

# 第一批罕见病目录

| 序号 | 中文名称                  | 英文名称  |
|----|-----------------------|---|
| 1  | 21-羟化酶缺乏症             | 21-Hydroxylase Deficiency                                     |
| 2  | 白化病                   | Albinism  |
| 3  | Alport 综合征            | Alport Syndrome   |
| 4  | 肌萎缩侧索硬化               | Amyotrophic Lateral Sclerosis                                 |
| 5  | Angelman 氏症候群 (天使综合征) | Angelman Syndrome   |
| 6  | 精氨酸酶缺乏症               | Arginase Deficiency   |
| 7  | 热纳综合征(窒息性胸腔失养症)       | Asphyxiating Thoracic Dystrophy (Jeune Syndrome)              |
| 8  | 非典型溶血性尿毒症             | Atypical Hemolytic Uremic Syndrome                            |
| 9  | 自身免疫性脑炎               | Autoimmune Encephalitis                                       |
| 10 | 自身免疫性垂体炎              | Autoimmune Hypophysitis                                       |
| 11 | 自身免疫性胰岛素受体病           | Autoimmune Insulin Receptoropathy (Type B insulin resistance) |
| 12 | $\beta$ -酮硫解酶缺乏症      | Beta-ketothiolase Deficiency                                  |
| 13 | 生物素酶缺乏症               | Biotinidase Deficiency  |
| 14 | 心脏离子通道病               | Cardic Ion Channelopathies                                    |

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| 15 | 原发性肉碱缺乏症               | Carnitine Deficiency  |
| 16 | Castleman 病            | Castleman Disease   |
| 17 | 腓骨肌萎缩症                 | Charcot-Marie-Tooth Disease                                 |
| 18 | 瓜氨酸血症                  | Citrullinemia   |
| 19 | 先天性肾上腺发育不良             | Congenital Adrenal Hypoplasia                               |
| 20 | 先天性高胰岛素性低血糖血症          | Congenital Hyperinsulinemic Hypoglycemia                    |
| 21 | 先天性肌无力综合征              | Congenital Myasthenic Syndrome                              |
| 22 | 先天性肌强直( 非营养不良性肌强直综合征 ) | Congenital Myotonia Syndrome (Non-Dystrophic Myotonia, NDM) |
| 23 | 先天性脊柱侧弯                | Congenital Scoliosis  |
| 24 | 冠状动脉扩张病                | Coronary Artery Ectasia                                     |
| 25 | 先天性纯红细胞再生障碍性贫血         | Diamond-Blackfan Anemia                                     |
| 26 | Erdheim-Chester 病      | Erdheim-Chester Disease                                     |
| 27 | 法布雷病                   | Fabry Disease   |
| 28 | 家族性地中海热                | Familial Mediterranean Fever                                |
| 29 | 范可尼贫血                  | Fanconi Anemia  |
| 30 | 半乳糖血症                  | Galactosemia  |
| 31 | 戈谢病                    | Gaucher' s Disease  |
| 32 | 全身型重症肌无力               | Generalized Myasthenia Gravis                               |

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| 33 | Gitelman 综合征        | Gitelman Syndrome   |
| 34 | 戊二酸血症 I 型           | Glutaric Acidemia Type I  |
| 35 | 糖原累积病 ( I 型、 II 型 ) | Glycogen Storage Disease (Type I、II )   |
| 36 | 血友病                 | Hemophilia  |
| 37 | 肝豆状核变性              | Hepatolenticular Degeneration(Wilson Disease)   |
| 38 | 遗传性血管性水肿            | Hereditary Angioedema (HAE)   |
| 39 | 遗传性大疱性表皮松解症         | Hereditary Epidermolysis Bullosa  |
| 40 | 遗传性果糖不耐受症           | Hereditary Fructose Intolerance   |
| 41 | 遗传性低镁血症             | Hereditary Hypomagnesemia   |
| 42 | 遗传性多发脑梗死性痴呆         | Hereditary Multi-infarct Dementia (Cerebral Autosomal Dominant Arteriopathy with Subcortical Infarcts and Leukoencephalopathy, CADASIL) |
| 43 | 遗传性痉挛性截瘫            | Hereditary Spastic Paraplegia   |
| 44 | 全羧化酶合成酶缺乏症          | Holocarboxylase Synthetase Deficiency   |
| 45 | 同型半胱氨酸血症            | Homocysteinemia   |
| 46 | 纯合子家族性高胆固醇血         | Homozygous  |

