Protein Tyrosine Nitration Control (Nitrotyrosine-BSA)

CATALOG NUMBER: STA-304 STORAGE: -80°C

QUANTITY AND CONCENTRATION: 100 µL of 0.1 mg/mL Nitrotyrosine-BSA in TBS, pH 7.4

SHELF LIFE: 2 years from receipt under proper storage conditions; avoid multiple freeze thaw cycles

Background

Nitric oxide (NO) was initially identified as a muscle relaxation factor, mediating its effects on smooth muscle by coordinating the heme moiety of guanylate cyclase, leading to enhanced production of cGMP. Besides smooth muscle relaxation, NO also influences a variety of biological systems including cell proliferation, apoptosis, neurotoxicity, Parkinson disease, extracellular matrix remodeling. Many of its effects are mediated through protein nitrosylation.

Peroxynitrite (ONOO⁻) is formed in the chemical reaction between nitric oxide and superoxide. ONOO⁻ nitrates free tyrosine and tyrosine residues in proteins; modifications that are used as markers of ONOO⁻ action under conditions of cellular damage and in numerous diseases.

This control is manufactured by nitration of bovine serum albumin *in vitro* using peroxynitrite as the NO donor.

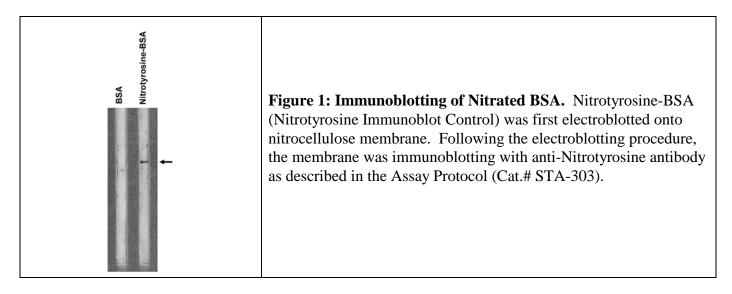
Methods

Dilute to 5 μ g/mL with reducing SDS sample buffer and heat for 5 min in boiling water. Load 20 μ L per lane for western blot analysis of protein tyrosine nitrosylation using Cell Biolabs' Nitrotyrosine Immunoblot Kit (STA-303) or anti-Nitrotyrosine antibody (STA-003 or STA-004).

Example of Results

The following figures demonstrate typical results. One should use the data below for reference only. This data should not be used to interpret actual results.





References

- 1. Halliwell, B., Zhao, K., and Whiteman, M. (1999) Free Radic. Res. 31, 651-669.
- 2. Greenacre, S. A. B., and Ischiropoulos, H. (2001) Free Radic. Res. 34, 541-581.

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