

**DISEASE:**  
**Hemochromatosis type 4**

|                     |                                                                                                                                                                                                                     |
|---------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <b>NAME:</b>        | Hemochromatosis type 4                                                                                                                                                                                              |
| <b>DESCRIPTION:</b> | Hemochromatosis type 4 (also called ferroportin disease) is a form of rare hereditary hemochromatosis (HH; see this term), a group of diseases characterized by excessive tissue iron deposition of genetic origin. |
| <b>ORPHACODE:</b>   | 139491                                                                                                                                                                                                              |
| <b>SYNONYMS:</b>    | Autosomal dominant hereditary hemochromatosis<br>Ferroportin disease<br>Hemochromatosis due to defect in ferroportin                                                                                                |
| <b>XREF(S):</b>     | <u>Orphanet</u><br><u>OMIM</u><br><u>MeSH</u><br><u>ICD-10</u>                                                                                                                                                      |
| <b>ANALYTE(S):</b>  | <u>SLC40A1</u><br><u>SLC40A1</u>                                                                                                                                                                                    |
| <b>CREATED:</b>     | 13 May 2019 - 01:02                                                                                                                                                                                                 |
| <b>CHANGED:</b>     | 01 May 2022 - 06:55                                                                                                                                                                                                 |

## RELATED CONTENT

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### Related Genetic Tests

- [Hemochromatosis \(17 genes\)](#)
- [Hemochromatosis hereditary type 4 \(SLC40A1 gene\)](#)
- [Hemochromatosis hereditary type 2 to type 5 \(5 genes\)](#)

### Related Laboratories

- [Centre de Génétique Humaine - CHU Sart-Tilman](#)
- [Centre de Génétique Humaine - Erasme ULB](#)
- [Centre de Génétique Médicale UCL](#)

### Related Analytes

- [solute carrier family 40 member 1](#)

### Related Gene Panels

- [Hemochromatosis \(17 genes\) - ULB](#)
- [Hemochromatosis hereditary \(types 2 to 5\) \(5 genes\) - UCL](#)

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