

**DISEASE:**  
**Familial dysfibrinogenemia**

<b>NAME:</b>	Familial dysfibrinogenemia
<b>DESCRIPTION:</b>	Familial dysfibrinogenemia is a coagulation disorder characterized by a bleeding tendency due to a functional anomaly of circulating fibrinogen.
<b>ORPHACODE:</b>	98881
<b>XREF(S):</b>	<u>Orphanet</u> <u>OMIM</u> <u>ICD-10</u>
<b>ANALYTE(S):</b>	<u>FGA</u> <u>FGB</u> <u>FGG</u>
<b>CREATED:</b>	13 May 2019 - 01:02
<b>CHANGED:</b>	22 Jun 2023 - 16:14

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## RELATED CONTENT

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### Related Genetic Tests

- [Trombosis - Hemostasis \(gene panel\)](#)

### Related Laboratories

- [Centrum Menselijke Erfelijkheid - KUL](#)

### Related Analytes

- [fibrinogen alpha chain](#)
- [fibrinogen beta chain](#)
- [fibrinogen gamma chain](#)

### Related Gene Panels

- [Trombosis - Hemostasis \(107 genes\) - KUL](#)

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