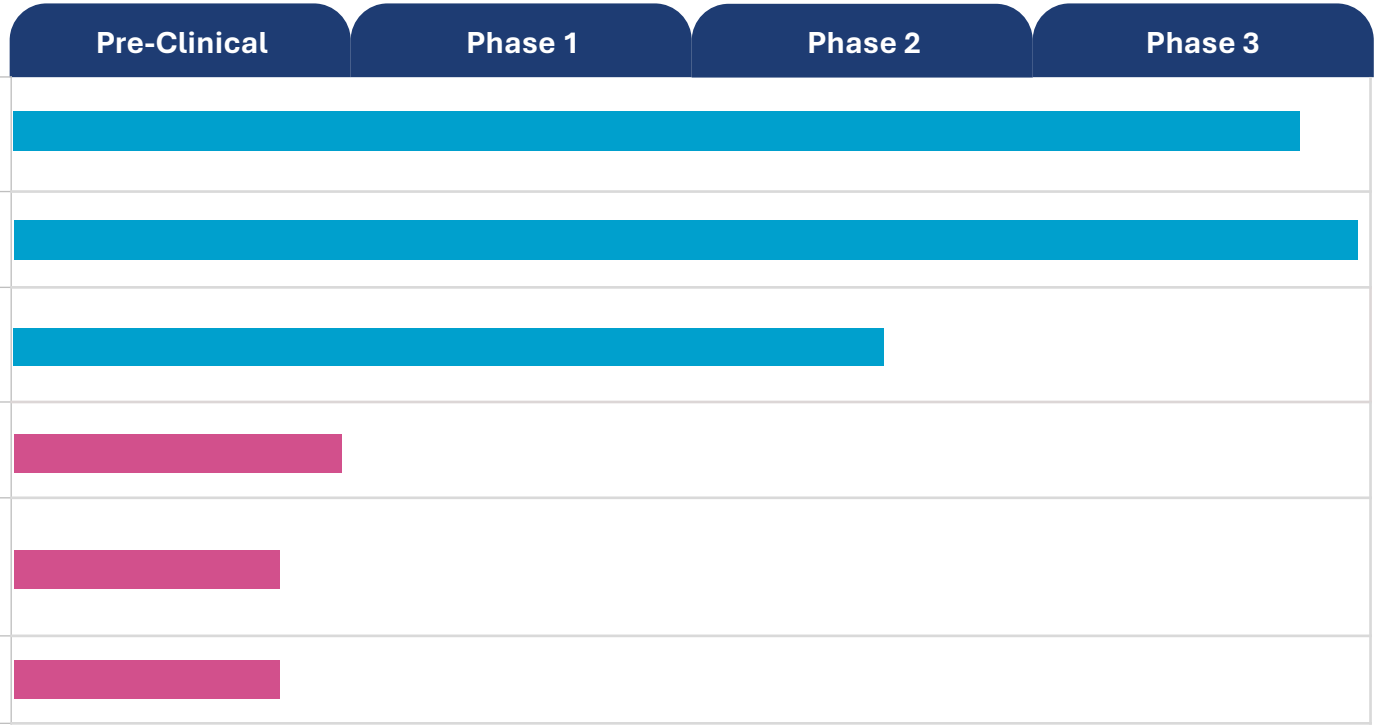


1.Targeting <5% of total product sales in 2026

2 Valneva Reports First Quarter 2026 Financial Results and Provides Corporate Updates

Valneva's Current R&D Portfolio

	Program	Design/Description
Clinical Programs	**LB6V / VLA15:** Lyme disease	World's most clinically advanced Lyme vaccine candidate; protein subunit-based
	VLA1553: Chikungunya	Post-marketing efficacy studies underway; live-attenuated vaccine
	S4V2: Shigellosis	Most advanced tetravalent bioconjugate vaccine candidate Potential first-in-class
Key Pre-Clinical Activities	**VLA2112:** EBV	Adjuvanted, protein subunit-based Large market, partnerable opportunity
	Enterotoxigenic *Escherichia coli* (ETEC)	Potential first ETEC vaccine candidate containing both heat-stable (ST) and heat-labile (LT) toxoid components
	Other	



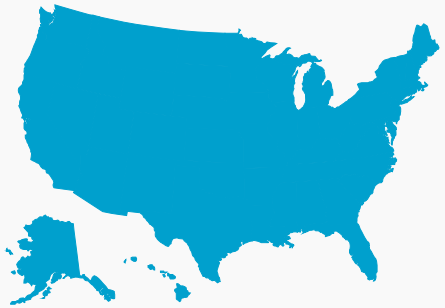
LB6V / VLA15

World’s leading Lyme Disease Vaccine Candidate

Lyme Disease Represents A Major Medical Need And Market Opportunity

No vaccine is currently available to prevent Lyme disease in humans

Commercial Opportunity



U.S: **87 million**

Population Living in Endemic Regions[1,2]

