

OPEN ACCESS

EDITED BY Kerim Munir, Boston Children's Hospital, United States

REVIEWED BY
Tuba Mutluer,
Koç University, Türkiye
Ali Evren Tufan,
Abant Izzet Baysal University, Türkiye
Mashudat Bello-Mojeed,
Federal Neuro Psychiatric Hospital, Nigeria

*CORRESPONDENCE
Ravi Shankar Reddy

✓ rshankar@kku.edu.sa

RECEIVED 08 October 2024 ACCEPTED 28 November 2024 PUBLISHED 16 December 2024

CITATION

Tedla JS, Asiri F, Reddy RS, Sangadala DR, Gular K and Kakaraparthi VN (2024) Assessing the quality of life in children with autism spectrum disorder: a cross-sectional study of contributing factors. *Front. Psychiatry* 15:1507856. doi: 10.3389/fpsyt.2024.1507856

COPYRIGHT

© 2024 Tedla, Asiri, Reddy, Sangadala, Gular and Kakaraparthi. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.

Assessing the quality of life in children with autism spectrum disorder: a cross-sectional study of contributing factors

Jaya Shanker Tedla, Faisal Asiri, Ravi Shankar Reddy*, Devika Rani Sangadala, Kumar Gular and Venkata Nagarai Kakaraparthi

Program of Physical Therapy, Department of Medical Rehabilitation Sciences, College of Applied Medical Sciences, King Khalid University, Abha, Saudi Arabia

Objective: The study aimed to assess the quality of life (QoL) in children with autism spectrum disorder (ASD) compared to typically developing peers, identify key influencing factors such as socio-demographic and comorbid conditions, and examine the impact of symptom severity on QoL outcomes.

Methods: In this cross-sectional study conducted in Saudi Arabia, 75 children with ASD were compared to 75 typically developing children matched for age and gender. QoL was evaluated using the Pediatric Quality of Life Inventory (PedsQL), while the severity of autism symptoms was assessed using the Autism Diagnostic Observation Schedule (ADOS). Additional variables, including sociodemographic factors, comorbid conditions, and family environment, were collected through structured interviews and clinical assessments. Statistical analyses, including independent samples t-tests, multiple linear regression, and ANOVA, were employed to compare QoL scores, identify predictors, and assess the impact of symptom severity on QoL outcomes.

Results: The mean overall QoL score for children with ASD was 57.86 (SD = 13.25) compared to 81.67 (SD = 10.89) for typically developing children (t = -10.56, p < 0.001, Cohen's d = 1.90). Socioeconomic status (β = -0.25, t = -5.00, p < 0.001), comorbid ADHD (β = -0.35, t = -5.83, p < 0.001), and parental mental health issues (β = -0.45, t = -9.00, p < 0.001) were significant predictors of lower QoL. ANOVA results showed that children with severe autism symptoms had the lowest QoL scores (mean = 40.12, SD = 15.67; F = 20.45, p < 0.001, η^2 = 0.45).

Conclusion: Children with ASD showed significantly lower QoL, particularly in social and school functioning, highlighting the need for targeted interventions addressing core symptoms and environmental and family factors to improve outcomes.

KEYWORDS

autism spectrum disorder, quality of life, pediatric, comorbid conditions, socioeconomic status, symptom severity

Introduction

Autism Spectrum Disorder (ASD) is a neurodevelopmental condition characterized by persistent challenges in social communication and interaction, along with restricted and repetitive behaviors or interests (1). Globally, there has been a notable rise in ASD prevalence, with recent estimates indicating approximately 1 in 54 children in the United States diagnosed with ASD (1). This increase has underscored the intricate challenges faced by individuals with ASD and their families (2). The origins of ASD are multifaceted, involving a blend of genetic and environmental factors (2). The diverse spectrum of symptoms and functional challenges associated with ASD significantly influences various aspects of quality of life (QoL), necessitating a comprehensive understanding of these impacts (3).

Comorbidities are highly prevalent in children with ASD and significantly impact their QoL (4). A study by Mutluer et al. (5) reported that up to 50% of children with ASD have co-occurring ADHD, while anxiety disorders affect approximately 40% of this population (6). These conditions exacerbate the challenges faced by children with ASD, leading to more pronounced difficulties in emotional regulation, social interaction, and daily functioning, further impacting their overall quality of life (6). Understanding the prevalence and impact of these comorbidities is essential for developing targeted interventions to improve their well-being, adaptive functioning, and overall quality of life.

