Nefrovisie
Richtlijnen · Registratie · Visitatie

ERA-EDTA PRD code	ERA-EDTA Primaire diagnose
	Unknown
00	Chronic renal failure; aetiology uncertain
	Glomerulonephritis
10	Glomerulonephritis; histologically NOT examined
11	Focal segmental glomerulosclerosis with nephrotic syndrome in children
12	IgA nephropathy (proven by immunofluorescence, not code 76 or 85)
13	Dense deposit disease; membrano-proliferative GN; type II (proven by immunofluorescence / electron microscopy)
14	Membranous nephropathy
15	Membrano-proliferative GN; type I (proven by immunofluorescence / electron microscopy, not code 84 or 89)
16	Crescentic (extracapillary) glomerulonephritis (type I, II, III)
17	Focal segmental glomerulosclerosis with nephrotic syndrome in adults
19	Glomerulonephritis; histologically examined
	Interstitial nephritis, including pyelonephritis, drug induced nephropathy and urolithiasis
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
31	Nephropathy (interstitial) due to analgesic drugs
32	Nephropathy (interstitial) due to cis-platinum
33	Nephropathy (interstitial) due to cyclosporin A
34	Lead induced nephropathy (interstitial)
39	Drug induced nephropathy (interstitial)
	Cystic kidney diseases
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
	Other congenital and hereditary kidney diseases
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
60	Renal hypoplasia (congenital) - type unspecified
61	Oligomeganephronic hypoplasia
63	Congenital renal dysplasia with or without urinary tract malformation
66	Syndrome of agenesis of abdominal muscles (Prune Belly)
	Renal vascular disease, excluding vasculitis
70	Renal vascular disease - type unspecified
71	Renal vascular disease due to malignant hypertension
72	Renal vascular disease due to hypertension
79	Renal vascular disease - due to other cause (not code 84-88)
	Diabetes Mellitus
80	Diabetes glomerulosclerosis or diabetic nephropathy - Type I
81	Diabetes glomerulosclerosis or diabetic nephropathy - Type II
	Other multisystem diseases
73	Renal vascular disease due to polyarteritis
74	Wegener's granulomatosis
82	Myelomatosis / light chain deposit disease
83	Amyloid

84	Lupus erythematosus
85	Henoch-Schoenlein purpura
86	Goodpasture's Syndrome
87	Systemic sclerosis (scleroderma)
88	Haemolytic Uraemic Syndrome (including Moschcowitz Syndrome)
89	Multi-system disease - other
	Others
90	Tubular necrosis (irreversible) or cortical necrosis (different from 88)
91	Tuberculosis
92	Gout
93	Nephrocalcinosis and hypercalcaemic nephropathy
94	Balkan nephropathy
95	Kidney tumour
96	Traumatic or surgical loss of kidney
99	Other identified renal disorders
75	Ischaemic renal disease/cholesterol embolism
76	Glomerulonephritis related to liver cirrhosis
78	Cryoglobulinaemic glomerulonephritis