NPHP1

Nephronophthisis, Joubert Syndrome, Senior-Loken Syndrome

Mode of Inheritance	Autosomal recessive
Renal Phenotype	 Nephronophthisis Age of onset: Childhood or early adolescence
Extra-renal Manifestations	 Growth retardation Retinitis pigmentosa Oculomotor apraxia Developmental delay Ataxia and impaired balance Cerebellar vermis hypoplasia
Pre-Transplant Management	Screening and management of extra-renal manifestations
Transplant Considerations	Tailor immunosuppression given low risk of recurrence post- transplant
Post-Transplant Management	 Low risk of disease recurrence Higher rate of post-transplant polyuria in patients with native kidneys (Hamiwka <i>Ped Transplant</i> 12:878, 2008)

https://www.omim.org/entry/607100