

Anti-Pro--BDNF antibody (STJ13100047)

GENERAL INFORMATION

Product Type Primary antibodies

Short Nz White Rabbit polyclonal antibody anti-Pro--BDNF is suitable for use in Immunohistochemistry and Western Blot research

Description 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 A dilution of 1:1000 is recommended. The optimal dilution should be determined by the end user.

Range

Formulation Shipped as lyophilised. Reconstitute in 100 ul of sterile water. Centrifuge to remove any insoluble material.

Isotype IgG

Storage Maintain the lyophilised/reconstituted antibodies frozen at-20C for long term storage and refrigerated at 2-8C for a shorter term. When

Instruction reconstituting, glycerol (1:1) may be added for an additional stability. Avoid freeze and thaw cycles.

TARGET INFORMATION

Gene ID 627 Gene Symbol BDNF

Immunogen A synthetic peptide from mouse pro BDNF conjugated to blue carrier protein was used as the antigen.

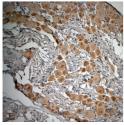
Uniprot ID BDNF_HUMAN

Immunogen

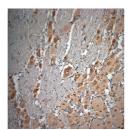
Region

Immunogen Sequence

Specificity Specific for pro BDNF.



on paraffin sections of rat DRG. The seed using Autoperfuser at a pressura with 300 ml 4% FA and further ing win sou in 470 FA and further post fixed reight before being processed for paraffin edding. HIER: Tris-EDTA, pH 9 for 20 min using mo PT Module. Blocking: 0.296 LFDM in TBST ed thru 0.2 um. Detection was done using Novolink polymer from Leica following manufacturer's uctions; DAB chromosop.



IHC-P on paraffin sections of rat DRG. The perfused using Autoperfuser at a press mmHg with 300 ml 4% FA and further mmHg wtn 30U ml 4% FA and turmen overnight before being processed fo embedding. HIER: Tris-EDTA, pH 9 for 20 Thermo PT Module. Blocking: 0.2% LFDN filtered thru 0.2 um. Detection was done usin HRP polymer from Leica following mar



paraffin sections of rat DRG. The animal was using Autoperfuser at a pressure of 130 titl 300 ml 49 FA and turther post fixed before being processed for paraffin g. HER. Tiss-EUR, ph 9 for 20 min using u. 0.2 um. Detection was done using Novolink mer from Leica following market clubter of the control of t

