Appendix 1 – Primary Kidney Disease Codes

1003	Adult nephrotic syndrome - no histology
1019	Nephrotic syndrome of childhood - steroid sensitive - no histology
1026	Congenital nephrotic syndrome (CNS) - no histology
1035	Congenital nephrotic syndrome (CNS) - Finnish type - no histology
1042	Congenital nephrotic syndrome (CNS) - Finnish type - histologically proven
1057	Congenital nephrotic syndrome (CNS) - diffuse mesangial sclerosis
1061	Congenital nephrotic syndrome (CNS) - focal segmental glomerulosclerosis (FSGS)
1074	Denys-Drash syndrome
1088	Congenital nephrotic syndrome (CNS) - congenital infection
1090	Minimal change nephropathy - no histology
1100	Minimal change nephropathy - histologically proven
1116	IgA nephropathy - no histology
1128	IgA nephropathy - histologically proven
1137	Familial IgA nephropathy - no histology
1144	Familial IgA nephropathy - histologically proven
1159	IgA nephropathy secondary to liver cirrhosis - no histology
1163	IgA nephropathy secondary to liver cirrhosis - histologically proven
1171	IgM - associated nephropathy
1185	Membranous nephropathy - idiopathic
1192	Membranous nephropathy - malignancy associated
1205	Membranous nephropathy - drug induced
1214	Membranous nephropathy - infection associated
1222	Mesangiocapillary glomerulonephritis type 1
1233	Mesangiocapillary glomerulonephritis type 2 (dense deposit disease)
1246	Mesangiocapillary glomerulonephritis type 3
1251	Idiopathic rapidly progressive (crescentic) glomerulonephritis
1267	Primary focal segmental glomerulosclerosis (FSGS)
1279	Familial focal segmental glomerulosclerosis (FSGS) - autosomal recessive - no histology
1280	Familial focal segmental glomerulosclerosis (FSGS) - autosomal recessive - histologically proven
1298	Familial focal segmental glomerulosclerosis (FSGS) - autosomal dominant - no histology
1308	Familial focal segmental glomerulosclerosis (FSGS) - autosomal dominant - histologically proven
1312	Focal segmental glomerulosclerosis (FSGS) secondary to obesity - no histology
1320	Focal segmental glomerulosclerosis (FSGS) secondary to obesity - histologically proven
1331	Diffuse endocapillary glomerulonephritis
1349	Mesangial proliferative glomerulonephritis
1354	Focal and segmental proliferative glomerulonephritis
1365	Glomerulonephritis - secondary to systemic disease
1377	Glomerulonephritis - histologically indeterminate
1383	Systemic vasculitis - ANCA negative - histologically proven
1396	Systemic vasculitis - ANCA positive - no histology
1401	Granulomatosis with polyangiitis - no histology
1417	Granulomatosis with polyangiitis - histologically proven
1429	Microscopic polyangiitis - histologically proven
1438	Churg-Strauss syndrome - no histology
1440	Churg-Strauss syndrome - histologically proven
1455	Polyarteritis nodosa
1464	Anti-Glomerular basement membrane (GBM) disease / Goodpasture's syndrome - no histology
1472	Anti-Glomerular basement membrane (GBM) disease / Goodpasture's syndrome - histologically proven
1486	Systemic lupus erythematosus / nephritis - no histology
1493	Systemic lupus erythematosus / nephritis - histologically proven
1504	Henoch-Schönlein purpura / nephritis - no histology
1515	Henoch-Schönlein purpura / nephritis - histologically proven
1527	Renal scleroderma / systemic sclerosis - no histology
1536	Renal scleroderma / systemic sclerosis - histologically proven
1543	Essential mixed cryoglobulinaemia - no histology
1558	Essential mixed cryoglobulinaemia - histologically proven
1562	Cryoglobulinaemia secondary to hepatitis C - no histology
1570	Cryoglobulinaemia secondary to hepatitis C - histologically proven
1589	Cryoglobulinaemia secondary to systemic disease - no histology