The severity of autism symptoms is another critical determinant of QoL in children with ASD. Symptom severity can vary widely among individuals, ranging from mild to severe, significantly impacting daily functioning and overall well-being (7). Children with milder symptoms may have better communication skills and social interactions, allowing for greater participation in typical childhood activities (8). In contrast, children with severe symptoms often experience profound difficulties in communication, social interaction, and behavior, which can lead to increased dependency on caregivers and reduced opportunities for social engagement (9). Studies have shown that higher severity of autism symptoms is associated with lower QoL, particularly in areas of social functioning and adaptive behaviors (9). This relationship underscores the importance of tailored interventions that address the specific needs of children based on the severity of their symptoms (9).

Despite the growing body of research on QoL in children with ASD, significant gaps remain in understanding its multidimensional nature (10). The educational outcomes and peer relationships are often underexplored, despite their critical role in overall well-being (11). Moreover, existing studies rarely examine how symptom severity disproportionately impacts specific QoL domains, such as the pronounced effects on social functioning compared to physical health (11). This study addresses these gaps by employing a comprehensive methodology that evaluates multiple QoL domains, incorporates standardized severity assessments, and considers key factors such as sociodemographics and comorbid conditions (12). By linking symptom severity to domain-specific QoL impacts, this research provides a nuanced understanding of how ASD affects daily life and overall well-being.

Recent research has highlighted critical limitations in traditional QoL measures for children with ASD, including their lack of specificity and cultural adaptability. Simpson et al. (13) review underscores the importance of developing ASD-specific scales to capture the unique challenges faced by this population. Similarly, the study by Chezan et al. (14) demonstrates how advanced statistical methodologies can enhance the validity of QoL assessments in ASD populations (14). Despite these advancements, many studies rely on cross-sectional data, limiting the ability to capture changes in QoL over time. Additionally, the underrepresentation of subgroups such as non-verbal children and the lack of diversity in participant samples have restricted the generalizability of findings, underscoring the need for a more inclusive approach (14). This study addresses these gaps by focusing on a comprehensive assessment of QoL and its determinants in a diverse cohort of children with ASD.

Given the critical importance of QoL for children with ASD, the study objectives were to evaluate the overall QoL in children with ASD and compare it with typically developing children, to identify key factors influencing QoL in children with ASD, including socio-demographic variables, comorbid conditions, and family environment, and to explore the relationship between ASD symptom severity and QoL. Specifically, we hypothesized that (1) children with ASD would have lower overall QoL compared to typically developing peers, (2) socio-demographic factors, comorbid conditions, and family environment would significantly impact QoL in children with ASD, and (3) greater severity of autism symptoms would be associated with lower QoL.

Materials and methods

Study setting and design

The study was conducted at Abha Tertiary Children's Hospital, a leading pediatric care center serving a diverse population in the southern region of Saudi Arabia. The hospital provides specialized neurodevelopmental and general pediatric services, ensuring access to a representative sample of children with ASD and typically developing peers. The demographic diversity of the patient population reflects a range of socioeconomic and cultural backgrounds, which enhances the generalizability of the findings to similar healthcare settings. Ethical guidelines as per the Declaration of Helsinki were strictly followed, including obtaining ethical approval (REC#234-2022) and securing written informed consent from parents, along with assent from the children involved.

Participants

The study included a total of 150 children, consisting of 75 children diagnosed with ASD and 75 typically developing children who served as a control group. An IQ threshold of 50 was selected for exclusion to minimize confounding effects related to severe intellectual disabilities, which can independently impact QoL and obscure the specific contributions of ASD-related factors (15). This

threshold also aligns with previous studies investigating QoL in children with ASD, ensuring comparability of findings. The sixmonth follow-up period was chosen to ensure clinical stability and consistency in participants' medical and developmental status, providing a reliable basis for QoL assessment (16). Typically developing children were confirmed through a combination of parent-reported developmental histories, clinical evaluations conducted by pediatricians, and standardized screening tools, such as the Ages and Stages Questionnaire (ASQ). To confirm the diagnosis of ASD, evaluations were conducted by a board-certified child and adolescent psychiatrist with expertise in neurodevelopmental disorders. The diagnosis was made according to the DSM-5 criteria, utilizing standardized diagnostic tools, including the Autism Diagnostic Observation Schedule (ADOS) and the Autism Diagnostic Interview-Revised (ADI-R).

The control group comprised typically developing children recruited from the general pediatric clinic of a specialist hospital. To ensure homogeneity, participants were screened through parent-reported developmental histories, standardized screening tools (Ages and Stages Questionnaire), and clinical evaluations conducted by pediatricians to confirm the absence of any developmental, neurological, or psychiatric conditions. Although matching and strict exclusion criteria were applied, including screening for developmental delays, neurological conditions, and psychiatric disorders using the Ages and Stages Questionnaire, challenges in ensuring complete adherence to these criteria are acknowledged. The reliance on parent-reported histories and clinical evaluations may have contributed to some inconsistencies, which are reflected in the results.