Europe: **223 million**

Population Living in Endemic Regions[1,2]

>$1 billion estimated global market[6]

Annual Burden of Disease

U.S. : **~476K cases**

Europe : **>132K cases**

Severe Manifestations[3]

10-30% cases develop

- Lyme carditis
- Lyme neuroborreliosis
- Lyme arthritis

Persistent Symptoms[4,5]

5-10% cases continue to have persistent symptoms following treatment

1 Kugeler et al. Emerging Infectious Disease, 2021 (doi.org/10.3201/eid2702.202731); **2** Davidson, A., Davis, J., Brestrich, G., Moisi, J., Jodar, L., & Stark, J. H. (in press) (2025). Lyme borreliosis incidence across Europe, 2015-2023: a surveillance-based review and analysis. Vector-borne and Zoonotic Diseases.; **3** Schwartz et al. Morbidity and Mortality Weekly Report Nov. 10, 2017; **4** Ursinus: https://www.thelancet.com/journals/lanepe/article/PIIS2666-7762(21)00119-8/fulltext; **5** Aucott, J.N., et al., Risk of post-treatment Lyme disease in patients with ideally-treated early Lyme disease: A prospective cohort study. Int J Infect Dis, 2022. 116: p. 230-237.; **6** Lyme Disease research and analysis conducted by an independent market research firm

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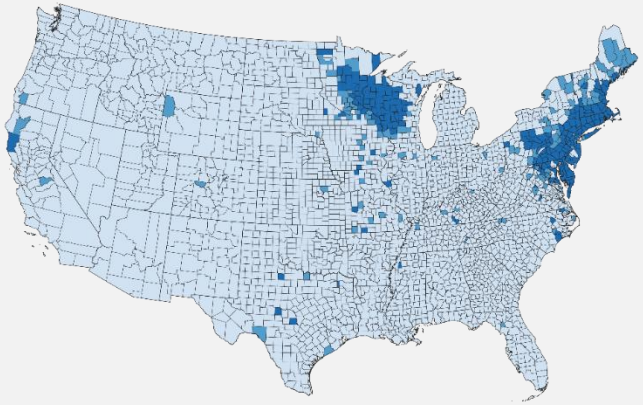
Lyme Disease Incidence in United States

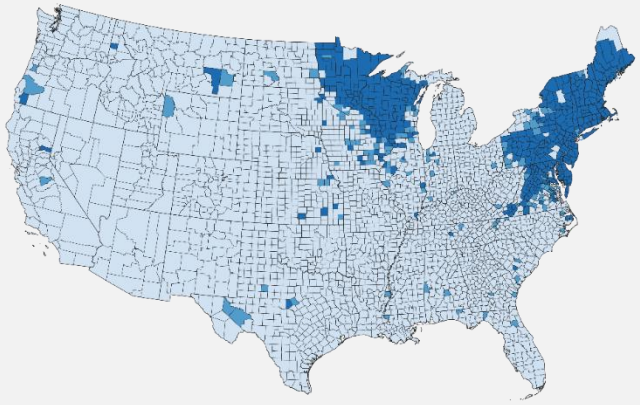
Incidence (cases per 100,000)
0 - 5
>5 - 10
>10

2002



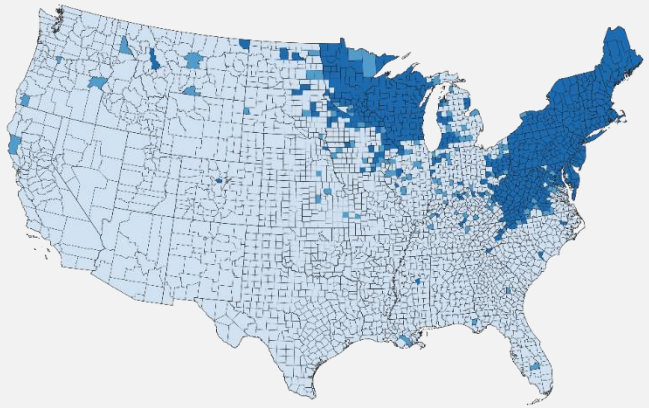
Focal areas of high incidence (>10 cases/100,000 persons) in the northeast and central Wisconsin and Minnesota

2012



Expansion of high incidence areas northward towards Canada and southward into the mid-Atlantic states

2022



Continued expansion of high incidence areas, particularly expansion into neighboring areas and states

Centers for Disease Control and Prevention, 2024: https://www.cdc.gov/lyme/data-research/facts-stats/surveillance-data-1.html

