RB&HH
SPECIALIST CARE

Clinical Genetics and Genomics Laboratory
Royal Brompton Hospital
Sydney Street
London SW3 6NP
Tel: 020 7352 8121 ext. 3009

Clinical Genetics Test Price List (Non-NHS)

Panel	Cost (£)
Aortopathy and connective tissue panels	
Alport syndrome, X-linked	750.00
Cutis laxa	950.00
Ehlers-Danlos syndrome (EDS)	1,050.00
Familial thoracic aortic aneurysm (FTAA)	1,050.00
Loeys-Dietz syndrome (LDS)	950.00
Marfan syndrome (MFS)	950.00
Weill-Marchesani syndrome	950.00
All Aortopathy and connective tissue genes	1,150.00
Arrhythmia panels	
Andersen-Tawil syndrome	750.00
Brugada syndrome (BrS)	950.00
Catecholaminergic polymorphic ventricular tachycardia (CPVT)	950.00
Long QT syndrome (LQTS)	1,050.00
Short QT syndrome	950.00
All Arrhythmia genes	1,150.00
Cardiomyopathy panels	
Arrhythmogenic right ventricular dysplasia/cardiomyopathy (ARVD/C)	1,150.00
Dilated cardiomyopathy (DCM)	1,350.00
Hypertrophic cardiomyopathy (HCM)	1,350.00
Laminopathy	750.00
Noncompaction cardiomyopathy (LVNC)	950.00
Fabry disease	700.00
All Cardiomyopathy genes	1,500.00
Familial Hypercholesterolemia	
Familial Hypercholesterolemia (FH)	700.00
Other cardiac conditions and genes	
Alagille syndrome	750.00
Carney complex	750.00
Heterotaxy/situs ambiguous	1,150.00
Holt-Oram syndrome	700.00
NKX2-5-related disorders	750.00
Noonan spectrum disorders	950.00
SALL4-related disorders	750.00
Vasculopathy panels	
Birt-Hogg-Dubé syndrome (Primary spontaneous pneumothorax)	700.00
Capillary malformation-arteriovenous malformation/Parkes-Weber syndrome	700.00
Hereditary Haemorrhagic Telangiectasia (HHT)	700.00
Homocystinuria	750.00
Megalencephaly Capillary Malformation syndrome	750.00
Microcephaly Capillary Malformation syndrome (MCAP)	750.00
Polycystic kidney disease	750.00
Venous Malformations	750.00
All Vasculopathy genes	950.00

RB&HH
SPECIALIST CARE

Clinical Genetics and Genomics Laboratory
Royal Brompton Hospital
Sydney Street
London SW3 6NP
Tel: 020 7352 8121 ext. 3009

Panel	Cost (£)
Bronchiectasis testing	
Cystic Fibrosis targeted mutation analysis - 36 most common *CFTR* mutations in EU populations	175.00
Sequencing of the *CFTR* gene	750.00
Non-CF Bronchiectasis	950.00
Primary Ciliary Dyskinesia (PCD)	1,000.00
All Bronchiectasis genes	1,350.00
Ciliopathy panels	
Joubert syndrome	1,000.00
Orofaciodigital syndrome	850.00
Short rib thoracic dysplasia (Jeune syndrome)	1,000.00
All Ciliopathy genes (including PCD)	1,350.00
Congenital Respiratory conditions panels	
Alveolar capillary dysplasia	550.00
Ataxia telangiectasia	550.00
Central Hypoventilation syndrome	750.00
Periventricular nodular heterotopia and lung disease	950.00
Primary pulmonary hypoplasia	750.00
Pulmonary alveolar microlithiasis	750.00
All Congenital Respiratory condition genes	1,050.00
Emphysema panels	
Alpha-1-Antitrypsin deficiency	400.00
All emphysema genes	750.00
Immunodeficiency panels	
Agammaglobulinemia	750.00
Autoimmune lymphoproliferative syndrome	700.00
Autoinflammation, antibody deficiency and immune dysregulation syndrome	700.00
Candidiasis, familial	800.00
Hyper-IgE recurrent infection	700.00
Immunodeficiency, common variable	1,050.00
Immunodysregulation, polyendocrinopathy and enteropathy	700.00
Susceptibility to Aspergillosis	700.00
All Immunodeficiency genes	1,150.00
Interstitial Lung Disease (ILD) panels	
Childhood ILD	875.00
Hermansky-Pudlak syndrome	875.00
Pulmonary fibrosis, familial	1,050.00
Tuberous sclerosis	750.00
All Interstitial Lung Disease (ILD) genes	1,350.00
Other panels	
Molecular autopsy (sudden cardiac death, SCD)	1,500.00
Pulmonary Hypertension	800.00
All Inherited Cardiac Conditions genes	1,600.00
All Inherited Respiratory Conditions genes	1,600.00
Opening up panel to next level (e.g. cardiomyopathy to all ICCs)	175.00
Familial variant testing (cascade testing) by Sanger sequencing, MLPA or ddPCR	200.00
Extract and store DNA	60.00
Bioinformatic CNV analysis from NGS data (without SNV analysis)	750.00