

DOI: 10.5455/msm.2020.32.93-98

Received: MAR 19 2020; Accepted: MAY 26, 2020

© 2020 Aikaterini Katsiana, Nikolaos Strimpakos, Ventoulis Ioannis, Eleni Kapreli Maria Sofologi, Eleni Bonti, Kotrotsiou Stilian, Anastasios Stalikas

This is an Open Access article distributed under the terms of the Creative Commons Attribution Non-Commercial License (<http://creativecommons.org/licenses/by-nc/4.0/>) which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

ORIGINAL PAPER

Mater Sociomed. 2020 Jun; 32(2): 93-98

# Health-related Quality of Life in Children with Autism Spectrum Disorder and Children with Down Syndrome

Aikaterini Katsiana<sup>1,2</sup>, Nikolaos Strimpakos<sup>3</sup>, Ventoulis Ioannis<sup>2</sup>, Eleni Kapreli<sup>3</sup>  
Maria Sofologi<sup>4</sup>, Eleni Bonti<sup>5</sup>, Kotrotsiou Stilian<sup>6</sup>, Anastasios Stalikas<sup>1</sup>

<sup>1</sup>Department of Psychology, Panteion University of Social and Political Sciences, Athens, Greece

<sup>2</sup>Department of Occupational Therapy, University of Western Macedonia, KEPTSE, Ptolemaida, Greece

<sup>3</sup>Department of Physiotherapy, University of Thessaly, Lamia, Greece

<sup>4</sup>Department of Psychology, Aristotle University of Thessaloniki, Greece

<sup>5</sup>1st Psychiatric Clinic, Medical School, Aristotle University of Thessaloniki, "Papageorgiou" General Hospital of Thessaloniki

<sup>6</sup>General Department, University of Thessaly, Larisa, Greece

## Corresponding authors:

Aikaterini Katsiana.  
Department of Psychology, Panteion University of Social and Political Sciences, Athens, Greece 5 Lioli street, 41222 Larissa, Greece. e-mail: [katerinakatsiana@gmail.com](mailto:katerinakatsiana@gmail.com). ORCID ID: <http://www.orcid.org/0000-0002-7826-1353>.

## ABSTRACT

**Introduction:** Measuring health-related quality of life (HRQoL) is very important for children with developmental disorders such as autism spectrum disorder (ASD) and Down syndrome (DS). However, no HRQoL studies found in the literature for the differences between children with ASD and children with DS. **Aim:** The aim of this study was to examine HRQoL in children with ASD and children with DS. **Methods:** The participants consisted of 206 children with ASD (61), DS (55) and typical development (TD) (90), aged 5-10 years old, after administering anonymous questionnaires to their parents-caregivers. The Pediatric Quality of Life Inventory™ 4.0- Parent Report (PedsQL) was used to measure HRQoL. One-way analysis of variance and  $\chi^2$  were applied for comparisons among groups. **Results:** TD group scored higher than ASD and DS in all comparisons. Post-hoc (Tukey) comparisons revealed that the statistically univariate effect was due to differences between the TD group and the other two groups, ASD and DS ( $p < 0.01$ ). The ASD group achieved significantly lower scores than DS in the emotional functioning scale. Post-hoc analysis did not reveal any significant differences between the DS and the ASD group in the physical health, psychosocial health and the total PedsQL summary scores. **Conclusions:** Children with ASD and DS had significantly lower HRQoL compared to a TD population, and this finding was not affected by age. Children with ASD demonstrated a significantly lower score in the emotional functioning scale than children with DS but are similar in the physical health scale. It is thus considered necessary to take the physical health scale into ac-

count when assessing and designing treatment for children with ASD. Future research studies should focus on HRQoL indicators that could serve as a standard diagnostic tool for the development of therapies and outcomes of assessment findings in ASD and DS.

