

## GBM biotinylated

### Antigen Specification

**Product Number:** 20700

**Description:**

Human  $\alpha 3$  chain of collagen IV; identical with the antigen called "glomerular basement membrane antigen" (GBM). Biotinylated recombinant antigen for in vitro research and manufacturing use only.

**Immunological function:**

Binds IgG-type human auto-antibodies.

**Origin:**

Recombinant. Expressed by recombinant baculovirus (*Autographa californica* multiple nuclear polyhedrosis virus; AcMNPV) infection of *Spodoptera frugiperda* Sf9 insect cells.

**Expression construct:**

cDNA coding for a minicollagen version of the human collagen IV  $\alpha 3$  chain fused to a hexa-histidine purification tag. The term minicollagen designates the removal of most of the epitope-less triplehelical collagenous region (situated between the N-terminal 7S domain and the C-terminal noncollagenous NC1 domain), which is a requirement for recombinant production of this antigen.

**Biochemical tests:**

Electrophoresis (purity > 80%).

**Calculated molecular weight:**

42 kDa (for protein component)

**Calculated isoelectric point:**

pH 8.9

**Immunological tests/Functionality:**

Functional Streptavidin-based ELISA test (analysis of positive/negative samples).

**Recommended buffer/storage and handling conditions:**

Recommendations for storage buffer: neutral to slightly alkaline pH; due to purification workup under denaturing conditions presence of up to 0.02% SDS (or similar detergents) may be required for maintaining solubility.

Storage conditions: -70°C or below.

Repeated freeze/thaw cycles should be avoided.

**CAUTION:** It has been reported that the immunodominant epitope of GBM is a cryptic epitope that is not easily accessible to the corresponding autoantibodies. It is necessary to treat the protein under non-reducing conditions with a denaturant such as urea to unmask the epitopes (see Hellmark et al. in Autoantibodies, Peter, J.B. and Shoefeld, Y., eds., Elsevier B.V., 1996, pp 291-298).

This GBM antigen product is produced in dissociated form and does not require additional unmasking of the epitope before coating.

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