



## Anti-SQSTM1 Antibody

**Alternative Names:** A170, DMRV, FTDALS3, NADGP, OSIL, p60, p62, p62B, PDB3, ZIP3, A170, FTDALS3, OSIL, p60, p62, p62B, PDB3, ZIP3, EBI3-associated protein of 60 kDa, EBIAP

**Catalogue Number:** AB18-10057

**Size:** 100ug

### Background Information

Sequestosome-1 (SQSTM1) is an autophagy receptor that binds ubiquitin and regulates activation of the nuclear factor kappa-B (NF- $\kappa$ B) signaling pathway. SQSTM1 functions as a scaffolding/adaptor protein with TNF receptor-associated factor 6 (TRAF6). Mutations in SQSTM1 are also observed in familial ALS cases associated with aberrant TDP-43 inclusions and additionally mutations in SQSTM1 may result in Paget disease of bone. SQSTM1 may also be involved in cell differentiation, apoptosis, immune response and regulation of K<sup>+</sup> channels.

### Product Information

<b>Antibody Type:</b>	Polyclonal	<b>Host:</b>	Rabbit
<b>Isotype:</b>	IgG	<b>Species Reactivity:</b>	Human Mouse Rat
<b>Immunogen:</b>	A synthesized peptide		
<b>Format:</b>	PBS with 0.03% Proclin300, 50% glycerol, pH7.3.		
<b>Storage Conditions:</b>	6 months: 4°C. Long-term storage: -20°C. Avoid multiple freeze and thaw cycles.		
<b>Applications:</b>	WB WB 1:3000.		

### Additional Information

<b>Subcellular location:</b>	Cytoplasm Nucleus	<b>MW:</b>	48kDa
<b>Gene ID</b>	8878	<b>Uniprot ID:</b>	Q13501



## References

1. Fecto, F. et al. SQSTM1 mutations in familial and sporadic amyotrophic lateral sclerosis. *Arch. Neurol.* 68, 1440–1446 (2011).