

All recorded native kidney biopsy diagnoses 2016

	ARI	XH	DGRI	GLAS	MONK	NINE	RAIG	RIE	VHK	Scotland 2016
IgA nephropathy - histologically proven	5	5	2	39	13	10	6	23	1	104
Tubulointerstitial nephritis - histologically proven	8	1	0	21	9	8	2	12	4	65
Microscopic polyangiitis - histologically proven	5	6	0	19	2	2	2	16	3	55
Membranous nephropathy - idiopathic	6	4	0	15	2	4	1	8	2	42
Primary focal segmental glomerulosclerosis (FSGS)	3	6	0	13	5	6	0	5	3	41
Minimal change nephropathy - histologically proven	1	4	0	14	2	1	1	6	2	31
Acute kidney injury	1	0	0	11	2	1	0	9	2	26
Granulomatosis with polyangiitis - histologically proven	2	3	0	13	0	4	0	1	0	23
Diabetic nephropathy in type II diabetes - histologically proven	4	1	0	9	4	0	1	4	0	23
Systemic lupus erythematosus / nephritis - histologically proven	0	0	0	13	1	1	3	4	0	22
Ischaemic nephropathy / microvascular disease - histologically proven	1	4	0	7	2	0	2	1	1	18
AL amyloid secondary to plasma cell dyscrasia	3	1	1	4	1	4	2	0	1	17
Henoch-Schönlein purpura / nephritis - histologically proven	3	0	0	5	3	0	1	3	0	15
Mesangial proliferative glomerulonephritis ^b	0	1	0	5	1	0	0	8	0	15
Chronic hypertensive nephropathy - histologically proven	3	2	0	5	1	0	1	2	0	14
Chronic kidney disease (CKD) / chronic renal failure (CRF) - aetiology uncertain / unknown - histologically proven	1	0	0	4	0	0	2	5	0	12
Glomerulonephritis - histologically indeterminate	1	0	0	4	4	0	0	2	0	11
Diabetic nephropathy in type I diabetes - histologically proven	3	2	0	0	3	0	0	0	1	9
Kidney biopsy result normal	0	0	0	1	1	2	0	5	0	9
Thin basement membrane disease	1	0	0	3	2	0	0	2	0	8
Drug-induced tubulointerstitial nephritis - histologically proven	0	4	0	1	0	0	0	2	0	7

Renal sarcoidosis - histologically proven	1	0	0	1	1	0	1	2	1	7
Other	1	0	0	2	0	0	2	1	0	6
Renal amyloidosis ^a	0	0	0	0	0	0	0	3	2	5
Anti-Glomerular basement membrane (GBM) disease / Goodpasture's syndrome - histologically proven	0	0	0	3	1	0	0	0	1	5
Cryoglobulinaemia secondary to systemic disease - histologically proven	1	0	0	0	2	1	1	0	0	5
Focal and segmental proliferative glomerulonephritis	0	0	0	2	2	0	0	1	0	5
Immunotactoid / fibrillary nephropathy	0	0	0	3	2	0	0	0	0	5
Tubulointerstitial nephritis with uveitis (TINU) - histologically proven	1	0	0	0	1	0	1	2	0	5
Glomerulonephritis - secondary to other systemic disease	1	0	0	0	0	0	0	3	0	4
Malignant hypertensive nephropathy / accelerated hypertensive nephropathy - histologically proven	0	0	0	3	0	0	1	0	0	4
Myeloma cast nephropathy - histologically proven	0	1	0	0	0	0	0	3	0	4
Systemic vasculitis - ANCA negative - histologically proven	0	1	0	0	0	0	0	1	1	3
Complement component 3 glomerulopathy	0	0	0	0	0	1	0	2	0	3
Atheroembolic renal disease - histologically proven	0	1	0	1	0	0	0	0	0	2
Focal segmental glomerulosclerosis (FSGS) secondary to obesity - histologically proven	0	0	0	1	0	0	1	0	0	2
Mesangiocapillary glomerulonephritis type 2 (dense deposit disease)	0	0	0	2	0	0	0	0	0	2
Nephropathy due to tacrolimus - histologically proven	0	0	0	1	0	0	0	1	0	2
Tubulointerstitial nephritis associated with autoimmune disease - histologically proven	0	0	0	0	0	2	0	0	0	2
AA amyloid secondary to chronic inflammation	0	0	0	0	1	0	0	0	0	1
Familial amyloid secondary to protein mutations - histologically proven	0	0	0	1	0	0	0	0	0	1

Calculus nephropathy / urolithiasis	0	0	0	1	0	0	0	0	0	1
Acute cortical necrosis	0	1	0	0	0	0	0	0	0	1
Fabry disease - histologically proven	0	0	0	0	0	1	0	0	0	1
Idiopathic rapidly progressive (crescentic) glomerulonephritis	0	0	0	0	0	0	0	0	1	1
Membranous nephropathy - malignancy associated	0	0	0	1	0	0	0	0	0	1
Light chain deposition disease	1	0	0	0	0	0	0	0	0	1
Mesangiocapillary glomerulonephritis type 1	1	0	0	0	0	0	0	0	0	1
Nephropathy due to analgesic drugs - histologically proven	0	0	0	0	0	0	0	1	0	1
Nephropathy due to lithium - histologically proven	0	0	0	0	0	0	0	1	0	1
Nephropathy related to HIV - histologically proven	1	0	0	0	0	0	0	0	0	1
Renal scleroderma / systemic sclerosis - histologically proven	0	0	0	0	0	0	0	1	0	1
Renal tuberculosis	0	1	0	0	0	0	0	0	0	1

^a Not including AL amyloidosis secondary to plasma cell dyscrasia or AA amyloidosis secondary to chronic inflammation

^b Not including IgA nephropathy or IgM nephropathy.