

ABGENT: A Leader of the market in Autophagy

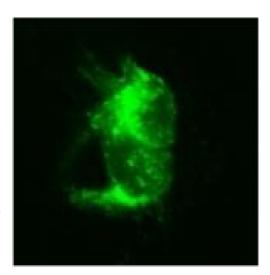
150 antibodies and peptides including almost all targets of autophagy

- Provide peptide induced autophagy
- Over 50 autophagy phospho-specific antibodies
- More than 100 antibodies about mTOR path
- Publish the reviews of pathway together with autophagy experts

AutoDOT

The new autophagy visualization dye!

- Superior to traditional monodansyl cadaverine staining
- Specific Staining of Autophagic Vacuoles
- Faster Penetration
- Higher Sensitivity/Greater Signal Endurance on Stored Slides
- Greater Resistance to Acid



Phosphorylation of Autophagy pathway

Paper published for Phosphorylation of Autophagy pathway:

"Regulation of the autophagy protein LC3 by phosphorylation"

Journal of JCB, 2010, volume 190, No.4:533-539

In this paper, we identified a new phosphorylation site on LC3 that reduces its recruitment and participation in autophagy. The result implicate LC3 phosphorylation as a novel switch that modulates its biological function in mammalian cells.

Our observations support the concept of a reserve pool of phosphorylated LC3 that can be rapidly recruited for autophagy in response to external stimuli such as nutrient deprivation or mitochondrial injury.



AutoDOT Example Staining Protocol

- 1. Grow cells overnight to approximately 70% confluency.
- 2. Induce autophagy in the test sample (protocol was tested using starvation-induced autophagy, but is expected to work with other forms of induction). Preserve a control sample where autophagy is not induced.
- 3. Prepare the staining solution by diluting stock AutoDOT 1:1000 in PBS.
- 4. Aspirate the media from the test and control cells and cover them in staining solution.
- 5. Incubate at 33 degrees C for 15 minutes.
- 6. Fix with 4% formaldehyde for 20 minutes.
- 7. Wash with PBS 3 times, 10 minutes each.
- 8. Immediately analyze and compare test and control samples by fluorescence microscopy using an inverted microscope equipped with a filter system (excitation filter: 380-420 nm, barrier filter: 450 nm).

For further protocols, consult J. Histochemistry & Cytochemistry 49(2): 177–185, 2001. AutoDOT is the trademarked name for the dye referred to as MDH in this citation.



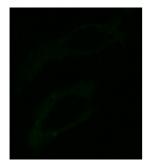
AutoDOT- The AutophagyVisualization Solution

Superior to traditional monodansyl cadaverine staining

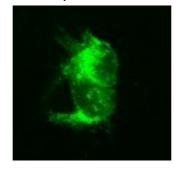
- √ Specific Staining of AutophagicVacuoles
- √ Faster Penetration
- √ Higher Sensitivity
- **V** Greater Signal Endurance on Stored Slides
- √ Greater Resistance to Acid
- √ Online Protocol

AutoDOT Staining in Mouse Cerebellar Cells

Untreated

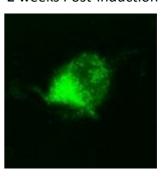


Same Day Induced Starvation



Stored Slides

2 weeks Post-induction



Citations: J. Histochemistry& Cytochemistry49(2): 177–185, 2001; AutoDOTis the trademarked name for the dye referred to as MDH in this citation

Cat No.	Concentration	Volume	Price
SM1000a	0.1M	50uL	\$50
SM1000b	0.1M	200 uL	\$150