

Product Data Sheet

Anti-CD35 Antibody

| Catalog # | Source | Reactivity | Applications | | | |
|--|------------|--|---|--|--|--|
| CPA9727 | Mouse | н | IH | | | |
| Description | Mou | Mouse monoclonal antibody to CD35 | | | | |
| Immunogen | KLH- | conjugated synthetic | peptide encompassing a sequence within human CD35. | | | |
| | The e | exact sequence is pro | prietary. | | | |
| Purification The antibody was purified by immunogen affinity chromatography. | | | | | | |
| Specificity Recognizes endogenous levels of CD35 protein. | | | | | | |
| Clonality | Mon | oclonal | | | | |
| Conjugation | | | | | | |
| Form | Mou | Mouse IgG1. Liquid in PBS containing 50% glycerol, 0.2% BSA and 0.01% sodium | | | | |
| | azide | 2. | | | | |
| Dilution | IH (1, | /100 - 1/300) | | | | |
| Gene Symbol | CR1 | | | | | |
| Alternative Na | ames C3BF | R; Complement recep | tor type 1; C3b/C4b receptor; CD35 | | | |
| Entrez Gene | 1378 | (Human) | | | | |
| SwissProt | P179 | 27 (Human) | | | | |
| Storage/Stabi | lity Shipp | oed at 4°C. Upon deliv | very aliquot and store at -20°C for one year. Avoid | | | |
| | freez | e/thaw cycles. | | | | |

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

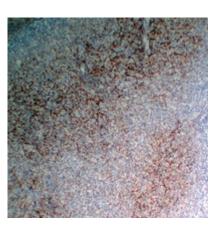
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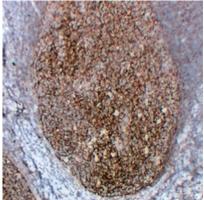
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Immunohistochemical analysis of CD35 staining in human follicular lymphoma formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubat



Immunohistochemical analysis of CD35 staining in human tonsil formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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