

HYPOGONADOTROPIC HYPOGONADISM GENE PANEL DG 3.4.0

(52 genes)

Releasedate: 19-04-2022

<i>Gene</i>	<i>TWIST covered >10x</i>	<i>TWIST covered >20x</i>	<i>Associated Phenotype description and OMIM disease ID</i>
ADCY3	100,0%	100,0%	No OMIM Disease ID
ANOS1	100,0%	100,0%	Hypogonadotropic hypogonadism 1 with or without anosmia (Kallmann syndrome 1), 308700
AXL	100,0%	100,0%	No OMIM Disease ID
BRCA2	100,0%	100,0%	Fanconi anemia, complementation group D1, 605724 Wilms tumor, 194070
CCDC141	100,0%	100,0%	No OMIM Disease ID
CHD7	100,0%	100,0%	Hypogonadotropic hypogonadism 5 with or without anosmia, 612370 CHARGE syndrome, 214800
CNGA2	100,0%	100,0%	No OMIM Disease ID
DCC	100,0%	100,0%	Mirror movements 1 and/or agenesis of the corpus callosum, 157600 Esophageal carcinoma, somatic, 133239 Colorectal cancer, somatic, 114500 Gaze palsy, familial horizontal, with progressive scoliosis, 2, 617542
DUSP6	100,0%	100,0%	Hypogonadotropic hypogonadism 19 with or without anosmia, 615269
ELP1	100,0%	100,0%	Dysautonomia, familial, 223900 Medulloblastoma, 155255
FEZF1	100,0%	100,0%	Hypogonadotropic hypogonadism 22, with or without anosmia, 616030
FGF17	100,0%	100,0%	Hypogonadotropic hypogonadism 20 with or without anosmia, 615270
FGF8	100,0%	100,0%	Hypogonadotropic hypogonadism 6 with or without anosmia, 612702
FGFR1	100,0%	100,0%	Pfeiffer syndrome, 101600 Hypogonadotropic hypogonadism 2 with or without anosmia, 147950 Jackson-Weiss syndrome, 123150 Hartsfield syndrome, 615465 Trigonocephaly 1, 190440 Osteoglophonic dysplasia, 166250 Encephalocraniocutaneous lipomatosis, somatic mosaic, 613001
FLRT3	100,0%	100,0%	Hypogonadotropic hypogonadism 21 with anosmia, 615271

FSHB	100,0%	100,0%	Hypogonadotropic hypogonadism 24 without anosmia, 229070
GNRH1	100,0%	100,0%	?Hypogonadotropic hypogonadism 12 with or without anosmia, 614841
GNRHR	100,0%	100,0%	Hypogonadotropic hypogonadism 7 without anosmia, 146110
GPR161	100,0%	100,0%	No OMIM Disease ID
HESX1	100,0%	100,0%	Pituitary hormone deficiency, combined, 5, 182230 Septo-optic dysplasia, 182230 Growth hormone deficiency with pituitary anomalies, 182230
HS6ST1	100,0%	100,0%	No OMIM Disease ID
IGSF10	100,0%	100,0%	No OMIM Disease ID
IL17RD	100,0%	100,0%	Hypogonadotropic hypogonadism 18 with or without anosmia, 615267
KISS1	100,0%	100,0%	?Hypogonadotropic hypogonadism 13 with or without anosmia, 614842
KISS1R	100,0%	100,0%	Hypogonadotropic hypogonadism 8 with or without anosmia, 614837 ?Precocious puberty, central, 1, 176400
KLB	100,0%	100,0%	No OMIM Disease ID
LEP	100,0%	100,0%	Obesity, morbid, due to leptin deficiency, 614962
LEPR	94,6%	94,6%	Obesity, morbid, due to leptin receptor deficiency, 614963
LHB	100,0%	100,0%	Hypogonadotropic hypogonadism 23 with or without anosmia, 228300
LHX3	100,0%	100,0%	Pituitary hormone deficiency, combined, 3, 221750
NROB1	100,0%	100,0%	Adrenal hypoplasia, congenital, 300200 46XY sex reversal 2, dosage-sensitive, 300018
NSMF	100,0%	100,0%	Hypogonadotropic hypogonadism 9 with or without anosmia, 614838
PALB2	100,0%	100,0%	Fanconi anemia, complementation group N, 610832
PCSK1	100,0%	100,0%	Obesity with impaired prohormone processing, 600955
PLXNA1	100,0%	100,0%	No OMIM Disease ID
POLG	100,0%	100,0%	Mitochondrial recessive ataxia syndrome (includes SANDO and SCAE), 607459 Mitochondrial DNA depletion syndrome 4B (MNGIE type), 613662 Mitochondrial DNA depletion syndrome 4A (Alpers type), 203700 Progressive external ophthalmoplegia, autosomal dominant 1, 157640 Progressive external ophthalmoplegia, autosomal recessive 1, 258450
PROK2	100,0%	100,0%	Hypogonadotropic hypogonadism 4 with or without anosmia, 610628
PROKR2	100,0%	100,0%	Hypogonadotropic hypogonadism 3 with or without anosmia, 244200
PROP1	100,0%	100,0%	Pituitary hormone deficiency, combined, 2, 262600
PTCH1	100,0%	100,0%	Basal cell carcinoma, somatic, 605462 Holoprosencephaly 7, 610828 Basal cell nevus syndrome, 109400
SEMA3A	100,0%	100,0%	No OMIM Disease ID
SEMA3E	100,0%	100,0%	?CHARGE syndrome, 214800

SMARCB1	100,0%	100,0%	Rhabdoid tumors, somatic, 609322 Coffin-Siris syndrome 3, 614608
SOX10	100,0%	100,0%	Waardenburg syndrome, type 4C, 613266 PCWH syndrome, 609136 Waardenburg syndrome, type 2E, with or without neurologic involvement, 611584
SOX2	100,0%	100,0%	Optic nerve hypoplasia and abnormalities of the central nervous system, 206900 Microphthalmia, syndromic 3, 206900
SPRY4	100,0%	100,0%	Hypogonadotropic hypogonadism 17 with or without anosmia, 615266
SUFU	100,0%	100,0%	Joubert syndrome 32, 617757 Medulloblastoma, desmoplastic, 155255 Basal cell nevus syndrome, 109400
TAC3	100,0%	100,0%	Hypogonadotropic hypogonadism 10 with or without anosmia, 614839
TACR3	100,0%	100,0%	Hypogonadotropic hypogonadism 11 with or without anosmia, 614840
TCF12	100,0%	100,0%	Craniosynostosis 3, 615314 Hypogonadotropic hypogonadism 26 with or without anosmia, 619718
TP53	91,7%	91,7%	Hepatocellular carcinoma, somatic, 114550 Breast cancer, somatic, 114480 Li-Fraumeni syndrome, 151623 Pancreatic cancer, somatic, 260350 Nasopharyngeal carcinoma, somatic, 607107 Bone marrow failure syndrome 5, 618165
WDR11	100,0%	100,0%	Hypogonadotropic hypogonadism 14 with or without anosmia, 614858

Gene symbols used follow HGNC guidelines: Gray KA, Yates B, Seal RL, Wright MW, Bruford EA. Nucleic Acids Res. 2015 Jan 43(Database issue):D1079-85.

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TWIST is the chemistry used for WES analysis.

Covered 10x describes the percentage of a gene's coding sequence that is covered at least 10x.

Covered 20x describes the percentage of a gene's coding sequence that is covered at least 20x.

Genes with coverage denoting NC are non-protein coding genes.

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 : April 19th , 2022.

This list is accurate for panel version DG 3.4.0

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