CHARACTERIZATION OF BEHAVIOR PROFILE AND SOCIAL COMPETENCE OF INDIVIDUALS WITH THE DEL22Q11.2 SYNDROME

Caracterização do perfil comportamental e de competência social de indivíduos com a síndrome del22q11.2

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ABSTRACT

Purpose: to characterize the behavioral problems and social competence of individuals with del22q11.2 syndrome and compare them with typically developing individuals, according to information from parents. **Methods:** participated in this study 24 parents of individuals of both sexes between 6 and 18 years, being 12 individuals with the syndrome del22q11.2 (sample group) and 12 individuals with typically developing (control group). The behavioral inventory "*Child Behavior Checklist (CBCL)*" was applied. **Results:** eight of the twelve patients with the syndrome were classified as "clinical" as the scales of behavior and Internalizing Problems. Five of the twelve individuals of sample group were "clinical" as the scales of behavior and Externalizing Problems. The skills of social competence, ten of the twelve individuals sample group were "clinical". **Conclusion:** individuals with the diagnostic of the del22q11.2 syndrome, according with opinion their parents present behavioral and social problems in different degrees of commitment. Comparing the groups, was observed statistically significant differences in variables of externalizing and internalizing behaviors. Therefore, we conclude that the sample group presents more abnormal behavior compared to the control group.

KEYWORDS: Speech, Language and Hearing Sciences; Behavior; Interpersonal Relations; Communication; Social Behavior Disorders; DiGeorge Syndrome

INTRODUCTION

Del22q11.2 syndrome (OMIM # 192430)¹ was first described in the 1950s, by Eva Sedlackova, and recognized in 1978 from the study published by Shprintzen and colleagues. In these studies, the authors described as an etiologic factor of this genetic condition the microdeletion on the long arm of chromosome 22 in the region q11.2², which

leads to a set of clinical features first observed in 12 individuals, hypernasality as a result of multiple defects, including cleft palate, cardiac anomalies, learning disabilities and behavioral disorders³.

This genetic condition has variable frequency, due to some cases are undiagnosed because they have mild phenotypic expression and clinical recognition difficult. Data from the United States reported frequency of 1: 2000 live births. Until the present moment, there are no national epidemiological data. Since this is a condition resulting from microdeletion, it is necessary for the diagnosis of del22q11.2 syndrome, a cytogenetic evaluation often performed by the technique of Fluorescence in situ hybridization (FISH) to detect the absence of chromosomal segment in the region 22q11. 2¹⁻³.

The phenotype of this syndrome presents wide variability among individuals, among the most common the presence of characteristic facial

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features: long face with prominent nose and broad nasal base, with square aspect; jaw alteration; retraction of the mandible; narrow palpebral fissures: minor ear alterations: velopharyngeal insufficiency and cleft palate; renal and skeletal abnormalities: cardiac abnormalities, representing about 90% of the cause of death in this syndrome^{4,5}; delayed acquisition of language⁶; learning problems with significant deficits in tasks of arithmetic⁷; and behavioral disorders8,9.

The behavioral phenotype of del22q11.2 syndrome has been described in the literature with problems with aggression, depression, and difficulty in interacting and similar social problems to disorders of the autistic spectrum^{10,11}.

Psychiatric symptoms, mainly schizophrenic disorders are among the behavioral problems frequently cited as part of the phenotype of del22q11.2^{12,13}. syndrome. The presence of schizophrenic symptoms has attributed to this genetic condition significant value as neurobiological research model of the genetic basis of schizophrenia, which has propelled the growing number of studies aimed at the characterization of the behavioral phenotype of this syndrome⁸⁻¹⁶.

The discussion about aggravation of psychiatric disorders according to age group is still contradictory. Some authors argue that psychiatric disorders did not differ in age in del22g11.28 syndrome, being identified both in children and in older individuals, but there are authors who argue that attention deficit disorder, anxiety, mood instability, hyperactivity, impulsivity, shyness or disinhibition aggravate according to age group¹⁷.

