

## DOCK8 gene

dedicator of cytokinesis 8

### Normal Function

The *DOCK8* gene provides instructions for making a member of the DOCK family of proteins. The proteins in this family act as guanine nucleotide exchange factors (GEFs). GEFs turn on (activate) proteins called GTPases, which play an important role in chemical signaling within cells. Signaling stimulated by DOCK family proteins are typically involved in the arrangement of the structural framework inside cells (the cytoskeleton). By controlling the shape of the cytoskeleton, DOCK family proteins play a role in cell structure and movement (migration).

The DOCK8 protein is found most abundantly in cells of the immune system. This protein plays a critical role in the survival and function of several types of immune system cells, including T cells, NK cells, and B cells. T cells and NK cells recognize and attack foreign invaders, such as viruses, to prevent infection. B cells produce proteins called antibodies, which attach to foreign particles and germs and mark them for destruction.

Through its function as a GEF, the DOCK8 protein helps maintain the structure and integrity of T cells and NK cells. It also aids in the movement of these immune system cells to sites of infection, particularly the skin. The DOCK8 protein is also involved in chemical signaling pathways that stimulate B cells to mature and produce antibodies. The protein is also involved in the normal development and survival of other types of immune system cells.

### Health Conditions Related to Genetic Changes

#### DOCK8 immunodeficiency syndrome

More than 130 mutations in the *DOCK8* gene have been found to cause DOCK8 immunodeficiency syndrome (also called autosomal recessive hyper-IgE syndrome or AR-HIES). DOCK8 immunodeficiency syndrome is an immune system disorder that causes recurrent severe infections of the skin and respiratory tract. Affected individuals may have other immune system problems, such as allergies, asthma, or an inflammatory skin disorder called eczema. Most of the mutations involved in DOCK8 immunodeficiency syndrome delete regions of DNA from the *DOCK8* gene. These deletions and other *DOCK8* gene mutations lead to production of an abnormally short

protein or production of no protein. As a result, affected individuals have little or no functional DOCK8 protein.

A shortage of DOCK8 protein impairs normal immune cell development and function. It is thought that T cells lacking DOCK8 protein cannot maintain their shape as they move through dense spaces, such as those found within the skin. The abnormal cells die too easily, resulting in reduced numbers of these cells. A shortage of T cells impairs the immune response to foreign invaders, accounting for the severe skin infections common in DOCK8 immunodeficiency syndrome. A lack of DOCK8 protein also impairs B cell maturation and the production of certain antibodies. Impairment of this type of immune response leads to recurrent respiratory tract infections in people with this disorder.

For unknown reasons, a reduction of DOCK8 protein results in higher-than-normal production of an immune system protein known as immunoglobulin E (IgE), which plays a role in allergic reactions. As a result, people with DOCK8 immunodeficiency syndrome have an increased risk of food and environmental allergies.

### **Other Names for This Gene**

- 1200017A24Rik
- dedicator of cytokinesis protein 8 isoform 1
- dedicator of cytokinesis protein 8 isoform 2
- dedicator of cytokinesis protein 8 isoform 3
- epididymis luminal protein 205
- FLJ00026
- FLJ00152
- FLJ00346
- HEL-205
- MRD2
- ZIR8

### **Additional Information & Resources**

#### Tests Listed in the Genetic Testing Registry

- Tests of DOCK8 ([https://www.ncbi.nlm.nih.gov/gtr/all/tests/?term=81704\[geneid\]](https://www.ncbi.nlm.nih.gov/gtr/all/tests/?term=81704[geneid]))

#### Scientific Articles on PubMed

- PubMed (<https://pubmed.ncbi.nlm.nih.gov/?term=%28%28DOCK8%5BTIAB%5D%29+OR+%28dedicator+of+cytokinesis+8%5BTIAB%5D%29%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1800+days%22%5Bdp>)

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### Catalog of Genes and Diseases from OMIM

- DEDICATOR OF CYTOKINESIS 8; DOCK8 (<https://omim.org/entry/611432>)

### Gene and Variant Databases

- NCBI Gene (<https://www.ncbi.nlm.nih.gov/gene/81704>)
- ClinVar ([https://www.ncbi.nlm.nih.gov/clinvar?term=DOCK8\[gene\]](https://www.ncbi.nlm.nih.gov/clinvar?term=DOCK8[gene]))

