



Immunic
THERAPEUTICS

Immunic Therapeutics

Developing Selective Oral Therapies in Immunology

NASDAQ: IMUX | August 2025

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→ Forward-looking statements included in this presentation are based on information available to Immunic as of the date of this presentation. Immunic does not undertake any obligation to update such forward-looking statements except as required by applicable law.

CLINICAL-STAGE BIOPHARMACEUTICAL COMPANY (NASDAQ: IMUX)

Dedicated to improving the lives of patients with
chronic inflammatory and autoimmune diseases



Innovative pipeline:

First in class oral drugs with unique
modes of actions for multiple sclerosis
and gastrointestinal diseases



Positive MS phase 2 data sets:

Underline neuroprotective effect of
Nurr1 activation by vidofludimus calcium



Large commercial opportunity:

\$3.7 billion peak sales potential for
vidofludimus calcium in MS



Experienced leadership team:

Successfully developed and
commercialized multiple medicines



Financials:

Cash balance of \$55.3 million
as of June 30, 2025

Leadership Team

Company is Led by an Experienced Management Team



Daniel Vitt,
PhD
Chief Executive
Officer



Jason Tardio,
MBA
President & Chief
Operating Officer



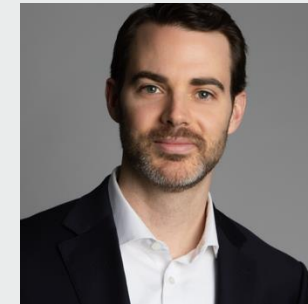
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MD, MBA
Chief Medical
Officer



Hella Kohlhof, PhD
Chief Scientific
Officer



Glenn Whaley, CPA
Chief Financial
Officer



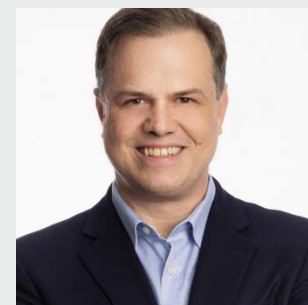
Patrick Walsh
Chief Business
Officer



Inderpal Singh
General Counsel



Werner Gladdines
Chief Development
Officer



Duane Nash,
MD, JD, MBA
Executive
Chairman

Advanced Clinical Pipeline

Well Differentiated Programs in Various Phases of Clinical Development

Program	Preclinical	Phase 1	Phase 2	Phase 3	Key Program Updates
Vidofludimus Calcium (IMU-838)*					✓ Phase 2 EMPhASIS trial in RRMS successfully completed, significantly reduced brain lesions, encouraging results in reducing disability worsening
	Relapsing Multiple Sclerosis (RMS) ENSURE 1 and ENSURE 2 Trials				✓ Interim analysis of ENSURE program completed, IDMC recommendation to continue trials as planned, both ENSURE trials fully enrolled in May 2025
					✓ CALLIPER trial successfully completed, substantial reductions in disability worsening
	Progressive Multiple Sclerosis (PMS) CALLIPER Trial				✓ Phase 2 CALDOSE-1 trial in UC completed, effective in 50 weeks maintenance phase
					▪ Top-line data for both ENSURE trials expected by end of 2026
IMU-856					✓ Phase 1/1b trial in healthy volunteers and celiac disease completed, first proof-of-concept in celiac disease
	Celiac Disease and other Gastrointestinal Disorders				✓ Dose-dependent increase of endogenous GLP-1 in post hoc analysis of phase 1b trial in celiac disease
IMU-381					▪ Further clinical testing in preparation
	Gastrointestinal Diseases				

■ Ongoing ■ Completed ■ In preparation or planned

RRMS: relapsing-remitting multiple sclerosis; IDMC: Independent Data Monitoring Committee; GLP-1: glucagon-like peptide-1

*Additional investigator-sponsored phase 2 RAPID_REVIVE trial of vidofludimus calcium in post COVID syndrome ongoing, sponsored by University Hospital Frankfurt



Vidofludimus Calcium in Multiple Sclerosis (MS)

Targeted to Elevate the Standard
of Care for the Full Spectrum of
Multiple Sclerosis Patients

Vidofludimus Calcium Has the Potential to Transform the Oral Multiple Sclerosis DMT Market

Designed to Combine the Best of Two Worlds: Neuroprotection and Relapse Prevention

First-in-class, dual mode of action approach designed to address the **full spectrum of disease**:

- Nurr1 activation provides **direct neuroprotective effects**
- DHODH inhibition is associated with **anti-inflammatory effects**

Oral DMT category: Aims for **best-in-class benefit / risk profile** by combining **strong efficacy** with **safety, tolerability**, and **once-daily** convenience

No first-dose or on-treatment monitoring makes it an **easy start or switch to therapy**

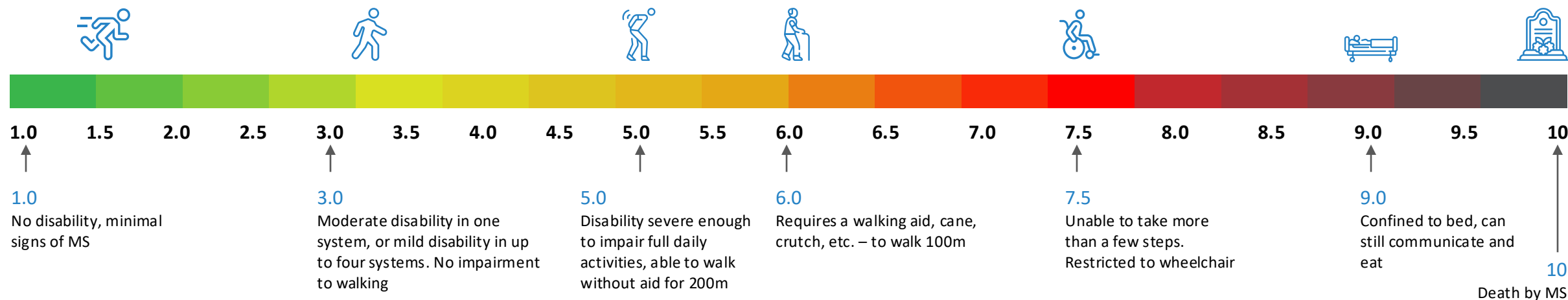
No anticipated black box warnings or serious infection risk (e.g., PML, malignancies, etc.)



→ **If approved, peak sales potential for vidofludimus calcium of \$3-7 billion^[1]**

DMT: disease-modifying therapy; Nurr1: nuclear receptor-related 1; DHODH: dihydroorotate dehydrogenase; PML: progressive multifocal leukoencephalopathy [1] Based on Immunic internal market research

Regardless of the Subtype, the Outcome of Every Patient Journey in Multiple Sclerosis Is Physical and/or Cognitive Disability



While over 15 anti-inflammatory treatments exist for relapsing multiple sclerosis, there is no therapy available that directly impacts the neurodegeneration driving disability progression

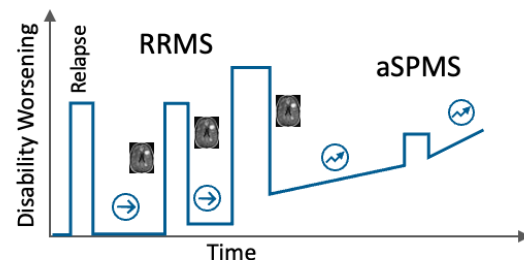


Vidofludimus Calcium Has the Potential to be the First and Only Oral DMT Approved for Both Relapsing and Progressive MS

Relapsing MS (RMS)

Relapsing MS

- Relapses and MRI lesions dominate clinical course (RRMS)
- Fewer relapses and lesions with continuous disability progression (aSPMS)



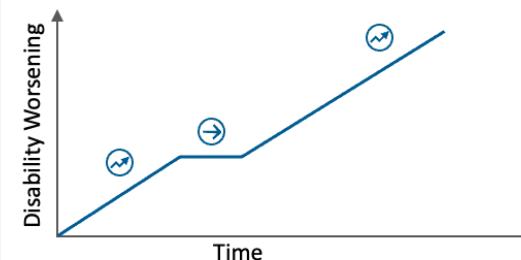
~900,000 patients diagnosed

DHODH Inhibition

Progressive MS (PMS)

Primary Progressive MS

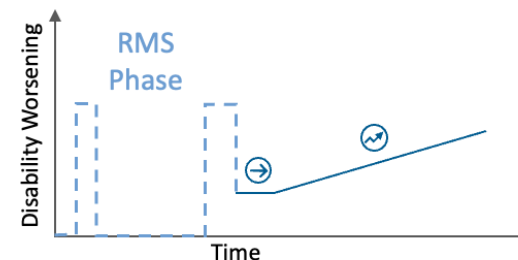
Disability worsening without relapses from the start



~120,000 patients diagnosed

Nurr1 Activation

Relapses have stopped, but disability progression continues



~175,000 patients diagnosed

Relapses and focal inflammation (RAW)

Smoldering disease and neurodegeneration (PIRA)

DMT: disease-modifying therapy; MS: multiple sclerosis; RRMS: relapsing-remitting MS; SPMS: secondary progressive MS; aSPMS: active SPMS; MRI: magnetic resonance imaging; DHODH: dihydroorotate dehydrogenase; Nurr1: nuclear receptor-related 1; RAW: relapse-associated worsening; PIRA: progression independent of relapse activity

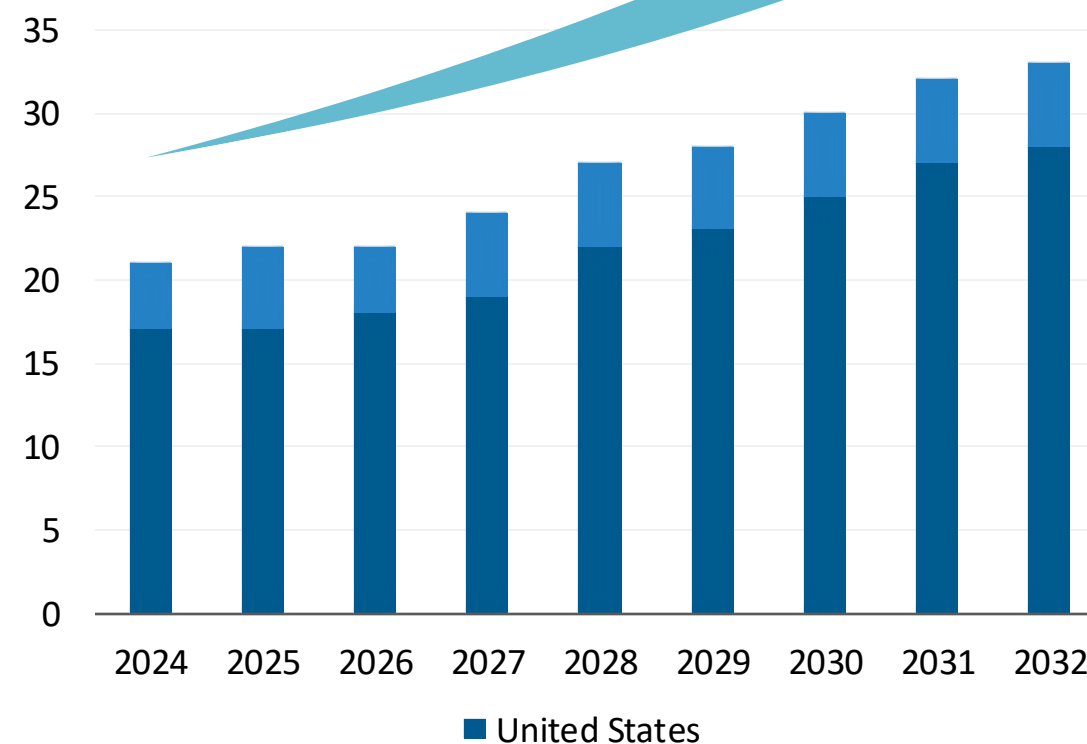
A Large and Growing Global Market Where Multiple Blockbusters Coexist

Many brands are generating in excess of \$1 billion in global annual sales in 2024^[1]

Ocrevus [®]	\$7.6 billion
Kesimpta [®]	\$3.2 billion
Tysabri [®]	\$1.7 billion
Tecfidera [®] & Vumerity [®]	\$1.6 billion
Mavenclad [®]	\$1.15 billion
Avonex [®] & Plegridy [®]	\$968 million
Rebif [®]	\$626 million
Gilenya [®]	\$552 million
Aubagio [®]	\$379 million
Briumvi [®]	\$310 million

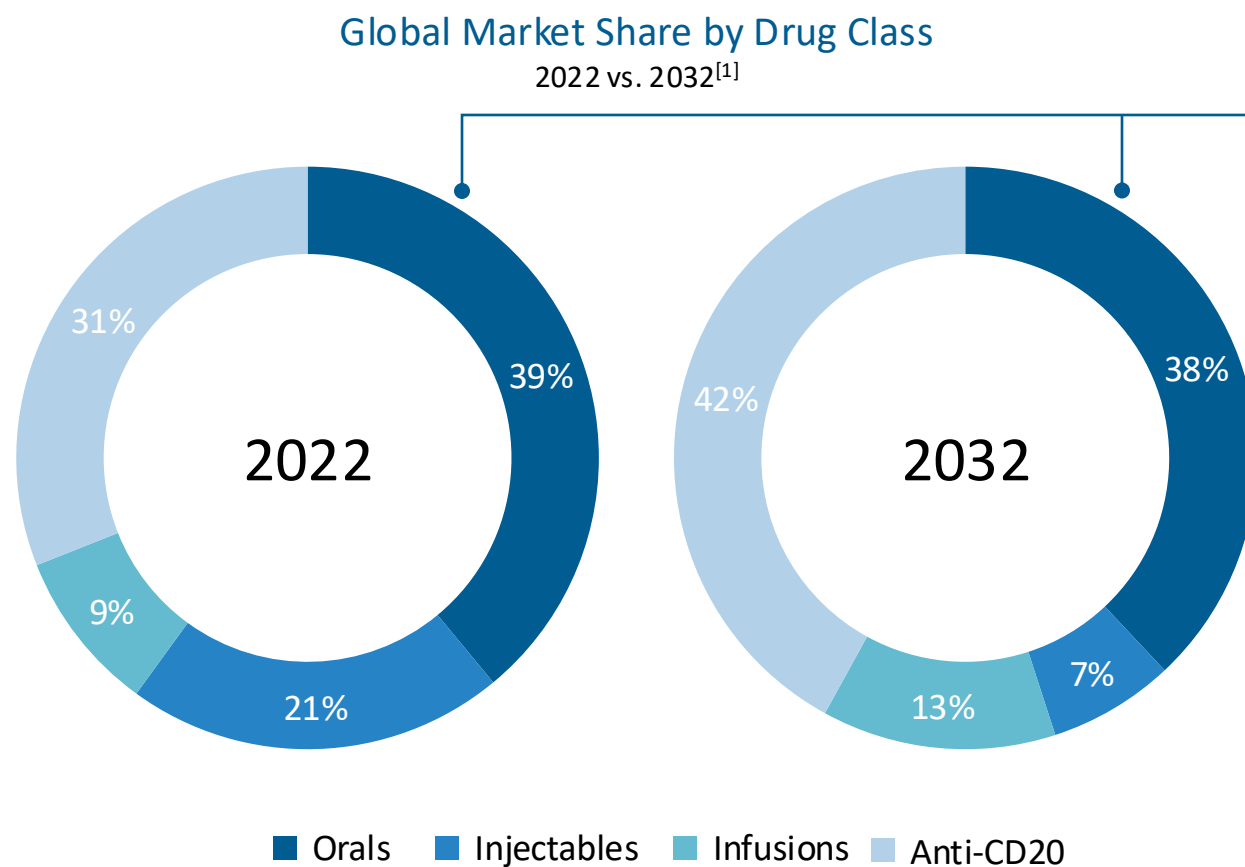
\$20 billion market today growing 4% y/y^[2]

Major market sales of MS therapies(\$ billion)



[1] Company public filings [2] Sales numbers in G7 countries (US, UK, Canada, Japan, Germany, France, Italy) in USD billion; Multiple Sclerosis Landscape and Forecast by Decision Resources Group Part of Clarivate

Oral DMTs Will Continue to Play a Big Role as Important Treatment Options



While anti-CD20 class of therapies continues to grow, oral class still expected to capture over 1/3 of the global market

- 42% of patients prefer oral medicines^[2]
- Early-line reliance on injectable therapies will continue to wane as the market shifts to using oral therapies earlier
- 15% of patients with PPMS and 25% of patients with non-active SPMS received oral treatments (off label)^[3]

[1] Sales numbers in G7 countries (US, UK, Canada, Japan, Germany, France, Italy) in USD billion; 2024 Multiple Sclerosis Landscape and Forecast by Decision Resources Group Part of Clarivate. [2] Jonker MF, et al. Med Decis Making. 2020 Feb;40(2):198-211 [3] Watson C, et al. Neurol Ther. 2023 Dec;12(6):1961-1979 / DMT: disease-modifying therapy; CD20: B lymphocyte cell-surface molecule; SPMS: secondary progressive MS; PPMS: primary progressive MS

