

EYA1

Branchio-Oto-Renal Syndrome (BOR)

Mode of Inheritance	<ul style="list-style-type: none">• Autosomal dominant• Variable expressivity and incomplete penetrance
Renal Phenotype	<ul style="list-style-type: none">• Variable, including: renal dysplasia/aplasia, polycystic kidneys, vesicoureteral reflux, and other structural anomalies
Extra-renal Manifestations	<ul style="list-style-type: none">• Facial features: long, narrow face• Sensorineural and/or conductive hearing loss• Preauricular pits• External ear and/or cochlear malformations• Branchial cleft fistulas or cysts
Pre-Transplant Management	<ul style="list-style-type: none">• Screening and management of extra-renal manifestations
Transplant Considerations	<ul style="list-style-type: none">• Careful screening of potential living related donors (inherited in an autosomal dominant manner)
Post-Transplant Management	<ul style="list-style-type: none">• Low risk of disease recurrence