**Keywords:** Health Related Quality of Life, Autism Spectrum Disorder, Down syndrome, children.

## 1. INTRODUCTION

Health-related quality of life (HRQoL) is an important measure not only of the physical health of the individuals but also of their mental health, as well as of the outcome of their treatment (1, 2). HRQoL is an approach to measuring health outcomes by evaluating an individual's psychosocial, emotional and physical well-being (3). In the pediatric Quality of life (QoL) literature, domains that have been systematically addressed include: physical functioning, emotional functioning, social functioning and school functioning (4.). Each domain can be broken down into indicators that are the actual factors measured. Utilizing QoL indicators as the standard for developing treatments and evaluating outcomes in autism is advantageous (e.g. social functioning can include indicators of the availability of social support, while emotional functioning may include happiness and mental health). Measuring health-related quality of life is very important for children with developmental disorders such as autism spectrum disorder (ASD) and Down syndrome (DS). Most quality of life studies focus on the parents of children with

ASD and other disorders or disabilities, whereas only few examine the quality of life of the children themselves (5). ASD is a neurodevelopmental disorder, a chronic lifelong condition that starts from childhood and affects the outcomes of adulthood, having an impact on the QoL of individuals with ASD, which in turn is not influenced by age and is lower compared to individuals without autism (5). ASD is characterized by difficulties in social and communication development in addition to repetitive behaviors and limited interests and thus creates difficulties in employment and socialization of these individuals. Children with ASD and their parents report significantly worse HRQoL than their peers with typical development in terms of both psychosocial and emotional health (7-9).

Children with ASD also exhibit greater rates of depression, stress and anxiety than typically developing (TD) children (10), (as well as poorer physical health (more sleep issues, gastrointestinal problems and allergies). They also have a worse QoL of life than peers with chronic conditions (5) and demonstrate higher rates of attention deficit-hyperactivity disorder (11) and psychiatric disorders such as disorganized behavioral disorders, anxiety and emotional disorders (4). To better understand the HRQoL of people with ASD, further studies on the difference between individuals with ASD and individuals with other chronic physical and psychiatric disorders and / or mental disability are needed (12). To our knowledge, no HRQoL studies have been performed on the differences between children with ASD and children with DS and very few studies have addressed the QoL of children with DS (13, 14). Children with DS have more emotional and behavioral problems but less anxiety and depression than TD children and therefore worse HRQoL than their typical peers (13, 15).

## 2. AIM

The purpose of this study was to investigate HRQoL in children with ASD and in children with DS in order to improve the provided services and treatment. We therefore hypothesized that, by means of traditional measures of HRQoL, children with ASD and DS would have poorer overall HRQoL than typically developing children and furthermore that children with ASD would demonstrate worse emotional and social functioning HRQoL scores than children with DS.

## 3. METHODS

### Participants

A convenient sample of caregivers and their children (N=206), consisted of three separate groups, one with ASD (N=61), one with DS (N=55), along with TD children serving as a control group (N=90), were recruited from kindergartens, primary schools, DS parents' associations, ASD parents' associations as well as from various special treatment and rehabilitation centers for children in Greece. The inclusion criteria were children aged between 5 and 10 years old with a diagnosis of ASD, according to the DSM-IV-TR (16) and DS. Children with ASD and DS should have no other developmental disorders such as physical disabilities, epilepsy, etc., while typically developing children shouldn't have been diagnosed with any learning

disabilities or chronic diseases. In order to participate in the study, parents needed to speak fluent Greek and give informed consent. Initially 280 questionnaires were handed out, of which 220 (78.6%) were returned. 14 of the answered questionnaires did not meet the inclusion criteria and were thus excluded. Eventually, the responses of 206 participants were taken into account. The study was approved by the National Institute for educational policy (PSE) and information leaflets, together with consent forms, were given to the participants recruited in the study.

### Procedures

All study participants were recruited within a year. The Pediatric Quality of Life Inventory™ 4.0 Generic Core Scales – Parent Report (PedsQL) questionnaire was used for the purpose of the study. Parents who met the inclusion criteria received a package containing an informative letter about the study and its purpose, a demographic questionnaire, the age-appropriate PedsQL questionnaire, the age-appropriate Vineland II- (domain “Daily Living Skills”) questionnaire and a consent form. The questionnaires were filled in anonymously with codes and were returned within one month.