Multiple MS Patient Segments Could Benefit from Vidofludimus Calcium



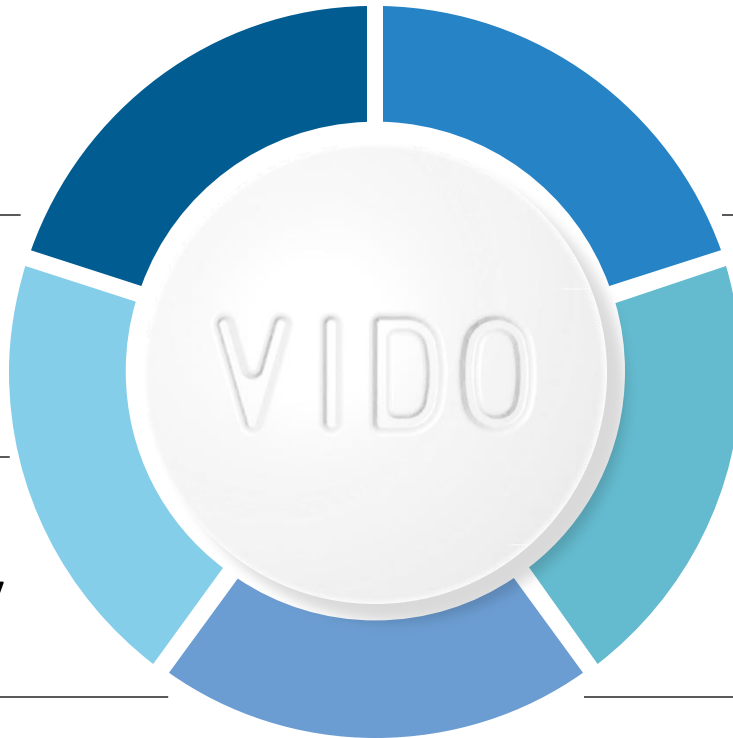
Newly diagnosed patients



Patients switching therapies due to disability worsening



Patients switching therapies due to tolerability or safety concerns



Older patients where immunosuppression is a concern



Untreated patients



Patients with progressive disease (naSPMS & PPMS)



naSPMS: non-active secondary progressive MS; PPMS: primary progressive MS

Vidofludimus Calcium: Derisked Near-Term Opportunity With \$3-7 Billion Peak Potential



Indication



Status



Clinical Evidence



Eligible Population



Next Milestones



Potential Peak Sales

RMS

Phase 3

78% reduction of new Gd+ lesions (Phase 2)

~900K

Phase 3 completion expected **end of 2026**

\$1-2B

naSPMS

Phase 3-ready

19% reduction of 24-week CDW (Phase 2)

~175K

End of phase 2 meeting with regulators **TBD**

\$1-2B

PPMS

Phase 3-ready

33% reduction of 24-week CDW (Phase 2)

~120K

End of phase 2 meeting with regulators **TBD**

\$2-3B

Patient and market size numbers sourced via internal Immunic analysis and 2024 Multiple Sclerosis Landscape and Forecast report by Decision Resources Group Part of Clarivate
RMS: relapsing MS; naSPMS: non-active secondary progressive MS; PPMS: primary progressive MS; Gd+: gadolinium-enhancing; CDW: confirmed disability worsening; K: thousand; B: billion

Global Market for PPMS Treatment Estimated to Be \$6+ Billion But Less Than Half of All Diagnosed Patients Are Treated Today

~120K

diagnosed PPMS patients
in the US & EU5

45%

of diagnosed PPMS patients
are currently on a DMT

~\$2.7B

in PPMS sales for the only
approved product



**Total global market for PPMS estimated to be \$6B+ and expected to grow
with the approval and increased availability of new medicines**

PPMS: primary progressive multiple sclerosis; DMT: disease-modifying therapy; K: thousand; B: billion / Patient and market size numbers sourced via internal Immunic analysis and 2024 Multiple Sclerosis Landscape and Forecast report by Decision Resources Group Part of Clarivate; EU5 countries: France, Germany, Italy, Spain, and United Kingdom; TD Cowen Therapeutic Categories Outlook Comprehensive Study – Multiple Sclerosis October 2024



Vidofludimus Calcium in Multiple Sclerosis (MS)

First-in-Class, Potent
Nurr1 Activator and
Selective DHODH Inhibitor

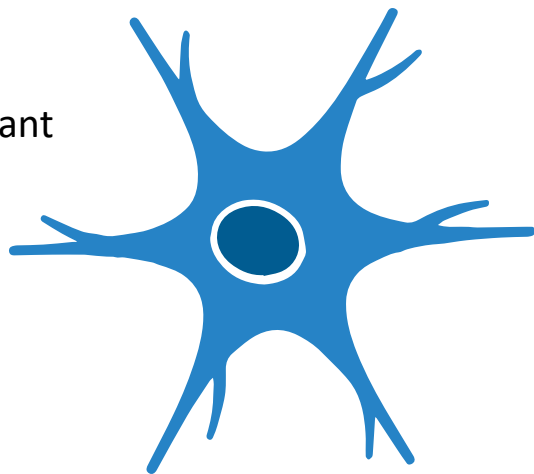
Vidofludimus Calcium Addresses Smoldering Neurodegeneration



First-in-Class Nurr1 Activator, Targeting Improvement of Physical and Mental Ability of Multiple Sclerosis Patients

Nurr1 Activator

- Direct and indirect **neuroprotective** effects
- Involved in protecting relevant neurons from cell death
- Known effects reducing activation of microglia and astrocytes
- Effect independent from focal inflammation

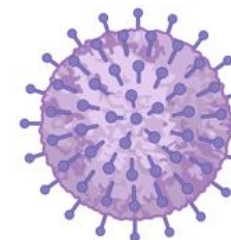


DHODH Inhibitor

- Selectively targets hyperactive immune cells
- Selective **anti-inflammatory** effects, reducing focal inflammation, magnetic resonance imaging lesions and relapses
- Broad-spectrum **antiviral** effects prevent reactivation of EBV and could stop cross reactive immune responses



Blocking of
Th17/Th1 cytokines



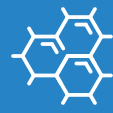
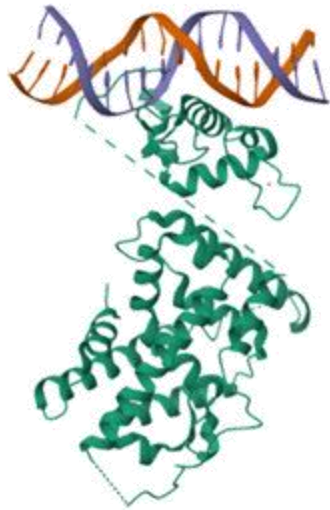
Vidofludimus Calcium Activates Nurr1, Shown to Increase Neuronal Survival



Nurr1 Binding

Nurr1 is a transcription factor binding to DNA^[1]

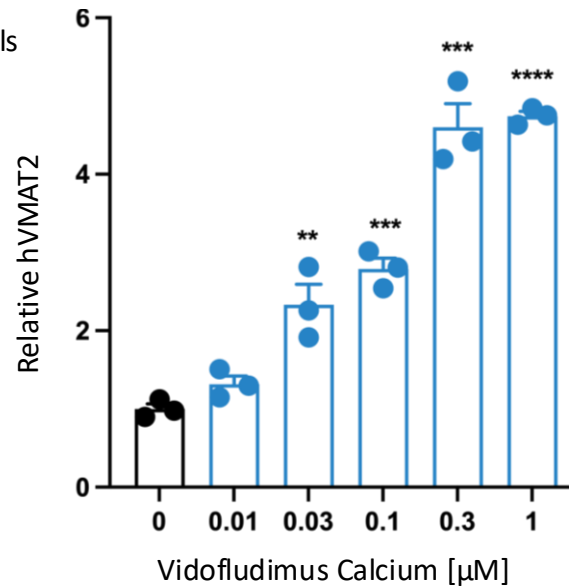
Vidofludimus calcium binds to and strongly activates Nurr1 activity with nM values



Gene Expression Regulation

Vidofludimus calcium induces a > 2-fold induction of target gene expression of VMAT2 at 30 nM concentration^[2]

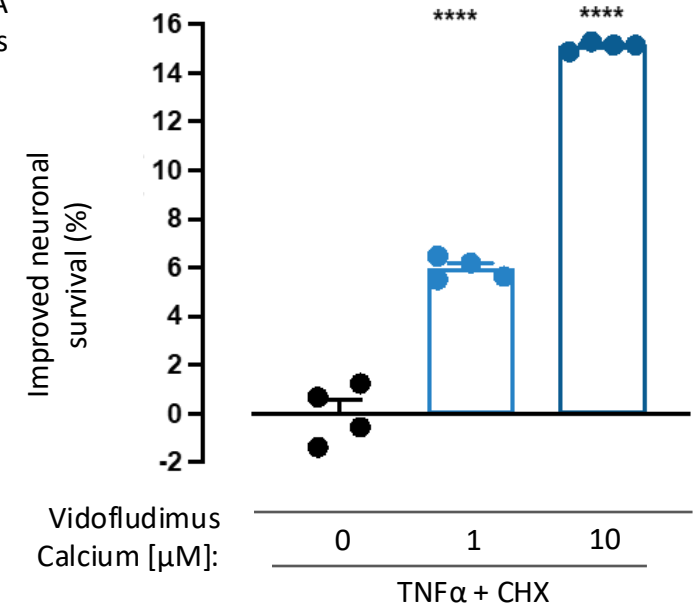
Human microglia cells (HMC3)



Improves Neuronal Survival

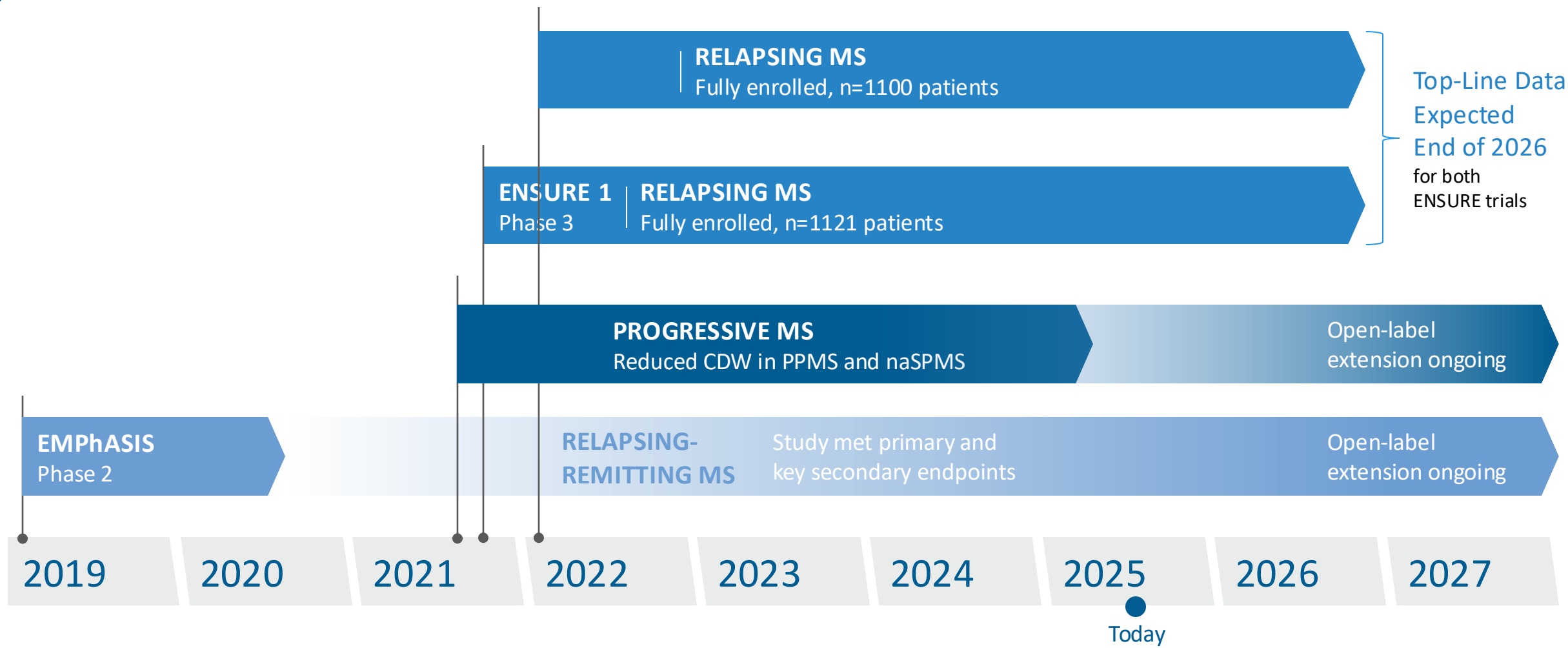
Vidofludimus calcium improves neuronal survival via Nurr1 activation^[3]

N2A cells



[1] Vieter et al., Journal of Medicinal Chemistry 2023 66 (9), 6391-6402 The related research project was funded by the German Federal Ministry of Education and Research under the grant number 03INT607AA; Structure: Zhao, M. et al. (2022) Proc Natl Acad Sci USA 119;
[2] Sun, Zuoming. City of Hope. 2023, unpublished [3] Unpublished data: Sun lab, City of Hope, Duarte; 2023 / Nurr1: nuclear receptor-related 1; DNA: deoxyribonucleic acid; VMAT2: vesicular monoamine transporter 2; DMSO: dimethyl sulfoxide; TNF: tumor necrosis factor

Vidofludimus Calcium: Clinical Trials Overview in Multiple Sclerosis (MS)



CDW: confirmed disability worsening; PPMS: primary progressive multiple sclerosis; naSPMS: non-active secondary progressive multiple sclerosis



Vidofludimus Calcium in Multiple Sclerosis (MS)

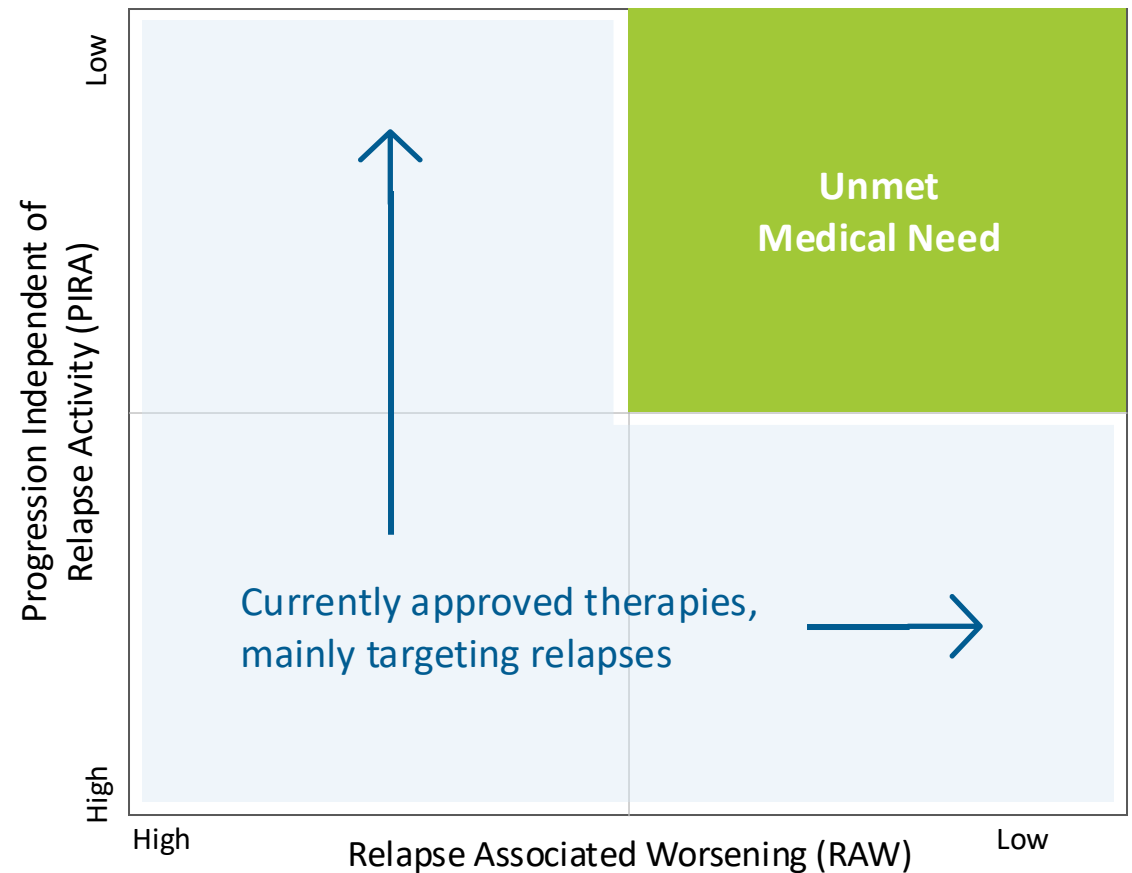
Development in Relapsing Multiple Sclerosis (RMS)

Vidofludimus Calcium Could be the First Treatment Option for Relapsing MS Fulfilling the Current Unmet Needs of Patients



Goals for New Relapsing Multiple Sclerosis Treatments

- Developing a new therapy offering:
 - Neuroprotection and effect on progression independent of relapse activity (PIRA)
 - Excellent safety and tolerability
 - Easy to use, convenient oral administration without complex screening requirements
- Developing a new therapy for newly diagnosed patients and as an excellent switch opportunity



EMPhASIS: Completed Phase 2 Trial in Relapsing-Remitting MS

NCT03846219



Coordinating Investigator

Robert J. Fox, M.D.
Cleveland Clinic



Double-Blind, Placebo-Controlled,
Randomized, Parallel-Group Trial

- Blinded main treatment period of 24 weeks
- Cohort 1: 30 and 45 mg or placebo QD
- Cohort 2: 10 mg or placebo QD
- Extended treatment period of up to 9.5 years ongoing to observe long-term safety is ongoing



