

Product Data Sheet

Purified anti-CRP (C-reactive protein)

Catalog # / Size: 620802 / 200 µl (20 Western blots)

Clone: Poly6208 Isotype: Rabbit IgG

Immunogen: Recombinant (partial)

Reactivity: Human

Preparation: The antibody was purified by antigen-affinity chromatography.

Formulation: This antibody is provided in phosphate-buffered solution, pH 7.2, containing

0.09% sodium azide and 50% glycerol.

Storage: Upon receipt, store frozen at -20° C.

Applications:

Applications: WB - Quality tested

Recommended Usage: Each lot of this antibody is quality control tested by Western blotting. Western

blotting, suggested working dilution(s): Use 10 µl per 5 ml antibody dilution buffer for each mini-gel. It is recommended that the reagent be titrated for

optimal performance for each application.

Description: CRP (C-reactive protein) is a member of the pentaxin family. CRP exists as a

homopentamer in a discoid arrangement (noncovalently attached). Individual subunits are approximately 24 kD, the molecular weight of the entire molecule is approximately 120 kD CRP is secreted and can be found in plasma. This protein promotes agglutination, phagocytosis, and complement fixation through Ca²⁺-dependent binding to phosphorylcholine. CRP is thought to scavenge nuclear material released from damaged circulating cells. CRP is the major protein involved in acute phase response to pathogens; both CRP and degradation products of CRP act as anti-inflammatory agents. CRP has been shown to interact with both DNA

and histones. The Poly6208 antibody recognizes human CRP and has been

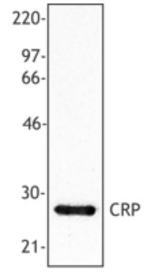
shown to be useful for Western blotting.

Antigen References: 1. Whitehead AS, et al. 1983. Science 221:69.
2. Tucci A, et al. 1983. J. Immunol. 131:2416.

3. Szalai AJ. 2002. Vascul. Pharmacol. 39:105 (review). 4. Chang MK, et al. 2002. P. Natl. Acad. Sci. USA 99:13043.

Related Products: Product Clone

Application ELÍSA, IHC, WB HRP Donkey anti-rabbit IgG (minimal x-reactivity) Poly4064



Recombinant human C-reactive protein (20 ng per lane) was resolved by electrophoresis, transferred to nitrocellulose, and probed with rabbit polyclonal antibody against CRP. Proteins were visualized using a donkey anti-rabbit secondary conjugated to HRP and a chemiluminescence detection system.



