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KEYWORDS Psychosocial adjustment; Congenital heart	Abstract Objectives: We aimed to study the psychosocial adjustment (PSA), psychiatric morbidity and quality of life of adolescents and young adults with congenital heart disease (CHD) to deter- mine which demographic and clinical variables negatively affect adjustment and which increase
disease; Psychiatric morbidity; Quality of life	resilience. <i>Methods:</i> The study included 74 patients with CHD, 41 male and 33 female, aged between 12 and 26 years (mean 18.76 ± 3.86). Demographic information and a complete clinical history were obtained. The participants were interviewed regarding social support, family environment,
	self-image and physical limitations. A standardized psychiatric interview was conducted, and self-report questionnaires were administered for assessment of PSA (Youth Self Report and Adult Self Report) and quality of life (World Health Organization Quality of Life – Short Version). A caregiver completed an observational version of the PSA questionnaire (Child Behavior Checklist
	or Adult Behavior Checklist). <i>Results</i> : Female participants showed more feelings of anxiety and depression (U=952.500; p=0.003), thought problems (U=929.500; p=0.005) and aggressive behavior (U=999.000; p=0.000). They also showed a higher rate of psychopathology. Patients with complex forms of CHD reported more thought problems (U=442.000; p=0.027) and internalization (U=429.000;

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p=0.021). Compared to the Portuguese population as a whole, participants showed better quality of life in the domains of social relationships (t=2.333; p=0.022) and environment (t=3.754; p=0.000). Patients who had undergone surgery had worse quality of life in physical terms (t=-1.989; p=0.050), social relationships (t=-2.012; p=0.048) and general quality of life (U=563.000; p=0.037), compared to those who were not operated. Better social support was associated with better quality of life in physical terms (t=3.287; p=0.002) and social relationships (t=3.669; p=0.000). Better school performance was also associated with better overall quality of life (U=457.000; p=0.046), less withdrawn behavior (U=812.500; p=0.031), fewer feelings of anxiety and depression (U=854.000; p=0.009), fewer attention problems (U=903.500; p=0.001), and lower scores for internalization (U=817.000; p=0.029) and externalization (U=803.500; p=0.042). Physical limitations had a detrimental effect on quality of life (U=947.500; p=0.001).

Discussion: Female participants were more prone to worse psychological adjustment and to psychopathology. Patients with complex forms of CHD showed worse PSA, as they need regular care, which restricts social contact with peers and family and integration in school and leisure activities. Patients who had undergone surgery showed worse quality of life as they often have long hospital stays, during which social activities are restricted, making it more difficult for them to develop a good social support network. They require close medical care, and the restrictions on their activities may be life-limiting. Their sense of survival may also be threatened.

Conclusions: Patients with CHD appear to be more prone to psychopathology and female patients are more likely to show worse PSA. Social support was shown to play a crucial role in buffering stress and promoting patients' adjustment.

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Ajustamento psicossocial, morbilidade psiquiátrica e qualidade de vida em adolescentes e jovens adultos com cardiopatias congénitas

Resumo

Objetivos: Caracterizar a população de adolescentes e jovens adultos com cardiopatias congénitas (cc) ao nível do seu ajustamento psicossocial e morbilidade psiquiátrica, bem como avaliar a qualidade de vida (QV), para perceber quais são as variáveis com impacto na vida e na adaptação à doença.

População e métodos: Participaram 74 pacientes com CC, sendo 41 do sexo masculino e 33 do sexo feminino, com idades entre 12-26 (média = 18,76 \pm 3,86). Foram recolhidos os dados demográficos e clínicos mais relevantes dos pacientes e foi aplicado um conjunto de instrumentos, incluindo uma entrevista semiestruturada sobre tópicos de suporte social, estilo de educação na família, autoimagem e limitações físicas, uma entrevista psiquiátrica estandardizada e questionários para avaliar o ajustamento psicossocial na forma de autorrelato (YSR e ASR) e relato dos cuidadores (CBCL e ABCL), bem como um questionário de avaliação da qualidade de vida (WHOQOL-BREF).

