Export to PDF

Full name:	Hemochromatosis (17 genes) - ULB
Laboratory:	Centre de Génétique Humaine - Erasme ULB
Created:	15 Jul 2019 - 10:46
Changed:	10 Mar 2022 - 09:31

Related Diseases

- Aceruloplasminemia
- Autosomal recessive sideroblastic anemia
- Congenital atransferrinemia
- Congenital dyserythropoietic anemia type I
- Congenital dyserythropoietic anemia type II
- FTH1-related iron overload
- Genetic hyperferritinemia without iron overload
- HJV or HAMP-related hemochromatosis
- Hemochromatosis type 4
- Hereditary hyperferritinemia-cataract syndrome
- IRIDA syndrome
- L-ferritin deficiency
- Microcytic anemia with liver iron overload
- Neuroferritinopathy
- Severe congenital hypochromic anemia with ringed sideroblasts
- Symptomatic form of hemochromatosis type 1
- TFR2-related hemochromatosis
- X-linked erythropoietic protoporphyria
- X-linked sideroblastic anemia

Related Analytes

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
ALAS2	100.00	1	NM_000032

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
CDAN1	100.00	1	NM_138477
CP	100.00	1	NM_000096
FTH1	100.00	1	NM_002032
FTL	100.00	1	NM_000146
HAMP	100.00	1	Core gene / NM_021175
HFE1	100.00	1	NM_000410
HJV	100.00	1	Core gene / NM_213653
HMOX1	100.00	1	NM_002133
SEC23B	100.00	1	NM_006363
SLC11A2	100.00	1	Core gene / NM_000617
SLC25A38	100.00	1	NM_017875
SLC40A1	100.00	1	NM_014585
STEAP3	100.00	1	NM_001008410
TF	100.00	1	NM_001063
TFR2	100.00	1	Core gene / NM_003227
TMPRSS6	100.00	1	NM_153609