

Anti-Hafnia alvei RABBIT POLYCLONAL AB-T156

Example of ELISA protocol used to test Hafnia alvei:

- 1. Coating of Hafnia alvei $(4\mu g/ml)$ in maxisorp well plates (Nunc) with a solution of sodium carbonate buffer 0.05M (pH 9.6), during sixteen hours at 4°C.
- 2. Saturation of well plates with of a solution of PBS (pH 7.3) containing 5g/l of BSA (Acros) and 0.5% of Tween (one hour at 37°C).
- 3. Wash with PBS containing 0.5% of Tween (PBS Tween) (three times).
- 4. Preabsorbed Hafnia alvei serum will be diluted (1/1,000-1/5,000) in PBS Tween containing 2.5g/l BSA, 200µl by well plate (incubating during 2 hours at 37°C).
- 5. Wash with PBS Tween (three times).
- 6. 200μl of peroxidase-labeled goat anti-rabbit (Jackson) diluted (1/10,000) in a solution of PBS Tween containing 2.5g/l of BSA, will be applied by well plate (during one hour at 37°C).
- 7. Well plates will be rinsed with PBS Tween (three times).
- 8. And finally the peroxidase will be developed by incubating 200µl by well plate of a citrate 0.1M/phosphate 0.2M (pH 5) solution containing 0.4% of OPD (Sigma) and 0.03% of hydrogen peroxide (Acros) for ten minutes in the dark, after that, we will stop the reaction by the addition of 50µl of 2M HCl.
- 9. The optical density will be measured at 492nm.

Example of Western blot protocol used to test Hafnia alvei:

Membrane Blocking, Antiboby Incubations and Detection of Proteins

- 1. Saturate the blot membrane with TBS + 5% Blocker for 1 hour at 37°C while mixing.
- 2. Wash the membrane twice for 5 minutes in TBS Tween at 37°C.
- 3. Incubate the membrane with the antibody diluted (1/1,000-1/2,000) in TBS 0.5% Blocker for 2 hours at 37° C.
- 4. Wash the membrane three times for 5 minutes in TBS Tween at 37°C.
- 5. Incubate with a biotinylated secondary antibody diluted 1:1000 in TBS 0.5% Blocker for 2 hours at 37°C.
- 6. Wash the membrane three times for 5 minutes in TBS Tween at 37°C.
- 7. Incubate with Streptavidin-HRP 1μ g/ml in TBS 0.5% Blocker for 2 hours at room temperature.
- 8. Wash the membrane three times for 5 minutes in TBS at 37°C.
- 9. Incubate in TBS (200ml) + (50mg DAB in 25ml methanol) + (150mg 4-chloro-1-naphtol in 25ml methanol) + 50µl H2O2 30% for a maximum of 30 minutes in the dark.
- 10. Stop the reaction by addition of distilled water.

Blocker = skim milk (Biorad 170-6404)

TBS = 20 mM Tris base, 0.5M NaCl, pH 7.5

TBS Tween = TBS + 0.05% Tween 20