# abcam

## Product datasheet

# HRP Anti-O-Linked N-Acetylglucosamine antibody [RL2] ab201995

2 References 画像数 2

#### 製品の概要

製品名 HRP Anti-O-Linked N-Acetylglucosamine antibody [RL2]

製品の詳細 HRP Mouse monoclonal [RL2] to O-Linked N-Acetylglucosamine

由来種 Mouse 標識 HRP

**アプリケーション 適用あり:** IHC-P, WB

種交差性 交差種: Human

ポジティブ・コントロール IHC-P: Normal human colon tissue. WB: Jurkat treated with 0 uM PugNAc and Jurkat treated with

4 mM glucosamine and 50 uM PugNAc (3 hours).

特記事項

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

#### 製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C.

Avoid freeze / thaw cycle. Store In the Dark.

**バッファー** pH: 7.40

Preservative: 0.1% Proclin 300 Solution

Constituents: PBS, 30% Glycerol (glycerin, glycerine), 1% BSA

精製度 IgG fraction ポリ/モノ モノクローナル

**クローン名** RL2 **アイソタイプ** lgG1

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アプリケーション	Abreviews	特記事項
IHC-P		1/100. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
WB		1/1000.

#### ターゲット情報

#### 関連性

Many cellular proteins, including nuclear pore, oncogene, cytoskeletal, heat shock, viral and transcription regulatory proteins contain single O-linked N-acetylglucosamine (O-GlcNAc) residues attached to serine or threonine residues. It has been observed that O-GlcNAc glycosylated proteins tend to be under phosphorylated relative to unglycosylated proteins and that O-GlcNAc bearing proteins tend to be found in multimeric complexes. This has led to the suggestion that O-GlcNAc glycosylation may obscure phosphorylation sites and acts as a signaling mechanism or mediator of signaling.

#### 画像



Western blot - HRP Anti-O-Linked N-Acetylglucosamine antibody [RL2] (ab201995) **All lanes :** HRP Anti-O-Linked N-Acetylglucosamine antibody [RL2] (ab201995) at 1/1000 dilution

Lane 1: Jurkat treated with 0 uM PugNAc

**Lane 2**: Jurkat treated with 4 mM glucosamine and 50 uM PugNAc (3 hours)

Lysates/proteins at 20 µg per lane.

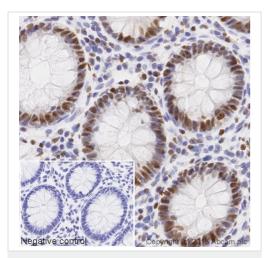
Performed under reducing conditions.

Observed band size: 100,65 kDa

Exposure time: 2 minutes

This blot was produced using a 4-12% Bis-tris gel under the MOPS buffer system. The gel was run at 200V for 50 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 3% milk before being incubated with ab201995 overnight at 4°C. Antibody binding

was visualised using ECL development solution ab133406.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - HRP Anti-O-Linked N-Acetylglucosamine antibody [RL2] (ab201995)

IHC image of O-linked N-Acetylglucosamine staining in a section of formalin-fixed paraffin-embedded normal human colon tissue\*, performed on a Leica BOND. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20mins. The section was then incubated with ab201995, 1/100 dilution, for 15 mins at room temperature. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX. The inset negative control image is taken from an identical assay without primary antibody.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.

\*Tissue obtained from the Human Research Tissue Bank, supported by the NIHR Cambridge Biomedical Research Centre.

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