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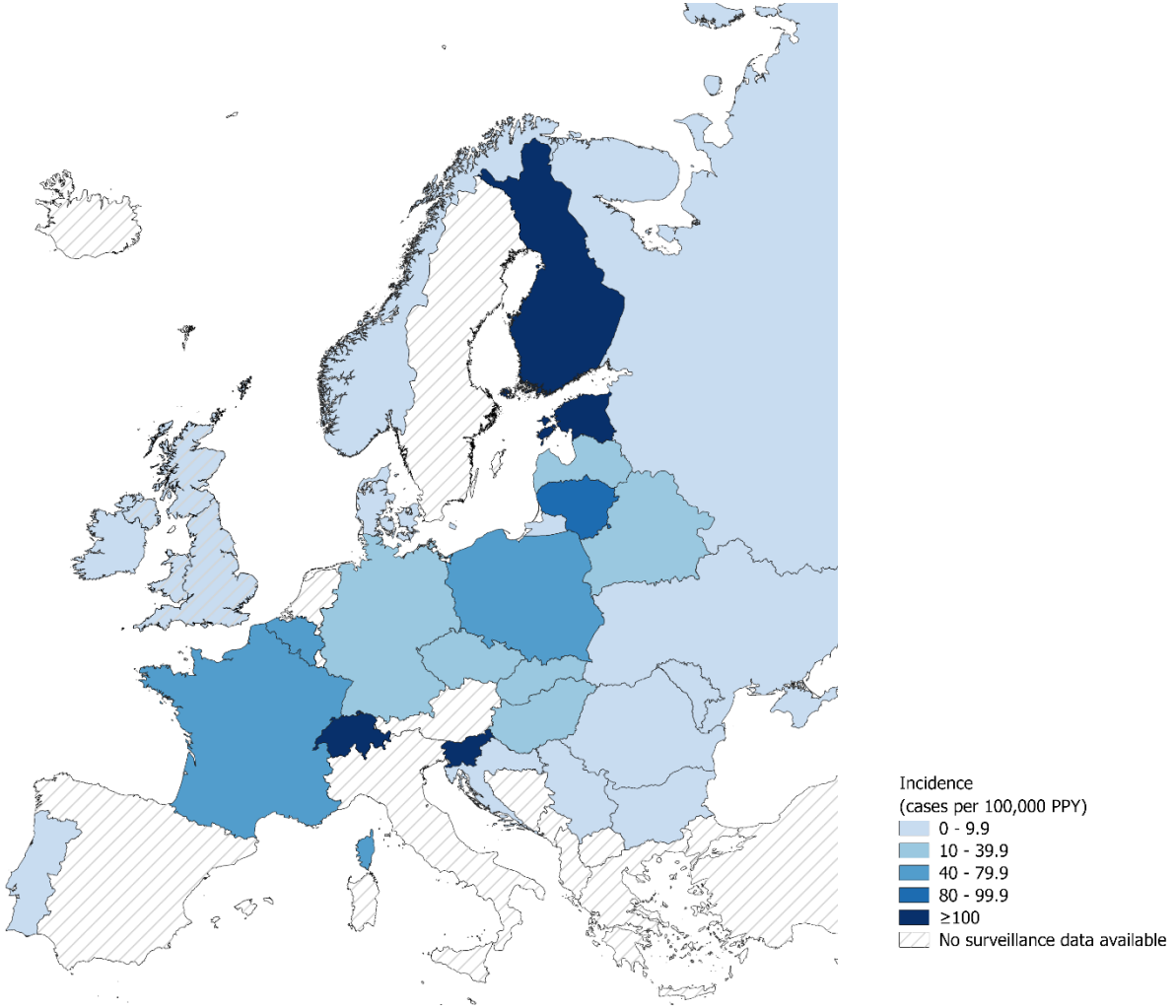
Average national incidence of Lyme borreliosis in the most recent three-year period* in Europe, 2015-2023



Alexander Davidson et al., Lyme Borreliosis Incidence Across Europe, 2015–2023: A Surveillance-Based Review and Analysis, Vector-Borne and Zoonotic Diseases (2025)
*Most recent three-year period of available data varies between countries
PPY = population per year

World's Leading Vaccine Candidate Against Lyme Disease

The only Lyme disease program in advanced clinical development today

Vaccine Highlights

Multivalent, recombinant proteins

Targets six most prevalent *Borrelia* serotypes causing Lyme disease in U.S. and Europe (>97% coverage)

Established mechanism of action

U.S. FDA Fast Track Designation

Market Opportunity

Exclusive, worldwide partnership[1]

Pfizer

>$1billion estimated global market[3]

Valneva eligible for upfront and milestone payments up to $408 million ($165 million received)

Tiered sales royalties 14-22%

Key Milestones

Reported strong efficacy in Phase 3 trial with no safety concerns identified[4]