The Child Behavior Checklist (CBCL) is one of the most used tools in the literature on the subject, since it enables, besides identifying problematic behaviors in specific populations, classify such problems due to the characteristics of the complaints in internalizing behaviors (e.g. problems become internalized as depressive thoughts, anxiety and somatic problems) and externalizing behaviors (e.g. related to the social context of individuals, such as social problems, thought problems, attention problems, presence of aggressive behavior and delinquency).

Behavioral studies using the CBCL in the population of individuals with del22q11, highlighted shifts in attention^{18,19}, changes in internalizing behavior and social competence²⁰. In another study, it was found through the application of behavioral inventory CBCL that individuals with the syndrome del22Q11.2 presented commitments in internalizing and externalizing behaviors when compared to a group of children with craniofacial abnormalities (ACF)²¹.

Therefore, this research proposes to characterize the behavioral problems and social skills of individuals with the syndrome del22g11.2 and compare them with individuals with typical development, according to information from parents.

METHODS

This study was approved by the Research Ethics Committee (nº 0706/ 2013) with humans from the institution. All participants signed a Free and Informed Consent before entering the study. It is noteworthy that all ethical criteria were applied respecting the Resolution 196/96 of the National Health Council (CNS/196) on Regulatory Guidelines and Standards for Research Involving Humans, and the recommendations of the Research Ethics Committee.

This is an experimental and cross-sectional study, which compares the information reported by parents of 24 individuals aged 6-18 years old of both genders. These individuals were divided into two subgroups: Sample Group (SG) and the Control Group (CG), namely:

Sample Group (SA): 12 individuals, seven girls (58.3%) and five boys (41.7%) with a diagnosis of del22g11.2 syndrome, positive for the deletion in the 22q11.2 region, using the technique of Fluorescent In situ Hybridization (FISH), patients in the Center for Study of Education and Health (CEES) UNESP Marília / SP or from the Hospital for Rehabilitation of Craniofacial Anomalies (HRAC), Bauru / SP. All patients in the sample perform dental treatment and speech therapy at HRAC and attend regular school.

Control group (CG): 12 individuals, paired with SA by gender and chronological age without clinical diagnosis of del22q11.2 syndrome, with negative history for genetic or psychiatric illness, no history of speech therapy or psycho-pedagogical treatment, and enrolled in public schools in the city of Marilia-SP.

For investigation of behavior problems and social skills, the instrument "Child Behavior Checklist for ages 6-18"22, Brazilian version of a questionnaire for parents²³. The CBCL consists of 138 items: 118 related to evaluation of behavior problems and 20 for the assessment of the child or adolescent social skills.

The behavioral inventory was applied through direct interviews with parents or guardians gathering information from 113 items related to behavior problems. The informant should classify the behavior: false or absent (score = 0); partially or sometimes true (score = 1); and very true or often present (score = 2). Only behavioral problems identified by parents within the past six months prior to the date of the questionnaire²³ were registered.

Behavior problems in CBCL are divided into eight behavioral scales: withdrawn, somatic complaints, anxious and depressed, social problems, thought problems, attention problems, rule-breaking behavior, delinquent and aggressive behavior. For each of these behaviors, the instrument provides a scoring system for rating scales, and the index should be <60 for non-clinical classification, \geq 60 \leq 63 for classification in borderline category and the clinical category> 63^{23} .

The sum of the first three groups form the Internalizing Behavior Scale (withdrawn, somatic complaints, anxious and depressed) and the remainder (social problems, thought and attention problems, delinquent and aggressive behavior) forms the Externalizing Behavior Scale. In the groups, the score for non-clinical category should be <67; for borderline category, $\ge67\le70$; and for clinical category, $>70^{23}$.

Social competence is divided into three scales (activities, social and school) that refer to the involvement of the child or youth in different activities: participation in group organization, relationship with people, independence in playing or working and school performance. Parents identified the

children as "below average," "above average" or "of average"²³.

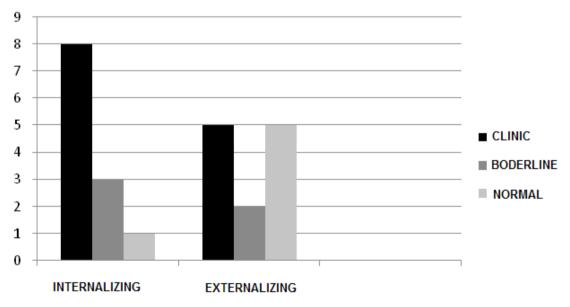
The data were analyzed using the statistical package SPSS (Statistical Package for Social Sciences), in its version 21.0, to obtain the results.