### Measures

Demographic variables were collected from a parent-completed questionnaire and included children's age, gender and chronic diseases, along with information regarding school type and parents' educational level. Measurement of HRQoL was performed by utilization of the Pediatric Quality of Life Inventory™ 4.0 Generic Core Scales – Parent Report (PedsQL) for ages 5-7 and 8-12. The Daily Living Skills Index of Vineland II (Parent/ Caregiver Rating Form) was used, as recommended in the literature (17), as an indication of the participants' developmental level (sample homogeneity). Due to troublesome communication and other related difficulties that are common among children with ASD and DS, the children's HRQoL was assessed using the parent-report version of the PedsQL.

### PedsQL

The PedsQL questionnaire is a tool for assessing the health-related quality of life of children aged 2-18 years old (18). It is a 23-item questionnaire which includes four age-appropriate versions and takes approximately 7-10 minutes to complete. The parent proxy-report format assesses the parent's perceptions of the child's QoL. The instructions in each question ask how much of a problem an item has been for the child during the past month. PedsQL has a five-point rating scale and it evaluates four distinct areas of health-related functioning: physical functioning, emotional functioning, social functioning and school functioning. Consequently, the last three scales are merged into a summary score for psychosocial health, while the physical functioning scale gives rise to the physical health summary score. Eventually the total PedsQL score consists of the sum of all the items over the number of items answered on all the scales. The items on the PedsQL questionnaire are converted into a 0-100 scale with higher scores indicating better HRQoL. The total PedsQL score, as well as the summary and scale scores, are computed when at least 50% of the items are completed. The parent-report version of the PedsQL exhibits good psychometric properties for

measuring HRQoL in healthy populations and in children with chronic diseases and acute health conditions (alpha 0.90- 0.92) (19) as well as in pediatric populations with psychiatric disorders (4)

The current study used the Greek version of PedsQL, which accordingly shows good psychometric properties (Cronbach alpha > 0.70). Factor analysis also manifested comparable results to the original version (20).

**Vineland II (VABS II)-Domain Daily Living Skills (DLS)**

The VABS II -Parent/ Caregiver Rating Form (21) assess the adaptive behavior and functional skills in people aged 0-90 years old. It has been also suggested for assessing adaptive behavior for people with ASD as well as with mental and developmental disorders (22). The subscale of DLS includes three sectors assessing skills for self-care, home care and living in the community. Its psychometric properties are very good and suitable for research purposes (23). The internal reliability (Spearman- Brown) for DLS is ranged between 0.88 and 0.94 (split-half method for different ages) (21) The Greek version of the questionnaire yielded almost identical reliability values.

**Statistical analysis**

Descriptive statistics of baseline characteristics were obtained using Statistical Package for Social Sciences, software (SPSS, version 20.0). Sample characteristics and presented means and standard errors were described for PedsQL scales. Among group (ASD, DS, TD) differences were compared using the ANOVA and chi-squared (x<sup>2</sup>) tests. Correlation between HRQoL scores and age was performed with Pearson correlation coefficient.

**4. RESULTS**

Table 1 summarizes the baseline demographic characteristics of the 206 children. Out of the 61 children with ASD, 25 (41%) were diagnosed with Autistic Disorder, 8 (13.1%) with Asperger’s Disorder and 28 (45.9%) with PDD-NOS. The age of the children ranged from 5 to 10 years old and there was a significant difference between all of them (F=5.65, p<0.01) mainly because the ASD group was slightly younger (differences between DS and TD were non-significant, p<0.05). Most children with ASD were boys (85.2%) as was expected (24). Among children with DS 60% were boys and 40% were girls while TD participants were