Variables

The outcome variable in this study was the quality of life (QoL) of children with ASD, assessed using the Pediatric Quality of Life Inventory (PedsQL), which measures QoL across physical health, emotional well-being, social functioning, and school functioning. Independent variables included socio-demographic factors (age, gender, socioeconomic status, and parental education), comorbid conditions (e.g., ADHD, anxiety), family environment (e.g., family structure, parental mental health), and ASD symptom severity. Additional variables assessed included type of schooling (mainstream or special education), daily living skills (independent or assisted), social skills, communication skills, and social inclusion.

The Pediatric Quality of Life Inventory (PedsQL) was used to measure QoL across four domains, using a 23-item Likert scale transformed to a 0–100 scale, with higher scores indicating better QoL. The Autism Diagnostic Observation Schedule, Second Edition (ADOS-2), assessed ASD symptom severity across communication, social interaction, and restricted behaviors. Symptom severity was categorized as mild, moderate, or severe using calibrated severity scores. Socio-demographic and family-related variables were collected through structured parent interviews and medical record reviews, while comorbid conditions were diagnosed based on DSM-5 criteria. Other variables, such as daily living skills,

communication, and social skills, were assessed through clinician evaluations and validated questionnaires.

Comorbid conditions, specifically the presence of ADHD and anxiety, were identified through medical records and parent interviews. The presence of these conditions was confirmed using standardized diagnostic criteria from the DSM-5 (17), as recorded in the child's medical history. Additionally, information regarding the family environment, including family structure (single-parent or two-parent households) and parental mental health status, was collected through structured interviews and reviewed from the medical records.

Parental mental health issues were assessed through structured clinical interviews conducted by trained clinicians, focusing on symptoms of anxiety, depression, and stress. Information was corroborated with medical records where available. Parents were categorized as having mental health issues if they reported clinically significant symptoms or had a formal diagnosis documented in their medical history. The types of therapy received by participants were categorized into three primary modalities: behavioral therapy, speech therapy, and occupational therapy. Many children received combined therapies, a common approach in ASD intervention. For analysis purposes, each child was assigned to the primary therapy type they were receiving most consistently, as reported by parents and verified through clinical records. Participants receiving significant, equally distributed components of two or more therapies were classified under a 'combined therapy' category.

Sample size calculation

The sample size calculation was conducted using G*Power statistical software to ensure sufficient power for detecting meaningful differences between groups. The primary statistical tests considered in the calculation included independent samples t-tests for comparing QoL scores between groups and ANOVA for examining the relationship between ASD symptom severity and QoL. A medium effect size (Cohen's d = 0.5) was chosen based on previous studies in similar populations, which reported medium to large effect sizes for QoL differences in children with ASD. This effect size was deemed appropriate given the anticipated variability in QoL scores across groups and the study's focus on identifying clinically significant differences. To achieve a power of 0.80 and a significance level of 0.05, a minimum of 64 participants per group was required, and additional participants were recruited to account for potential dropouts and incomplete data.

Data analysis

Data were analyzed using IBM SPSS Statistics, Version 24.0. Descriptive statistics summarized the demographic and clinical characteristics of the sample, and the data's normal distribution validated the use of parametric tests. For Objective 1, independent samples t-tests were used to compare mean QoL scores across all domains between children with ASD and typically developing

children. QoL was analyzed as a continuous variable. Multiple linear regression analysis was performed for Objective 2 to identify independent factors associated with QoL, with all predictors, including socio-demographic factors, comorbid conditions, family environment, and ASD symptom severity, entered into the model simultaneously. Beta coefficients (β) indicated the direction and magnitude of relationships, and multicollinearity checks (e.g., variance inflation factor analysis) ensured variables were not excessively correlated. For Objective 3, Pearson correlation coefficients (r) explored the relationship between ASD symptom severity (measured continuously via ADOS) and QoL scores. Regression analysis further evaluated the collective impact of independent variables while controlling for confounding factors. To examine group differences in QoL scores across mild, moderate, and severe ASD symptom categories, ANOVA was used, followed by Tukey's HSD post-hoc tests. All statistical analyses were twotailed, with a significance level of p < 0.05.

