Human genes associated with disease / syndromes by missense mutations

In the list below, some variants are known to cause detrimental phenotypes, others are known to be neutral, or a phenotype is currently not known (neutral or deleterious). All the proteins below are known to be associated with disesase / syndromes when mutations disrupt their function. The condition is followed by the protein name; in parenthesis is the official human gene abbreviation; then the accession number for the version of the human protein to use.

Nomenclature example for AA variants: I65T – at position 65 (65^{th} amino acid in the protein), there is usually or normally an I (isoleucine); in the variant, there is a T (threonine). Note that this may or may not affect the function of the protein.

Condition/Syndrome: Protein (enzyme) name (Human gene abbreviation) – Accession No.
List of variants

Phenylketonuria: Phenylalanine hydroxylase (PAH) - P00439
D59Y
I65T
R158Q
Q160P
L255V
R261Q
M276K
P281L
P314S
Y343F
S349P
K363N
R408W

Segawa Syndrome, Dopa-responsive Dystonia, Infantile Parkinsonism: Tyrosine Hydroxylase (TH) - P07101
R233H
L236P
T276P
G280R
E284V
V306I
T314M
R337H
C359F
I440T

Tay Sachs disease: Beta-Hexosaminidase A (HEXA) - P06865
L39R
S59L
L127R
R170Q
R178H
V192A

K197T
H204R
S293I
G367D
W420C
R424H

Sandhoff disease, GM2-Gangliosidosis: Beta-hexosaminidase B (HEXB) - P07686
[clinically indistinguishable from Tay-Sachs]
W57C
R100P
H113R
T150N
K217T
Y266D
P417L
I420V
Y456S
L498F
R505Q

Ehlers-Danlos syndrome with short stature and limb anomalies (EDSSLA): 4-Beta-Galactosyltransferase (B4GALT7) - Q9UBV7
S40L
L41P
H136L
R141W
A180T
A186D
L206P
C214Y
D229N
D260G
R270C
C307Y

Gaucher disease: Acid beta-glucocerebrosidase aka Beta-glucosidase (GBA) - P04062
S281N
L353R
V391L
T408M
N409S
P415R
A423V
K480N
L483P
S504P

Severe combined immunodeficiency (aka 'Bubble boy disease'): Adenosine deaminase (ADA) - P00813
H15D
P24T
G74V
P116S
W117R
R156H
Q158H
L193P
G216R
S291L
R313W
A329V

Glycine Encephalopathy: Aminomethyltransferase (AMT) - P48728
C28W
P25T
H42R
G47R
R62P
S117L
D197H
G269D
N275K
Q317P
R320H

BH4-deficient hyperphenylalaninemia (HPA) D [HPABH4D]: Pterin carbinolamine dehydratase (PCBD1) - P61457
D14Y
N19K
K36R
H39R
F40L
L60P
H63P
D42G
T79I
C82R
A83T
S86A
R88Q
R88W
I90T
E97K

BH4-deficient hyperphenylalaninemia (HPA) C [HPABH4C]: Quinoid dihydropteridine reductase (QDPR) - P09417

G23D
W36R
K55T
A91T
W108G
H120R
A139S
Y150C
H158Y
L164V
S193P