RaDaR Diagnoses and Cohorts

The following table shows which cohort to enter each patient into on RaDaR

Diagnosis	RaDaR Cohort
Adenine Phosphoribosyltransferase Deficiency (APRT-D)	APRT Deficiency
AH amyloidosis	MGRS
AHL amyloidosis	MGRS
AL amyloidosis	MGRS
Alport Syndrome Carrier - Female heterozygote for X-linked Alport Syndrome (COL4A5)	Alport
Alport Syndrome Carrier - Heterozygote for autosomal Alport Syndrome (COL4A3, COL4A4)	Alport
Alport Syndrome	Alport
Anti-Glomerular Basement Membrane Disease (Goodpastures)	Vasculitis
Atypical Haemolytic Uraemic Syndrome (aHUS)	aHUS
Autoimmune distal renal tubular acidosis	Tubulopathy
Autosomal recessive distal renal tubular acidosis	Tubulopathy
Autosomal recessive proximal renal tubular acidosis	Tubulopathy
Autosomal Dominant Polycystic Kidney Disease (ARPKD)	ADPKD
Autosomal Dominant Tubulointerstitial Kidney Disease (ADTKD)	ADTKD
Autosomal Recessive Polycystic Kidney Disease (ARPKD)	ARPKD/NPHP
Bartters Syndrome	Tubulopathy
BK Nephropathy	BK Nephropathy
C3 Glomerulopathy	MPGN
C3 glomerulonephritis with monoclonal gammopathy	MGRS
Calciphylaxis	Calciphylaxis
Crystalglobulinaemia	MGRS
Crystal-storing histiocytosis	MGRS
Cystinosis	Cystinosis
Cystinuria	Cystinuria
Dense Deposit Disease (DDD)	MPGN
Dent Disease	Dent & Lowe
Denys-Drash Syndrome	INS
Dominant hypophosphatemia with nephrolithiasis or osteoporosis	Tubulopathy
Drug induced Fanconi syndrome	Tubulopathy
Drug induced hypomagnesemia	Tubulopathy
Drug induced Nephrogenic Diabetes Insipidus	Tubulopathy
Epilepsy, Ataxia, Sensorineural deafness, Tubulopathy (EAST) Syndrome	Tubulopathy
Fabry Disease	Fabry

Diagnosis	RaDaR Cohort
Familial Hypomagnesaemia with hypercalciuria and nephrocalcinosis	Tubulopathy
Familial primary hypomagnesemia with hypocalcuria	Tubulopathy
Familial primary hypomagnesemia with normocalcuria EGF	Tubulopathy
Familial renal glucosuria	Tubulopathy
Fanconi Renotubular syndrome 1 (FRTS1)	Tubulopathy
Fanconi Renotubular syndrome 2 (FRTS2)	Tubulopathy
Fanconi Renotubular syndrome 3 (FRTS3)	Tubulopathy
Fibrillary Glomerulonephritis	MGRS
Fibromuscular Dysplasia	Fibromuscular Dysplasia
Focal Segmental Glomerulosclerosis (FSGS)	INS
Generalized pseudohypoaldosteronism type 1	Tubulopathy
Giant Vessel Arteritis	Vasculitis
Gitelman Syndrome	Tubulopathy
Glomerulocystic Disease	HNF1b
Heavy metal induced Fanconi syndrome	Tubulopathy
Hepatocyte Nuclear Factor-1 Beta Mutations (HNF1B)	HNF1b
Hereditary renal hypouricemia	Tubulopathy
Hereditary hypophosphatemic rickets with hypercalciuria	Tubulopathy
Hyperuricaemic Nephropathy	ADTKD
IgA Nephropathy	IgA Nephropathy
IgA Vasculitis (Henoch Schonlein)	Vasculitis
Immunotactoid/Glomerulonephritis with Organised Microtubular Monoclonal Immunoglobulin Deposits (GOMMID)	MGRS
Inherited/Genetic Diabetes Mellitus Type II (MODY)	HNF1b
Inherited Renal Cancer Syndrome	Renal Cancer Inherited
Intracapillary monoclonal IgM without cryoglobulin	MGRS
Intraglomerular/capillary lymphoma/leukaemia	MGRS
Isolated autosomal dominant hypomagnesemia, Glaudemans type	Tubulopathy
Large Vessel Vasculitis	Vasculitis
Liddle Syndromes	Tubulopathy
Light chain cast nephropathy	MGRS
Light chain proximal tubulopathy, crystalline	MGRS
Light chain proximal tubulopathy, non crystalline	MGRS
Lowe Syndrome	Dent & Lowe
Medium Vessel Vasculitis	Vasculitis
Medullary Cystic Kidney Disease	ADTKD
Membranous Nephropathy	Membranous Nephropathy
Membranoproliferative Glomerulonephritis (MPGN)	MPGN

Diagnosis	RaDaR Cohort
Minimal Change Nephropathy	INS
Mitochondrial Disease	Mitochondrial
Monoclonal Immunoglobulin Deposition Disease (MIDD; includes Light Chain Deposition Disease - LCDD; Heavy Chair Deposition Disease - HCDD; and Light and Heavy Chain Deposition Disease - LHCDD)	MGRS
Multicystic Dysplastic Kidneys	HNF1b
Nail Patella Syndrome	INS
Nephrogenic diabetes insipidus	Tubulopathy
Nephrogenic syndrome of inappropriate antidiuresis	Tubulopathy
Nephronophthisis	ARPKD/NPHP
Oncogenic osteomalacia	Tubulopathy
Osteopetrosis with renal tubular acidosis	Tubulopathy
Pregnancy and Chronic Kidney Disease	Pregnancy
Pregnancy & Lupus Nephritis	Pregnancy
Pregnancy in a Renal Transplant Recipient	Pregnancy
Primary hypomagnesemia with secondary hypocalcemia	Tubulopathy
Primary Hyperoxaluria	Hyperoxaluria
Primary Renal Fanconi syndrome	Tubulopathy
Proliferative glomerulonephritis with monoclonal immunoglobulin deposits (PGNMID)	MGRS
Proximal tubulopathy without crystals	MGRS
Pseudohypoaldosteronism type 2A	Tubulopathy
Pseudohypoaldosteronism type 2B	Tubulopathy
Pseudohypoaldosteronism type 2C	Tubulopathy
Pseudohypoaldosteronism type 2D	Tubulopathy
Pseudohypoaldosteronism type 2E	Tubulopathy
Pure Red Cell Aplasia	PRCA
Renal Cysts & Diabetes Syndrome	HNF1b
Shiga Toxin Associated Haemolytic Uraemic Syndrome (HUS)	STEC HUS
Renal pseudohypoaldosteronism type 1	Tubulopathy
Retroperitoneal Fibrosis	Retroperitoneal Fibrosis
Small Vessel Vasculitis (ANCA Associated)	Vasculitis
Steroid Resistant Nephrotic Syndrome (SRNS)	INS
Steroid Sensitive Nephrotic Syndrome (SSNS)	INS
Thin Basement Membrane Nephropathy	Alport
Thrombotic Microangiopathy with monoclonal gammopathy	MGRS
Tuberous Sclerosis	Tuberous Sclerosis
Type 1 cryoglobulinaemic Glomerulonephritis	MGRS
Unclassified Monoclonal Gammopathy of Renal Significance	MGRS
Uromodulin-Associated Nephropathy (Familial Juvenile Hyperuricaemic Nephropathy)	ADTKD
Variable Vessel Vasculitis	Vasculitis
Vasculitis	Vasculitis

NephroS patients should be entered under **INS** first and then **NephroS**