Results

A total of 150 participants were recruited for the study, comprising 75 children with ASD and 75 typically developing children. The representativeness of the control group may be limited due to the recruitment setting, which was a specialist hospital. Although stringent screening ensured the inclusion of typically developing children, this population may not fully reflect the diversity of typically developing children in the general population. All recruited participants completed the study, resulting in a 100% response rate with no

drop-outs. The children in both groups were matched for age and gender, ensuring comparability. The ASD group had a mean age of 10.50 years (SD = 2.50), while the typically developing group had a mean age of 10.30 years (SD = 2.40). The majority of participants in both groups were male, with 66.67% in the ASD group and 64.00% in the typically developing group.

Children with ASD exhibit notable differences in various demographic and clinical characteristics compared to typically developing children (Table 1). The age distribution is similar between the two groups, but a higher proportion of children with ASD are male (66.67%) compared to typically developing children (64.00%). Socioeconomic status and parental education levels are comparable between the groups. Notably, children with ASD have higher instances of comorbid conditions, with 66.67% affected, and a significant portion of these children require special education (60.00%) compared to typically developing children, most of whom attend mainstream schools (86.67%). Children with ASD also display lower levels of daily living skills, social skills, and communication skills, with 73.33% needing assistance for daily living and 40.00% being nonverbal. The observed findings in Table 1 may partially reflect limitations in the rigorous exclusion of children with undiagnosed developmental, neurological, or psychiatric conditions, despite the use of standardized screening tools.

Children diagnosed with ASD exhibit markedly diminished QoL in comparison to their typically developing peers across various domains (see Table 2). In terms of physical health, the average score for children with ASD was 65.32, significantly lower than the mean score of 85.67 for typically developing children, indicating a mild to moderate effect size of 1.75 and a highly

TABLE 1 Demographic and clinical characteristics of children with ASD and typically developing children.

Characteristics	Children with ASD (n=75)	Typically Developing Children (n=75)	p-value
Age (years)	10.50 ± 2.50	10.30 ± 2.40	0.651
Gender (Male/Female)	50/25 (66.67%/33.33%)	48/27 (64.00%/36.00%)	0.732
Socioeconomic Status (Low/Middle/High)	20/40/15 (26.67%/53.33%/20.00%)	18/42/15 (24.00%/56.00%/20.00%)	0.924
Parental Education (High School/College/Graduate)	10/45/20 (13.33%/60.00%/26.67%)	8/47/20 (10.67%/62.67%/26.67%)	0.855
Severity of ASD Symptoms (Mild/Moderate/Severe)	20/40/15 (26.67%/53.33%/20.00%)	N/A	N/A
Comorbid Conditions (Yes/No)	50/25 (66.67%/33.33%)	N/A	N/A
Family Structure (Single-parent/Two-parent)	20/55 (26.67%/73.33%)	15/60 (20.00%/80.00%)	0.612
Parental Mental Health Issues (Yes/No)	30/45 (40.00%/60.00%)	N/A	N/A
Type of Schooling (Mainstream/Special Education)	30/45 (40.00%/60.00%)	65/10 (86.67%/13.33%)	0.001
Therapy Received (Behavioral/Speech/Occupational)	40/25/10 (53.33%/33.33%/13.33%)	N/A	N/A
Daily Living Skills (Independent/Assisted)	20/55 (26.67%/73.33%)	60/15 (80.00%/20.00%)	0.003
Social Skills (High/Medium/Low)	15/40/20 (20.00%/53.33%/26.67%)	40/20/15 (53.33%/26.67%/20.00%)	0.002
Communication Skills (Verbal/Non-verbal)	45/30 (60.00%/40.00%)	70/5 (93.33%/6.67%)	0.011
School Support Services (Yes/No)	35/40 (46.67%/53.33%)	60/15 (80.00%/20.00%)	0.005
Peer Relationships (Good/Average/Poor)	10/30/35 (13.33%/40.00%/46.67%)	45/20/10 (60.00%/26.67%/13.33%)	0.001
Social Inclusion (Yes/No)	50/25 (66.67%/33.33%)	65/10 (86.67%/13.33%)	0.001

ASD, Autism Spectrum Disorder.

N/A, Not Available.

TABLE 2 Comparison of overall quality of life between children with ASD and typically developing children

Quality of Life Domains	Children with ASD (Mean ± SD)	Typically Developing Children (Mean <u>+</u> SD)	t-value	p-value	Cohen's d
Physical Health	65.32 ± 12.45	85.67 ± 10.12	-9.87	<0.001	1.75
Emotional Well-being	58.76 ± 14.32	78.23 ± 11.45	-8.76	<0.001	1.60
Social Functioning	50.22 ± 15.67	82.34 ± 10.76	-11.45	<0.001	2.10
School Functioning	55.13 ± 13.89	80.45 ± 11.23	-9.34	<0.001	1.80
Overall Quality of Life	57.86 ± 13.25	81.67 ± 10.89	-10.56	<0.001	1.90