It was used for statistical analysis, the Mann-Whitney Test and Spearman correlation analysis, and the significance value (p) calculated in less than 5% (0,050 / 0,005).

RESULTS

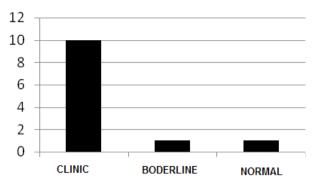
The data from the sample group indicated that on the scales of behavior and Internalizing Problem 66.7% of the individuals showed behavior rated as "clinical", with a lower percentage for "borderline" (25%) and "normal" (8.3%). As for the Externalizing Problems scale, there is equivalence between the behaviors classified as "clinical" and "normal" (41.7% of the individuals each) and lower percentage for borderline behavior (16.7%) (Figure 1).

As for skills of social competence, 83.3% of the individuals with del22q11.2 syndrome were classified as "clinical", 8.3% as "borderline" and 8.3% as "normal" (Figure 2).



Legend: Clinical> 70, and Borderline ≥67≤70 N ormal <67 0-9: number of individuals with the syndrome

Figure 1 – internalizing and externalizing behaviours in group sample



Legend: Clinical> 70, and Borderline ≥67≤70 Normal <67 0-12: number of individuals with the syndrome

Figure 2 – Social competence in group sample

By comparing the sample groups (SG) and control (CG) using the Mann-Whitney test, there were significant differences among them (Table 1, Figure 3). It was registered a statistically significant difference between CG and SG for the variables "internalizing behaviors", "withdrawn", problems", "attention problems" and "aggressive behavior."

In order to better characterize the behavior problems presented by the SG and CG and check if they established a relation with each other, the Spearman correlation analysis (Table 2) was performed.

According to the data obtained for behavioral problems, it was noted that complaints of "anxious and depressed" presented positive relation to complaints of "aggressive behavior." The same happened in the relation "anxious and depressed" with "the rule-breaking behavior".

The variable "somatic complaints" had a positive relationship with "social problems" and "thought problems" where the variables "social problems" and "thought problems" also had a positive relation with each other.

The variable "anxious and depressed" had a positive relation to "activities", related to social competence, and also with Internalizing Behaviors ("thought problems", "withdrawn" and "somatic complaints").

In order to verify possible differences between genders, a statistical correlation analysis was performed, in which it was not found a statistically significant difference for behavior problems and social competence in terms of male and female individuals in the SG (Figures 3 and 4).

Table 1 – Behavioral problems of sample group and control group

		М	SD	Min	Max	Med.	р	
Anxiety / Depressed	SG	62,17	7,68	50	76	65	0,664	
	CG	59,67	7,98	50	72	58		
Withdrawn	SG	67,42	8,85	54	85	68	0,013*	
	CG	57,67	7,34	50	77	57		
Somatic Complaints	SG	61,58	8,82	50	78	63	0,116	
	CG	56,08	4,66	50	64	55,5		
Social Problems	SG	68,83	7,69	61	83	67	<0,001**	
	CG	56,17	5,09	50	66	56,5		
Thought Problems	SG	60,58	7,56	50	71	60	0,021	
	CG	54,00	5,72	50	69	51		
Attention Problems	SG	65,83	6,9	52	77	65	0,001**	
	CG	54,08	5,38	50	66	52		
Hule-Breaking Behavior	SG	55,75	5,61	50	66	55,5	0,276	
	CG	53,00	3,41	50	60	51		
Agressive Behavior	SG	61,42	6,40	50	69	64	0,020*	
	CG	54,67	5,31	50	67	52		
Internalizing	SG	65,92	6,81	48	72	68,5	0,019*	
	CG	56,67	8,42	48	72	53		
Externalizing	SG	59,08	7,41	44	67	61,5	0,052*	
	CG	53,00	5,89	46	67	51		

Mann-Whitney Test p<.05*, p<.005**

Legend: M= Mean, SD= standard deviation, Min= Mínimum, Max= Maximum, Med= Median, (p)= significance, SG = sample group, CG = control group

