| TABLE S5. Angiogenesis-related rare diseases associated to a single gene | | | |
|---|---|---|---|
| | | | |
| **nº** | **Orpha ID** | **Rare diseases** | **Opha genes ID** |
| 1 | 3453 (C) | Autoimmune polyendocrinopathy type 1 | 119562 |
| 2 | 109 (B) | Bannayan-Riley-Ruvalcaba syndrome | 118128 |
| 3 | 136 (C) | CADASIL syndrome | 123860 |
| 4 | 137667 (A) | Capillary malformation-arteriovenous malformation | 118231 |
| 5 | 178029 (C) | Central diabetes insipidus | 118943 |
| 6 | 179 (C) | Chorioretinopathy, Birdshot type | 218376 |
| 7 | 190 (C) | Coats disease | 123713 |
| 8 | 99977 (A) | Esophageal squamous cell carcinoma | 159233 |
| 9 | 324 (B) | Fabry disease | 122153 |
| 10 | 733 (A) | Familial adenomatous polyposis | 123393 |
| 11 | 523 (A) | Familial leiomyomatosis | 121823 |
| 12 | 337 (C) | Fibrodysplasia ossificans progressiva | 117759 |
| 13 | 2092 (C) | Focal dermal hypoplasia | 159255 |
| 14 | 221126 (C) | Fowler syndrome | 227058 |
| 15 | 77260 (C) | Gaucher disease, type 2 | 122039 |
| 16 | 77261 (C) | Gaucher disease, type 3 | 122039 |
| 17 | 83454 (A) | Glomuvenous malformation | 122172 |
| 18 | 855 (C) | Hashimoto struma | 159574 |
| 19 | 90308 (B) | Klippel-Trenaunay syndrome | 119516 |
| 20 | 1571 (C) | Knobloch syndrome | 120702 |
| 21 | 65285 (A) | Lhermitte-Duclos disease | 118128 |
| 22 | 530 (C) | Lipoid proteinosis | 121261 |
| 23 | 69078 (A) | Liposarcoma | 159615 |
| 24 | 98292 (A) | Mastocytosis | 122862 |
| 25 | 1332 (A) | Medullary thyroid carcinoma | 118274 |
| 26 | 54370 (C) | Membranoproliferative glomerulonephritis | 119363 |
| 27 | 565 (C) | Menkes disease | 118879 |
| 28 | 2573 (C) | Moyamoya disease | 138831 |
| 29 | 2451 (B) | Mucocutaneous venous malformations | 120020 |
| 30 | 824 (B) | Myelofibrosis with myeloid metaplasia | 122727 |
| 31 | 99967 (A) | Myxoid liposarcoma | 159615 |
| 32 | 94058 (C) | Neovascular glaucoma | 83299 |
| 33 | 649 (C) | Norrie disease | 123713 |
| 34 | 85212 (C) | Perinatal-lethal Gaucher disease | 122039 |
| 35 | 2869 (B) | Peutz-Jeghers syndrome | 119884 |
| 36 | 729 (B) | Polycythemia vera | 122727 |
| 37 | 758 (C) | Pseudoxanthoma elasticum | 117658 |
| 38 | 90050 (C) | Retinopathy of prematurity | 123713 |
| 39 | 797 (C) | Sarcoidosis | 122427 |
| 40 | 54057 (C) | Thrombotic thrombocytopenic purpura | 117776 |
| 41 | 892 (B) | Von Hippel-Lindau disease | 120467 |
| 42 | 903 (C) | Von Willebrand disease | 120487 |
| 43 | 51636 (C) | WHIM syndrome | 120914 |
| 44 | 905 (C) | Wilson disease | 118882 |
| 45 | 913 (B) | Zollinger-Ellison syndrome | 123196 |
| (A), (B), (C) correspond to subset A, B and C respectively. | | | |