ASD, Autism Spectrum Disorder; SD, Standard Deviation; t-value, t-test value; p-value, Probability value; Cohen's d, Effect size.

significant t-value of -9.87 (p < 0.001). Similarly, emotional wellbeing scores were notably lower among children with ASD, averaging 58.76 compared to 78.23 for their typically developing counterparts, with a Cohen's d of 1.60 and a t-value of -8.76 (p < 0.001). Social functioning showed the most substantial disparity, where children with ASD scored 50.22 on average, in contrast to 82.34 for their typically developing peers, resulting in a Cohen's d of 2.10 and a t-value of -11.45 (p < 0.001). School functioning followed a similar pattern, with children with ASD scoring 55.13 versus 80.45 for typically developing children, with an effect size of 1.80 and a t-value of -9.34 (p < 0.001). The overall QoL was markedly lower for children diagnosed with ASD, evidenced by a mean score of 57.86 compared to 81.67 for typically developing children. This finding yielded a substantial Cohen's d of 1.90 and a highly significant t-value of -10.56 (p < 0.001).

The multiple regression analysis revealed several significant factors influencing the quality of life (QoL) in children with ASD (Table 3 and Figure 1). Lower socioeconomic status (β = -0.25, p < 0.001), comorbid ADHD (β = -0.35, p < 0.001), comorbid anxiety (β = -0.40, p < 0.001), single-parent family structure (β = -0.20, p < 0.001), and parental mental health issues (β = -0.45, p < 0.001) were associated with lower QoL scores. Conversely, higher parental education was positively associated with QoL (β = 0.30, p < 0.001).

Table 4 and Figure 2 provides detailed insights into the relationship between the severity of autism symptoms and the QoL in children diagnosed with ASD. Children with mild ASD symptoms exhibit notably higher overall QoL (mean = 70.45) compared to those with moderate (mean = 55.23) and severe symptoms (mean = 40.12), with all comparisons demonstrating strong statistical significance (p < 0.001). Scores for physical health, emotional well-being, social functioning, and school functioning decline progressively with increasing symptom severity. Effect sizes

 (η^2) ranging from 0.35 to 0.45 underscore the substantial impact of symptom severity on these QoL domains.

Discussion

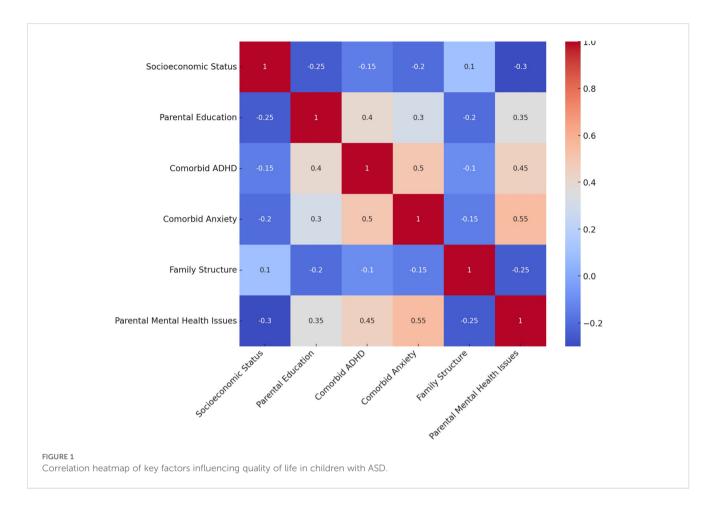
In this study, we aimed to assess the QoL in children with ASD and compare it with typically developing children, identify key factors influencing their QoL, and explore the relationship between the severity of autism symptoms and QoL. The differences in mean QoL scores between children with ASD and typically developing peers were statistically significant across all domains, with p-values <0.001. However, the effect sizes, as measured by Cohen's d, ranged from 0.60 to 0.79, indicating moderate effects rather than large effects, based on standard guidelines. These moderate effect sizes suggest that while the differences are meaningful, other factors may contribute to QoL disparities and warrant further exploration. Key factors influencing the QoL in children with ASD include socioeconomic status, parental education, comorbid ADHD and anxiety, family structure, and parental mental health issues, all of which were significantly associated with variations in QoL. Furthermore, the severity of autism symptoms was inversely related to QoL, with children experiencing mild symptoms reporting higher QoL than those with moderate or severe symptoms, highlighting the substantial impact of symptom severity on physical health, emotional well-being, social functioning, and school functioning. These findings emphasize the need for targeted interventions to address these factors and improve the QoL for children with ASD.

