# abcam

## Product datasheet

## Recombinant Human HPRT protein ab97411

## 2 Images

**Description** 

Product name Recombinant Human HPRT protein

Purity > 95 % SDS-PAGE.

ab97411 is purified using conventional chromatography techniques.

**Expression system** Escherichia coli

Accession P00492

Protein length Full length protein

Animal free No

**Nature** Recombinant

**Species** Human

**Sequence** *MGSSHHHHHH SSGLVPRGSH* MATRSPGVVI

SDDEPGYDLD LFCIPNHYAE DLERVFIPHG LIMDRTERLA

RDVMKEMGGH HIVALCVLKG GYKFFADLLD

YIKALNRNSD RSIPMTVDFI RLKSYCNDQS TGDIKVIGGD DLSTLTGKNV LIVEDIIDTG KTMQTLLSLV RQYNPKMVKV

ASLLVKRTPR SVGYKPDFVG FEIPDKFVVG YALDYNEYFR DLNHVCVISE TGKAKYKA

Predicted molecular weight 27 kDa

Amino acids 1 to 218

Tags His tag N-Terminus

## **Specifications**

Our **Abpromise guarantee** covers the use of **ab97411** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

**Applications** SDS-PAGE

Mass Spectrometry

Western blot

Mass spectrometry MALDI-TOF-TOF

Form Liquid

Preparation and Storage

#### Stability and Storage

Shipped at 4°C. Upon delivery aliquot and store at -20°C or -80°C. Avoid repeated freeze / thaw

cycles.

pH: 8.00

Constituents: 0.316% Tris HCl, 20% Glycerol (glycerin, glycerine)

#### **General Info**

**Function**Converts guanine to guanosine monophosphate, and hypoxanthine to inosine monophosphate.

Transfers the 5-phosphoribosyl group from 5-phosphoribosylpyrophosphate onto the purine. Plays

a central role in the generation of purine nucleotides through the purine salvage pathway.

Pathway Purine metabolism; IMP biosynthesis via salvage pathway; IMP from hypoxanthine: step 1/1.

Involvement in disease Defects in HPRT1 are the cause of Lesch-Nyhan syndrome (LNS) [MIM:300322]. LNS is

characterized by complete lack of enzymatic activity that results in hyperuricemia,

choreoathetosis, mental retardation, and compulsive self-mutilation.

Defects in HPRT1 are the cause of gout HPRT-related (GOUT-HPRT) [MIM:300323]; also known as HPRT-related gout or Kelley-Seegmiller syndrome. Gout is characterized by partial enzyme

activity and hyperuricemia.

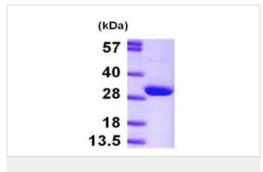
Sequence similarities

Belongs to the purine/pyrimidine phosphoribosyltransferase family.

**Cellular localization** 

Cytoplasm.

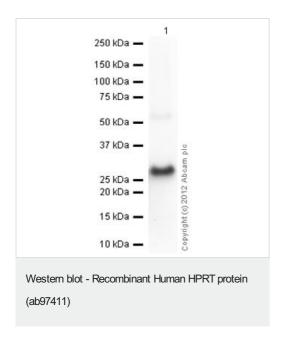
## **Images**



SDS-PAGE - Recombinant Human HPRT protein

(ab97411)

15% SDS-PAGE showing ab97411 at approximately 26.7kDa (3 $\mu$ g).



Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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