EMA submission validated; Pfizer expects regulatory decisions in the next 12 months

Valneva eligible for up to $143m in early commercialization milestones

1 Pfizer and Valneva Initiate Phase 3 Study of Lyme Disease Vaccine Candidate VLA15; **3** Lyme Disease research and analysis conducted by an independent market research firm; 4. Pfizer and Valneva Announce Lyme Disease Vaccine Candidate Demonstrates Strong Efficacy in Phase 3 VALOR Trial

Phase 3: Randomized Placebo-Controlled Field Efficacy Study

Population: ~9,400 evaluable participants ≥5 years of age at high risk of Lyme disease (LD) (by residence and occupational/ recreational activities) in U.S., Canada and Europe (randomization approx. 1:1 VLA15/Placebo and 2:1 N. America/EU)

Primary endpoint: Rate of confirmed LD cases[2] after 2nd consecutive tick season (i.e., after completion of full vaccination series 3+1)

Secondary endpoints include rate of confirmed LD cases[1] after 1st tick season (i.e., after completion of primary vaccination series) amongst other secondary endpoints as defined in Phase 3 protocol

1 Cases are evaluated and confirmed by an Endpoint Adjudication Committee

LB6V (formerly VLA15) Demonstrates Strong Efficacy in Phase 3[1]

Results strengthen confidence – MAA validated by EMA[2]; regulatory decisions expected in the next 12 months[3]

Vaccine Efficacy

Pre-specified Analysis (Season 2)	Efficacy (95% CI)
From Day 28 post-dose 4	73% (16, 94)
From Day 1 post-dose 4	75% (22, 94)

