abcam

Product datasheet

# HRP Anti-3-Nitrotyrosine antibody [7A12AF6] ab198491

1 References 1 Image

## Overview

| | |
|---|---|
| **Product name** | HRP Anti-3-Nitrotyrosine antibody [7A12AF6] |
| **Description** | HRP Mouse monoclonal [7A12AF6] to 3-Nitrotyrosine |
| **Host species** | Mouse |
| **Conjugation** | HRP |
| **Tested applications** | **Suitable for:** WB |
| **Species reactivity** | **Reacts with:** Cow, Human<br>**Predicted to work with:** Mouse, Rat |
| **Immunogen** | Chemical/ Small Molecule. This information is proprietary to Abcam and/or its suppliers. |
| **Positive control** | WB: Nitrated liver homogenate and nitrated bovine serum albumin. |
| **General notes** | 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.<br>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 |

## Properties

| | |
|---|---|
| **Form** | Liquid |
| **Storage instructions** | 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. |
| **Storage buffer** | pH: 7.40<br>Preservative: 0.1% Proclin 300 Solution<br>Constituents: 30% Glycerol (glycerin, glycerine), 1% BSA, PBS |
| **Purity** | Proprietary Purification |
| **Purification notes** | The antibody was produced in vitro using hybridomas grown in serum-free medium, and then purified by biochemical fractionation. Purity >95% by SDS-PAGE. |
| **Clonality** | Monoclonal |
| **Clone number** | 7A12AF6 |

| | |
|---|---|
| **Isotype** | IgG2b |
| **Light chain type** | kappa |

## Applications

**The Abpromise guarantee** Our **Abpromise guarantee** covers the use of ab198491 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

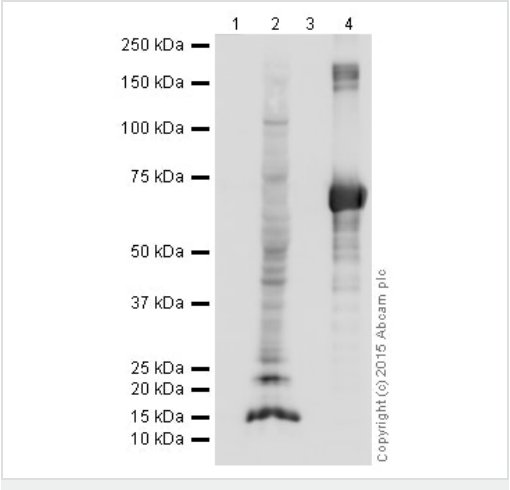
| Application | Abreviews | Notes |
|---|---|---|
| **WB** | | 1/5000. |

## Target

**Relevance**

Protein tyrosine nitration results in a post-translational modification that is increasingly receiving attention as an important component of nitric oxide signaling. While multiple nonenzymatic mechanisms are known to be capable of producing nitrated tyrosine residues, most tyrosine nitration events involve catalysis by metalloproteins such as myeloperoxidase, eosinophilperoxidase, myoglobin, the cytochrome P-450s, superoxide dismutase and prostacyclin synthase. Various studies have shown that protein tyrosinenitration is limited to specific proteins and that the process is selective. For example, exposure of human surfactant protein A, SP-A, to oxygen-nitrogen intermediates generated by activated alveolar macrophages resulted in specific nitration of SP-A at tyrosines 164 and 166, while addition of 1.2 mMCO 2 resulted in additional nitration at tyrosine 161. The presence of nitrotyrosine-containing proteins has shown high correlation to disease states such as atherosclerosis, Alzheimer's disease, Parkinson's disease and amyotrophic lateral sclerosis.

## Images




Western blot - HRP Anti-3-Nitrotyrosine antibody [7A12AF6] (ab198491)

**All lanes :** HRP Anti-3-Nitrotyrosine antibody [7A12AF6] (ab198491) at 1/5000 dilution

**Lane 1 :** Human liver tissue lysate - total protein (**ab29889**)
**Lane 2 :** Nitrated Human Liver Homogenate Standard (**ab131380**)
**Lane 3 :** Bovine Serum Albumin
**Lane 4 :** Bovine Serum Albumin (nitrated ) (**ab131379**)

Lysates/proteins at 10 µg per lane.

Developed using the ECL technique.

Performed under reducing conditions.

**Exposure time:** 1 second

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 2% Bovine Serum Albumin before being incubated with ab198491 overnight at 4°C. Antibody binding was visualised using ECL development solution **ab133406**.



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