

Product datasheet for AR05053PU-N

Hyaluronidase Bovine Protein

Product data:

Product Type: Recombinant Proteins

Description: Hyaluronidase bovine protein, 0.5 g

Species: **Bovine**

Concentration: lot specific

Buffer: Presentation State: Purified

State: Liquid purified protein

Bioactivity: Biological: A unit is defined as the amount of enzyme that liberates one micromole of N-

acetylglucosamine per minute at 37°C and pH 4.0.

Specific: 315 U/mg protein

Preparation: Liquid purified protein

ELISA. **Applications:**

Protein Description: Hyaluronidase catalyzes the depolymerization of mucopolysaccharides, hyaluronic acid, and

the chondroitin sulfates A and C. The enzyme is widely distributed in animal tissues but is

found in great concentrations in the Bovine and Ovine testes.

Storage: Store the protein at -20°C.

Avoid repeated freezing and thawing.

Stability: Shelf life: one year from despatch

RefSeq: NP 149349

Locus ID: 3373

UniProt ID: Q12794, A0A024R2X3

Cytogenetics: 3p21.31

Synonyms: HYAL-1; LUCA1; MPS9; NAT6



OriGene Technologies, Inc. 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com



Hyaluronidase Bovine Protein - AR05053PU-N

Summary: This gene encodes a lysosomal hyaluronidase. Hyaluronidases intracellularly degrade

hyaluronan, one of the major glycosaminoglycans of the extracellular matrix. Hyaluronan is thought to be involved in cell proliferation, migration and differentiation. This enzyme is active at an acidic pH and is the major hyaluronidase in plasma. Mutations in this gene are associated with mucopolysaccharidosis type IX, or hyaluronidase deficiency. The gene is one

of several related genes in a region of chromosome 3p21.3 associated with tumor

suppression. Multiple transcript variants encoding different isoforms have been found for

this gene. [provided by RefSeq, Jul 2008]

Protein Families: Secreted Protein

Protein Pathways: Glycosaminoglycan degradation, Lysosome, Metabolic pathways