衛生福利部國民健康署

107 年 9 月罕見疾病通報個案統計表

統計期間:個案數至 107 年 10 月 16 日／死亡數至 107 年 6 月 21 日

製表時間：107 年 10 月 16 日

分類序號	疾病名稱	ICD10 碼	個案數	死亡數
A 先天性代謝異常				
◎A1 尿素循環代謝異常 Urea cycle disorders （高血氨症）				
01	Congenital urea cycle disorders(先天性尿素循環代謝障礙)	E72.20	72	21
02	Citrullinemia(瓜胺酸血症)	E72.23	78	2
03	Nitroacetylglutamate synthetase deficiency,NAG synthetase deficiency(乙醯穀胺酸合成酶缺乏症)	E72.29	0	0
04	Ornithine transcarbamylase deficiency(鳥胺酸氨甲醯基轉移酶缺乏症)	E72.4	23	2
05	Hyperornithinemia-Hyperammonemia-Homocitrullinuria Syndrome(高鳥胺酸血症-高氨血症-高瓜胺酸血症症候群)	E72.4	2	0
◎ A2 胺基酸/有機酸代謝異常 Amino acid metabolic disorders / Organic acidemias				
01	Amino acid metabolic disorders(Aminoacidopathies)(胺基酸代謝疾病)	E72.8	11	3
02	Homocystinuria(高胱胺酸血症)	E72.11	34	1
03	Hypermethioninemia(高甲硫胺酸血症)	E72.19	29	0
04	Nonketotic hyperglycinemia(非酮性高甘胺酸血症)	E72.51	26	13
05	Phenylketouria(苯酮尿症)	E70.0	276	1
06	Tetrahydrobiopterin deficiency(四氫基喋呤缺乏症)	E70.1	5	0
07	Hereditary tyrosinemia(遺傳性高酪胺酸血症)	E70.21	11	1
08	Maple syrup urine disease(楓糖尿症)	E71.0	39	10
09	Organic acidemias(有機酸血症)	E71.118	20	0
10	Isovaleric academia(異戊酸血症)	E71.110	13	2
11	Glutaric aciduria type Ⅰ、Ⅱ(戊二酸尿症，第一型、第二型)	E72.3 E71.313	79	8
12	Propionic academia(丙酸血症)	E71.121	17	4
13	Methylmalonic acidemia (甲基丙二酸血症)	E71.120	60	10
14	3-Hydroxy-3-methyl-glutaric acidemia(3-氫基-3-甲基戊二酸血症)	E71.118	6	0
15	PAH type PKU combine with Sucrase-isomaltase deficiency(典型苯酮尿症合併蔗糖酶同麥芽糖酶缺乏症)	E74.31 E70.0	2	0
16	Hyperlysinemia(高離氨基酸血症)	E72.3	0	0
17	Histidinemia(組胺酸血症)	E70.41	0	0
18	3-Methylcrotonyl-CoA carboxylase deficiency(三甲基巴豆醯	E71.19	50	0

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	輔酶 A 羧化酵素缺乏症)			
19	Multiple carboxylase deficiency(多發性羧化酶缺乏症)	D81.819	5	0
20	Hyperprolinemia(高脯胺酸血症)	E72.59	2	0
21	Aromatic L-amino acid decarboxylase deficiency(芳香族 L-胺基酸類脫羧基酶缺乏症)	E70.9	44	14
22	Tyrosine hydroxylase deficiency(酪胺酸羥化酶缺乏症)	E70.20	9	1
23	Cobalamin C defect （ Methylmalonic aciduria and Homocystinuria, cbl C type）(甲基丙二酸血症併高胱胺酸血症 (Cbl C 型))	E71.120 +E72.11	1	0
◎ A3 脂質儲積				
01	Gaucher's disease(高雪氏症)	E75.22	39	4
02	GM1/GM2 gangliosidosis(GM1/GM2 神經節苷脂儲積症)	E75.19 E75.00	6	5
03	Fabry disease(Fabry 氏症)	E75.21	313	16
04	Niemann-Pick disease(Niemann-Pick 氏症，鞘髓磷脂儲積症)	E75.240 E75.241 E75.242 E75.243 E75.248 E75.249	26	12
05	Metachromatic Leukodystrophy（MLD）(MLD 症候群)	E75.25	19	8
06	Globoid Cell Leukodystrophy (Krabbe's disease)(球細胞腦白質失養症)	E75.23	1	0
07	Infantile form Lysosomal Acid Lipase Deficiency （Wolman Disease）(嬰兒型溶酶體酸性脂肪酶缺乏症（又稱伍爾曼氏症）)	E75.5	1	0
◎A4 碳水化合物代謝異常				
01	Galactosemia(半乳糖血症)	E74.21	26	0
02	Glycogen storage disease(肝醣儲積症)	E74.09 E74.01 E74.02 E74.03 E74.09 E74.04	219	41
03	Glut（Glucose Transport）1 deficiency syndrome(腦血管屏障葡萄糖輸送缺陷)	E74.8	4	0
◎ A5 脂肪酸氧化異常				
01	Fatty acid oxidation defect(脂肪酸氧化作用缺陷)	E71.30 E71.310 E71.311 E71.312 E71.313 E71.314 E71.318 E71.32 E71.39	39	3

