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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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Datasheet for C300-0050

Guinea Pig Complement (Fresh Frozen)**Overview**

Description:	Guinea Pig Complement (Fresh Frozen) - C300-0050
Item No.:	C300-0050
Size:	50 mL
Applications:	Cellular Assay, ELISA, FC, Purification
Origin:	Guinea Pig

Product Details

Background:	Special processing techniques are used to yield products with high complement activity and low background cytotoxicity. Guinea Pig Complement is suitable for CFT and SRH.
Synonyms:	Complement system, tissue macrophages, blood monocytes, protease C3-convertase, mannose-binding lectin pathway, C3, C3a, C3b, C5a, C5b, C6, C7, C8, and polymeric C9, cascade cleavage and activation events, recruit inflammatory cells, anaphylatoxin
Species of Origin:	Guinea Pig

Target Details

Relevant Links:	<ul style="list-style-type: none">C300 SDS
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Application Details

Suggested Applications:	Cellular Assay, ELISA, FC, Purification (Based on references)
Application Note:	pH: normal Immunoelectrophoresis: normal Hemoglobin: normal IgG Concentration: normal

Assay Dilutions: All assays should be optimized by the user. Recommended dilutions (if any) may be listed below.

Tissue Data

Tissue Type: Complement

Sex: Mixed

Strain: Guinea Pig - Mixed

Formulation

Physical State: Fresh Frozen

Concentration: 80 mg/mL by Refractometry

Buffer: None

Sterility: Non-sterile

Preservative: None

Stabilizer: None

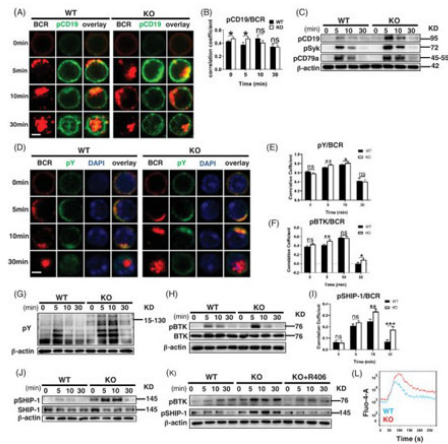
Shipping & Handling

Shipping Condition: Dry Ice

Storage Condition: Store Guinea Pig Complement at -70° C prior to opening. Aliquot contents and freeze at -70° C or below. Use aseptic technique to maintain sterility when opening product. Avoid cycles of freezing and thawing. Centrifuge product if not completely clear after standing at room temperature. COMPLEMENT IS A TEMPERATURE SENSITIVE PRODUCT. IMPROPER STORAGE WILL INACTIVATE COMPLEMENT ACTIVITY.

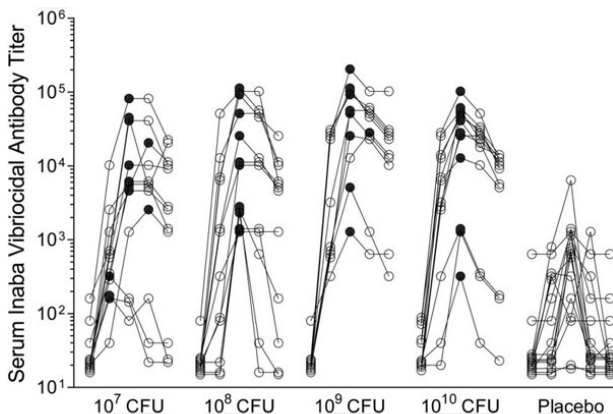
Expiration: Expiration date is one (1) year from date of receipt.

Images



Immunofluorescence Microscopy

Ccr2-KO mice exhibit enhanced BCR proximal signalling. B cells were purified from splenic mononuclear cells by incubation of anti-Thy-1 and guinea pig complement (p/n C300-0500) for 30 min. Purified splenic B cells were incubated with AF546-F(ab')₂-anti-mouse-Ig (M + G) at 4°C for 30 min and activated at 37°C for 5, 10 and 30 min, confocal microscopy (CFm) was performed. Cells were incubated with biotin-conjugated F(ab')₂-anti-mouse-Ig (M + G) and streptavidin, then activated at 37°C for 5, 10 and 30 min, western blotting was performed. (A) Representative CFm images of phosphorylated CD19 (pCD19) and BCR (60× objective, scale bar = 2.5 μm). (B) Colocalization between pCD19 and BCR. (C) Western blotting of pCD19, pSyk, pCD79a expression in B cells. (D) Representative CFm images of pY and BCR (60× objective, scale bar = 2.5 μm). (E) Colocalization between pY and BCR. (F) Colocalization between pBTK and BCR. (G) Western blotting of pY expression in B cells. (H) Western blotting of pBTK and BTK expression in B cells. (I) Colocalisation between pSHIP-1 and BCR. (J) Western blotting of pSHIP-1 and SHIP-1 expression in B cells. (K) Western blotting of pBTK and pSHIP-1 in WT B cells, Ccr2-KO B cells and Ccr2-KO B cells treated with 5μM R406. (L) Representative image of intracellular Ca²⁺ flux kinetics in WT and Ccr2-KO B cells following stimulation with 10 μg/ml biotin-conjugated F(ab')₂ anti-mouse Ig (M + G). All images were representative images from three independent experiments. The number of cells analyzed for each parameter in CFm assay was 30–50. Error bars were shown as mean (± SD). *p < .05, **p < .01, ***p < .001, ns: no statistical significance. Fig 2. PMID: 35875970.