Resultados: O sexo feminino relatou mais ansiedade/depressão (u = 952,500; p = 0,003), alterações do pensamento (u = 929,500; p = 0,005) e comportamento agressivo (u = 999,000; p = 0,000). Pacientes com CC complexas relataram mais alterações de pensamento (u = 442,000; p = 0,027) e internalização (u = 429,000; p = 0,021). Comparativamente com a população portuguesa, os nossos participantes apresentaram melhor QV nos domínios relações sociais (t = 2,333; p = 0,022) e ambiente (t = 3,754; p = 0,000). Os pacientes que não foram submetidos a intervenções cirúrgicas revelaram melhor QV nos domínios físico (t = -1,989; p = 0,050), de relações sociais (t = -2,012; p = 0,048) e QV geral (u = 563,000; p = 0,037). A presença de um melhor suporte social está relacionada com uma melhor QV nos pacientes em todos os domínios avaliados, com destaque para o físico (t = 3,287; p = 0,002) e relações sociais (t = 3,669; p = 0,000). Um melhor desempenho académico encontrava-se também associado com uma melhor perceção de QV global (u = 457 000; p = 0,046), com menos tendência ao isolamento (u = 812.500; p = 0.031), menos sentimentos de ansiedade e de depressão (u = 854 000; p = 0.009), menos problemas de atenção (u = 903 500; p = 0,001), e valores mais baixos de internalização (u = 817 000; p = 0,029) e de externalização (u = 803 500; p = 0,042). As limitações físicas tinham um efeito nocivo no QV (u = 947 500; p = 0,001).

Discussão: As participantes do sexo feminino evidenciaram pior ajustamento psicossocial e maior propensão à psicopatologia. As CC complexas exigem mais cuidados regulares, limitando o contacto social com pares, a integração escolar e de lazer, e dificultando a adaptação.

PALAVRAS-CHAVE

Ajustamento psicossocial; Cardiopatias congénitas; Morbilidade psiquiátrica; Qualidade de vida



A realização de cirurgias influencia a QV dos pacientes, pela exigência de mais cuidados médicos, induzindo maiores limitações na sua vida diária, o que dificulta a aquisição de uma rede de suporte social satisfatória.

Conclusões: Os pacientes com CC evidenciaram uma maior tendência a apresentar psicopatologia, sendo o sexo feminino mais predisposto a sofrer com os efeitos do desajustamento psicossocial. O suporte social revelou um papel preponderante no ajustamento dos doentes. © 2011 Sociedade Portuguesa de Cardiologia. Publicado por Elsevier España, S.L. Todos os direitos reservados.

Introduction

Congenital heart disease (CHD) is the result of abnormalities that occur during gestation leading to malformations of the heart or blood vessels.¹

Developments in medical and surgical treatment have improved survival, and 85–90% of children with CHD now live to adulthood with a reasonable quality of life.¹⁻³

As advances in treatment at various levels have enabled CHD patients to live increasingly longer and with fewer limitations, the question of their adjustment to their condition has attracted increasing attention.⁴ It is thus essential to understand the effects of CHD at all levels, as the literature shows conflicting results.⁵⁻⁸

While some studies have reported a tendency for behavioral and emotional problems in these patients, ^{5,9,10} others have revealed adaptive psychological functioning in adults with CHD.¹¹ Similarly, in some studies more severe CHD appears to be associated with worse psychosocial adjustment and psychiatric morbidity, ^{6,9} but others do not support this finding.⁹

There have also been conflicting results in terms of quality of life, with some studies reporting worse psychological well-being and quality of life compared to the healthy population,^{4,12} and others finding no correlation between the presence of CHD and lower quality of life, with similar results to those found in healthy subjects.^{13,14}

This study aimed to assess psychosocial adjustment, psychiatric morbidity and quality of life in a population of adolescents and young adults with CHD, with or without surgical repair, in order to clarify the impact of certain demographic and clinical variables on increasing resilience and adaptation or worsening adjustment and well-being.

Methods

The study included 74 patients with CHD, who were approached to participate in the study on a single occasion during attendance at the pediatric cardiology or cardiology outpatient clinics. Adult patients or caregivers in the case of minors gave their informed consent after all aspects of the study had been explained.

Of the 74 patients, 41 were male and 33 were female. The mean age of the participants was 18.76 ± 3.86 years (range 12–26).

CHD was complex in 27, moderate in 13 and mild in 34. With regard to residual lesions, four had severe, 18 moderate and 52 mild lesions. Seventeen participants (23%) had been diagnosed with at least one form of psychopathology during their lives.

