All recorded native biopsy diagnoses 2015

	Scotland 2015	ARI	XH	DGRI	GLAS	MONK	NINE	RAIG	RIE	VHK
IgA nephropathy	101	4	8	0	30	11	16	4	25	3
Membranous nephropathy - idiopathic	73	13	5	4	22	4	8	4	10	3
Tubulointerstitial nephritis	61	3	6	0	16	4	13	2	15	2
Insufficient tissue for histological diagnosis	43	16	2	1	4	3	4	0	11	2
Diabetic nephropathy in type I or type II diabetes - histologically proven	42	3	3	1	15	10	2	3	4	1
Systemic lupus erythematosus / nephritis	39	7	2	0	12	4	1	3	10	0
Microscopic polyangiitis - histologically proven	34	2	2	1	8	1	4	7	7	2
Granulomatosis with polyangiitis	33	8	2	0	11	3	5	2	1	1
Primary focal segmental glomerulosclerosis (FSGS)	31	7	1	0	7	1	5	1	8	1
Minimal change nephropathy	28	2	1	1	12	1	2	0	5	4
Mesangiocapillary glomerulonephritis - not type 2	18	2	0	0	7	3	0	0	4	2
Other	17	7	0	0	7	0	1	0	1	1
Acute kidney injury (acute tubular necrosis)	14	2	0	0	6	2	0	0	4	0
Focal and segmental proliferative glomerulonephritis	13	0	0	0	5	2	0	0	6	0
AL amyloid secondary to plasma cell dyscrasia	12	1	3	1	4	1	0	0	2	0
Chronic kidney disease (CKD) / chronic renal failure (CRF) - aetiology uncertain / unknown	12	3	0	0	1	0	0	0	8	0
Chronic hypertensive nephropathy	11	1	0	0	1	0	5	3	1	0
Drug-induced tubulointerstitial nephritis	11	2	2	0	0	0	0	2	5	0
Henoch-Schönlein purpura / nephritis	10	1	2	2	0	3	2	0	0	0
Ischaemic nephropathy / microvascular disease	10	0	1	0	2	1	2	0	4	0

Kidney biopsy normal	8	1	2	0	2	1	1	1	0	0
Renal amyloidosis [a]	7	4	1	0	1	0	1	0	0	0
Complement component 3 glomerulopathy	7	0	0	0	4	0	0	0	3	0
Glomerulonephritis - histologically indeterminate	7	0	0	0	2	1	1	0	1	2
Myeloma cast nephropathy - histologically proven	7	3	0	0	0	0	0	1	2	1
Light chain deposition disease	6	0	0	0	3	0	0	1	2	0
Mesangial proliferative glomerulonephritis	6	1	0	0	0	1	2	0	2	0
Renal sarcoidosis - histologically proven	6	0	0	0	1	2	0	1	2	0
Glomerulonephritis - secondary to other systemic disease	5	0	0	0	0	0	0	0	5	0
Idiopathic rapidly progressive (crescentic) glomerulonephritis	5	0	0	0	2	0	0	0	3	0
Thin basement membrane disease	5	3	1	0	0	0	0	1	0	0
Diagnosis not recorded	4	0	1	0	0	0	0	2	1	0
Diffuse endocapillary glomerulonephritis	4	0	0	0	1	0	0	1	2	0
Mesangiocapillary glomerulonephritis type 3	4	1	0	0	2	0	0	0	1	0
Anti-Glomerular basement membrane (GBM) disease / Goodpasture's syndrome - histologically proven	3	0	0	0	0	0	0	1	2	0
Immunotactoid / fibrillary nephropathy	3	0	0	0	2	1	0	0	0	0
Malignant hypertensive nephropathy / accelerated hypertensive nephropathy - histologically proven	3	0	0	0	3	0	0	0	0	0
Nephropathy due to ciclosporin - histologically proven	3	2	1	0	0	0	0	0	0	0
AA amyloid secondary to chronic inflammation	2	0	0	0	1	0	0	0	0	1
Alport syndrome - histologically proven	2	0	0	0	0	1	0	1	0	0
Atheroembolic renal disease - histologically proven	2	0	0	0	2	0	0	0	0	0

Cryoglobulinaemia secondary to systemic disease - histologically proven	2	0	0	0	0	1	0	0	1	0
Sickle cell nephropathy - histologically proven	2	0	0	0	1	0	0	0	1	0
Tubulointerstitial nephritis associated with autoimmune disease - histologically proven	2	0	0	0	1	0	0	1	0	0
Tubulointerstitial nephritis with uveitis (TINU) - histologically proven	2	1	0	0	1	0	0	0	0	0
Acute pyelonephritis	1	0	0	0	1	0	0	0	0	0
Atypical haemolytic uraemic syndrome (HUS) - diarrhoea negative	1	1	0	0	0	0	0	0	0	0
Churg-Strauss syndrome - histologically proven	1	0	0	0	0	0	0	0	1	0
Systemic vasculitis - ANCA negative - histologically proven	1	0	0	0	0	0	0	1	0	0
Fabry disease - histologically proven	1	0	0	0	0	1	0	0	0	0
Familial interstitial nephropathy - histologically proven	1	0	0	0	0	0	0	0	1	0
Focal segmental glomerulosclerosis (FSGS) secondary to obesity - histologically proven	1	0	0	0	0	1	0	0	0	0
Membranous nephropathy - drug induced	1	0	0	0	0	0	0	0	1	0
Nephropathy due to lithium - histologically proven	1	0	0	0	0	0	0	0	1	0
Nephropathy related to HIV - histologically proven	1	0	0	0	0	0	0	0	1	0
Acute cortical necrosis	0	0	0	0	0	0	0	0	0	0
Essential mixed cryoglobulinaemia - histologically proven	0	0	0	0	0	0	0	0	0	0
Cryoglobulinaemia secondary to hepatitis C - histologically proven	0	0	0	0	0	0	0	0	0	0
Membranous nephropathy - malignancy associated	0	0	0	0	0	0	0	0	0	0
IgM - associated nephropathy	0	0	0	0	0	0	0	0	0	0
Mesangiocapillary glomerulonephritis type 2	0	0	0	0	0	0	0	0	0	0

(dense deposit disease)										

[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.