The findings indicate that children with ASD experience a significantly lower QoL across various domains, including physical health, emotional well-being, social functioning, and school

TABLE 3 Multiple regression analysis of factors influencing quality of life in children with ASD.

Variable	Regression Coefficient (β)	Standard Error (SE)	t-value	p-value
Socioeconomic Status	-0.25	0.05	-5.00	<0.001
Parental Education	0.30	0.04	7.50	<0.001
Comorbid ADHD	-0.35	0.06	-5.83	<0.001
Comorbid Anxiety	-0.40	0.05	-8.00	<0.001
Family Structure (Single-parent)	-0.20	0.04	-5.00	<0.001
Parental Mental Health Issues	-0.45	0.05	-9.00	<0.001

ASD, Autism Spectrum Disorder; SE, Standard Error; t-value, t-test value; p-value, Probability value; β : Beta Coefficient.



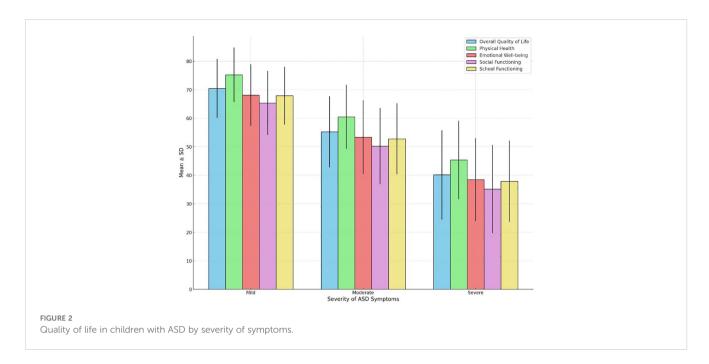
functioning, compared to typically developing children (18). Several factors may contribute to these disparities. Children with ASD often face challenges in physical health due to co-occurring conditions, dietary restrictions, and lower levels of physical activity (19). Emotional well-being is impacted by the higher prevalence of anxiety, depression, and difficulty in processing emotions (20). Social functioning deficits arise from impaired social communication skills and difficulties in forming peer relationships (21). School functioning is adversely affected by the need for specialized educational support and the challenges in coping with the academic and social demands of a mainstream school environment (22). These cumulative challenges significantly lower the overall QoL for children with ASD. These results are consistent with previous studies that have documented similar findings. For instance, Kuhlthau et al. (2010) reported that children with ASD have poorer health-related QoL compared to their peers, primarily due to social and emotional difficulties (23). Additionally, research by Eapen et al. (24) highlighted the impact of comorbid conditions such as anxiety and ADHD on the overall QoL in children with ASD (24). The findings of our study align with these previous observations, underscoring the need for comprehensive interventions that address not only the core symptoms of ASD but also the associated physical, emotional, and social challenges to improve the QoL for these children (25).

The study identifies several key factors that significantly influence the QoL in children with ASD. Socioeconomic status is a notable factor, with lower socioeconomic status correlating with reduced QoL, likely due to limited access to resources, healthcare, and educational opportunities (26). Higher parental education levels are associated with better QoL, possibly because educated parents are better equipped to navigate the healthcare and educational systems to secure necessary services for their children (27). Comorbid conditions, particularly ADHD and anxiety, substantially reduce the QoL due to their

TABLE 4 Relationship between severity of autism symptoms and quality of life in children with ASD.

Severity of ASD Symptoms	Number of Chil- dren (n)	Overall Quality of Life (Mean ± SD)	Physical Health (Mean <u>+</u> SD)	Emotional Well-being (Mean <u>+</u> SD)	Social Functioning (Mean <u>+</u> SD)	School Functioning (Mean <u>+</u> SD)	F- value	p- value	Effect Size (η²)
Mild	25	70.45 ± 10.34	75.23 ± 9.56	68.12 ± 10.78	65.34 ± 11.23	67.89 ± 10.12	15.34	<0.001	0.35
Moderate	30	55.23 ± 12.45	60.45 ± 11.23	53.34 ± 12.90	50.23 ± 13.34	52.78 ± 12.45	18.23	<0.001	0.40
Severe	20	40.12 ± 15.67	45.34 ± 13.78	38.45 ± 14.56	35.12 ± 15.45	37.89 ± 14.23	20.45	<0.001	0.45

