

**DESCRIPTION**

**Source** *E. coli*-derived human C1qL4 protein  
Gly105-Asp238, with an N-terminal Met and 6-His tag  
Accession # Q86Z23

**N-terminal Sequence Analysis** Met

**Predicted Molecular Mass** 16 kDa

**SPECIFICATIONS**

**SDS-PAGE** 13 kDa, reducing conditions

**Activity** Measured by its binding ability in a functional ELISA.  
When Recombinant Human C1qL4 is immobilized at 2 µg/mL, 100 µL/well, the concentration of Recombinant Human BAI2 Fc Chimera (Catalog # 9338-BA) that produces 50% of the optimal binding response is 2-10 µg/mL.

**Endotoxin Level** <0.10 EU per 1 µg of the protein by the LAL method.

**Purity** >95%, by SDS-PAGE visualized with Silver Staining and quantitative densitometry by Coomassie® Blue Staining.

**Formulation** Lyophilized from a 0.2 µm filtered solution in Tris, NaCl and TCEP with Trehalose. See Certificate of Analysis for details.

**PREPARATION AND STORAGE**

**Reconstitution** Reconstitute at 500 µg/mL in water.

**Shipping** The product is shipped with polar packs. Upon receipt, store it immediately at the temperature recommended below.

**Stability & Storage** Use a manual defrost freezer and avoid repeated freeze-thaw cycles.

- 12 months from date of receipt, -20 to -70 °C as supplied.
- 1 month, 2 to 8 °C under sterile conditions after reconstitution.
- 3 months, -20 to -70 °C under sterile conditions after reconstitution.

**BACKGROUND**

C1qTNF11 (CTRP11), also known as C1qL4, is an approximately 29 kDa member of the C1qTNF family of secreted proteins (1, 2). Mature human C1qTNF11/C1qL4 contains two distinct domains: a collagen-like region and one C1q-like domain, and can form disulfide-linked heteromers with C1qTNF14/C1qL1 (3, 4). The C1qTNF11/C1qL4 gene is conserved in human, chimpanzee, dog, mouse, rat, zebrafish, and frog. Within the C1q-like domain, human C1qTNF11/C1qL4 shares 100% amino acid (aa) sequence identity with that of mouse, monkey and rat. C1qTNF11/C1qL4 is expressed predominantly in testis and adipocytes, and also at a lower level in skeletal muscle and kidney (4), hippocampus, and cerebral cortex (5). Similar to C1qTNF13/C1qL3, C1qTNF10/C1qL2, and C1qTNF14/C1qL1, C1qTNF11/C1qL4 binds to BAI3 in the cerebral cortex and on cerebellar Purkinje cells (5-8). C1qTNF11/C1qL4 binding to BAI3 induces the formation and maintenance of excitatory synapses between climbing fibers and parallel fibers with Purkinje cells (5, 7). Our in house assay shows that C1qTNF11/C1qL4 also binds to BAI-2. In addition, C1qTNF11/C1qL4 has been reported to promote angiogenesis via activation of the ERK pathway in HUVEC cells mediated by BAI3 (9). Conversely, in adipocytes, C1qTNF11/C1qL4 treatment decreased ERK1/2 signaling and inhibited adipogenesis (4).

**References:**

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