Of the total study population, 40 had been diagnosed during the neonatal period, 28 in the first year of life, three between the ages of one and three, and three between the ages of three and six.

Of the participants who had undergone surgery, the first intervention had been performed during the neonatal period in two, in the first year of life in 24, between the ages of one and three in 15, between the ages of three and six in 16, and between the ages of six and 12 in three.

With regard to level of education, nine had completed the second cycle, 27 the third cycle, 33 secondary education, and five had a university degree. Most participants (69) were single, two were married, two were in a civil union and one was divorced.

Sixty-four caregivers accompanying patients also participated in the study by completing a questionnaire; 10 refused to participate or were not present at the time of application of the protocol and subsequently expressed no desire to participate.

The instruments used in the study included a personal record, a semi-structured interview, a standardized psychiatric interview (Schedule for Affective Disorders and Schizophrenia – Lifetime version [SADS-L]), the Youth Self Report (YSR) and Child Behavior Checklist (CBCL) for patients aged under 18, and the Adult Self Report (ASR) and Adult Behavior Checklist (ABCL) for those aged 18 or over. The short version of the World Health Organization Quality of Life questionnaire (WHOQOL-Bref) was also applied.

Each participant's personal record covered demographic variables, including civil status, educational level and profession, and clinical history, including diagnosis, severity and type of CHD, description of surgery, residual lesions and drug therapy.

The semi-structured interview, consisting of 38 multiplechoice or short-answer questions, covered areas such as social support, family environment, functional limitations, school performance and self-image.

The standardized psychiatric interview SADS-L was used to assess the presence of psychopathology.

YSR and ASR are self-report questionnaires designed to characterize a child's or adult's functioning, respectively. For the purposes of this study, responses were analyzed on the scales related to withdrawal, somatic complaints, anxiety and depression, thought problems, attention problems, rule-breaking behavior, aggressive behavior, internalization and externalization. Responses on the YSR scale related to social problems were not considered since the ASR does not have an equivalent scale.¹⁵ The CBCL and ABCL are designed to be completed by someone who knows the patient well (parent or caregiver), describing their perception of a patient's functioning and behavior.

Since the YSR/ASR and CBCL/ABCL are similar instruments, the scores were combined in order to obtain a larger number of responses and thus greater representativeness, the total scores for each scale being used in the statistical analysis.

WHOQOL-Bref is a quality of life self-report questionnaire that can be applied in healthy subjects or individuals with any type of disease.¹⁶ It consists of 26 questions, with responses rated on a five-point Likert scale, in which 5 corresponds to the highest rating, except in the case of the negatively formulated questions 3, 4 and 26, for which 5 is the lowest rating.

The statistical analysis was performed using the IBM Social Package for the Social Sciences (SPSS), version 19.0. The Student's t test was used for parametric variables and the Mann-Whitney test for nonparametric variables.

Results

Table 1 shows differences in self-reported psychosocial adjustment between the sexes.

Table 2 shows differences in psychosocial adjustment between participants with complex CHD and those with moderate or mild forms.

Table 3 shows differences in psychosocial adjustment between participants who had undergone surgery and those who did not require surgical intervention, as assessed by caregivers.

Table 4 shows differences in psychosocial adjustment between participants with satisfactory school performance compared to those with poor performance.

Table 5 shows the prevalence of diagnosed psychopathology compared to the references values of other studies.

Table 1	Mann-Whitney test for gender differences in self-
reported	psychosocial adjustment.

ASR/YSR	M (n=41) Mean	F (n=33) Mean	U	p
Withdrawal	36.30	38.98	725.500	0.591
Somatic complaints	32.39	43.85	886.000	0.021
Anxiety/ depression	30.77	45.86	952.500	0.003
Thought problems	31.33	45.17	929.500	0.005
Attention problems	33.41	42.58	844.000	0.067
Rule-breaking behavior	36.78	38.39	706.000	0.744
Aggressive behavior	29.63	47.27	999.000	0.000
Internalization	31.18	45.35	935.500	0.005
Externalization	34.18	41.62	812.500	0.139

ASR: adult self report; F: female; M: male; U: Mann-Whitney test; YSR: youth self report.

