Entera Bio Ltd.

Global Leader in Oral Peptide Therapeutics

Disclaimer

Various statements in this presentation are "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995. All statements (other than statements of historical facts) in this presentation regarding our prospects, plans, financial position, business strategy and expected financial and operational results may constitute forward-looking statements. Words such as, but not limited to, "anticipate," "believe," "can," "could," "expect," "estimate," "design," "goal," "intend," "may," "might," "objective," "plan," "predict," "project," "target," "likely," "should," "will," and "would," or the negative of these terms and similar expressions or words, identify forward-looking statements. Forward-looking statements are based upon current expectations that involve risks, changes in circumstances, assumptions and uncertainties. Forward-looking statements should not be read as a guarantee of future performance or results and may not be accurate indications of when such performance or results will be achieved.

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Entera Mission and Core Values:

- To develop transformative, first-in-class oral peptide treatments
- To invest in therapeutic spaces that have been ignored and require urgent attention

- To drive value for patients, clinicians and healthcare ecosystems smartly and expeditiously
- To think outside the box, innovate and cultivate our science, strategy and people everyday

Entera Company Background

First-in-Class Oral Peptides Designed to Empower Patients and Unlock Clinician Access

Proprietary N-Tab® Science and Platform

Innovative first-in-class oral peptides

Large (4kD+) linear, hydrophilic targets

Simple once a day tablet treatment

Robust protein design and modification capability via OPKO partnership

Corporate

Global interdisciplinary team

Nasdaq: ENTX, 49.2M shares outstanding

Lead Asset – EB613 (Osteoporosis)

Disruptive Therapeutic Asset

First oral tablet to democratize anabolic (bone building) treatment

Osteoporosis is a women centric condition with persistent treatment gap

Validated MOA, PTH(1-34), teriparatide

~$4–6B+ US revenue potential

Robust Phase 2 Data Package

Derisked 12M Phase 3 Regisrational Program Pursuant to June 2026 FDA Alignment

Diversified Pipeline

Rare Endocrine: Hypoparathyroidism

Metabolic /Obesity

GI Inflammation: Short Bowel Syndrome

Additional oral peptide development via partnership model where N-Tab® can be combined with unique targets

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Entera Oral Peptide Tablet Pipeline

Program	Indication	Target	Preclinical	Phase 1	Phase 2	Phase 3	Partner
EB613	Osteoporosis	PTH 1-34, teriparatide					
EB612	Hypoparathyroidism	LA-PTH 1-34					OPKO (Entera - 50% Ownership)
EB618	Obesity / Metabolic	GLP-1 & Glucagon Agonist					OPKO (Entera - 40% Ownership)
GLP-2	Short Bowel Syndrome	Long Acting GLP-2					OPKO
EB613	Stress Fractures	PTH 1-34					Investigator Sponsored Trial

Entera Core Team

Bringing Together Scientific Excellence to Deliver Better Treatment Outcomes Globally

Miranda Toledano
Chief Executive Officer and Board Member

Gregory Burshtein, PhD
Chief of Research and Development

Felicia Cosman, MD
Chair, Scientific Advisory Board and Chief Clinical Advisor (Endocrinology)

Riccardo P. Camisasca
Chief Medical Officer

Steven R. Goldstein, MD
Chief Clinical Advisor (Gynecology)

Michal Kushnir
Director of Program Management

Michele McDermott, MD
EVP Medical Affairs

Helen Pentikis, PhD
Clinical Pharmacology Specialist

Rachel B. Wagman, MD, FACE, FACP
Chief Clinical Advisor

Cherin Smith, PMP
EVP Clinical Operations

Osnat Bar-Peled, PhD
Executive Director of IP

Constantin Itin, RPh, PhD
VP of Preclinical Development

Dana Yaacov, CPA
Chief Financial Officer

Anke Hoppe
VP of Clinical Operations

Hillel Galitzer, PhD
Chief Operating Officer

N-Tab® Proprietary Platform

Barriers to Oral Peptide Delivery

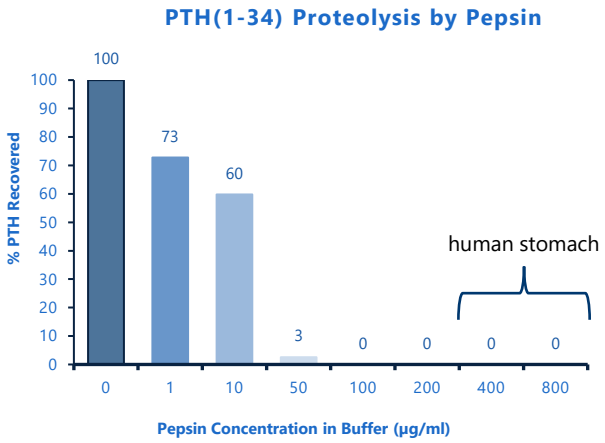
Proteolytic Degradation

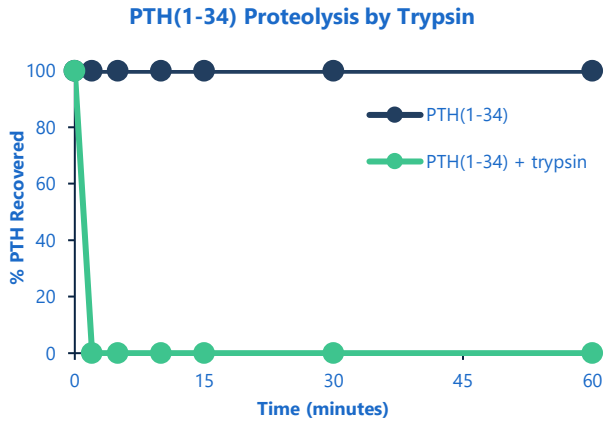
Stomach:

- Pepsin-mediated peptide cleavage
- Acidic pH → protein denaturation

Small Intestine:

- Trypsin & α-chymotrypsin degradation
- Brush-border metabolism





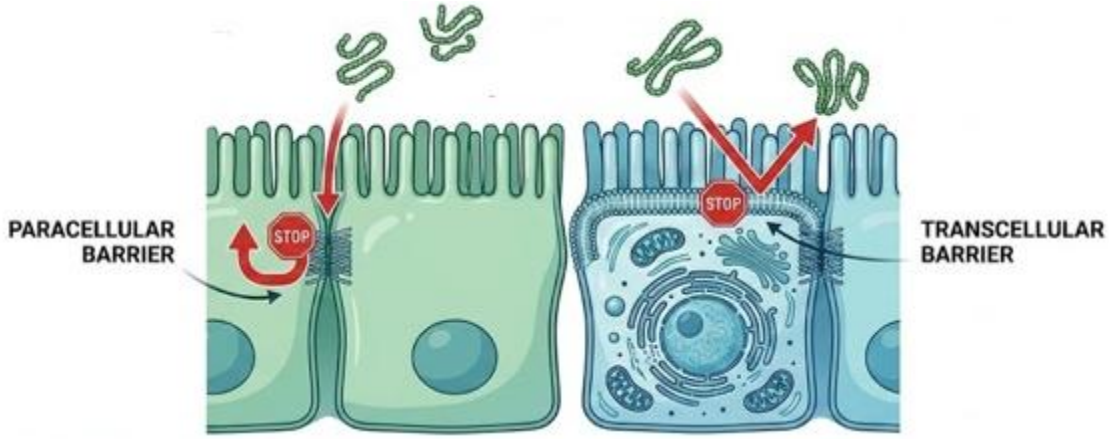
100%

peptide breakdown within few min of luminal exposure

Epithelial Permeability Barrier

Paracellular: Too large for tight junction transport

Transcellular: too hydrophilic to cross the hydrophobic core of enterocyte barrier



~0%

typical oral bioavailability for unformulated peptides

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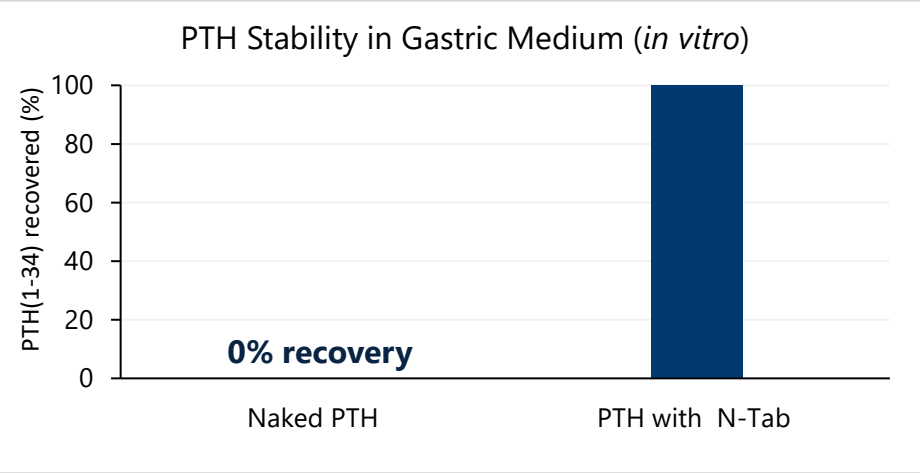
Unlocking Oral Peptide Absorption: Two Synergistic Mechanisms of the N-Tab® Platform

The N-Tab® platform overcomes enzymatic degradation and enhances permeability for improved bioavailability

1 Proteolysis Inhibition

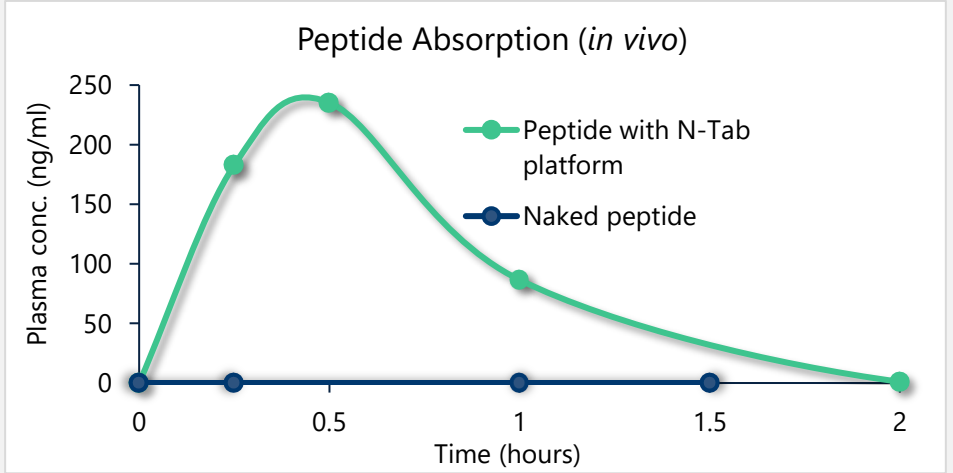
Protease Activity is Blocked.
Peptide Survives.



The naked peptide is completely degraded in the stomach within 5 minutes. N-Tab platform locally stabilizes the peptide

2 Permeability Enhancement

Membrane Fluidity Transiently Increased.
Drug Absorbed Systemically.



The N-Tab platform transiently increases enterocytes membrane fluidity, enabling transcellular permeation of the peptide through gastric wall

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EB613 Oral PTH (1-34), Teriparatide Tablet Treatment

First Once Daily Oral Osteoanabolic for Post-Menopausal Women with Osteoporosis

EB613: First Oral Anabolic (Bone Building) Tablet

- ✓ Oral tablet with identical API as blockbuster SC injection, Forteo® (Eli Lilly, approved since 2002, ~$1.7BN peak sales)
- ✓ Phase 2 showed comparable efficacy/safety as Forteo
- ✓ Unique dual MOA rapidly stimulates osteoblasts to increase new bone formation (PINP) and suppresses bone resorption (CTX)
- ✓ Oral supports earlier osteoanabolic intervention in line with guidelines
- ✓ Patients prefer oral tx in silent asymptomatic condition
- ✓ Potentially widens access from endo/rheum specialists (~15% Rx) to primary care/Gyn (~80% Rx)
- ✓ Potential for preferred tiering addresses huge healthcare burden
- ✓ Derisked asset, Phase 3 ready with efficient path to registration

Our Mission is to Democratize Anabolic Therapy so Every Woman can Protect her Bone Health and Deter the Risk of Fracture

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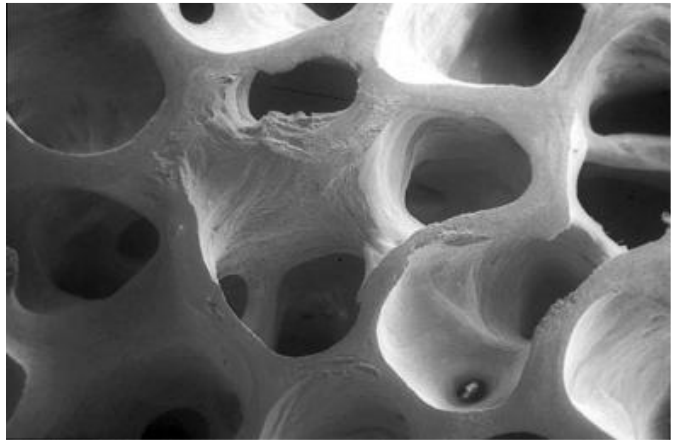
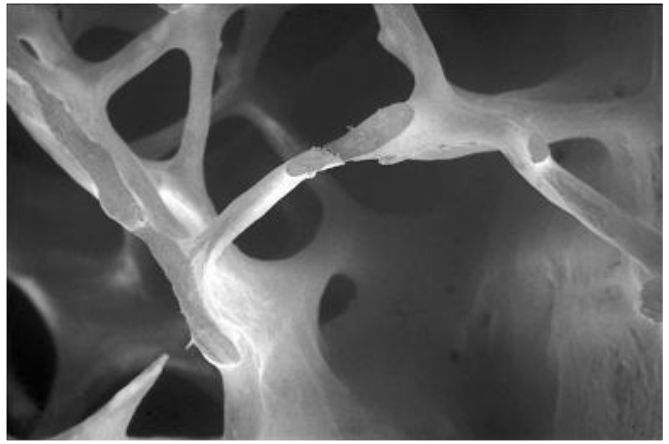
Osteoporosis is the Most Common Metabolic Bone Disease Worldwide





Dysregulated Bone Remodeling:

Increased Osteoclast Activity ↑
resorption (CTX biomarker)

Reduced Bone Formation ↓
formation (P1NP biomarker)

- Characterized by Low Bone Mass, Deteriorated Skeletal Microarchitecture, Compromised Bone Strength, and Increased Risk of Fracture
- Postmenopausal women are at highest risk of developing osteoporosis-related fractures, especially in the first 5-10 years after menopause
- Multiple guidelines recommend anabolic (bone forming) treatment as frontline treatment because of its superiority in rapidly increasing BMD and lowering fracture risk vs. anti-resorptive treatment
- The 3 approved anabolic drugs (Forteo®, Tymlos®, Evenity®) require daily/"in-office" monthly subcutaneous (SC) injections which limits both clinician and patient access