Safety

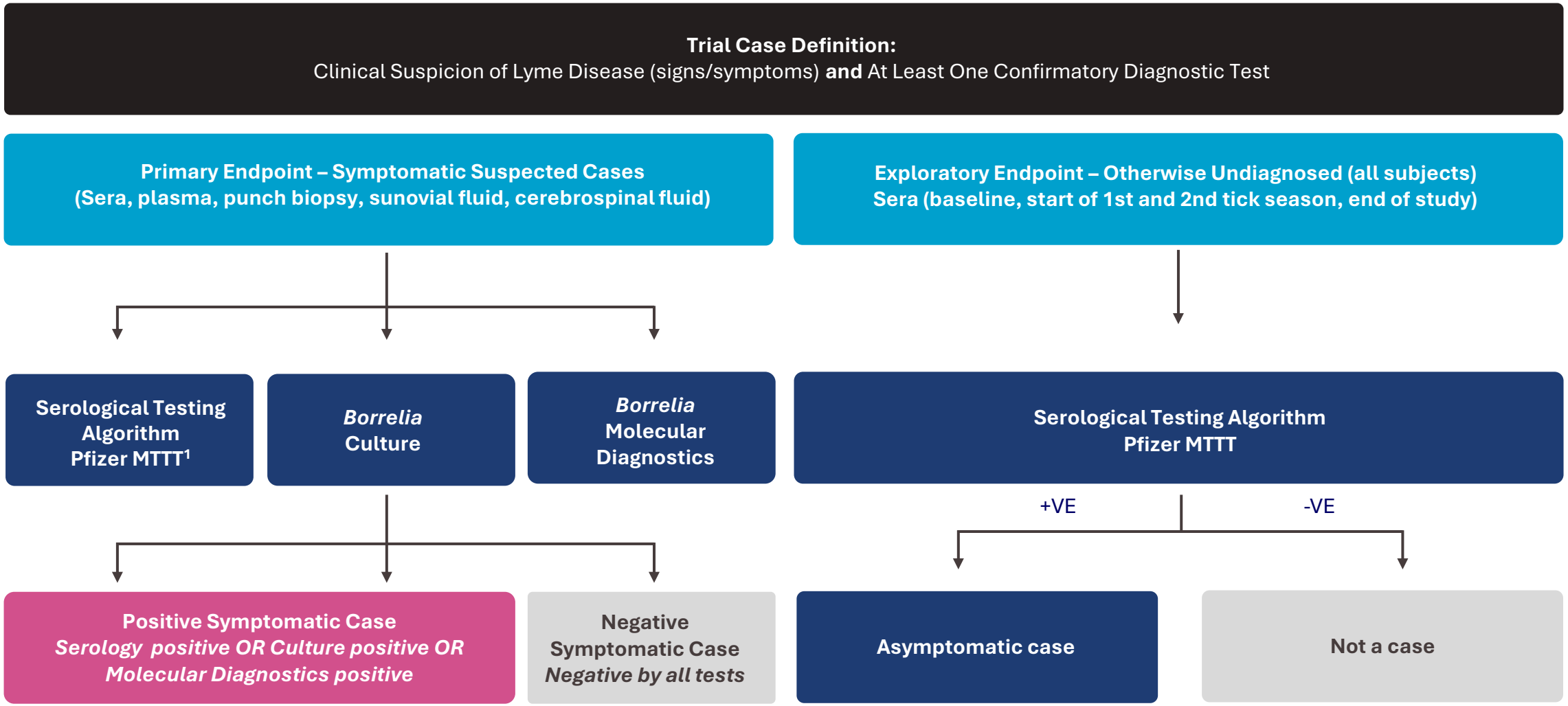
Vaccine candidate was well tolerated

No safety concerns identified at time of analysis

- Fewer than anticipated Lyme disease cases were accrued over the study period, and the pre-determined statistical criterion (95% confidence interval lower bound >20) was not met in the first pre-specified analysis (primary endpoint)
- Given the clinically meaningful efficacy and the fact that the 95% confidence interval lower bound was >20 in the second pre-specified analysis, Pfizer is confident in LB6V's potential and is optimistic that the vaccine candidate will obtain registration[4]

1 Pfizer and Valneva Announce Lyme Disease Vaccine Candidate Demonstrates Strong Efficacy in Phase 3 VALOR Trial; 2 Pfizer and Valneva's Lyme Disease Vaccine Candidate Marketing Authorization Application Validated by European Medicines Agency; 3 https://s206.q4cdn.com/795948973/files/doc_financials/2026/q2/Q2-2026-Earnings-Charts-FINAL.pdf ;
4 https://s206.q4cdn.com/795948973/files/doc_events/2026/Jun/08/PFE-USQ_Transcript_2026-06-08.pdf

Lyme vaccine Phase 3 diagnostic algorithms



1. Modified 2-tier testing

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LB6V is a Compelling Opportunity in an Underserved Market

First-Mover Advantage in Highly Receptive Market

Only Lyme disease vaccine candidate in late-stage clinical development

First potential Lyme vaccine in nearly 30 years

Differentiated Product Profile

Proven MoA with broader coverage: multivalent (six key serotypes)

Modern, state of the art recombinant protein vaccine

Compelling Target Population and Use Case

Broad addressable population (age 5+)

High and growing disease burden in high-risk areas

Exploding tick population in target markets

Strategic Fit with Pfizer's Vaccine Franchise

Leverages Pfizer's established commercial capabilities in adult and pediatric vaccines

Attractive Commercial Dynamics

Prophylactic vaccine model supports predictable demand and repeat dosing

Potential inclusion in routine immunization schedules for high-risk areas

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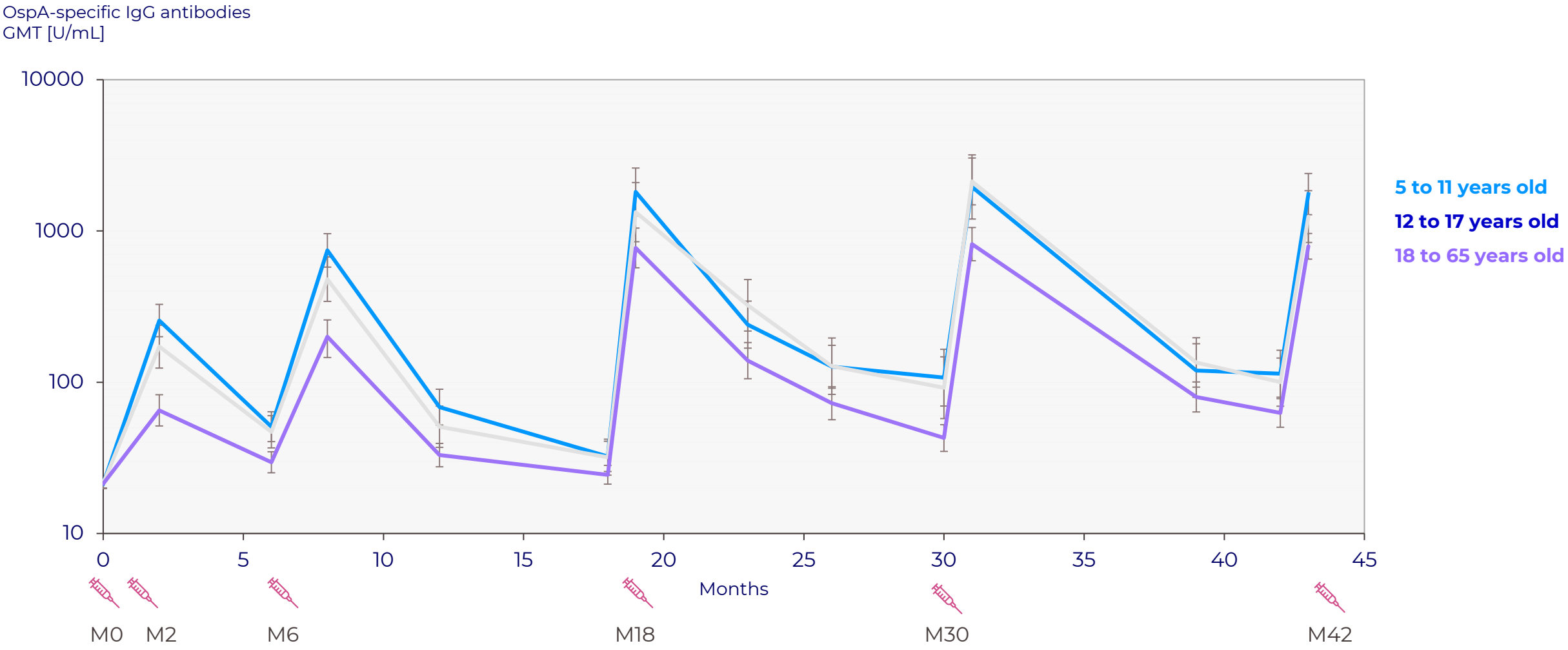
Study VLA15-221: Booster doses significantly increase OspA antibody titers



