


Fraser Syndrome

Subjects: Genetics

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Definition

Fraser syndrome is a rare disorder that affects development starting before birth. Characteristic features of this condition include eyes that are completely covered by skin and usually malformed (cryptophthalmos), fusion of the skin between the fingers and toes (cutaneous syndactyly), and abnormalities of the genitalia and the urinary tract (genitourinary anomalies). Other tissues and organs can also be affected. Depending on the severity of the signs and symptoms, Fraser syndrome can be fatal before or shortly after birth; less severely affected individuals can live into childhood or adulthood.

1. Introduction

Cryptophthalmos is the most common abnormality in people with Fraser syndrome. Both eyes are usually completely covered by skin, but in some cases, only one eye is covered or one or both eyes are partially covered. In cryptophthalmos, the eyes can also be malformed; for example, the eyeballs may be fused to the skin covering them, or they may be small (microphthalmia) or missing (anophthalmia). Eye abnormalities typically lead to impairment or loss of vision in people with Fraser syndrome. Affected individuals can have other problems related to abnormal eye development, including missing eyebrows or eyelashes or a patch of hair extending from the side hairline to the eyebrow.

Cutaneous syndactyly typically occurs in both the hands and the feet in Fraser syndrome. In most people with this feature, the skin between the middle three fingers and toes are fused, but the other digits can also be involved. Other abnormalities of the hands and feet can occur in people with Fraser syndrome.

Individuals with Fraser syndrome can have abnormalities of the genitalia, such as an enlarged clitoris in females or undescended testes (cryptorchidism) in males. Some affected individuals have external genitalia that do not appear clearly female or male (ambiguous genitalia).

The most common urinary tract abnormality in Fraser syndrome is the absence of one or both kidneys (renal agenesis). Affected individuals can have other kidney problems or abnormalities of the bladder and other parts of the urinary tract.

A variety of other signs and symptoms can be involved in Fraser syndrome, including heart malformations or abnormalities of the voicebox (larynx) or other parts of the respiratory tract. Some affected individuals have facial abnormalities, including ear or nose abnormalities or an opening in the upper lip (cleft lip) with or without an opening in the roof of the mouth (cleft palate).

2. Frequency

Fraser syndrome affects an estimated 1 in 200,000 newborns. The condition occurs in approximately 1 in 10,000 fetuses that do not survive to birth.

3. Causes

Mutations in the *FRAS1*, *FREM2*, or *GRIP1* gene can cause Fraser syndrome. *FRAS1* gene mutations are the most common cause, accounting for about half of cases of Fraser syndrome. *FREM2* and *GRIP1* gene mutations are each found in a small percentage of cases.

The *FRAS1* and *FREM2* proteins (produced from the *FRAS1* and *FREM2* genes, respectively) are part of a group of proteins called the FRAS/FREM complex. The *GRIP1* protein (produced from the *GRIP1* gene)

ensures that FRAS1 and FREM2 get to the correct location of the cell to form the FRAS/FREM complex.

The FRAS/FREM complex is found in basement membranes, which are thin, sheet-like structures that separate and support cells in many tissues. This complex is particularly important during development before birth. One of the complex's roles is to anchor the top layer of skin by connecting its basement membrane to the layer of skin below. The FRAS/FREM complex is also involved in the proper development of other organs and tissues, including the kidneys, although the mechanism is unclear.

Mutations in any of these genes prevent formation of the FRAS/FREM complex. Lack of this complex in the basement membrane of the skin leads to detachment of the top layer of skin, causing blisters to form during development. These blisters likely impair the proper formation of certain structures before birth, leading to cryptophthalmos and cutaneous syndactyly. It is unknown how lack of the FRAS/FREM complex leads to kidney and genital abnormalities and other problems in Fraser syndrome.

3.1. The Genes Associated with Fraser Syndrome

- FRAS1
- FREM2
- GRIP1

4. Inheritance

This condition is inherited in an autosomal recessive pattern, which means both copies of the gene in each cell have mutations. The parents of an individual with an autosomal recessive condition each carry one copy of the mutated gene, but they typically do not show signs and symptoms of the condition.

5. Other Names for This Condition

- cryptophthalmos syndactyly syndrome
- cryptophthalmos syndrome
- cryptophthalmos with other malformations
- Fraser's syndrome
- Fraser-Francois syndrome
- Meyer-Schwickerath syndrome
- Ullrich-Feichtiger syndrome

References

1. Barisic I, Odak L, Loane M, Garne E, Wellesley D, Calzolari E, Dolk H, AddorMC, Arriola L, Bergman J, Bianca S, Boyd PA, Draper ES, Gatt M, Haeusler M, Khoshnood B, Latos-Bielenska A, McDonnell B, Pierini A, Rankin J, Rissmann A, Queisser-Luft A, Verellen-Dumoulin C, Stone D, Tenconi R. Fraser syndrome: epidemiological study in a European population. *Am J Med Genet A*. 2013 May;161A(5):1012-8. doi: 10.1002/ajmg.a.35839.
2. McGregor L, Makela V, Darling SM, Vrontou S, Chalepakis G, Roberts C, Smart N, Rutland P, Prescott N, Hopkins J, Bentley E, Shaw A, Roberts E, Mueller R, Jadeja S, Philip N, Nelson J, Francannet C, Perez-Aytes A, Megarbane A, Kerr B, Wainwright B, Woolf AS, Winter RM, Scambler PJ. Fraser syndrome and mouse blebbed phenotype caused by mutations in FRAS1/Fras1 encoding a putative extracellular matrix protein. *Nat Genet*. 2003 Jun;34(2):203-8.
3. Petrou P, Makrygiannis AK, Chalepakis G. The Fras1/Frem family of extracellular matrix proteins: structure, function, and association with Fraser syndrome and the mouse bleb phenotype. *Connect Tissue Res*. 2008;49(3):277-82. doi: 10.1080/03008200802148025.
4. Slavotinek AM, Tift CJ. Fraser syndrome and cryptophthalmos: review of the diagnostic criteria and evidence for phenotypic modules in complex malformation syndromes. *J Med Genet*. 2002 Sep;39(9):623-33. Review.
5. van Haelst MM, Maiburg M, Baujat G, Jadeja S, Monti E, Bland E, Pearce K; Fraser Syndrome Collaboration Group, Hennekam RC, Scambler PJ. Molecular study of 33 families with Fraser syndrome: new data and mutation review. *Am J Med Genet A*. 2008 Sep 1;146A(17):2252-7. doi: 10.1002/ajmg.a.32440.
6. van Haelst MM, Scambler PJ; Fraser Syndrome Collaboration Group, Hennekam RC. Fraser syndrome: a clinical study of 59 cases and evaluation of diagnostic criteria. *Am J Med Genet A*. 2007 Dec 15;143A(24):3194-203.
7. Vogel MJ, van Zon P, Brueton L, Gijzen M, van Tuil MC, Cox P, Schanze D, Kariminejad A, Ghaderi-Sohi S, Blair E, Zenker

M, Scambler PJ, Ploos van AmstelHK, van Haelst MM. Mutations in GRIP1 cause Fraser syndrome. J Med Genet. 2012May;49(5):303-6. doi: 10.1136/jmedgenet-2011-100590.

Keywords

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