

## Project Code Reference List

When completing the DCS<sup>1</sup> and SLF<sup>1</sup> ('project' field), please use the code for the Rare Diseases OR Research Tissue Bank cohort the participant is being recruited to.

| <b>BioResource – Research Tissue Bank</b> |  | <b>CODE</b> |
|-------------------------------------------|--|-------------|
| Generic                                   |  | RTB-GEN     |
| Inflammatory Bowel Disease                |  | RTB-IBD     |
| Mental Health                             |  | RTB-MHS     |
| Strides                                   |  | RTB-STR     |
| NAFLD                                     |  | RTB-NAF     |
| IMID                                      |  | RTB-IMI     |

| <b>BioResource – Rare Diseases Cohort</b>            |  | <b>CODE</b> |
|------------------------------------------------------|--|-------------|
| Achondroplasia and Hypochondroplasia                 |  | RDC-ACH     |
| Acute Coronary Syndromes: SCAD, CAE and FMD          |  | RDC-ACS     |
| Alpha 1 Antitrypsin Deficiency                       |  | RDC-AAT     |
| Arthrogryposis                                       |  | RDC-AMC     |
| Atypical Haemolytic Uraemic Syndrome                 |  | RDC-AHU     |
| Autoimmune Hepatitis                                 |  | RDC-AIH     |
| Autosomal Dominant Parkinson's Disease               |  | RDC-ADP     |
| Autosomal Dominant Tubulointerstitial Kidney Disease |  | RDC-ATK     |
| Birdshot Chorioretinopathy                           |  | RDC-BCR     |
| Bleeding and Platelet Disorders                      |  | RDC-BPD     |
| Cerebral Small Vessel Disease                        |  | RDC-CSV     |
| CHARGE Syndrome                                      |  | RDC-CHG     |
| Ciliopathies: Bardet Biedl and Alström Syndrome      |  | RDC-CIL     |
| Congenital Hyperinsulinism                           |  | RDC-CHI     |
| Cutaneous Lymphoma                                   |  | RDC-CTL     |
| Dystonia                                             |  | RDC-DYS     |
| Familial Interstitial Pneumonia                      |  | RDC-FIP     |
| Giant Cell Arteritis                                 |  | RDC-GCA     |
| Haemoglobinopathies                                  |  | RDC-HBP     |
| Haemophilia Centres Project***                       |  | RDC-HCP     |
| Histiocytic Disorders                                |  | RDC-HCD     |
| IgA Nephropathy                                      |  | RDC-IAN     |
| Imprinting Disorders cohort                          |  | RDC-IMP     |
| Inherited Optic Neuropathies                         |  | RDC-ION     |
| Inherited Retinal Dystrophy                          |  | RDC-IRD     |

*...continued...*

| BioResource – Rare Diseases Cohort                                        | CODE    |
|---------------------------------------------------------------------------|---------|
| Intrahepatic Cholestasis of Pregnancy                                     | RDC-ICP |
| Membranoproliferative Glomerulonephritis and C3 Glomerulopathy            | RDC-PMG |
| Multiple Endocrine Neoplasia Type 1 and Pancreatic Neuroendocrine Tumours | RDC-ECT |
| Multiple Primary Malignant Tumours                                        | RDC-MMT |
| Multiple System Atrophy                                                   | RDC-MSA |
| Narcolepsy - Type 1 and Type 2                                            | RDC-NRC |
| Neurofibromatosis type 1                                                  | RDC-NF1 |
| Neurofibromatosis type 2                                                  | RDC-NF2 |
| Next Generation Children***                                               | RDC-NGC |
| Ocular Maldevelopment: Microphthalmia, Anophthalmia and Ocular Coloboma   | RDC-MAC |
| Overgrowth Disorders                                                      | RDC-POD |
| Paediatric Neurodevelopmental Disorders                                   | RDC-PND |
| Pregnancy in Chronic Kidney Disease                                       | RDC-CKD |
| Primary Biliary Cholangitis                                               | RDC-PBC |
| Primary Immunodeficiency                                                  | RDC-PID |
| Primary Sclerosing Cholangitis                                            | RDC-PSC |
| Primary Sjogrens Syndrome                                                 | RDC-PSJ |
| Rare Inherited Neurological Disorders                                     | RDC-IND |
| Refractoriness First-line Treatment for Blood Cell Autoimmunity           | RDC-BCA |
| Renal Systemic Lupus Erythematosus                                        | RDC-RLE |
| SAPHO (synovitis, acne, pustulosis, hyperostosis, osteitis)               | RDC-SAP |
| Septo-optic Dysplasia                                                     | RDC-SOD |
| Severe Hyperemesis Gravidarum                                             | RDC-SHG |
| Stem Cell and Myeloid Disorders                                           | RDC-SMD |
| Steroid Resistant Nephrotic Syndrome                                      | RDC-SNS |
| Systemic Autoinflammatory Disorders - AOSD and uSAID                      | RDC-SAD |
| Systemic Sclerosis                                                        | RDC-SSC |
| Turner's Syndrome                                                         | RDC-TUS |
| Type 2 Diabetes in Children                                               | RDC-PDC |
| Wolfram Syndrome                                                          | RDC-WOL |

## Notes

This list will be updated and re-circulated to incorporate changes and / or new projects.

\*\*\*Projects recruiting through specific sites/centres.