Protein G / Agarose

IgG binding protein

Catalog code: gel-agg-2, gel-agg-5 https://www.invivogen.com/protein-g-agarose

For research use only

Version 19B11-MM

ANITIDODY

PRODUCT INFORMATION

Contents

Protein G / Agarose is available in two quantities:

- 2 ml Protein G / Agarose provided as a 50% v/v gel slurry in 20% v/v ethanol (total volume 4 ml): gel-agg-2
- 5 ml Protein G / Agarose provided as a 50% v/v gel slurry in 20% v/v ethanol (total volume 10 ml); gel-agg-5

Storage and stability

Protein G / Agarose is shipped at room temperature. Store at 4° C. Product is stable for 12 months when properly stored in regeneration and storage buffer (20% (v/v) ethanol in phosphate buffered saline). **DO NOT FREEZE.**

Note: Protein G / Agarose can be reused at least 10 times.

DESCRIPTION

Protein G is an immunoglobulin-binding protein expressed in group C and G Streptococcal bacteria. It is a 65-kDa (G148 protein G) and a 58 kDa (C40 protein G) cell surface protein that has found application in purifying antibodies through its binding to the Fc region¹. Protein G binds to most mammalian IgGs through the Fc region, but some binding also occurs through the Fab region. Native protein G also binds albumin, however, because serum albumin is a major contaminant of antibody sources, the albumin binding site has been removed from recombinant forms of Protein G^2 .

Protein G / Agarose from InvivoGen uses the recombinant form of protein G lacking the albumin-binding region.

In Protein G / Agarose, the recombinant protein G is coupled to beads using a leak-resistant chemistry that provides a support with minimal non-specific binding.

Binding capacity of Protein G:

- 20 mg of human IgG per ml of gel
- 15 mg of rabbit IgG per ml of gel
- 6 mg of mouse IgG per ml of gel

1. Sjobring U. *et al.*, **1991.** Streptococcal Protein G. Gene structure and protein binding properties. J Biol Chem. 140:1194-1197. **2. Goward CR.** *et al.*, **1990.** Expression and purification of truncated recombinant streptococccal Protein G. Biochem J. 267:171-177.

ANTIBODY	PROTEIN G BINDING
Human Total IgG	++++
Human IgG1	++++
Human IgG2	++++
Human IgG3	++++
Human IgG4	++++
Human IgA	-
Human IgM	-
Human IgD	-
Human Fab	+
Human ScFv	-
Mouse Total IgG	++++
Mouse IgG1	+++
Mouse IgG2a	++++
Mouse IgG2b	++++
Mouse IgG3	++++
Mouse IgM	-
Rat Total IgG	+++
Rat IgG1	+++
Rat IgG2a	++++
Rat IgG2b	+
Rat IgG2c	++++
Cow Total IgG	++++
Cow IgG1	++++
Cow IgG2	++++
Goat Total IgG	++++
Goat IgG1	++++
Goat IgG2	++++
Sheep Total IgG	++++
Sheep IgG1	++++
Sheep IgG2	++++
Horse Total IgG	++++
Horse IgG(ab)	-
Horse IgG(c)	-
Horse IgG(T)	++++
Rabbit Total IgG	++++
Guinea Pig TotalIgG	+
Pig Total IgG	+
Dog Total IgG	+
Cat Total IgG	+
Chicken Total IgY	-
O	

DDOTEIN C DIVIDING

InvivoGen USA (Toll-Free): 888-457-5873 InvivoGen USA (International): +1 (858) 457-5873 InvivoGen Europe: +33 (0) 5-62-71-69-39 InvivoGen Hong Kong: +852 3622-3480

E-mail: info@invivogen.com



METHOD

Purification of immunoglobulins using immobilized Protein G / Agarose

Buffers

Equilibration and wash buffer: 10 mM sodium phosphate, 150 mM sodium chloride, pH 7.2

Elution buffer: 0.1 M glycine, pH 2-3

Neutralization buffer: 0.75 M sodium phosphate or 1 MTRIS, pH 7.5-9 **Regeneration and storage buffer:** 20% (v/v) ethanol in phosphate

buffered saline

Immunoglobulin purification procedure:

- 1. Pack $\dot{1}$ ml of immobilized Protein G / Agarose into a suitable column
- 2. Perform all chromatography steps at a flow rate of 0.5-1 ml/min, or under gravity flow.
- 3. Equilibrate the column with 5 ml of Equilibration and Wash Buffer.

Optional:

In the presence of certain denaturing agents such as urea or guanidine chloride, we recommend to dialyze sample against 100 volumes of equilibration and wash buffer.

- 4. Filter the sample using a 0.2 µm filter.
- 5. Load the sample onto the column.
- 6. Wash the column with 10 ml of equilibration and wash buffer.
- 7. Elute the column with 10 ml of elution buffer.
- $8.\,\mbox{Immediately adjust}$ the eluate to pH 7.5 by adding neutralization buffer.
- 9. Wash the column with 10 ml of equilibration and wash buffer. 10. Store Protein G / Agarose in regeneration and storage buffer at $4^{\circ}\mathrm{C}$

Note: The procedure outlined above can be scaled up or down as desired.

RELATED PRODUCTS

Catalog code
pfuse-hchg1
pfuse-hchg2
pfuse-hchg3
pfuse-hchg4
pfuse-mchg1
pfuse-mchg2a
pfuse-mchg2b
pfuse-mchg3
gel-pdm-2
gel-protl-2



E-mail: info@invivogen.com