

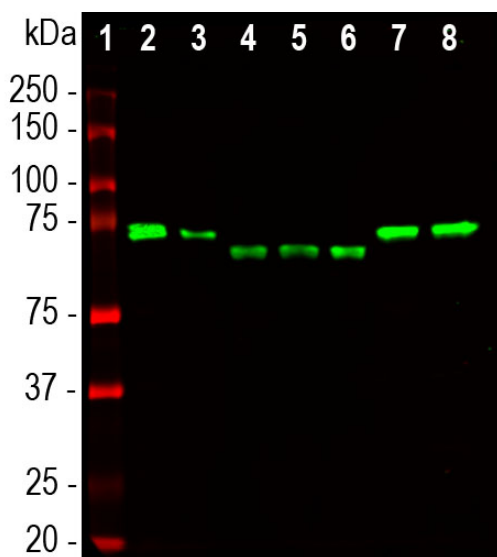
Ordering Information
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HGNC Name: UBQLN2
UniProt: Q9UHD9
RRID: AB_2572390
Immunogen: Human ubiquilin 2 expressed in and purified from *E. coli*
Format: Purified antibody at 1mg/mL in 50% PBS, 50% glycerol plus 5mM Na₂S₂O₃
Storage: Store at 4°C for short term, for longer term at -20°C. Avoid freeze / thaw cycles.
Recommended dilutions:
 WB: 1:1,000-1:2,000. IF/ICC and IHC: 1:1,000.

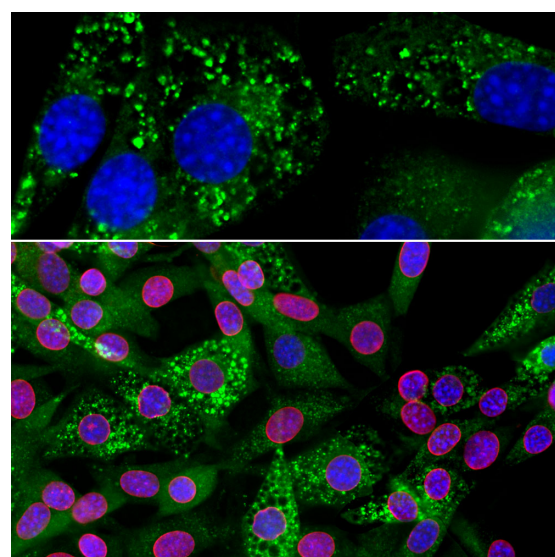
References:

1. Kleijnen MF, et al. The hPLIC proteins may provide a link between the ubiquitination machinery and the proteasome. *Molec. Cell* 6:409-19 (2000).
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8. Rutherford NJ, et al. Ubiquilin-1 and -2 are Predominantly Associated with Huntingtin Inclusions. *Brain Res.* 1524:62-73 (2013)

Applications	Host	Isotype	Molecular Wt.	Species Cross-Reactivity
WB, IF/ICC, IHC	Mouse	IgG1	66-68 kDa	Hu, Rt, Ms



Western blot analysis of different tissue and cell lysates using mouse mAb to ubiquilin 2, MCA-6H9, dilution 1:1,000 in green: [1] protein standard (red), [2] NIH-3T3, [3] C6, [4] HEK293, [5] HeLa, [6] SH-SY5Y, [7] rat whole brain, and [8] mouse whole brain. The band at 65-70kDa corresponds to ubiquilin 2 protein, which is known to differ between the human and rodent proteins.



Immunofluorescent analysis of an NIH-3T3 cell culture stained with mouse mAb to ubiquilin 2, MCA-6H9, dilution 1:1,000 in green, and costained with chicken pAb to lamin A/C, CPCA-Lamin A/C dilution 1:5,000 in red. The blue is DAPI staining of nuclear DNA. The cells were treated with 50µM of chloroquine, an inhibitor of autophagy, for 16 hours prior to staining. The MCA-6H9 antibody reveals punctate staining of ubiquilin 2 protein accumulated in lysosomes in the cytoplasm, while the lamin A/C antibody stains the nuclear lamina.

Background:

Ubiquilin 2, also known as PLIC2 and Chap1, is one of 4 members of the ubiquilin protein family, which regulate the degradation of cellular proteins through proteasome or autophagy-like pathways (1-3). All ubiquilins contain an N-terminal ubiquitin-like (UBL) domain and a C-terminal ubiquitin-associated (UBA) domain, while the central part of the molecules are highly variable. The UBL domains bind subunits of the proteasome, and the UBA domains binds to polyubiquitin chains that are typically conjugated onto proteins marked for proteosomal degradation. Mutations in the ubiquilin 2 gene leading to protein point mutations are implicated in certain forms of **amyotrophic lateral sclerosis (ALS)** and **Frontotemporal lobar degeneration (FTLD)** (4). Increased length of GC rich hexanucleotide repeats in a non-coding region of the *C9orf72* gene is the most common known genetic defect seen in patients suffering from ALS and FTLD (5,6). These patients have a distinct ubiquilin 2 pathology (7).

The MCA-6H9 antibody was made against full length recombinant human Ubiquilin 2. It can be used to track this protein by ELISA, on western blots and for IF, ICC and IHC (for IHC see data under "Additional Info" tab). A partial characterization of this antibody is described in a peer reviewed publication showing that ubiquilin 2 accumulations are seen on the inclusions of both Huntington's disease mouse models and patients (8). This publication shows that MCA-6H9 has little or no cross reactivity to the closely related ubiquilin 1.

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Abbreviation Key:

mAb—Monoclonal Antibody pAb—Polyclonal Antibody WB—Western Blot IF—Immunofluorescence ICC—Immunocytochemistry
 IHC—Immunohistochemistry E—ELISA Hu—Human Mo—Monkey Do—Dog Rt—Rat Ms—Mouse Co—Cow Pi—Pig Ho—Horse Ch—Chicken
 Dr—D. rerio Dm—D. melanogaster Sm—S. mutans Ce—C. elegans Sc—S. cerevisiae Sa—S. aureus Ec—E. coli.