

Product datasheet for AM26775LE-N

Product datasifeet for Awizo//SLE-IV

Mouse IgG2a Isotype Control

Product data:

Product Type: Isotype Controls
Applications: FC, ICC/IF, IP, WB

Host: Mouse Isotype: IgG2a

Clonality: Monoclonal Immunogen: Mineral oils

Specificity: This antibody reacts with an unknown epitope. It does not react with a variety of resting,

activated, live, and fixed mouse, rat and human tissues. The reagent is intended as isotype control for flow cytometry analysis to establish the amount of non-specific antibody binding. For your particular experiment, use the same concentration of this isotype control antibody as the recommended working concentration of the antigen-specific antibody. Also, when working with prediluted antibodies, dilute the isotype control to the same concentration as is the concentration of the antigen-specific antibody in the prediluted antibody solution you are using. If under particular experimental conditions the background signal of the isotype

control is too high (usually when working concentrations of used antibodies are above 10 μ g per ml of incubation mixture), change the conditions of your experiment to reduce the

background.

Buffer: Azide free phosphate buffered saline (PBS), approx. pH 7.4; 0.2 µm filter sterilized

Concentration: 1 mg/ml

Purification: Protein-A affinity chromatography

Conjugation: Unconjugated

Storage: Store undiluted at 2-8°C.

DO NOT FREEZE!

Shelf life: one year from despatch.



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Background:

The specificity of staining by monoclonal antibodies to target antigens should be verified by establishing the amount of non-specific antibody binding. Especially at higher concentration (more than 15 μ g/ml) the antibody staining usually has consignable background. To this end a non-reactive immunoglobulin of the same isotype is included as a negative control for each specific monoclonal antibody used in a particular immunoassay. The monoclonal antibody MOPC-173, generated against an undefined antigen, does not react specifically with mouse, rat and human samples, and hence all the background that could be observed when working with this antibody would be a result of general nonspecific interactions between an mouse $\lg G2a$ molecule and the respective sample under the particular conditions. This shall help the customer to set up the experimental conditions so that the nonspecific binding of any antibody is abolished.