

Journal of Clinical & Experimental **Cardiology**

Open Access

Particular Aspects in Pediatric Congestive Heart Failure

Angela Butnariu¹ and Gabriel Samaşcal^{2*}

¹Department of Pediatrics III, Iuliu Haţieganu, University of Medicine and Pharmacy Cluj-Napoca, Romania ²Department of Immunology, Iuliu Haţieganu, University of Medicine and Pharmacy Cluj-Napoca, Romania

Congestive heart failure (CHF) assumes high material costs and a significant death rate. In the U.S., CHF is a major public health problem, with more than 900.000 hospitalizations per year and more than 250.000 deaths annually. Most CHF cases occur in adults, so that the statistics presented, primarily interested in the adult population. In children, the scope of the problem is less well defined. Data from the American Registry of pediatric cardiomyopathy suggest an annual incidence of 1.13 per 100.000 children cardiomyopathy, most developing CHF. The mortality rate at 2 years after diagnosis is 13.6 % in the forms of dilated cardiomyopathy [1].

CHF in children presents important charactersitic features from the adult congestive failure, from the physiopathological and mostly from the etiopathogenical point of view. While the congestive heart failure at adult age is due to ischemia in 60-70 % of cases, congestive heart failure at children is, in most cases, a consequence of either congenital heart diseases (CHD) which remained unoperated, undergone a palliative operation or presented postsurgery complications or of one cardiomyopathy. Chronic CHF may occur in children with biventricular circulation (systolic or diastolic dysfunction), into cardiac structural abnormalities with right ventricle as systemic ventricle and in the so-called univentricular heart.

We mark the fact that the incidence of the congenital heart diseases is of 8/1000 alive new-born children and that, in Romania, are born, annually, between 800 and 1300 children having a CHD. Out of these more then 20 % develop at least once congestive cardiac failure during their first childhood, in accordance with our information.

So far have been developed excellent guidelines for CHF diagnosis and therapy in adult patients, but the same can not be said about pediatric CHF. Given the significant differences between adults, children and adolescents with CHF, there is little reason to believe that the adult guidelines are directly applicable to children. However, many attitudes in the diagnosis and therapy of pediatric CHF are extrapolated from adult. Most of the studies concerning the paediatric CHF diagnosis and therapy are extended to adult cardiac failure although it is proven the absence of complet propinquity.

Diagnosis and prognosis issue: So far there are few criterions to clearly define the paediatric CHF, especially at small age; its assessment and prognosis evaluation. At present, the pediatric CHF diagnosis is based especially on clinical assessment, and on ultrasound functional findings. The latter, is mainly used to evaluate the left ventricular (LV) functional parameters since the diastolic disorder is difficult to assess

Clinically, the functional evaluation of CHF at children age is made accordind to the Ross classification (an adaptation of the NYHA classification for children). Ross classification allows the evaluation of the state of severity at infant and small child. It is used on daily basis by the Canadian Cardiovascular Society as the official CHF evaluation system in the case of pediatric patients and it has been assumed by numerous pediatric cardiologists, including us. It is important to emphasize that this classification system includes, also, the growth failure, as another indicator in the assessment of the child's CHF.

The echocardiographic parameters most frequently used for

the global evaluation of the left ventricular systolic function are: the shortening fraction, the ejection fraction measured in TM and 2D or through Doppler examination. These indicators depend on the cardiac loading conditions and do not reflect the myocardic fibre intrinsic value.