Osteoporosis Related Fractures Afflict More Women than Heart Attack, Breast Cancer and Stroke Combined Yet a Treatment Gap Persists Globally

200M

Women worldwide are affected by osteoporosis

1 in 3

Women and 1 in 5 men over the age of 50 will suffer an osteoporosis related fracture

54M

Americans at risk

~13M

Diagnosed

~60%

Do not accept or have access to currently available treatment options

>2M

Osteoporosis fractures (US) affect quality of life, disability, and premature death

>20%

of hip-fracture patients die within one year of the event

2019

Date of Last Approved Osteoporosis Drug

Phase 3 Registrational Program Included ~11,300 subjects

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EB613 Addresses the Treatment Gap in the Osteoporosis Patient Journey

~13M diagnosed with osteoporosis in the US, with estimated less than ~40% treated

There is need for novel products with enhanced efficacy, tolerability, and ease of use – FDA (2025)*

Anti-Resorptive

1st Line Bisphosphonates Fosamax®

~60%

Oral, Highly Accessible

~50% discontinue within 12 month due to AEs acid reflux, concerns over ONJ, AFF

RANK-Ligand Inhibitor (SC) Prolia®

~20%

SC injection every 6 months

Risk of Multiple Vertebral Fractures upon discontinuation, ONJ, AFF

EB613

- **First PTH Anabolic Tablet**
  Facilitates earlier bone building intervention to prevent fractures
- **High Accessability to Patients and Clinicians**
  Primary Care, GYN, Endo, Rheum
- **Ease of Administration Critical**
  In a silent progressive asymptomatic disease patients strongly prefer oral tablets
- **Mitigated Risk**
  Addresses patients' fear of injections and AEs such as Osteonecrosis of the Jaw (ONJ) and Atypical Femoral Fracture (AFF)

Injectable Anabolic

PTH Receptor Agonist Forteo®, Tymlos®

~9%

Daily SC injection

Many HCPs do not have in-office resources to bill for injectable drugs

Anti-Sclerostin Evenity®

~6%

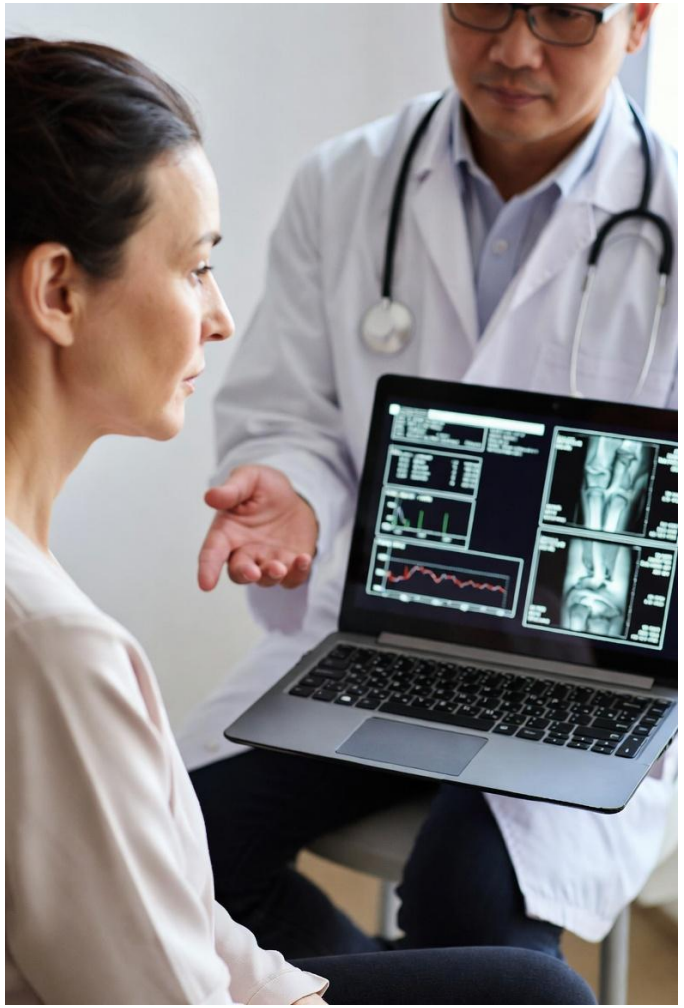
2 monthly "in-office" injections

CVD black box, Requires resources to buy and bill for and administer

*FDA Full Qualification Package Integrated Review, BMD as Validated Surrogate Endpoint for Post-Menopausal Women with Osteoporosis., December 19 2025
AFF=Atypical Femoral Fracture; ONJ=Osteonecrosis of the jaw

Why an Oral Osteoanabolic?

- 1 in 3 women over age 50 will experience a fracture in their lifetime
  Yet more than 80% will not receive treatment after a fracture
- Clinical guidelines recommend osteoanabolics as frontline treatment ("replace lost bone first")
  Yet injectable anabolics make up <~15% of estimated Rx
- Patients **less willing to take daily injections** in silent, asymptomatic disease
- There are major barriers to prescribing currently approved injectable anabolics only available as **"buy and bill"** and through **"specialty pharmacies"**
- Medical practices **cannot afford to purchase and manage reimbursement** for injectable osteoanabolics
- Prior authorizations and in office injections place an **unacceptable economic and staffing burden** on healthcare providers



EB613 Oral PTH (1-34) Tablet Treatment

Clinical Overview

EB613: First Once Daily Anabolic Tablet Treatment for Osteoporosis

Oral PTH(1-34) in a daily tablet (QD), LD Forteo®/Forsteo®

Thesis	EB613 is intended to provide an oral anabolic treatment earlier in an osteoporosis patient's journey Aligns with endocrinology/ women's health focus – bone health, menopause, longevity portfolios
Dual MOA	Rapid stimulation of osteoblast activity to increase new bone formation (PINP) with rapid suppression of bone resorption (CTX)
Clinical Efficacy	Placebo-controlled Phase 2 study in 161 post-menopausal women produced a rapid onset of action with significant increases in BMD at both cortical bone (femoral neck and hip region) and cancellous bone (spine)[1]
Safety Profile	AES in line with PTH agonists: headache, nausea, and dizziness
Opportunity	Three approved anabolic drugs require daily/monthly subcutaneous (SC) injections which limits patient access. Oral administration also expands access for HCPs who manage osteoporosis patients including gynecologists, women's health specialists and primary care providers as well as endocrinologists and rheumatologists
Target Patients	Sample target patients include postmenopausal women with: - T-scores $\leq$ -2.5 at total hip, femoral neck or lumbar spine with no fracture history - low bone mass or osteoporosis and history of nonvertebral fracture more than 2 years ago - low bone mass or osteoporosis and radiographic evidence of vertebral fracture
FDA Plan	505(b)(2) Pathway: A single 12-month placebo-controlled Phase 3 registrational study with BMD as primary endpoint and scientific bridge to LD Forteo, to be supported by a 12-month extension study

1. Tripto-Shkolnik et al., J Bone Miner Res 2024

EB613 Phase 2 Clinical Study in Postmenopausal Women with Osteoporosis

Key Inclusion Criteria

50+ years old

3+ years postmenopause

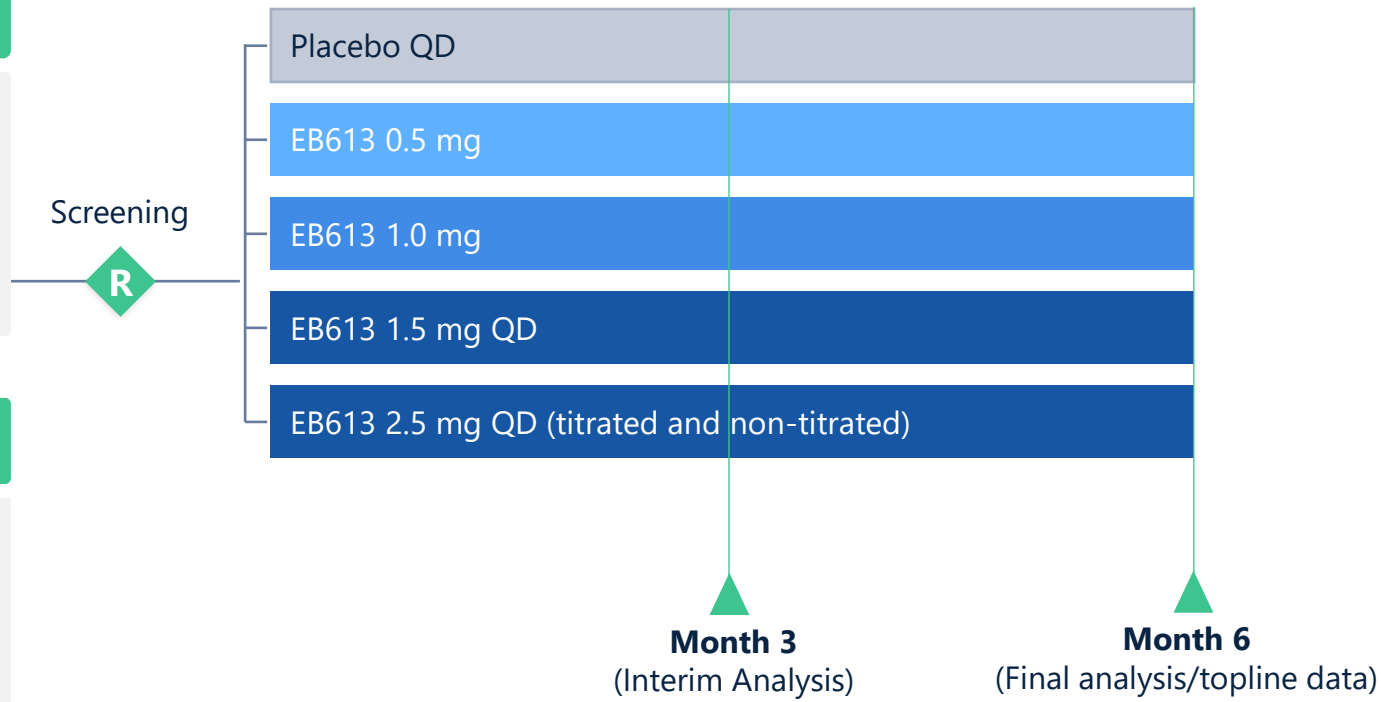
Low bone mass (≤-2.0 in at least one site)

High risk; no prior fracture

Key Exclusion Criteria

Osteoporosis treatment within last 2 years

Severe osteoporosis that precludes placebo



Primary Endpoint

Serum PINP % change from baseline (placebo-adjusted) at Month 3

Secondary Endpoints

BMD % change from baseline

Serum OC and CTX % change from baseline at Months 1, 2, 3, and 6

Serum PINP % change at Months 1, 2, and 6

Plasma PTH(1-34) at T15 min

6-month, randomized, dose-ranging, placebo-controlled study in postmenopausal women with osteoporosis met primary and secondary endpoints

Conducted at 4 sites; Enrollment: 161 patients (118 active, 43 placebo)

Tripto-Shkolnik et al., J Bone Miner Res 2024

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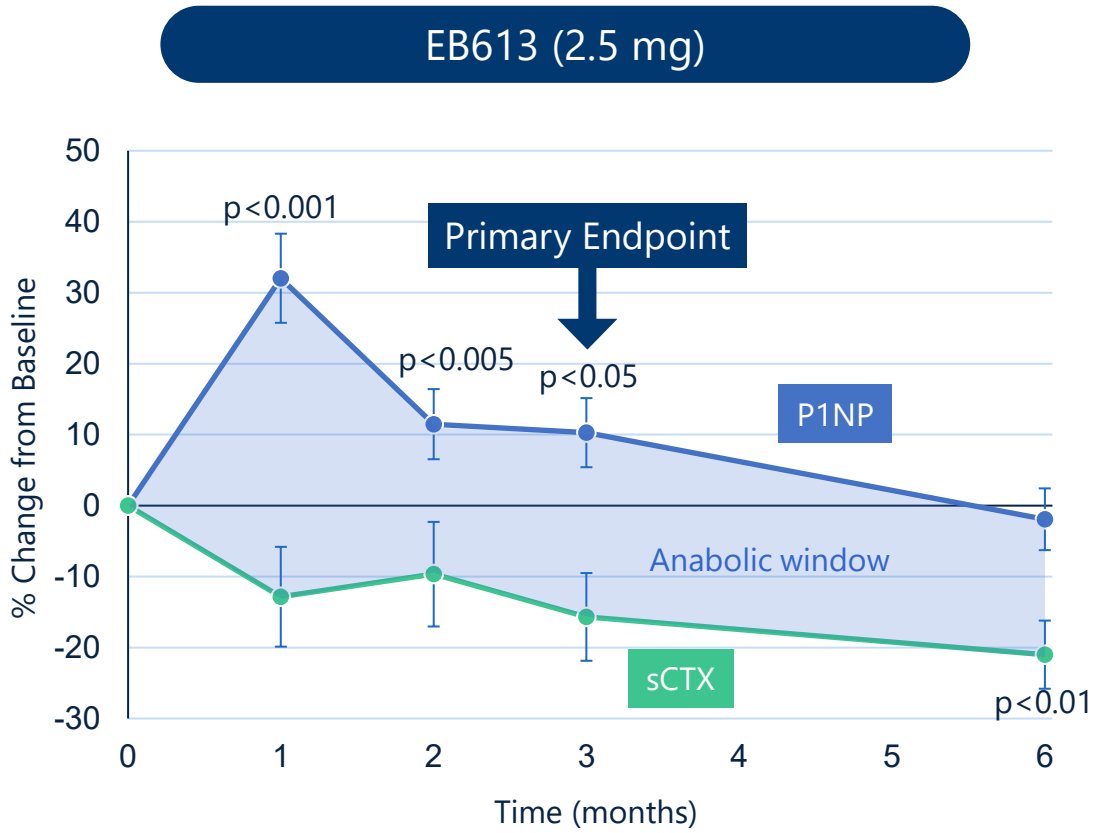
EB613 Induces Bone Formation (P1NP) while Reducing Resorption (CTX)

EB613's dual mechanism stimulates new bone formation on trabecular and cortical (periosteal and/or endosteal) bone surfaces by preferential stimulation of osteoblastic activity over osteoclastic activity



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Editorial

ASBMR The American Society for Bone and Mineral Research

OXFORD

A novel oral hPTH(1-34) unveils the promise of modeling-based anabolism with no increase in bone remodeling

