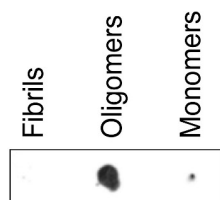
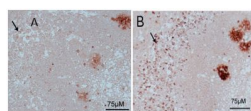


Product no **AS13 2716****Anti-mAB-M | Mouse anti-human Abeta protein (3-10) region, oligomer-specific (clone 2D10,F6)****Product information****Immunogen** | Synthetic peptide chosen from human Abeta protein (3-10) region, oligomer specific**Host** | Mouse**Clonality** | Monoclonal**Subclass/isotype** | 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 store 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 Peroxidase 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 MW** | 4.5 kDa**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.**Selected references** | [Meilandt et al. \(2019\)](#). Characterization of the selective in vitro and in vivo binding properties of crenezumab to oligomeric A[βeta]. *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

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.).