

## Product datasheet

# Anti-Caspase-8 p10 antibody [C502S] ab2553

## 2 References

### Overview

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|----------------------------|--|
| <b>Product name</b>        | Anti-Caspase-8 p10 antibody [C502S]  |
| <b>Description</b>         | Mouse monoclonal [C502S] to Caspase-8 p10  |
| <b>Host species</b>        | Mouse  |
| <b>Specificity</b>         | This antibody recognizes the 10 kDa small subunit of Caspase 8. Does not detect full-length caspase 8. |
| <b>Tested applications</b> | <b>Suitable for:</b> IP, WB  |
| <b>Species reactivity</b>  | <b>Reacts with:</b> Human  |
| <b>Immunogen</b>           | Synthetic peptide corresponding to Human Caspase-8 p10 (N terminal).                                   |

### Properties

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|-----------------------------|---|
| <b>Form</b>                 | Liquid  |
| <b>Storage instructions</b> | Shipped at 4°C. Upon delivery aliquot and store at -20°C or -80°C. Avoid repeated freeze / thaw cycles. |
| <b>Storage buffer</b>       | Preservative: 0.02% Thimerosal (merthiolate)<br>Constituents: 50% Glycerol, 1% BSA, PBS                 |
| <b>Purity</b>               | Immunogen affinity purified   |
| <b>Clonality</b>            | Monoclonal  |
| <b>Clone number</b>         | C502S   |
| <b>Myeloma</b>              | unknown   |
| <b>Isotype</b>              | IgG1  |
| <b>Light chain type</b>     | unknown   |

### Applications

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Our [Abpromise guarantee](#) covers the use of **ab2553** 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                            |
|-------------|-----------|----------------------------------|
| IP          |           | Use a concentration of 10 µg/ml. |
| WB          |           | Use a concentration of 1 µg/ml.  |

## Target

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|---|--|
| <b>Function</b>                         | <p>Most upstream protease of the activation cascade of caspases responsible for the TNFRSF6/FAS mediated and TNFRSF1A induced cell death. Binding to the adapter molecule FADD recruits it to either receptor. The resulting aggregate called death-inducing signaling complex (DISC) performs CASP8 proteolytic activation. The active dimeric enzyme is then liberated from the DISC and free to activate downstream apoptotic proteases. Proteolytic fragments of the N-terminal propeptide (termed CAP3, CAP5 and CAP6) are likely retained in the DISC. Cleaves and activates CASP3, CASP4, CASP6, CASP7, CASP9 and CASP10. May participate in the GZMB apoptotic pathways. Cleaves ADPRT. Hydrolyzes the small-molecule substrate, Ac-Asp-Glu-Val-Asp-AMC. Likely target for the cowpox virus CRMA death inhibitory protein. Isoform 5, isoform 6, isoform 7 and isoform 8 lack the catalytic site and may interfere with the pro-apoptotic activity of the complex.</p> |
| <b>Tissue specificity</b>               | <p>Isoform 1, isoform 5 and isoform 7 are expressed in a wide variety of tissues. Highest expression in peripheral blood leukocytes, spleen, thymus and liver. Barely detectable in brain, testis and skeletal muscle.</p>   |
| <b>Involvement in disease</b>           | <p>Caspase-8 deficiency</p>  |
| <b>Sequence similarities</b>            | <p>Belongs to the peptidase C14A family.<br/>Contains 2 DED (death effector) domains.</p>  |
| <b>Domain</b>                           | <p>Isoform 9 contains a N-terminal extension that is required for interaction with the BCAP31 complex.</p>   |
| <b>Post-translational modifications</b> | <p>Generation of the subunits requires association with the death-inducing signaling complex (DISC), whereas additional processing is likely due to the autocatalytic activity of the activated protease. GZMB and CASP10 can be involved in these processing events.<br/>Phosphorylation on Ser-387 during mitosis by CDK1 inhibits activation by proteolysis and prevents apoptosis. This phosphorylation occurs in cancer cell lines, as well as in primary breast tissues and lymphocytes.</p>   |
| <b>Cellular localization</b>            | <p>Cytoplasm.</p>  |

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