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[2]Infectious and Immune Diseases Axis, CHU de Québec-Université Laval Research Centre, Quebec, Quebec G1V 4G2, Canada

An ideal osteoanabolic treatment would be one that could optimize the impact on modeling (formation) while producing lesser changes in resorption (remodeling) – JBMR, June 2024

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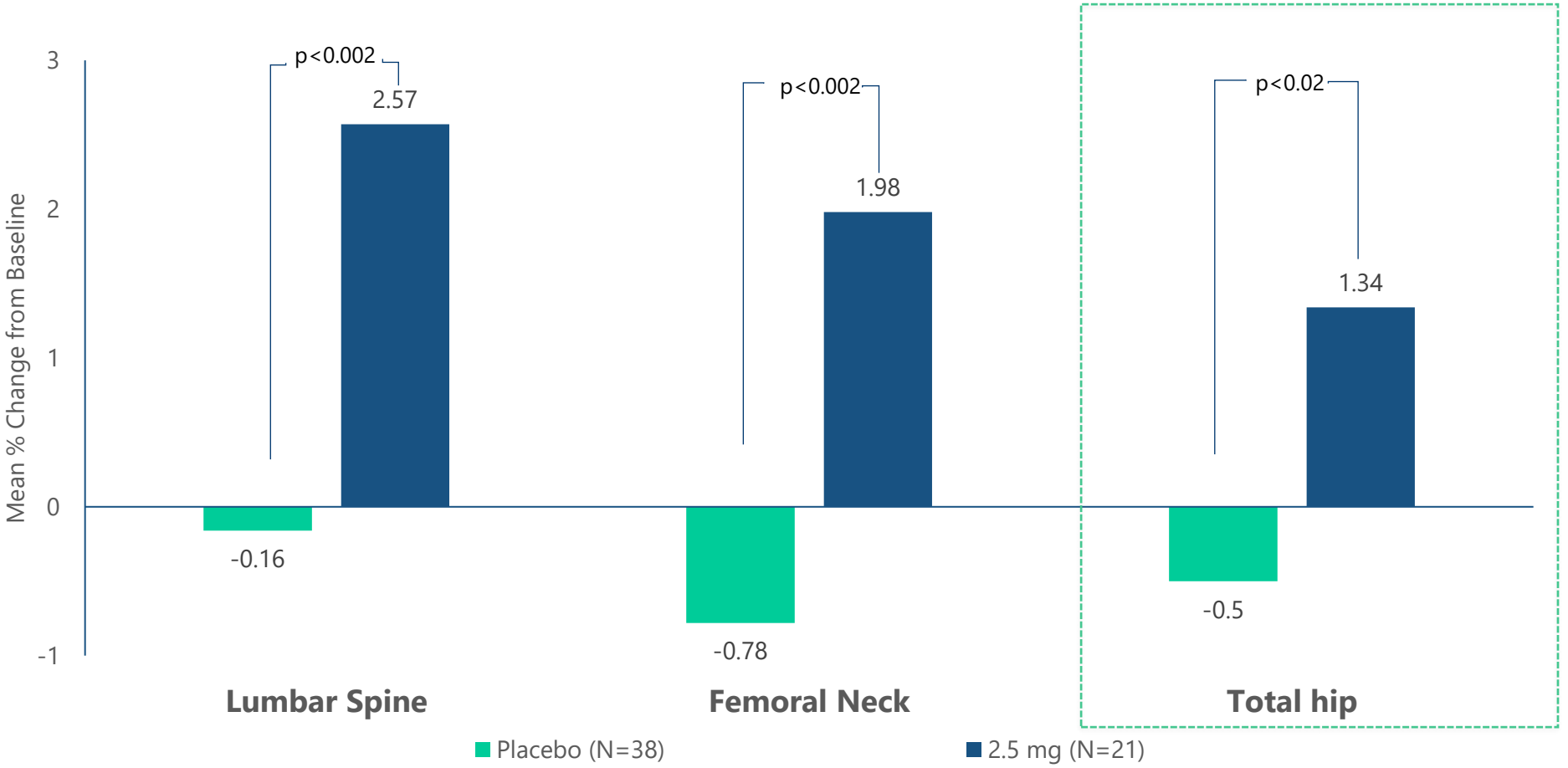
6 Months Treatment with EB613 2.5 mg Produced Significant Increases in BMD at All Skeletal Sites



	Lumbar Spine	Femoral Neck	Total Hip
Placebo Adjusted	2.73	2.76	1.84

Tripto-Shkolnik et al., J Bone Miner Res 2024

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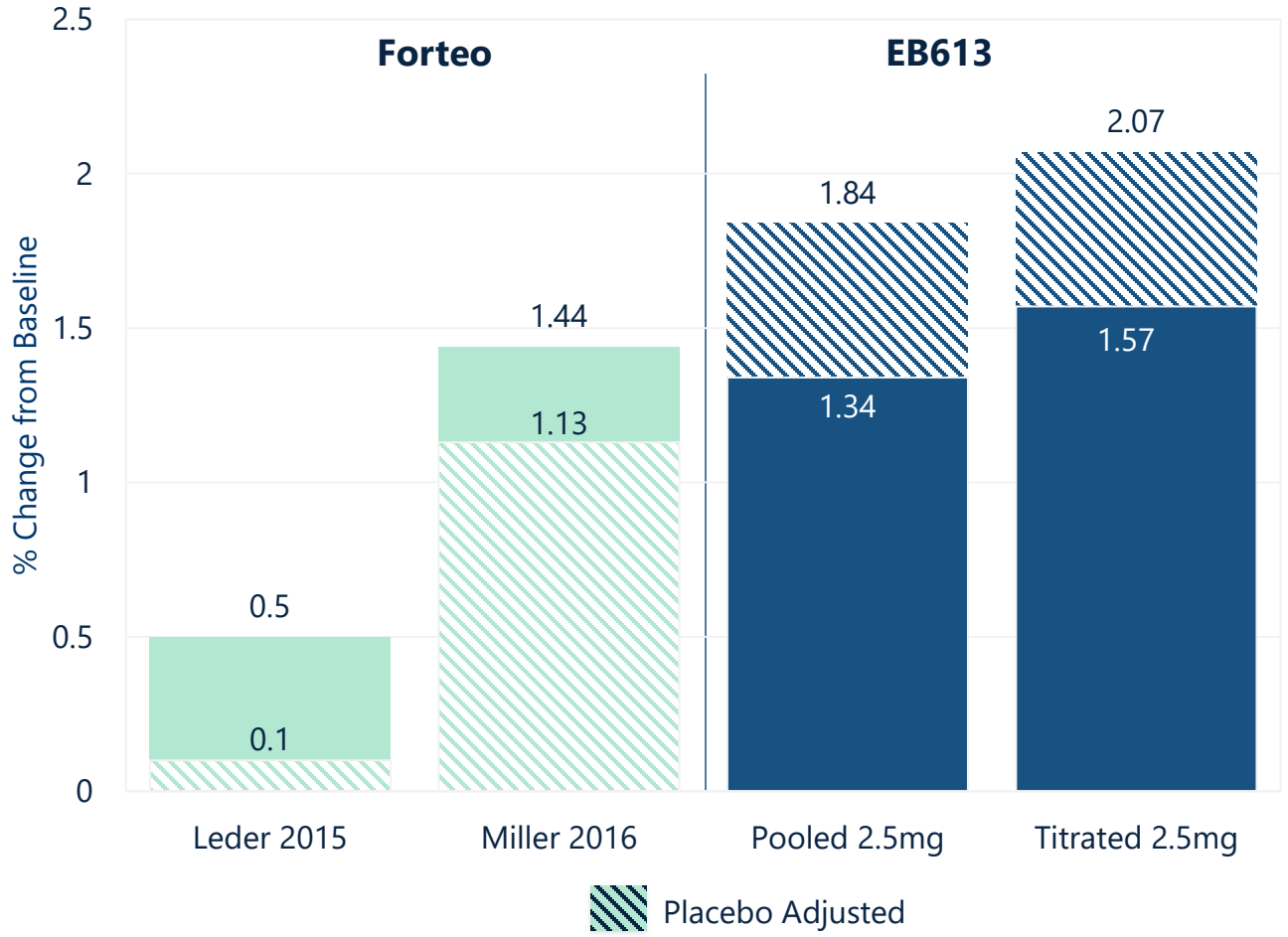
EB613 Produced Comparable Total Hip BMD Increases as Forteo® Published 6 Month Data



- Total Hip BMD is an FDA Efficacy Endpoint as of 12/19/2025
- 6 months of daily Forteo injections resulted in a 0.1 – 1.13% increase in Total Hip BMD[1]
- 6 months of daily Oral EB613 2.5mg resulted in a 1.84 – 2.07% increase in Total Hip BMD[2]

1. Forteo® data based on Leder BZ et.al. JCEM (2015) N=39, and Miller et.al. JAMA (2016) N= 754
2. EB613 data based on Tripto-Shkolnik L et.al. JBMR (2024) Pooled 2.5mg N=21, Titrated 2.5mg N=14

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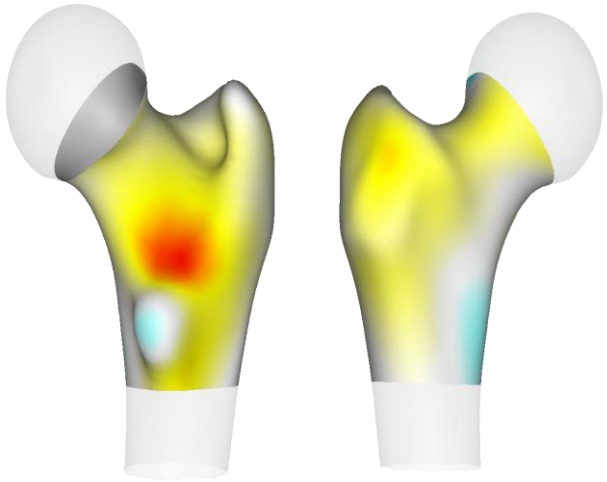
Effects of EB613 on Trabecular and Cortical Bone Using 3D-DXA

Average Cortical Surface BMD % Change from Baseline to Month 6 for EB613 and Placebo

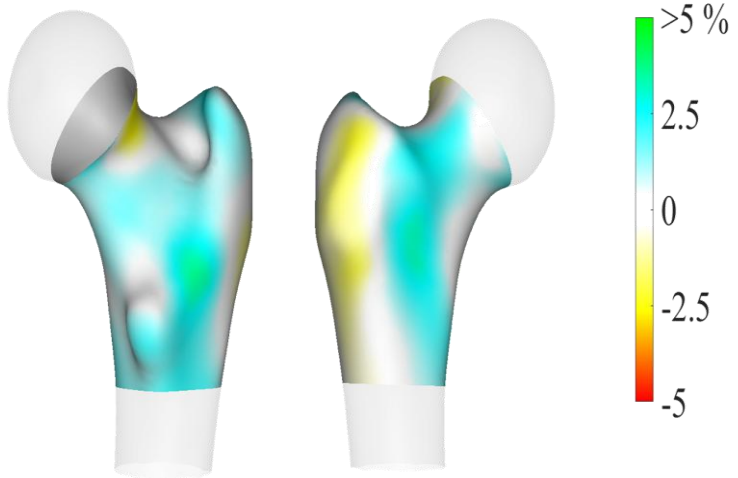
- EB613 showed an early effect on both trabecular and cortical bone of the proximal femur consistent with the dual mechanism of increased formation and decreased resorption
- Comparable assessment with Forteo (teriparatide) and Tymlos (abaloparatide) on cortical bone at 6 months[1]

Placebo (N = 38)



Posterior View Anterior View

EB613 (N = 21)



Posterior View Anterior View

Cortical sBMD (mg/cm^2) is calculated as the multiplication of the cortical thickness and the cortical volumetric BMD

1. Winzenrieth R et al., Osteoporos Int 2021; 32: 75-583.

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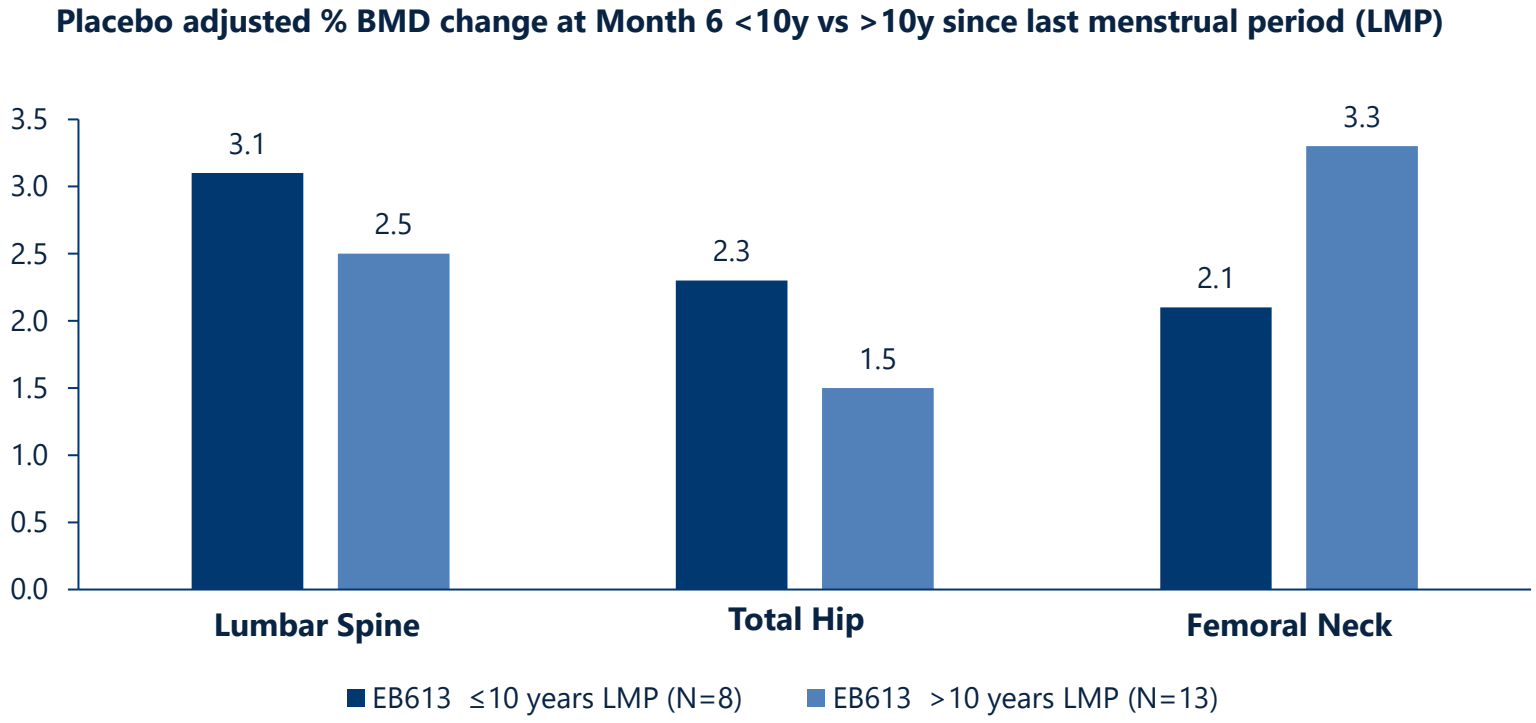
Significant BMD Improvements in Early Postmenopausal Women at 6 Months

