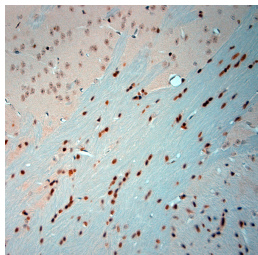


## Rabbit antibody to Olig2

|                       |   |
|-----------------------|---|
| <b>Code</b>           | OSO00005W   |
| <b>ID Tag</b>         | Rb3496-171119-WS  |
| <b>Unit size</b>      | 100 ul  |
| <b>Immunogen</b>      | A synthetic peptide from human Olig2 conjugated to blue carrier protein was used as the antigen. The peptide is homologous in rat and mouse.  |
| <b>Conjugate</b>      | Unconjugated antibody   |
| <b>Also known</b>     | Basic helix loop helix protein class B 1, BHLHB1, olig2, OLIGO2, Oligodendrocyte specific bHLH transcription factor 2, Oligodendrocyte transcription factor 2, PRKCBP2, Protein kinase C binding protein 2, Protein kinase C binding protein RACK17, RACK17 |
| <b>Host</b>           | NZ white rabbit   |
| <b>Purity</b>         | Whole serum   |
| <b>Clonality</b>      | Polyclonal  |
| <b>Isotype</b>        | Polyclonal, whole serum   |
| <b>Applications</b>   | IHC, WB, Flow Cyt. A dilution of 1 : 250 is recommended. The optimal dilution should be determined by the end user. Not yet tested in other applications.   |
| <b>Specificity</b>    | Specific for Olig2.   |
| <b>Spes X-react.</b>  | Human, rat, mouse, marmoset. Other species not yet tested.  |
| <b>Format</b>         | Lyophilised   |
| <b>Reconstitution</b> | Reconstitute in 100 Åµl of sterile water. Centrifuge to remove any insoluble material.  |
| <b>Storage</b>        | 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.             |
| <b>Expiry Date</b>    | 12 months after reconstitution  |
| <b>Shipping</b>       | This item will be shipped to you at ambient temperature in a lyophilised form.  |
| <b>Limitation</b>     | For research use only   |



IHC-P on paraffin sections of rat 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 µm.

Detection was done using Novolink HRP polymer from Leica following manufacturers instructions; DAB chromogen: Candela DAB chromogen from Osenses.

Primary antibody: dilution 1: 250, incubated 30 min at RT using Autostainer.

Sections were counterstained with Harris Hematoxylin.