

**RAG2 Antibody (C-term) Blocking peptide**  
**Synthetic peptide**  
**Catalog # BP12445b****Specification**

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**RAG2 Antibody (C-term) Blocking peptide - Product Information**Primary Accession [P55895](#)**RAG2 Antibody (C-term) Blocking peptide - Additional Information****Gene ID** 5897**Other Names**

V(D)J recombination-activating protein 2, RAG-2, RAG2

**Format**

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

**Storage**

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

**Precautions**

This product is for research use only. Not for use in diagnostic or therapeutic procedures.

**RAG2 Antibody (C-term) Blocking peptide - Protein Information****Name** RAG2**Function**

Core component of the RAG complex, a multiprotein complex that mediates the DNA cleavage phase during V(D)J recombination. V(D)J recombination assembles a diverse repertoire of immunoglobulin and T- cell receptor genes in developing B and T-lymphocytes through rearrangement of different V (variable), in some cases D (diversity), and J (joining) gene segments. DNA cleavage by the RAG complex occurs in 2 steps: a first nick is introduced in the top strand immediately upstream of the heptamer, generating a 3'-hydroxyl group that can attack the phosphodiester bond on the opposite strand in a direct transesterification reaction, thereby creating 4 DNA ends: 2 hairpin coding ends and 2 blunt, 5'-phosphorylated ends. The chromatin structure plays an essential role in the V(D)J recombination reactions and the presence of histone H3 trimethylated at 'Lys-4' (H3K4me3) stimulates both the nicking and hairpinning steps. The RAG complex also plays a role in pre-B cell allelic exclusion, a process leading to expression of a single immunoglobulin heavy chain allele to enforce clonality and monospecific recognition by the B-cell antigen receptor (BCR) expressed on individual B-lymphocytes. The introduction of DNA breaks by the RAG complex on one immunoglobulin allele induces ATM- dependent repositioning of the other allele to pericentromeric heterochromatin, preventing accessibility to the RAG complex and recombination of the second allele. In the RAG complex, RAG2 is not the catalytic component but is required for all known catalytic activities mediated by RAG1. It probably acts as a sensor of chromatin state that recruits the RAG complex to H3K4me3 (By similarity).

**Cellular Location**

Nucleus.

**Tissue Location**

Cells of the B- and T-lymphocyte lineages.

**RAG2 Antibody (C-term) Blocking peptide - Protocols**

Provided below are standard protocols that you may find useful for product applications.

- [Blocking Peptides](#)

**RAG2 Antibody (C-term) Blocking peptide - Images****RAG2 Antibody (C-term) Blocking peptide - Background**

This gene encodes a protein that is involved in the initiation of V(D)J recombination during B and T cell development. This protein forms a complex with the product of the adjacent recombination activating gene 1, and this complex can form double-strand breaks by cleaving DNA at conserved recombination signal sequences. The recombination activating gene 1 component is thought to contain most of the catalytic activity, while the N-terminal of the recombination activating gene 2 component is thought to form a six-bladed propeller in the active core that serves as a binding scaffold for the tight association of the complex with DNA. A C-terminal plant homeodomain finger-like motif in this protein is necessary for interactions with chromatin components, specifically with histone H3 that is trimethylated at lysine 4. Mutations in this gene cause Omenn syndrome, a form of severe combined immunodeficiency associated with autoimmune-like symptoms.

**RAG2 Antibody (C-term) Blocking peptide - References**

Davila, S., et al. Genes Immun. 11(3):232-238(2010)  
Couedel, C., et al. J. Clin. Invest. 120(4):1337-1344(2010)  
Hosgood, H.D. III, et al. Occup Environ Med 66(12):848-853(2009)  
Liang, X.S., et al. Br. J. Haematol. 146(4):418-423(2009)  
Ameratunga, R., et al. N. Z. Med. J. 122(1304):46-53(2009)