Limbach Genetics
Medizinische Genetik Mainz

Test Portfolio

We are still in the process of developing our services and are continually expanding our test range. The provision of expert counselling services for the various indications is extremely important to us. Please find below our established panels in the field of nephrogenetics.

In the event that a test you require is not listed, please contact us and we will find a solution. Should you send us a request for a test that is not yet offered by our facilities in Mainz, we will make sure that your samples are processed by a certified partner laboratory of comparable superior quality.

Please contact us if you have questions about our services in diagnostics. We are happy to provide assistance.

Indication-specific NGS-Panel (nephrogenetic)	
A	
	Agenesie, renal
	Alagille-Syndrom
	Alport-Syndrom
	Alström-Syndrom
	Aminoazidurie
	ARC (Arthrogrypose-Nierenfunktionsstörung-Cholestase)-Syndrom
	Atypisches hämolytisch-urämisches Syndrom (aHUS)
B	
	Bardet-Biedl-Syndrom (BBS)
	Bartter-Syndrom
	Birt-Hogg-Dubé-Syndrom
	BOR (branchio-oto-renales)-Syndrom
C	
	C1q-Defizienz
	CAKUT (Congenital Anomalies of the Kidney and Urinary Tract)
	Cobalamin-Mangel
	Cystinose
	Cystinurie
D	
	Dent-Krankheit
	Denys-Drash-Syndrom
	Diabetes insipidus
	Dysplasie, renal

E	
	Ellis-van-Creveld-Syndrom (EVC)
F	
	Fokal-segmentale Glomerulosklerose (FSGS)
	Frasier-Syndrom
G	
	Gitelman-Syndrom
H	
	Hämaturie
	Heterotaxie
	Hyperkalzämie
	Hyperoxalurie
	Hypomagnesiämie
	Hypophosphatämische Rachitis/Phosphatdiabetes
	Hypoplasie, renal
I	
	Imerslund-Gräsbeck-Syndrom
J	
	Jeune-Syndrom
	Joubert-Syndrom
L	
	Liddle-Syndrom
M	
	Mainzer-Saldino-Syndrom
	Meckel-Gruber-Syndrom
N	
	Nephronophthise (NPHP)
	Nephropathie vom Typ der dünnen Basalmembran (TBMN)
	Nephrotisches Syndrom
	Nierensteinerkrankungen (Nephrolithiasis)
O	
	Oro-fazio-digitales Syndrom (OFD)
P	
	Peroxisomen-Biogenese Störungen
	Polyzystische Lebererkrankung (PCLD)
	Polyzystische Nierenerkrankung, autosomal-dominant (ADPKD)
	Polyzystische Nierenerkrankung, autosomal-rezessiv (ARPKD)

	Primäre Ziliäre Dyskinesie (PCD)
	Primärer Aldosteronismus
	Proteinurie
	Proximale Tubulopathie
	Pseudohypoaldosteronismus
R	
	Renal-tubuläre Dysgenesie (RTD)
	Renales Fanconi-Syndrom
	Renal-tubuläre Azidose (RTA)
S	
	Senior-Loken-Syndrom
	Sensenbrenner-Syndrom
T	
	Tuberöse Sklerose (TSC)
	Tubulo-interstitielle Nierenerkrankung, autosomal-dominant (ADTKD)
U	
	Urolithiasis
V	
	VACTERL-Assoziation
	Von Hippel-Lindau-Erkrankung (VHL)
W	
	Wilms-Tumor
X	
	Xanthinurie
Z	
	Ziliopathie
	Zellweger-Syndrom
	Zystische Nierenerkrankung/Zystennieren