	Autism Spectrum Disorder (N=61)	Down Syndrome (N=55)	Typical Development (N=90)	
Mean age (SD)	6.5 (1.30)	7.3 (1.51)	7.2 (1.36)	F=5,65**
Gender				
Male %	52 (85)	33 (60)	40 (44.4)	χ <sup>2</sup> =25,38***
Female %	9 (14)	22 (40)	50 (55.6)	
Mother’s education level				
Primary	3(4.9%)	2(3.6%)	1(1.1%)	χ <sup>2</sup> =16.37**
Secondary	18(27.3%)	15(27.3%)	13(14.6%)	
Post-secondary	14(22.9%)	17(30.9%)	15(16.9%)	
Higher	26(42.7%)	21(38.2%)	60(67.4%)	
Father’s education level				
Primary	4(6.6%)	4(7.4%)	2(2.3%)	χ <sup>2</sup> =19.84**
Secondary	27(44.3%)	11(20.4%)	19(35.2%)	
Post-secondary	8(13.1%)	19(35.2%)	19(21.1%)	
Higher	22(36%)	20(37%)	47(54.1%)	
School type				
Public	52 (85.2%)	45 (83.3%)	89 (98.9%)	
Private	9 (14.8%)	9 (16.7%)	1 (1.1%)	

Table 1. Demographic characteristics of the children with Autism Spectrum Disorder (ASD), Down syndrome (DS) and Typical Development (TD) and their parents. Note: \*p < .05, \*\* p < .01, \*\*\* p < .001

	N	Mean	SD	Range	p
Autism Spectrum Disorder	61	71,31	15,64	38 – 134	NS
Down Syndrome	55	72	11,31	46 – 67	
Typical Development	90	115,82	15,9	85 – 158	

Table 2. Participants’ developmental level (sample homogeneity): Daily living skills Index (VABS II). NS=non-significant

44.4% and 55.6% respectively. The gender differences between the groups were statistically significant (x<sup>2</sup> = 25.38, p<0.001) (Table 1). Table 2 is presenting participants’ developmental level according to children’ Daily Living skills with non-significant differences between ASD and DS children (p>0.05) showing sample homogeneity. The majority of the children were attending a public school with about 15% of children with ASD and DS were going to a private one. Parents’ education level varied among groups for both mothers and fathers (p<0.01) with higher education yielding

	Autism Spectrum Disorder (N=61)		Down Syndrome (N=55)		Typical Development (N=90)		F	η <sup>2</sup>	P
	M.O	SD	M.O	SD	M.O	SD			
Parent- Proxy Report (PedsQL)									
Physical Functioning	74,29	17,61	73,98	17,5	90,90	14,05	27,39	.21	.001
Emotional Functioning	69,02	22,23	77,11	17,86	85,33	14,74	15,01	.13	.001
Social Functioning	54,40	24,83	61,84	18,03	90,28	14,25	76,37	.43	.001
School Functioning	65	20,35	67	15,71	88,22	13,5	47,13	.32	.001

Table 3. Mean PedsQL scale scores for children with Autism Spectrum Disorder (ASD), Down syndrome (DS) and Typical Development (TD).

Parent-Proxy Report (PedsQL)	Autism Spectrum Disorder (N=61)		Down Syndrome (N=55)		Typical Development (N=90)		F	$\eta^2$	p
	M.O	SD	M.O	SD	M.O	SD			
Physical health	74.29	(17.61)	73.98	(17.5)	90.90	(14.05)	27.39	.21	.001
Psychosocial health	62.81	(18.95)	68.65	(14.18)	87.94	(11.97)	59.22	.37	.001
Total score	65.68	(17.57)	69.98	(14.03)	88.68	(11.49)	56.59	.36	.001

Table 4. Mean PedsQL summary and total scores for children with Autism Spectrum Disorder (ASD), Down Syndrome (DS) and Typical Development (TD)

	Physical health	Psychosocial health	PedsQL Total score
Autism Spectrum Disorder	.08	.07	.08
Down Syndrome	.20	.06	.11
Typical Development	.17	.11	.14

Table 5. Correlation analysis (Pearson r) of HRQoL with age for children with Autism Spectrum Disorder (ASD), Down Syndrome (DS) and Typical Development (TD)

the highest percentage (Table 1).

