



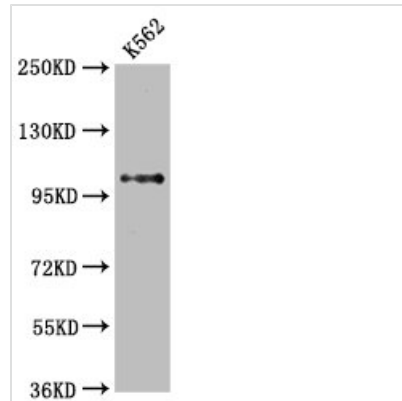
DGCR8 Recombinant Monoclonal Antibody

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|----------------------------|--|
| Product Code | CSB-RA845175A0HU |
| Abbreviation | Microprocessor complex subunit DGCR8 |
| Storage | Upon receipt, store at -20°C or -80°C. Avoid repeated freeze. |
| Uniprot No. | Q8WYQ5 |
| Immunogen | A synthesized peptide derived from human DGCR8 |
| Species Reactivity | Human |
| Tested Applications | ELISA, WB; Recommended dilution: WB:1:500-1:5000 |
| Relevance | <p>Component of the microprocessor complex that acts as a RNA- and heme-binding protein that is involved in the initial step of microRNA (miRNA) biogenesis. Component of the microprocessor complex that is required to process primary miRNA transcripts (pri-miRNAs) to release precursor miRNA (pre-miRNA) in the nucleus. Within the microprocessor complex, DGCR8 function as a molecular anchor necessary for the recognition of pri-miRNA at dsRNA-ssRNA junction and directs DROSHA to cleave 11 bp away from the junction to release hairpin-shaped pre-miRNAs that are subsequently cut by the cytoplasmic DICER to generate mature miRNAs (PubMed:26027739, PubMed:26748718). The heme-bound DGCR8 dimer binds pri-miRNAs as a cooperative trimer (of dimers) and is active in triggering pri-miRNA cleavage, whereas the heme-free DGCR8 monomer binds pri-miRNAs as a dimer and is much less active. Both double-stranded and single-stranded regions of a pri-miRNA are required for its binding (PubMed:15531877, PubMed:15574589, PubMed:15589161, PubMed:16751099, PubMed:16906129, PubMed:16963499, PubMed:17159994). Specifically recognizes and binds N6-methyladenosine (m6A)-containing pri-miRNAs, a modification required for pri-miRNAs processing (PubMed:25799998). Involved in the silencing of embryonic stem cell self-renewal (By similarity).</p> |
| Form | Liquid |
| Conjugate | Non-conjugated |
| Storage Buffer | Rabbit IgG in phosphate buffered saline , pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. |
| Purification Method | Affinity-chromatography |
| Isotype | Rabbit IgG |
| Clonality | Monoclonal |
| Alias | Microprocessor complex subunit DGCR8, DiGeorge syndrome critical region 8, DGCR8, C22orf12, DGCRK6, LP4941 |
| Immunogen Species | Homo sapiens (Human) |
| Research Area | Epigenetics and Nuclear Signaling |
| Gene Names | DGCR8 |



Clone No. 2H2

Image



Western Blot

Positive WB detected in: K562 whole cell lysate

All lanes: DGCR8 antibody at 2.65μg/ml

Secondary

Goat polyclonal to rabbit IgG at 1/50000 dilution

Predicted band size: 87, 33, 83 KDa

Observed band size: 100 KDa

Description

The recombinant DGCR8 antibody is a monoclonal antibody made in vitro using the DGCR8 antibody genes that are typically expressed from a plasmid in a stable mammalian cell line. The genes coding for the DGCR8 antibody will ultimately assemble into a fully functional antibody after translation. The synthesized antibody is the recombinant antibody against DGCR8. It underwent purification using affinity-chromatography. This recombinant DGCR8 antibody is suitable for use in the ELISA, WB to detect the DGCR8 protein from Human.

DGCR8 is an allele located in the q11.2 region of human chromosome 22. It is associated with DiGeorge Syndrome (DGCS), so it is named DGCR83. DGCR8 is an RNA-binding protein involved in the synthesis of microRNAs. In the synthesis of microRNAs, it binds pri-microRNAs and promotes the cleavage of Drosha by interacting with Drosha and stabilizing microprocessors, which regulate the production of microRNAs and affect the proliferation, migration, and invasion of tumors^{4,5}. The high expression of DGCR8 promotes the occurrence, development, and metastasis of cancer, including kidney clear cell carcinoma, thyroid carcinoma, and ovarian cancer. DGCR8 is abnormally expressed in TNBC specimens. Besides, the knockdown of DGCR8 inhibited cell migration and invasion in TNBC cells, while the overexpression of DGCR8 promoted cell migration and invasion in TNBC cells. The above results indicated that DGCR8 promotes the metastasis of TNBC and may act as an oncogene.