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# Monoclonal

# Llama IgG, heavy chain (and/or Camelide IgG, heavy chain) purification kit (Code: LIKhG-FF KIT)

Price: 300 EUR/KIT

## KIT CONTENT

(sufficient for 8 purifications with 200 ml of cell culture supernatant and/or ascites fluid/each)

Llama IgG, heavy chain Binding Gel (Sepharose<sup>™</sup> fast flow) (Code : LIKhG-FF) : 5 ml gel column. Binding capacity: approx. 10 mg Llama IgG heavy chain/ml wet gel.

Purity: 95% by SDS-PAGE

Maximum pressure: 3 bars (43 psi, 0.3 MPa).

Gel life: approx. 50 cycles with routine regeneration.

- Llama IgG Binding Buffer (Code: BBLG) 2x concentrated: 1000 ml. Add 1000 ml of distilled water to have a total of 2000 ml before use.
- Llama IgG Elution Buffer (Code: EBLG) 4x concentrated: 125 ml. Add 375 ml of distilled water to have a total of 500 ml before use..
- Llama IgG Precipitating Agent (Code: PALG): 8 x 1 sachet of sufficient quantity for precipitating all IgG from 200 ml of Llama cell culture supernatant and/or ascites fluid.

#### **INSTRUCTIONS FOR USE**

- 1. Add with mild agitation 1 sachet of Precitating Agent (PALG) to 200 ml of Llama cell culture supernatant (and/or ascites fluid) for 15 minutes. Stop the agitation and allows to stand for 30 minutes at 4°C. Ce ntrifuge at 3000 g for 10 minutes. Discard the supernant from the pellet. Dissolve the pellet in 30 ml of Binding Buffer (BBLG). Such a sample is ready to be loaded into the column.
- 2. Equilibrate the column with 20 ml of Llama IgG Binding Buffer (BBLG).
- 3. Load the sample prepared in point 1 into the column prepared in point 2 at a flow rate of 30 ml/hour.
- 4. Wash the column with 200 ml of Llama IgG Binding Buffer (BBLG).
- 5. Elute the Llama IgG with the Llama IgG Elution Buffer (Code: EBLG) until the O.D. at 280nm of the eluent reaches the baseline level. Collect 10 fractions of 5 ml elution volume.
- 6. If you want an important concentration of Llama IgG without loss of its content, use our Protein concentration kit (Code: PC KIT).
- 7. Assay the elution fractions obtained in point 5, using the most appropriate system (SDS-PAGE, immunodiffusion, radioimmunoassay, Elisa...)

## REGENERATION OF THE LLAMA IGG BINDING GEL It is recommended to regenerate the gel after every 5 cycles of use.

1. Wash the column with 10x volumes of NaOH 0.1M.

- 2. Wash the column with 10x volumes of distilled water.
- 3. Equilibrate the column 10x volumes of PBS (50 mM K<sub>2</sub>HPO<sub>4</sub>, 150mM NaCl) pH 7.4.
- 4. Store the column at 4℃ in the presence of NaN 3 0.1% (w/v).
- 5. For the next use, see INSTRUCTION FOR USE as described above.

If you need sterile materials, the regeneration can be carried out as follows. STERILE REGENERATION OF THE LLAMA IGG BINDING GEL

## (GEL SANITIZATION) **AFTER EVERY 5 CYCLES OF USE**

- 1. Wash 1 volume of gel column with 5 volumes of acetic acid 1 M.
- 2. Wash this column with 10 volumes of sterile distilled water.
- 3. Wash this column with 5 volumes of NaOH 1M.
- 4. Wash this column with 10 volumes of sterile distilled water.
- 5. Wash this column with 10 volumes of PBS (50 mM K<sub>2</sub>HPO<sub>4</sub>, 150mM NaCl) pH 7.4; NaN<sub>3</sub> 0.1%(w/v).
  6. The sterile gel column is now ready to be re-used.

Affiland s.a. Rue de l'Yser, 304 B-4430 ANS (BELGIUM) **2**: +32 - 4 - 366 33 94 R.C. Liège: 212 804 TVA/VAT : BE 0479 907 203

Fax: +32 - 4 - 246 15 06