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|    | 症                     | Hypercholesterolemia   |
| 47 | 亨廷顿舞蹈病                | Huntington Disease   |
| 48 | HHH 综合征               | Hyperornithinaemia-Hyperammon<br>aemia-Homocitrullinuria<br>Syndrome |
| 49 | 高苯丙氨酸血症               | Hyperphenylalaninemia  |
| 50 | 低碱性磷酸酶血症              | Hypophosphatasia   |
| 51 | 低磷性佝偻病                | Hypophosphatemic Rickets   |
| 52 | 特发性心肌病                | Idiopathic Cardiomyopathy  |
| 53 | 特发性低促性腺激素性性<br>腺功能减退症 | Idiopathic Hypogonadotropic<br>Hypogonadism                          |
| 54 | 特发性肺动脉高压              | Idiopathic Pulmonary Arterial<br>Hypertension                        |
| 55 | 特发性肺纤维化               | Idiopathic Pulmonary Fibrosis  |
| 56 | IgG4 相关性疾病            | IgG4 related Disease   |
| 57 | 先天性胆汁酸合成障碍            | Inborn Errors of Bile Acid Synthesis                                 |
| 58 | 异戊酸血症                 | Isovaleric Acidemia  |
| 59 | 卡尔曼综合征                | Kallmann Syndrome  |
| 60 | 朗格汉斯组织细胞增生症           | Langerhans Cell Histiocytosis  |
| 61 | 莱伦氏综合征                | Laron Syndrome   |
| 62 | Leber 遗传性视神经病变        | Leber Hereditary Optic   |

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|----|---------------------|---|
|    |                     | Neuropathy  |
| 63 | 长链 3-羟酰基辅酶 A 脱氢酶缺乏症 | Long Chain 3-hydroxyacyl-CoA Dehydrogenase Deficiency |
| 64 | 淋巴管肌瘤病              | Lymphangiomyomatosis (LAM)                            |
| 65 | 赖氨酸尿蛋白不耐受症          | Lysinuric Protein Intolerance                         |
| 66 | 溶酶体酸性脂肪酶缺乏症         | Lysosomal Acid Lipase Deficiency                      |
| 67 | 枫糖尿症                | Maple Syrup Urine Disease                             |
| 68 | 马凡综合征               | Marfan Syndrome                                       |
| 69 | McCune-Albright 综合征 | McCune-Albright Syndrome                              |
| 70 | 中链酰基辅酶 A 脱氢酶缺乏症     | Medium Chain Acyl-CoA Dehydrogenase Deficiency        |
| 71 | 甲基丙二酸血症             | Methylmalonic Acidemia                                |
| 72 | 线粒体脑肌病              | Mitochondrial Encephalomyopathy                       |
| 73 | 黏多糖贮积症              | Mucopolysaccharidosis                                 |
| 74 | 多灶性运动神经病            | Multifocal Motor Neuropathy                           |
| 75 | 多种酰基辅酶 A 脱氢酶缺乏症     | Multiple Acyl-CoA Dehydrogenase Deficiency            |
| 76 | 多发性硬化               | Multiple Sclerosis                                    |
| 77 | 多系统萎缩               | Multiple System Atrophy                               |
| 78 | 肌强直性营养不良            | Myotonic Dystrophy                                    |
| 79 | N-乙酰谷氨酸合成酶缺乏        | N-acetylglutamate Synthase                            |