Regarding the presence of chronic diseases, almost half of DS children (44.5%) had a chronic problem such as heart disease (16.4%), respiratory disease (9.1%) or other health problems (20%) while only 19.7% of children with ASD presented with chronic health problems mainly concerning the respiratory system (9.8%)

Table 3 shows possible differences among the three groups across the various PedsQL scales. The one-way analysis revealed statistically significant differences among the three groups in all four domains. The post-hoc comparisons (Turkey) indicated that the statistically univariate effect was due to differences between the typically developing group and the other two groups, with the TD group consistently scoring higher in all four domains. Interestingly, the DS and the ASD group did not differ from each other with regard to the physical functioning, social functioning and school functioning scale. The only comparison that detected a statistically significant difference was the one concerning the emotional functioning scale; as anticipated, the ASD group had significantly lower scores than their peers with DS.

Two composite scores were calculated, namely the psychosocial health summary score (representing the emotional, social and school functioning scales) and the total PedsQL score (the sum of all items over the number of items answered on all scales). Also included in Table 4 is the physical health summary score which corresponds to the physical functioning score presented in Table 3.

One-way analyses of variance were conducted to examine possible differences among the three groups of children on the two summary scores of the PedsQL, as well as on the total composite score of the scale (total PedsQL score). In all comparisons, statistically significant differences were found. As expected, the TD group scored higher in all 3 comparisons. The significant differences between the TD group and the other two groups (DS and ASD) were subsequently confirmed in the post-hoc (Turkey) analyses

performed. Interestingly, these post-hoc analyses did not reveal any significant differences between the DS and the ASD group. Nevertheless, as shown in Table 4, the ASD group scored lower than the DS group in the psychosocial health summary score and, by extension, in the total PedsQL score, reflecting the often cited lower quality of life that ASD children have in these domains.

The correlation analysis of HRQoL with age showed no statistically significant relationship both in each separate HRQoL category (physical health and psychosocial health) and in the total PedsQL score for all three groups of children (Table 5).

### 5. DISCUSSION

The results of the present study support our initial hypothesis that children with ASD and DS would have poorer HRQoL than TD children. The parents of children with ASD and DS reported poorer HRQoL scores for their children than scores reported by parents of TD children. In fact, significant differences were observed between these two groups in all HRQoL scales (physical, emotional, social and school functioning scales, as well as total HRQoL score).

The findings of this study are similar to those of a previous meta-analysis of ten studies (from 2004 to 2012) for children with ASD, which concluded that the observed poorer QoL was not affected by age (6) and are also in accordance with studies showing poorer physical health, mental health and overall QoL of children with ASD (4, 5, 11). ASD is a neurological disorder characterized by irregular social interaction and communication, repetitive behaviors and limited interests, impinging on many areas of the children's development and adversely affecting their functioning. Therefore, compared to TD children, children with ASD are expected to experience significant problems related to psychological, social and emotional health. Deficits in social skills and communication are evident (25).

Children with ASD may also have higher rates of depression, stress, and anxiety (10). In this study, parents reported that the main problems of their children with ASD involved aspects of the social functioning. Similar results were reported in the study by Kuhlthau et al., (2010). In addition to the social and emotional difficulties, ASD children often face difficulties in school life, learning and academic functioning. Moreover, according to the literature review, there are reports of some mild neurological findings related to ASD, such as motor impairment and motor developmental delay (26). Frequently, sleep disorders, gastrointestinal problems and various allergies coexist in ASD children (27) therefore affecting their physical health. As a result, developmental delays and difficulties in the above mentioned areas can cause poorer HRQoL in children with ASD.