### **References**

- Crawford G, Enders A, Gileadi U, Stankovic S, Zhang Q, Lambe T, Crockford TL, Lockstone HE, Freeman A, Arkwright PD, Smart JM, Ma CS, Tangye SG, Goodnow CC, Cerundolo V, Godfrey DI, Su HC, Randall KL, Cornall RJ. DOCK8 is critical for the survival and function of NKT cells. *Blood*. 2013 Sep 19;122(12):2052-61. doi:10.1182/blood-2013-02-482331. Epub 2013 Aug 8. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/23929855>) or Free article on PubMed Central (<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3778549/>)
- Engelhardt KR, McGhee S, Winkler S, Sassi A, Woellner C, Lopez-Herrera G, Chen A, Kim HS, Lloret MG, Schulze I, Ehl S, Thiel J, Pfeifer D, Veelken H, Niehues T, Siepermann K, Weinspach S, Reisli I, Keles S, Genel F, Kutukculer N, Camcioglu Y, Somer A, Karakoc-Aydiner E, Barlan I, Gennery A, Metin A, Degerliyurt A, Pietrogrande MC, Yeganeh M, Baz Z, Al-Tamemi S, Klein C, Puck JM, Holland SM, McCabe ER, Grimbacher B, Chatila TA. Large deletions and point mutations involving the dedicator of cytokinesis 8 (DOCK8) in the autosomal-recessive form of hyper-IgE syndrome. *J Allergy Clin Immunol*. 2009 Dec;124(6):1289-302.e4. doi:10.1016/j.jaci.2009.10.038. Erratum In: *J Allergy Clin Immunol*. 2010 Mar;125(3):743. Kutuculer, Necil [corrected to Kutukculer, Necil]. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/20004785>) or Free article on PubMed Central (<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2818862/>)
- Harada Y, Tanaka Y, Terasawa M, Pieczyk M, Habiro K, Katakai T, Hanawa-Suetsugu K, Kukimoto-Niino M, Nishizaki T, Shirouzu M, Duan X, Uruno T, Nishikimi A, Sanematsu F, Yokoyama S, Stein JV, Kinashi T, Fukui Y. DOCK8 is a Cdc42 activator critical for interstitial dendritic cell migration during immuneresponses. *Blood*. 2012 May 10;119(19):4451-61. doi: 10.1182/blood-2012-01-407098. Epub 2012 Mar 28. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/22461490>) or Free article on PubMed Central (<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3418773/>)
- Jabara HH, McDonald DR, Janssen E, Massaad MJ, Ramesh N, Borzutzky A, Rauterl, Benson H, Schneider L, Baxi S, Recher M, Notarangelo LD, Wakim R, Dbaiho G, Dasouki M, Al-Herz W, Barlan I, Baris S, Kutukculer N, Ochs HD, Plebani A, Kanariou M, Lefranc G, Reisli I, Fitzgerald KA, Golenbock D, Manis J, Keles S,

Ceja R, Chatila TA, Geha RS. DOCK8 functions as an adaptor that links TLR-MyD88 signaling to B cell activation. *Nat Immunol.* 2012 May 13;13(6):612-20. doi:10.1038/ni.2305. Erratum In: *Nat Immunol.* 2022 May;23(5):815. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/22581261>) or Free article on PubMed Central (<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3362684/>)

- Lambe T, Crawford G, Johnson AL, Crockford TL, Bouriez-Jones T, Smyth AM, Pham TH, Zhang Q, Freeman AF, Cyster JG, Su HC, Cornall RJ. DOCK8 is essential for T-cell survival and the maintenance of CD8+ T-cell memory. *Eur J Immunol.* 2011 Dec;41(12):3423-35. doi: 10.1002/eji.201141759. Epub 2011 Nov 10. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/21969276>) or Free article on PubMed Central (<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3517112/>)
- Randall KL, Lambe T, Johnson AL, Treanor B, Kucharska E, Domaschewicz H, Whittle B, Tze LE, Enders A, Crockford TL, Bouriez-Jones T, Alston D, Cyster JG, Lenardo MJ, Mackay F, Deenick EK, Tangye SG, Chan TD, Camidge T, Brink R, Vinuesa CG, Batista FD, Cornall RJ, Goodnow CC. Dock8 mutations cripple B cell immunological synapses, germinal centers and long-lived antibody production. *Nat Immunol.* 2009 Dec;10(12):1283-91. doi: 10.1038/ni.1820. Epub 2009 Nov 8. Erratum In: *Nat Immunol.* 2010 Jul;11(7):644. Johnson, Andy [corrected to Johnson, Andy L]. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/19898472>) or Free article on PubMed Central (<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3437189/>)
- Su HC, Jing H, Angelus P, Freeman AF. Insights into immunity from clinical and basic science studies of DOCK8 immunodeficiency syndrome. *Immunol Rev.* 2019 Jan;287(1):9-19. doi: 10.1111/imr.12723. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/30565250>) or Free article on PubMed Central (<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6350515/>)
- Zhang Q, Dove CG, Hor JL, Murdock HM, Strauss-Albee DM, Garcia JA, Mandl JN, Grodick RA, Jing H, Chandler-Brown DB, Lenardo TE, Crawford G, Matthews HF, Freeman AF, Cornall RJ, Germain RN, Mueller SN, Su HC. DOCK8 regulates lymphocyte shape integrity for skin antiviral immunity. *J Exp Med.* 2014 Dec 15;211(13):2549-66. doi: 10.1084/jem.20141307. Epub 2014 Nov 24. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/25422492>) or Free article on PubMed Central (<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4267229/>)
- Zhang Q, Jing H, Su HC. Recent Advances in DOCK8 Immunodeficiency Syndrome. *J Clin Immunol.* 2016 Jul;36(5):441-9. doi: 10.1007/s10875-016-0296-z. Epub 2016 May 20. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/27207373>) or Free article on PubMed Central (<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4914394/>)

## Genomic Location

The *DOCK8* gene is found on chromosome 9 (<https://medlineplus.gov/genetics/chromosome/9/>).

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