Trial Met Key
Efficacy and Safety Endpoints

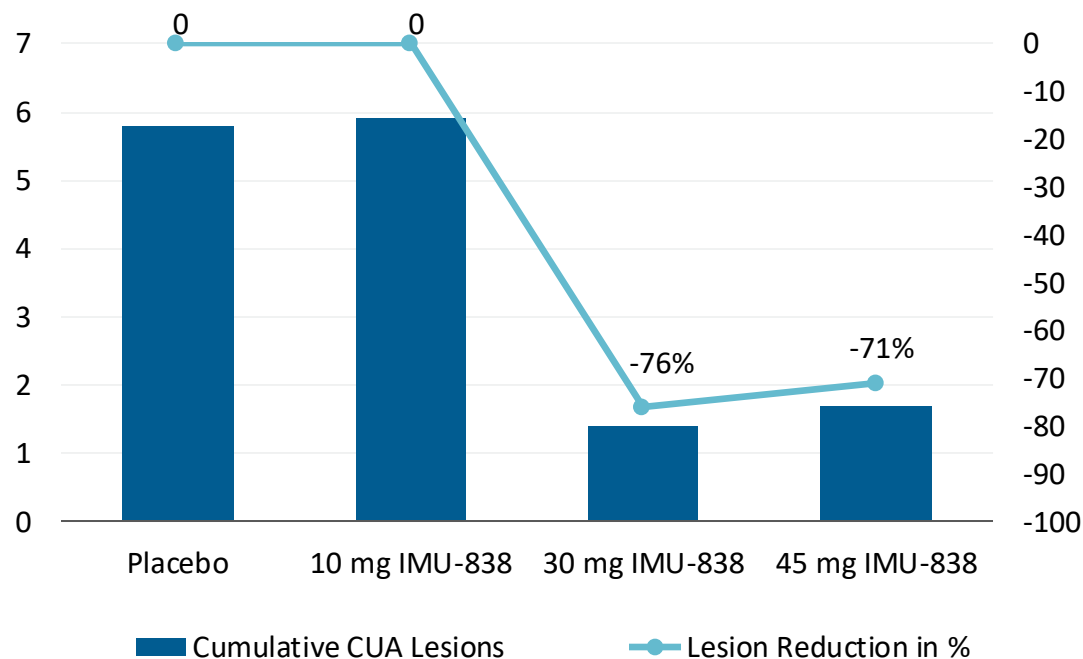
- Randomized 268 patients in 36 centers across four European countries
- Vidofludimus calcium showed strong activity in relapsing-remitting MS population
 - Primary and key secondary endpoints met with high statistical significance: strong reduction of MRI lesion activity
 - Reduced serum NfL concentrations
 - Signal in preventing confirmed disability worsening
- Vidofludimus calcium's safety profile was similar to placebo
 - No general safety signals observed
 - Low discontinuation rates, considerably lower than placebo

MS: multiple sclerosis; QD: quaque die = once-daily; MRI: magnetic resonance imaging; NfL: neurofilament light chain

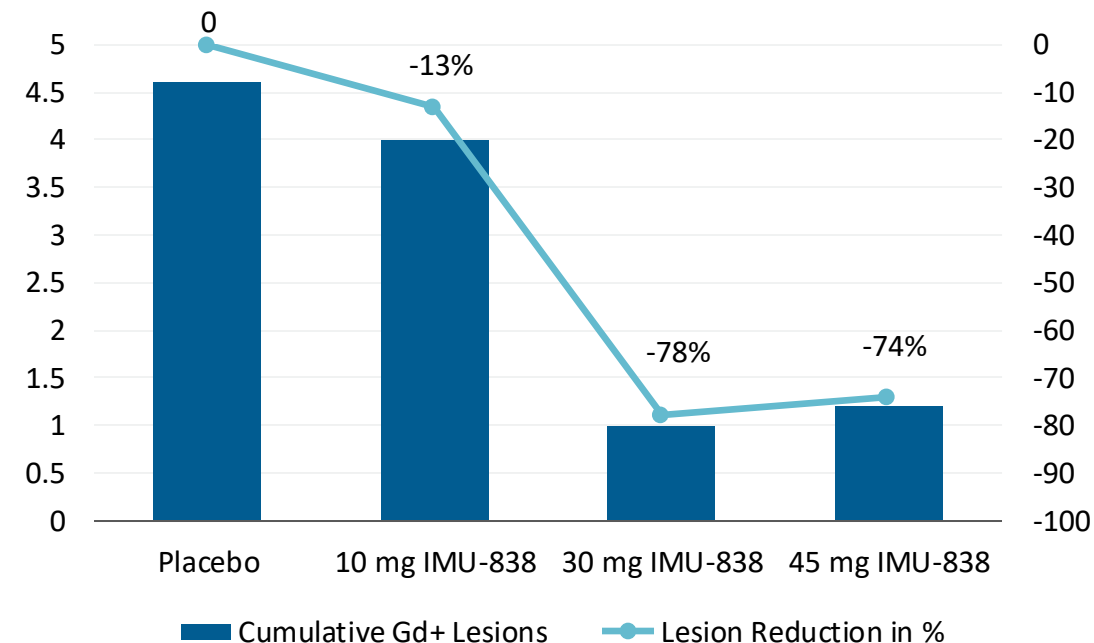
EMPhASIS: Strong Reduction of MRI Lesion Activity

Primary Endpoint Hit With High Statistical Significance, Pooled Cohorts 1 & 2

Reduction in Cumulative CUA Lesions up to Week 24



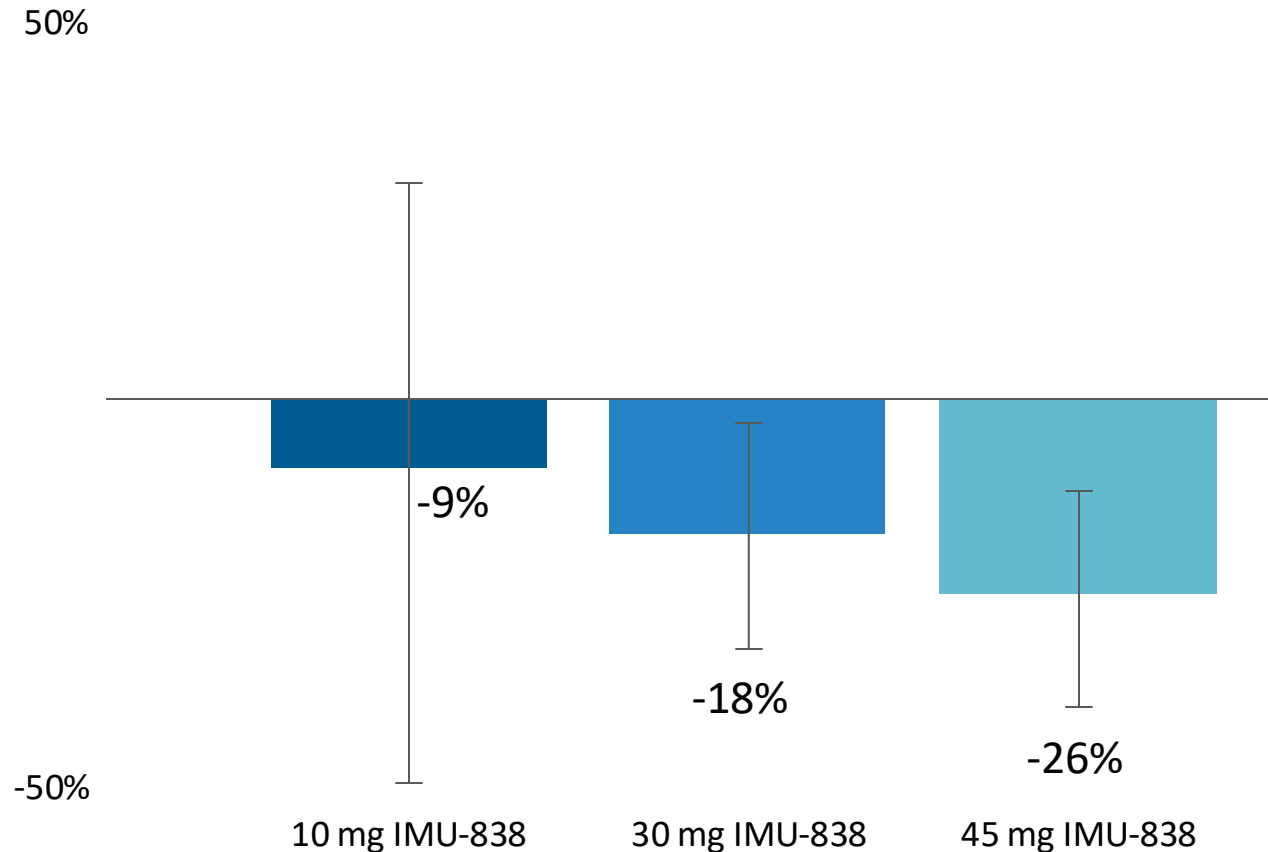
Reduction in Gd+ Lesions up to Week 24



Primary and key secondary endpoints of cumulative number of new CUA lesions up to week 24 met with high statistical significance (primary 45 mg vs. placebo: $p = 0.0002$ / key secondary 30 mg vs. placebo: $p < 0.0001$)

As Cohort 2 only allowed MRI machines of 1.5T, pooled data of Cohorts 1 & 2 only include patients that were evaluated at MRI field strength of 1.5 Tesla. Modified full analysis set C1/C2 (N10 = 47, N30 = 65, N45 = 66, NPBO C1 = 59, NPBO C2 = 12). Data displayed are as adjusted mean values. Estimates are adjusted for baseline volume of T2 lesions and baseline number of Gd+ lesions (0, >=1) using a generalized linear model with a negative binomial distribution and a logarithmic link function. Log transformation of time from first investigational medicinal product (IMP) dose to date of last MRI assessment with non-missing values is used as offset term / RMS: relapsing-remitting multiple sclerosis; MRI: magnetic resonance imaging; CUA: cumulative unique active, Gd+: gadolinium-enhancing

EMPhASIS: Reduction of Serum NfL Concentrations Observed Versus Placebo After 24 Weeks, Pooled Cohorts 1 & 2



Vidofludimus calcium showed remarkable reduction in NfL levels in all active doses tested compared with placebo

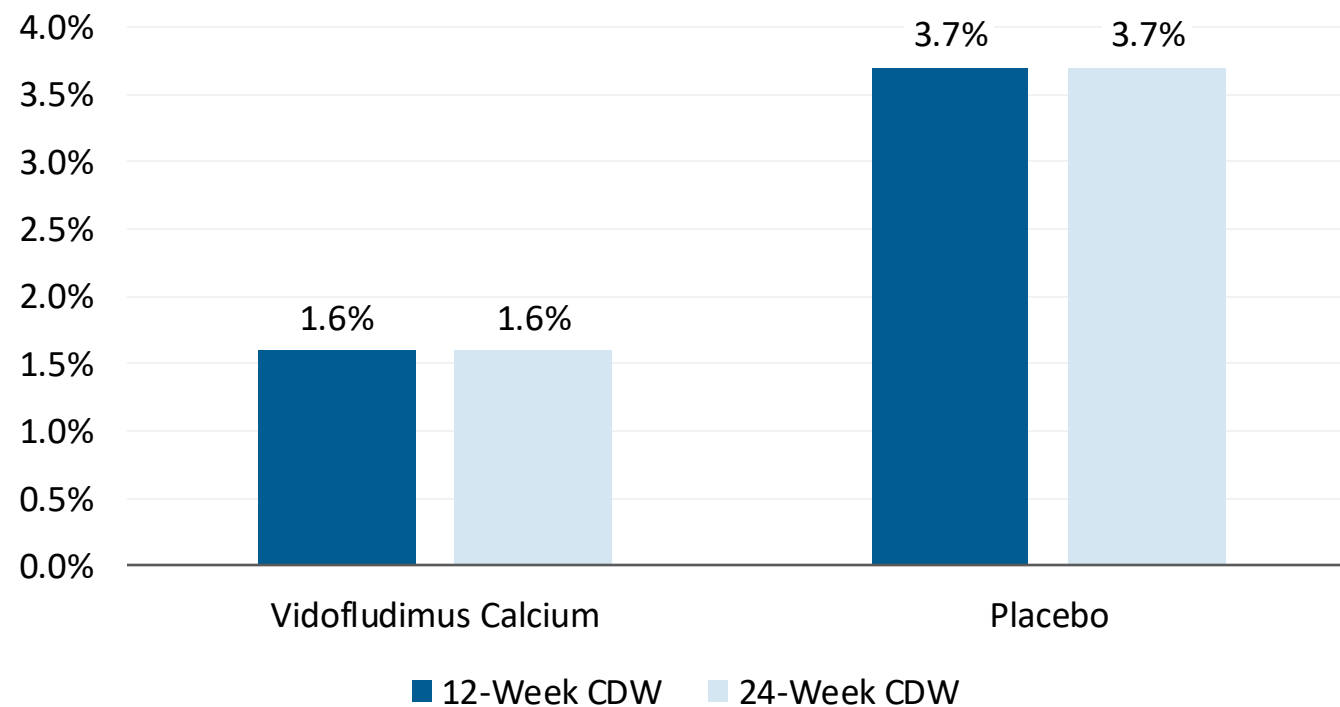
- Clear dose-response relationship in NfL reduction
- Higher doses expected to show stronger neuroprotective effects

Displayed are median values of differences between percentage change of serum neurofilament light chain concentration (Hodges-Lehmann estimation), treatment vs. placebo
Data shows 10 mg versus placebo for Cohort 2 and 30/45 mg versus placebo for Cohort 1; NFL: neurofilament light chain

EMPhASIS: Reduced Confirmed Disability Worsening Events

End of 24-Week Blinded Treatment Period

CDW Events at the End of the 24-Week Blinded Treatment Period



CDW: confirmed disability worsening; EDSS: Expanded Disability Status Scale

Only disability worsenings with a trigger point during the 24-week blinded treatment period are considered. The EDSS increases during the blinded treatment phase were subsequently confirmed during open-label extension phase of the trial. Patients at risk in this analysis are 187 for vidofludimus calcium (pooling 10, 30 and 45 mg data) and 81 for placebo. The trigger event is an EDSS progression defined as an increase in the EDSS compared to Baseline of at least 1.5 points if Baseline EDSS = 0, of at least 1.0 points if Baseline EDSS of 1-5, or of at least 0.5 points if Baseline EDSS ≥ 5.5

12-week CDW: The confirmation event is at least 77 days after the trigger event. At the confirmation event and each assessment between trigger and possible confirmation event, EDSS must be at least as high as at the trigger event.

24-week CDW is defined analogously, the only difference being the time interval between trigger event and confirmation visit, which is at least 161 days.

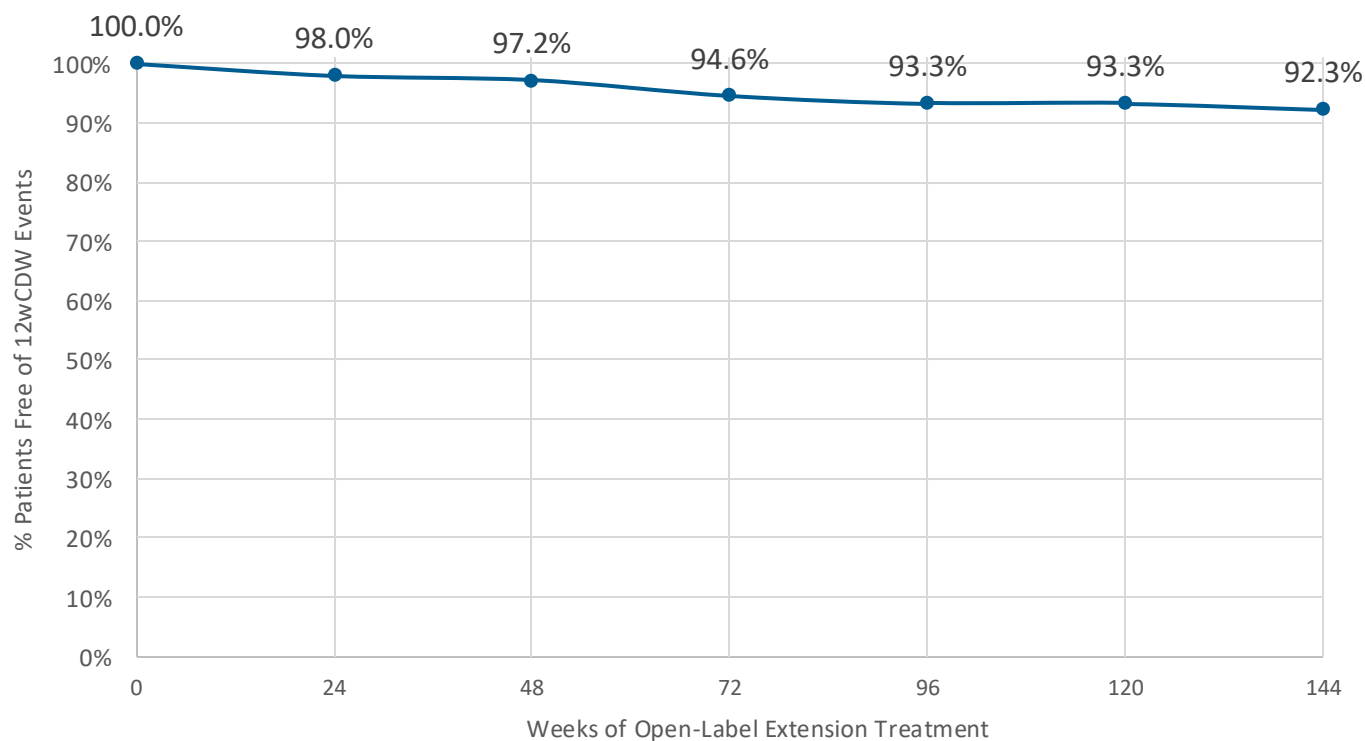
Full analysis set pooled cohorts 1&2 (N10 = 47, N30 = 71, N45 = 69, NPBO C1 = 69, NPBO C2 = 12)

