

Anti-FceR1 α (human IgE receptor) antibody, mouse monoclonal (CRA2)

Product code	72-005
Size	100 μg
Storage	-20°C
Concentration	1.0 mg/ml
Buffer	PBS- with 50% glycerol
Purity	Purified IgG fraction with protein A from hybridoma cell culture medium.
Immunogen	Purified recombinant extracellular portion of human FceR1a (corresponding to amino acids Met-26-197, where signal peptide is 1-25)
Isotype	Mouse IgG1 κ
Reactivity	Human
Special notes	Epitope: Amino acids 110-197 of FcεR1α (Ref 3)
Application	 Western blotting (~1μg/ml) (Ref 2, 3) Flow-Cytometry (Ref 1,2) Immunohistochemistry (Paraffin and Frozen) and immunocytochemistry (Ref 4) Inhibition of binding of IgE with FcεR1α (Ref 2) Titration of IgE-bound fraction of the FcεR1α using CRA1 and CRA2 antibodies (Ref 2) Stimulation of serotonin release from human platelets. (Ref 1)
Background	FceR1 α is subunit of the high affinity receptor for IgE to which IgE directly binds. FceR1 is a tetrameric complex consisting of one α , one β and two γ subunits. The latter two subunits are required for signal transduction activity. The FceR1 α complex plays an important role in triggering allergic responses. The CRA2 (AER24) monoclonal antibody reacts with the FceR1 α subunit on a region that overlaps the region of the IgE binding site, thus it competes with IgE for the receptor binding. Since the CRA1 (AER37) monoclonal antibody reacts with the site different from the IgE binding site on FceR1 α , it does not compete with IgE for the receptor binding. Combining the two antibodies, one can quantitatively measure the amounts of the IgE-bound FceR1 α .
Data Link	UniProtKB/Swiss-Prot P12319 (FCERA_HUMAN)

PROCEDURES. NOT FOR MILITARY USE.



Data Images: 72-005 Anti-FceR1a (human IgE receptor) antibody, mouse monoclonal (CRA2)

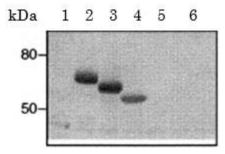


Fig.1 Epitope mapping of clone CRA2 of anti-FceR1a monoclonal antibody by western blotting.

Samples are maltose binding protein fused truncated.

Extra-cellular domain of FceR1a expressed in E.coli.

1.MalE-LacZ

2.26-197

3.68-197

4. 26-109

5. 26-153

6.68-153

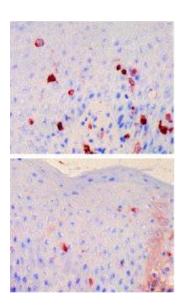


Fig.2 Immunohistochemical staining of skin sections from atopic dermatitis lesional skin with anti-FceR1a antibodies.

Aceton-fixed cryostat sections were incubated with either anti- FceR1a antibody clone CRA1 (above) of CRA2 (below)and positive reactions were visualized using the LLSAB kit (Dako, Denmark). CRA1 recognize non-IgE binding site of FceR1a while CRA2 recognize IgE binding site. Thus CRA2 can not bind to IgE-bound FceRIa.



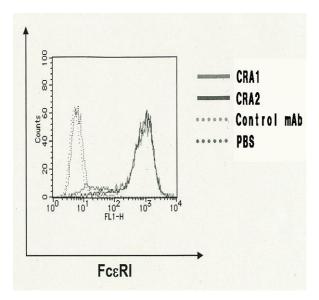


Fig.3 Flow-cytometry of CHO/Fc ϵ R1 α cells with CRA1 and CRA2 antibodies

CHO cells were transfected with plasmid expressing human FceRIa. The second antibody is FITC-conjugated anti-mouse IgG2b antibody.

References: Anti-FceR1a monoclonal antibody (CRA2) has been used in the following publications

- 1. Takai T *et al* "Epitope analysis and primary structures of variable regions of anti-human FcepsilonRI monoclonal antibodies, and expression of the chimeric antibodies fused with human constant regions" *Biosci Biotechnol Biochem* 64:1856-1867(2000) PMID: <u>11055388</u>
- 2. Takai T *et al* "Direct expression of the extracellular portion of human FcepsilonRIalpha chain as inclusion bodies in Escherichia coli "*Biosci Biotechnol Biochem* 65:79-85 (2001) PMID: <u>11272849</u>
- 3. Hasegawa S *et al.* "Functional Expression of the High Affinity Receptor for IgE (FceRI) in Human Platelets and Its' Intracellular Expression in Human Megakaryocytes" Blood 93: 2543-2551 (1999) PMID: 10194433
- 4. Goto T *et al.* "Enhanced expression of the high-affinity receptor for IgE (Fc(epsilon)RI) associated with decreased numbers of Langerhans cells in the lesional epidermis of atopic dermatitis" <u>J</u> Dermatol Sci. 27:156-61 (2001) PMID: 11641054

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