Table 2Mann-Whitney test for differences in self-reportedpsychosocial adjustment according to severity of CHD.

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ASR/YSR	Complex (n=27) Mean	Moderate/ mild (n=47) Mean	U	р
Withdrawal	43.54	34.03	471.500	0.065
Somatic complaints	43.93	33.81	461.000	0.049
Anxiety/ depression	38.59	36.87	605.000	0.740
Thought problems	44.63	33.40	442.000	0.027
Attention problems	35.00	38.94	702.000	0.446
Rule-breaking behavior	42.56	34.60	498.000	0.119
Aggressive behavior	43.22	34.21	480.000	0.081
Internalization	45.11	33.13	429.000	0.021
Externalization	34.85	39.02	706.000	0.422
-				

ASR: adult self report; U: Mann-Whitney test; YSR: youth self report.

It should be noted that the prevalence of psychopathology among female participants was more than twice as high as in males (33.3% and 14.6%, respectively, a statistically significant difference).

Table 6 compares results in the study population with reference values for the Portuguese population as a whole for all quality of life domains.

Table 7 shows differences in quality of life between participants who had undergone surgery and those who did not require surgical intervention.

Table 8 shows differences in quality of life between those with good or poor social support.

Table 9 shows differences in quality of life between participants with satisfactory or poor school performance.

It should be noted that 51.4% of participants had failed exams at school (mean 1.74 ± 0.86 years).

Discussion

In this study, female participants presented significantly more somatic complaints, higher indices of anxiety and depression, thought problems and aggressive behaviors, and a higher degree of internalization, and hence worse psychosocial adjustment.

The gender differences observed in CHD patients may be linked to the presence of a surgical scar on the chest, affecting body image and self-esteem and contributing to feelings of insecurity and uncertainty in personal and sexual relations. Another factor leading to insecurity in these patients may be fears that the disease could affect their reproductive capacity and childbirth.^{5,6,20}

Female patients also tend to be more concerned about their bodies than males,⁵ and be more likely to experience anxiety and depression when faced with situations that hinder interpersonal relationships, reflected in a greater

ABCL/CBCL (CR)	Surgery (n=54)	No surgery (n=11)	U	р
	Mean	Mean		
Withdrawal	32.67	34.64	315.000	0.751
Somatic complaints	30.87	43.45	412.000	0.041
Anxiety/depression	31.35	41.09	386.000	0.117
Thought problems	30.94	43.14	408.500	0.040
Attention problems	33.21	31.95	285.500	0.840
Rule-breaking behavior	32.88	33.59	303.500	0.907
Aggressive behavior	34.06	27.82	240.000	0.317
Internalization	32.70	34.45	313.000	0.779
Externalization	32.96	33.18	299.000	0.972

Table 3 Mann-Whitney test for differences in psychosocial adjustment between patients with and without surgical intervention, as reported by caregivers.

ABCL: adult behavior checklist; CBCL: child behavior checklist; CR: caregiver report; U: Mann-Whitney test.

Table 4Mann-Whitney test for differences in self-reported psychosocial adjustment between participants with satisfactoryvs. poor school performance.

ASR/YSR	Poor school performance (n=26) Mean	Satisfactory school performance (n=48) Mean	U	p
Withdrawal	44.75	33.57	812.500	0.031
Somatic complaints	38.94	36.72	661.500	0.667
Anxiety/depression	46.35	32.71	854.000	0.009
Thought problems	43.52	34.24	780.500	0.070
Attention problems	48.25	31.68	903.500	0.001
Rule-breaking behavior	40.25	36.01	695.500	0.410
Aggressive behavior	41.29	35.45	722.500	0.263
Internalization	44.92	33.48	817.000	0.029
Externalization	44.40	33.76	803.500	0.042

ASR: adult self report; U: Mann-Whitney test; YSR: youth self report.

Table 5	Prevalence of diagnosed psychopathology com-
pared to r	eferences values of various other populations.