Post-hoc EB613 Phase 2 analysis: BMD outcomes by time since last menstrual period (LMP)



For younger high-risk women without a prior fracture, BMD is the single most important predictor of osteoporotic fractures

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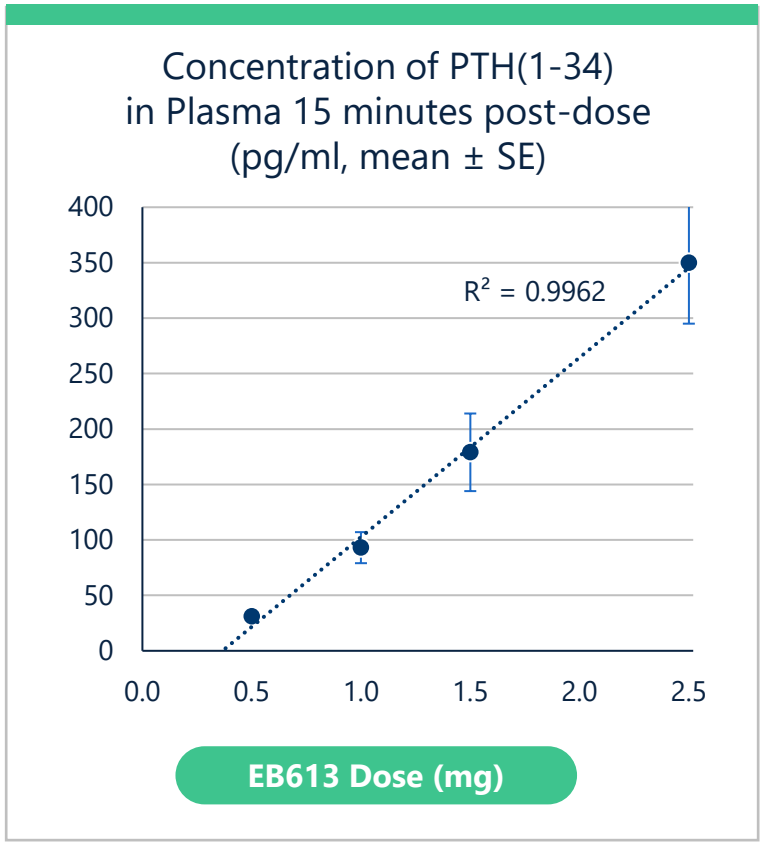
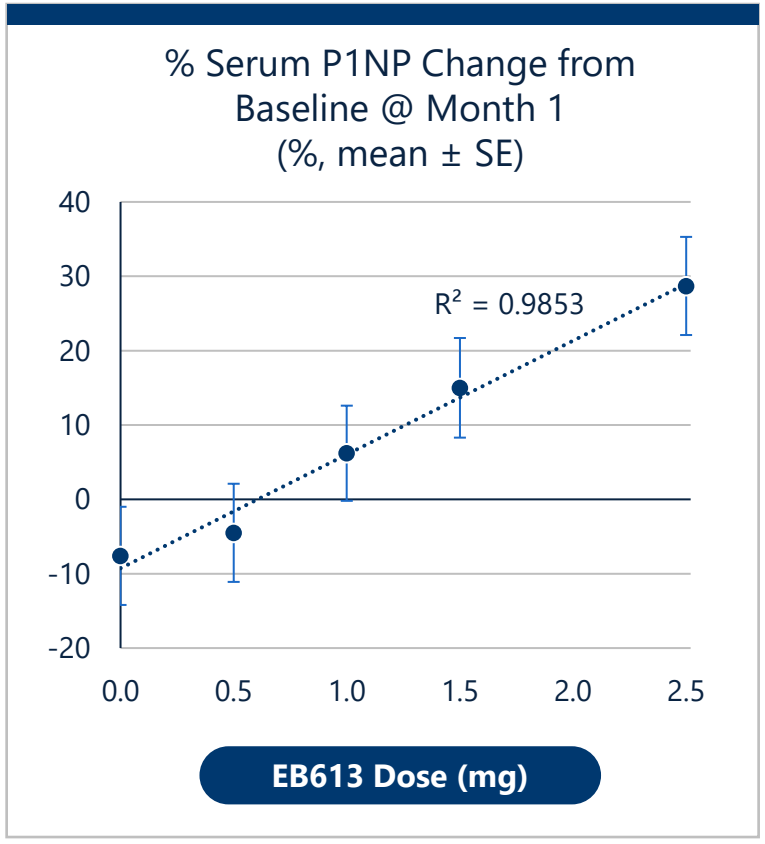
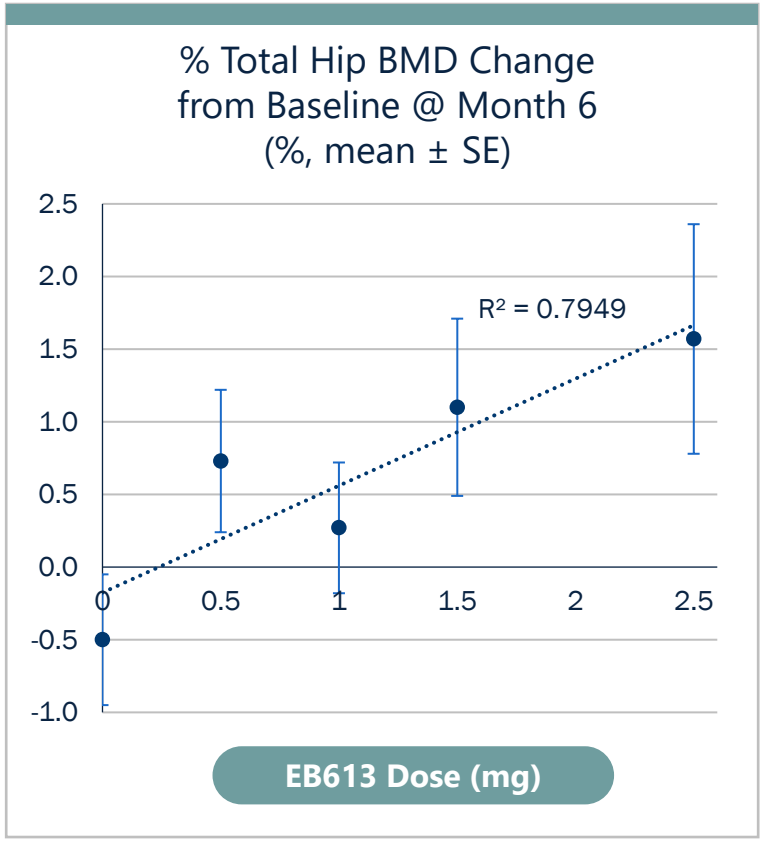
EB613 Showed Linear Dose Response Across PTH Exposure, P1NP Biomarker, and BMD







EB613 produced a statistically significant BMD dose response in lumbar spine BMD ($p<0.0001$), femoral neck BMD ($p<0.002$), and total hip BMD ($p<0.008$)

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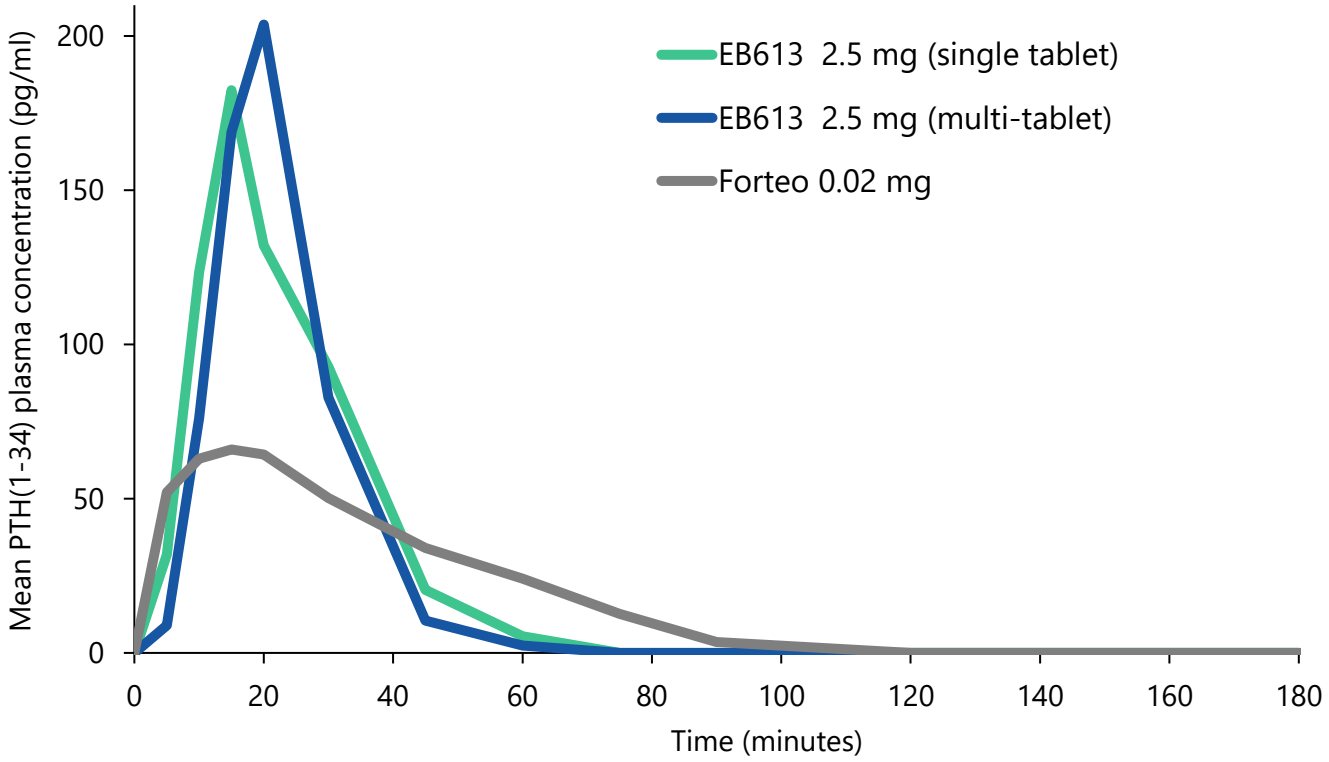
EB613 Phase 2 Safety Profile Consistent with PTH Agonists

Most Common Treatment Emergent AE (≥5% of participants)	EB613 Treated (N=118) n (%)
Headache	21 (17.8)
Nausea	18 (15.3)
Dizziness	13 (11.0)
Nasopharyngitis	7 (5.9)
Back pain	7 (5.9)
Palpitation	6 (5.1)
Dyspepsia	6 (5.1)
Presyncope	6 (5.1)

- Similar AE profile to that reported for Forteo® and other PTH agonists
- Mechanistic symptoms of orthostatic hypotension
  - headache, nausea, and dizziness
- EB613 was not associated with serum calcium increases or hypercalcemia adverse events
- 2.5 mg dose with titration (1.5 mg for 1 month, 2.0 mg for the next month and 2.5 mg during months 3 to 6) well tolerated
- No serious AEs related to EB613

Tripto-Shkolnik et al., J Bone Miner Res 2024

Phase 1 Bridging Study Supports Advancement of EB613 Single Tablet into Phase 3



Similar T_{max}, C_{max}, and total systemic exposure (AUC) were observed for EB613 formulations

AUC of EB613 2.5 mg is comparable to Forteo SC, but duration of exposure is slightly shorter

