

# LLGL1 (D2B5A) Rabbit mAb

✓ 100 µl  
(10 western blots)

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

Applications W, IF-IC Endogenous	Species Cross-Reactivity* H, Mk	Molecular Wt. 130 kDa	Isotype Rabbit IgG**
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**Background:** In *Drosophila*, lethal giant larvae (*lgl*), discs large (*dlg*), and scribble (*scrib*) act as tumor suppressor genes. Their loss of function in flies causes neoplastic overgrowth of larval brain tissue and imaginal epithelial cells hallmarked by disruption of the cytoskeletal network and cellular polarity (1,2). The human homolog of the *Drosophila lgl* protein, lethal giant larvae protein homolog 1 (LLGL1), is a cytoskeletal protein implicated in regulating cellular organization, migration, and cell polarity (3). As in *Drosophila*, decreased expression of LLGL1 correlates with an increased incidence of cellular overgrowth and malignant transformation (4-6). In mammalian epithelial cells, LLGL1 redistributes from the cytoplasm to regions of cell-cell contact, allowing the establishment and maintenance of a polarized morphology (7). LLGL1 also plays a role in the formation of epithelial junctions via its direct interactions with PAR6 and aPKC, the latter of which has been shown to phosphorylate LLGL1 at Ser663, thus restricting its localiza-

tion to the basolateral region of the cell (8). LLGL1 may also play an additional, unrealized role in cellular development and differentiation as indicated by the fact that *Drosophila lgl* has been implicated in controlling self-renewal and differentiation of progenitor cells (9). Recent studies in mice have suggested that a mammalian LLGL1 homolog that does not have tumor suppressor-like activity, LLGL2, is required for proper polarized invasion of trophoblasts and efficient branching morphogenesis during placental development (10).

**Specificity/Sensitivity:** LLGL1 (D2B5A) Rabbit mAb recognizes endogenous levels of total LLGL1 protein. This antibody does not cross-react with LLGL2.

**Source/Purification:** Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Leu757 of human LLGL1 protein.

Entrez-Gene ID #3996  
Swiss-Prot Acc. #Q15334

**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 antibody.

\*Species cross-reactivity is determined by western blot.

\*\*Anti-rabbit secondary antibodies must be used to detect this antibody.

## Recommended Antibody Dilutions:

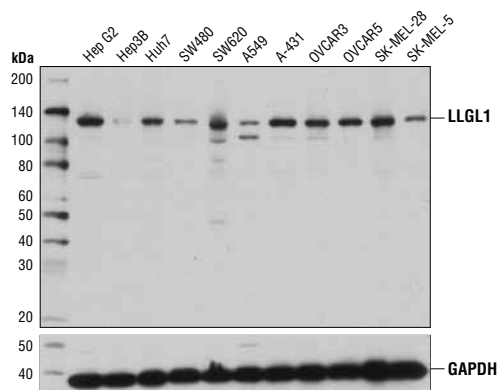
Western blotting	1:1000
Immunofluorescence (IF-IC)	1:50

For product specific protocols please see the web page for this product at [www.cellsignaling.com](http://www.cellsignaling.com).

Please visit [www.cellsignaling.com](http://www.cellsignaling.com) for a complete listing of recommended complementary products.

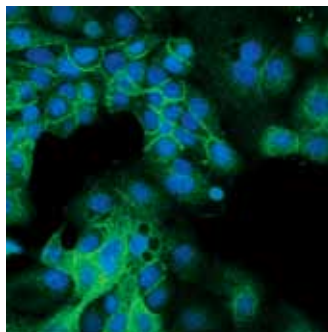
## Background References:

- (1) Agrawal, N. et al. (1995) *Dev Biol* 172, 218-29.
- (2) Massimi, P. et al. (2008) *Exp Cell Res* 314, 3306-17.
- (3) Strand, D. et al. (1995) *Oncogene* 11, 291-301.
- (4) Schimanski, C.C. et al. (2005) *Oncogene* 24, 3100-9.
- (5) Kuphal, S. et al. (2006) *Oncogene* 25, 103-10.
- (6) Lu, X. et al. (2009) *Clin Cancer Res* 15, 3287-96.
- (7) Misch, A. et al. (2002) *Mol Biol Cell* 13, 158-68.
- (8) Yamanaka, T. et al. (2003) *Curr Biol* 13, 734-43.
- (9) Mechler, B.M. et al. (1985) *EMBO J* 4, 1551-7.
- (10) Sripathy, S. et al. (2011) *Mol Cell Biol* 31, 2920-33.

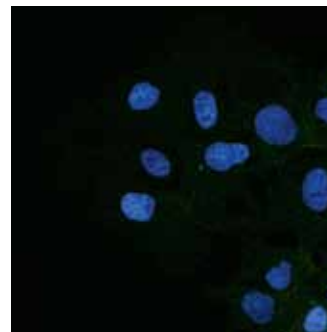


Western blot analysis of extracts from various cell lines using LLGL1 (D2B5A) Rabbit mAb (upper) or GAPDH (D16H11) XP® Rabbit mAb #5174 (lower).

## OVCAR3



## Hep3B



Confocal immunofluorescent analysis of OVCAR3 (positive; left) and Hep3B (weak; right) cells using LLGL1 (D2B5A) Rabbit mAb (green). Blue pseudocolor = DRAQ5® #4084 (fluorescent DNA dye).

**IMPORTANT:** For western blots, incubate membrane with diluted antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween-20 at 4°C with gentle shaking, overnight.

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**Applications Key:** W—Western IP—Immunoprecipitation IHC—Immunohistochemistry ChIP—Chromatin Immunoprecipitation IF—Immunofluorescence F—Flow cytometry E-P—ELISA-Peptide  
**Species Cross-Reactivity Key:** 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.