Regarding children with DS, not enough studies have examined their QoL. In the current study, children with DS also showed a lower HRQoL than their TD peers, in all domains and regardless of age. These results are consistent with those of two previous studies (14, 15). Presumably, this is an expected finding since these children have

developmental deficits (average developmental age lower than their TD peers), as well as more behavioral and emotional problems.

The results of the current study indicate that DS has a stronger impact on all aspects of emotional functioning and also that DS children experience greater difficulties in social functioning than in other domains. Their social functioning is lower than that of TD children and this may be due to their behavioral problems and intellectual disability. The QoL associated with school functioning also seems to be low and this could be attributed to their intellectual disability and dysfunction. Indeed, the latter is supported by a previous study (28). The QoL associated with physical functioning is also shown to be adversely affected and this may be related to the chronic diseases that coexist in DS children (mainly respiratory diseases, congenital heart disorders, thyroid dysfunctions, etc.), as well as the delay in their motor development.

Our hypothesis in this study, that children with ASD would achieve poorer QoL scores than children with DS, was not confirmed for all scales of HRQoL except for emotional functioning. Given that emotional impairment is a defining characteristic of ASD but not of DS, it is not surprising that children with ASD were found to have lower emotional functioning scores than children with DS. Children with ASD have deficits in the cognitive processing of emotions, both with themselves and with others. They actually face difficulties in both identifying and describing emotions. They are also more frequently depressed than TD children (10), whereas, in a previous study, children with DS demonstrated better scores than TD children in the anxiety/depression scale (14). In addition, children with ASD have more difficulty in recognizing facial expressions compared to children with other developmental disorders, e.g. with schizophrenic disorder, and this poses an additional family burden (29). Similar to children with cognitive disabilities and children with speech disorders, children with ASD exhibit greater difficulty in recognizing expressions and emotions such as sadness, joy, anger, surprise, etc. (30).

Although the group of children with ASD scored lower in all HRQoL scales, there did not appear to be any statistically significant differences in the physical health, psychosocial health and the total PedsQL summary scores, when compared to their peers with DS. Given the hypotonia of the children with DS, as well as their physical condition and the various health problems they usually encounter (respiratory, heart diseases, etc.) as in present study, it was expected that children with ASD would have a better quality of life related to physical health. However, what may be surprising is the fact that children with ASD scored as poorly as children with DS in the physical health scale, especially considering the fact that ASD has not traditionally been thought to have a substantial impact on physical health, while many other chronic diseases of children with DS do. These results reinforce the view of a previous study (5) that the physical health of children with ASD should be taken into consideration based on the finding that their physical health was similar to that of children with chronic diseases. It is therefore necessary to take the physical health scale into account when assessing and

designing treatment for children with ASD. Empowering and maintaining the physical health of children with ASD may need to be addressed intensively, in conjunction with the psycho-emotional health (emotional and social functioning), which is an integral part of the ASD. Comparison of HRQoL across diagnostic groups might help clinicians and families understand their experience with ASD in the context of other developmental disorders. A better understanding of similarities across diagnostic groups may also result in a greater collaboration among advocacy groups for improving HRQoL in children with DS. Despite the advances in early detection, intensive intervention and therapeutic approaches, QoL in children with ASD and DS remains poor. Few studies have utilized specific QoL indicators in order to evaluate treatment outcomes in children with ASD and DS, leaving this area of study largely untapped (1).

The present study has several limitations. First, reports of HRQoL are based on parent proxy-report and not on child self-report due to the severe cognitive and communication issues faced by a majority of our study sample; most children would not have been able to reliably self-report. Although we used a validated methodology for children, this does not exclude the possibility of parenthood subjectivity through providing reports on behalf of their children. Second, despite the fact that the sample was selected at random and represented geographical areas across Greece, parental participation in the study was voluntary and the study participants constituted a convenience sample. Moreover, although in the present study the effect of socioeconomic status of parents on children HRQoL was not measured since it is not usual in children QoL studies and because of personal data protection limitations it would be interesting future studies to examine this factor also.

