

Product datasheet

Anti-C2 antibody [5C3] ab17451

2 References

Overview

| | |
|----------------------------|---|
| Product name | Anti-C2 antibody [5C3] |
| Description | Mouse monoclonal [5C3] to C2 |
| Host species | Mouse |
| Specificity | This antibody reacts with C2 and a subfraction of C2 believed to be C2b |
| Tested applications | Suitable for: ELISA, WB |
| Species reactivity | Reacts with: Human |
| Immunogen | Full length native complement component C2 protein isolated from human plasma. |
| Epitope | Epitope specificity differs from that of ab17452 as determined by inhibition ELISA. |

Properties

| | |
|-----------------------------|---|
| Form | Liquid |
| Storage instructions | Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles. |
| Storage buffer | pH: 7.40 Constituents: 0.0268% PBS, 2.9% Sodium chloride |
| Purity | Protein A purified |
| Clonality | Monoclonal |
| Clone number | 5C3 |
| Myeloma | x63-Ag8.653 |
| Isotype | IgG2a |
| Light chain type | kappa |

Applications

Our [Abpromise guarantee](#) covers the use of **ab17451** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

| Application | Abreviews | Notes |
|-------------|-----------|----------|
| ELISA | | 1/12000. |

| Application | Abreviews | Notes |
|-------------|-----------|--|
| WB | | Use at an assay dependent dilution. Predicted molecular weight: 102 kDa. |

Target

| | |
|-------------------------------|--|
| Function | Component C2 which is part of the classical pathway of the complement system is cleaved by activated factor C1 into two fragments: C2b and C2a. C2a, a serine protease, then combines with complement factor 4b to generate the C3 or C5 convertase. |
| Involvement in disease | Defects in C2 are the cause of complement component 2 deficiency (C2D) [MIM:217000]. A deficiency of the complement classical pathway associated with the development of autoimmune disorders, mainly systemic lupus erythematosus. Skin and joint manifestations are common and renal disease is relatively rare. Patients with complement component 2 deficiency are also reported to have recurrent or invasive infections. |
| Sequence similarities | Belongs to the peptidase S1 family. Contains 1 peptidase S1 domain. Contains 3 Sushi (CCP/SCR) domains. Contains 1 VWFA domain. |
| Cellular localization | Secreted. |

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