VLA15-221: Analysis 4 Table 14.4.3.2.4: ELISA: GMTs for OspA ST1 Specific IgG Antibodies by Visit (Full Analysis Set);
Analysis 5. Tables 14.5.3.1.4-9: ELISA: GMTs for OspA ST1-ST6 Specific IgG Antibodies by Visit (Per-Protocol Analysis Set)

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Valneva's and Pfizer's Collaboration to Co-Develop and Commercialize LB6V

Established	April 2020
Updated	June 2022; Equity Investment of $95 Million by Pfizer; Phase 3 cost split 40/60% (Valneva/Pfizer)*
Rationale	Maximize Lyme disease opportunity by leveraging Pfizer's outstanding development and commercial expertise
Scope	Pfizer leading late-stage development and will have sole control over global commercialization
Key Financial Terms	Valneva eligible to receive up to $408 million ($165 million received) • $130 million upfront payment (received) • $35 million in development milestone payments (received) • $143 million in early commercialization milestones • $100 million in cumulative sales milestones Tiered sales royalties ranging 14-22%
Co-development costs	Valneva portion fulfilled in H1 2024

* As of 1st May 2022

IXCHIQ® / VLA1553

A Highly Differentiated single-shot Chikungunya Vaccine

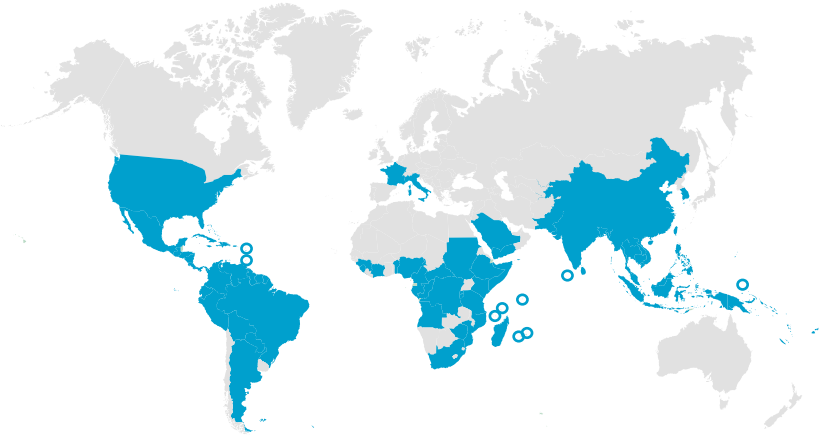
Chikungunya: A Major Public Health Threat

Mosquito-transmitted outbreak disease with potentially debilitating consequences

Aedes aegypti

Aedes albopictus

- Often causes **large, explosive outbreaks**
- Affecting **up to 75%** of the local population[1]
- Substantial quality-of-life and **health-economic impact**[2]
- Nearly half (43%) of those infected develop chronic symptoms[3]
- ~Major outbreaks in China, India, and Indian Ocean in 2025
- La Reunion/Mayotte outbreaks led to 700 locally acquired cases in France[4]
- 2025 cases >double in UK vs 2024[5]



75% of world population lives in areas at-risk of chikungunya

Identified in **>110 countries** across five continents

1. Staples et al. CDC Yellow Book 2020, Chapter 4; 2. The global health and economic burden of chikungunya from 2011 to 2020: a model-driven analysis on the impact of an emerging vector-borne disease; 3. Bettis et al, PLOS Neglected Tropical Diseases 16(1): e0010069. 4. 2025 data as of Oct. 8th 2025: https://www.ecdc.europa.eu/en/chikungunya-virus-disease/surveillance-and-updates/seasonal-surveillance; 5. https://doi.org/10.1136/bmj.r1746

