

Product no **AS16 3113****Transthyretin 56-61, amyloid specific (mouse monoclonal)****Product information**

<b>Immunogen</b>	Recombinant protein corresponding to the Human wild type Transthyretin. GPTGTGESKPLMVKVLDAVRGSPAINVAVHVFRKAADDTWEPFASGKTSESGELH GLTTEEEFVEGIYKVEIDTKSYWKALGISPFHEHAEEVFTANDSGPRRYTIAALLSPYS YSTTAVVTNPKE The epitope has been mapped to residue 56-61
<b>Host</b>	Mouse
<b>Clonality</b>	Monoclonal
<b>Subclass/isotype</b>	IgG1
<b>Purity</b>	Affinity purified in PBS pH 7.4.
<b>Format</b>	Lyophilized
<b>Quantity</b>	100 µg
<b>Reconstitution</b>	Add 100 µl sterile water to reconstitute to 1 mg/ml
<b>Storage</b>	Store lyophilized/reconstituted at 4 °C, 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.

**Application information**

<b>Recommended dilution</b>	1:1000 (ELISA), 1:500 (IHC), 1:1000 (WB)
<b>Expected   apparent MW</b>	155
<b>Confirmed reactivity</b>	Human Transthyretin Amyloids
<b>Not reactive in</b>	No confirmed exceptions from predicted reactivity are currently known
<b>Additional information</b>	Specifically reactive to the amyloid form of human Transthyretin. Epitope mapped to residue 56-61 which remains buried within the native fold of transthyretin but becomes exposed within its amyloid form. It has been suggested that that two distinct mechanisms of TTR-amyloidosis exists. The first, most common seen in wild type TTR Amyloidosis, consists of the full length TTR. Whereas the other type of amyloidosis mainly consists of the C-terminal region of the protein and is more common in mutant versions of TTR. Mouse IgG1 Anti-Transthyretin 56-61 (Amyloid Specific) epitope is located at the C-terminal strand of cleaved TTR and is suitable to detect amyloid formation derived from the C-terminal.
<b>Selected references</b>	<a href="#">Goldsteins et al. (1999)</a> . Exposure of cryptic epitopes on transthyretin only in amyloid and in amyloidogenic mutants. Proc Natl Acad Sci U S A. 1999 Mar 16; 96(6): 3108–3113