Psychopathology	n=74	%
No	57	77
Yes	17	23
Reference values		10 ^a
		25 ^b
		19.4 ^c
		37.9 ^d
		18.1 ^e
		25.2 ^f
		47.4 ^g

^a Percentage of adult population worldwide with mental and behavioral disorders, as assessed by the World Health Organization.¹⁷

^b Percentage of population with some form of mental disorder according to the results of the ESEMeD project in six European countries.¹⁸

c,d,e,f,g Percentage of population with some form of mental disorder in Spain, France, Italy, Germany and the USA, respetively.¹⁹ tendency for internalization.^{5,6} They thus have a higher risk of presenting emotional and behavioral problems than their male counterparts.²¹

Patients with complex CHD more often report somatic complaints and thought problems and have higher indices of internalization than those with moderate or mild CHD. This suggests that adolescents and young adults with complex CHD present worse psychosocial adjustment, which may be related to their need for more medical care throughout their lives, while those with mild or moderate CHD are more able to lead a similar life to healthy individuals of the same age.^{6,22}

Since complex CHD presents a higher risk than mild or moderate CHD, it is understandable that adolescents and young adults with a complex form should report more somatic complaints, as they are more alert to the signs produced by their bodies. This will tend to interfere with their thought processes, making them more anxious about possible complications (anxiety is one component of the internalization scale).

When assessed by caregivers, individuals who had not undergone surgery more often presented somatic complaints and thought problems, which suggests that surgical repair

	Reference values ^a		Study population		t	р
	Mean	SD	Mean	SD		
Physical	77.49	12.27	75.29	13.516	-1.400	0.166
Psychological	72.38	13.50	73.34	13.924	0.590	0.557
Social relationships	70.42	14.54	74.91	16.573	2.333	0.022
Environmental	64.89	12.24	70.65	13.209	3.754	0.000
General	71.51	13.30	72.47	15.89	0.518	0.606

Table 6 Means and standard deviation for the four quality of life domains and general quality of life in the study participants and the Portuguese population as a whole.

^a Portuguese healthy population.¹⁶ SD: standard deviation; t: Student's t test.

Table 7 Student's t test for the four quality of life domains and Mann-Whitney test for general quality of life, comparing patients who had undergone surgery with those who did not.

	Surgery (n=60)		No surgery (n=14)		t	р
	Mean	SD	Mean	SD		
Physical	15.81	2.224	17.06	1.567	-1.989	0.050
Psychological	15.46	2.304	16.71	1.557	-1.928	0.058
Social relationships	15.69	2.634	17.24	2.409	-2.012	0.048
Environmental	15.14	2.207	16.04	1.525	-1.443	0.153
	Mean		Mean		U	р
General	35.12		47.71		563.000	0.037

SD: standard deviation; t: Student's t test; U: Mann-Whitney test.

Table 8Student's t test for the four quality of life domains and Mann-Whitney test for general quality of life, comparingparticipants with good or poor social support.

	Good social support (n=54)		Poor social support (n=20)		t	р
	Mean	SD	Mean	SD	_	
Physical	16.52	2.011	14.77	2.083	3.287	0.002
Psychological	16.16	2.167	14.45	1.935	3.094	0.003
Social relationships	16.62	2.143	14.27	3.145	3.669	0.000
Environmental	15.70	1.901	14.25	2.348	2.725	0.008
	Mean		Mean		U	р
General	41.52		26.65		323.000	0.005

SD: standard deviation; t: Student's t test; U: Mann-Whitney test.

Table 9Student's t test for the four quality of life domains and Mann-Whitney test for general quality of life, comparingparticipants with satisfactory or poor school performance.

	Satisfactory school performance (n=48)		Poor school performance (n=26)		t	р
	Mean	SD	Mean	SD		
Physical	16.42	2.192	15.36	1.968	2.045	0.045
Psychological	16.06	2.218	15.03	2.127	1.950	0.055
Social relationships	16.15	2.327	15.67	3.185	0.752	0.455
Environmental	15.73	2.210	14.52	1.708	2.413	0.018
	Mean		Mean		U	р
General	40.98		31.08		457.000	0.046

SD: standard deviation; t: Student's t test; U: Mann-Whitney test.

plays an important role in improving quality of life in CHD patients. $^{9,23}\!$

Patients with CHD require regular medical care, which means that many have to miss school, which in turn leads to poor performance, anxiety and depression, attention problems, and higher indices of internalization and externalization; this is reflected in our study's results, which show that patients with poor school performance presented worse psychosocial adjustment.^{1,23-26}

