Mouse Monoclonal Antibody to

LC₃ (microtubule-associated protein1 light chain 3B) clone 5F10

Order No ·	0231-100/LC3-5E1
Urder No.:	0231-100/LC3-3F1

Size (µg) Lot No.:

0231S

100





0

Autophagy is an alternative process of proteasomal degradation for some long-lived proteins or organelles. Alterations in the autophagic-lysosomal compartment have been linked to neuronal death in many neurodegenerative disorders as well as in transmissible neuronal pathologies (prion diseases). Genetic studies in yeast have shown that Autophagy-defective Gene-8 (Atg-8) represents a specific marker for autophagy. Among the four families of mammalian Atg8-related proteins only LC3 (microtubule-associated protein1 light chain 3) is expressed at sufficient high levels and efficiently recruited to autophagic vesicles in cells and tissues. During autophagy the cytoplasmic form, LC3-I is processed and recruited to autophagosomes, where LC3-II is generated by site specific proteolysis near to the C-terminus. Autophagic vacuoles have been also reported frequently in cardiomyopathies or muscle cells exposed to different experimental settings.

Mab LC3-5F10 specifically recognizes both forms of endogenous LC3, the cytoplasmic LC3-I (18 kDa) as well as the lipidated form generated during autophagosome and autophagolysosome formation: LC3-II (16 kDa). Immunocytochemical staining of cells with LC3-5F10 mab reveals the specific punctate distribution of endogenous LC3-II as a hallmark of autophagic activity.

Purification:	The antibody was purified from serum-free cell culture supernatant by subsequent ultrafiltration and size exclusion chromatography.
Formulation:	Lyophilized from 1ml PBS / 0.09% Na-azide / PEG and Sucrose.
Reconstitution:	Reconstitute with 1ml H ₂ O (15 min, RT).
Stability:	For long-term storage, freeze lyophilizate upon arrival (-20°C). Upon reconstitution, aliquote and freeze in liquid nitrogen; reconstituted antibody can be stored frozen at -80°C up to 1 year. Thaw aliquots at 37°C. Thawed aliquots may be stored at 4°C up to 3 months. Avoid repeated freeze / thaw cycles
Positive Control:	#0911: Cell lysate from untreated Neuro 2A
Immunoblotting:	0.5 µg/ml for HRPO/ECL detection <u>Recommended blocking buffer:</u> Casein/Tween 20 based blocking and blot incubation buffer, e.g. nanoTools product #3031-500/CPPT or #3031-3000/CPPT.
Immunoprecipitation:	ND
Immunocytochemistry:	Use at 1-10 µg/ml (paraformaldehyd/methanol fixation)
ELISA:	ND

All products are supplied for research and investigational use only. Not for use in humans or laboratory animals.

mab to LC3 #0260-100/LC3-2G6 mab to LC3 #0261-100/LC3-5H3 mab to Beclin #0240-100/Beclin-12B4

Alzheimer Disease

mab to BA4 (1-40), C-Terminus #0060-100/bA4(40)-5C3

mab to β A4 (1-42), C-Terminus #0061-100/bA4(42)-8G

mab to βA4 (1-40/42), C-Terminus #0062-100/bA4(40/42)-9F1

mab to BA4 (1-43), C-Terminus #0095-100/bA4(43)-6G12

mab to βA4, N-Terminus #0064-100/bA4N-19H5

mab to βA4, N-Terminus #0084-100/bA4N-19H11

mab to BA4, N-Terminus #0197-100/bA4N-11H3

For monoclonal antibodies against PKB/akt, and SAPK/jnk, please refer to our website at www.nanotools.de



Endogenous LC-3 punctae detected with anti LC-3 mab 5F10.

The majority of LC-3 was diffusely localized in unstimulated COS-7 cells, wheres punctated signals of LC-3 increase after induction of autophagy by vinblastin stimulation for 2 hr. Cells were fixed with paraformaldehyd followed by methanol treatment. Cells were permeabilized with 0.3% TritonX100. Endogenous LC-3 was detected with mab 5F10. (Images by courtesy of I. Ciechomska and A. Tolkovsky, University of Cambridge, UK).



