## Serum insulin like Growth Factor-I and Growth Hormone Levels in Children with Congenital Heart Disease, Relationship with Nutritional Status, Cyanosis and Left Vent Functions

Thesis

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### List of Abbreviations

**ACEI** : Angiotension converting enzyme inhibitor

A<sub>m</sub> wave : Late diastolic mitral annular velocity

**AO** : Aorta

A wave : Late diastolioc waveASD : Atrial septal defectAV : Atrio-ventricular valve

**AV** : Atrio-ventriculo

AVSD : Atrio-ventricular septal defect
AVVR : Atrio-ventricular venous return

Awave : Early diastolic
BMI : Body Mass Index

CAHD : Congenital acyanotic heart diseaseCCHD : Congenital cyanotic heart disease

CHDs : Congenital heart diseasesCHF : Congestive heart failureDHEA : Dehydroepiandrosterone

**DT** : Deceleration time

**D-TGA** : Dextro-transposition of great arteries

**EF** : Ejection fraction

**E**<sub>m</sub> wave : Early diastolic mitral annular velocity

E wave : Early diastolic wave FS : Fractional shortening

**GHD** : Growth hormone deficiency

**GH-RH** : Growth hormone-releasing hormone

HGH : Human growth hormoneIGF-1 : Insulin-like growth factor-1

IGF-1R : Insulin-like growth factor-1 receptorIGF-BPs : Insulin-like growth factor binding protein

**IQ** : Intelligence quotient

IVCT : Isovolumetric contraction time
 IVRT : Iso volumetric relaxation time
 L-TGA : Left- transposition of great arteries

**LV** : Left ventricular

LVEDd : Left ventricular end diastolic diameterLVESd : Left ventricular end systolic diameter

**LVMI** : Left ventricular mass index

**LVMPI** : Left ventricular myocardial performance index

**MPI** : Myocardial performance index

MR : Mitral regurge

MRI : Magnetic resonance imaging

PDA : Patent ducts arteriosus
PS : Pulmonary stenosis

**PVR** : Pulmonary vascular resistance

**PWT** : Posterior wall thickness

**rhGH** : Recombinant human growth hormone

**RV** : Right ventricle

**RVMI** : Right ventricular mass index

**SD** : Standard deviation

 $S_m$  wave : Peak systolic mitral annular velocity

**SVR** : Systemic vascular resistance

**SWT** : Septal wall thickness

**TDI** : Tissue Doppler imaging

**TGA** : Transposition of great arteries

**TOF** : Tetralogy of Fallot

**VSD** : Ventricular septal defect



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#### Introduction

Congenital heart diseases refers to any anatomic defect in the heart and major blood vessels that is present at birth (*Smith*, 2001).

Cardiac abnormalities occur with an incidence of 0.8 per 1000 live birth and represent 25% of all congenital malformations (*Nemer et al.*, 2006). Congenital cardiovascular malformations (CCVM) present any society with an enormous burden of grief and expense; about eight percent of all deaths during the first year of life are due to CCVM and account for a third or more of infants deaths due to birth defects, more than that for any other congenital anomaly (*Bailey and Berry*, 2005).

Failure to thrive and protein- energy malnutrition are well-recognized complications of Congenital heart diseases (CHDs) (*Soliman et al.*, 2001).

They are related to repeated respiratory infections, increased Oxygen consumption rate and changes induced by chronic hypoxia (*Soliman et al.*, 2001). Hypoxia causes feeding difficulties, insufficient caloric intake, intestinal anoxia and venous congestion (*Soliman et al.*, 2001), which causes hypermetabolism, reduction in nutrient ingestion, intestinal malabsorption of nutrient and malnutrition (*Varan*, 1999).

Insulin- like growth factors (IGF) are growth hormones related peptides that play a major role in anabolic and mitogenic activities (*Soliman*, 2000). The combination of decreased IGF-1

level and increased basal GH levels is associated with malnutrition and hypermetabolic states (*Bentham*, 1993).

IGF are pronounced to have also effects in the pathogenesis of failure to thrive seen in CHD (*Soliman et al.*, 2001).

At present, there is a growing body of evidence implicating growth hormone (GH) and/or insulin-like growth factor-1 (IGF-1) in the intricate cascade of events connected with the regulation of heart development and hypertrophy (*Lombardi*, 1997).

### **Aim of the Work**

The aim of this work is to evaluate serum insulin- like growth factor IGF-1 and GH levels in children with Congenital heart diseases (CHDs) and to determine their relationship with nutritional status and left ventricular global function in these patients.

# Congenital Heart Diseases (CHDS)

#### **Definition:**

Congenital heart diseases (CHDs) refer to any abnormality in cardiocirculatory structure or function that is present at birth even if it is discovered later (*Park*, 2008).

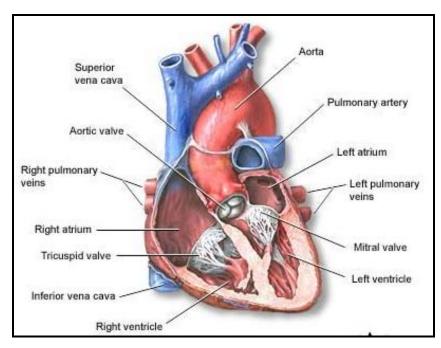


Fig. (1): Structure of a normal human heart (Park, 2008).

#### The embryology of congenital heart disease:

The embryological development of the heart is an awesome and complex process that occurs between third and ninth weeks of gestation (*Suddaby*, 1999).

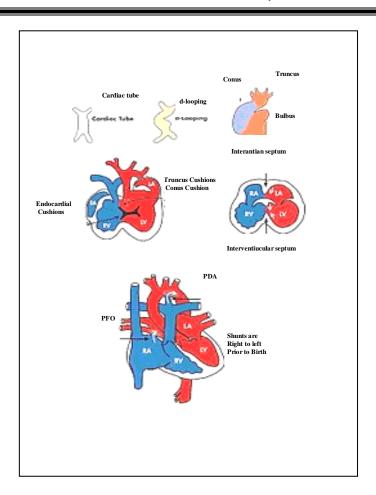


Fig.(2): CVS embryology (Heiden, 2010).

The heart begins as a single tube that septates into two tubes and begins to twist rightward onto itself, called "d" looping. This tube will form an "S" shaped structure that will eventually form all the structure of the heart and begin pumping blood by the fourth week of life. The superior (top) portion of the tube will start to balloon out and will begin to form the atria. Meanwhile, the inferior (bottom) portion of the tube will balloon out and begin to form the ventricles .The middle portion of this tube will