1591	Cryoglobulinaemia secondary to systemic disease - histologically proven
1602	Primary reflux nephropathy - sporadic
1618	Familial reflux nephropathy
1625	Congenital dysplasia / hypoplasia
1639	Multicystic dysplastic kidneys
1641	Renal dysplasia due to fetal ACE-inhibitor exposure
1656	Glomerulocystic disease
1660	Congenital pelvi-ureteric junction obstruction
1673	Congenital vesico-ureteric junction obstruction
1687	Posterior urethral valves
1694	Syndrome of agenesis of abdominal muscles - prune belly syndrome
1706	Congenital neurogenic bladder
1710	Bladder exstrophy
1723	Megacystis-megaureter
1734	Oligomeganephronia
1747	Renal papillary necrosis
1752	Acquired obstructive uropathy / nephropathy
1768	Acquired obstructive nephropathy due to neurogenic bladder
1775	Obstructive nephropathy due to prostatic hypertrophy
1781	Obstructive nephropathy due to prostate cancer
1799	Obstructive nephropathy due to bladder cancer
1809	Obstructive nephropathy due to other malignancies
1813	Idiopathic retroperitoneal fibrosis
1821	Retroperitoneal fibrosis secondary to malignancies
1832	Calculus nephropathy / urolithiasis
1845	Calcium oxalate urolithiasis
1850	Enteric hyperoxaluria
1866	Magnesium ammonium phosphate (struvite) urolithiasis
1878	Uric acid urolithiasis
1884	Tubulointerstitial nephritis - no histology
1897	Tubulointerstitial nephritis - histologically proven
1907	Familial interstitial nephropathy - no histology
1911	Familial interstitial nephropathy - histologically proven
1924	Tubulointerstitial nephritis associated with autoimmune disease - no histology
1930	Tubulointerstitial nephritis associated with autoimmune disease - histologically proven
1948	Tubulointerstitial nephritis with uveitis (TINU) - no histology
1953	Tubulointerstitial nephritis with uveitis (TINU) - histologically proven
1969	Renal sarcoidosis - no histology
1976	Renal sarcoidosis - histologically proven
1982	Aristolochic acid nephropathy (Balkan / Chinese herb / endemic nephropathy) - no histology
1995	Aristolochic acid nephropathy (Balkan / Chinese herb / endemic nephropathy) - histologically proven
2005	Drug-induced tubulointerstitial nephritis - no histology
2014	Drug-induced tubulointerstitial nephritis - histologically proven
2022	Nephropathy due to analgesic drugs - no histology
2033	Nephropathy due to analgesic drugs - histologically proven
2046	Nephropathy due to ciclosporin - no histology
2051	Nephropathy due to ciclosporin - histologically proven
2067	Nephropathy due to tacrolimus - no histology
2079	Nephropathy due to tacrolimus - histologically proven
2080	Nephropathy due to aminoglycosides - no histology
2098	Nephropathy due to aminoglycosides - histologically proven
2108	Nephropathy due to amphotericin - no histology
2112	Nephropathy due to amphotericin - histologically proven
2120	Nephropathy due to cisplatin - no histology
2131	Nephropathy due to cisplatin - histologically proven
2149	Nephropathy due to lithium - no histology
2154	Nephropathy due to lithium - histologically proven
2165	Lead induced nephropathy - no histology
2177	Lead induced nephropathy - histologically proven
2183	Acute urate nephropathy - no histology
2196	Acute urate nephropathy - histologically proven
2203	Chronic urate nephropathy - histologically proven
2219	Radiation nephritis
2226	Renal / perinephric abscess
2235	Renal tuberculosis
2242	Leptospirosis
2257	Hantavirus nephropathy
2261	Xanthogranulomatous pyelonephritis
2274	Nephropathy related to HIV - no histology
2288	Nephropathy related to HIV - histologically proven
2290	Schistosomiasis
2300	Other specific infection
2316	Diabetic nephropathy in type I diabetes - no histology