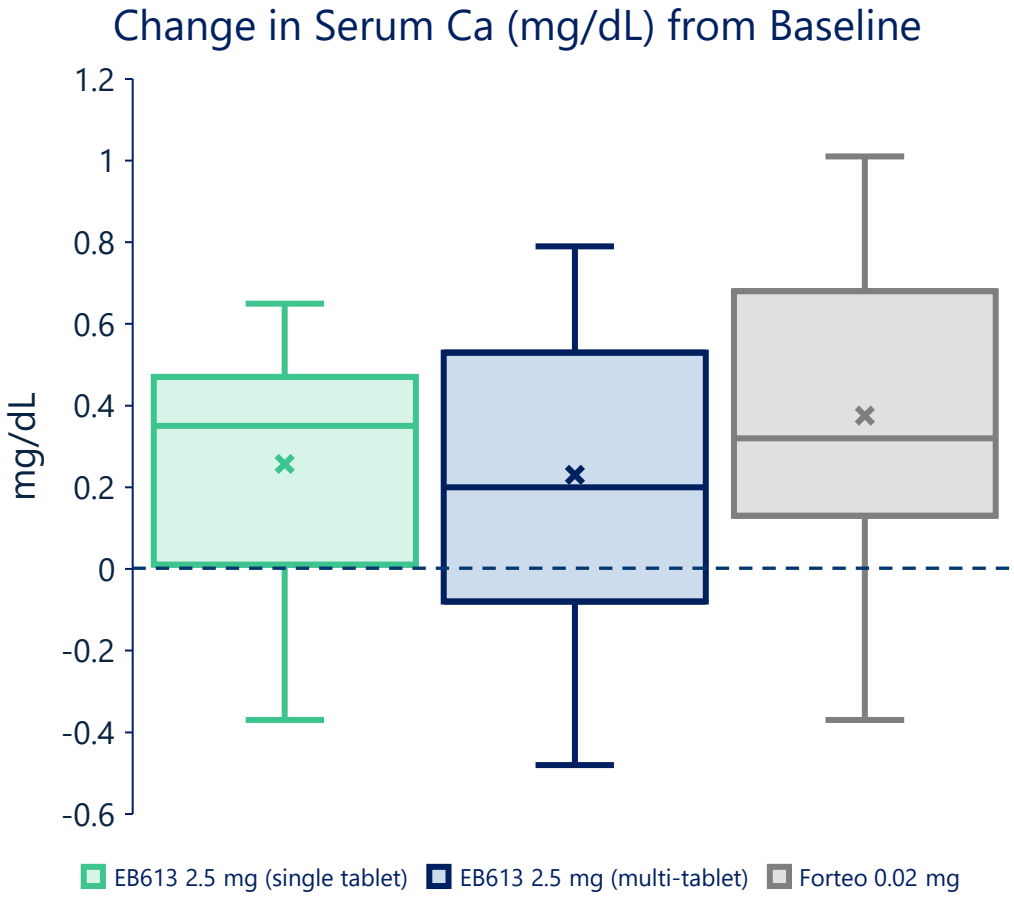
Multi-tablet 2.5 mg EB613

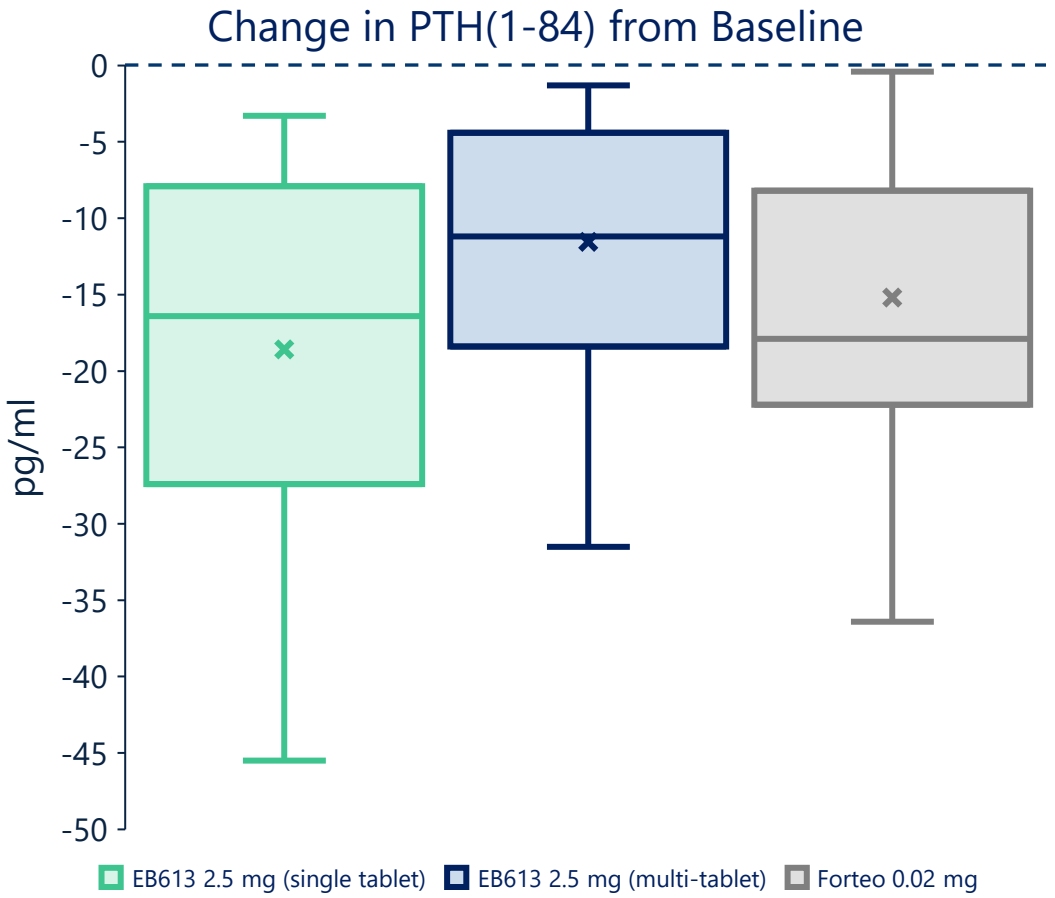
Single tablet 2.5 mg EB613

EB613 single tablet showed similar PK profile to multi-tablet EB613 treatment tested in phase 2 study

Burshtein et. al., Transforming Anabolic Treatments for Osteoporosis: New Clinical Data Supports a Single EB613 Tablet [Oral PTH(1-34)] as the Final Candidate for a Phase 3 Study, Endo 2026 Oral Presentation

Phase 1 Bridging Study - Comparable Calcemic Effects and Suppression of Endogenous PTH





Serum Ca and PTH(1-84) were measured 3 hours following EB613 dose and 4 hours following Forteo. The box represents the interquartile range (25th–75th percentile), with the line inside the box indicating the median. The mean is shown as an **x**. The whiskers extend to the minimum and maximum values within the typical range.

Burshtein et. al., Transforming Anabolic Treatments for Osteoporosis: New Clinical Data Supports a Single EB613 Tablet [Oral PTH(1-34)] as the Final Candidate for a Phase 3 Study, Endo 2026 Oral Presentation

Entera

Phase 1 Bridging Study Supports Advancement of EB613 Single Tablet into Phase 3

- Single EB613 tablet showed similar PK profile to multi-tablet EB613 treatment tested in phase 2 study
- Comparable calcemic effects and consistent suppression of endogenous PTH were shown for both oral EB613 treatments and for Forteo
- Safety profile of EB613 is consistent with Forteo

- A single daily oral PTH tablet (EB613) could make anabolic treatment far more acceptable to many patients and health care providers and make a substantial impact toward reducing the treatment gap in patients with osteoporosis

Overview of Safety for EB613

Phase 1 and phase 2

- In the clinical development program, the final single-tablet formulation and the multi-unit formulations have been administered collectively to a total of 270 subjects in phase 1 (n=117) and phase 2 (n=153) studies in:
  - healthy volunteers
  - postmenopausal women with low BMD or osteoporosis
  - male and female patients with hypoparathyroidism in doses up to 9mg daily
- EB613 has been well-tolerated, and no new drug-related AEs were identified in the Phase 1 and Phase 2 studies
- The incidence of mechanistic AEs, which are common to PTH agonists including both teriparatide and abaloparatide, including hypercalcemia, tachycardia, and orthostatic hypotension, were similar to or lower with EB613 as compared to Forteo in Phase 1 studies

Safety Profile of EB613 vs Forteo in Comparative Phase 1 studies

EB613 has been compared with Forteo in three phase 1 studies, N = 102 subjects received both drugs

	Forteo	EB613[1]
Number of study drug administrations	87	432
Treatment related AEs	11% (10)	5% (20)
Symptoms of orthostatic hypotension[2]	2.3% (2)	2.5% (11)
Nausea: the most frequent AE of EB613	2.3% (2)	1.4% (6)
Hypercalcemia: the most frequent AE of Forteo	3% (3)	<1% (2)

[1]All EB613 doses were included in the safety analysis

[2]Events of orthostatic hypotension symptoms including dizziness, headache, palpitations, tachycardia, nausea

ENT-01-2011 CSR, ENT-02-2013 CSR, ENT-11-2023 preliminary [1]safety data for the final 15-subject cohort of ENT-11-2023 study are still pending formal analysis

EB613
Phase 3 Design and Proposed Registrational Strategy

FDA Qualification of Total Hip BMD as Regulatory Endpoint, December 19 2025

U.S. FOOD & DRUG ADMINISTRATION

DRUG DEVELOPMENT TOOL
QUALIFICATION DETERMINATION
DDT-BMQ-000054

December 19, 2025

Currently available treatments for osteoporosis have limitations with respect to both efficacy and safety. There is need for novel products with enhanced efficacy, tolerability, and ease of use. The

This FQP for the surrogate efficacy endpoint proposes in the context of use (COU) that "the percentage change in total hip bone mineral density (BMD) assessed by dual-energy X-ray absorptiometry (DXA) can be used as a surrogate endpoint for the assessment of investigational anti-osteoporosis drug treatments for post-menopausal women at risk for osteoporotic fracture".

current approval paradigm using fracture endpoint faces multiple challenges: 1) it requires trials of long duration (2-5 years); 2) Placebo-controlled trials in high-risk patients are no longer ethical due to the availability of effective treatments, necessitating active controls or enrollment of lower-risk patients; 3) Large sample sizes are required to adequately power studies.

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EB613: June 2026 FDA Alignment

FDA Feedback Clears the Path for EB613's Phase 3 Registrational Study

> *"The FDA accepted Entera's plan to conduct a single, randomized, double-blind, placebo-controlled, Phase 3 trial in approximately 750 postmenopausal women with osteoporosis, with a primary endpoint of percent change from baseline in total hip BMD at Month 12 to support a potential New Drug Application (NDA) submission for EB613 for the treatment of women with post-menopausal osteoporosis. The proposed NDA package will also include Entera's scientific bridge analysis with Forteo® (teriparatide SC injection, Eli Lilly) under the 505(b)(2) pathway, and a transiliac crest bone biopsy sub-study in a subset of patients."*

Source: Entera's press release, June 22, 2026

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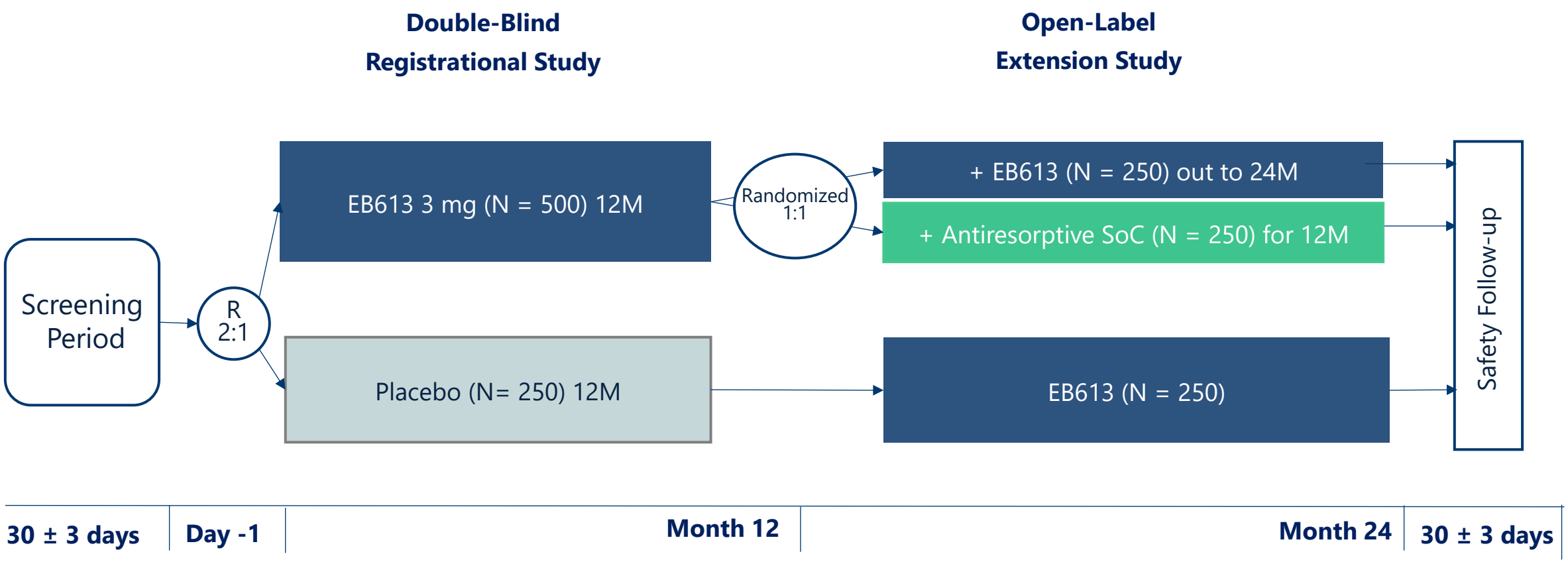
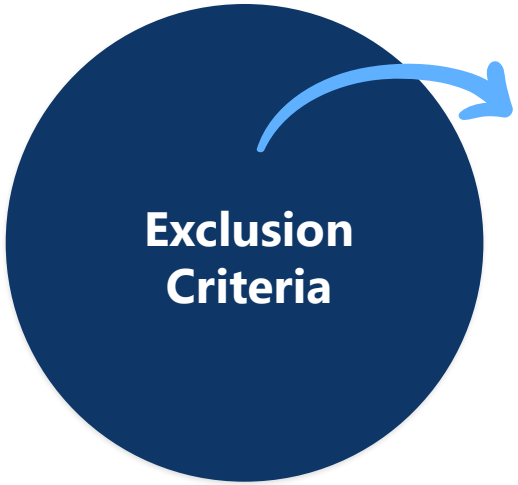
EB613 Phase 3 Development Plan



Key Eligibility Criteria

- **Ambulatory women ≥50 and ≤ 90 years of age**
- **Postmenopausal ≥5 years**
- **BMD T-score ≤ - 2.50 at lumbar spine, total hip, or femoral neck**
- **Eligible fracture types:**
  - One mild vertebral fracture
  - Non-vertebral fractures (other than hip or pelvis) >2 years



- **Very high risk for fracture, defined as one of the following**
  - Prior fracture of hip or pelvis, >1 mild vertebral fracture, or any moderate-severe vertebral fracture
  - Other non-vertebral fracture in the last two years, or
  - BMD T-score <-3.50 at lumbar spine, total hip, or femoral neck
- **Appropriate exclusions/washout periods for other osteoporosis medications**

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EB613 Patient and Clinical Surveys

EB613: Driving Value Across All Key Stakeholders

Healthcare Provider Surveys

GYNs, PCPs, Rheums and Endos likely to prescribe

Oral administration and known mechanism supports wide adoption

Patient Surveys

55% interested in oral bone-building therapy

70% Prioritize increasing bone density

47% Currently untreated despite high need

Payers & Market Access

Oral formulation expected to improve adherence, reduce costs

Differentiated profile supports favorable access

Robust Commercial Opportunity

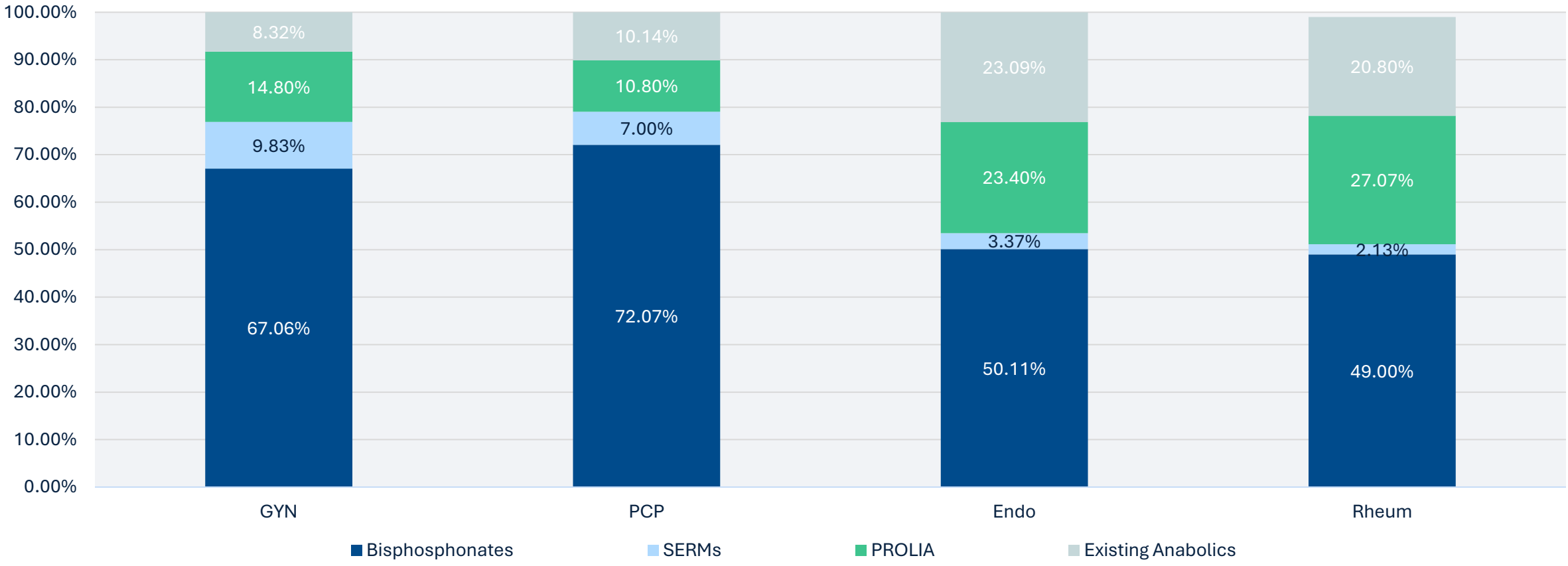
First and Only Once-Daily Oral Anabolic in Development