The increasing filling pressures in the left ventricule represents the common path towards CHF, no matter its etiology. For this reason, the left ventricle diastolic function is an important step in the diagnosis and therapy assessment process. The 2D and TM findings have a low specificity and, as a result, are no loger in use. During the last years, the Doppler ultrasound examination has proved to be very useful in noninvasive evaluation of the left ventricular diastolic function and it is, in the same time, very accesible in the adult cardiology as well. Nowadays, most of the information regardind the left ventricular protodiastola is drawn from the transmitral tide and the myocardic velocity evaluation using tissuue Doppler [2,3]. The tissue Doppler is still rarely used in paediatrics and the physiological standards have not been established yet. However, the few publications on this subject consider tissue Doppler imaging as rewarding in paediatric cardiology [4,5] and , for all that, one of our research objectives is to evaluate, using tissue Doppler, the left ventricular diastolic function in children with CHF. Various biological markers have been considered, during the past few years, as possible indicators of CHF. The most important of them are the compounds derived from compensatory neurohumoural activity such as natriuretic peptides. Its blood concentration level varies in correlation with the overloading volume or pressure degree in the heart.

Natriuretic Peptides in Paediatric Practice

The plasmatic natriuretic peptides concentration is normally high during the first days of life and its values decrease then, gradually, remaining relatively constant during childhood [6].

Children having a congenital heart disease which leads to increased intracardiac volumes and pressures, also present levels of the mentioned neurohormones and these values are inverse proportional to the heart's functional capacity. Limited studies suggest that these hormones may even be markers in the paediatric cyanotic, obstructive and inflamatory diseases, as well as paediatric heart failure [10]. They could also be used in the postoperative evaluation of CHD in infants [11,12].

*Corresponding author: Dr. Gabriel Samasca - Department of Immunology, Croitorilor Street, 19-21 No, Iuliu Haţieganu, University of Medicine and Pharmacy Cluj-Napoca, Romania, E-mail: Gabriel.Samasca@umfcluj.ro

Received December 24, 2011; Accepted January 20, 2012; Published January 24, 2012

Citation: Butnariu A, Samaşcal G (2012) Particular Aspects in Pediatric Congestive Heart Failure. J Clinic Experiment Cardiol 3:e104. doi:10.4172/2155-9880.1000e104

Copyright: © 2012 Butnariu A, et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Authors	Nornal values of NT-proBNP (pg/ml)
Mir et al. [7]	103,1 (0-9 years; n = 79)
Dulac et al. [8]	< 100 (5 days-11 years; n=34)
Wu et al. [9]	556,5 (2 months-2 years; n=20)
Butnariu ş.c.	148 (0-3 years; n=16)

Legend: n-number of patients

Table 1: Natriuretic peptide normal values in pediatrics.

Myocardic injury markers: In paediatrics, ischemia is not suspected at a patient with CHF, in contrast with the adult's situation. Increased cardiac troponin is a specific marker of the myocardic injury at both children and adult pacients, but, in paediatrics, it is not currently used [13]. At some children the troponin level should be determined since some of the cardiomyopathies associated with CHF and even some congenital heart diseases are accompanied by injuries of the myocytes.

Inflammatory markers: It has been proven the release of different inflammatoty markers in CHF [14,15]. Their correlation with the CHF etiology, with NYHA functional class and with the left ventricle function echocardiographic paramaters is not well studied in children.

Therapeutical Issues in the Paediatric CHF

There is a large literature addressing CHF treatment for adult patients, with a much smaller literature concerning CHF therapy in children. Excellent guidelines for adult patient have been published, but given the significant differences between adult and pediatric patient with HF, there is little reason to believe that these guidelines are directly applicable to children [16]. Most pediatric patients with CHF can be treated with a combination of three types of drugs: an inhibitor of angiotensin converting enzyme (ACEI), a loop diuretic and a antialdosteronic. The angiotensin conversion enzyme inhibitors along with the diuretics represent the first line medication. If needed, digitalics can be associated. Beta-blokers such as carvedilol are recommended in paediatric CHF which does not improve under conventional therapy with ACEI, diuretics, digitalics. For special situations, generally using other drugs or circulatory support devices. Drug therapeutic attitude should take into account the type of HF, chronic or acute and also the etiological aspects and hemodynamics [17,18].

