

Anti-Native-NGF antibody (STJ13100189)

STJ13100189

GENERAL INFORMATION

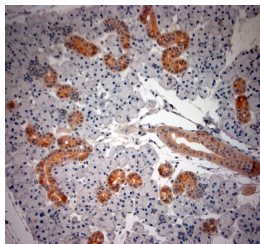
Product Type	Primary antibodies
Short Description	Nz White Rabbit polyclonal antibody anti-Native-NGF is suitable for use in Immunohistochemistry and Western Blot research applications.
Applications	IHC, WB
Host/Source	NZ White Rabbit
Reactivity	Mouse, Rat, Human

PRODUCT PROPERTIES

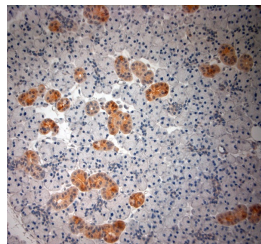
Clonality	Polyclonal
Clone ID	
Concentration	
Conjugation	Unconjugated
Purification	Whole serum
Dilution Range	A dilution of 1:2000 is recommended for WB and 1:1000 to 1:2000 for IHC-P; for inhibition of biological activities a dilution of up to 1:100 is recommended and for in vivo 1-20 µl per gram body weight. The optimal dilution should be determined by the
Formulation	Shipped as lyophilised. Reconstitute in 150 µl of sterile water. Centrifuge to remove any insoluble material.
Isotype	IgG
Storage Instruction	Maintain the lyophilised/reconstituted antibodies frozen at -20°C for long term storage and refrigerated at 2-8°C for a shorter term. When reconstituting, glycerol (1:1) may be added for an additional stability. Avoid freeze and thaw cycles.

TARGET INFORMATION

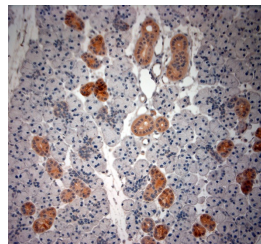
Gene ID	4803
Gene Symbol	NGF
Uniprot ID	NGF_HUMAN
Immunogen	Native beta-nerve growth factor (Ngfb, beta-NGF) purified from mouse submandibular gland
Immunogen Region	
Specificity	Specific for NGF
Immunogen Sequence	



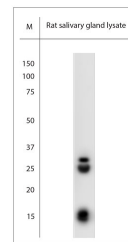
IHC-P on paraffin sections of rat salivary gland. The animal was perfused using Autoperfuser at a pressure of 130 mmHg with 300 ml of Davidsons modified fixative (4% formaldehyde also works nicely) before being processed for paraffin embedding. HIER: Tris-EDTA, pH 9 for 20 min using Thermo PT Module. Blocking: 0.2% LFDM in TBST filtered thru 0.2 µm. Detection was done using Novolink HRP polymer from Leica following manufacturer's instructions; DAB chromogen. Primary antibody dilution 1: 1000, incubated 30 min at RT using Autostainer. Sections were counterstained with Harris Hematoxylin.



IHC-P on paraffin sections of rat salivary gland. The animal was perfused using Autoperfuser at a pressure of 130 mmHg with 300 ml of Davidsons modified fixative (4% formaldehyde also works nicely) before being processed for paraffin embedding. HIER: Tris-EDTA, pH 9 for 20 min using Thermo PT Module. Blocking: 0.2% LFDM in TBST filtered thru 0.2 µm. Detection was done using Novolink HRP polymer from Leica following manufacturer's instructions; DAB chromogen. Primary antibody dilution 1: 1000, incubated 30 min at RT using Autostainer. Sections were counterstained with Harris Hematoxylin.



IHC-P on paraffin sections of rat salivary gland. The animal was perfused using Autoperfuser at a pressure of 130 mmHg with 300 ml of Davidsons modified fixative (4% formaldehyde also works nicely) before being processed for paraffin embedding. HIER: Tris-EDTA, pH 9 for 20 min using Thermo PT Module. Blocking: 0.2% LFDM in TBST filtered thru 0.2 µm. Detection was done using Novolink HRP polymer from Leica following manufacturer's instructions; DAB chromogen. Primary antibody dilution 1: 1000, incubated 30 min at RT using Autostainer. Sections were counterstained with Harris Hematoxylin.



WB on rat salivary gland lysate. Blocking: 1% LFDM for 30 min at RT; primary antibody dilution 1: 1000 incubated at 4C overnight.

This product is suitable for in-vitro studies under the RESEARCH USE ONLY [RUO] licence. This product must not be used as for diagnostic or other medical purposes.

St John's Laboratory Ltd, Knowledge Dock Business Centre, University Way, London, E16 2RD | Tel: 0208 223 3081