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IXCHIQ®: Focused on Confirming Efficacy/Safety and Expanding Access

Robust clinical program supported by CEPI grant

Pilot Vaccination Campaign Ongoing (Brazil)

To serve as the basis for post-marketing commitment studies

- Launched in February 2026 with partner Instituto Butantan in select municipalities in Brazil
- Adults aged 18 – 59 years ▶ objective to achieve 20 – 40% coverage within the target population; ~50,000 vaccinated to date

Post-Marketing Effectiveness Studies

To confirm effectiveness and to optimize description of the safety profile

- Observational effectiveness study (VLA1553-402) and prospective safety cohort study (VLA1553-406) in Brazil
- Special population studies: observational pregnancy surveillance (VLA1553-403); Study in HIV-infected individuals[1]
- Pragmatic randomized controlled effectiveness and safety study: adults and adolescents in endemic countries (VLA1553-404)

Ensuring Greater Access

To address unmet medical needs in endemic countries

- Expanding network of manufacturing and distribution partners in low-and-middle-interest countries (LMICs)
- May 2026: Locally manufactured vaccine approved in Brazil (VLA1555 / “Butantan-chik”); expected to be incorporated into Brazil’s public health system

1 Valneva is participating as part of a European consortium with grant funding to evaluate IXCHIQ® vaccination in healthy and HIV infected volunteers in Africa

S4V2

World’s Most Clinically Advanced Tetravalent Shigella Vaccine Candidate

S4V2: Opportunity to Develop First-in-Class Vaccine for a Life-Threatening Disease

Tetravalent bioconjugate vaccine with potential to cover up to ~85% of shigellosis infections[1]

Vaccine Highlights

World's most clinically advanced tetravalent Shigella vaccine candidate

Exclusive global license from (LMTB)[2]

Includes four most common pathogenic Shigella bacteria serotypes: *S. flexneri* 2a, 3a, 6, and *S. sonnei*

Positive initial Phase 1/2 clinical data reported[3]

Awarded FDA Fast Track designation

Market Opportunity

Global market expected to exceed $500 million annually[4]

Second-leading cause of fatal diarrheal disease; Up to estimated 165 million cases and 600,000 deaths annually[5]

Identified as a priority vaccine by World Health Organization (WHO)[6]

Valneva has worldwide commercialization rights upon potential approval

Key Milestones

Phase 2 infant study launched in 2025; data expected in Q3 2026

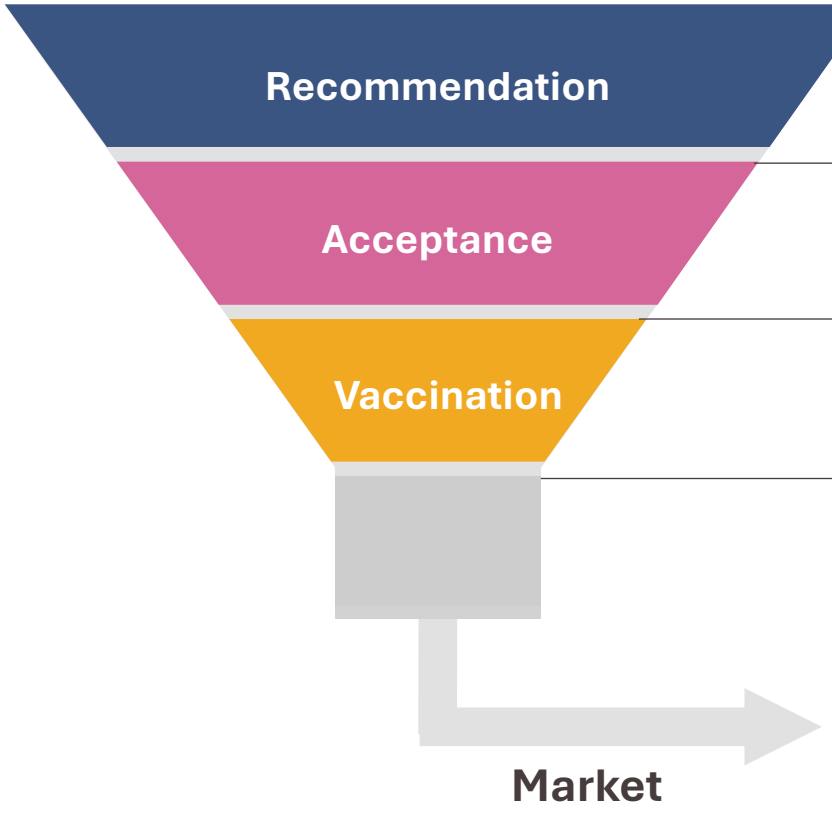
Ongoing Phase 2b CHIM[7] study aiming to provide early look at potential efficacy; data expected in Q3 2026