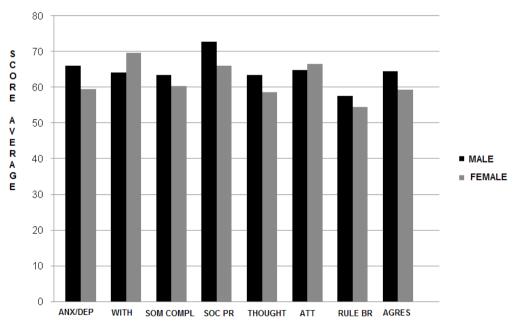
Table 2 – Degree of relationship between the variables (sample group)

		Anx / Depr	Withdrawn	Somatic Complains	Social Problems	Thought	Attention	Rule- Breaking Behavior	Agressive Behavior	Internaliz	Externaliz
Anx / Depr	Coef. Correl. (r)						,	,			
	Sig. (p)										
Withdrawn	Coef. Correl. (r)	-0,339									
	Sig. (p)	0,282									
Somatic Complains	Coef. Correl. (r)	+0,529	-0,112								
	Sig. (p)	0,077	0,728								
Social Problems	Coef. Correl. (r)	+0,327	-0,353	+0,549							
	Sig. (p)	0,300	0,261	0,065							
Thought	Coef. Correl. (r)	+0,247	-0,069	+0,454	+0,384						
	Sig. (p)	0,439	0,831	0,138	0,217						
Attention	Coef. Correl. (r)	+0,137	-0,110	+0,332	+0,637	+0,450					
	Sig. (p)	0,672	0,733	0,292	0,026*	0,143					
Rule- Breaking Behavior	Coef. Correl. (r)	+0,153	-0,250	+0,565	+0,576	+0,481	+0,263				
	Sig. (p)	0,634	0,433	0,056	0,050	0,113	0,409				
Agressive Behavior	Coef. Correl. (r)	+0,695	-0,419	+0,550	+0,556	+0,456	+0,372	+0,643			
	Sig. (p)	0,012*	0,175	0,064	0,061	0,136	0,234	0,024*			
Internaliz	Coef. Correl. (r)	+0,629	+0,093	+0,895	+0,480	+0,377	+0,107	+0,462	+0,474		
	Sig. (p)	0,028*	0,775	< 0,001**	0,114	0,227	0,741	0,130	0,119		
Externaliz	Coef. Correl. (r)	+0,420	-0,340	+0,670	+0,618	+0,543	+0,360	+0,933	+0,855	+0,551	
	Sig. (p)	0,173	0,279	0,017*	0,032*	0,068	0,250	< 0,001**	< 0,001**	0,063	
Total	Coef. Correl. (r)	+0,467	-0,340	+0,723	+0,866	+0,665	+0,577	+0,759	+0,671	+0,655	+0,813
	Sig. (p)	0,126	0,280	0,008*	< 0,001**	0,018*	0,050	0,004**	0,017*	0,021*	0,001**

Spearman Correlation p <.05, * P <.005 **

Legend: Cor. Coef. (r)=Correlation coefficient, (p)= significance

Anx/depr: anxious and depressed



Legend: Anx/Dep = Anxious and depressed, With = Withdrawn, Som Compl = Somatic Complains, Soc Pr = Social Problems, Thought = Thought Problems, Att = Attention Problems, Rule Br = Rule-Breaking Behavior, Agres = Aggressive Behavior, Mann-Whitney Test, p<.05*, p<.005**.

Figure 3 - Behavior problems as a function of gender for group sample

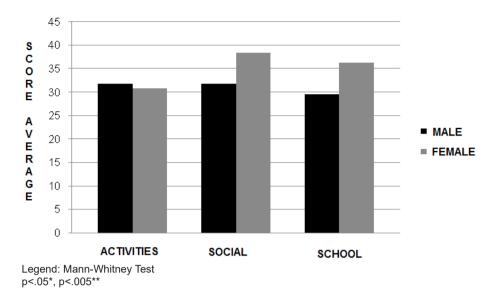


Figure 4 – Social competence as a function of gender for sample group

DISCUSSION

This was the first Brazilian paper that characterized the behavioral profile and social competence of individuals to del22q11.2 syndrome compared to typically developing individuals. These findings confirm international studies on behavioral problems using the CBCL in this population¹⁸⁻²⁰.