With regard to psychiatric morbidity, 23% of our subjects had been diagnosed with a mental disorder. Based on the World Health Organization estimate of 10% for the adult population worldwide, this suggests that adolescents and young adults with CHD are more prone to psychopathology.¹⁷

However, the prevalence of 23% in our study population was comparable to that reported for the general population in six European countries (25%).¹⁸ Similar results were also observed for lifetime prevalence of mental disorders in Spain (19.4%), Italy (18.1%) and Germany (25.2%), although the figures were much higher in France (37.9%) and the USA (47.4%).¹⁹

It would be useful to compare the prevalence of psychopathology among our patients with estimates for the general Portuguese population, but such figures are not yet available, and this is one of the major limitations of our study.

With regard to the gender distribution of psychopathology, its prevalence was more than twice as high in female participants, a finding that is consistent with their worse psychosocial adjustment and with reported prevalences of mental disorders, particularly anxiety and depression, in population studies in various countries.

Our population of CHD patients reported a better quality of life compared to the reference values for the healthy Portuguese population in the social relationships and environment domains, while other authors have found a worse quality of life than for the normal population, particularly in the WHOQOL-Bref physical and general quality of life domains, as well as in the psychological and social domains, albeit less marked.¹²

Individuals who had not undergone any form of surgery reported a better quality of life in physical, social relationships and general terms on self-assessment than those who had. While most CHD patients who undergo surgical repair succeed in functioning normally in most areas of their lives, there are certain developmental issues in cases of complex or moderate disease, which make long-term planning more difficult for these patients.²⁷

Social support was shown to have an important impact on quality of life, individuals with greater support presenting better quality of life in all the domains assessed, particularly in physical, psychological, social relationships, environmental and general terms.

Social support, at the individual, family or community level, has a major impact as a mediator between stress and psychological well-being; patients' perception of its availability is extremely important in counteracting the effects of the stress caused by both physical and mental health problems.²⁸

Individuals with a satisfactory school performance present a better quality of life in physical, environmental and general terms, as well as in psychological adjustment. School performance can be severely affected by CHD, as patients often require hospitalization and may have restrictions on activities with their peers, both of which mean they are regularly excluded from school activities.^{1,29}

Conclusions

The study's results appear to show that adolescents and young adults with CHD are prone to emotional and behavioral problems. Female patients were more likely to have worse psychosocial adjustment, as were those with complex CHD, those who had not undergone surgical intervention (according to caregivers), and those with poor school performance.

With regard to psychiatric morbidity, the data for this population appear to indicate a slightly higher likelihood of suffering some form of mental disorder.

By contrast, the study participants reported a better quality of life than the general Portuguese population, highlighting the importance of social support.

Nevertheless, despite the role of various types of social support, the fact that CHD is diagnosed at an early stage of life means that patients have to draw on all their skills to deal with the challenges of developmental milestones and to live a normal life, albeit with certain limitations, many of which can be overcome by coping strategies.

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Conflicts of interest

The authors have no conflicts of interest to declare.

References

- Nousi D, Christou A. Factors affecting the quality of life in children with congenital heart disease. Health Sci J. 2010;2:94–100.
- Claessens P, Moons P, Casterlé B, et al. What does it mean to live with a congenital heart disease? A qualitative study on the lived experiences of adult patients. Eur J Cardiovasc Nurs. 2005;4:3–10.
- Kasper D, Braunwald E, Fauci A, et al. Harrison medicina interna. 16th ed. Rio de Janeiro: McGraw-Hill Interamericana do Brasil Ltd.; 2006.
- Spijkerboer A, Utens E, de Koning W, et al. Health-related quality of life in children and adolescents after invasive treatment for congenital heart disease. Qual Life Res. 2006;15: 663-73.
- Rijen E, Utens E, Roos-Hesselink J, et al. Longitudinal development of psychopathology in an adult congenital heart disease cohort. Int J Cardiol. 2005;99:315–23.
- Rijen E, Utens E, Roos-Hesselink J, et al. Medical predictors for psychopathology in adults with operated congenital heart disease. Eur Heart J. 2004;25:1605–13.
- Hülser K, Dubowy K, Knobl H, et al. Developmental outcome and psychosocial adjustment in children after surgery for congenital heart disease during infancy. J Reprod Infant Psychol. 2007;25:139–51.