Estimated 38% patient share across specialties

Source: Entera market research

Oral Bisphosphonates and Prolia Dominate Across All Clinician Groups (76%), While Current Use Of Anabolic Therapies Remains Limited (~15%)

Consistent prescribing patterns across major treating clinicians



[Q7] Of your postmenopausal patients with osteoporosis who are currently using a prescription medication, please indicate the percent use of each of the following treatment options in the past 6 months:

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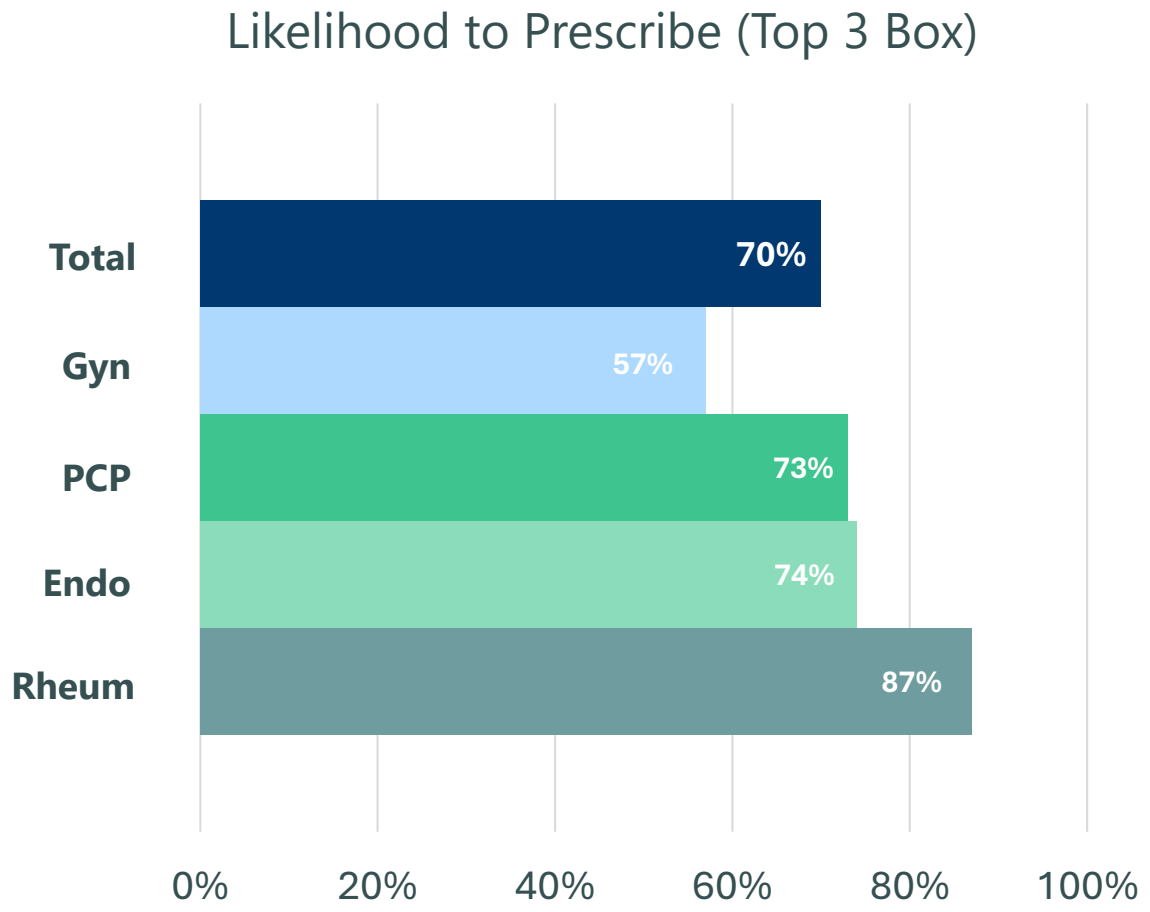
EB613 – A New Oral Anabolic Agent Poised To Transform Osteoporosis Treatment

- Physicians who rated their likelihood to prescribe as 8, 9, or 10 are represented visually
- ENDOs are extremely enthusiastic about willingness to prescribe EB613

Overwhelming prescriber confidence in EB613 across all physician segments



[Q25] How likely are you to prescribe it on a scale of 1 to 10 where 1 is “not at all likely to prescribe” and 10 is “extremely likely to prescribe”?

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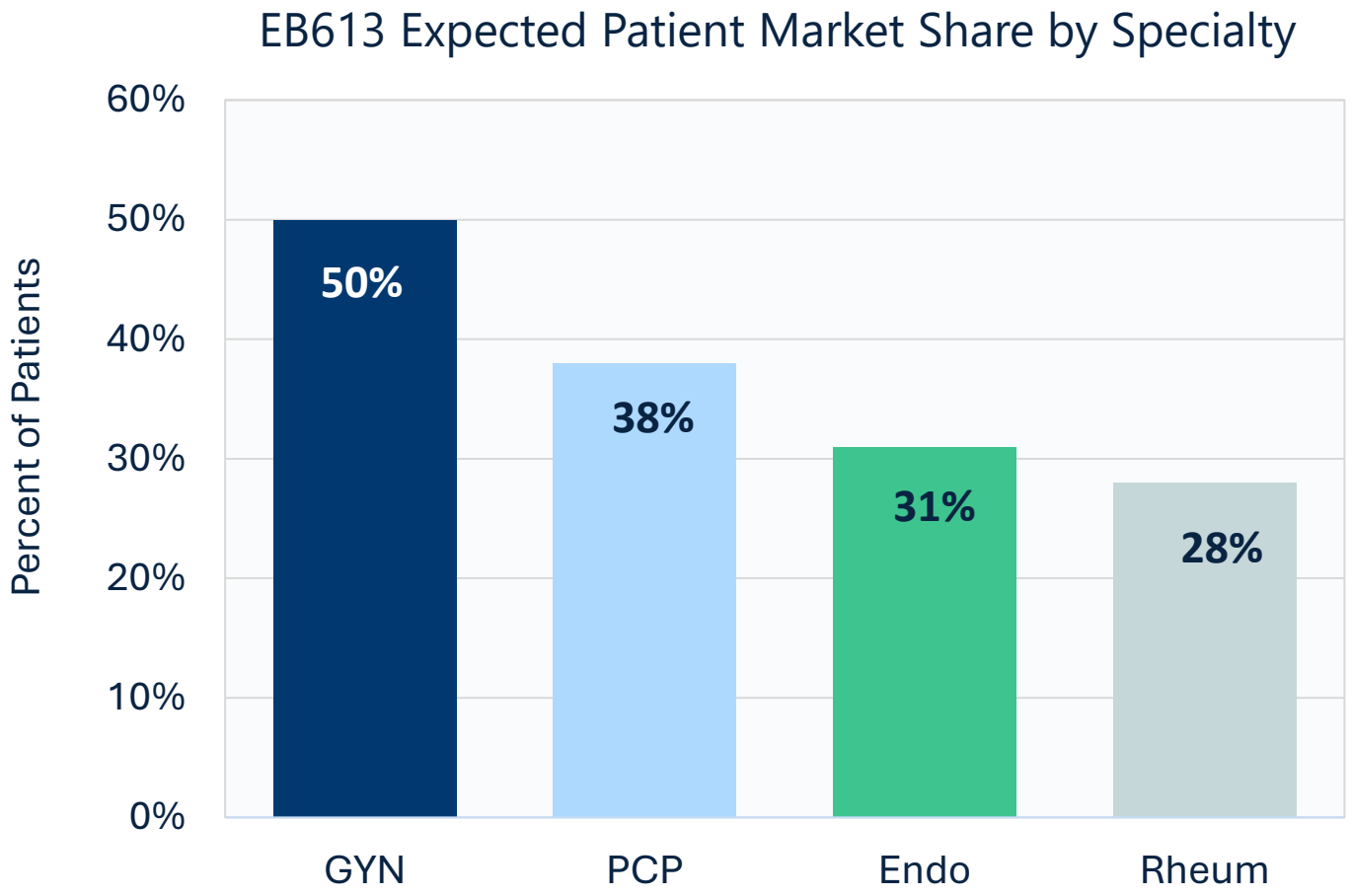
Across All Specialties EB613 Demonstrates Strong Adoption Potential



This broad acceptance across specialties, with a **38%** overall patient market share, underscores the potential for EB613 to become a game-changing treatment in osteoporosis.

These results signal a significant shift and readiness of HCPs to integrate EB613 into clinical practice.

[Q26] Considering your total postmenopausal osteoporosis population, what percentage of patients would you prescribe Product E for?

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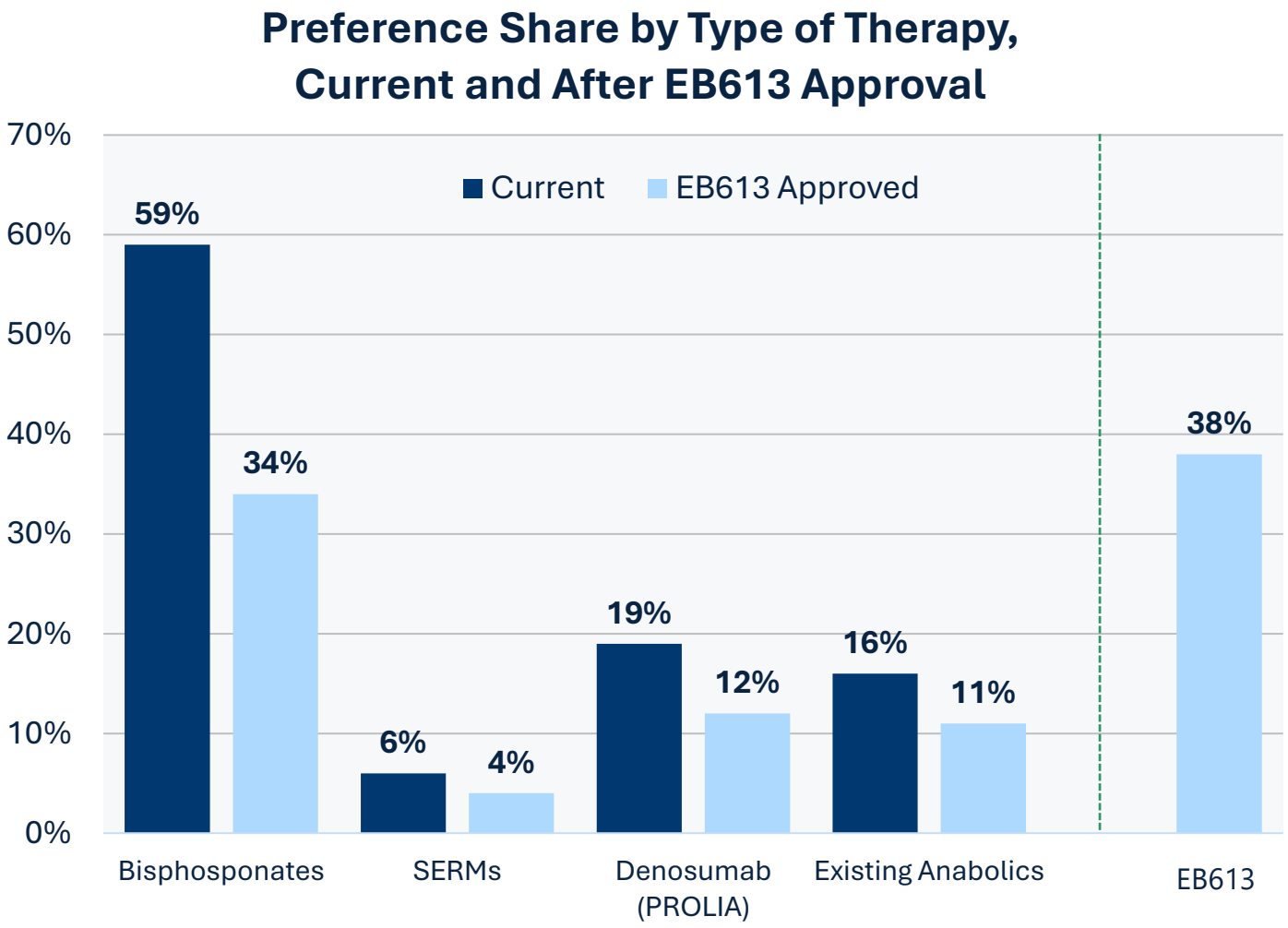
EB613 Is Poised To Take Share From All Established Therapies

- EB613 is likely to become a preferred oral anabolic option in osteoporosis treatment, displacing both oral bisphosphonates, denosumab, and existing anabolics
- ENDOs report that nearly half (47%) of patients rejecting anabolics would be open to taking EB613, showing strong potential for this oral alternative
- GYNs report that over half (56%) of patients who previously refused injectable anabolic treatments would choose EB613



Source: Market research survey, 2024 by Fit2Market

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2023 Survey: OB/GYN and GYN Practices Are the Natural Hubs for Advocacy of Women's Bone Health

At the 2023 global menopause society conference, a 19-question, self-administered survey assessed osteoporosis knowledge and treatment practices of 129 responding physicians from OB/GYN (63%) and GYN (37%) practice specialties.

