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-20°C

#54831

PhosphoPlus® Beclin-1 (Ser30) Antibody Duet



Cell Signaling
TECHNOLOGY®

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Entrez-Gene ID #8678
UniProt ID #Q14457

New 04/21

For Research Use Only. Not For Use In Diagnostic Procedures.

Products Included	Product #	Quantity	Mol. Wt.	Isotype
Phospho-Beclin-1 (Ser30) (E1C4X) Rabbit mAb	35955	100 µl	60 kDa	Rabbit IgG
Beclin-1 (D40C5) Rabbit mAb	3495	100 µl	60 kDa	Rabbit IgG

See www.cellsignal.com for individual component applications, species cross-reactivity, dilutions, and additional application protocols.

Description: PhosphoPlus® Duets from Cell Signaling Technology (CST) provide a means to assess protein activation status. Each Duet contains an activation-state and total protein antibody to your target of interest. These antibodies have been selected from CST's product offering based upon superior performance in specified applications.

Background: Autophagy is a catabolic process for the autophagosomal-lysosomal degradation of proteins activated in response to nutrient deprivation and in neurodegenerative conditions (1). One of the proteins critical to this process is Beclin-1, the mammalian orthologue of the yeast autophagy protein Apg6/Vps30 (2). Beclin-1 can complement defects in yeast autophagy caused by loss of Apg6 and can also stimulate autophagy when overexpressed in mammalian cells (3). Mammalian Beclin-1 was originally isolated in a yeast two-hybrid screen for Bcl-2 interacting proteins and has been shown to interact with Bcl-2 and Bcl-xL, but not with Bax or Bak (4). While Beclin-1 is generally ubiquitously expressed, research studies have shown it is monoallelically deleted in 40-75% of sporadic human breast and ovarian cancers (5). Beclin-1 is localized within cytoplasmic structures including the mitochondria, although overexpression of Beclin-1 reveals some nuclear staining and CRM1-dependent nuclear export (6). Investigators have demonstrated that Beclin-1^{-/-} mice die early in embryogenesis and Beclin-1^{-/-} mice have a high incidence of spontaneous tumors. Stem cells from the null mice demonstrate an altered autophagic response, although responses to apoptosis appeared normal (7). Researchers have also found that overexpression of Beclin-1 in virally infected neurons *in vivo* resulted in significant protection against Sindbis virus-induced disease and neuronal apoptosis (4).

ULK1 phosphorylates Beclin-1 at Ser30 to induce autophagy in response to amino acid starvation, mTORC1 inhibition, and hypoxia (8).

Specificity/Sensitivity: Phospho-Beclin-1 (Ser30) (E1C4X) Rabbit mAb recognizes endogenous levels of Beclin-1 protein only when phosphorylated at Ser30. Beclin-1 (D40C5) Rabbit mAb detects endogenous levels of total Beclin-1 protein.

Source/Purification: Monoclonal antibody is produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Ser30 of human Beclin-1 protein, or with a synthetic peptide corresponding to residues surrounding Thr72 of human Beclin-1 protein.

Storage: Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at -20°C. *Do not aliquot the antibodies.*

Background References:

- (1) Reggiori, F. and Klionsky, D.J. (2002) *Eukaryot Cell* 1, 11-21.
- (2) Kametaka, S. et al. (1998) *J Biol Chem* 273, 22284-91.
- (3) Liang, X.H. et al. (1999) *Nature* 402, 672-6.
- (4) Liang, X.H. et al. (1998) *J Virol* 72, 8586-96.
- (5) Aita, V.M. et al. (1999) *Genomics* 59, 59-65.
- (6) Liang, X.H. et al. (2001) *Cancer Res* 61, 3443-9.
- (7) Yue, Z. et al. (2003) *Proc Natl Acad Sci USA* 100, 15077-82.
- (8) Park, J.M. et al. (2018) *Autophagy* 14, 584-597.

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Applications: W—Western IP—Immunoprecipitation IHC—Immunohistochemistry ChIP—Chromatin Immunoprecipitation IF—Immunofluorescence F—Flow cytometry E-P—ELISA-Peptide **Species Cross-Reactivity:** H—human M—mouse R—rat Hm—hamster Mk—monkey Mi—mink C—chicken Dm—D. melanogaster X—Xenopus Z—zebrafish B—bovine Dg—dog Pg—pig Sc—S. cerevisiae Ce—C. elegans Hr—Horse All—all species expected Species enclosed in parentheses are predicted to react based on 100% homology.