

HSP CAUSATIVE GENES
<i>ALS2</i> (Alsin)[1,2], <i>ALDH18A1</i> (Aldehyde dehydrogenase 18 family member A1)[3,4], <i>AP4B1</i> (Adaptor Related Protein Complex 4 Subunit Beta 1)[5], <i>AP4E1</i> (Adaptor Related Protein Complex 4 Subunit Epsilon 1)[5,6], <i>AP4M1</i> (Adaptor Related Protein Complex 4 Subunit Mu 1)[7], <i>AP4S1</i> (Adaptor Related Protein Complex 4 Subunit Sigma 1)[8], <i>AP5Z1</i> (Adaptor Related Protein Complex 5 Subunit Zeta 1)[9], <i>ARL6IP1</i> (ADP Ribosylation Factor Like GTPase 6 Interacting Protein 1)[1], <i>ATAD3A</i> (ATPase Family AAA Domain Containing 3A)[10], <i>ATL1</i> (Atlastin GTPase 1)[11], <i>B4GALNT1</i> (Beta-1,4-N-Acetyl-Galactosaminyltransferase 1)[12], <i>BICD2</i> (BICD Cargo Adaptor 2)[13], <i>BSCL2</i> (BSCL2 Lipid Droplet Biogenesis Associated, Seipin)[14], <i>C19orf12</i> (Chromosome 19 Open Reading Frame 12)[15], <i>CAPN1</i> (Calpain 1)[16], <i>CCT5</i> (Chaperonin Containing TCP1 Subunit 5)[17], <i>CYP2U1</i> (Cytochrome P450 Family 2 Subfamily U Member 1)[18], <i>CYP7B1</i> (Cytochrome P450 Family 7 Subfamily B Member 1)[19], <i>DDHD1</i> (DDHD Domain Containing 1)[20,21], <i>DDHD2</i> (DDHD Domain Containing 2)[18], <i>ERLIN2</i> (ER Lipid Raft Associated 2)[22], <i>ENTPD1</i> (Ectonucleoside Triphosphate Diphosphohydrolase 1)[1], <i>EPT1</i> (Ethanalaminephosphotransferase 1)[23], <i>ERLIN1</i> (ER Lipid Raft Associated 1)[1], <i>FARS2</i> (Phenylalanyl-TRNA Synthetase 2, Mitochondrial)[24], <i>FA2H</i> (Fatty Acid 2-Hydroxylase)[25], <i>GBA2</i> (Glucosylceramidase Beta 2)[18], <i>IBA57</i> (Iron-Sulfur Cluster Assembly Factor IBA57)[26], <i>KCNA2</i> (Potassium Voltage-Gated Channel Subfamily A Member 2)[27], <i>KIAA0196/WASHC5</i> (WASH Complex Subunit 5)[28], <i>KIF1A</i> (Kinesin Family Member 1A)[29], <i>KIF5A</i> (Kinesin Family Member 5A)[30], <i>KY</i> (Kyphoscoliosis Peptidase)[31], <i>L1CAM</i> (L1 Cell Adhesion Molecule)[32,33], <i>LYST</i> (Lysosomal Trafficking Regulator)[34], <i>NIPA1</i> (NIPA Magnesium Transporter 1)[35,36], <i>NT5C2</i> (5'-Nucleotidase, Cytosolic II)[1], <i>PCYT2</i> (Phosphate Cytidyltransferase 2, Ethanalamine)[37], <i>PLP1</i> (Proteolipid Protein 1)[38], <i>PNPLA6</i> (Patatin Like Phospholipase Domain Containing 6)[39,40], <i>RAB3GAP2</i> (RAB3 GTPase Activating Non-Catalytic Protein Subunit 2)[41,42], <i>REEP1</i> (Receptor Accessory Protein 1)[43], <i>REEP2</i> (Receptor Accessory Protein 2)[44], <i>RTN2</i> (Reticulon 2)[45], <i>SETX</i> (Senataxin)[46], <i>SLC33A1</i> (Solute Carrier Family 33 Member 1)[47], <i>SPAST</i> (Spastin)[48], <i>SPG11</i> (SPG11 Vesicle Trafficking Associated, Spatacsin)[49], <i>SPG21/ACP33</i> (SPG21 Abhydrolase Domain Containing, Maspardin)[50], <i>SPG7</i> (SPG7 Matrix AAA Peptidase Subunit, Paraplegin)[51], <i>SPG80/UBAP1</i> (Ubiquitin Associated Protein 1)[52], <i>TECPR2</i> (Tectonin Beta-Propeller Repeat Containing 2)[53], <i>TFG</i> (Trafficking From ER To Golgi Regulator)[54], <i>TUBB4A</i> (Tubulin Beta 4A Class IVa)[55], <i>VCP</i> (Valosin Containing Protein)[56], <i>VPS37A</i> (VPS37A Subunit Of ESCRT-1)[57], <i>ZFYVE26</i> (Zinc Finger FYVE-Type Containing 26)[58,59], <i>ZFYVE27</i> (Zinc Finger FYVE-Type Containing 27)[60]
