IRON DISORDERS PANEL DG-4.0.0 (54 GENES)

Gene	*Twist X2 covered >10x*	*Twist X2 covered >20x*	*WGS covered >10x*	*WGS covered >20x*	*Associated Phenotype description and OMIM disease ID*
ABCB10	100.0%	100.0%	99.9%	96.9%	
ABCB7	99.8%	99.3%	98.3%	74.8%	Anemia, sideroblastic, with ataxia, 301310
ACVR1	100.0%	99.9%	100.0%	98.4%	Fibrodysplasia ossificans progressiva, 135100
ALAS2	100.0%	99.8%	98.3%	72.9%	Anemia, sideroblastic, 1, 300751;Protoporphyria, erythropoietic, X-linked, 300752
ATP4A	100.0%	100.0%	100.0%	98.1%	
BMP6	100.0%	100.0%	99.9%	95.4%	{Iron overload, susceptibility to}, 620121
CALR	100.0%	100.0%	100.0%	99.2%	Myelofibrosis, somatic, 254450;Thrombocythemia, somatic, 187950
CCL2	100.0%	100.0%	100.0%	95.6%	{Mycobacterium tuberculosis, susceptibility to}, 607948;{HIV-1, resistance to}, 609423;{Spina bifida, susceptibility to}, 182940;{Coronary artery disease, modifier of},

CDAN1	100.0%	100.0%	99.9%	96.7%	Dyserythropoietic anemia, congenital, type Ia, 224120
CDIN1	100.0%	99.9%	100.0%	99.0%	Dyserythropoietic anemia, congenital, type Ib, 615631
CP	100.0%	100.0%	100.0%	98.7%	Aceruloplasminemia, 604290
CYBRD1	100.0%	100.0%	100.0%	98.3%	
EXOC6	100.0%	100.0%	100.0%	97.3%	
FECH	100.0%	100.0%	100.0%	99.1%	Protoporphyria, erythropoietic, 1, 177000
FTH1	100.0%	100.0%	100.0%	98.1%	Neurodegeneration with brain iron accumulation 9, 620669;?Hemochromatosis, type 5, 615517
FTL	100.0%	100.0%	100.0%	96.5%	Hyperferritinemia-cataract syndrome, 600886;L-ferritin deficiency, dominant and recessive, 615604;Neurodegeneration with brain iron accumulation 3, 606159
FXN	100.0%	100.0%	100.0%	95.9%	Friedreich ataxia with retained reflexes, 229300;Friedreich ataxia, 229300

GATA1	100.0%	100.0%	97.4%	68.7%	Leukemia, megakaryoblastic, with or without Down syndrome, somatic, 159595;Thrombocytopenia, X-linked, with or without dyserythropoietic anemia, 300367;Anemia, X-linked, with/without neutropenia and/or platelet abnormalities, 300835;Thrombocytopenia with beta-thalassemia, X-linked, 314050;Hemolytic anemia due to elevated adenosine deaminase, 301083
GLRX5	100.0%	100.0%	100.0%	97.1%	Anemia, sideroblastic, 3, pyridoxine-refractory, 616860;Spasticity, childhood-onset, with hyperglycinemia, 616859
HAMP	100.0%	100.0%	100.0%	99.8%	Hemochromatosis, type 2B, 613313
HEPH	99.8%	99.2%	98.0%	72.5%	
HFE	100.0%	100.0%	100.0%	98.3%	Hemochromatosis, type 1, 235200
HJV	100.0%	100.0%	100.0%	98.6%	Hemochromatosis, type 2A, 602390

HMOX1	100.0%	100.0%	100.0%	99.8%	Heme oxygenase-1 deficiency, 614034;{Pulmonary disease, chronic obstructive, susceptibility to}, 606963
HSCB	100.0%	100.0%	100.0%	97.6%	?Anemia, sideroblastic, 5, 619523
HSPA9	100.0%	100.0%	100.0%	98.5%	Even-plus syndrome, 616854;Anemia, sideroblastic, 4, 182170
JAK2	100.0%	100.0%	100.0%	98.3%	{Budd-Chiari syndrome, somatic}, 600880;Myelofibrosis, somatic, 254450;Erythrocytosis, somatic, 133100;Leukemia, acute myeloid, somatic, 601626;Thrombocythemia 3, 614521;Polycythemia vera, somatic, 263300
KIF23	100.0%	100.0%	100.0%	98.4%	Anemia, congenital dyserythropoietic, type IIIA, 105600
KLF1	100.0%	100.0%	100.0%	98.5%	Blood group--Lutheran inhibitor, 111150;Dyserythropoietic anemia, congenital, type IV, 613673;[Hereditary persistence of fetal hemoglobin], 613566