 $ASD, \ Autism \ Spectrum \ Disorder; \ SD, \ Standard \ Deviation; \ F-value, \ F-test \ value; \ p-value, \ Probability \ value; \ \eta^2, \ Eta \ Squared \ (Effect \ Size).$



additional psychological and behavioral burdens (28). The impact of family structure is evident, with children from single-parent families experiencing a lower QoL, likely due to reduced social and financial support (29). Furthermore, parental mental health issues significantly decrease the QoL in children with ASD, as these issues can affect the overall family environment and the level of support provided to the child (30). These findings are consistent with previous research, further validating the results of this study. For instance, Mathew et al. (31) found that children with ASD from lower socioeconomic backgrounds had poorer health outcomes and reduced QoL, highlighting the influence of socioeconomic status (31). Similarly, Koukouriki et al. (32) reported that higher parental education levels are linked to better health-related QoL in children with ASD, supporting the current study's findings (32). The detrimental effects of comorbid conditions such as ADHD and anxiety on QoL have been well-documented by D'Agati et al. (33), who emphasized the additional challenges these conditions pose (33). The impact of family structure is also supported by research from Blackstock et al. (34), who observed that single-parent families face more significant difficulties, affecting the child's well-being (34). Lastly, the influence of parental mental health issues on child outcomes has been substantiated by studies like those of Martin et al. (35), which underscore the importance of addressing parental mental health to improve the QoL in children with ASD (35).

The study reveals a clear relationship between the severity of autism symptoms and the QoL in children with ASD (36). Children with mild ASD symptoms report a significantly higher overall QoL (mean = 70.45) compared to those with moderate (mean = 55.23) and severe symptoms (mean = 40.12), with all comparisons showing strong statistical significance (p < 0.001). As symptom severity increases, scores in physical health, emotional well-being, social functioning, and school functioning all decline markedly (37). This trend can be attributed to the greater challenges faced by children with more severe symptoms, including more pronounced difficulties in communication, social interaction, and behavior, which in turn

affect their ability to engage in daily activities, form relationships, and succeed in academic settings (38). The effect sizes (η^2) range from 0.35 to 0.45, indicating a substantial impact of symptom severity on these domains of QoL. These findings are corroborated by previous research. For instance, Yerys et al. (36) found that higher severity of autism symptoms is associated with lower QoL, particularly in areas of social functioning and adaptive behaviors (36). Similarly, studies by Stole et al. (39) and Young et al. (40) support the notion that children with more severe autism symptoms experience greater difficulties that negatively impact their overall well-being and daily functioning (39, 40). These studies highlight the cumulative burden of severe autism symptoms on various aspects of life, emphasizing the need for targeted interventions that address the specific needs of children based on their symptom severity (41). This alignment with previous research underscores the reliability and validity of the current study's findings, reinforcing the importance of a nuanced approach to support and intervention for children with ASD.

The findings of this study align with prior research, such as those by Eapen et al. (42) and Mavroeidi et al. (43), which reported lower QoL in children with ASD compared to typically developing peers. However, the unique socio-cultural context of the study population in Saudi Arabia may influence generalizability. Cultural factors, including familial structures and societal stigma, can shape the experiences of children with ASD differently than in Western settings. The extended family support networks in Saudi Arabia may enhance emotional wellbeing, while limited access to specialized services exacerbates challenges in school and social functioning. Comparisons with Western studies, where inclusive policies and broader resources often lead to better school functioning outcomes, highlight the need for region-specific strategies. Future research should examine QoL across diverse socio-cultural settings to better understand the influence of contextual factors.

This study highlights the urgent need for targeted interventions to address the significantly lower QoL observed in children with ASD compared to their typically developing peers. Key findings emphasize

the importance of addressing socioeconomic disparities through equitable access to resources, such as subsidized special education, inclusive teacher training, and individualized education plans. Community-based programs, including parent-led support groups and peer-mediated social skills interventions, can further bridge gaps in service access. To mitigate the impact of parental mental health issues, family-centered approaches such as mental health counseling, stress management workshops, respite care, and strengthened community support are essential. The influence of socioeconomic status, parental education, comorbid conditions, and family environment underscores critical areas for clinical focus, while the correlation between symptom severity and QoL highlights the need for personalized treatment plans.

This study provides valuable insights into the QoL of children with ASD, but several limitations should be noted. The reliance on parentreported QoL measures may introduce subjective bias, as parents' perceptions may not fully capture the child's lived experiences. Although standardized tools like the PedsQL were used to mitigate this, future studies should incorporate direct child assessments for a more comprehensive perspective. The sample was exclusively drawn from Abha Tertiary Children's Hospital, limiting the generalizability of findings to other regions or healthcare settings and potentially missing broader cultural and socioeconomic diversity. The study did not evaluate specific ASD symptoms, such as restrictive behaviors or sensory issues, which are critical determinants of QoL, limiting the depth of insights. Additionally, the recruitment of typically developing children from the outpatient clinic of a specialist hospital may not reflect the general population, and challenges in rigorously excluding undiagnosed developmental or psychiatric conditions remain. Future research should include community-based recruitment, more comprehensive assessments, and broader population diversity to enhance representativeness and understanding.