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02	Carnitine deficiency syndrome, primary(原發性肉鹼缺乏症)	E71.41	143	5
03	Medium-chain acyl-coenzyme A dehydrogenase deficiency（MCAD）(中鏈脂肪酸去氫酵素缺乏症)	E71.311	8	0
04	Short-chain acyl-CoA dehydrogenase deficiency(短鏈脂肪酸去氫酶缺乏症)	E71.312	5	1
◎ A6 粒線體代謝異常				
01	Mitochondrial defect(粒線體缺陷)	E88.40	304	52
02	Kearns-Sayre syndrome(Kearns-Sayre 氏症候群)	H49.811 H49.812 H49.813 H49.819	4	1
03	Leigh disease(Leigh 氏童年期腦脊髓病變)	G31.82	23	8
04	MELAS(MELAS 症候群)	E88.41	91	18
05	Mitochondrial Neurogastrointestinal Encephalopathy Syndrome(MNGIE 症候群粒線體性神經胃腸腦病變症候群)	E88.49	3	1
06	Pyruvate dehydrogenase deficiency(丙酮酸鹽脫氫酶缺乏症)	E74.4	4	0
07	Barth Syndrome(巴氏症候群)	E78.71	3	0
◎ A7 溶小體代謝異常				
01	Cystinosis(胱胺酸血症)	E72.04	5	0
02	Mucopolysaccharidoses(黏多醣症)	E76.3	174	67
03	Fucosidosis(岩藻糖代謝異常（儲積症）)	E77.1	2	1
04	Sialidosis(涎酸酵素缺乏症)	E77.1	30	4
05	Mucolipidosis(黏脂質症)	E77.1 E77.0 E75.11	9	0
06	Neuronal ceroid lipofuscinosis(神經元蠟樣脂褐質儲積症)	E75.4	5	3
07	Multiple sulfatase deficiency(多發性硫酸脂酶缺乏症)	E75.29	2	0
◎ A8 膽固醇及脂質代謝異常 Cholesterol and Lipid metabolism				
01	Homozygous familial hypercholesterolemia(同合子家族性高膽固醇血症)	E78.0	47	2
02	Familial Hyperchylomicronemia(家族性高乳糜微粒血症)	E78.3	13	0
03	Sitosterolemia(豆固醇血症 〈植物性〉)	E78.0	4	0
◎A9 礦物離子缺陷				
01	Wilson's disease(威爾森氏症)	E83.01	569	38
02	Menkes syndrome(Menkes 症候群)	E83.09	12	5
03	Molybdenum cofactor deficiency(鉬輔酶缺乏症)	E61.5	3	1
◎A10 過氧化體代謝異常				
01	Zellweger syndrome(Zellweger 氏症候群)	E71.510	3	2
02	Adrenoleukodystrophy(腎上腺腦白質失養症)	E71.511 E71.520	64	20

