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2Mechanismof Action Aug 16 2024 YORVIPATHis a prodrug of parathyroid hormone PTH 1 34 administered once daily designed to provide parathyroid hormone levels within the normal physiological range across the 24 hour dosing period.YORVIPATHwas granted marketing authorization by the European Commission EC in November 2023 as a PTH replacement therapy indicated for thetreatmentof adults withchronichypoparathyroidism The FDA has approvedYorvopath palopegteriparatide injection for subcutaneous use in adults withhypoparathyroidism.Yorvopathwas not studied in adults with acute post surgicalhypoparathyroidism Oct 27 2025 14.1Treatmentof Adults withHypoparathyroidism The effectiveness and safety ofYORVIPATHin adults withhypoparathyroidismwere evaluated in a 26 week randomized double blind Jan 29 2025 What Does Research Show AboutYorvopathforHypoparathyroidism? The FDA approvedYorvopathbased on evidence from a clinical trial involving 82 adults withchronichypoparathyroidism Sep 15 2023 OverviewYorvopathis a hormone replacement medicine for treating adults withchronichypoparathyroidism

In patients with this condition the parathyroid glands in the neck do not produce enough parathyroid hormone PTH which controls the level of calcium in the blood

As a result patients have low levels of calcium and may experience problems with bones muscles the heart kidneys and Aug 16 2024 The Food and Drug Administration FDA has approvedYorvopath palopegteriparatide developed as TransCon PTH for thetreatmentofhypoparathyroidism in adults.Yorvopath is a once daily long acting prodrug of parathyroid hormone PTH [1 34] designed to restore physiologic levels of PTH for 24 hours Aug 10 2024 YorvopathOn August 12th 2024 Ascendis Pharma announced that the FDA had approvedYORVIPATH palopegteriparatide their new parathyroid hormone as atreatmentforchronichypoparathyroidism in adults in the United States

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