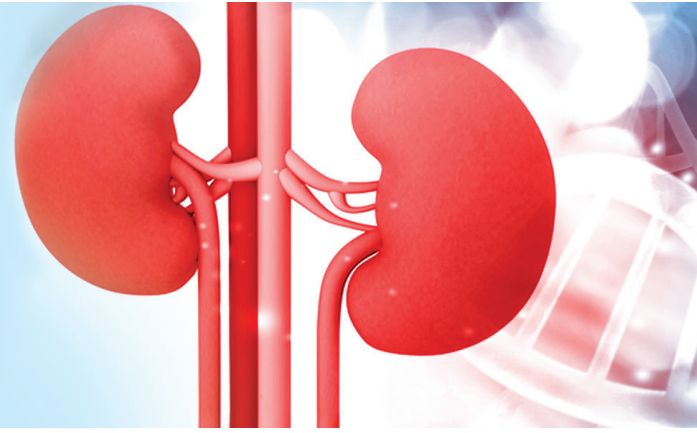


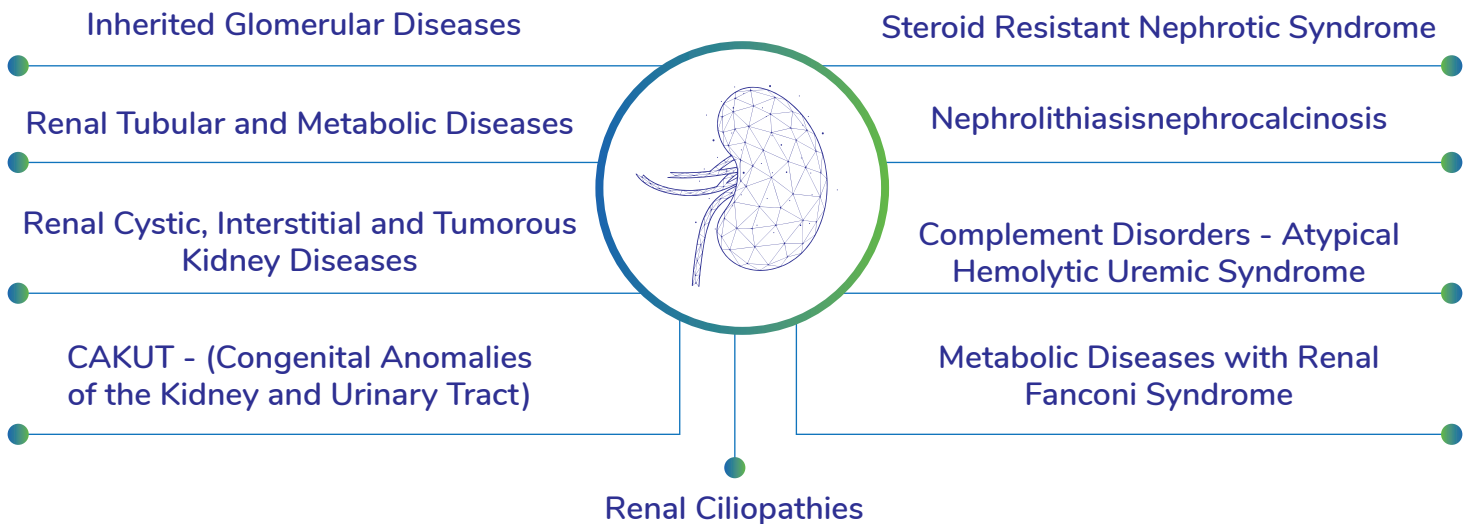
Nephro Genetics

Comprehensive Gene Panels for Inherited Renal Diseases



- Inherited renal diseases (IRD) are among the leading causes of early-onset chronic kidney disease
- IRDs account for ~10–15% of kidney transplant cases in adults
- Genetic testing enables the early detection of disease causal variant which helps in timely treatment & management strategies

Inherited Kidney Diseases



20% of cases of CKD are thought to be due to genetic forms of renal disease³.

Genetic renal disorders affects 5-15% of the adult population¹.

To date, more than 600 genes have been implicated in monogenic kidney diseases

Who should get Tested

Diagnosed with cystic kidney diseases, tubulopathies or monogenic glomerulopathies

With family history

Syndromic disease, with extra renal manifestations

Congenital kidney anomalies

Unknown aetiology with <25–30 years of age of onset

Hematuria/ proteinuria without a definitive diagnosis

Interstitial nephropathy or chronic kidney disease without a known cause

Unusual disease course

To guide therapeutic decisions

If an individual belongs to a high risk population

Testing Modalities

Next-Generation Sequencing (NGS)

Multiplex Ligation-dependent Probe Amplification (MLPA)



Chromosomal Microarray

Test Details

MGM106	Polycystic Kidney Disease gene panel
MGM104	Bartter syndrome gene panel
MGM103	Alport syndrome gene panel
MGM153	Joubert Syndrome gene panel
MGM308	Hemolytic Uremic Syndrome - HUS (deletion duplication analysis)
MGM368	Nephrotic Syndrome gene panel
MGM107	Primary Hyperoxaluria gene panel
MGM244	Bardet-Biedl Syndrome gene panel
MGM326	Trio Exome Sequencing
MGM2572	ExomeMAX
MGM2741	KaryoTrack

To know more about relevant genetic testing options, please write to us at techsupport@medgenome.com



Genetic Counselling: Free pre & post test genetic counselling with our expert and certified genetic counsellors is available

Why MedGenome

- Unmatched experience of 350,000+ exomes
- South Asia largest database with 5,000,000+ variants
- Recommended by 4000+ hospitals
- Trusted by 10000+ clinicians
- Consistent proficiency testing through "External Quality Monitoring Programs"
- Two levels of analysis and review by clinical geneticists
- Expert genetic counselling available pre and post test

References : Pubmed: 24980890; 35460632, 34264297

Talk to the Experts: [1800 103 3691](tel:18001033691)

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