HSP ASSOCIATED GENES
<i>ABCD1</i> (ATP Binding Cassette Subfamily D Member 1)[61], <i>ADAR</i> (Adenosine Deaminase RNA Specific)[62], <i>ALDH3A2</i> (Aldehyde Dehydrogenase 3 Family Member A2)[63], <i>AMPD2</i> (Adenosine Monophosphate Deaminase 2)[1], <i>ARSA</i> (Arylsulfatase A)[64], <i>ARSI</i> (Arylsulfatase I precursor)[1], <i>ATP13A2</i> (ATPase Cation Transporting 13A2)[65], <i>CPT1C</i> (Carnitine Palmitoyltransferase 1C)[66], <i>ELOVL1</i> (ELOVL Fatty Acid Elongase 1)[64], <i>FLRT1</i> (Fibronectin Leucine Rich Transmembrane Protein 1)[1], <i>GALC</i> (Galactosylceramidase)[64], <i>GCH1</i> (GTP Cyclohydrolase 1)[67], <i>GRN</i> (Granulin Precursor)[68], <i>HSPD1/HSP60</i> (Heat Shock Protein Family D (Hsp60) Member 1)[69,70], <i>IFIH1</i> (Interferon Induced With Helicase C Domain 1)[62], <i>KIF1C</i> (Kinesin Family Member 1C)[71], <i>MAG</i> (Myelin-associated glycoprotein)[1,72], <i>MARS</i> (Methionyl-TRNA Synthetase)[1], <i>MFN2</i> (Mitofusin-2)[73], <i>MT-ATP6</i> (Mitochondrially Encoded ATP Synthase Membrane Subunit 6)[74], <i>OPA1</i> (Dynamin-like 120 kDa protein)[73], <i>OPA3</i> (Outer Mitochondrial Membrane Lipid Metabolism Regulator OPA3)[75], <i>PGAP1</i> (Post-GPI Attachment To Proteins Inositol Deacylase 1)[1], <i>PLA2G6</i> (Phospholipase A2 Group VI)[76], <i>POLR3A</i> (RNA Polymerase III Subunit A)[77], <i>SERAC1</i> (Serine Active Site Containing 1)[64], <i>SLC16A2</i> (Solute Carrier Family 16 Member 2)[78,79], <i>SPART</i> (Spartin)[80], <i>SPG20</i> (Troyer syndrome)[63], <i>TRPV4</i> (Transient Receptor Potential Cation Channel Subfamily V Member 4)[81], <i>USP8</i> (Ubiquitin Specific Peptidase 8)[1], <i>WDR48</i> (WD Repeat Domain 48)[1], <i>ZFR</i> (Zinc Finger RNA Binding Protein)[1]
HSP LINKED GENES
<i>C12orf65</i> (Probable peptide chain release factor C12orf65, mitochondrial)[82], <i>CYP27A1</i> (Cytochrome P450 Family 27 Subfamily A Member 1)[64], <i>GAD1</i> (Glutamate Decarboxylase 1)[83,84], <i>GJC2</i> (Gap Junction Protein Gamma 2)[85], <i>KLC2</i> (Kinesin Light Chain 2)[82], <i>RIPK5/DSTYK</i> (Dual serine/threonine and tyrosine protein kinase)[86], <i>UCHL1</i> (Ubiquitin C-Terminal Hydrolase L1)[87]
PLS GENES
<u>CAUSATIVE</u> : <i>ALS2</i> [2,88] <u>ASSOCIATED</u> : <i>ALS15</i> (Ubiquilin 2)[89], <i>C9orf72</i> (C9orf72-SMCR8 Complex Subunit)[90], <i>DCTN1</i> (Dynactin Subunit 1)[90,91], <i>ERLIN2</i> (ER Lipid Raft Associated 2)[92], <i>PARK2</i> (Parkin)[90,93] <u>LINKED</u> : <i>FIG4</i> (FIG4 Phosphoinositide 5-Phosphatase)[94], <i>SPG7</i> (SPG7 Matrix AAA Peptidase Subunit, Paraplegin)[95]

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