Study Number ☐☐☐☐☐☐☐

Date of Initial Echo (DD/MMM/YYYY) ☐☐/☐☐☐/☐☐☐☐

Initial Echo CD Received?
☐No
☐Yes
☐Unknown

Initial Echo Report Received?
☐No
☐Yes
☐Unknown

Date of Post-op Echo (DD/MMM/YYYY) ☐☐/☐☐☐/☐☐☐☐

Post-op Echo CD Received?
☐No
☐Yes
☐Unknown

Post-op Echo Report Received?
☐No
☐Yes
☐Unknown

Date of 1 Year Follow-up Echo (DD/MMM/YYYY) ☐☐/☐☐☐/☐☐☐☐

1 Year Follow-up Echo CD Received?
☐No
☐Yes
☐Unknown

1 Year Follow-up Echo Report Received?
☐No
☐Yes
☐Unknown

Timing of Diagnosis
☐Prenatal
☐Perinatal
☐Postnatal
☐Not in Report

Date of Diagnosis (DD/MMM/YYYY) ☐☐/☐☐☐/☐☐☐☐

First Admission to CHSS Institution Weight (kg) _____

First Admission to CHSS Institution Height (cm) _____

First Admission to CHSS Institution BSA (m2) _____

Gestational Age (weeks) _____

Birth Weight (kg)_____

Birth Height (cm)_____

Birth BSA (m2)_____

Institutional Diagnosis:
☐Balanced
☐Unbalanced
☐Not Stated

If Unbalanced, then
☐Right Dominant
☐Left Dominant

Intended Surgical Strategy
☐1
☐1.5
☐2
☐PA band and wait
☐Palliative shunt

Actual Surgical Strategy
☐1
☐1.5
☐2
☐PA band and wait
☐Palliative shunt
☐Not in Report

Associated Non Cardiac Anomalies
☐None
☐Anal Atresia (imperforate anus)
☐Congenital Diaphragmatic Hernia
☐Gastroschisis
☐Hirschprungs
☐Intestinal Malrotation
☐Omphalocele
☐Transesophageal Fistula

Genetic/Chromosomal Anomalies
☐No
☐Yes
☐Not in Report

☐Not Applicable

If yes, specify

☐Mucopolysaccharidosis type IH (Hurler syndrome)
☐Mucopolysaccharidosis type IH/S (Hurler-Scheie syndrome)
☐Mucopolysaccharidosis type II (Hunter syndrome)
☐Mucopolysaccharidosis type IS (Scheie syndrome)
☐Noonan Syndrome
☐Patau Syndrome (Trisomy 13)
☐Rethore Syndrome (Trisomy 9)
☐Rubella
☐Rubinstein-Taybi Syndrome
☐Short QT Syndrome
☐Situs Inversus
☐Smith-Lemli-Opitz Syndrome
☐Turner Synrdome (45XO)
☐VACTERL syndrome (VACTER/VATER/VATERR Syndrome)
☐VACTERL-H syndrome (VATER association with hydrocephalous) (Briard-Evans Syndrome)
☐Warkany Syndrome (Trisomy 8)
☐Williams Syndrome (Williams-Beuren Syndrome)
☐Wolff-Parkinson-White Syndrome (WPW Syndrome)
☐Wolf-Hirschhorn Syndrome
☐Other syndromic abnormality
☐No chromosomal abnormality identified
☐Alagille syndrome (intrahepatic biliary duct agenesis)
☐Apert syndrome
☐Brugada syndrome (Sudden unexplained nocturnal death syndrome) (SUNDS)
☐Cardiofaciocutaneous syndrome
☐Carpenter syndrome
☐Cat-eye syndrome
☐CHARGE Association
☐Cornelia de Lange syndrome
☐Costello syndrome
☐Cri-du-chat syndrome
☐Deletion 10p syndrome
☐Deletion 8p syndrome
☐DiGeorge syndrome (velocardiofacial syndrome) (conotruncal anomaly face syndrome) (22q11 deletion)
☐Down syndrome (Trisomy 21)
☐Ellis-van Creveld syndrome

☐Goldenhar syndrome
☐Heterotaxy syndrome
☐Heterotaxy syndrome, Asplenia syndrome
☐Edwards syndrome (Trisomy 18)
☐Heterotaxy syndrome, Polysplenia syndrome
☐Holt-Oram syndrome
☐Jacobsen syndrome
☐Kabuki syndrome
☐Klinefelter syndrome (XXY Syndrome)
☐LEOPARD syndrome
☐Loeys-Dietz syndrome
☐Long QT syndrome (Ward Romano syndrome)
☐Marfan syndrome
☐Marfan-like syndrome
☐Fetal alcohol syndrome (FAS)
☐Fetal drug exposure

If other anomalies, please specify: