

GO - Biological Process

GO:0007186 : G protein-coupled receptor signaling pathway
(https://www.ebi.ac.uk/QuickGO/term/GO:0007186)
GO:0001835 : blastocyst hatching (https://www.ebi.ac.uk/QuickGO/term/GO:0001835)
GO:0051482 : positive regulation of cytosolic calcium ion concentration involved in phospholipase C-activating G protein-coupled signaling pathway
(https://www.ebi.ac.uk/QuickGO/term/GO:0051482)
GO:0035025 : positive regulation of Rho protein signal transduction
(https://www.ebi.ac.uk/QuickGO/term/GO:0035025)

GO - Cellular Component

GO:0005886 : plasma membrane (https://www.ebi.ac.uk/QuickGO/term/GO:0005886)
GO:0005887 : integral component of plasma membrane
(https://www.ebi.ac.uk/QuickGO/term/GO:0005887)
GO:0043231 : intracellular membrane-bounded organelle
(https://www.ebi.ac.uk/QuickGO/term/GO:0043231)

Yes (https://www.gephebase.org/search-criteria?/and+Presumptive Null="Yes"#gephebase-summary-title) Presumptive Null
Coding (https://www.gephebase.org/search-criteria?/and+Molecular Type="Coding"#gephebase-summary-title) Molecular Type
Deletion (https://www.gephebase.org/search-criteria?/and+Aberration Type="Deletion"#gephebase-summary-title) Aberration Type
1-9 bp Deletion Size
2 deletions - one at position 172-175 delAACT and one at position 177 delG - cause frameshift at codon 58 (PTC +31) Molecular Details of the Mutation
Association Mapping (https://www.gephebase.org/search-criteria?/and+Experimental Evidence="Association Mapping"#gephebase-summary-title) Experimental Evidence
Disruption of P2RY5, an orphan G protein-coupled receptor, underlies autosomal recessive woolly hair. (2008) (https://pubmed.ncbi.nlm.nih.gov/18297072) Main Reference
Shimomura Y; Wajid M; Ishii Y; Shapiro L; Petukhova L; Gordon D; Christiano AM Authors
The genetic determinants of hair texture in humans are largely unknown. Several human syndromes exist in which woolly hair comprises a part of the phenotype; however, simple autosomal recessive inheritance of isolated woolly hair has only rarely been reported. To identify a gene involved in controlling hair texture, we performed genetic linkage analysis in six families of Pakistani origin with autosomal recessive woolly hair (ARWH; OMIM 278150). All six families showed linkage to chromosome 13q14.2-14.3 (Z = 17.97). In all cases, we discovered pathogenic mutations in P2RY5, which encodes a G protein-coupled receptor and is a nested gene residing within intron 17 of the retinoblastoma 1 (RB1) gene. P2RY5 is expressed in both Henle's and Huxley's layers of the inner root sheath of the hair follicle. Our findings indicate that disruption of P2RY5 underlies ARWH and, more broadly, uncover a new gene involved in determining hair texture in humans. Abstract
Additional References

RELATED GEPHE

1 (Trichohyalin) (https://www.gephebase.org/search-criteria?/or+Taxon ID="9606"/and+Trait=Hair type/and+groupHaplotypes=true#gephebase-summary-title) Related Genes
4 (https://www.gephebase.org/search-criteria?/or+Gene Gephebase="P2RY5"/and+Taxon ID="9606"/or+Gene Gephebase="P2RY5"/and+Taxon ID="9606"#gephebase-summary-title) Related Haplotypes

EXTERNAL LINKS

COMMENTS