References

- Rosenthal D, Chrisant M, Edens E, Mahony L, Canter C, et al. (2004) International Society for Heart and Lung Transplantation: Practice guidelines for management of heart failure in children. J Heart Lung Transplant 23: 1313-1333.
- Garcia MJ, Thomas JD, Klein AL (1998) New Doppler echocardiographic applications for the study of diastolic function. J Am Coll Cardiol 32: 865-875.
- 3. Ommen SR, Nishimura RA, Appleton CP, Miller FA, Oh JK, et al. (2000)

Clinical utility of Doppler echocardiography and tissue Doppler imaging in the estimation of left ventricular filling pressures: a comparative simultaneous Doppler-catheterization study. Circulation 102: 1788-1794.

- Rumeau P, Acar P, Paranon S, Bassil R, Cournot M, et al. (2007) Evaluation of left ventricular diastolic function in children by Doppler tissue imaging. Arch Mal Coeur Vaiss 100: 405-410.
- Friedberg MK, Silverman NH, Dubin AM, Rosenthal DN (2007) Mechanical dyssynchrony in children with systolic dysfunction secondary to cardiomyopathy: a Doppler tissue and vector velocity imaging study. J Am Soc Echocardiogr 20: 756-763.
- Soldin SJ, Soldin OP, Boyajian AJ, Taskier MS (2006) Pediatric BNP and NT-BNP reference intervals. Clin Chim Acta 1: 304-308.
- Mir TS, Flato M, Falkenberg J, Haddad M, Budden R, et al. (2006) Plasma concentrations of N-terminal brain natriuretic peptide in healthy children, adolescents, and young adults: effect of age and gender. Pediatr Cardiol 27: 73-77.
- Dulac Y, Zabalawi A, Taktak A, Plat G, Bassil R, et al. (2006) B-natriuretic peptide and cardiological emergencies in childhood. Arch Mal Coeur Vaiss 99: 477-481.
- Wu YR, Chen SB, Huang MR, Zhang YQ, Sun K, et al. (2005) Diagnostic value of plasma concentration of pro-brain natriuretic peptide in congestive heart failure in pediatric patients with ventricular septal defects. Zhonghua Er Ke Za Zhi 43: 161-164.
- Nir A, Nasser N (2005) Clinical value of NT-ProBNP and BNP in pediatric cardiology. J Card Fail 11: S76-S80.
- Cannesson M, Bionda C, Gostoli B, Raisky O, di Filippo S, et al. (2007) Time Course and Prognostic Value of Plasma B-type Natriuretic Peptide Concentration in Neonates Undergoing the Arterial Switch Operation. Anesth Analg 104: 1059-1065.
- Walsh R, Boyer C, LaCorte J, Parnell V, Sison C, et al. (2008) N-terminal B-type natriuretic peptide levels in pediatric patients with congestive heart failure undergoing cardiac surgery. J Thorac Cardiovasc Surg 135: 98-105.
- Muniz A E (2004) Elevated Cardiac Troponin I in a 9-Week-Old Infant. Ped Emerg Care 20: 674-676.
- Sanchez-Lázaro IJ, Almenar L, Reganon E, Vila V, Martínez-Dolz L, et al. (2007) Inflammatory markers in stable heart failure and their relationship with functional class. Int J Cardiol 129: 388-393.
- Vila V, Martinez-Sales V, Almenar L, Sanchez-Lazaro I, Villa P, et al. (2007) Inflammation, endothelial dysfunction and angiogenesis markers in chronic heart failure patients. Int J Cardiol 130: 276-277.
- Heart Failure Society of America (HFSA) practice guidelines (1999) HFSA guidelines for management of patients with heart failure caused by left ventricular systolic dysfunction--pharmacological approaches. J Card Fail 5: 357-382.
- 17. James N, Smith M (2005) Treatment of heart failure in children. Current Paediatrics 7: 539-548.
- Margossian R (2008) Contemporany management of pediatric heart failure. Expert Rev Cardiovasc Ther 2: 187-197.