



Parathyroid Hormone (PTH) (C-Terminal) Antibody - With BSA and Azide

Mouse Monoclonal Antibody [Clone PTH/1174] Catalog # AH12155

Product Information

ApplicationIHC, IF, FCPrimary AccessionP01270Other Accession5741, 37045ReactivityHumanHostMouseClonalityMonoclonal

Isotype Mouse / IgG2b, kappa

Clone Names PTH/1174
Calculated MW 12861

Additional Information

Gene ID 5741

Other Names Parathyroid hormone, PTH, Parathormone, Parathyrin, PTH

Application Note IHC~~1:100~500 IF~~1:50~200 FC~~1:10~50

Storage Store at 2 to 8°C.Antibody is stable for 24 months.

Precautions Parathyroid Hormone (PTH) (C-Terminal) Antibody - With BSA and Azide is

for research use only and not for use in diagnostic or therapeutic procedures.

Protein Information

Name PTH {ECO:0000303|PubMed:35932760, ECO:0000312|HGNC:HGNC:9606}

Function Parathyroid hormone elevates calcium level by dissolving the salts in bone

and preventing their renal excretion (PubMed: <u>11604398</u>, PubMed: <u>35932760</u>). Acts by binding to its receptor, PTH1R, activating G protein-coupled receptor

signaling (PubMed: 18375760, PubMed: 35932760). Stimulates

[1-14C]-2-deoxy-D-glucose (2DG) transport and glycogen synthesis in

osteoblastic cells (PubMed:21076856).

Cellular Location Secreted

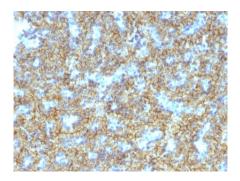
Background

Epitope of this MAb maps in the C-terminus of PTH, a hormone produced by the parathyroid gland that regulates the concentration of calcium and phosphorus in extracellular fluid. This hormone elevates blood Ca2+ levels by dissolving the salts in bone and preventing their renal excretion. It is produced in the parathyroid gland as an 84 amino acid single chain polypeptide. It can also be secreted as N-terminal truncated fragments or C-terminal fragments after intracellular degradation, as in case of hypercalcemia. Defects in this gene are a cause of familial isolated hypoparathyroidism (FIH); also called autosomal dominant hypoparathyroidism or autosomal dominant hypocalcemia. FIH is characterized by hypocalcemia and hyperphosphatemia due to inadequate secretion of parathyroid hormone. Symptoms are seizures, tetany and cramps. FIH exist both as autosomal dominant and recessive forms of hypoparathyroidism.

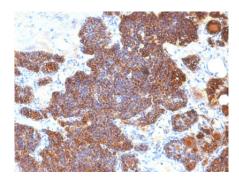
References

Watson, P.H. and Hanley, D.A. 1993. Parathyroid hormone: regulation of synthesis and secretion. Clin. Invest. Med. 16: 58-77.

Images



Formalin-fixed, paraffin-embedded human Parathyroid stained with PTH Monoclonal Antibody (PTH/1174).



Formalin-fixed, paraffin-embedded human Parathyroid stained with PTH Monoclonal Antibody (PTH/1174).

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.