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|    | 症                | Deficiency                                     |
| 80 | 新生儿糖尿病           | Neonatal Diabetes Mellitus                     |
| 81 | 视神经脊髓炎           | Neuromyelitis Optica                           |
| 82 | 尼曼匹克病            | Niemann-Pick Disease                           |
| 83 | 非综合征性耳聋          | Non-Syndromic Deafness                         |
| 84 | Noonan 综合征       | Noonan Syndrome                                |
| 85 | 鸟氨酸氨甲酰基转移酶缺乏症    | Ornithine Transcarbamylase Deficiency          |
| 86 | 成骨不全症 (脆骨病)      | Osteogenesis Imperfecta (Brittle Bone Disease) |
| 87 | 帕金森病(青年型、早发型)    | Parkinson Disease (Young-onset , Early-onset)  |
| 88 | 阵发性睡眠性血红蛋白尿      | Paroxysmal Nocturnal Hemoglobinuria            |
| 89 | 黑斑息肉综合征          | Peutz-Jeghers Syndrome                         |
| 90 | 苯丙酮尿症            | Phenylketonuria                                |
| 91 | POEMS 综合征        | POEMS Syndrome                                 |
| 92 | 卟啉病              | Porphyria                                      |
| 93 | Prader-Willi 综合征 | Prader-Willi Syndrome                          |
| 94 | 原发性联合免疫缺陷        | Primary Combined Immune Deficiency             |

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| 95  | 原发性遗传性肌张力不全             | Primary Hereditary Dystonia                            |
| 96  | 原发性轻链型淀粉样变              | Primary Light Chain Amyloidosis                        |
| 97  | 进行性家族性肝内胆汁淤积症           | Progressive Familial Intrahepatic Cholestasis          |
| 98  | 进行性肌营养不良                | Progressive Muscular Dystrophy                         |
| 99  | 丙酸血症                    | Propionic Acidemia                                     |
| 100 | 肺泡蛋白沉积症                 | Pulmonary Alveolar Proteinosis                         |
| 101 | 肺囊性纤维化                  | Pulmonary Cystic Fibrosis                              |
| 102 | 视网膜色素变性                 | Retinitis Pigmentosa                                   |
| 103 | 视网膜母细胞瘤                 | Retinoblastoma   |
| 104 | 重症先天性粒细胞缺乏症             | Severe Congenital Neutropenia                          |
| 105 | 婴儿严重肌阵挛性癫痫 (Dravet 综合征) | Severe Myoclonic Epilepsy in Infancy (Dravet Syndrome) |
| 106 | 镰刀型细胞贫血病                | Sickle Cell Disease                                    |
| 107 | Silver-Russell 综合征      | Silver-Russell Syndrome                                |
| 108 | 谷固醇血症                   | Sitosterolemia   |
| 109 | 脊髓延髓肌萎缩症( 肯尼迪病 )        | Spinal and Bulbar Muscular Atrophy (Kennedy Disease)   |
| 110 | 脊髓性肌萎缩症                 | Spinal Muscular Atrophy                                |
| 111 | 脊髓小脑性共济失调               | Spinocerebellar Ataxia                                 |
| 112 | 系统性硬化症                  | Systemic Sclerosis                                     |

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| 113 | 四氢生物蝶呤缺乏症            | Tetrahydrobiopterin Deficiency                       |
| 114 | 结节性硬化症               | Tuberous Sclerosis Complex                           |
| 115 | 原发性酪氨酸血症             | Tyrosinemia  |
| 116 | 极长链酰基辅酶 A 脱氢酶<br>缺乏症 | Very Long Chain Acyl-CoA<br>Dehydrogenase Deficiency |
| 117 | 威廉姆斯综合征              | Williams Syndrome                                    |
| 118 | 湿疹血小板减少伴免疫缺<br>陷综合征  | Wiskott-Aldrich Syndrome                             |
| 119 | X-连锁无丙种球蛋白血症         | X-linked Agammaglobulinemia                          |
| 120 | X-连锁肾上腺脑白质营养<br>不良   | X-linked Adrenoleukodystrophy                        |
| 121 | X-连锁淋巴增生症            | X-linked Lymphoproliferative<br>Disease              |

注：本目录中的 121 种罕见病不受本合同主险条款第十条责任免除中关于“遗传性疾病、先天性畸形、变形或染色体异常”的限制。