LARS2	100.0%	100.0%	100.0%	99.1%	Perrault syndrome 4, 615300;Hydrops, lactic acidosis, and sideroblastic anemia, 617021
LPIN2	99.3%	99.2%	100.0%	98.6%	Majeed syndrome, 609628
MPL	100.0%	100.0%	100.0%	98.8%	Myelofibrosis with myeloid metaplasia, somatic, 254450;Amegakaryocytic thrombocytopenia, congenital, 1, 604498;Thrombocythemia 2, 601977
NCOA4	100.0%	100.0%	100.0%	97.9%	
NDUFB11	99.7%	97.9%	88.1%	61.0%	Linear skin defects with multiple congenital anomalies 3, 300952;?Mitochondrial complex I deficiency, nuclear type 30, 301021
PANK2	100.0%	100.0%	100.0%	98.6%	Neurodegeneration with brain iron accumulation 1, 234200
PUS1	100.0%	100.0%	100.0%	98.5%	Myopathy, lactic acidosis, and sideroblastic anemia 1, 600462
SEC23B	100.0%	100.0%	100.0%	98.4%	?Cowden syndrome 7, 616858;Dyserythropoietic anemia, congenital, type II, 224100
SF3B1	100.0%	100.0%	100.0%	98.2%	Myelodysplastic syndrome, somatic, 614286

SFXN4	100.0%	100.0%	100.0%	96.9%	Combined oxidative phosphorylation deficiency 18, 615578
SLC11A2	100.0%	100.0%	100.0%	98.8%	Anemia, hypochromic microcytic, with iron overload 1, 206100
SLC19A2	100.0%	100.0%	100.0%	99.5%	Thiamine-responsive megaloblastic anemia syndrome, 249270
SLC25A37	100.0%	100.0%	100.0%	98.8%	
SLC25A38	100.0%	100.0%	100.0%	99.2%	Anemia, sideroblastic, 2, pyridoxine-refractory, 205950
SLC40A1	100.0%	100.0%	100.0%	98.9%	Hemochromatosis, type 4, 606069
SLC46A1	100.0%	100.0%	100.0%	98.7%	Folate malabsorption, hereditary, 229050
STEAP3	100.0%	100.0%	100.0%	99.2%	?Anemia, hypochromic microcytic, with iron overload 2, 615234
TF	100.0%	100.0%	100.0%	99.1%	Atransferrinemia, 209300
TFR2	100.0%	100.0%	100.0%	97.7%	Hemochromatosis, type 3, 604250
TFRC	95.5%	95.5%	100.0%	98.5%	Immunodeficiency 46, 616740
TMEM14C	100.0%	100.0%	100.0%	98.7%	
TMPRSS6	100.0%	100.0%	100.0%	99.0%	Iron-refractory iron deficiency anemia, 206200

TRNT1	92.0%	91.9%	100.0%	98.7%	Sideroblastic anemia with B-cell immunodeficiency, periodic fevers, and developmental delay, 616084;Retinitis pigmentosa and erythrocytic microcytosis, 616959
UROS	100.0%	100.0%	100.0%	98.2%	Porphyria, congenital erythropoietic, 263700
YARS2	100.0%	100.0%	100.0%	97.6%	Myopathy, lactic acidosis, and sideroblastic anemia 2, 613561

Gene symbols used follow HGCN guidelines: Gray KA, Yates B, Seal RL, Wright MW, Bruford EA. Nucleic Acids Res. 2015 Jan 43(Database issue):D1079-85.
TWIST X2 Covered 10x describes the percentage of a gene's coding sequence that is covered at least 10x when analyzed by WES using TWIST X2 chemistry.
TWIST X2 Covered 20x describes the percentage of a gene's coding sequence that is covered at least 20x when analyzed by WES using TWIST X2 chemistry.
srWGS GRCh38 Covered 10x describes the percentage of a gene's coding sequence that is covered at least 10x when analyzed by WGS mapped against GRCh38.
srWGS GRCh38 Covered 20x describes the percentage of a gene's coding sequence that is covered at least 20x when analyzed by WGS mapped against GRCh38.
non-protein coding genes are covered, but as coverage statistics are based on protein coding regions, statistics could not be generated.
OMIM release used for OMIM disease identifiers and descriptions : March 17th, 2023.
This list is accurate for panel version DG 4.0.0

Ad 1. "No OMIM phenotype" signifies a gene without a current OMIM association Ad 2. OMIM phenotype descriptions between {} signify risk factors