

Parathyroid Hormone (PTH) (ABT44R) rabbit mAb

Catalog No: YM7188

Reactivity: Human;

Applications: IHC; ELISA

Target: Parathyroid Hormone

Fields: >>Neuroactive ligand-receptor interaction;>>Parathyroid hormone synthesis,

secretion and action;>>Endocrine and other factor-regulated calcium

reabsorption;>>Rheumatoid arthritis

Gene Name: PTH

Protein Name : Parathyroid hormone (PTH) (Parathormone) (Parathyrin)

Human Gene Id: 5741

Human Swiss Prot

No:

Immunogen: Synthesized peptide derived from human Parathyroid Hormone (PTH) AA

range:50-115

P01270

Specificity: This antibody detects endogenous levels of Parathyroid Hormone

Formulation: PBS, 50% glycerol, 0.05% Proclin 300, 0.05%BSA

Source: Monoclonal, Rabbit IgG1, Kappa

Dilution: IHC 1:100-500, ELISA 1:5000-20000

Purification: Recombinant Expression and Affinity purified

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Background: This gene encodes a member of the parathyroid family of proteins. The encoded

preproprotein is proteolytically processed to generate a protein that binds to the parathyroid hormone/parathyroid hormone-related peptide receptor and regulates



blood calcium and phosphate levels. Excess production of the encoded protein, known as hyperparathyroidism, can result in hypercalcemia and kidney stones. On the other hand, defective processing of the encoded protein may lead to hypoparathyroidism, which can result in hypocalcemia and numbness. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Oct 2015],

Function:

disease:Defects in PTH are a cause of familial isolated hypoparathyroidism (FIH) [MIM:146200]. FIH exist both as autosomal dominant and recessive forms of hypoparathyroidism.,function:PTH elevates calcium level by dissolving the salts in bone and preventing their renal excretion.,online information:Parathyroid hormone entry,similarity:Belongs to the parathyroid hormone family.,

Subcellular Location :

Cytoplasmic, Membranous

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