## 6. CONCLUSION

The present study explored the HRQoL in children with ASD and in children with DS. Children with ASD and DS had significantly lower HRQoL compared to a group of TD children, irrespective of age. When compared to data from children with DS, children with ASD demonstrated a worse HRQoL in the emotional functioning scale, but did not exhibit differing scores neither in the total HRQoL and the psychosocial health score nor in the social functioning, school functioning and physical functioning scales. These results indicate that the physical health of children with ASD should consistently be taken into consideration. Contrary to DS, ASD has not traditionally been thought to have a substantial impact on physical health. It is therefore essential to take physical health into account whenever assessing the QoL as well as when designing treatments for children with ASD. Knowing which domains of HRQoL are affected in children with ASD and DS can help clinicians focus on particular HRQoL domains during the diagnostic process and to define adequate treatment goals. A future research step must emphasize on HRQoL indicators in order to build up a future assessment and diagnostic tool in an attempt to develop a variety of therapies for ASD and DS children. A plethora of therapies that support HRQoL can contribute to greater happiness and overall well-being for children with ASD and DS in their environment.

- **Declaration of Patient Consent:** The authors certify that they obtained all appropriate patient consent forms.
- **Author's contribution:** All authors contributed equally in the conduction of the study and the preparation of the manuscript and they were overseeing these contents of the article, had full access and responsibility of the data. Manuscript preparation and final proof reading was made by the first and second author.
- **Conflict of interest:** None declared.
- **Financial support and sponsorship:** Nil"