- Signal in preventing 12-week and 24-week confirmed disability worsening events as compared to placebo
- Confirmatory data will be obtained in phase 3 ENSURE clinical program

EMPhASIS: Low Rates of Confirmed Disability Worsening Events

Open-Label Extension Period, 196 Patients Reaching 144 Weeks of Treatment

Proportion of patients free of 12-week confirmed disability worsening after up to 144 weeks of open-label extension vidofludimus calcium treatment



- At week 144, 92.3 % of patients remained free of 12-week CDW
- 29 CDW events confirmed at 12 weeks following trigger event through week 144
- Of these, 13 events (44.8%) defined as relapse-associated worsening (RAW), while only 4 (13.8%) as progression independent of relapse activity (PIRA)
- Low discontinuation rate with 196 out of 254 patients reaching 144 weeks of OLE treatment
- Vidofludimus calcium continued to demonstrate favorable safety and tolerability profile

CDW: confirmed disability worsening; OLE: open-label extension

Vidofludimus Calcium: Unrivaed Safety and Tolerability Profile Observed in Multiple Clinical Trials

- Safety profile similar to placebo: no general safety signals observed in clinical trials so far
- No increased rates of diarrhea, neutropenia, or alopecia
- No increased rates of infections and infestations or hematology values
- Drug exposure tested in approximately 2,700 human subjects and patients, to date, with data available up to 5.5 years
- Low rates of adverse events
- No signals for hepatotoxicity or elevations of liver enzymes and no Hy’s law cases observed to date



Vidofludimus Calcium’s Safety Profile to Date is Unique

	PML risk	Increased number of infections	Vaccination limitations	Gastrointestinal toxicities, incl. diarrhea	Cardiovascular risks, incl. blood pressure	Lymphopenia	Neutropenia	Risk of liver injury	Increased risk of cancer	Macular edema
Vidofludimus Calcium	●	●	●	●	●	●	●	●	●	●

● Favorable profile

PML: progressive multifocal leukoencephalopathy

EMPhASIS: Vidofludimus Calcium Well-Tolerated With Adverse Events Similar to Placebo

Safety & Tolerability Endpoints	Placebo	Vidofludimus Calcium 30 mg	Vidofludimus Calcium 45 mg
Any treatment-emergent adverse event	44%	45%	41%
Treatment-emergent adverse events occurring in >5% of total patients by preferred term			
Headache	6%	4%	6%
Nasopharyngitis	4%	4%	7%
Treatment-emergent adverse events occurring in 2%-5% of total patients by preferred term			
Upper respiratory tract infection	4%	3%	0%
Viral respiratory tract infection	4%	0%	3%
Treatment-emergent adverse events occurring in >1 to <2% of total patients by preferred term			
Back pain	3%	1%	0%
ALT increase	3%	1%	0%
Influenza	3%	0%	1%
Liver enzymes elevated	1%	1%	3%
Nausea	1%	1%	3%
Bronchitis	1%	0%	3%
Alopecia	0%	4%	1%
Fatigue	0%	3%	3%
Rash	0%	3%	3%
Cystitis	0%	1%	4%
Treatment-emergent adverse events by severity			
Mild	33%	41%	30%
Moderate	12%	16%	23%
Severe	1%	0%	0%
Series adverse events	1%	3%	0%
Treatment discontinuation for any reason	7%	3%	6%
Treatment-emergent adverse events leading to treatment discontinuation	4%	0%	3%

The observed adverse events were generally mild in nature.

There were very few adverse events with medium and high incidence rate.

EMPhASIS: Patients Feel Well-Treated With Vidofludimus Calcium



Reflected in **Low Discontinuation Rates** for Vidofludimus Calcium-Treated RRMS Patients, Considerably Lower Than Placebo*

	Vidofludimus Calcium	Glatiramer Acetate ^[1]	Aubagio® ^[2]	Tecfidera® ^[3]	Gilenya® ^[4]	Zeposia® ^[5]
Administration	Oral	Injectable	Oral	Oral	Oral	Oral
Daily Dose	30 mg QD	20 mg QD	14 mg QD	240 mg TID	1.25 mg QD	1 mg QD
Treatment Period	24 weeks	9 months	36 weeks	24 weeks	6 months	24 weeks
Active Treatment	2.8%	5.9%	19.3%	15.6%	5.4%	2.3%
Placebo	7.2%	5.8%	6.6%	9.2%	6.5%	3.4%

*The table summarizes the data on treatment/study discontinuation rates of the commercial dose in phase 2 trials of RRMS drugs. If the commercial dose was not included in the phase 2 trials, the dose closest to the commercial dose was shown. This high-level comparison is provided for illustrative purposes only, is based on publicly available data and does not purport to be a comprehensive comparison or depiction of the other trials. Larger data sets than presented in this presentation are publicly available for certain of the compounds included on this slide. Please note that these results are taken from placebo-controlled trials, and these medications have not been tested in head-to-head assessments.

[1] Comi et al. Ann Neurol. 2001;49(3):290-297 [2] O'Connor et al. Neurology. 2006;66(6):894-900 [3] Kappos et al. Lancet. 2008;372(9648):1463-1472 [4] Kappos et al. N Engl J Med. 2006;355(11):1124-1140 [5] Cohen JA, Arnold DL, Comi G, et al. Lancet Neurol. 2016;15(4):373-381; QD: quaque die = once-daily; TID: ter in die = three times daily; RRMS: relapsing-remitting multiple sclerosis

ENSURE: Ongoing Pivotal Phase 3 Trials in Relapsing MS

NCT05134441 & NCT05201638



Coordinating Investigator

Robert J. Fox, M.D.
Cleveland Clinic



Included Patient Population: Relapsing Forms of MS

- Adult patients aged 18 to 55 years
- Established diagnosis of MS (revised McDonald criteria 2017)
- Confirmed relapsing MS (1996 Lublin criteria^[1])
- Active disease as defined by Lublin 2014
- EDSS score at screening between 0 to 5.5

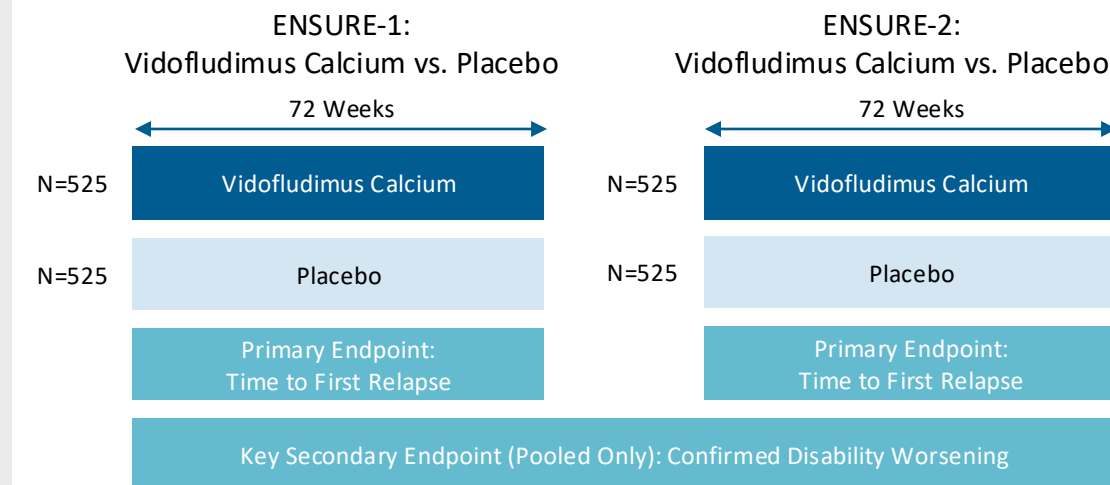
[1] Lublin FD, et al. Neurology. 2014;83(3):278-286

MS: multiple sclerosis; EDSS: Expanded Disability Status Scale; QD: quaque die = once-daily



Two Multicenter, Randomized, Double-Blind Phase 3 Trials

- More than 100 sites in 15 countries in each trial, including the United States, India, Middle East and North Africa, Latin America, Central and Eastern Europe
- Randomization to 30 mg vidofludimus calcium or placebo QD
- Positive interim analysis: Unblinded IDMC recommended continuing trial without changes, including no need for a potential upsizing
- Enrollment completed: 1,121 patients in ENSURE-1; 1,100 patients in ENSURE-2
- Top-line data for both ENSURE trials expected by end of 2026





Vidofludimus Calcium in Multiple Sclerosis (MS)

Development in Progressive Multiple Sclerosis (PMS)

Huge Unmet Medical Need Exists in PPMS, An Underdiagnosed and Tougher to Treat Patient Population

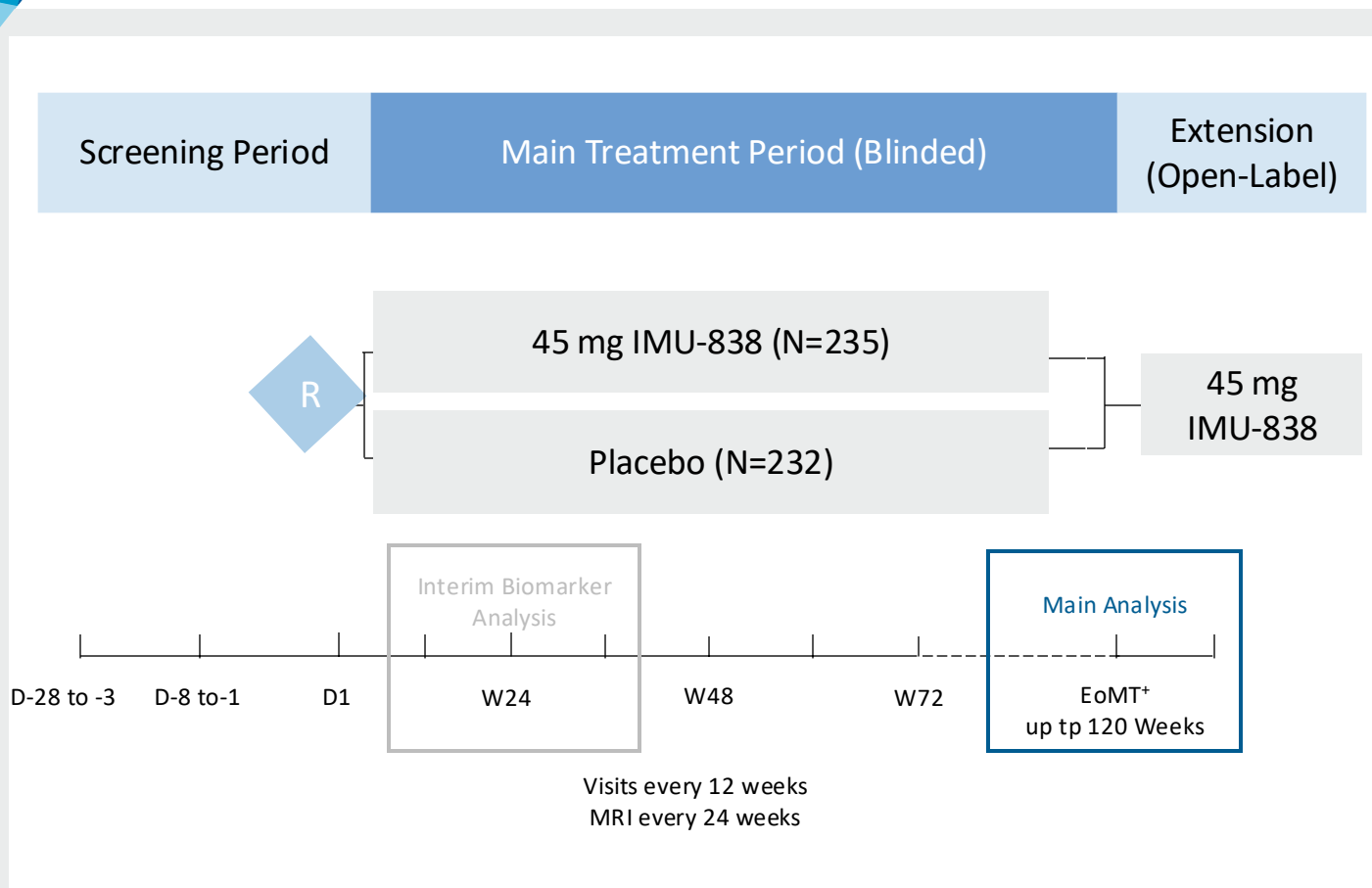


- PPMS, which affects **10-15% of people diagnosed with MS**, is characterized by a steady worsening of neurological function from the beginning of the disease, without distinct relapses or periods of remission
- Compared with RMS, PPMS is clinically associated with **greater symptom severity** and **functional impairment**, **higher rates of unemployment** and **hospitalization**, **greater economic burden**, and a more **substantial impact on health-related quality of life**
- **~120,000 patients diagnosed** (US & EU5), of which only ~54,000 (45%) are currently treated by disease-modifying therapies
- **Underdiagnosed and undertreated**, due to lack of safe, effective and convenient treatments (only one approved therapy)

PPMS: primary progressive multiple sclerosis; RMS: relapsing multiple sclerosis / Gross HJ, Watson C. Neuropsychiatr Dis Treat. 2017;13:1349–1357; National Multiple Sclerosis Society website: <https://www.nationalmssociety.org/understanding-ms/what-is-ms/types-of-ms/primary-progressive-ms>; Patient numbers sourced via internal Immunic analysis and 2024 Multiple Sclerosis Landscape and Forecast report by Decision Resources Group Part of Clarivate; EU5 countries: France, Germany, Italy, Spain, and United Kingdom

CALLIPER: Phase 2 Clinical Trial in Progressive Multiple Sclerosis

NCT05054140



Coordinating Investigator: Robert J. Fox, M.D., Cleveland Clinic



Multicenter, Randomized, Double-Blind, Placebo-Controlled Phase 2 Trial

- 467 adult patients, aged 18 to 65 years, enrolled at more than 70 sites in North America, Western, Central and Eastern Europe
 - PPMS or SPMS diagnosis (revised McDonald criteria 2017)
 - EDSS score at screening between 3.0 to 6.5
 - No relapse in last 24 months before randomization
 - Evidence of disability progression
- Randomization to 45 mg vidofludimus calcium or placebo QD
- Blinded main treatment period up to 120 weeks
- Optional, approximately 8-year, open-label extension period

EoMT: end of main treatment period, either at Week 120 or when last enrolled patient reached Week 72

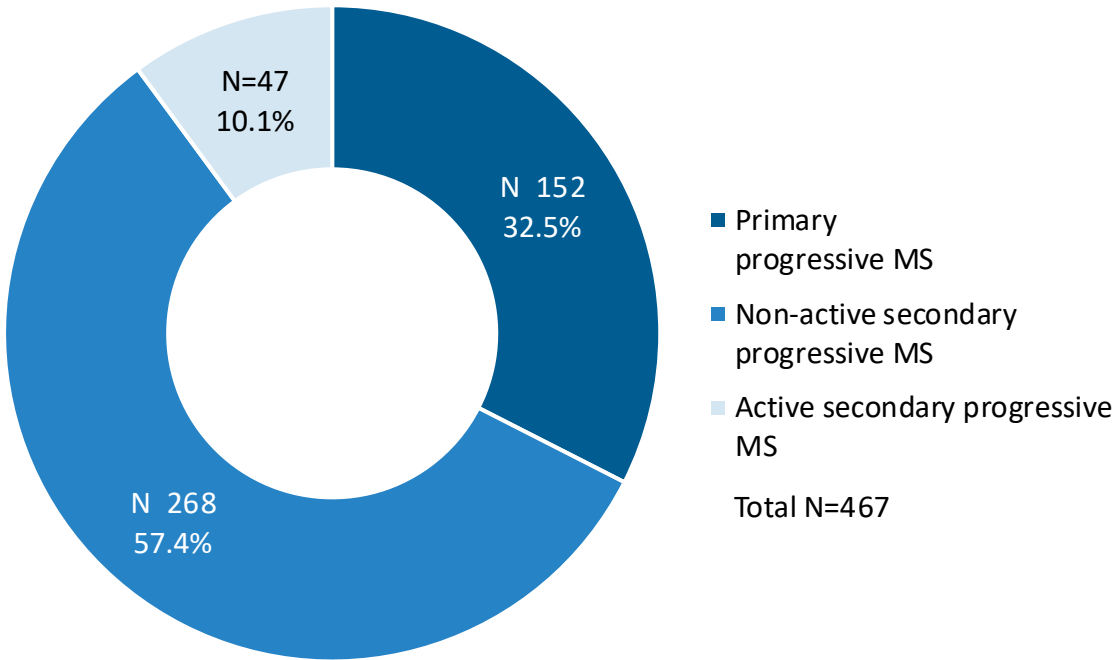
R: randomization; D: day; W: week; EoMT: end of main treatment period; MRI: magnetic resonance imaging; PPMS: primary progressive multiple sclerosis; SPMS: secondary progressive multiple sclerosis; EDSS: Expanded Disability Status Scale; QD: quaque die = once-daily

CALLIPER: Patient Demographics and Baseline Characteristics

Total Study Population of 467 Enrolled Patients



Progressive Disease Subtypes



Baseline Characteristics

Baseline Patient Characteristics	Total (N=467)
Age [years], median (min-max)	51.0 (21-65)
Gender (n and % female)	302 (64.7%)
Race (n and % White)	460 (98.7%)
BMI [kg/m^2], median (min-max)	24.85 [14.0 - 46.6]
SDMT [points], median (min-max)	40 [8-80]
EDSS at Visit 1, median (min-max)	5.5 [2.5-6.5]
MS relapses during last 24 months, median (min-max)	0.0 [0-1]
Gd+ lesions at baseline MRI (%)	16.3%
Treatment duration, median	589 days