Through the analyzes performed, it can be highlighted the majority of complaints made by parents or quardians from the group with the 22q11.2 syndrome referred to deficits in social skills (83.3%), this fact confirms the findings of studies with the same population^{24,25}.

On literature, social skill impairments are associated with changes in school performance of children and adolescents in addition to the ability to relate with family and other individuals of the same age²⁵. It was observed the sample group had higher rates of attention problems than the control group, confirming reports of studies in which the prevalence of attention deficit disorders and attention deficit hyperactivity disorder is approximately ten times more in individuals with del22q11.2 than in the general population 18,26,27. There was a statistically significant difference between the sample group (SG) and the control group (CG) with respect to school performance. Assuming that attention is a basic skill for success in school, school failure may be explained by attentional deficits of this population, and other factors, such as intellectual disabilities28.

Anxiety problems were identified in this population as part of internalizing behavior problems, showing that 8 of the 12 individuals who participated in this study del22q11.2 classified as "clinical", i.e. out of the expected range for normality. One of the main of schizophrenia and most common found in del22g11.2 syndrome is anxiety9. The high incidence of schizoid symptoms in this particular genetic condition has been described as a neurobiological model of the investigation of genetic basis of schizophrenia that it was reported a subtype of del22q11.2 - Schizophrenia14.

All the individuals were categorized with behaviors beyond expectations in at least one of the scales of the CBCL. The data obtained allowed to observe statistically significant differences between CG and SG for the variables "withdrawn", "social problems", "attention problems", "aggressive behavior" and "internalizing behaviors" (e.g. anxious and depressed), thus confirming the hypothesis by previous studies that the presence of anxiety and depression would be frequent in children and adolescents with a diagnosis of del22q11.2 syndrome, which would cause impairments in adaptive functioning, e.g., a lack of ability of social nature in different areas 17,29.

The sample group also had higher rates of aggressiveness when compared to the control group, and the aggressiveness a feature commonly described in del22q11.2 syndrome14. It was also observed the reported problems with anxiety and depression, thought problems and rule-breaking behavior presented positive relation to complaints of aggressiveness for the subjects of this study¹⁹.

It was also proposed in this study to verify the possible distinctions based on gender, which it was observed there is no difference between the behavior problems and social competence between the male and female genders in the sample group, reinforcing the findings of Klaassen et al ²⁰. Paradoxically, there are other researchers, who found higher proportions of behavior problems for boys, expressed by clinically significant scores in "internalizing behaviors" (withdrawn, somatic complaints, anxious and depressed) and about thought problems regarding the girls^{30,31}.

CONCLUSION

Parents or guardians of individuals with diagnosis of del22q11.2 syndrome when responding to

behavioral inventory CBCL, reported the presence of behavioral and social competence in different degrees of commitment.

When performed to compare the groups we observe statistically significant differences in variables of externalizing behaviors (i.e. aggressive behavior, social and attention problems) and internalizing behaviors (i.e. withdrawn, anxious and depressed). Thus, we conclude that the sample group presents more abnormal behavior compared to the control group.

RESUMO

Objetivo: caracterizar os problemas comportamentais e de competência social de indivíduos com a síndrome del22q11.2 e compará-los com indivíduos com desenvolvimento típico, segundo informação dos pais. Métodos: participaram desta pesquisa 24 pais de indivíduos de ambos os gêneros, entre seis e 18 anos, sendo 12 indivíduos com a síndrome del22q11.2 (grupo amostral) e 12 indivíduos com desenvolvimento típico (grupo controle). Foi aplicado o inventário comportamental "Child Behavior Checklist (CBCL)". Resultados: oito dos 12 indivíduos com a síndrome foram classificados como "clínico" nas escalas de comportamento e Problemas Internalizantes; cinco dos 12 indivíduos do grupo amostral foram classificados como "clínico" quanto às escalas de comportamento e Problemas Externalizantes. Nas habilidades de competência social, dez dos 12 indivíduos do grupo amostral foram classificados como "clínico". Conclusão: indivíduos com diagnóstico da síndrome del22q11.2 apresentaram, segundo opinião dos pais, problemas comportamentais e de competência social, em diferentes graus de comprometimento. Quando realizada a comparação entre os grupos pode-se observar diferenças estatisticamente significantes em variáveis dos comportamentos externalizantes e dos comportamentos internalizantes. Desta forma, concluí-se que o grupo amostral apresenta comportamentos mais alterados quando comparados ao grupo controle.