89% of respondents

routinely include bone health issues in their standard GYN care in menopausal and perimenopausal patients.

- 92% order DXA scans
- 82% are familiar with FRAX

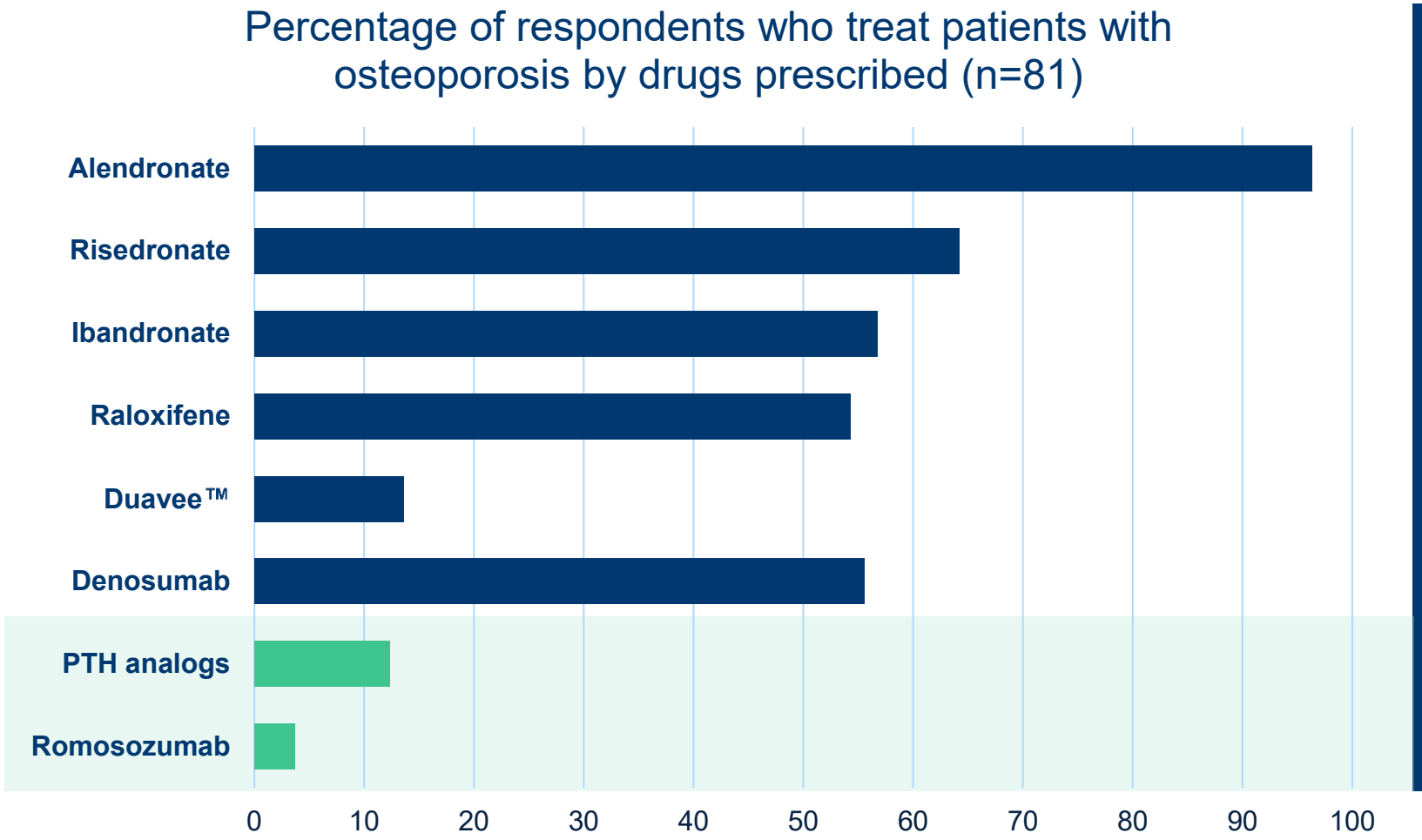
71% of respondents

treat patients to **prevent** osteoporosis.

64% of respondents

treat patients **with** osteoporosis.

GYN Osteoporosis Treatment Selection Reflects the Underutilization and Limited Market Penetration of Injectable Anabolics



Of the **64% of respondents** who treat patients with osteoporosis,

only **12%** prescribe injectable anabolics.

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Evolving the Route of Administration (Such as an Oral Anabolic Minitablet) Dramatically Shifts OB/GYN and GYN Willingness to Prescribe

Availability of an oral anabolic would shift the use of anabolics

In treaters of osteoporosis (n=75),

from **12%** to **92%.**

Of treaters who **do not use** injectable anabolics (n=63),

92% would prescribe an oral anabolic.

Of treaters **using** injectable anabolics (n=10),

100% would prescribe an oral anabolic.

Even in those who do not treat osteoporosis (n=28),

from **0%** to **50%.**

EB612 Program

First Daily Oral LA-PTH(1-34) Replacement Therapy for the Treatment of Hypoparathyroidism

In Partnership with OPKO

Hypoparathyroidism: Significant Unmet Need Remains

Current therapies leave the majority of patients inadequately controlled or burdened by daily injections

>200K

Patients (US + EU)

Predominantly women

~75%

Post-surgical cases

Thyroid /Parathyroid surgery

Daily

Injection burden

Palopegteriparatide (Yorvipath, 2024)

Conventional Therapy - Standard of Care

High-dose oral calcium (up to 3 g/day) + active vitamin D analogs

Goal: maintain serum Ca in low-normal range

But fails to address:

- ✗ Restore physiological Ca–phosphate homeostasis
- ✗ Prevent hypercalciuria and kidney disease
- ✗ Normalize patient quality of life - fatigue, cramps, cognitive symptoms persist

Current Approved / in Clinical Development

Natpara - rhPTH 1-84 (Takeda)	Discontinued	**Daily SC injection**; withdrawn due to manufacturing issues (rubber particulates)
Yorvipath - Palopegteriparatide (Ascendis)	Approved	**Once-daily SC injection**
Eneboparatide (AZN/Alexion)	Regulatory Filing Pending	**Once-daily SC injectable** PTH analog, 31.1% achieved composite primary endpoint at week 24; immunogenicity in majority of patients
Canvuparatide (MBX Biosciences)	Phase 3 (to be initiated)	**Once-weekly SC injection**
SEP-479 (Septerna)	Phase 1 (data expected in late 2026/early 2027)	Oral **small molecule** PTHR1 (Previous molecule SEP786 discontinued February 2025 due to safety)

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EB612 Profile

Oral Long-acting PTH(1-34) Tablet for Hypoparathyroidism

MOA: First-in-Class Oral, Long-Acting PTH(1-34) Protein Replacement Therapy Tablet

Dosage Form: Daily Single Tablet

Pharmacokinetics and Pharmacodynamics: >24 hours exposure following a single dose with a robust PD response

Production process: Scalable pharmaceutical production process

Shelf-life: Long shelf-life at convenient storage conditions

Oral Peptide

Tablet friendly for patient compliance
Differentiation vs small molecules (Safety, DDI)

Sustain PTH Exposure

Prolonged plasma half-life >72h
Potential low mg daily dose

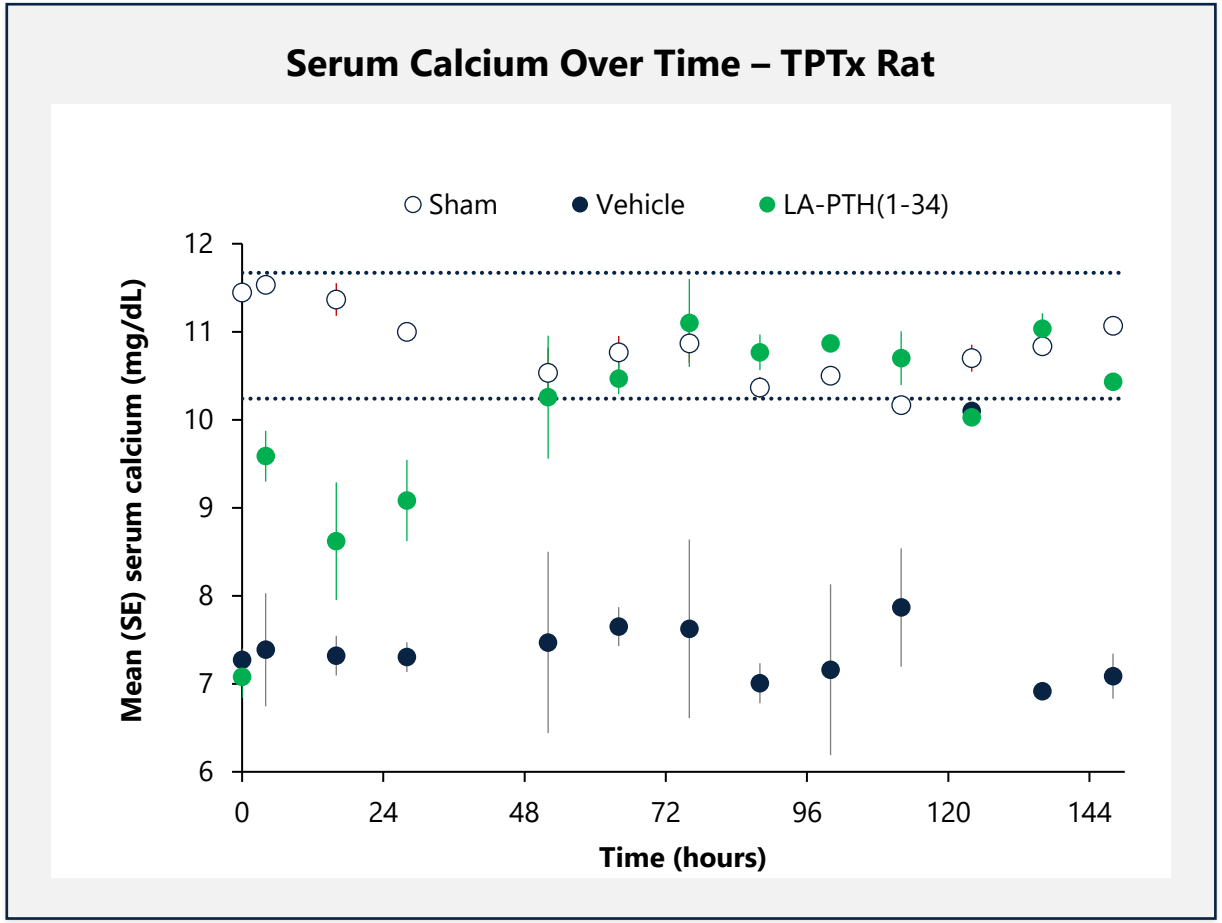
Cross-Species Preclinical Validation

Pharmacodynamic effect in TPTx rat and two large animal species - providing a robust translational package for IND-enabling studies

Well Tolerated

Favorable early tolerability signal

EB612: LA-PTH(1-34) Variant Normalizes Serum Calcium in TPTx Rat Model



- The thyroparathyroidectomized (TPTx) rat is the universally accepted gold-standard model of surgical hypoparathyroidism
- The drug was administered intravenously to TPTx male SD rats on a daily basis for one week (n=9 in each arm).
- **Full Calcium Normalisation**
  EB612 restored serum calcium and reduced phosphate to levels comparable to sham control animals
- Calcemic effects observed in the TPTx rat model are consistent with those reported in similar studies with other clinically validated PTH-replacement therapies for hypoparathyroidism

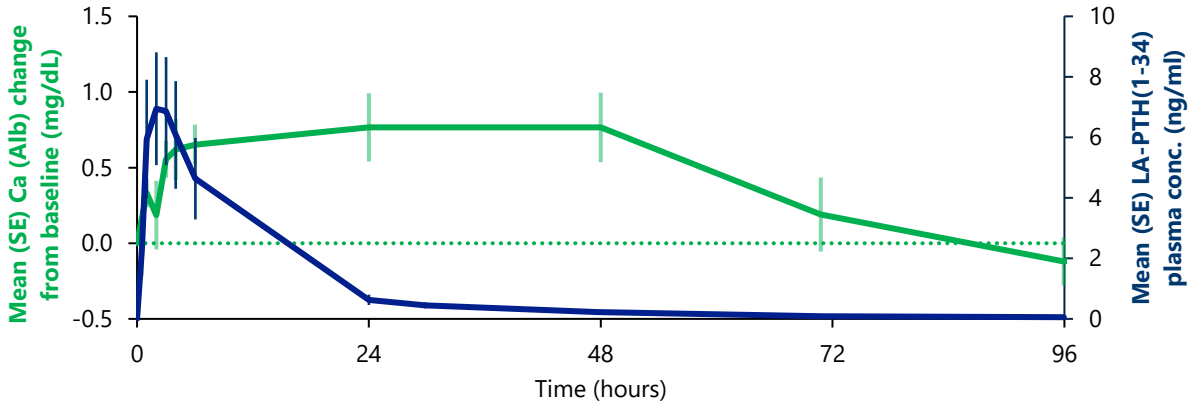
Burshtein et. al., Pre-Clinical Results for EB612: First-in-Class Oral Long-Acting PTH(1-34) Analog as Hormone Replacement Tablet for Patients with Hypoparathyroidism, [Poster presentation]. ENDO 2026: The Endocrine Society Annual Meeting, Chicago, IL, USA.