ELISA

Individual serum Inaba vibriocidal responses for each of the four dosages (10⁷, 10⁸, 10⁹, or 10¹⁰ CFU) or placebo are indicated. Within each dosage group, the five circles denote (from left to right) the following five time points: baseline and 7, 10, 14, and 28 days postvaccination. A closed circle indicates the peak response for an individual. The vibriocidal antibody assay compares the amount of *V. cholerae* growth achieved in a 96-well plate when mixed with guinea pig complement (p/n C300-0050) of a standard activity and serial dilutions of the heat-inactivated human serum samples, all assayed in duplicate. FIG 1. PMID: 25410205.

References

- Gu H et al. Ultra-high static magnetic fields cause immunosuppression through disrupting B-cell peripheral differentiation and negatively regulating BCR signaling. *MedComm (2020)*. (2023)
- Guan F et al. GSDMA3 deficiency reprograms cellular metabolism and modulates BCR signaling in murine B cells. *iScience*. (2023)
- Zhu, Y et al. Involvement of MST1/mTORC1/STAT1 activity in the regulation of B-cell receptor signalling by chemokine receptor 2. *Clinical and Translational Medicine* (2022)
- Luo L et al. Abelson tyrosine kinase controls BCR signalling and B-cell differentiation by promoting B-cell metabolism. *Immunology*. (2022)
- Yang L et al. CCL2 regulation of MST1-mTOR-STAT1 signaling axis controls BCR signaling and B-cell differentiation. *Cell Death Differ*. (2021)
- Li N, Jiang P, Chen A, et al. CX3CR1 positively regulates BCR signaling coupled with cell metabolism via negatively controlling actin remodeling. *Cell Mol Life Sci*. (2020)
- Jing Y. et. al. STING couples with PI3K to regulate actin reorganization during BCR activation. *IMMUNOLOGY* (2020)
- Yang JS et al. IgM specific to lipopolysaccharide of *Vibrio cholerae* is a surrogate antibody isotype responsible for serum vibriocidal activity. *PLoS One*. (2019)
- Hosseini SM et al. Transcriptome profiling of bovine inner cell mass and trophectoderm derived from in vivo generated blastocysts. *BMC Dev Biol*. (2015)
- Chen WH et al. Safety and immunogenicity of escalating dosages of a single oral administration of peru-15 pCTB, a candidate live, attenuated vaccine against enterotoxigenic *Escherichia coli* and *Vibrio cholerae*. *Clin Vaccine Immunol*. (2014)
- Zhang J et al. Development and characterization of an infectious cDNA clone of the modified live virus vaccine strain of equine arteritis virus. *Clin Vaccine Immunol*. (2012)
- Ozawa M et al. Importance of culture conditions during the morula-to-blastocyst period on capacity of inner cell-mass cells of bovine blastocysts for establishment of self-renewing pluripotent cells. *Theriogenology*. (2012)
- LaFleur RL et al. One-year duration of immunity induced by vaccination with a canine Lyme disease bacterin. *Clin Vaccine Immunol*. (2010)
- Yang JS et al. A duplex vibriocidal assay to simultaneously measure bactericidal antibody titers against *Vibrio cholerae* O1 Inaba and Ogawa serotypes. *J Microbiol Methods*. (2009)
- Yang JS et al. A semi-automated vibriocidal assay for improved measurement of cholera vaccine-induced immune responses. *J Microbiol Methods*. (2007)
- Whitaker-Menezes D et al. An epithelial target site in experimental graft-versus-host disease and cytokine-mediated cytotoxicity is defined by cytokeratin 15 expression. *Biol Blood Marrow Transplant*. (2003)
- Lynch JM et al. Increased protection against pneumococcal disease by mucosal administration of conjugate vaccine plus interleukin-12. *Infect Immun*. (2003)

- Jones SC et al. Post-hematopoietic cell transplantation control of graft-versus-host disease by donor CD4+ 25+ T cells to allow an effective graft-versus-leukemia response. *Biol Blood Marrow Transplant.* (2003)
- Patterson AE et al. Infusion of select leukemia-reactive TCR Vbeta+ T cells provides graft-versus-leukemia responses with minimization of graft-versus-host disease following murine hematopoietic stem cell transplantation. *Biol Blood Marrow Transplant.* (2001)
- Schnaper HW et al. Identification and initial characterization of concanavalin A- and interferon-induced human suppressor factors: evidence for a human equivalent of murine soluble immune response suppressor (SIRS). *J Immunol.* (1984)
- Rittenberg MB et al. In vitro initiated secondary anti-hapten response. 3. Separable roles of hapten and carrier in immune paralysis. *Immunohistochemistry.* (1972)

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