# Conditions Cured:

### Acute Leukaemias
- Acute Lymphoblastic Leukaemia (ALL)
- Acute Myelogenous Leukaemia (AML)
- Acute Biphenotypic Leukaemia
- Acute Undifferentiated Leukaemia

### Chronic Leukaemias
- Chronic Myelogenous Leukaemia (CML)
- Chronic Lymphocytic Leukaemia (CLL)
- Juvenile Chronic Myelogenous Leukaemia (JCML)
- Juvenile Myelomonocytic Leukaemia (JMML)

### Myelodysplastic Syndromes
- Myelodysplastic Syndrome (MDS)
- Refractory Anaemia (RA)
- Refractory Anaemia with Ringed Sideroblasts (RARS)
- Refractory Anaemia with Excess Blasts (RAEB)
- Refractory Anaemia with Excess Blast in Transformation (RAEB-T)
- Chronic Myelomonocytic Leukaemia (CMML)

### Haematopoietic Cell Disorders
- Aplastic Anaemia (Severe)
- Congenital Dyserythropoietic Anaemia
- Fanconi Anaemia
- Paroxysmal Nocturnal Haemoglobinuria (PNH)
- Pure Red Cell Aplasia
- Acute Myelofibrosis
- Agnogenic Myeloid Metaplasia (Myelofibrosis)

### Lymphoproliferative Disorders
- Non-Hodgkin's Lymphoma
- Hodgkin's Diseases

### Phagocyte Disorders
- Chediak-Higashi Syndrome
- Chronic Granulomatous Disease

### Plasma Cell Disorders
- Multiple Myeloma
- Plasma Cell Leukaemia
- Waldenstrom's Macroglobulinemia

### Other Malignancies
- Ewing Sarcoma
- Neuroblastoma
- Testicular Cancer

### Neurological Disorders
- Cerebral Palsy

### Inherited Metabolic Disorders
- Aspartylglucosaminuria
- Adrenoleukodystrophy
- Alpha-mannosidosis
- Congenital Erythropoietic Porphyria
- Fucosidosis
- Gangliosidosis
- Gaucher's Disease
- Hunter Syndrome
- Hurler Syndrome
- Hurler-Scheie Syndrome
- I-cell Disease
- Infantile Ceroid Lipofuscinosis
- Krabbe Disease
- Lesch-Nyhan Syndrome
- Metachromatic Leukodystrophy
- Maroteaux-Lamy Syndrome
- Morquio Syndrome
- Mucopolysaccharidosis
- Niemann-Pick Disease
- Sandhoff Disease
- Sanfilippo Disease
- Sialidosis
- Tay Sachs Disease
- Wolman Disease

### Histiocytic Disorders
- Familial Erythrophagocytic Lymphohistiocytosis
- Histiocytosis-X
- Haemophagocytosis

### Inherited Immune System Disorders
- Ataxia-Telangiectasia
- Kostmann Syndrome
- Myelokathexis
- Leukocyte Adhesion Deficiency
- DiGeorge Syndrome
- Bare Lymphocyte Syndrome
- Omenn Syndrome
- Severe Combined Immunodeficiency (SCID)
- SCID with Adenosine Deaminase Deficiency
- SCID with absence of T & B Cells
- SCID with absence of T Cells, Normal B Cell
- Common Variable Immunodeficiency
- Wiskott-Aldrich Syndrome

### Inherited Haemoglobinopathies
- Thalassaemia
- Sickle cell disease

### Inherited Platelet Abnormalities
- Amegakaryocytosis / Congenital Thrombocytopenia