

Introduction

Introduction and Overview

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AL/AH Amyloidosis

AA Amyloidosis

AFib Amyloidosis

AGel Amyloidosis

ALECT2 Amyloidosis

AApoAI Amyloidosis

ApoAII Amyloidosis

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*Autosomal Dominant FSGS Due to *INF2* Mutations*

**APOL1*-Related Glomerular Disease*

Schimke Immuno-Osseous Dysplasia

Genetic Storage and Lipid Diseases

Lecithin-Cholesterol Acyltransferase Deficiency

APOE Lipoprotein Glomerulopathy

Type III Hyperlipoproteinemia

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Thrombotic Microangiopathy, Autoimmune
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Multicentric Castleman Disease
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Glomerulopathy of Hereditary Multiple Exostoses
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Fibromuscular Dysplasia
Neurofibromatosis
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Renal Artery Thrombosis
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Overview and Classification of Tubulointerstitial Diseases

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Antibrush Border Autoantibody Tubulointerstitial Nephritis

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Light Chain Proximal Tubulopathy Without Crystals

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Drug-Induced Acute Interstitial Nephritis

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Osmotic Tubulopathy

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Acute Phosphate Nephropathy

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Calcineurin Inhibitor Toxicity

mTOR Inhibitor Toxicity

Vancomycin-Induced Cast Nephropathy

Warfarin Nephropathy

Illicit Drugs and Opioids

Toxic Injury

Lead and Other Heavy Metal Toxins

Aristolochic Acid Nephropathy

Balkan Endemic Nephropathy

Ethylene Glycol Toxicity

Argyria

Autosomal Dominant Tubulointerstitial Kidney Disease

*Autosomal Dominant Tubulointerstitial Kidney Disease, *MUC1*-Related*

*Autosomal Dominant Tubulointerstitial Kidney Disease, *UMOD*-Related*

*Autosomal Dominant Tubulointerstitial Kidney Disease, *REN*-Related*

*Autosomal Dominant Tubulointerstitial Kidney Disease, *HNF1B*-Related*

Genetic Crystal Deposition Diseases

Primary Hyperoxaluria

2,8-Dihydroxyadeninuria

Cystinosis

Uric Acid Nephropathy/Gout

Genetic Transport Diseases

Bartter Syndrome

Dent Disease

Oculocerebrorenal Syndrome of Lowe

Other Genetic Diseases Affecting Tubules

Methylmalonic Acidemia

Systemic Karyomegaly

Mitochondriopathies

Miscellaneous Tubulointerstitial Diseases

Nephrocalcinosis
Secondary Oxalosis
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Extramedullary Hematopoiesis
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Bacterial Infections of the Kidney
Acute Pyelonephritis
Chronic Pyelonephritis
Xanthogranulomatous Pyelonephritis
Malakoplakia
Tuberculosis
BCG Granulomatous Interstitial Nephritis
Leprosy
Megalocytic Interstitial Nephritis
Nocardiosis
Leptospirosis
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Fungal, Rickettsial, and Parasitic Infections of the Kidney
Mucormycosis
Candidiasis
Histoplasmosis
Coccidioidomycosis
Blastomycosis
Paracoccidioidomycosis
Aspergillosis
Cryptococcosis
Microsporidiosis
Rickettsial Infections
Toxoplasmosis
Hydatidosis
Viral Infections of the Kidney
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Adenovirus Infection

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Dysplasia/Hypoplasia/Agenesis

Oligomeganephronia

Ectopia, Malrotation, Duplication, Fusion, Supernumerary Kidney

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Autosomal Recessive Polycystic Kidney Disease

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Chronic T-Cell-Mediated Rejection

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Chronic Antibody-Mediated Rejection

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De Novo Membranous Glomerulonephritis

Anti-GBM Disease in Alport Syndrome

Engraftment Syndrome

Nonimmunologic Injury

Acute Allograft Ischemia

Hyperperfusion Injury

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Lymphocele

Transplant Renal Artery Stenosis

Renal Artery or Vein Thrombosis

Posttransplant Lymphoproliferative Disease

BK Polyomavirus Neoplasia

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Alport Collagen IV Immunofluorescence

C4d Immunohistochemistry/Immunofluorescence

Polyomavirus Detection in Tissue

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