- Utens E, Bieman H, Verhulst F, et al. Psychopathology in young adults with congenital heart disease. Eur Heart J. 1998;19:647–51.
- 9. Latal B, Helfricht S, Fischer J, et al. Psychological adjustment and quality of life in children and adolescents following openheart surgery for congenital heart disease: a systematic review. BMC Pediatrics. 2009;9:1–10.
- Birkeland A, Rydberg A, Hägglö B. The complexity of the psychosocial situation in children and adolescents with heart disease. Acta Paediatr. 2005;94:1495–501.
- Kovacs A, Sears S, Saidi A. Biopsychosocial experiences of adults with congenital heart disease: review of the literature. Am Heart J. 2005;150:193–201.
- Rose M, Köhler K, Köhler F, et al. Determinants of quality of life of patients with congenital heart disease. Qual Life Res. 2005;14:35–43.
- Fekkes M, Kamphuis RP, Ottenkamp J, et al. Health-related quality of life in young adults with minor congenital heart disease. Psychol Health. 2001;16:239–50.
- Moons P, van Deyk K, Marquet K, et al. Individual quality of life in adults with congenital heart disease: a paradigm shift. Eur Heart J. 2005;26:298–307.
- Achenbach TM, Rescorla LA. Manual for the ASEBA adult forms & profiles. Burlington, VT: University of Vermont. Research Center for Children, Youth and Families; 2003. p. 1–12.
- Vaz Serra A, Canavarro M, Simões M, et al. Estudos psicométricos do instrumento de avaliação da Qualidade de Vida da Organização Mundial de Saúde (WHOQOL-Bref) para português de Portugal. Psiquiatria Clínica. 2006;27:41–9.
- 17. World Health Organization. Prevention of mental disorders: effective interventions and policy options. Geneva: World Health Organization; 2004. Available from: http:// www.who.int/mental_health/evidence/en/prevention_of_ mental_disorders_sr.pdf [accessed 02.12.10].
- Alonso J, Angermeyer MC, Bernert S, et al. Prevalence of mental disorders in Europe: results from the European study of the epidemiology of mental disorders (ESEMeD) project. Acta Pyschiatr Scand. 2004;109:21–7.

- Kessler R, Angermeyer M, Anthony J, et al. Lifetime prevalence and age-of-onset distributions of mental disorders in the World Health Organization's World Mental Health Survey Initiative. World Psychiatry. 2007;6:168–76.
- Lyon M, Kuehl K, McCarter R. Transition to adulthood in congenital heart disease: missed adolescent milestones. J Adolesc Health. 2006;39:121–4.
- 21. Bellinger D, Newburger J. Neuropsychological, psychosocial, and quality-of-life outcomes in children and adolescents with congenital heart disease. Prog Pediatr Cardiol. 2010;29: 87–92.
- Casey F, Sykes D, Craig B, et al. Behavioral adjustment of children with surgically palliated complex congenital heart disease. J Pediatr Psychol. 1993;21:335–52.
- 23. Spijkerboer A, Utens E, Bogers A, et al. Long-term behavioural and emotional problems in four cardiac diagnostic groups of children and adolescents after invasive treatment for congenital heart disease. Int J Cardiol. 2008;125:66–73.
- Wernovsky G. Current insights regarding neurological and developmental abnormalities in children and young adults with complex congenital cardiac disease. Cardiol Young. 2006;16:92–104.
- Hövels-Gürich H, Konrad K, Wiesner M, et al. Long term behavioural outcome after neonatal arterial switch operation for transposition of the great arteries. Arch Dis Child. 2002;87:506–10.
- 26. Majnemer A, Limperopoulos C, Shevell M, et al. Developmental and functional outcomes at school entry in children with congenital heart defects. J Pediatr. 2008;153:55-60.
- 27. Foster E, Graham T, Reid G, et al. Task force 2: special health care needs of adults with congenital heart disease. J Am Coll Cardiol. 2001;37:1161–98.
- Tak Y, McCubbin M. Family stress, perceived social support and coping following the diagnosis of a child's congenital heart disease. J Adv Nurs. 2002;39:190–8.
- 29. Birks Y, Sloper P, Lewin R, et al. Exploring health-related experiences of children and young people with congenital heart disease. Health Expect. 2007;10:16–29.