Valneva to determine next development steps based on clinical data and future R&D strategy

1. https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8878964/pdf/vaccines-10-00212.pdf; **2.** Valneva and LimmaTech Enter into a Strategic Partnership to Accelerate the Development of the World's Most Clinically Advanced Tetravalent Shigella Vaccine Candidate; **3** 20240221_LimmaTech_Shigella-Interim-Data-PR_Final.pdf (lmtbio.com); **4** LEK 2024; Appox. 7 years after launch; **5.** *Shigellosis* | CDC Yellow Book 2024; **6.** Immunization, Vaccines and Biologicals (who.int); **7.** Controlled Human Infection Model

Commercial Assessment of Shigella Vaccine

Shigella vaccine market estimated to peak at ~€500 million[1]



	Traveler (~23%)	Children in endemic countries (~76%)	Military (<1%)
Recommendation	20-50% (Med-high risk destinations)	30-100% (OOP-reimbursement)	70-90% (Med-high risk destinations
Acceptance	30% (OOP)	15 - 60% 50-90% (reimbursement)	90%
Vaccination	2-5%	9-11%[2] (OOP) 15-50%[2] (reimbursement)	100%
Market	Europe, North America, Japan, South Korea and Singapore with €80-85m from the U.S. alone	Non-Gavi endemic countries: public funding anticipated, driven by high disease burden notably in India, Brazil, Indonesia and Gavi supported countries. Gavi-supported countries, vaccines could cover over 10m children per year	U.S. representing c.50% of the revenue

Source: Market Study: LEK 2024, 1 Appox. 7 years after launch; 2 Converted to vaccination rate by applying the yearly vaccination penetration every year over a cohort of 5 years

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Financial Guidance & Outlook

Financial Outlook

2026 Sales Guidance

- Product Sales: €135 - €150 million (reiterated); Commercial business expected to remain cash-flow positive
- Total Revenues: €145 - €160 million
- New DoD expected in the coming months

Financial Outlook

- Continued cash flows from proprietary commercialized vaccines
- Restructuring program implemented with positive P&L and cash flow impacts expected from H2 2026
- Product gross margins expected to improve in H2 2026 following non-recurrent effects H1 2026
- Potential for financial self-sustainability starting in 2027 subject to successful Lyme disease vaccine regulatory approval and commercialization

Our Next Phase as a Leading Vaccine Biotech Company

Potential LB6V Success Would Offer Strategic Growth Opportunities

Leverage core strengths in vaccine development to deliver greater long-term value

Key initiatives:

Build scale in R&D pipeline post-LB6V exit

- Strategic in-licensing/M&A to augment clinical-stage pipeline
- Curate a risk-balanced portfolio of innovative specialty, life-cycle and high-value vaccine assets

Expand vaccine focus beyond vector-borne diseases

- Target new assets based on defined criteria (ongoing)
- Advance EBV, ETEC/enteric disease candidates; strategic focus on reducing antimicrobial resistance (AMR)

Optimize integrated operations

- Control value chain by investing into enhanced end-to-end capabilities
- Structure commercial model to generate cash

Thank you
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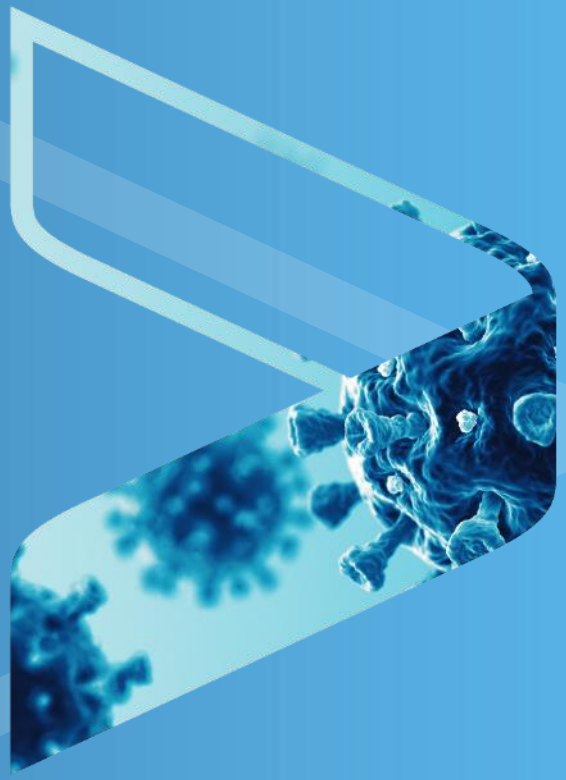
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