2328	Diabetic nephropathy in type I diabetes - histologically proven
2337	Diabetic nephropathy in type II diabetes - no histology
2344	Diabetic nephropathy in type II diabetes - histologically proven
2359	Chronic hypertensive nephropathy - no histology
2363	Chronic hypertensive nephropathy - histologically proven
2371	Malignant hypertensive nephropathy / accelerated hypertensive nephropathy - no histology
2385	Malignant hypertensive nephropathy / accelerated hypertensive nephropathy - histologically proven
2392	Ageing kidney - no histology
2407	Ischaemic nephropathy - no histology
2411	Ischaemic nephropathy / microvascular disease - histologically proven
2424	Renal artery stenosis

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Appendix 1 – Primary Kidney Disease Codes

2430	Atheroembolic renal disease - no histology
2448	Atheroembolic renal disease - histologically proven
2453	Fibromuscular dysplasia of renal artery
2469	Renal arterial thrombosis / occlusion
2476	Renal vein thrombosis
2482	Cardiorenal syndrome
2495	Hepatorenal syndrome
2509	Renal amyloidosis
2513	AA amyloid secondary to chronic inflammation
2521	AL amyloid secondary to plasma cell dyscrasia
2532	Familial amyloid secondary to protein mutations - no histology
2545	Familial amyloid secondary to protein mutations - histologically proven
2550	Familial AA amyloid secondary to familial Mediterranean fever / TRAPS (Hibernian fever) - no histology
2566	Familial AA amyloid secondary to familial Mediterranean fever / TRAPS (Hibernian fever) - histologically proven
2578	Myeloma kidney - no histology
2584	Myeloma cast nephropathy - histologically proven
2597	Light chain deposition disease
2606	Immunotactoid / fibrillary nephropathy
2610	Haemolytic uraemic syndrome (HUS) - diarrhoea associated
2623	Atypical haemolytic uraemic syndrome (HUS) - diarrhoea negative
2634	Thrombotic thrombocytopenic purpura (TTP)
2647	Haemolytic uraemic syndrome (HUS) secondary to systemic disease
2652	Congenital haemolytic uraemic syndrome (HUS)
2668	Familial haemolytic uraemic syndrome (HUS)
2675	Familial thrombotic thrombocytopenic purpura (TTP)
2681	Nephropathy due to eclampsia
2699	Sickle cell nephropathy - no histology
2702	Sickle cell nephropathy - histologically proven
2718	Autosomal dominant (AD) polycystic kidney disease
2725	Autosomal dominant (AD) polycystic kidney disease type I
2739	Autosomal dominant (AD) polycystic kidney disease type II
2741	Autosomal recessive (AR) polycystic kidney disease
2756	Alport syndrome - no histology
2760	Alport syndrome - histologically proven
2773	Benign familial haematuria
2787	Thin basement membrane disease
2794	Cystic kidney disease
2804	Medullary cystic kidney disease type I
2815	Medullary cystic kidney disease type II
2827	Uromodulin-associated nephropathy (familial juvenile hyperuricaemic nephropathy)
2836	Nephronophthisis
2843	Nephronophthisis - type 1 (juvenile type)
2858	Nephronophthisis - type 2 (infantile type)
2862	Nephronophthisis - type 3 (adolescent type)
2870	Nephronophthisis - type 4 (juvenile type)
2889	Nephronophthisis - type 5
2891	Nephronophthisis - type 6
2901	Primary Fanconi syndrome
2917	Tubular disorder as part of inherited metabolic diseases
2929	Dent disease
2938	Lowe syndrome (oculocerebrorenal syndrome)
2940	Inherited aminoaciduria
2955	Cystinuria
2964	Cystinosis
2972	Inherited renal glycosuria
2986	Hypophosphataemic rickets X-linked (XL)
2993	Hypophosphataemic rickets autosomal recessive (AR)
3000	Primary renal tubular acidosis (RTA)
3016	Proximal renal tubular acidosis (RTA) - type II
3028	Distal renal tubular acidosis (RTA) - type I
3037	Distal renal tubular acidosis with sensorineural deafness - gene mutations