DESCRITORES: Fonoaudiologia; Comportamento; Relações Interpessoais; Comunicação; Transtornos do Comportamento Social; Síndrome de DiGeorge

REFERENCES

- 1. Online Mendelian Inheritance in Man, OMIM (TM). McKusick-Nathans Institute for Genetic Medicine, Johns Hopkins University (Baltimore, MD) and National Center for Biotechnology Information, National Library of Medicine (Bethesda, MD). Disponível em: http://www.ncbi.nlm.nih.gov/omim/. Acesso em: fevereiro/2014.
- 2. Shprintzen RJ. Velo-cardio-facial syndrome: 30 years of study. Dev Disabil Res Rev. 2008;14:3-10.
- 3. Shprintzen RJ, Goldberg RB, Lewin ML, Sidoti EJ, Berkman MD, Argamaso RV et al. A new syndrome involving cleft palate, cardiac anomalies,

typical facies, and learning disabilities: velo-cardio-facial syndrome. Cleft Palate J. 1978;15(1):56-62.

- 4. Sandrin-Garcia P, Abramides DVM, Martelli LR, Ramos ES, Richieri-Costa A, Passos GAS. Fluorescence in situ hybridization (FISH) screening for the 22q11.2 deletion in patients with clinical features of velocardiofacial syndrome but without cardiac anomalies. Genet Mol Biol. 2007;30(1):21-4.
- 5. Ryan AK, Goodship JA, Wilson DI, Philip N, Levy A, Seidel H et al. Spectrum of clinical features associated with interstitial chromosome 22ql1 deletions: a European collaborative study. J Med Genet.1997;34(10):798-804.

- 6. Solot CB, Knightly C, Handler SD, Gerdes M, McDonald-McGinn DM, Moss E et al. Communication Disorders in the 22q11.2 Microdeletion Syndrome J. Commun Disord. 2000;33:187-204.
- 7. De Smedt B. Mathematical learning disabilities in children with 22q11.2 deletion syndrome: a review. Dev Disabil Res Rev. 2009;15(1):4-10.
- 8. Bassett AS, Marshall CR, Lionel AC, Chow EW, Scherer SW. Copy number variations and risk for schizophrenia in 22q11.2 deletion syndrome. Hum Mol Gen. 2008;17(24):445-53.
- 9. Philip N, Bassett AS. Cognitive, behavioural and psychiatric phenotype in 22q11.2 deletion syndrome. Behav Genet. 2011:41(3):403-12.
- 10. Fine SE, Weissman A, Gerdes M, Pinto-Martin J, Zackai EH, McDonald-McGinn DM et al. Autism Spectrum Disorders and Symptoms in Children with Molecularly Confirmed 22q11.2 Deletion Syndrome. J Autism Dev Disord. 2005;35(4):461.
- 11. Antshel KM. Autistic Spectrum Disorders in Velo-cardio Facial Syndrome (22g11.2 Deletion). J Autism Dev Disord. 2007;37:1776-86.
- 12. Antshel KM, Shprintzen R, Fremont W, Higgins AM, Faraone SV, Kates WR. Cognitive and psychiatric predictors to psychosis in velocardiofacial syndrome: a 3-year follow-up study. J Am Acad Child Adolesc Psychiatry. 2010;49(4):333-44.
- 13. Kates WR. Neuroanatomic predictors to prodromal psychosis in velocardiofacial syndrome (22q11.2 deletion syndrome): a longitudinal study. Biol Psychiatry. 2011;69:945-52.
- 14. Sinderberry B, Brown S, Hammond P, Stevens AF, Schall U, Murphy DGM. et al. Subtypes in 22q11.2 deletion syndrome associated with behaviour and neurofacial morphology. Res Dev Disabil. 2013; 34:116-25.
- 15. Aneja A, Fremont WP, Antshel KM, Faraone SV, AbdulSabur N, Higgins AM et al. Manic symptoms and behavioral dysregulationin youth with velocardiofacial syndrome (22q11.2 deletion syndrome). J Child Adolescent Psychopharmacol. 2007;17(1):105-14.
- 16. WoodinM, Wang PP, Aleman D, McDonald-McGinn D, Zackai E, Moss E. Neuropsychological profile of children and adolescents with the 22q11.2 microdeletion.Genet Med. 2001;3(1):34-9.
- 17. Briegel W, Schneider M, Schwab KO. 22q11.2 deletion syndrome: behaviour problems of children and adolescents and parental stress. J Compilation. 2008;34(6)795-800.

