

This product is for research use only (not for diagnostic or therapeutic use)

contact: support@agrisera.com

Agrisera AB | Box 57 | SE-91121 Vännäs | Sweden | +46 (0)935 33 000 | www.agrisera.com

## Product no AS13 2716

# Anti-mAB-M | Mouse anti-human Abeta protein (3-10) region, oligomer-specific (clone 2D10,F6) **Product information**

Immunogen Synthetic peptide chosen from human Abeta protein (3-10) pregion, oligomer specific

Host Mouse

Clonality Monoclonal

**Subclass/isotype** IgG1, kappa light chain (clone number 2D10,F6)

**Purity** Immunogen affinity purified serum in PBS pH 7.4.

Format Lyophilized

Quantity 50 μg

**Reconstitution** For reconstitution add 50 μl of sterile water

Storage

For short-term storage, please add sodium azide and srote at +4°C. For long-term storage, store lyophilized/reconstituted at -20°C; once reconstituted make aliquots to avoid repeated freeze-thaw cycles. Please remember to spin the tubes briefly prior to opening them to avoid any losses that might occur from material adhering to the cap or sides of the tube.

Additional information Immunolocalization: human tissue was paraffin-embedded and sectioned. De-waxed and rehydrated in an ethanol gradient. Antigens were retrieved in sodium citrate buffer (pH 6) at 95 °C for 1 h. The tissue sections were separately incubated for 1 h at RT with primary antibody and antibody binding was visualized with IgG Preoxidase Reagent Kit.

This antibody is specific for human Amyloid-Beta oligomers.

### **Application information**

**Recommended dilution** 10 μg/ml (IL), 1-2 μg/ml (Dot), 2-4 μg/ml (ELISA capture)

Expected | apparent 4.5 kDa MW

Confirmed reactivity Human

Not reactive in No confirmed exceptions from predicted reactivity are currently known

Additional information Due to location of antigen used to elicit this antibody in 3-10 region, it should bind to full length APP.

Meilandt et al. (2019). Characterization of the selective in vitro and in vivo binding properties of crenezumab to Selected references oligomeric A[beta]. Alzheimers Res Ther. 2019 Dec 1;11(1):97. doi: 10.1186/s13195-019-0553-5.

Brännström et al. (2014). A Generic Method for Design of Oligomer-Specific Antibodies. PLoS ONE. DOI:

10.1371/journal.pone.0090857.

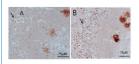
### **Application examples**

### Dot blot



Dot blot reaction of the binding capacity of mAB-M to fibrils, monomers and oligomers. Equal amounts of each sample were spotted on a nitrocellulose membrane and then dried. The membrane was blocked with 5% non-fat milk before incubated for 1 h with anti-mAB-M (25nM) and then with secondary antibody, anti-mouse HRP-conjugated (1:1500). The membrane was washed with PBS containing 0.25% Tween-20 before detection using ECL prime (GE Healthcare).

#### **Immunolocalization**





This product is for research use only (not for diagnostic or therapeutic use)

contact: support@agrisera.com

Agrisera AB | Box 57 | SE-91121 Vännäs | Sweden | +46 (0)935 33 000 | www.agrisera.com

IHC used to illustrate the lack of binding of mAB-M to plaques. Tissue sections from the human AD hippocampus were de-waxed and rehydrated in ethanol and then incubated with AS08 357 (A) and mAB-M(B) at RT for 1h. The immunoreactivity was detected with the anti-mouse Peroxidase Reagent Kit (ImmPRESS, Vector Laboratories, Inc.) and then developed using the ImmPACT AEC Peroxidase Substrate kit (Vector Laboratories, Inc.).