Baseline characteristics initially assessed by the investigators when patients entered screening based on history. These data summarize the disease subtype as assessed per diagnosis at screening visit 1. A small number of patients changed their subtype (in particular from non-active to active disease) due to events during the screening period. Definition non-active SPMS (according to CALLIPER protocol): no evidence of relapse in the last 24 months before randomization, AND patients showing no evidence of Gd+ MRI lesions in the brain or spinal cord in the last 12 months; definition nonrelapsing SPMS: no evidence of relapse in the last 24 months before randomization / BMI: body mass index; SDMT: Symbol Digit Modalities Test; EDSS: Expanded Disability Status Scale; Gd+: gadolinium-enhancing; MRI: magnetic resonance imaging; N: number of patients evaluated



CALLIPER successfully demonstrated the neuroprotective potential of vidofludimus calcium in PMS patients

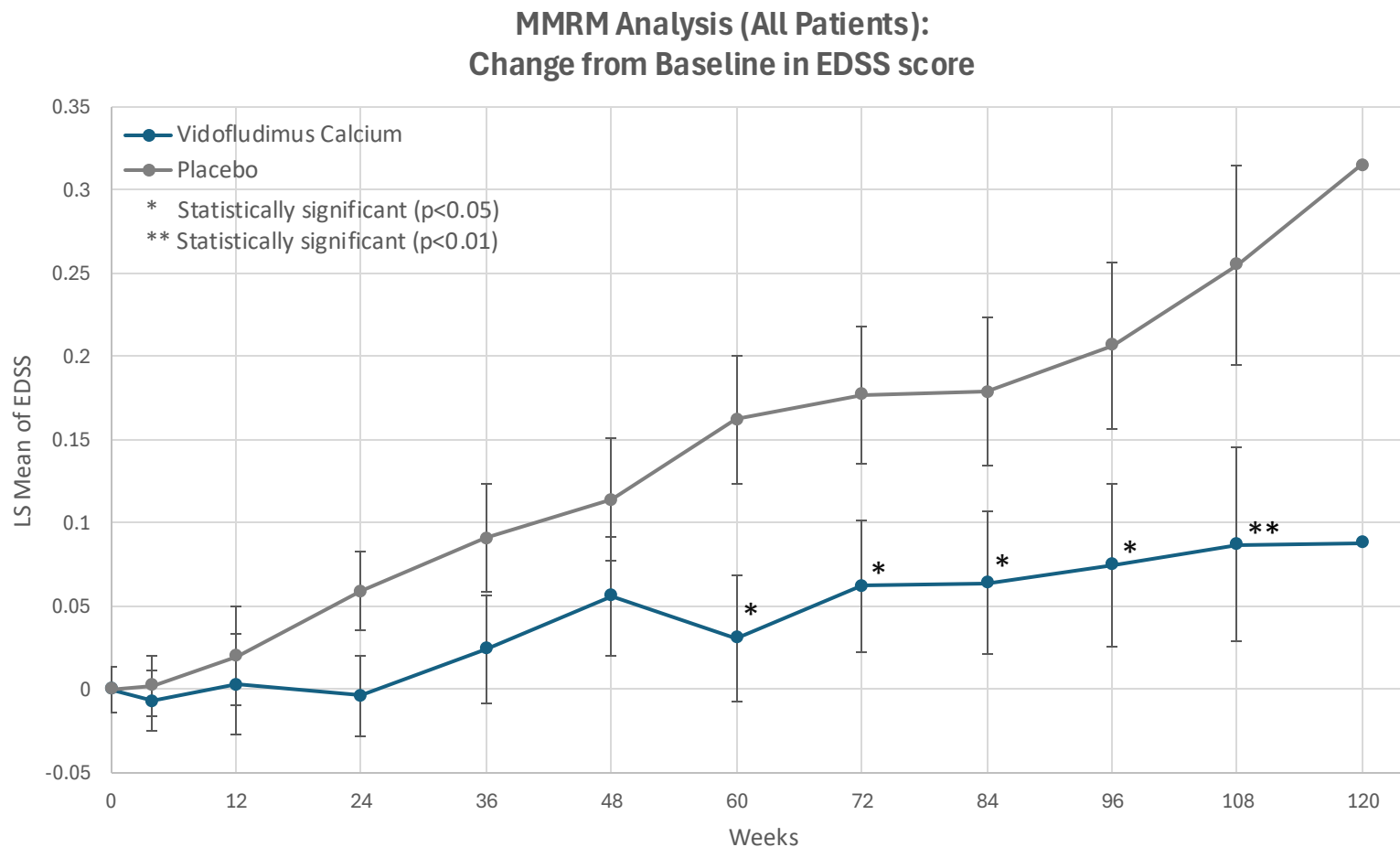
Clinically meaningful risk reduction of confirmed disability worsening of 24% in overall PMS population and even more prominent 31% reduction in PPMS population

CALLIPER: Vidofludimus Calcium Reduced **Time to 24wCDW** in Overall Study Population and All Subtypes Compared to Placebo

	Overall CALLIPER Patient Population (N=467)	PPMS (N=152)	naSPMS (N=268)	aSPMS (N=47)
HR (Kaplan Meier)	0.762	0.687	0.808	0.663
95 % CI	[0.479; 1.210]	[0.330; 1.430]	[0.418, 1.564]	[0.315, 2.974]
p-value	0.249	0.315	0.527	0.592
Risk Reduction of tt24wCDW	23.8%	31.3%	19.2%	33.7%

24wCDW: 24-week confirmed disability worsening; tt: time to; PPMS: primary progressive multiple sclerosis; naSPMS: non-active secondary progressive multiple sclerosis; aSPMS: active secondary progressive multiple sclerosis; HR: hazard ratio; CI: confidence interval / based on intent-to-treat population (ITT), patients analyzed as randomized; 24-week confirmed disability worsening based on EDSS scale (24wCDW), total of 73 events for 24wCDW based on EDSS, 70 events observed and 3 events imputed after 12-week confirmation before end of study (performed as sensitivity analysis); data displayed for the reduction of risk of time to 24wCDW; disease subtype as per diagnosis at screening visit 1

CALLIPER: Vidofludimus Calcium Significantly Reduced EDSS Increase from Baseline Compared with Placebo



EDSS: expanded disability status scale; MMRM: mixed models for repeated measure; LS: least square / MMRMs analysis: For the calculation of LS means based on the MMRM, patients with baseline and at least one post-baseline visit are considered. Missing values are calculated based on the analysis set. Estimates are adjusted for stratification factors used at baseline randomization (disease type and baseline EDSS value). 2-sided p-value is presented. Error bars show the standard error of the LS Mean. Data are based on group level analysis for overall CALLIPER population, vidofludimus calcium N=235, placebo N=232.

Comparison CALLIPER Versus ORATORIO Trials in PPMS Population

	ORATORIO*	CALLIPER
	(N=732)	(N=152)
Mean Age (Years)	44.6	47.4
Female (N,%)	361 (49.3%)	93 (61.1%)
EDSS - Mean	4.7	4.9
EDSS - Median	4.5	4.5
Gd+ Lesions at Baseline MRI (N,%)	26.6%	17.8%
Relative Risk Reduction of Time to 24wCDW Active Over Placebo	25%	31%

* Clinical Review Report: Ocrelizumab (Ocrevus): (Hoffmann-La Roche Limited): Indication: Management of adult patients with early primary progressive multiple sclerosis as defined by disease duration and level of disability, in conjunction with imaging features characteristic of inflammatory activity [Internet]. Ottawa (ON): Canadian Agency for Drugs and Technologies in Health; 2018 May. Results. Available from: <https://www.ncbi.nlm.nih.gov/books/NBK533357/>
PPMS: primary progressive multiple sclerosis; EDSS: Expanded Disability Status Scale; Gd+: gadolinium-enhancing lesions found on T1-weighted MRI images; MRI: magnetic resonance imaging; tt24wCDW: time to 24-week confirmed disability worsening; N: number of patients evaluated

CALLIPER: Vidofludimus Calcium Substantially Reduced 24wCDW in Patients Without Gd+ Lesions at Baseline

Time to 24wCDW With No Evidence of Gd+ Lesions at Baseline

Group	Number of Patients	HR	95 % CI	p-value	RRR tt24wCDW
All Patients	391	0.663	[0.394, 1.115]	0.120	33.7%
PPMS	125	0.656	[0.294; 1.464]	0.303	34.4%
naSPMS	250	0.702	[0.346, 1.422]	0.326	29.8%



- Precisely the patients who were largely shown to **not benefit from current anti-inflammatory therapies**
- These **clinical effects underlines neuroprotective effect** of Nurr1 activation by vidofludimus calcium

24wCDW: 24-week confirmed disability worsening based on the EDSS (expanded disability status scale) score; tt: time to; Gd+: gadolinium-enhancing; HR: hazard ratio; CI: confidence interval; RRR: relative risk reduction; disease subtype as per diagnosis at screening visit 1
24wCDW is defined as patients with worsening in EDSS sustained for at least 22 weeks (154 days) given the visit window +/- 7 days. Confirmed disability progression event status was imputed for participants who completed the trial, met the criteria for confirmed disability progression sustained for at least 12 weeks, and continued to meet the criteria for disability progression according to the EDSS score through the final trial assessment but did not reach the 24-week confirmation visit. Total of 73 events for 24wCDW based on EDSS, 70 events observed and 3 events imputed after 12-week confirmation before end of study (performed as sensitivity analysis).

CALLIPER: Top-Line Data Confirmed Favorable Safety and Tolerability Profile of Vidofludimus Calcium Observed in Previous Clinical Trials

Number of Patients With Any TEAE and SAE

	Vidofludimus Calcium N=235	Placebo N=233
Any TEAE, n(%)	163 (69.4)	159 (68.5)
Any SAE n(%)	19 (8.1)	15 (6.5)

- No new safety signals identified
- Occurrence of TEAEs and SAEs with similar frequency in both treatment arms

Five Most Common TEAE Events

	Vidofludimus Calcium	Placebo	Total
Urinary tract infection	161	152	313
Upper respiratory infection	57	49	106
Headache	16	42	58
Back pain	11	24	35
Fall	15	17	32

Most Common SAE Events (all SAE with total incidence >1)

	Vidofludimus Calcium	Placebo	Total
Pyelonephritis	1	1	2
Femoral neck fracture	0	2	2
Femur fracture	0	2	2
Vertigo	2	0	2

TEAE: treatment-emergent adverse event; SAE: serious adverse event; N: number of patients evaluated

Safety Population contains any patient who received at least 1 dose of study drug, Vidofludimus calcium (N=235), Placebo (N=232), Total (N=467). All other SAE not listed had only single occurrences in the CALLIPER trial.

Positive Data from Phase 2 CALLIPER Trial of Vidofludimus Calcium in Progressive Multiple Sclerosis



- Clinically meaningful risk reduction of 24wCDW by 24% in overall study population; even more prominent 31% reduction in high unmet need population of PPMS
- Remarkable 34% reduction of 24wCDW in patients without baseline inflammatory lesions in overall study population
- Confirmed favorable safety and tolerability observed in previous clinical trials; no new safety signals identified
- As of April 2025, more than 375 patients continue to be treated in open-label extension phase of CALLIPER trial
- Underlines Nurr1 activation as new mode of action for preventing neurodegeneration in MS and substantiates impact on disability accumulation by both PIRA and RAW
- Further de-risks ongoing phase 3 ENSURE program with potential to offer relapsing MS patients an oral, safe and neuroprotective treatment early in the disease

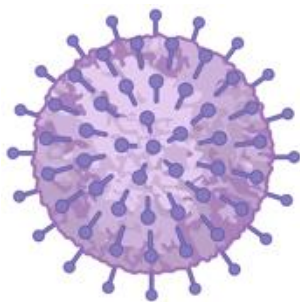
Vidofludimus Calcium: DHODH Inhibition Provides Broad-Spectrum Antiviral Activity Against Different Pathogenic Viruses



Inhibits Epstein-Barr Virus (EBV)
Replication and Reactivation

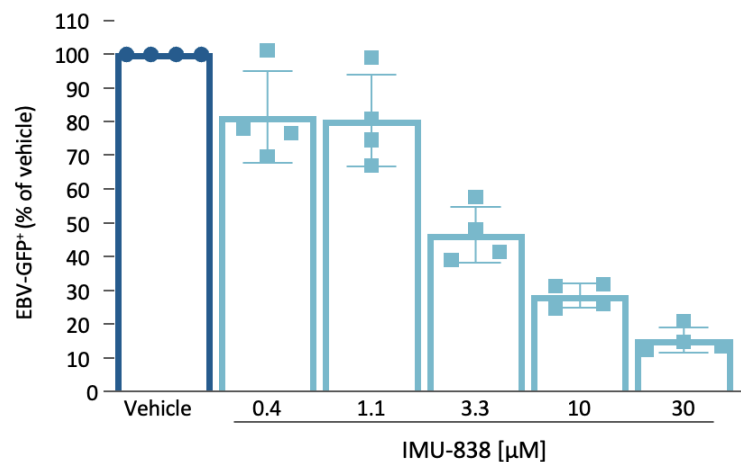
By targeting the host cell metabolism, vidofludimus calcium has shown to be active against different RNA and DNA viruses *in vitro*

- Shows antiviral activity with EC₅₀ values in single digit μM range
- Including strong anti-EBV activity



Showed Dose-Dependent
Inhibition of EBV Reactivation

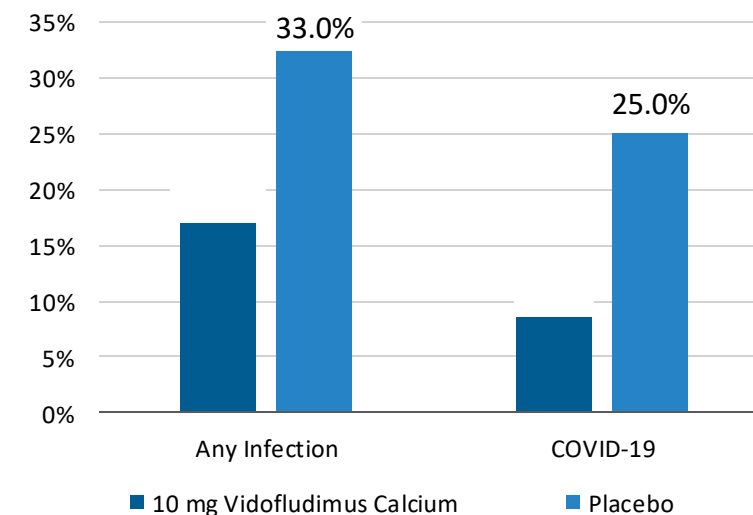
Anti-Akata-BX1-EBV-GFP stimulated with hIgG



Decreased Number of Opportu-
nistic SARS-CoV-2 Infections

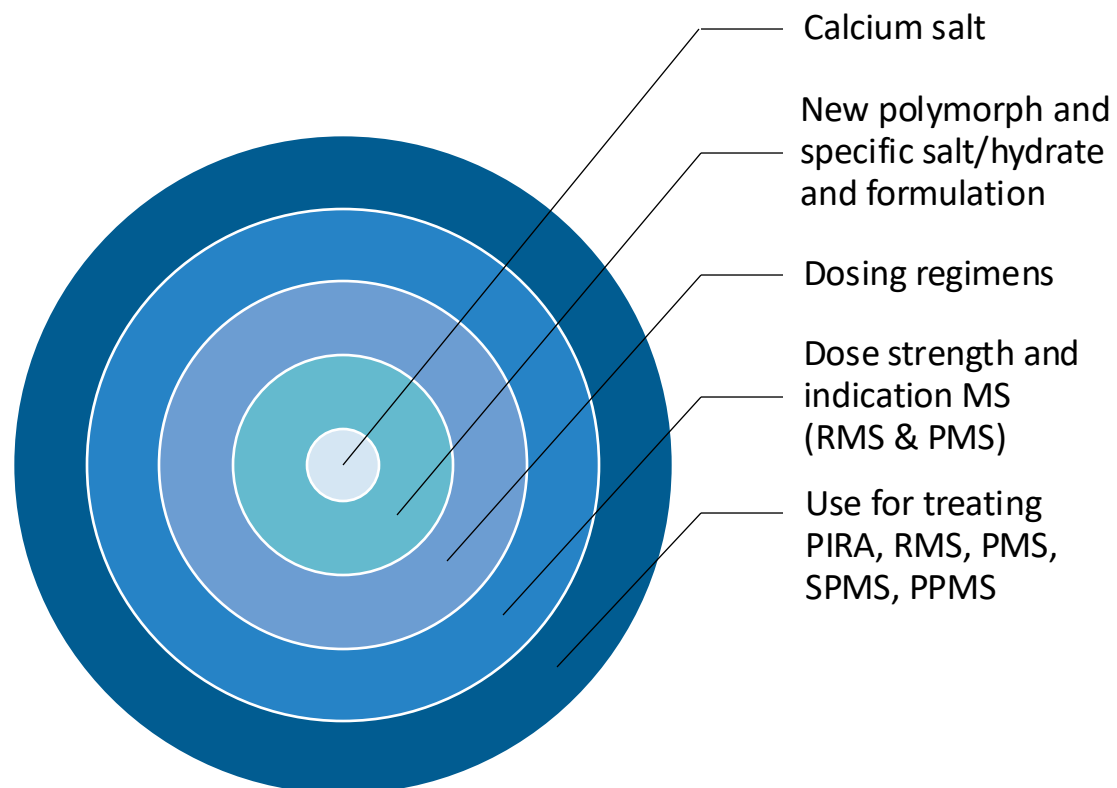
Vidofludimus calcium showed interesting hints for clinical anti-SARS-CoV-2 activity in the phase 2 EMPHASIS trial in RRMS