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		E71.521 E71.528 E71.529		
03	Rhizomelic Chondrodysplasia Punctata(肢近端型點狀軟骨發育不良)	E71.540	1	0
◎A11 其他代謝異常				
01	Porphyria(紫質症)	E80.20 E80.21 E80.29	105	3
02	Lesch-Nyhan syndrome(Lesch-Nyhan 氏症候群)	E79.1	13	3
03	Sulfite oxidase deficiency(亞硫酸鹽氧化酶缺乏)	E72.19	10	3
04	Carbohydrate-deficiency glycoprotein syndrome(碳水化合缺乏醣蛋白症候群)	E77.8	11	2
05	Trimethylaminuria(三甲基胺尿症)	E72.52	11	0
06	Congenital generalized lipodystrophy(先天性全身脂質營養不良症)	E88.1	29	7
07	Cerebrotendinous Xanthomatosis(腦腱性黃瘤症)	E75.5	6	2
08	Hypophosphatasia(低磷酸酯酶症)	E83.39 E83.31	4	0
09	Beta-Ketothiolase Deficiency(Beta 硫解酶缺乏症)	E71.19	0	0
10	Biotinidase Deficiency(生物素酶缺乏症)	D81.810	5	0
B 腦部或神經系統病變				
01	Multiple sclerosis(多發性硬化症)	G35	1,729	103
02	Amyotrophic lateral sclerosis (ALS) (肌萎縮性側索硬化症)	G12.21	1,313	557
03	Ataxia telangiectasia(共濟失調微血管擴張症候群)	G11.3	9	3
04	Huntington disease(又稱 Huntington's chorea) (亨丁頓氏舞蹈症)	G10	338	59
05	Rett syndrome(瑞特氏症候群)	F84.2	139	5
06	Spinal muscular atrophy(脊髓性肌肉萎縮症)	G12.9	458	62
07	Spinocerebellar ataxia (脊髓小腦退化性動作協調障礙)	G11.1	1,239	221
08	Tuberous sclerosis(結節性硬化症)	Q85.1	619	21
09	Congenital insensitivity to pain with anhidrosis（CIPA）(先天性痛不敏感症合併無汗症)	L74.4	10	2
10	Neurofibromatosis type Ⅱ(神經纖維瘤症候群第二型)	Q85.02	78	5
11	Alexander disease (Alexander 氏病)	E75.29	4	0
12	Stiffperson syndrome(僵體症候群)	G25.82	16	2
13	Hereditary spastic paraplegia(遺傳性痙攣性下身麻痺)	G11.4	112	2
14	Joubert syndrome(Joubert 氏症候群（家族性小腦蚓部發育不全）)	Q04.3	9	0
15	Pelizaeus-Merzbacher Disease(Pelizaeus-Merzbacher 氏症（慢性兒童型腦硬化症）)	E75.29	17	0

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16	Charcot Marie Tooth Disease(Charcot Maire Tooth 氏症(進行性神經性腓骨萎縮症))	G60.0	331	7
17	Kennedy Disease(甘迺迪氏症(脊髓延髓性肌肉萎縮症))	G12.20 G12.21 G12.22 G12.29	75	5
18	Familial Amyloidotic Polyneuropathy(家族性澱粉樣多發性神經病變)	E85.1	113	36
19	Moebius syndrome(Moebius 症候群)	Q87.0	18	6
20	Mcleod syndrome(Mcleod 症候群)	J43.0	1	1
21	Aicardi-Goutieres syndrome(Aicardi-Goutieres 症候群)	G31.89	1	0
22	Proteus Syndrome(普洛提斯症候群)	Q87.3	1	0
23	Methyl CpG binding protein 2 Duplication Syndrome（MECP2 Duplication Syndrome）(MECP2 綜合症候群)	Q99.8	4	0
24	Cerebro-Costo-Mandibular Syndrome(腦肋小頜症候群)	Q87.89	3	0
25	Dravet Syndrome (DS)(Dravet 症候群)	G40.311	26	0
26	Vanishing White Matter Disease(腦白質消失症)	G37.8	1	0
C 呼吸循環系統病變				
01	Idiopathic Infantile Arterial Calcification(特發性嬰兒動脈硬化症)	Q28.8	2	1
02	Cystic fibrosis (囊狀纖維化症)	E84.9	19	4
03	Primary Pulmonary Hypertension（PPH）(原發性肺動脈高壓)	I27.0	447	90
04	Holt-Oram Syndrome(Holt-Oram 氏症候群)	Q87.2	6	0
05	Andersen syndrome(Andersen 氏症候群（心節律障礙暨週期性麻痺症候群；鉀離子通道病變）)	E74.09	3	0
06	Hereditary Hemorrhagic Telangiectasia(遺傳性出血性血管擴張症)	I78.0	25	3
07	Asphyxiating thoracic dystrophy(窒息性胸腔失養症)	Q77.2	0	0
08	Congenital Central Hypoventilation Syndrome(先天性中樞性換氣不足症候群)	G47.35	10	0
D 消化系統病變				
01	Progressive intrahepatic cholestasis（PFIC）(進行性家族性肝內膽汁滯留症)	K83.1	18	4
02	Inborn errors of bile acid synthesis(先天性膽酸合成障礙)	E78.70	3	0
03	α1- Antitrypsin deficiency(α1-抗胰蛋白酶缺乏症)	E88.01	0	0
04	Congenital Interstitial Cell of Cajal Hyperplasia with Neuronal Intestinal Dysplasia(先天性 Cajal 氏間質細胞增生合併腸道神經元發育異常)	Q43.8	1	0
05	Alagille Syndrome(阿拉吉歐症候群)	Q44.7	12	0