## REFERENCES

- Burgess AF, Gutstein SE. Quality of life for people with autism: Raising the standard for evaluating successful outcomes. *Child and Adolescent Mental Health*. 2007; 12: 80-86.
- Kamp-Becker I, Schroder J, Remschmidt H, Bachmann CJ. Health-related quality of life in adolescents and young adults with high functioning autism-spectrum disorder. *Psychosoc Med*. 2010; 7: 1-10.
- Fayers P, Machin D. *Quality of Life: The assessment, analysis and interpretation of patient-reported outcomes*. 2nd ed. US:Wiley; 2007.
- Bastiaansen D, Koot HM, Bongers IL, Varni JW, Verhulst FC. Measuring quality of life in children referred for psychiatric problems: psychometric properties of the PedsQL 4.0 generic core scales. *Qual.Life Res*. 2004; 13: 489-495.
- Kuhlthau K, Orlich F, Hall TA, Sikora D, Kovacs EA, Delahaye J et al. Health-Related Quality of Life in children with autism spectrum disorders: results from the autism treatment network. *J Autism Dev Disord*. 2010; 40 (6): 721-729.
- Van heijst BF, Geurts HM. Quality of life in autism across the lifespan: a meta-analysis. *Autism*. 2015; 19: 158-167.
- de Vries M, Geurts H. Influence of autism traits and executive functioning on quality of life in children with an autism spectrum disorder. *J Autism Dev Disord* 2015; 45: 2734–2743.
- Ikeda E, Hinckson E, Krägeloh C. Assessment of quality of life in children and youth with autism spectrum disorder: a critical review. *Qual Life Res*. 2014; 23(4): 1069-1085.
- Ten Hoopen LW, de Nijs PF, Duvekot J, Greaves-Lord K, Hillegers MHJ, Brouwer WBF et al. Children with an autism spectrum disorder and their caregivers: Capturing health-related and care-related quality of life. *J Autism Dev Disord*. 2020; 50(1): 263-277.
- Gurney JG, McPheeters ML, Davis MM. Parental report of health conditions and health care use among children with and without autism: National survey of children's health. *Arch Pediatr Adolesc Med*. 2006; 160: 825-830.
- Lee LC, Harrington RA, Louie BB, Newschaffer CJ. Children with autism: quality of life and parental concerns. *J Autism Dev Disord*. 2008; 38: 147-1160.
- Kose S, Erermis S, Ozturk O, Ozbaran B, Demiral N, Bildik T. et al. Health- related quality of life in children with autism spectrum disorders: The clinical and demographic related factors in Turkey. *Res Autism Spectr Disord*. 2013; 7: 213-220.
- Bertoli M, Biasini G, Calignano MT, Celani G, De Grossi G, Digilio MC. et al. Needs and challenges of daily life for people with Down syndrome residing in the city of Rome, Italy. *J Intellect Disabil Res*. 2011; 55: 801-820.
- Van Gameren-Oosterom HB, Fekkes M, Buitendijk SE, Mo- hangoo AD, Bruil J, Van Wouwe JP. Development, problem behavior, and quality of life in a population based sample of eight-year-old children with Down syndrome. *PLoS One*. 2011; 6(7): e21879.
- Shields N, Leonard HM, Munteanu SA, Bourke J. Parent-reported health-related quality of life of children with Down syndrome: A descriptive study. *Dev Med Child Neurol*. 2018; 60(4).
- American Psychiatric Association. *Diagnostic and Statistical Manual of Mental Disorders*. 4th ed. Washington: DC: Author; 2000.
- Lane AE, Young RL, Baker AE, Angley MT. Sensory processing subtypes in autism: association with adaptive behavior. *J Autism Dev Disord*. 2010; 40: 112-122.
- Varni JW, Seid M, Rode CA. The PedsQLTM: Measurement model for the Pediatric Quality of Life Inventory TM. *Med Care*. 1999; 37: 126-139.
- Varni JW, Seid M, Kurtin PS. PedsQL 4.0: reliability and validity of the Pediatric Quality of Life Inventory version 4.0 generic core scales in healthy and patient populations. *Med Care*. 2001; 39: 800-812.
- Gkoltsiou, K, Dimitrakaki C, Tzavara C, Papaevangelou V, Varni JW, Tountas Y. Measuring health-related quality of life in Greek children: psychometric properties of the Greek version of the Pediatric Quality of Life Inventory(TM) 4.0 Generic Core Scales. *Qual Life Res*. 2008; 17: 299-305.
- Sparrow SS, Cicchetti D, Balla D. *Vineland Adaptive Behavior Scales*. 2nd ed. Minneapolis: Pearson Assessment; 2005.
- Mazefsky CA, Williams DL, Minshew NJ. Variability in Adaptive Behavior in Autism: Evidence for the Importance of Family History. *J Abnorm Child Psychol*. 2008; 36(4): 591-599.
- Kraijer D. Review of adaptive behavior studies in mentally retarded persons with autism/pervasive developmental disorder. *J Autism Dev Disord*. 2000; 30: 39-47.
- Kogan MD, Blumberg SJ, Schieve LA, Boyle CA, Perrin JM, Ghandour RM, et al. Prevalence of parent-reported diagnosis of autism spectrum disorder among children in the US, 2007. *Pediatrics*. 2009; 124: 1395-1403.
- American Psychiatric Association. *Diagnostic and statistical Manual of Mental Disorders*. 5th ed. Washington: DC: Author; 2013.
- De Bruin EI, de Nijs PF, Verheij F, Hartman CA, Ferdinand RF. Multiple complex developmental disorder delineated from PDD-NOS. *J Autism Dev Disord*. 2007; 37: 1181-1191.
- Cotton S, Richdale A. Brief report: parental descriptions of sleep problems in children with autism, Down syndrome, and Prader-Willi syndrome. *Res Dev Disabil*. 2006; 27: 151-161.
- Weijerman, ME, de Winter JP. Clinical practice: The care of children with Down syndrome. *Eur J Pediatr*. 2010; 169: 1445-1452.
- Bolte S, Poustka, F. The recognition of facial affect in autistic and schizophrenic subjects and their first-degree relatives. *Psychol Med*. 2003; 33(5): 907-915.
- Gross TF. The perception of four basic emotions in human and nonhuman faces by children with autism and other developmental disabilities. *J Abnorm Child Psychol*. 2004; 32: 469-480.