- Number of reported COVID-19 cases Cohort 2:



Left: Eur J Clin Invest. 2020;50:e13366 / middle: Marshall et al., Poster ECTRIMS 2021 / right: Immunic data; DHODH: dihydroorotate dehydrogenase; RNA: ribonucleic acid; DNA: deoxyribonucleic acid; EC50: half-maximal effective concentration; EBV: Epstein-Barr virus; hIgG: human immunoglobulin G; SARS-CoV-2: severe acute respiratory syndrome coronavirus; COVID-19: coronavirus disease 2019; RRMS: relapsing-remitting multiple sclerosis

Several Layers of Patents Protecting Vidofludimus Calcium



Eight Independent Patent Families Protecting Vidofludimus Calcium

- IP for superior calcium salt and specific polymorph of the drug product
 - Additional patent directed to specific polymorph matching the only polymorph in the drug product granted in the US and other jurisdictions
- Broad IP for all salts directed to dosing regimens, covers all label-relevant dosing schemes, granted in the US and Japan
- Dose strengths subject of another granted patent in the US
- Use of vidofludimus for treating PIRA as well as other neurodegenerative diseases, also including biomarker-based subgroups, filed in 2023
- Another level of protection expected by data exclusivity based on vidofludimus calcium's classification as New Chemical Entity (NCE)

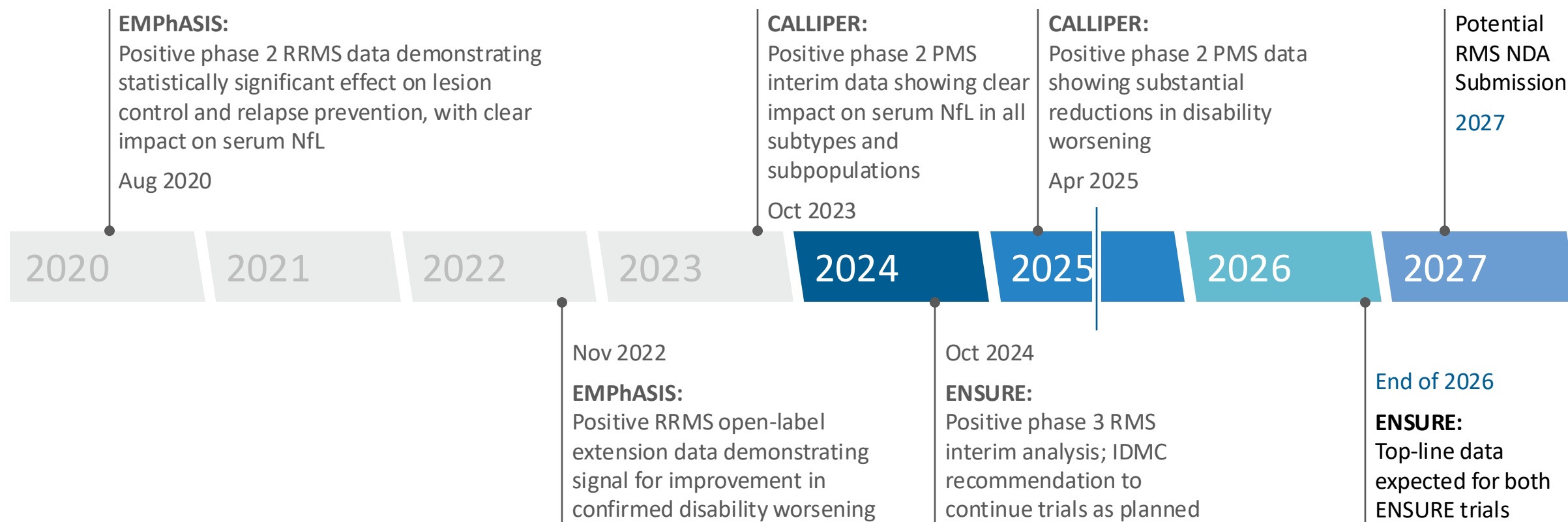


Patent portfolio expected to provide exclusivity into 2041 in the US, unless extended further

IP: intellectual property; MS: multiple sclerosis; RMS: relapsing MS; PMS: progressive MS; SPMS: secondary progressive MS; PPMS: primary progressive MS; PIRA: progression independent of relapse activity

Vidofludimus Calcium in Multiple Sclerosis

Consistent and Differentiated Results to Date Support Straightforward Path Towards Potential Regulatory Approvals



Although we currently believe that each of these goals is achievable, they are each dependent on numerous factors, most of which are not under our direct control and can be difficult to predict. We plan to periodically review this assessment and provide updates of material changes as appropriate. / MS: multiple sclerosis; RRMS: relapsing-remitting MS; RMS: relapsing MS; PMS: progressive MS; NfL: neurofilament light chain



IMU-856

Restoring a Healthy Gut through Renewal of the Bowel Wall

IMU-856 Targets Physiological Intestinal Epithelial Regeneration and Restoration of Gut Cell Function



- Innovative oral therapeutic approach potentially applicable to a broad range of gastrointestinal disorders



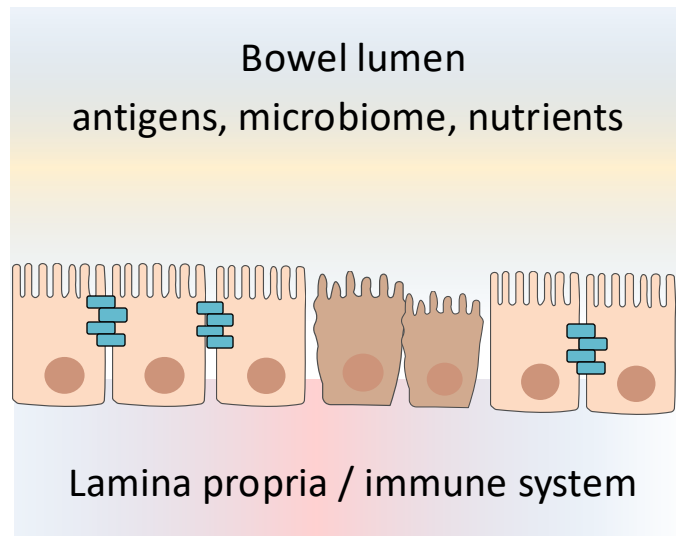
- Targets physiological intestinal epithelial regeneration, including gut hormon-producing cells



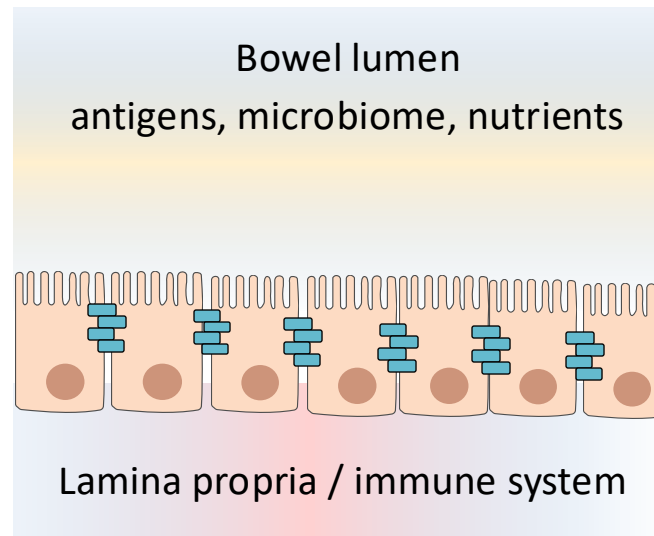
- Designed to strengthen gut wall integrity and function without immunosuppression

Once-Daily, Oral IMU-856 Aims to Regenerate the Gut Wall and Barrier Function by a New Innovative Targeted Mechanism

Damaged Gut Wall



Healthy Gut Wall



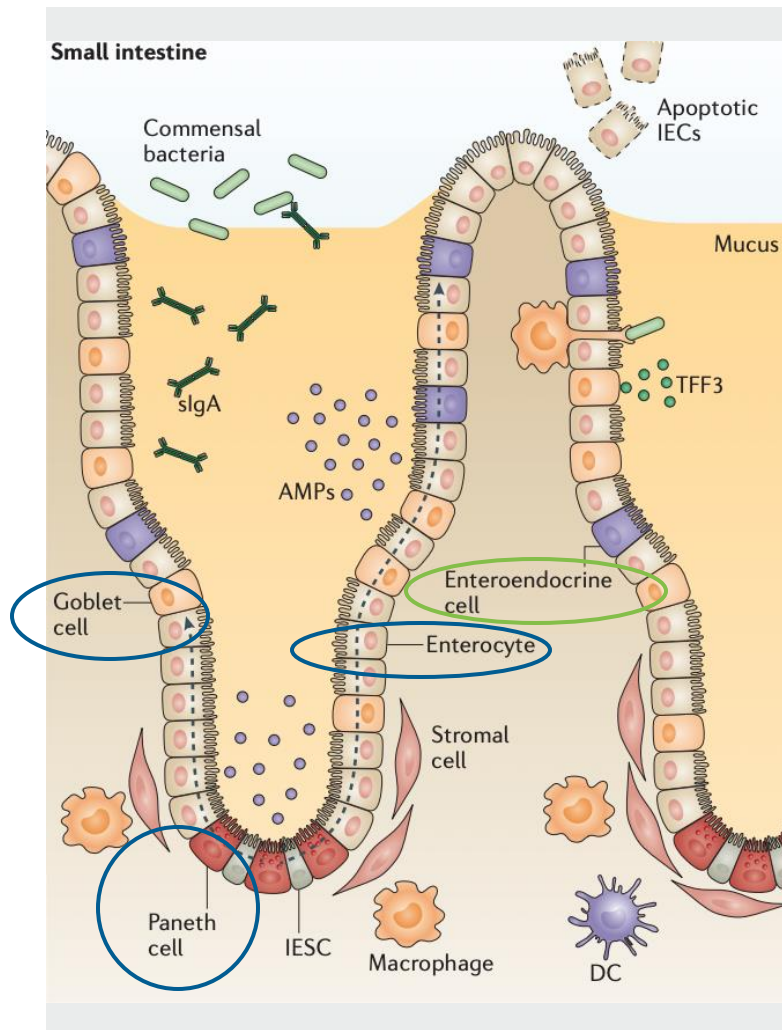
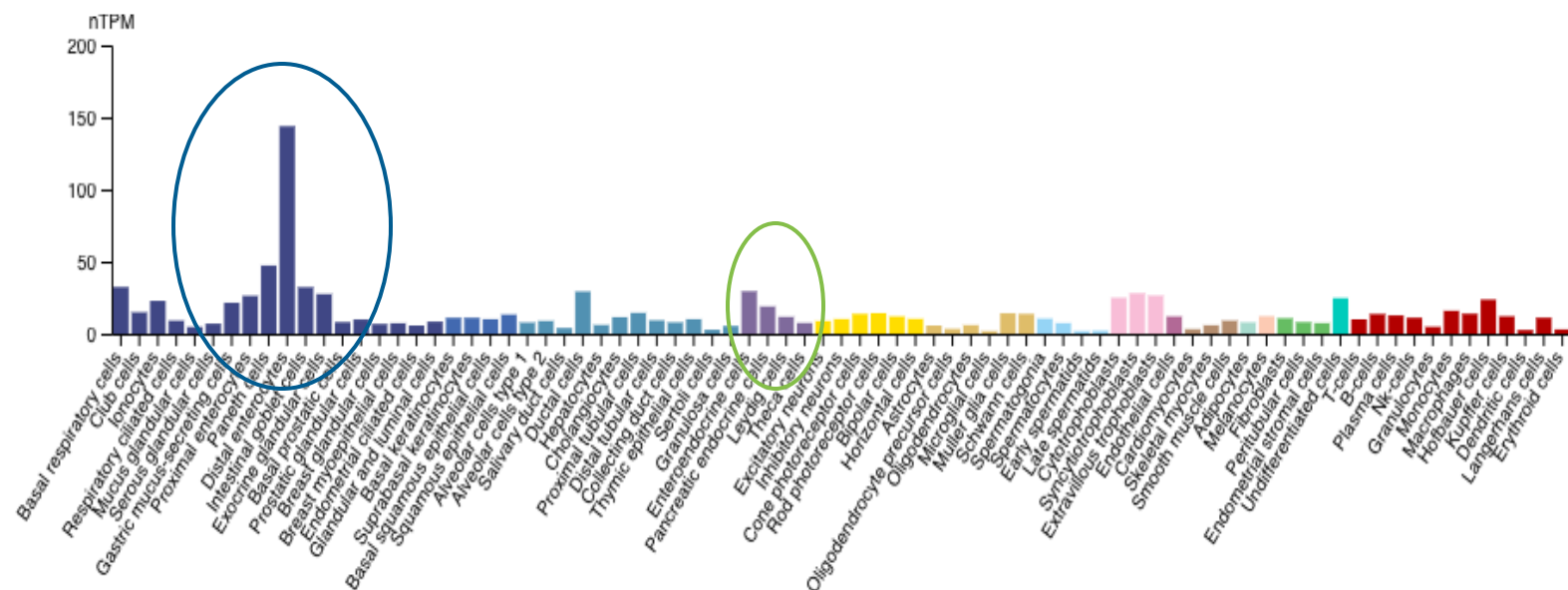
IMU-856:

- First-in-class modulator of sirtuin 6 (SIRT6), targets physiological intestinal epithelial regeneration and restoration of barrier function
- Provides protection and enhances transport of nutrients
- This new approach avoids immunosuppression

SIRT6 Target Is Highly Expressed in Gut Epithelial Cells



Highest mRNA Expressions in Paneth Cells, Enterocytes, Goblet Cells and Enteroendocrine Cells such as L-Cells



Left: <https://www.proteinatlas.org/> // Right: Peterson, L., Artis, D. Nat Rev Immunol 14, 141–153 (2014)
SIRT: sirtuin; mRNA: messenger ribonucleic acid; nTPM: normalized transcript per million

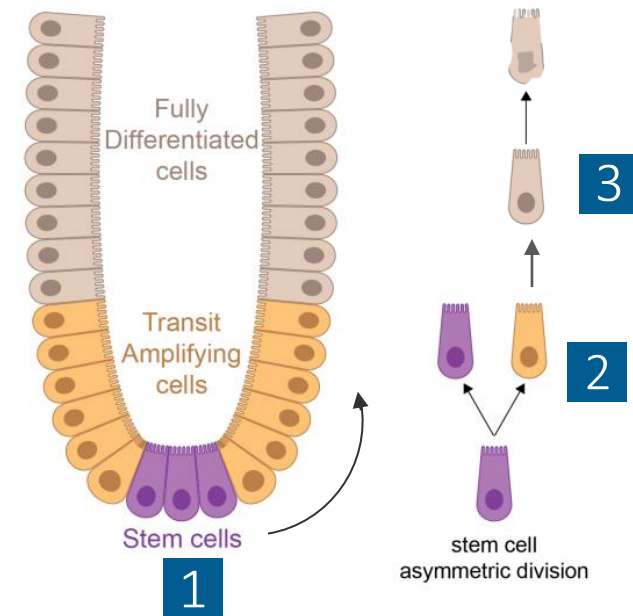
IMU-856 Enhances the Natural Regenerative Process in the Gut

Gut wall renewal is a normal physiological process

1. Regeneration begins in the crypts, where intestinal stem cells are located
2. Stem cells undergo asymmetric division thereby producing fully differentiated epithelial gut cells and renewing intestinal stem cells
3. These new epithelial cells are renewing the lining of crypts and villi to maintain healthy gut and proper intestinal barrier

→ IMU-856 is an epigenetic regulator which enhances this natural tissue renewal phenotype

Asymmetric cell division renews stem cells and regenerates the gut wall



Adapted from Mamis K et al., Proc. R. Soc. B. 290:20231020 (2023)



IMU-856: Additional Pharmacological Effect

Dose-Dependent Increase of GLP-1 in Patients

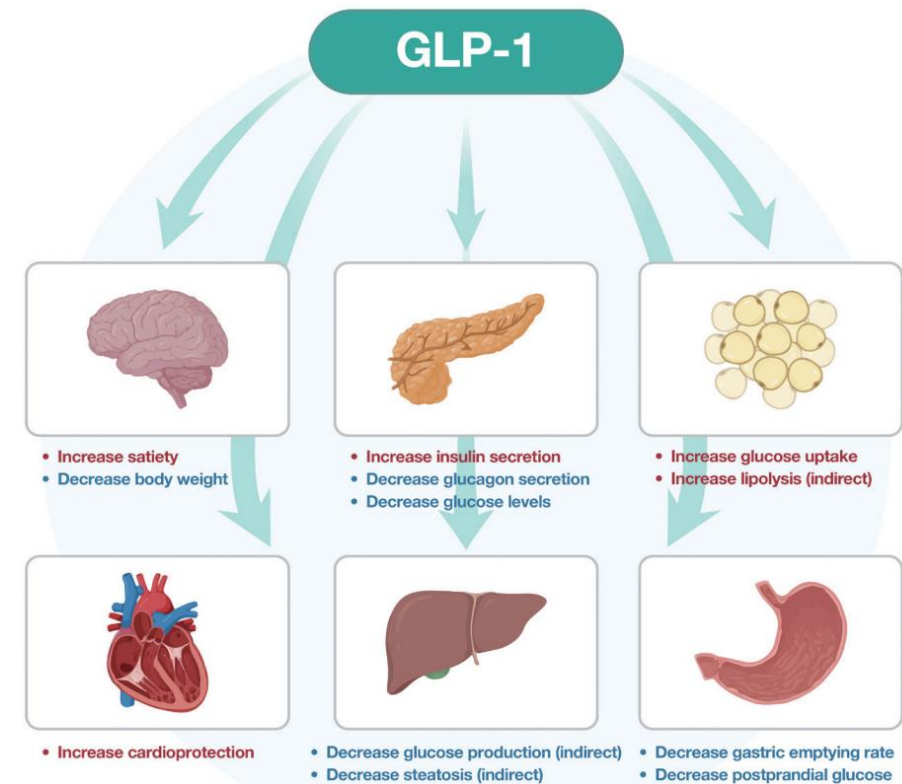
Intestinal Production of GLP-1 Mediates Effects on Body Weight