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E 腎臟泌尿系統病變				
01	Lowe syndrome(Lowe 氏症候群)	E72.03	19	2
02	Bartter's syndrome(Bartter 氏症候群)	E26.81	81	0
03	Autosomal recessive polycystic kidney disease(體染色體隱性多囊性腎臟疾病)	Q61.19	6	0
F 皮膚病變				
01	Hereditary epidermolysis bullosa (遺傳性表皮分解性水泡症)	Q81.9	76	6
02	Lchthyosis, lamellar recessive(層狀魚鱗癬（自體隱性遺傳型))	Q80.2	21	2
03	Collodion baby(膠膜兒)	Q80.2	1	0
04	Harlequin ichthyosis(斑色魚鱗癬)	Q80.4	2	1
05	Bullous Congenital ichthyosiform erythoderma（epidermolytic hyperkeratosis）(水泡型先天性魚鱗癬樣紅皮症（表皮鬆解性角化過度症）)	Q80.3	26	0
06	Ectodermal Dysplasias(外胚層增生不良症)	Q82.4	90	0
07	Meleda disease(Meleda 島病)	Q82.8	8	0
08	Darier's disease(Darier 氏症（毛囊角化病）)	Q82.8	28	1
09	Dyskeratosis Congenita(先天性角化不全症)	Q82.8	1	0
10	Diffuse Non-epidermolytic Palmoplantar Keratoderma type Unna-Thost(皮膚過度角化症雅司病)	Q82.8	10	0
11	Incontinentia Pigmenti(色素失調症)	Q82.3	49	0
12	Netherton Syndrome(Netherton 症候群)	Q80.3	6	1
G 肌肉病變				
01	Duchenne muscular dystrophy(裘馨氏肌肉失養症)	G71.0	353	59
02	Nemaline Rod Myopathy(Nemaline 線狀肌肉病變)	G71.2	27	2
03	Schwartz Jampel syndrome(Schwartz Jampel 氏症候群)	G71.13	1	0
04	Myotonic dystrophy(肌肉強直症)	G71.11	198	34
05	Facioscapulohumeral muscular dystrophy(面肩胛肱肌失養症)	G71.0	127	5
06	Myotubular Myopathy(肌小管病變)	G71.2	16	3
07	Becker Muscular Dystrophy(貝克型肌肉失養症)	G71.0	50	5
08	Freeman-Sheldon syndrome(Freeman-Sheldon 氏症候群)	Q87.0	1	0
09	Limb-girdle muscular dystrophy(肢帶型肌失養症)	G71.0	55	0
10	Congenital Muscular Dystrophy(先天性肌失養症)	G71.0	17	0
11	Central Core Disease(中心軸空肌病)	G71.2	2	0
12	Multiminicore Disease(多微小軸空肌病)	G71.2	1	0
13	Emery–Dreifuss Muscular Dystrophy (EDMD)(Emery–Dreifuss 肌失養症)	G71.0	2	0
H 骨及軟骨病變				

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01	Achondroplasia(軟骨發育不全症)	Q77.4	392	7
02	Osteogenesis imperfecta(成骨不全症)	Q78.0 M88.0 M88.1 M88.811 M88.812 M88.819 M88.821 M88.822 M88.829 M88.831 M88.832 M88.839 M88.841 M88.842 M88.849 M88.851 M88.852 M88.859 M88.861 M88.862 M88.869 M88.871 M88.872 M88.879 M88.88 M88.89 M88.9	348	9
03	Primary Paget disease(原發性變形性骨炎)		3	0
04	Cleidocraninal dysplasia(鎖骨顱骨發育異常)	Q74.0	40	0
05	Fibrodysplasia Ossificans Progressiva(進行性骨化性肌炎)	M61.10 M61.111 M61.112 M61.119 M61.121 M61.122 M61.129 M61.131 M61.132 M61.139 M61.141 M61.142 M61.143 M61.144 M61.145 M61.146 M61.151 M61.152 M61.159 M61.161 M61.162 M61.169 M61.171 M61.172 M61.173 M61.174 M61.175	11	0

