

All recorded native kidney biopsy diagnoses 2017

	ARI	XH	DGRI	GLAS	MONK	NINE	RAIG	RIE	VHK	Scotland 2017
IgA nephropathy - histologically proven	4	5	1	18	7	2	6	17	6	66
Tubulointerstitial nephritis - histologically proven	7	4	0	25	3	7	4	8	5	63
Microscopic polyangiitis - histologically proven	4	4	1	13	1	2	1	12	3	41
Membranous nephropathy - idiopathic	4	2	0	17	2	4	1	5	2	37
Minimal change nephropathy - histologically proven	2	1	1	10	4	4	2	7	1	32
Primary focal segmental glomerulosclerosis (FSGS)	5	2	0	7	0	7	2	7	1	31
Diabetic nephropathy in Type II diabetes - histologically proven ^a	2	4	1	11	5	3	1	3	0	30
Ischaemic nephropathy / microvascular disease - histologically proven	0	2	0	14	3	2	0	3	1	25
Acute kidney injury	1	4	1	3	3	0	0	11	0	23
Chronic kidney disease (CKD) / chronic renal failure (CRF) - aetiology unclear	7	1	0	2	4	2	2	3	2	23
Systemic lupus erythematosus / nephritis - histologically proven	3	2	0	4	3	4	2	3	2	23
Chronic hypertensive nephropathy - histologically proven	1	0	1	4	3	0	0	10	0	19
Granulomatosis with polyangiitis - histologically proven	1	2	1	6	1	0	0	2	0	13
Renal amyloidosis ^a	2	1	0	4	0	0	1	4	1	13
Mesangial proliferative glomerulonephritis	1	1	1	3	0	0	0	4	1	11
Henoch-Schönlein purpura / nephritis - histologically proven	1	0	0	0	1	1	2	2	2	9
Light chain deposition disease	3	0	0	0	0	0	1	5	0	9
Mesangiocapillary glomerulonephritis type 1	0	0	1	1	0	1	2	2	0	7
Focal and segmental proliferative glomerulonephritis	0	0	0	6	0	0	0	0	0	6
Myeloma cast nephropathy - histologically proven	0	0	0	4	0	0	1	1	0	6
Anti-Glomerular basement membrane (GBM) disease / Goodpasture's syndrome	0	0	0	1	0	3	1	0	0	5
Diffuse endocapillary glomerulonephritis	0	0	1	2	0	0	1	1	0	5
Glomerulonephritis - histologically indeterminate	0	0	0	1	2	0	0	2	0	5
Immunotactoid / fibrillary nephropathy	0	0	0	5	0	0	0	0	0	5
Malignant hypertensive nephropathy / accelerated hypertensive nephropathy	0	1	0	2	0	1	0	0	1	5
Glomerulonephritis - secondary to other systemic disease	1	0	0	2	1	0	0	0	0	4
Thin basement membrane disease	1	0	0	2	1	0	0	0	0	4
Renal amyloidosis	0	0	0	0	0	0	1	1	1	3
Atypical haemolytic uraemic syndrome (HUS) - diarrhoea negative	0	1	1	1	0	0	0	0	0	3
Idiopathic rapidly progressive (crescentic) glomerulonephritis	0	0	0	0	0	0	0	3	0	3

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Tubulointerstitial nephritis with uveitis (TINU) - histologically proven	0	1	0	0	1	0	0	0	1	3
Systemic vasculitis - ANCA negative - histologically proven	0	0	0	1	0	0	0	1	0	2
Essential mixed cryoglobulinaemia - histologically proven	1	0	0	0	1	0	0	0	0	2
Cryoglobulinaemia secondary to systemic disease - histologically proven	0	0	0	0	1	0	0	1	0	2
Membranous nephropathy - drug induced	0	1	0	0	0	1	0	0	0	2
Tubulointerstitial nephritis associated with autoimmune disease - histologically proven	1	0	0	0	0	0	1	0	0	2
AA amyloid secondary to chronic inflammation	0	0	0	0	0	0	0	1	0	1
Acute pyelonephritis	0	0	0	0	0	0	1	0	0	1
Churg-Strauss syndrome - histologically proven	0	0	0	0	0	0	0	1	0	1
Acute cortical necrosis	0	0	0	0	0	0	0	1	0	1
Complement component 3 glomerulopathy	0	0	0	0	0	0	0	1	0	1
Focal segmental glomerulosclerosis (FSGS) secondary to obesity - histologically proven	0	1	0	0	0	0	0	0	0	1
IgA nephropathy secondary to liver cirrhosis - histologically proven	0	0	0	0	0	0	0	1	0	1
Membranous nephropathy - infection associated	0	0	0	0	0	1	0	0	0	1
Membranous nephropathy - malignancy associated	0	0	0	1	0	0	0	0	0	1
Insufficient renal tissue for diagnosis	0	0	0	0	1	0	0	0	0	1
Medullary cystic kidney disease type II	0	0	0	0	0	0	0	1	0	1
Mesangiocapillary glomerulonephritis type 2 (dense deposit disease)	0	0	0	1	0	0	0	0	0	1
Renal sarcoidosis - histologically proven	0	0	0	1	0	0	0	0	0	1