

O-GlcNAc (6D93): sc-71736

BACKGROUND

O-GlcNAc (O-linked N-acetylglucosamine) is a form of protein glycosylation found exclusively in the nucleus and cytoplasm of eukaryotic cells. Many proteins are modified at their serine and threonine hydroxyl groups by the attachment of O-GlcNAc. Proteins that regulate trafficking into and out of the nuclear pore are extensively O-GlcNAcylated. Phosphorylated O-GlcNAc proteins form reversible multimeric complexes with other proteins and these associations are often regulated by phosphorylation. O-GlcNAc proteins may play a key role in pathogenesis of tumors and various cancer cells. O-GlcNAc residues regulate the assembly of the preinitiation complex and are therefore important in transcriptional initiation. Cytoskeletal and membrane O-GlcNAc proteins maintain erythrocyte cell shape and regulate the degradation of proteins responsible for lesions in Alzheimer's disease.

REFERENCES

- Haltiwanger, R.S., et al. 1992. Glycosylation of nuclear and cytoplasmic proteins. Purification and characterization of a uridine diphospho-N-acetylglucosamine:polypeptide β -N-acetylglucosaminyltransferase. *J. Biol. Chem.* 267: 9005-9013.
- Kreppel, L.K., et al. 1997. Dynamic glycosylation of nuclear and cytosolic proteins. Cloning and characterization of a unique O-GlcNAc transferase with multiple tetratricopeptide repeats. *J. Biol. Chem.* 272: 9308-9315.
- Lubas, W.A., et al. 1997. O-linked GlcNAc transferase is a conserved nucleocytoplasmic protein containing tetratricopeptide repeats. *J. Biol. Chem.* 272: 9316-3624.
- Shafi, R., et al. 2000. The O-GlcNAc transferase gene resides on the X chromosome and is essential for embryonic stem cell viability and mouse ontogeny. *Proc. Natl. Acad. Sci. USA* 97: 5735-5739.
- Akimoto, Y., et al. 2003. Localization of the O-GlcNAc transferase and O-GlcNAc-modified proteins in rat cerebellar cortex. *Brain Res.* 966: 194-205.
- Chen, D., et al. 2005. Identification of secret agent as the O-GlcNAc transferase that participates in Plum pox virus infection. *J. Virol.* 79: 9381-9387.
- Leavy, T.M. and Bertozzi, C.R. 2007. A high-throughput assay for O-GlcNAc transferase detects primary sequence preferences in peptide substrates. *Bioorg. Med. Chem. Lett.* 17: 3851-3854.

SOURCE

O-GlcNAc (6D93) is a mouse monoclonal antibody raised against pore complex-lamina fraction purified from liver nuclear envelopes of rat origin.

PRODUCT

Each vial contains 200 μ g IgG₁ kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

STORAGE

Store at 4° C, ****DO NOT FREEZE****. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

APPLICATIONS

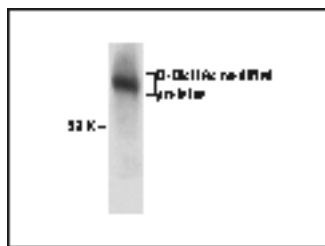
O-GlcNAc (6D93) is recommended for detection of O-GlcNAc in a broad range of species, including mammals, insects, worms, plants and filamentous fungi, by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)] and immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500).

Positive Controls: HeLa nuclear extract: sc-2120, A549 cell lysate: sc-2413 or mouse brain extract: sc-2253.

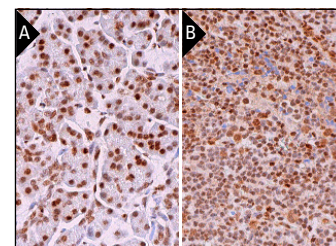
RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgG κ BP-HRP: sc-516102 or m-IgG κ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker[™] Molecular Weight Standards: sc-2035, UltraCruz[®] Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use m-IgG κ BP-FITC: sc-516140 or m-IgG κ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz[®] Mounting Medium: sc-24941 or UltraCruz[®] Hard-set Mounting Medium: sc-359850.

DATA



O-GlcNAc (6D93): sc-71736. Western blot analysis of O-GlcNAc modified proteins in HeLa cell lysate.




O-GlcNAc (6D93): sc-71736. Immunoperoxidase staining of formalin fixed, paraffin-embedded human pancreas tissue showing nuclear staining of exocrine glandular cells and Islets of Langerhans (A). Immunoperoxidase staining of formalin fixed, paraffin-embedded human tonsil tissue showing nuclear staining of cells in non-germinal center (B).

SELECT PRODUCT CITATIONS

- Sakaidani, Y., et al. 2010. O-GlcNAc modification of the extracellular domain of Notch receptors. *Methods Enzymol.* 480: 355-373.
- Borghgraef, P., et al. 2013. Increasing brain protein O-GlcNAc-ylation mitigates breathing defects and mortality of Tau.P301L mice. *PLoS ONE* 8: e84442.

RESEARCH USE

For research use only, not for use in diagnostic procedures.



See **O-GlcNAc (RL2): sc-59624** for O-GlcNAc antibody conjugates, including AC, HRP, FITC, PE, Alexa Fluor[®] 488 and Alexa Fluor[®] 647.