- 18. Lajiness-O'Neill R, Beaulieu I, Asamoah A, Titus JB, Bawle E, Ahmad S et al. The neuropsychological phenotype of velocardiofacial syndrome (VCFS): Relationship to psychopathology. Arch Clin Neuropsychol. 2006;21:175-84.
- 19. Niklasson L, Rasmussen P, Óskarsdóttir S, Gillberg C. Autism, ADHD, mental retardation and behavior problems in 100 individuals with 22q11 deletion syndrome. Res Devel Desabil. 2009;30:763-73.
- 20. Klaassen P, Duijff S, Veye S, Vorstman J, Beemer F, Sinnema G. Behavior in Preschool Children with the 22q11.2 deletion syndrome. AM J Med Gen. 2012;161(1):94-101.
- 21. Jansen PW. Duiiff SN. Beemer FA. Vorstman JA, Klaassen PW, Morcus ME et al. Behavioral problems in relation to intelligence in children with 22q11.2 deletion syndrome: a matched control study. Am J Med Genet A. 2007;143:574-80.
- 22. Achenbach TM. Manual for the child behavior checklist/ 4-18 and profile. Burlington VT. University of Vermont, Department of Psychiatry, 1991.
- 23. Bordin IA, Mari JJ, Caeiro MF. Versão brasileira do "Child Behavior Checklist for ages 6-18". ASEBA: University of Vermont, 2010.
- 24. Ferro MR, Abramides DVM, Veronez FS, Tavano LD. Souza SRB. De-Vitto LPM etal. Habilidades sociais em pacientes com síndrome velocardiofacial. Arg Ciênc Saúde. 2008;15(4):157-62.
- 25. Looman WS, Thurmes AK, O'conner-Von SK. Quality of Life among Children with Velocardio Syndrome.Cleft Palate Craniofac 2010;47(3):273-83.
- 26.Zagursky K, Weller RA, Jessani N, Abbas J, Weller EB. Prevalence of ADHD in Children with Velocardiofacial Syndrome: A Preliminary Report. Curr Psychiatry Rep. 2006;8(2):102-7.
- 27. Shashi V, Veerapandiyan A, Kwapil T, Keshavan E, Hooper S. Social skills and associated psychopathology in children with chromosome 22q11.2 deletion syndrome: implications for interventions. J Intellect Disabil Res. 2012;56(9):865-78.
- 28. Hooper SR, Curtiss K, Schoch K, Keshavan MS, Allen A, Shashi V. A longitudinal examination of the psychoeducational, neurocognitive, and psychiatric functioning in children with 22q11.2 deletion syndrome.Res Dev Disabil. 2013;34(5):1758-69.
- 29.Fabbro A, Rizzi E, Schneider M, Debbane M, Eliez S. Depression and anxiety disorders in children and adolescents with velo-cardio-facial syndrome (VCFS).Eur Child Adolesc Psychiatry. 2012;21(7):379-85.

- 30. Young AS, Shashi V, Schoch K, Kwapil T, Hooper SR. Discordance in Diagnoses and Treatment of Psychiatric Disorders in Children and Adolescents with 22q11.2 deletion syndrome. Asian J Psychiatr. 2011;1-4(2):119-24.
- 31. Sobin C, Kiley-Brabeck K, Monk SH, Khuri J, Karayiorgou M. Sex differences in the behavior of children with the 22q11 deletion syndrome. Psychiatry Res. 2009;166(1):24-34.

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