产品名称: Proteasome 20S alpha 5/PSMA5 Rabbit

Monoclonal Antibody 产品货号: AMRe87371



产品概述 (Summary)

产品名称 (Production Name) Proteasome 20S alpha 5/PSMA5 Rabbit Monoclonal Antibody

描述 (**Description**) Recombinant rabbit monoclonal antibody

宿主 (Host) Rabbit 应用 (Application) WB,IHC

种属反应性 (Reactivity) Human, Mouse, Rat

产品性能 (Performance)

偶联物 (Conjugation)Unconjugated修饰 (Modification)Unmodified

同种型 (Isotype) IgG

克隆 (Clonality) Monoclonal 形式 (Form) Liquid

Store at 4°C short term. Aliquot and store at -20°C long term. Avoid 存放说明 (Storage)

freeze/thaw cycles.

Supplied in 50mM Tris-Glycine(pH 7.4), 0.15M NaCl, 40% Glycerol, 0.01%

储存溶液 (Buffer) sodium azide and 0.05% protective protein. Stable for 12 months from date

of receipt.

纯化方式 (Purification) Affinity Purification

免疫原信息 (Immunogen)

基因名 (Gene Name) Proteasome 20S alpha 5/PSMA5

别名 (Alternative Names) PSC5; ZETA

基因 ID (Gene ID) 5686 蛋白 ID (SwissProt ID) P28066.

产品应用 (Application)

稀释比 (Dilution Ratio) WB 1:1000-1:5000,IHC 1:100-1:200

蛋白分子量 (Molecular Weight) Calculated MW:26 kDa; Observed MW:26 kDa

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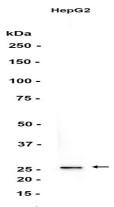


研究背景 (Background)

The proteasome is a multicatalytic proteinase complex with a highly ordered ring-shaped 20S core structure. The core structure is composed of 4 rings of 28 non-identical subunits; 2 rings are composed of 7 alpha subunits and 2 rings are composed of 7 beta subunits. Proteasomes are distributed throughout eukaryotic cells at a high concentration and cleave peptides in an ATP/ubiquitin-dependent process in a non-lysosomal pathway. An essential function of a modified proteasome, the immunoproteasome, is the processing of class I MHC peptides. This gene encodes a member of the peptidase T1A family, that is a 20S core alpha subunit. Multiple alternatively spliced transcript variants encoding two distinct isoforms have been found for this gene. [provided by RefSeq, Dec 2010]

研究领域 (Research Area)

图片 (Image Data)



Western blot analysis of extracts from HepG2 cells using AMRe87371 at 1:1000.

注意事项 (Note)

For research use only.

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