

DCXR Gene

Subjects: **Genetics & Heredity**

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Dicarbonyl and L-Xylulose Reductase: The DCXR gene provides instructions for making a protein called dicarbonyl and L-xylulose reductase (DCXR), which plays multiple roles in the body.

genes

1. Normal Function

One of its functions is to perform a chemical reaction that converts a sugar called L-xylulose to a molecule called xylitol. This reaction is one step in a process by which the body can use sugars for energy. There are two versions of L-xylulose reductase in the body, known as the major isoform and the minor isoform. The *DCXR* gene provides instructions for making the major isoform, which converts L-xylulose more efficiently than the minor isoform. It is unclear if the minor isoform is produced from the *DCXR* gene or another gene.

Another function of the DCXR protein is to break down toxic compounds called alpha-dicarbonyl compounds. These compounds, which are byproducts of certain cellular processes or are found in foods in the diet, must be broken down so they do not damage cells.

The DCXR protein is also one of several proteins that get attached to the surface of sperm cells as they mature. DCXR is involved in the interaction of a sperm cell with an egg cell during fertilization.

2. Health Conditions Related to Genetic Changes

2.1 Essential Pentosuria

At least two mutations in the *DCXR* gene cause a condition called essential pentosuria, which is found almost exclusively in individuals with Ashkenazi Jewish ancestry. Affected individuals have high levels of L-xylulose in their urine, but they have no associated health problems. The gene mutations involved in this condition lead to the production of altered DCXR proteins that are quickly broken down. Without this protein, L-xylulose is not efficiently converted to xylitol, and the excess sugar is released in the urine. Only the major isoform of L-xylulose reductase is affected by these mutations, but the minor isoform cannot compensate for the loss of DCXR's function in breaking down L-xylulose. It is thought that other processes are able to break down toxic alpha-dicarbonyl compounds, likely accounting for the lack of symptoms in people with essential pentosuria.

Males with essential pentosuria appear to have normal reproductive function, despite studies that show that a shortage of DCXR protein attached to sperm cells can be associated with the inability to have biological children (infertility).

3. Other Names for This Gene

- carbonyl reductase 2
- carbonyl reductase II
- DCR
- dicarbonyl/L-xylulose reductase
- HCR2
- HCRII
- KIDCR
- kidney dicarbonyl reductase
- L-xylulose reductase
- L-xylulose reductase isoform 1
- L-xylulose reductase isoform 2
- P34H
- SDR20C1
- short chain dehydrogenase/reductase family 20C, member 1
- sperm surface protein P34H
- XR

References

1. Boué F, Blais J, Sullivan R. Surface localization of P34H an epididymalprotein, during maturation, capacitation, and acrosome reaction of humanspermatozoa. *Biol Reprod.* 1996 May;54(5):1009-17.
2. Ebert B, Kisiela M, Maser E. Human DCXR - another 'moonlighting protein'involved in sugar metabolism, carbonyl detoxification, cell adhesion and malefertility? *Biol Rev Camb Philos Soc.* 2015 Feb;90(1):254-78. doi:10.1111/brv.12108.
3. Lane AB. On the nature of L-xylulose reductase deficiency in essentialpentosuria. *Biochem Genet.* 1985 Feb;23(1-2):61-72.
4. Lee SK, Son le T, Choi HJ, Ahnn J. Dicarbonyl/L-xylulose reductase (DCXR): Themultifunctional pentosuria enzyme. *Int J Biochem Cell Biol.* 2013Nov;45(11):2563-7. doi: 10.1016/j.biocel.2013.08.010.
5. Nakagawa J, Ishikura S, Asami J, Isaji T, Usami N, Hara A, Sakurai T, Tsuritani K, Oda K, Takahashi M, Yoshimoto M, Otsuka N, Kitamura K. Molecularcharacterization of mammalian dicarbonyl/L-xylulose reductase and itslocalization in kidney. *J Biol Chem.* 2002 May 17;277(20):17883-91.
6. Pierce SB, Spurrell CH, Mandell JB, Lee MK, Zeligson S, Bereman MS, Stray SM, Fokstuen S, MacCoss MJ, Levy-Lahad E, King MC, Motulsky AG. Garrod's fourthinborn error of metabolism solved by the identification of mutations causingpentosuria. *Proc Natl Acad Sci U S A.* 2011 Nov 8;108(45):18313-7. doi:10.1073/pnas.1115888108.

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