

## Rabbit antibody to NPY (20-60)

| Code<br>ID Tag<br>Unit size | OSN00111W<br>Rb3748-080620-WS<br>100 ul   |
|-----------------------------|---|
| Immunogen                   | A synthetic peptide from aa region 20-60 of mouse NPY conjugated to blue carrier protein was used as the antigen. The antigen is homologous in many other species including human, rat, zebrafish and xenopus.                                  |
| Conjugate                   | Unconjugated antibody   |
| Also known                  | Neuropeptide Y, Neuropeptide tyrosine, C-flanking peptide of NPY, CPON  |
| Host                        | NZ white rabbit   |
| Purity                      | Whole serum   |
| Clonality                   | Polyclonal  |
| Isotype                     | Polyclonal, whole serum   |
| Applications                | IHC, WB. A dilution of 1: 250 to 1: 1000 is recommended. The optimal dilution should be determined by the end user. Not yet tested in other applications.   |
| Specificity                 | Specific for NPY  |
| Spcs X-react.               | Human, mouse, rat, marmoset. Other species not yet tested.  |
| Format                      | Lyophilised   |
| Reconstitution              | Reconstitute in 100 ul of sterile water. Centrifuge to remove any insoluble material.   |
| Storage                     | Maintain the lyophilised/reconstituted antibodies frozen at -20C for long term storage and refrigerated at 2-8C for a shorter term. When reconstituting, glycerol (1:1) may be added for an additional stability. Avoid freeze and thaw cycles. |
| Expiry Date                 | 12 months after reconstitution  |
| Shipping                    | This item will be shipped to you at ambient temperature in a lyophilised form.  |
| Limitation                  | For research use only   |



IHC-P on paraffin sections of mouse brain.

The animal was perfused using Autoperfuser at a pressure of 130 mmHg with 300 ml 4% FA 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 um.

Detection was done using Novolink HRP polymer from Leica following manufacturers instructions; DAB chromogen: Candela DAB chromogen from Osenses. Primary antibody: dilution 1: 500, incubated 30 min at RT using Autostainer.

Sections were counterstained with Harris Hematoxylin.