GLP-1: Glucagon-Like Peptide-1

- Peptide hormone generated through enzymatic breakdown of proglucagon
- Endocrine hormone, secreted by **enteroendocrine L-cells** located in the distal jejunum, ileum, and colon in response to **nutrient ingestion** and neuroendocrine stimulation
- Typical physiological **increase** in GLP-1 levels in healthy humans **after a meal is 2-3 times**
- GLP-1 increase leads to slow gut motility, lower food intake, increase satiety and induce insulin secretion

Main Physiologic Effects of GLP-1



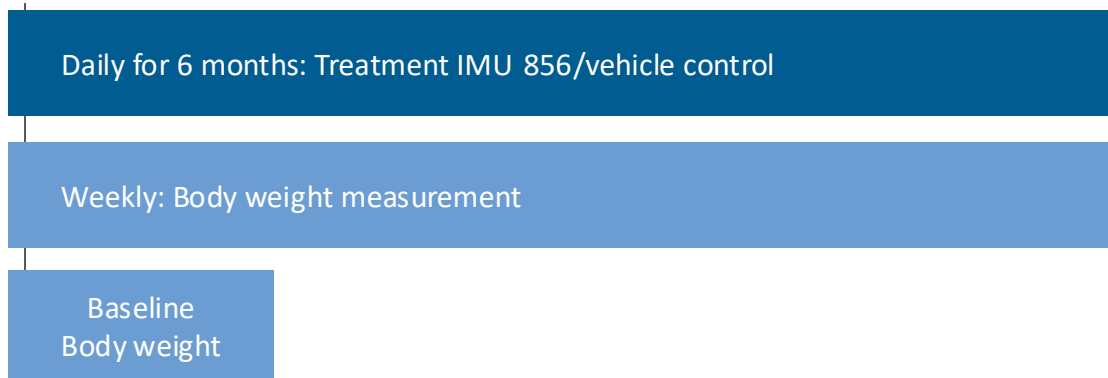
Left: Review Zheng, Z., Zong, Y., Ma, Y. et al. Sig Transduct Target Ther 9, 234 (2024); right: Jakubowska A, Roux CWL, Viljoen A. Endocrinol Metab (Seoul). 2024 Feb;39(1):12-22

IMU-856: Effects on Body Weight in Preclinical Experiment and on Blood GLP-1 Levels in Celiac Disease Clinical Trial



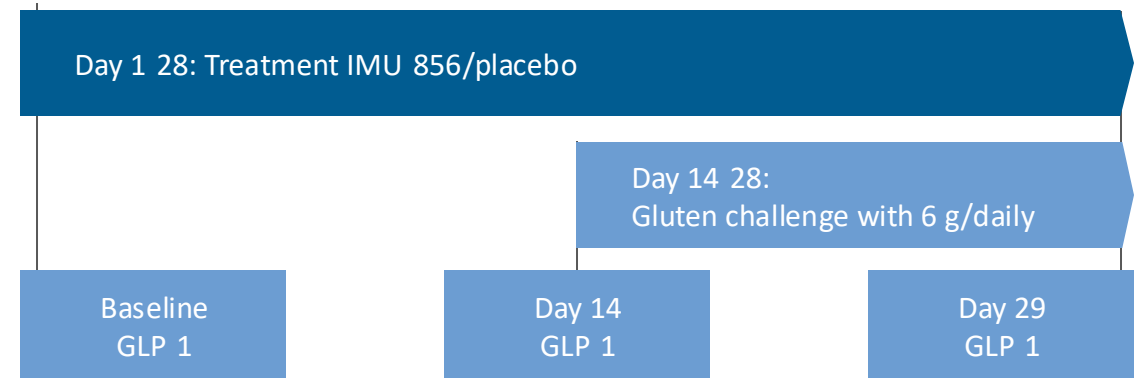
6-Months *In Vivo* Study

- Regulated GLP study^[1] to support clinical development
- Daily oral treatment of rats^[2] for 6 months
- Dosing: 0 (vehicle), 10, 25, 75 mg/kg/day of IMU-856
- Weekly body weight measurement



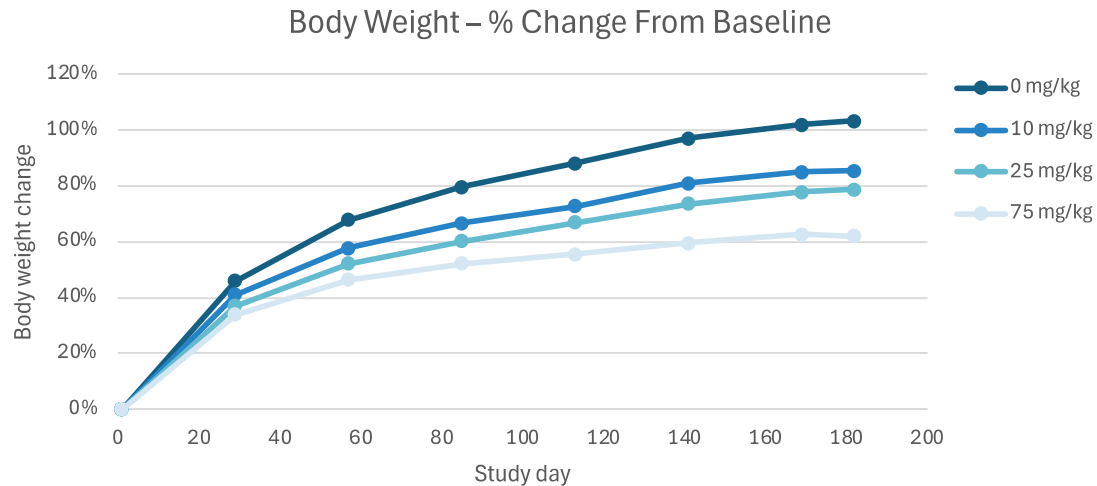
Phase 1b Clinical Trial of IMU-856

- Designed to explore effects of gluten challenge in a celiac disease patient population
- Total of 43 patients enrolled (IMU-856: N=29)
- Dosing: 80 and 160 mg QD of IMU-856, or placebo
- Double-blind treatment period of 28 days, 13 days without and 15 days with 6 g daily gluten challenge
- Patients measured post hoc for plasma GLP-1 concentrations



[1] according to ICH M3(R2) [2] Wistar Han rats / GLP-1: glucagon-like peptide-1; GLP: Good Laboratory Practice; QD: quaque die = once-daily; ICH: International Council for Harmonisation of Technical Requirements for Pharmaceuticals for Human Use

In a 6-Months *In Vivo* Study, IMU-856 Dose-Dependently Reduced Weight Gain



Reduction of weight gain in
% compared to vehicle control

10 mg/kg	-18 %
25 mg/kg	-23 %
75 mg/kg	-40 %

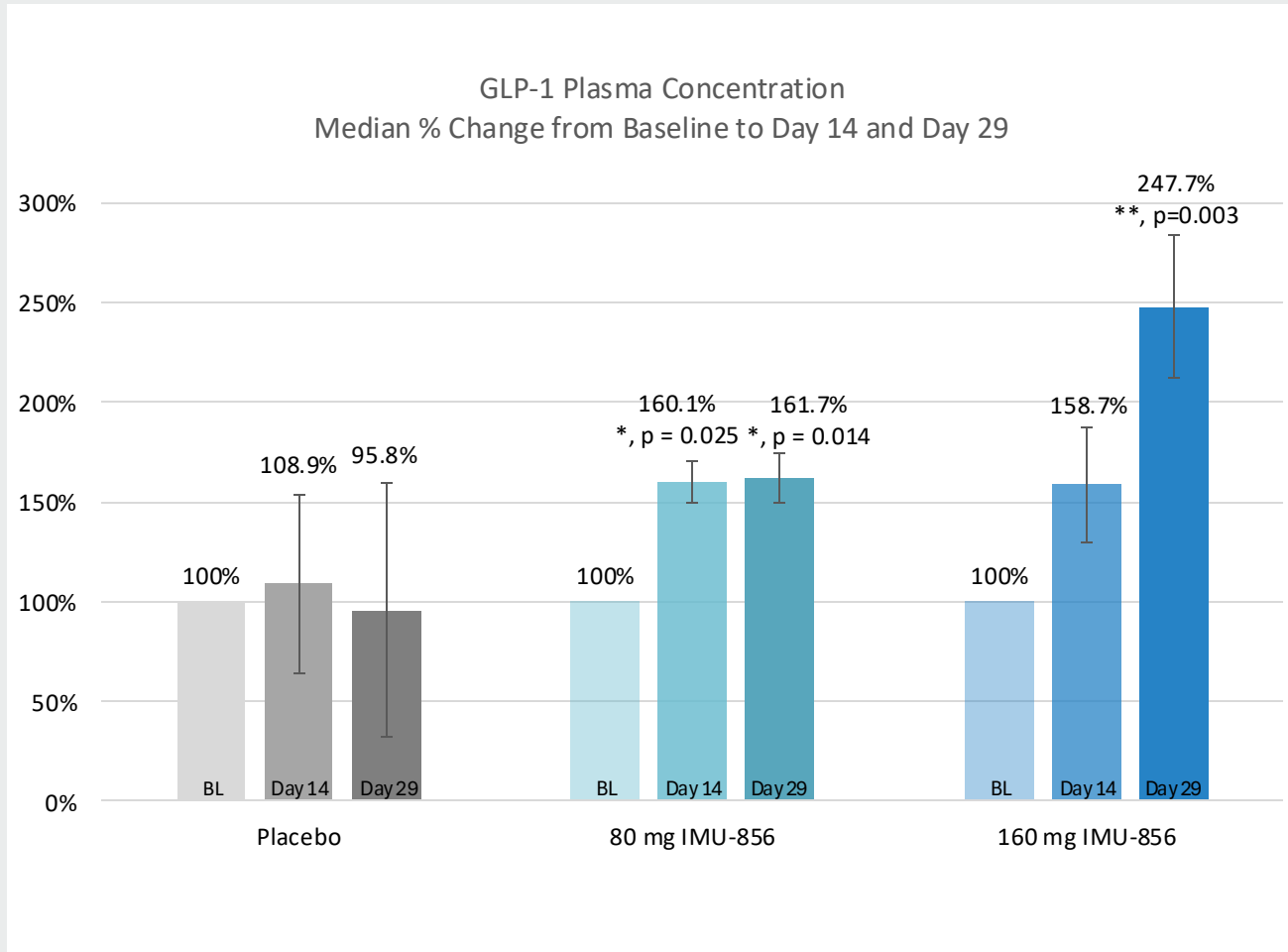
- Dose-dependent effect on body weight gain
- Linked to reduced food consumption
- Effect in both males and females
- No effect on general health condition



IMU-856 reduced body weight gain in a dose-dependent fashion up to -40 % compared to vehicle control

Reduced body weight gain observed in 6-month toxicology study. Rats were 7-8 weeks old at study start and were expected to gain weight over the course of the study. Data show less weight gain in IMU-856 treated animals in connection with reduced food consumption.

Confirmation of Effects as Part of Phase 1b Clinical Trial: IMU-856 Dose-Dependently Increased GLP-1 in Celiac Disease Patients



28-day phase 1b clinical trial of IMU-856 in celiac disease

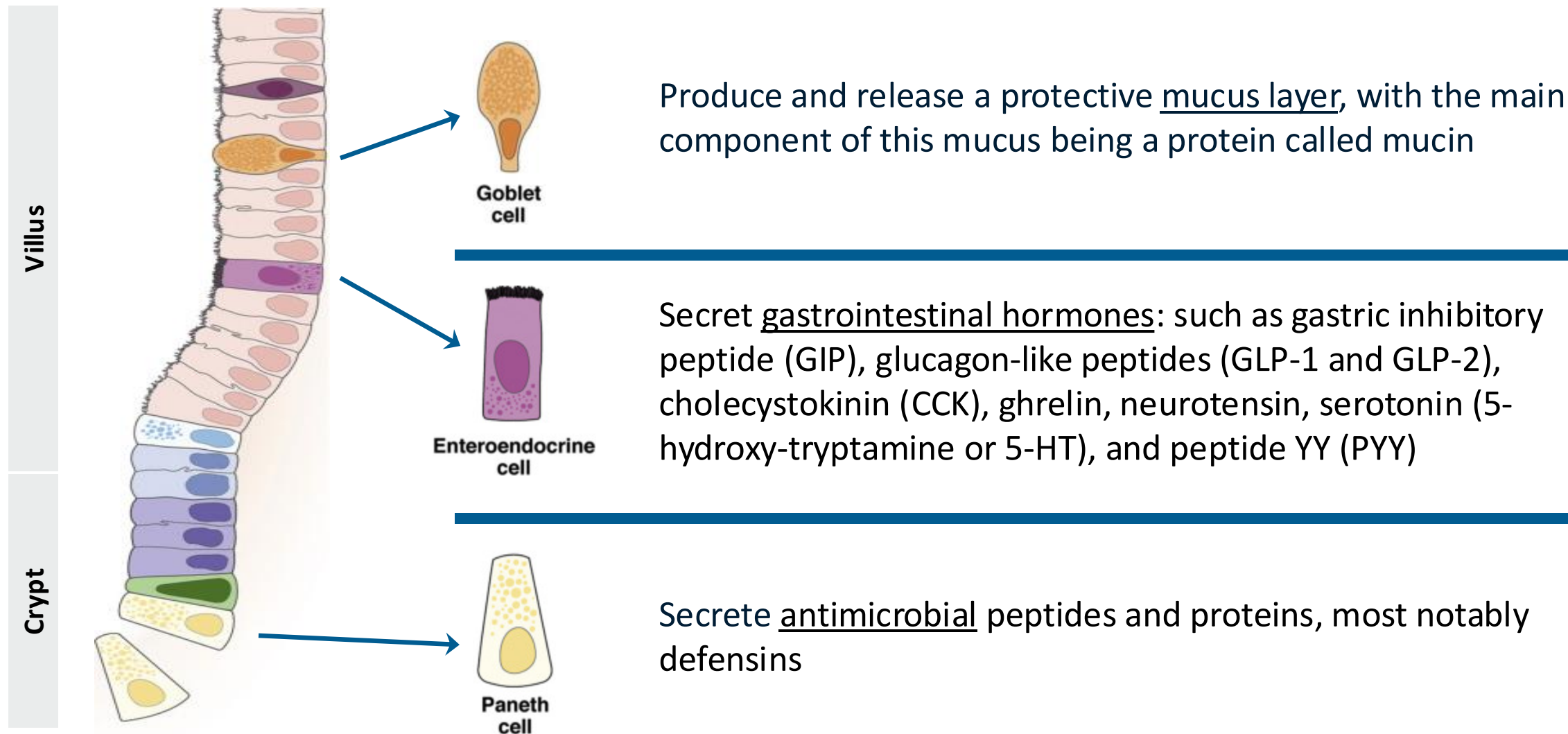
- Patients measured for plasma GLP-1 concentrations: N=11 (placebo), N=13 (80 mg IMU-856), N=13 (160 mg IMU-856)
- Baseline: Day 1, N=37 over all arms
- Day 14: before start of challenge, N=36
- Day 29: after last treatment on Day 28, N=32
- Morning baseline levels under fasting conditions

- **Dose-dependent increase of endogenous GLP-1 levels of up to 2.5 times versus placebo control**
- **Typical physiological increase in GLP-1 levels in healthy humans after a meal is also 2-3 times**

Statistics: two-sided Mann-Whitney U, treatment vs. placebo at Day 14 and Day 29 / GLP-1: glucagon-like peptide-1; BL: baseline

Main Secretory Epithelial Cells of the Small Intestine and Colon Epithelium

All Have Been Shown to Express SIRT6 Target



Meyer AR, Brown ME, McGrath PS, Dempsey PJ. Cell Mol Gastroenterol Hepatol. 2022;13(3):843-856 / SIRT: sirtuin

IMU-856: A Novel Mechanism Offering Potential to Go Beyond Existing GLP-1, GLP-2, GIP Mimetics



SIRT6 Targeting Approach IMU-856

- **Functional improvement** of enteroendocrine and other epithelial cells through increasing physiologic cell regeneration in gut wall
- Secretion of the **physiological GLP-1** protein and possible increase of secretion of **multiple incretins** (currently being investigated)
- Improvement of gut barrier and functionality in general
- **Oral** administration, small molecule



Incretin Mimetics GLP-1, GLP-2, GIP

- Providing **synthetic peptides** that mimic the natural hormones secreted by enteroendocrine cells
- Targets **one or two target incretins** only (at this point)
- **Injectable**, peptide

SIRT: sirtuin; GLP: glucagon-like peptide; GIP: glucose-dependent insulin-tropic polypeptide

Obesity Market Expected to Reach More Than \$170 Billion Globally by 2031^[1]



Unmet Needs Still Exist to Address This Growing Medical Challenge

- Obesity and overweight are among the fastest growing and most prevalent chronic human conditions in the world affecting ~2.5 billion adults worldwide^[2]
- The economic impact of obesity and overweight in the United States is estimated to be \$706 billion, increasing to \$2.6 trillion by 2060^[3]
- GLP-1 receptor agonist class has revolutionized obesity treatment but there are still **unmet needs for novel mode of actions, oral administration**, increased tolerability and greater efficacy
- Current drugs in development are mainly peptidomimetics – with challenges in oral administration

