Appendix 1 ERA-EDTA Primary Renal Diagnosis Codes and Groupings

Group 1 Primary Glomerulonephritis

10 Glomerulonephritis; histologically NOT examined
11 Focal segmental glomerulosclerosis with nephrotic syndrome in children
12 IgA nephropathy (proven by immunofluorescence, not 85)
13 Dense deposit disease; membranoproliferative GN; type II (proven by immunofluorescence and/or electron microscopy)
14 Membranous nephropathy
15 Membranoproliferative GN; type I (proven by immunofluorescence and/or electron microscopy-not code 84 or 89)
16 Crescentic (extra-capillary) glomerulonephritis (type I, II, III)
17 Focal segmental glomerulosclerosis with nephrotic syndrome in adults
19 Glomerulonephritis; histologically examined, not given above

Group 2 Interstitial Nephropathies

20 Pyelonephritis cause not specified
21 Pyelonephritis associated with neurogenic bladder
22 Pyelonephritis due to congenital obstructive uropathy with/without vesico-ureteric reflux
23 Pyelonephritis due to acquired obstructive uropathy
24 Pyelonephritis due to vesico-ureteric reflux without obstruction
25 Pyelonephritis due to urolithiasis
29 Pyelonephritis due to other cause
30 Interstitial nephritis (not pyelonephritis) due to other cause, or unspecified (not mentioned below)
31 Interstitial nephropathy due to analgesic drugs
32 Interstitial nephropathy due to cis-platinum
33 Interstitial nephropathy due to cyclosporin A
34 Lead induced interstitial nephropathy
39 Drug induced interstitial nephropathy not mentioned above
40 Cystic kidney disease-type unspecified
41 Polycystic kidneys; adult type (dominant)
42 Polycystic kidneys; infantile (recessive)
43 Medullary cystic disease; including nephronophthisis
49 Cystic kidney– disease-other specified type
50 Hereditary/Familial nephropathy-type unspecified
51 Hereditary nephritis with nerve deafness (Alport's Syndrome)
52 Cystinosis
53 Primary oxalosis
54 Fabry's disease
59 Hereditary nephropathy-other specified type
61 Oligomeganephronic hypoplasia
63 Congenital renal dysplasia with/without urinary tract malformation
66 Syndrome of agenesis of abdominal muscles (Prune Belly)
92 Gout nephropathy (urate)
93 Nephrocalcinosis and hypercalcaemic nephropathy

Group 3 Multisystem Diseases

70 Renal vascular disease-type unspecified
71 Renal vascular disease due to malignant hypertension (No PRD)
72 Renal vascular disease due to hypertension (No PRD)
73 Renal vascular disease due to polyarteritis
74 Wegeners Granulomatosis
75 Ischaemic renal disease / cholesterol embolisation
76 Glomerulonephritis related to liver cirrhosis
78 Cryoglobulinaemic glomerulonephritis
79 Renal vascular disease-due to other cause (not given above and not code 84-88)
82 Myelomatosis/light chain deposit disease
83 Amyloid
84 Lupus erythematosus
85 Henoch-Schonlein purpura
86 Goodpasture's Syndrome
87 Systemic sclerosis (scleroderma)
88 Haemolytic uraemic Syndrome (including Moschcowitz Syndrome)
89 Multi-system disease-other (not mentioned above)
90 Tubular necrosis (irreversible) or cortical necrosis (different from 88)
91 Tuberculosis
94 Balkan nephropathy
95 Kidney tumour
96 Traumatic or surgical loss of kidney

Group 4 Diabetes

80 Diabetic glomerulosclerosis or diabetic nephropathy

Group 5 Not Known and Other

00 Chronic renal failure; aetiology uncertain/unknown/unavailable
60 Renal hypoplasia (congenital)-type unspecified
99 Other identified renal disorders