Top level metadata in RaDaR, updated 19/04/2021

Rare Disease Cohort	Patient Demographics	Primary Diagnoses	Lab results and Observations	Medications Link
Alport Syndrome	842	684	623	414
Atypical Haemolytic Uraemic Syndrome	270	217	215	158
Autosomal Dominant Polycystic Kidney Disease	7351	5036	5664	3995
Autosomal Dominant Tubulointerstitial Kidney Disease (FUAN)	203	165	146	105
Autosomal Recessive Polycystic Kidney Disease/Nephronophthisis	221	165	179	118
BK Nephropathy	34	28	29	27
Calciphylaxis	49	40	36	29
Cystinosis	146	133	131	90
Cystinuria	446	401	295	91
Dent Disease and Lowe Syndrome	60	51	31	20
Fabry Disease	45	30	32	15
Fibromuscular Dysplasia	39	28	25	7
HNF1b Mutations	81	65	55	28
Hyperoxaluria	119	97	85	56
Idiopathic Nephrotic Syndrome	3907	3407	3123	2478
IgA Nephropathy	3915	2889	3020	2435
Membranoproliferative Glomerulonephritis / Dense Deposit Disease	1083	852	850	691
Membranous Nephropathy	2302	1876	1605	1243
Monoclonal Gammopathy of Renal Significance	158	135	106	71
Pregnancy	656	466	427	266
Retroperitoneal Fibrosis	141	122	98	83
STEC-associated HUS	162	145	92	28
Tuberous Sclerosis	224	184	120	77
Tubulopathy	339	296	185	125
Vasculitis	4428	3599	2807	2321