EB612: Long-Lasting Serum Calcium Elevation in Large Animal Models

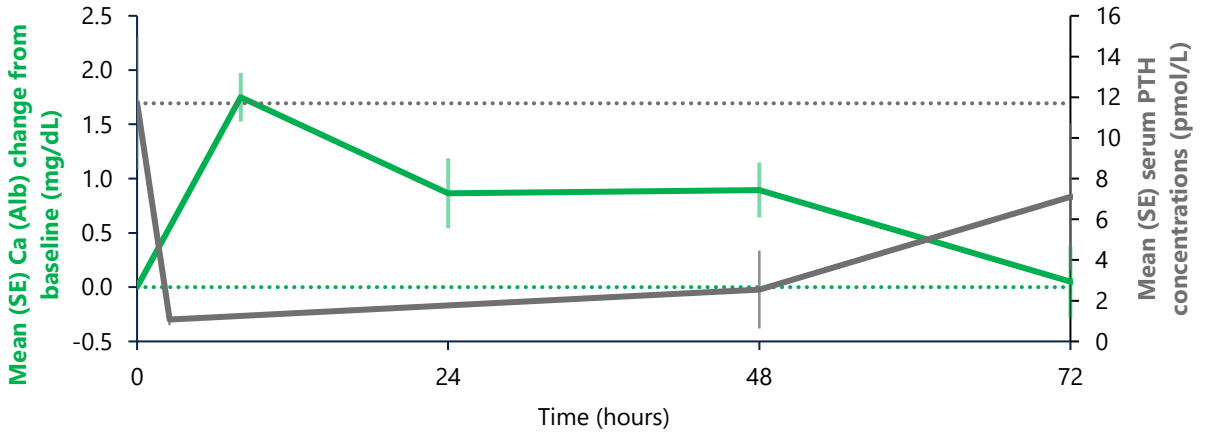
Cross-species validation

- **Rapid onset** - A single EB612 tablet achieved maximal plasma levels 2 to 3 hours post-dose
- **Prolonged PK profile** - The drug remained detectable in plasma for approx. three days
- **Sustained calcemic effect** ~48-72h supports QD dosing
- **Consistently lowering endogenous PTH(1-84)**
- **Well tolerated** in single dose, large animal studies

Minipig model



Non-human Primate model



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GLP1/Glucagon and GLP2 Programs

In Partnership with OPKO

First Oral Once Daily OXM (GLP-1/Glucagon Dual Agonist) Tablet for Obesity and Metabolic Disorders

>1 billion people suffer from obesity globally; market is estimated to grow to $100B by 2030

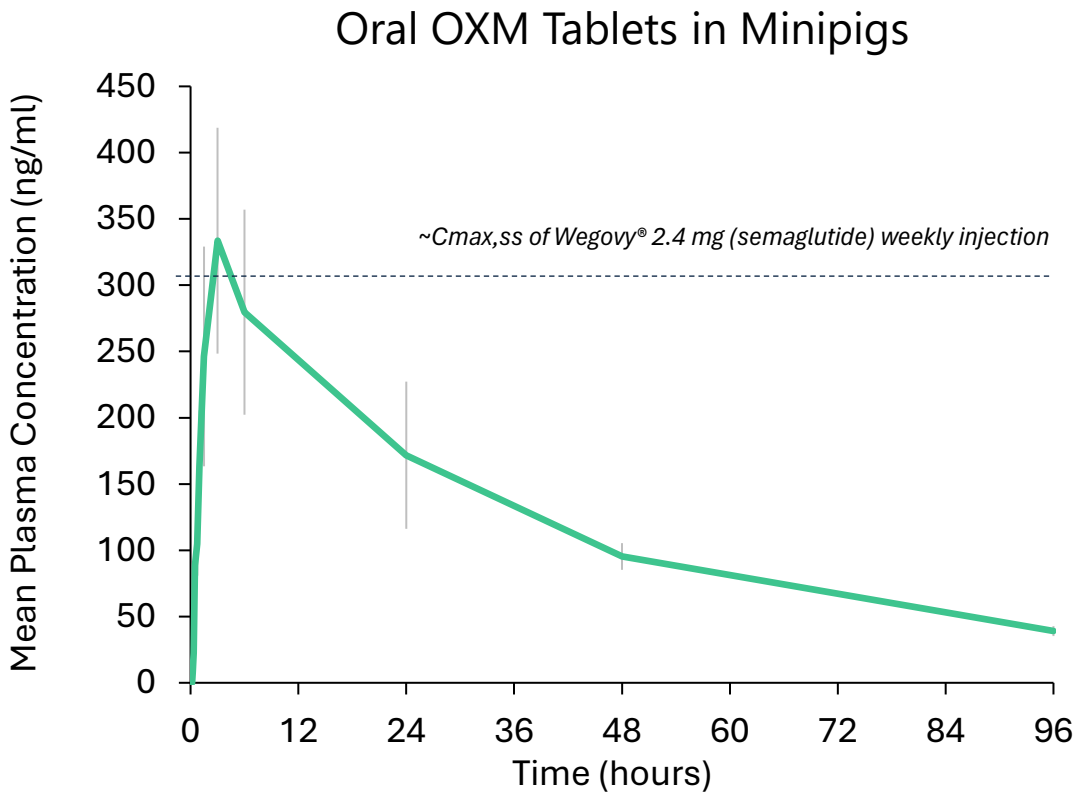
SUN-680. First-in-Class Oral Dual GLP-1/Glucagon Agonist for Patients with Obesity and Metabolic Disorders: In Vivo Pharmacokinetic and Pharmacodynamic Results

Gregory Burshtein[1], Constantin Itin[1], Daniel Pery[1], Eli Reichman[1], Hillel Galitzer[1], Michal Kushnir[1], Ahuva Bar-Ilan[2], Moran Golan[2], Laura Moschcovich[2], Miri Zakar[2], Amit Rivkin[2], Moran Levy[2], Miranda Toledano[1], Jane Hsiao[3]

[1] Entera Bio Ltd., Jerusalem, Israel, [2]OPKO Biologics Ltd. Kiryat Gat, Israel, [3]OPKO Health Inc. Miami, FL, USA

Entera Bio

OPKO



Extended Plasma Half-Life: Biologic half-life consistent with the half-life reported for semaglutide in a similar animal model

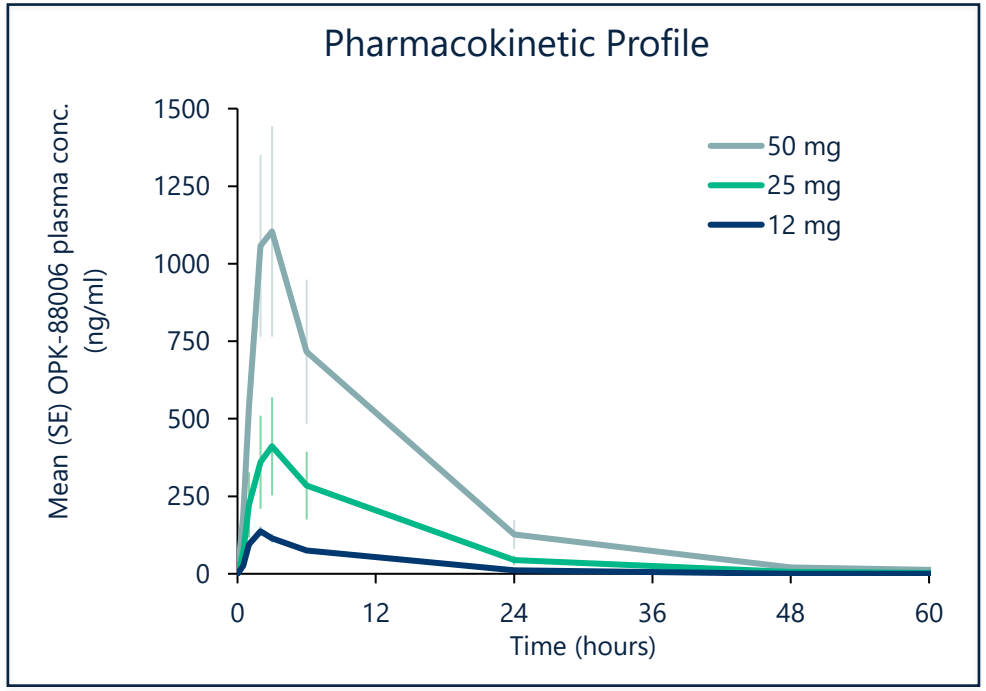
Robust Oral Bioavailability: Plasma concentrations following a single oral dose were similar to the reported clinical steady state concentration following a 2.4 mg subcutaneous dose of Wegovy (semaglutide)

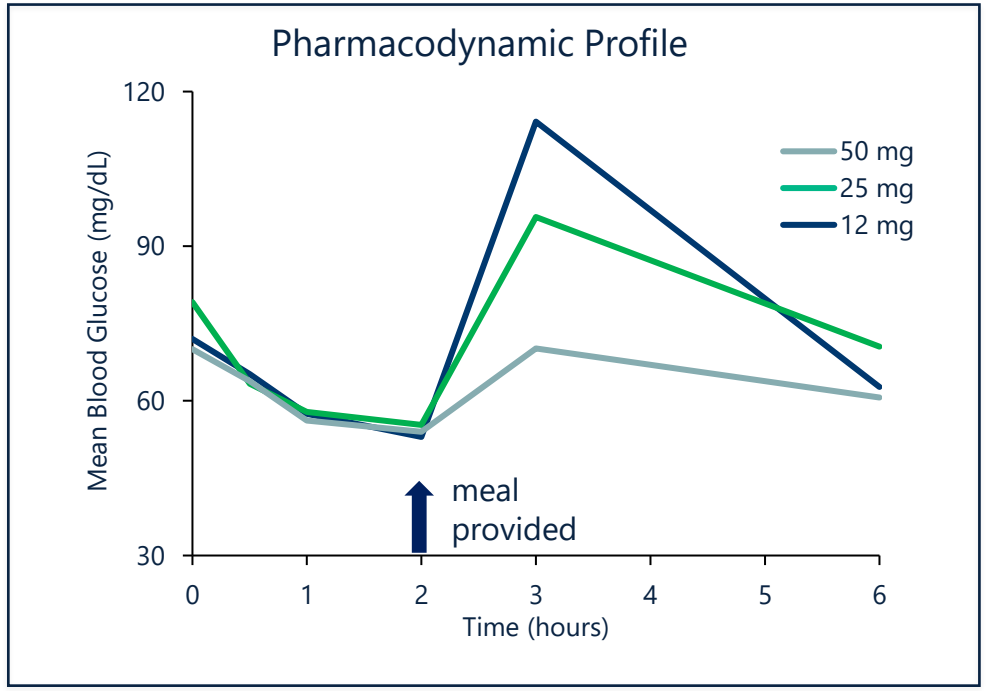
Pharmacologic Effect: Statistically significant reduction in plasma glucose levels compared with placebo in rats

Favorable Profile: PK profile and bioavailability observed in vivo appears suitable for daily tablet dosing using Entera's N-Tab® technology platform

N-Tab® Oral OXM IND filing expected pursuant to SC OXM filing (2026E)

EB618: Dose-Proportional PK/PD Effect in Non-Human Primate DRF Study





- **Robust bioavailability -** dose-proportional systemic exposure and low variability
- **Well tolerated** - no safety concerns were identified, at doses exceeding the anticipated clinical dose range by more than tenfold (mg/kg body weight)
- **A dose-proportional pharmacologic effect** on postprandial blood glucose levels

Burshtein et. al., EB618, First-in-Class Oral Tablet of Dual GLP-1/Glucagon Receptor Agonist for Patients with Obesity and Metabolic Disorders: Results from PK-PD Study in Non-Human Primates, [Poster presentation]. ENDO 2026: The Endocrine Society Annual Meeting, Chicago, IL, USA.

First Oral Once Daily GLP-2 Tablet for Patients with SBS

Oral GLP-2 offers less-invasive administration with titratable dosing for personalized treatment in this rare and diverse condition ~30k patients across US and EU

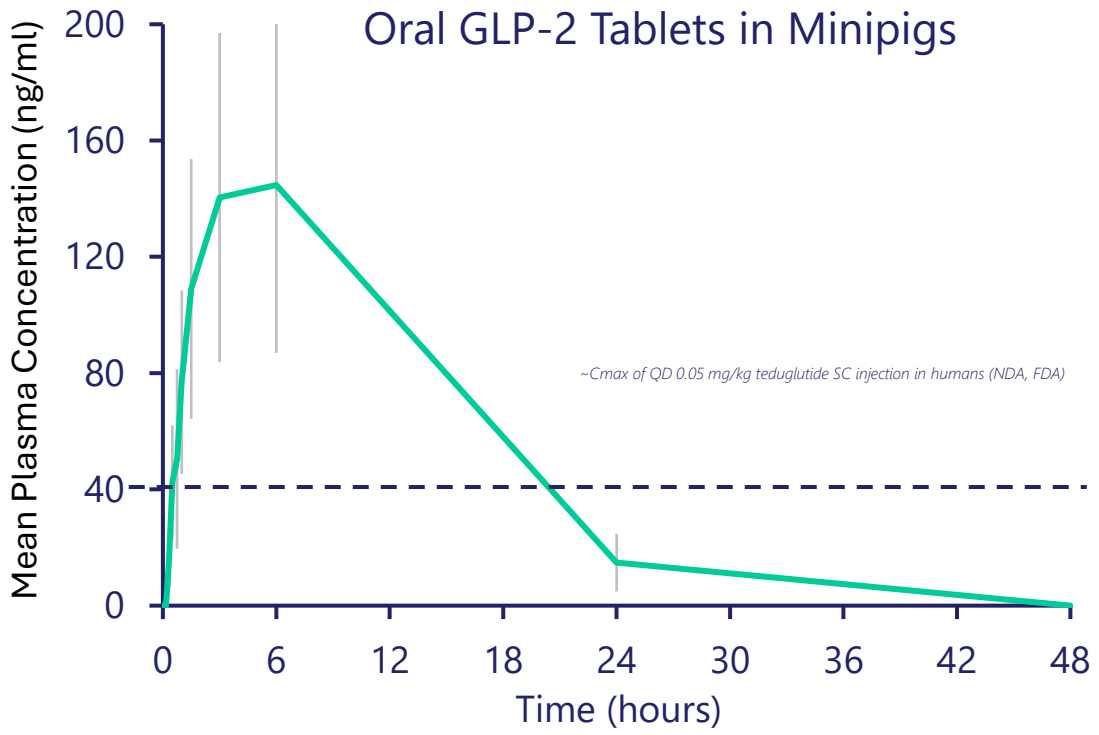
Entera Bio
OPKO

A FIRST- IN-CLASS ORAL GLP-2 ANALOG FOR TREATMENT OF SHORT BOWEL SYNDROME

LB055

Gregory Burshtein[1], Constantin Itin[1], Daniel Pery[1], Eli Reichman[1], Hillel Galitzer[1], Michal Kushnir[1], Ahuva Bar-Ilan[2], Moran Golan[2], Laura Moschcovich[2], Miri Zakar[2], Amit Rivkin[2], Moran Levy[2], Miranda Toledano[1], Jane Hsiao[3]

[1] Entera Bio Ltd., Jerusalem, Israel, [2] OPKO Biologics Ltd. Kiryat Gat, Israel, [3] OPKO Health Inc. Miami, FL, USA



Extended Plasma Half-Life: Approximately 15 hours, an 18-fold improvement over teduglutide (Gattex®), the only approved GLP-2 therapy

Robust Oral Bioavailability: Peak plasma concentrations reached ~200 ng/ml (Cmax), substantially exceeding the reported Cmax of daily teduglutide subcutaneous injection in humans

Prolonged Exposure: Systemic exposure (AUC ~2 h*µg/ml) was maintained for more than 24 hours with relatively low variability, supporting once-daily oral dosing

Pharmacologic effect: Rodent repeat-dose PK/PD studies showed clear pharmacologic activity in intestinal tissue

Favorable Safety Profile: No signs of toxicity were observed in preclinical studies

Thank you

IR@enterabio.com