3044	Nephrogenic diabetes insipidus
3059	Lesch Nyhan syndrome - hypoxanthine guanine phosphoribosyl transferase deficiency
3063	Phosphoribosyl pyrophosphate synthetase (PRPPS) superactivity
3071	Alagille syndrome
3085	Bartter syndrome
3092	Gitelman syndrome
3102	Liddle syndrome
3118	Apparent mineralocorticoid excess
3125	Glucocorticoid suppressible hyperaldosteronism
3139	Inherited / genetic diabetes mellitus type II
3141	Pseudohypoaldosteronism type 1
3156	Pseudohypoaldosteronism type 2 (Gordon syndrome)
3160	Familial hypocalciuric hypercalcaemia
3173	Familial hypercalciuric hypocalcaemia
3187	Familial hypomagnesaemia
3194	Primary hyperoxaluria
3207	Primary hyperoxaluria type I
3211	Primary hyperoxaluria type II
3224	Fabry disease - no histology
3230	Fabry disease - histologically proven
3248	Xanthinuria
3253	Nail-patella syndrome
3269	Rubinstein-Taybi syndrome
3276	Tuberous sclerosis
3282	Von Hippel-Lindau disease
3295	Medullary sponge kidneys
3305	Horse-shoe kidney
3314	Frasier syndrome
3322	Branchio-oto-renal syndrome
3333	Williams syndrome
3346	Townes-Brocks syndrome
3351	Lawrence-Moon-Biedl / Bardet-Biedl syndrome
3367	Mitochondrial cytopathy
3379	Familial nephropathy
3380	Acute kidney injury
3398	Acute kidney injury due to hypovolaemia
3403	Acute kidney injury due to circulatory failure
3419	Acute kidney injury due to sepsis
3426	Acute kidney injury due to rhabdomyolysis
3435	Acute kidney injury due to nephrotoxicity
3442	Acute cortical necrosis
3457	Acute pyelonephritis
3461	Kidney tumour
3474	Renal cell carcinoma - histologically proven
3488	Transitional cell carcinoma - histologically proven
3490	Wilms tumour - histologically proven
3501	Mesoblastic nephroma - histologically proven
3517	Single kidney identified in adulthood
3529	Chronic kidney disease (CKD) / chronic renal failure (CRF) caused by tumour nephrectomy
3538	Chronic kidney disease (CKD) / chronic renal failure (CRF) due to traumatic loss of kidney
3540	Chronic kidney disease (CKD) / chronic renal failure (CRF) due to donor nephrectomy
3555	Chronic kidney disease (CKD) / chronic renal failure (CRF) - aetiology uncertain / unknown - no histology
3564	Chronic kidney disease (CKD) / chronic renal failure (CRF) - aetiology uncertain / unknown - histologically proven
3572	Haematuria and proteinuria - no histology
3604	Nephrotic syndrome of childhood - steroid resistant - no histology
3615	Nephrotic syndrome of childhood - no trial of steroids - no histology
3627	Renal cysts and diabetes syndrome
3636	Chronic urate nephropathy - no histology
3643	Chronic renal failure due to systemic infection
3658	Renal coloboma syndrome
3662	Hypercalcaemic nephropathy
3670	Retroperitoneal fibrosis secondary to peri-aortitis
3689	Retroperitoneal fibrosis secondary to drugs
3691	Renal failure
3708	Chronic renal failure
3712	Isolated haematuria - no histology
3720	Isolated proteinuria - no histology
3731	Primary hyperoxaluria type III
3749	Glomerulonephritis - no histology
3754	Focal segmental glomerulosclerosis (FSGS) secondary to HIV
3765	Focal segmental glomerulosclerosis (FSGS) secondary to lithium
3777	Focal segmental glomerulosclerosis (FSGS) secondary to sickle cell
3783	Renal papillary necrosis caused by diabetes
3796	Renal papillary necrosis caused by analgesics
3806	Renal papillary necrosis caused by sickle cell
3810	Kidney stones due to ARPT deficiency
3823	Infiltration by lymphoma - histologically proven
3834	Nephropathy due to pre-eclampsia
3847	Systemic vasculitis - ANCA negative - no histology
3852	Systemic vasculitis - ANCA positive - histologically proven

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