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06	Split-hand/ Split-foot malformation（SHFM）(裂手裂足症)	M61.176 M61.177 M61.178 M61.179 M61.18 M61.19 Q71.60 Q71.61 Q71.62 Q71.63 Q72.70 Q72.71 Q72.72 Q72.73	13	0
07	Osteopetrosis(骨質石化症)	Q78.2	22	7
08	Pseudoachondroplastic dysplasia(假性軟骨發育不全)	Q77.8	13	0
09	Multiple Epiphyseal Dysplasia(多發性骨骺發育不全症)	Q78.3	10	0
I 結締組織病變				
01	Ehlers Danlos syndromeⅣ(先天結締組織異常第四型)	Q79.6	19	0
J 血液疾病				
01	Thalassemia major(重型海洋性貧血)	D56.0 D56.1	383	12
02	Thrombasthenia(血小板無力症)	D69.1	30	1
03	Homozygous proetin C deficiency(同基因合子蛋白質 C 缺乏症)	D68.59	6	0
04	Paroxysmal Nocturnal Hemoglobinuria(陣發性夜間血紅素尿症)	D59.5	89	5
05	Atypical Hemolytic Uremic Syndrome(非典型性尿毒溶血症候群)	D59.3	18	3
K 免疫疾病				
01	Chronic primary granulomatous disease(原發性慢性肉芽腫病)	D71	21	1
02	Congenital Hyper IgE syndrome(先天性高免疫球蛋白 E 症候群)	D82.4	14	4
03	Bruton’s agammaglobulinemia(布魯頓氏低免疫球蛋白血症)	D80.0	20	2
04	Wiskott- Aldrich Syndrome(Wiskott- Aldrich 氏症候群)	D82.0	22	7
05	Severe combined immunodeficiency(嚴重複合型免疫缺乏症)	D81.0 D81.1 D81.2 D81.9	20	2
06	Complement Component 8 deficiency(補體成份 8 缺乏症)	D84.1	2	0
07	IPEX Syndrome(IPEX 症候群)	E31.0	1	1
08	Hyper-IgM syndrome(高免疫球蛋白 M 症候群)	D80.5	7	1
09	Interferon γ receptor 1 deficiency (γ 干擾素受體 1 缺陷)	D84.8	3	0
10	Hereditary Angioedema (HAE)(遺傳性血管性水腫)	D84.1	8	0

