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Datasheet for ABIN4998199 anti-C1QL2 antibody (Alexa Fluor 680)



| Overview | |
|-------------------|---|
| Quantity: | 100 μL |
| Target: | C1QL2 |
| Reactivity: | Human, Mouse, Rat |
| Host: | Rabbit |
| Clonality: | Polyclonal |
| Conjugate: | This C1QL2 antibody is conjugated to Alexa Fluor 680 |
| Application: | Western Blotting (WB), Immunofluorescence (Paraffin-embedded Sections) (IF (p)) |
| Product Details | |
| Immunogen: | KLH conjugated synthetic peptide derived from human C1QL2/C1QTNF10 |
| Isotype: | lgG |
| Cross-Reactivity: | Human, Mouse, Rat |
| Purification: | Purified by Protein A. |
| Target Details | |
| Target: | C1QL2 |
| Alternative Name: | C1QTNF10 (C1QL2 Products) |
| Background: | Synonyms: C1q and tumor necrosis factor related protein 10, C1q domain containing protein, C1QL2, C1QL2_HUMAN, C1QTNF10, Complement C1q-like protein 2, Complement component 1, q subcomponent-like 2, CTRP10, gliacolin like. Background: C1qL2, also known as CTRP10 or C1QTNF10, is a 287 amino acid secreted protein |
| | |

Order at www.antibodies-online.com | www.antikoerper-online.de | www.anticorps-enligne.fr | www.antibodies-online.cn International: +49 (0)241 95 163 153 | USA & Canada: +1 877 302 8632 | support@antibodies-online.com Page 1/2 | Product datasheet for ABIN4998199 | 03/06/2024 | Copyright antibodies-online. All rights reserved.

| | that contains one C1q domain and one collagen-like domain. C1qL2 belongs to a large family of |
|---------------------|---|
| | multimeric proteins with a signature globular domain homologous to C1QA. These proteins |
| | also share structural homology with TNF family members. The gene that encodes C1qL2 |
| | consists of approximately 2,653 bases and maps to human chromosome 2q14.2. Consisting of |
| | 237 million bases, chromosome 2 encodes over 1,400 genes and makes up approximately 8 $\%$ |
| | of the human genome. A number of genetic diseases are linked to genes on chromosome 2. |
| | Harlequin icthyosis, a rare and morbid skin deformity, is associated with mutations in the |
| | ABCA12 gene. The lipid metabolic disorder sitosterolemia is associated with ABCG5 and |
| | ABCG8. An extremely rare recessive genetic disorder, Alstr syndrome, is due to mutations in the |
| | ALMS1 gene. |
| Gene ID: | 165257 |
| Application Details | |
| Application Notes: | IF(IHC-P) 1:50-200 |
| Restrictions: | For Research Use only |
| Handling | |
| Format: | Liquid |
| Concentration: | 1 μg/μL |
| Buffer: | Aqueous buffered solution containing 0.01M TBS (pH 7.4) with 1 % BSA, 0.03 % Proclin300 and |
| | 50 % Glycerol. |
| Preservative: | ProClin |
| Precaution of Use: | This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE, which should be |
| | handled by trained staff only. |
| Storage: | -20 °C |
| Storage Comment: | Store at -20°C. Aliquot into multiple vials to avoid repeated freeze-thaw cycles. |
| Expiry Date: | 12 months |
| | |