[1] GlobalData Pharma DECODED, Feb. 11th 2025 "Obesity: Seven-Market Drug Forecast and Market Analysis – Update" [2] <https://www.who.int/news-room/fact-sheets/detail/obesity-and-overweight#:~:text=In%202022%2C%202.5%20billion%20adults%20aged%2018%20years%20and%20older,1990%20to%2020%25%20in%202022>
[3] <https://data.worldobesity.org/economic-impact-new/countries/US.pdf>

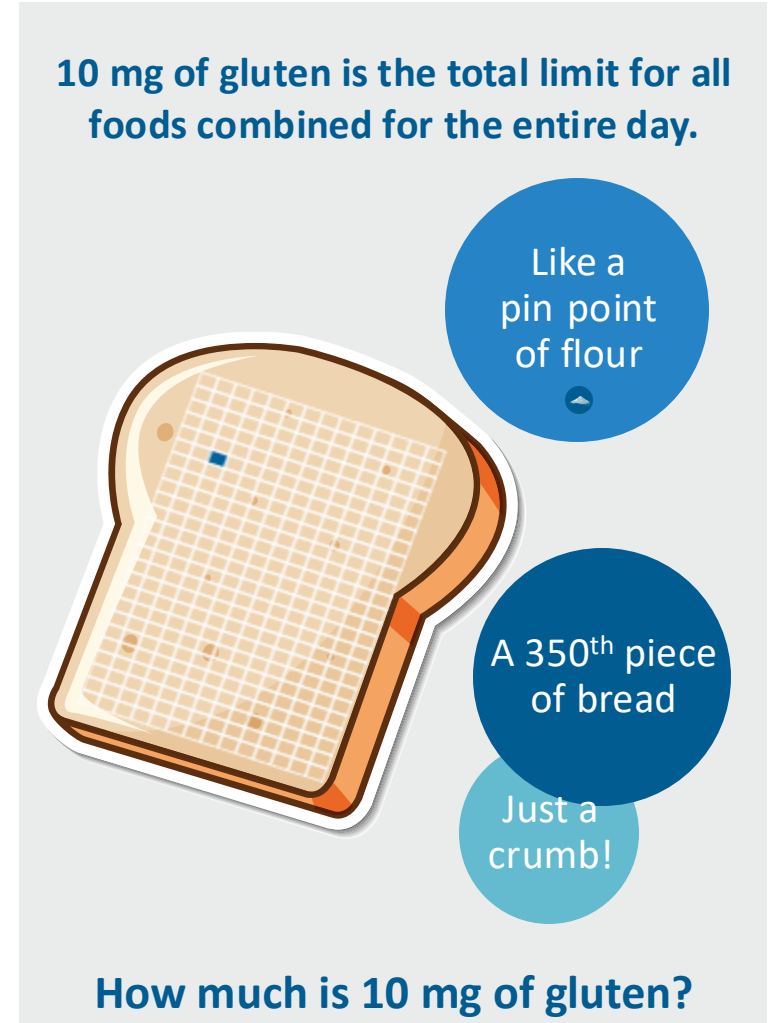


IMU-856 in Celiac Disease

Demonstrated Clinical
Proof-of-Concept in a
Phase 1b Clinical Trial

Celiac Disease Currently Has No Adequate Treatment Options

- Two million patients diagnosed with celiac disease in the US; more than one million more undiagnosed^[1,2]
- Most studies report between **24% and 47%**^[3-8] of patients with signs and symptoms of ongoing active celiac disease (OACD) **despite a gluten-free diet**, most likely due to continuous (inadvertent) gluten exposure
- **Only established therapeutic option is a life-long strict adherence to a gluten-free diet**^[9], which involves complete avoidance of proteins from wheat, barley, and rye
- Gluten challenge is an accepted concept for clinical trials in celiac disease



[1] Singh et al., Clinical Gastroenterology and Hepatology 2018;16:823–836 [2] Choung et al., Mayo Clin Proc. 2016 Dec 5:S0025-6196(16)30634-6 [3] Lebowitz et al., Aliment Pharmacol Ther. 2014 March ; 39(5): 488–495 [4] Lanzini et al., Aliment Pharmacol Ther. 2009; 29(12):1299–308 [5] Ciacci et al., Digestion. 2002; 66(3):178–85 [6] Selby et al., Scand J Gastroenterol. 1999; 34(9):909–14 [7] Rubio-Tapia et al., Am J Gastroenterol. 2010; 105(6):1412–20 [8] Sharkey et al., Aliment Pharmacol Ther. 2013; 38(10):1278–91 [9] <https://nationalceliac.org/celiac-disease-questions/understanding-gluten-levels/> (text and picture)

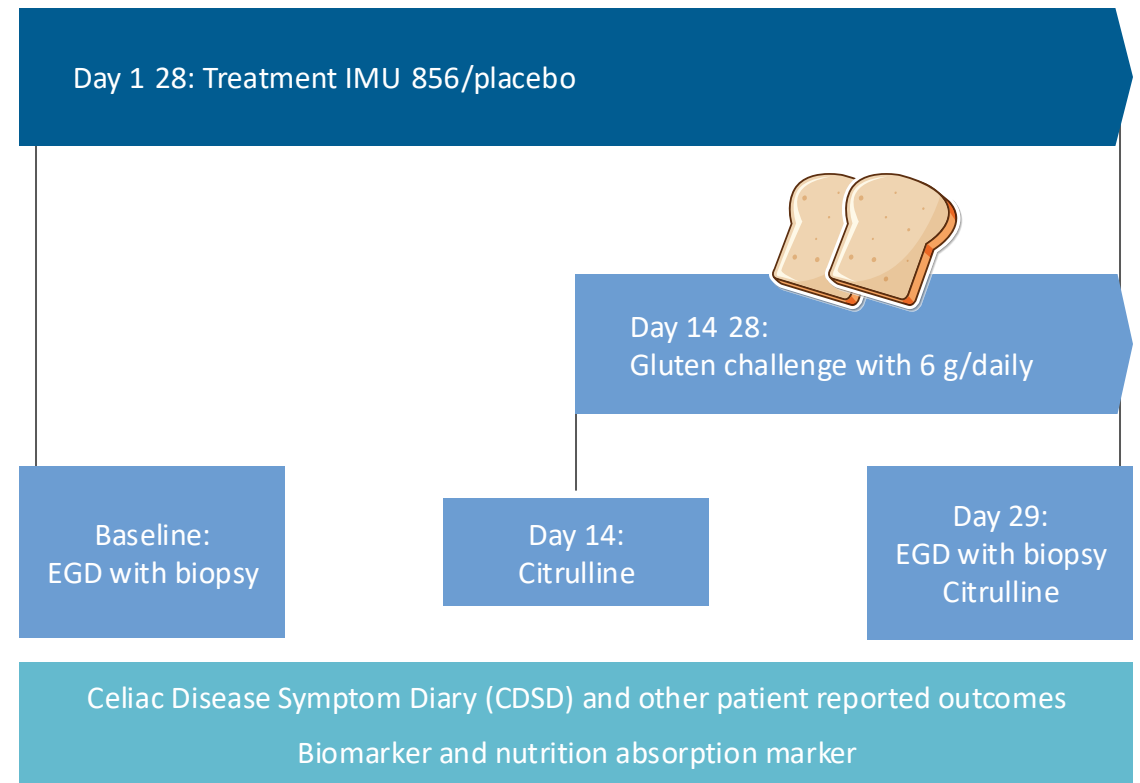
IMU-856 Demonstrated Clinical Proof-of-Concept in a Phase 1b Clinical Trial in Celiac Disease



Proof-of-Concept Study Designed as a Gluten Challenge Trial

- **Celiac disease used as disease model to provide clinical proof-of-activity of IMU-856 in a 28-day trial setting**
- Designed to explore effects of gluten challenge in a celiac disease patient population
- Dosing: 80 and 160 mg QD of IMU-856, or placebo
- 43 patients enrolled (IMU-856: N=29)
- Assessed safety, tolerability, pharmacokinetics, and pharmacodynamics of IMU-856
- Proof-of-concept: measured histological changes, blood biomarkers of epithelial mass, nutrient uptake and disease-related symptoms

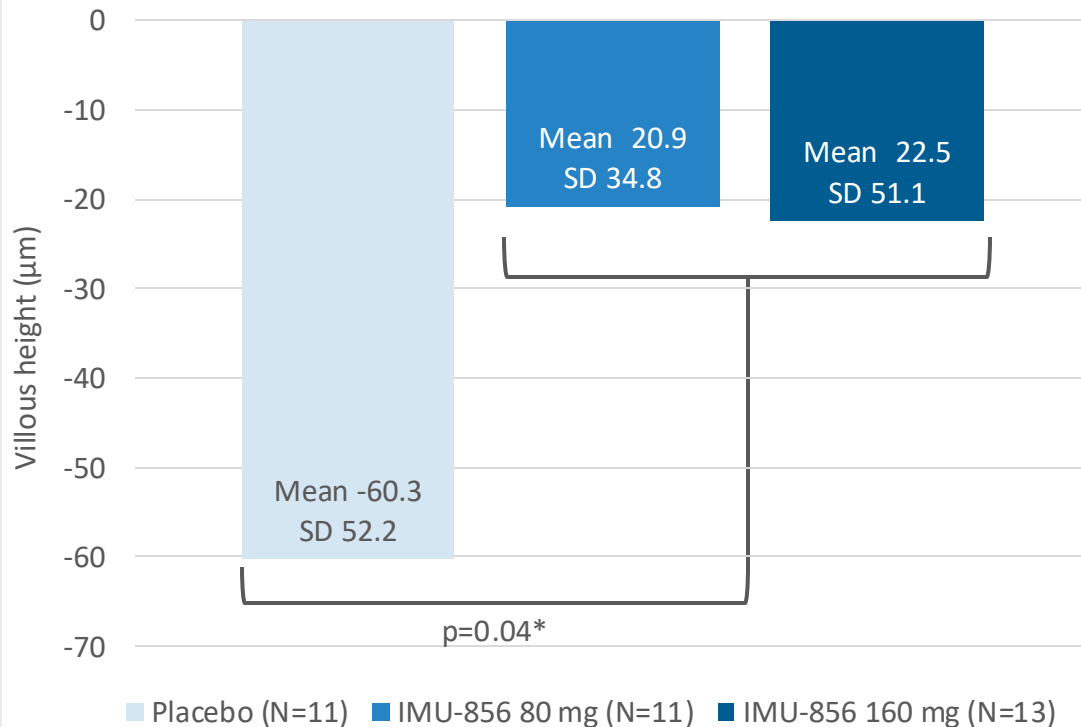
Flow Chart of Phase 1b Clinical Trial in Celiac Disease



QD: quaque die = once-daily; EGD: esophagogastroduodenoscopy

IMU-856 Protected Against Gluten-Induced Decrease in Villous Height as Compared to Placebo

Absolute change in villous height (μm) between Baseline and Day 29



Day 1 28: Treatment IMU 856/placebo

Day 14 28:
Gluten challenge with 6 g/daily

Baseline:
EGD with biopsy

Visit 6 / Day 29:
EGD with biopsy

- Substantial protection for IMU-856 treatment groups as compared to placebo
- Reached statistical significance* for this objective readout which is known to be relevant to influence future medical complications of celiac disease
- Assessed by central pathology laboratory and blinded pathology reader

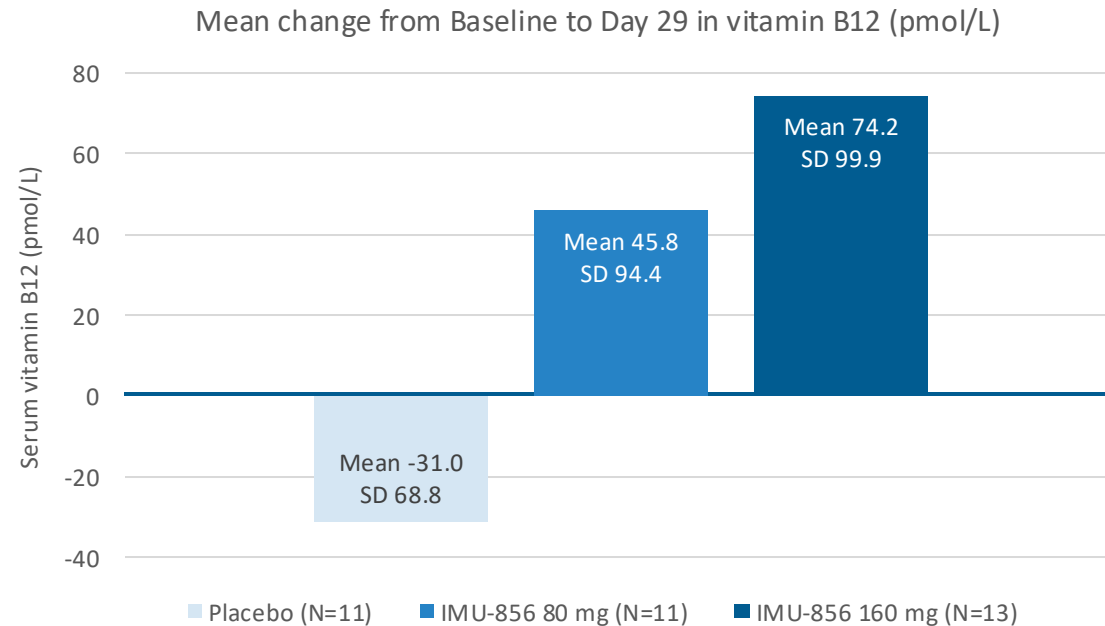
* Wilcoxon Two-Sample Test comparison between pooled IMU-856 groups and placebo, performed as post-hoc exploratory statistical analysis

Disease Analysis Set: N=35/43 included in histology analysis set. 8 patients not included in this analysis due to early termination. Gluten Challenge for 15 days with 6g daily. Central pathology laboratory: Jilab Inc. Tampere, Finland
EGD: esophagogastroduodenoscopy; SD: standard deviation

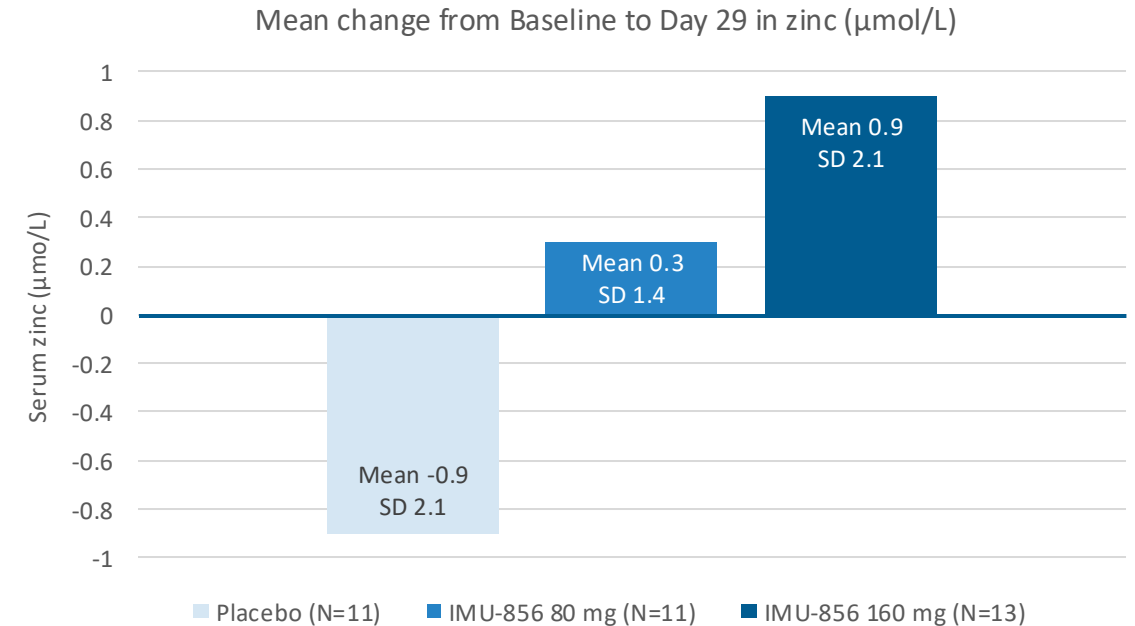
IMU-856 Improved Uptake of Actively Transported Essential Nutrients Vitamin B12 and Zinc



Vitamin B12



Zinc



SD: standard deviation



Immunic Therapeutics

Summary

Summary: Vidofludimus Calcium Is a Derisked Near-Term Opportunity



Innovative clinical pipeline: First in class oral drugs with unique modes of actions for multiple sclerosis and gastrointestinal diseases in various phases of clinical development



Relapsing MS opportunity is meaningful and de risked:

Oral category going to remain a large portion of overall MS market; peak sales potential for vidofludimus calcium of \$1-2 billion

Currently available oral therapies have limitations in benefit/risk profile; there is need for improvement

Vidofludimus calcium has the potential to address these shortcomings and transform the oral MS DMT market

ENSURE program: Two identical phase 3 clinical trials, designed to achieve potential regulatory approval of vidofludimus calcium in relapsing MS in a low-risk study design; top-line data for both ENSURE trials expected by end of 2026



Progressive MS provides tremendous upside opportunity:

High unmet medical need market: No approved therapies for non-active SPMS; one approved therapy for PPMS (infusion)

Peak sales potential for vidofludimus calcium of \$3-5 billion across respective indications

Phase 2 CALLIPER trial successfully demonstrated neuroprotective potential of vidofludimus calcium in progressive MS patients

Results to be discussed with healthcare authorities to determine appropriate next steps



Financials:

Cash position: \$55.3 million (as of June 30, 2025), shares outstanding: 98,650,590 (as of July 31, 2025)

Thank You!



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