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L 內分泌疾病				
01	Kenny-Caffey syndrome(Kenny-Caffey 氏症候群)	Q87.1	1	0
02	Pseudohypoparathyroidism(假性副甲狀腺低能症)	E20.1	63	0
03	X-linked hypophosphatemic rickets(性連遺傳型低磷酸鹽佝僂症)	E83.31	123	2
04	Laron syndrome（Laron Dwarfism）(Laron 氏侏儒症候群)	E34.3	5	0
05	Bardet-Biedl syndrome(Bardet-Biedl 氏症候群)	Q87.89	37	3
06	Alsrtom Syndrome(Alstrom 氏症候群)	Q87.89	19	3
07	Persistent hyperinsulinemic hypoglycemia of infancy (PHHI)(持續性幼兒型胰島素過度分泌低血糖症)	E16.1	72	5
08	Wolfram syndrome(Wolfram 氏症候群)	E88.9	10	4
09	McCune Albright syndrome(McCune Albright 氏症候群)	Q78.1	22	1
10	Campomelic dysplasia with autosomal sex reversal(短指發育不良及性別顛倒)	Q99.8	1	1
11	ACTH resistance(腎上腺皮促素抗性)	E27.49	1	0
12	25-Hydroxyvitamin D 1-Alpha-Hydroxylase Deficiency(第一型遺傳性維生素 D 依賴型佝僂症)	E83.32	15	0
13	Congenital adrenal hypoplasia(先天性腎上腺發育不全)	Q89.1	19	0
14	Kallmann syndrome(Kallmann 氏症候群)	E23.0	49	1
15	Permanent Neonatal Diabetes Mellitus(永久性新生兒糖尿病)	P70.2	1	0
M 先天畸形症候群				
01	Aarskog-Scott syndrome(Aarskog-Scott 氏症候群)	Q87.1	11	0
02	Waardenburg syndrome (瓦登伯格氏症候群)	E70.8	78	2
03	Apert syndrome(愛伯特氏症)	Q87.0	30	1
04	Smith-Lemli-Opitz syndrome(Smith-Lemli-Opitz 氏症候群)	E78.72	5	3
05	Larsen syndrome(Larsen 氏症候群（顎裂-先天性脫位症候群）)	Q74.8	2	1
06	Beckwith Wiedemann syndrome(Beckwith Wiedemann 氏症候群)	Q87.3	60	1
07	Crouzon syndrome(Crouzon 氏症候群)	Q75.1	67	4
08	Fraser syndrome(Fraser 氏症候群)	Q87.0	1	0
09	Multiple pterygium syndrome(多發性翼狀膜症候群)	Q79.8	10	1
10	Cornelia de Lange syndrome(Cornelia de Lange 氏症候群)	Q87.1	60	5
11	Hallerman-Streiff Syndrome(海勒曼-史德萊夫氏症候群)	Q87.0	6	0
12	Kabuki syndrome(歌舞伎症候群)	Q89.8	54	1
13	Oto-Palato-Digital syndrome(耳-齶-指（趾）症候群)	Q87.0	3	0
14	Conradi-Hunermann syndrome(Conradi-Hunermann 氏症候群)	Q77.3	2	0

分類序號	疾病名稱	ICD10 碼	個案數	死亡數
15	Treacher Collins Syndrome(Treacher Collins 氏症候群)	Q75.4	24	0
16	Robinow Syndrome(Robinow 氏症候群)	Q87.1	5	1
17	Pfeiffer syndrome(Pfeiffer 氏症候群)	Q87.0	7	4
18	Pantothenate Kinase Associated Neurodegeneration（PKAN）(泛酸鹽激酶關聯之神經退化性疾病)	G23.0	12	2
19	Nail-Patella Syndrome(指（趾）甲髕骨症候群)	Q87.2	4	0
20	Cardiofaciocutaneous Syndrome(CFC 症候群)	Q87.89	6	0
21	Peters-Plus syndrome(Peters-Plus 症候群)	Q13.4	2	0
22	Nager Syndrome(Nager 症候群)	Q75.4	2	0
23	CHARGE Syndrome(CHARGE 症候群)	Q89.8	8	0
24	White-Sutton syndrome(懷特-薩頓症候群)	Q99.8 F84.8 F78.0	1	0
N 染色體異常				
01	Angelman syndrome(Angelman 氏症候群)	Q93.5	68	1
02	DiGeorge's syndrome(DiGeorge's 症候群)	D82.1	193	13
03	Prader-Willi syndrome(Prader-Willi 氏症候群)	Q87.1	313	26
04	W A G R syndrome（Wilms' tumor-Aniridia-Genitourinary Anomalies-mental Retardation）(威爾姆氏腫瘤、無虹膜、性器異常、智能障礙症候群（ＷＡＧＲ症候群）)	Q87.89	10	0
05	Miller Dieker syndrome(Miller Dieker 症候群)	Q93.88	17	4
06	Rubinstein-Taybi syndrome(Rubinstein-Taybi 氏症候群)	Q87.2	37	0
07	Williams Syndrome(威廉斯氏症候群)	Q93.89	257	3
08	Von Hippel-Lindau disease(Von Hippel-Lindau 症候群)	Q85.8	20	2
09	Branchio-Oto-Renal Syndrome （ BOR Syndrome ）(Branchio-Oto-Renal 症候群（BOR 症候群）)	Q87.89	3	0
Z 其他未分類或不明原因				
01	Cockayne syndrome(Cockayne 氏症候群)	Q87.1	28	19
02	Hutchinson Gilford progeria syndrome(早老症)	E34.8	7	4
03	Tricho-hepato-enteric syndrome(髮-肝-腸症候群)	Q87.9	2	1
04	Stargardt's disease(Stargardt 's 氏症)	H35.50	0	0
05	Occult Macular Dystrophy；OMD(隱匿性黃斑部失養症)	H35.50	2	0