



Entera Bio Announces Additional Presentation of EB613 Phase 3 Study Design at the ASBMR 2026 Annual Meeting

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TEL AVIV, Israel, Oct. 08, 2026 (GLOBE NEWSWIRE) -- Entera Bio Ltd. (NASDAQ: ENTX) ("Entera" or the "Company"), a leader in the development of oral peptides, today announced that an additional presentation highlighting the clinical study design for its planned Phase 3 registrational trial of EB613 (oral PTH(1-34) tablet) will be presented at the American Society for Bone and Mineral Research (ASBMR) 2026 Annual Meeting, taking place October 9-12, 2026, at the John B. Hynes Veterans Memorial Convention Center in Boston, Massachusetts.

The additional presentation, titled "EB613 [Oral PTH(1-34)]: The First Oral Anabolic Tablet for the Treatment of Osteoporosis: Phase 3 Study Design," has been accepted as a late-breaking poster presentation (Presentation Number: Sun-522) during Late Breaking Poster Session II on Sunday, October 11, 2026. Led by Dr. Felicia Cosman, Professor of Medicine at Columbia University and an internationally recognized osteoporosis authority, the presentation will outline the study design and clinical methodology for Entera's planned single, randomized, double-blind, placebo-controlled Phase 3 registrational study evaluating EB613 in postmenopausal women with osteoporosis.

This presentation is in addition to the Company's Phase 2 clinical data for EB613 which was selected for a Plenary Poster presentation at the meeting, evaluating treatment effects in postmenopausal women with osteoporosis at high risk of fracture.

Presentation Details:

Abstract Title: EB613 [Oral PTH(1-34)]: The First Oral Anabolic Tablet for the Treatment of Osteoporosis: Phase 3 Study Design

Session Title: Late Breaking Poster Session II

Presentation Number: Sun-522

Session Date/Time: Sunday, October 11, 2026, 2:00-3:30 p.m. ET

Abstract Title: EB613, Oral PTH(1-34) Tablet, Shows Markedly Enhanced Anabolic Effects in Postmenopausal Women with Osteoporosis at High Risk of Fracture: Insights from Phase 2 Trial

Session Title: Welcome Reception and Plenary Poster Session (Plenary Poster Presentation)

Presentation Number: Fri-026

Session Date/Time: Friday, October 9, 2026, 5:30-7:00 p.m. ET

Session Title: Poster Session II

Presentation Number: Sun-026

Session Date/Time: Sunday, October 11, 2026, 2:00-3:30 p.m. ET

About EB613

Substantial evidence supports the efficacy of anabolic therapies over anti-resorptive agents for lowering fracture risk in osteoporosis patients at high risk. However, all available anabolic therapies are administered by subcutaneous (SC) injection and used in a minority of eligible patients. Entera's EB613 program (oral PTH(1-34), teriparatide) is being developed as the first oral, once-daily anabolic tablet treatment for osteoporosis. Entera completed a Phase 2, 6-month, 161-patient, placebo-controlled study that met all biomarker and BMD endpoints without significant safety concerns in women with postmenopausal osteoporosis or low BMD (JBMR 2024). EB613 produced rapid dose-proportional increases in biochemical markers of bone formation, reductions in markers of bone resorption, and increases in lumbar spine, total hip, and femoral neck BMD. The effects of EB613 on trabecular and cortical bone using 3D-DXA showed increases with EB613 compared to placebo on a variety of indices, including integral volumetric BMD and trabecular volumetric BMD, cortical thickness, and cortical surface BMD. Mechanistically, the findings suggest that bone strengthening and fracture resistance may occur rapidly with EB613. Furthermore, the data are consistent with that of published subcutaneous teriparatide at the 6-month time point.

About Entera

Entera is a clinical stage company focused on developing oral peptide and protein replacement therapies for significant unmet medical needs where an oral tablet form holds the potential to transform the standard of care. The Company leverages a disruptive and proprietary technology platform (N-Tab®) and its pipeline of first-in-class oral peptide programs. The Company's most advanced product candidate, EB613 (oral PTH(1-34)), is being developed as the first oral, osteoanabolic (bone building) once-daily tablet for osteoporosis. A placebo-controlled, dose-ranging Phase 2 study of EB613 tablets (n = 161) met primary (PD/bone turnover biomarker) and secondary endpoints (BMD). Entera is also developing the first oral Long

Acting PTH(1-34) tablet as a replacement therapy for patients with hypoparathyroidism (EB612), the first oral oxyntomodulin, a dual targeted GLP1/glucagon peptide tablet for the treatment of obesity and metabolic syndromes; and the first oral GLP-2 tablet as an injection-free alternative for patients suffering from rare malabsorption conditions such as short bowel syndrome in collaboration with OPKO Health, Inc. For more information on Entera, visit www.enterabio.com or follow us on [LinkedIn](#), [Twitter](#), and [Facebook](#).

Cautionary Statement Regarding Forward Looking Statements

Various statements in this press release 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 press release regarding our prospects, plans, financial position, business strategy, clinical development activities, collaboration arrangements and expected financial and operational results are 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. Important factors that could cause actual results to differ materially from those reflected in Entera's forward-looking statements include, among others: changes in the interpretation of clinical data; results of our clinical trials; the FDA's interpretation and review of our results from and analysis of our clinical trials; unexpected changes in our ongoing and planned preclinical development and clinical trials, the timing of and our ability to make regulatory filings and obtain and maintain regulatory approvals for our product candidates; the potential disruption and delay of manufacturing supply chains; loss of available workforce resources, either by Entera or its collaboration and laboratory partners; impacts to research and development or clinical activities that Entera may be contractually obligated to provide; overall regulatory timelines; the size and growth of the potential markets for our product candidates; the scope, progress and costs of developing Entera's product candidates; Entera's reliance on third parties to conduct its clinical trials; Entera's ability to establish and maintain development and commercialization collaborations; Entera's operation as a development stage company with limited operating history; Entera's competitive position with respect to other products on the market or in development for the treatment of osteoporosis, hypoparathyroidism, short bowel syndrome, obesity, metabolic conditions and other disease categories it pursues; Entera's ability to continue as a going concern absent access to sources of liquidity; Entera's ability to obtain and maintain regulatory approval for any of its product candidates; Entera's ability to comply with Nasdaq's minimum listing standards and other matters related to compliance with the requirements of being a public company in the United States; Entera's intellectual property position and its ability to protect its intellectual property; and other factors that are described in the "Cautionary Statement Regarding Forward-Looking Statements," "Risk Factors" and "Management's Discussion and Analysis of Financial Condition and Results of Operations" sections of Entera's most recent Annual Report on Form 10-K filed with the SEC, as well as Entera's subsequently filed Quarterly Reports on Form 10-Q and Current Reports on Form 8-K. There can be no assurance that the actual results or developments anticipated by Entera will be realized or, even if substantially realized, that they will have the expected consequences to, or effects on, Entera. Therefore, no assurance can be given that the outcomes stated or implied in such forward-looking statements and estimates will be achieved. Entera cautions investors not to rely on the forward-looking statements Entera makes in this press release. The information in this press release is provided only as of the date of this press release, and Entera undertakes no obligation to update or revise publicly any forward-looking statements, whether as a result of new information, future events or otherwise, except to the extent required by law.

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