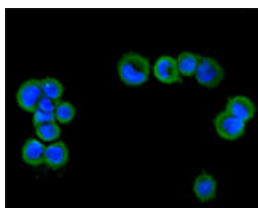


## Sheep antibody to TRPM4

<b>Code</b>	OST00033W
<b>ID Tag</b>	Sh238-1-280310-WS
<b>Unit size</b>	100 µl
<b>Immunogen</b>	A synthetic peptide from human TRPM4 conjugated to blue carrier protein was used as the antigen.
<b>Conjugate</b>	Unconjugated antibody
<b>Also known</b>	Transient receptor potential cation channel subfamily M member 4, long transient receptor potential channel 4, hTRPM4, melastatin-4, calcium-activated non-selective cation channel 1, TRPM4, TRPM4B, FLJ20041
<b>Host</b>	Sheep
<b>Purity</b>	Whole serum
<b>Clonality</b>	Polyclonal
<b>Isotype</b>	Polyclonal, whole serum
<b>Applications</b>	IHC, WB. A dilution of 1 : 300 to 1 : 2000 is recommended. The optimal dilution should be determined by the end user. Not yet tested in other applications.
<b>Specificity</b>	Specific for TRPM4.
<b>Spcs X-react.</b>	Human. Other species not yet tested.
<b>Format</b>	Lyophilised
<b>Reconstitution</b>	Reconstitute in 100 µl of sterile water. Centrifuge to remove any insoluble material.
<b>Storage</b>	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.
<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



A cell cytospin of human lung cancer cell line A549 was fixed with cold acetone for 90 seconds and air-dried. Cells were incubated with the blocking buffer (PBS containing 5% FCS) for 30 minutes at room temperature. Cells were then washed once in PBS and incubated with primary antibody, diluted 1:100 in the blocking buffer, for 30 minutes. Slides were washed 3X in PBS and incubated with Goat anti-sheep conjugated to Alexa-488, diluted 1:200 in blocking buffer, for 30 minutes at room temperature in the dark. Slides were washed as above and mounting media (10% Glycerol in PBS) containing Hoechst 33258 1 µg/ml was used for nuclear counterstaining. Fluorescent cell staining were analysed using a Olympus microscope and the analysis LS Research software.