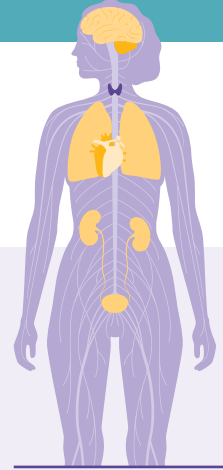




HypoPARAthyroidism (Also called "hypoPARA")



What Is HypoPARAthyroidism?

Defined as hypocalcemia in the presence of an undetectable, low, or insufficient parathyroid hormone (PTH) level.

Confirmed by blood tests on 2 occasions, at least 2 weeks apart.

Abnormalities that support diagnosis:

- Hyperphosphatemia
- Low serum level of 1,25 dihydroxy Vitamin D
- History of thyroid or neck surgery
- High 24-hour urine calcium levels
- Caution - Biotin (vitamin B7) intake can lead to falsely low PTH values in a few assays

Postsurgical hypoPARA is considered permanent (chronic) if it persists >12 months after surgery.

Types of HypoPARAthyroidism

Postsurgical (78% of cases)

Nonsurgical (less common)

- Genetic
- Idiopathic
- Functional (high or low serum magnesium)
- Destruction of glands (i.e. copper overload, hemochromatosis, radiation therapy)

Note: PseudohypoPARAthyroidism is a rare inherited disorder that mimics hypoPARAthyroidism, characterized by resistance to PTH. Instead of having low PTH levels, people with pseudohypoPARAthyroidism have elevated levels of PTH.



Body Systems Affected by HypoPARAthyroidism



Renal

- Nephrocalcinosis
- Kidney stones
- Chronic kidney disease
- Elevated serum phosphate level



Peripheral Nervous

- Paresthesia
- Muscle cramps
- Tetany



Neuropsychiatric

- Cognitive dysfunction
- Poor quality of life
- Symptoms of anxiety and depression
- Poor memory
- Brain fog



Central Nervous

- Seizures
- Brain calcifications
- Parkinsonism or dystonia



Cardiovascular

- Arrhythmias
- Hypocalcemia-associated dilated cardiomyopathy



Respiratory

- Laryngospasm
- Bronchospasm or wheezing



Ophthalmological

- Cataracts
- Papilledema



Dental

- Altered tooth morphology

Causes of HypoPARA¹

- 78% Postsurgical
- 9% Other Causes
- 7% Genetic
- 6% Idiopathic



¹ Clarke BL, Brown EM, Collins MT, et al. Epidemiology and diagnosis of hypoparathyroidism. J Clin Endocrinol Metab. 2016;101(6):2284-2299. doi:10.1210/jc.2015-390

Conventional Therapy

Goal of conventional therapy with calcium and active vitamin D is to raise serum calcium into the lower half of or just below the normal reference range, alleviate symptoms of hypocalcemia, avoid hypercalciuria, and maintain normal serum phosphate level.

Routine Monitoring

Every 3-4 months

- Serum calcium (albumin corrected or ionized)
- Magnesium
- Serum creatinine/eGFR
- Phosphate

Every 6-24 months

- 24-hour urine calcium and creatinine
- 25 OH(D)

Baseline Tests

- Renal imaging with ultrasound and or x-ray (KUB)
- Eye exam

Repeat Tests

- Renal imaging if patient has kidney stones or kidney disease and or high 24-hour urine calcium levels (>400mg/day and or low urine citrate levels (<300 mg/day), calcifications or stones on imaging, or declining renal function.
- Ophthalmologic exam if visual symptoms

- **Calcium citrate or calcium carbonate**
 - Not more than 500mg–600mg per dose.
 - Best taken with meals to control serum phosphate levels.
- **Vitamin D analogues: calcitriol or alphacalcidol**
 - Consider cholecalciferol or ergocalciferol to maintain 25-hydroxyvitamin D (25(OH)D) levels in normal range.

Additional Tests

- **DXA BMD** (Dual X-ray absorptiometry bone mineral density) is not needed routinely and may in fact not reflect bone strength accurately given most hypoPARA patients tend to have overly dense bones without a high fracture risk.
- **Risk of worsening hypocalcemia:** In hypoPARA patients who also have osteoporosis, BE VERY CAUTIOUS USING BISPHOSPHONATE RX's (such as Reclast®, Fosamax®, Actonel®, or Boniva®) as well as Prolia® (denosumab) DUE TO THE RISK OF HYPOCALCEMIA!



This document gives an overview of basic facts about hypoPARAthyroidism, its diagnosis, and typical treatment options. While this brochure contains important information about hypoPARAthyroidism, **the patient's individual course of testing, treatment, and follow-up may vary for many reasons.**

HypoPARAthyroidism Association

Our mission is to improve the lives of people impacted by hypoPARAthyroidism through education, support, research, and advocacy. **Learn more at www.hypopara.org.**

Board

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Medical Advisory Board

We are fortunate to have a distinguished Medical Advisory Board comprised of professionals with world-recognized expertise in hypoPARAthyroidism. They provide valuable counsel and support of HPA goals in education, treatment, and research.

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Resources

PubMed PMID: [26943719](https://pubmed.ncbi.nlm.nih.gov/26943719/)

PubMed PMID: [26135962](https://pubmed.ncbi.nlm.nih.gov/26135962/)

PubMed PMID: [30540559](https://pubmed.ncbi.nlm.nih.gov/30540559/)

This brochure combines the significant efforts of HPA's Medical Advisory Board and Board members. We greatly appreciate everyone's expertise and support.

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