Conclusion

In conclusion, this study provides compelling evidence that children with ASD experience a significantly lower QoL compared to their typically developing peers, across all measured domains including physical health, emotional well-being, social functioning, and school functioning. Key factors such as socioeconomic status, parental education, comorbid ADHD and anxiety, family structure, and parental mental health issues substantially influence these outcomes. Additionally, the severity of autism symptoms is inversely related to QoL, with greater symptom severity correlating with more pronounced impairments. These findings highlight the critical need for targeted, multifaceted interventions that address both the core symptoms of ASD and the associated socio-demographic and familial factors, to enhance the overall well-being and life satisfaction of children with ASD.

Data availability statement

The data supporting the findings of this study are available in the "ZENODO" repository at DOI: 10.5281/zenodo.13381005.

Ethics statement

The studies involving humans were approved by King Khalid University Ethics Committee. The studies were conducted in accordance with the local legislation and institutional requirements. Written informed consent for participation in this study was provided by the participants' legal guardians/next of kin. Written informed consent was obtained from the individual(s) for the publication of any potentially identifiable images or data included in this article.

Author contributions

JT: Conceptualization, Data curation, Formal analysis, Resources, Writing – original draft, Writing – review & editing. FA: Conceptualization, Data curation, Formal analysis, Funding acquisition, Methodology, Project administration, Resources, Writing – original draft, Writing – review & editing. RR: Conceptualization, Data curation, Formal analysis, Investigation, Methodology, Writing – original draft, Writing – review & editing. DS: Conceptualization, Data curation, Methodology, Writing – original draft, Writing – review & editing. KG: Conceptualization, Data curation, Formal analysis, Methodology, Writing – original draft, Writing – review & editing. VK: Conceptualization, Data curation, Formal analysis, Methodology, Writing – original draft, Writing – review & editing.

Funding

The author(s) declare financial support was received for the research, authorship, and/or publication of this article. King Salman Center for Disability Research, No KSRG- 2022-005.

Acknowledgments

The authors extend their appreciation to the King Salman Center for Disability Research for funding this work through Research Group No KSRG- 2022-005.

Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Generative AI statement

The author(s) declare that no Generative AI was used in the creation of this manuscript.

Publisher's note

All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article, or claim that may be made by its manufacturer, is not guaranteed or endorsed by the publisher.

References

- 1. Joon P, Kumar A, Parle M. What is autism? $\it Pharmacol~Rep.~(2021)~73:1255-64.$ doi: 10.1007/s43440-021-00244-0
- 2. Gandhi T, Lee CC. Neural mechanisms underlying repetitive behaviors in rodent models of autism spectrum disorders. *Front Cell Neurosci.* (2021) 14:592710. doi: 10.3389/fncel.2020.592710
- 3. Tobin MC, Drager KD, Richardson LF. A systematic review of social participation for adults with autism spectrum disorders: Support, social functioning, and quality of life. *Res Autism Spectr Disord*. (2014) 8:214–29. doi: 10.1016/j.rasd.2013.12.002
- 4. Sáez-Suanes GP, Álvarez-Couto M. Factors associated with quality of life in adults with autism spectrum disorder: A systematic review. *Rev J Autism Dev Disord.* (2021) 9:1–13. doi: 10.1007/s40489-021-00254-x
- 5. Mutluer T, Aslan Genç H, Özcan Morey A, Yapici Eser H, Ertinmaz B, Can M, et al. Population-based psychiatric comorbidity in children and adolescents with autism spectrum disorder: A meta-analysis. *Front Psychiatry*. (2022) 13:856208. doi: 10.3389/fpsyt.2022.856208
- 6. Popow C, Ohmann S, Plener P. Practitioner's review: Medication for children and adolescents with autism spectrum disorder (ASD) and comorbid conditions. *Neuropsychiatrie.* (2021) 35:113–34. doi: 10.1007/s40211-021-00395-9
- 7. Baj J, Flieger W, Flieger M, Forma A, Sitarz E, Skórzyńska-Dziduszko K, et al. Autism spectrum disorder: Trace elements imbalances and the pathogenesis and severity of autistic symptoms. *Neurosci Biobehav Rev.* (2021) 129:117–32. doi: 10.1016/j.neubiorev.2021.07.029
- 8. Rieske RD, Matson JL. Parental age at conception and the relationship with severity of autism symptoms. *Dev neurorehabilitation*. (2020) 23:265–70. doi: 10.1080/17518423.2019.1645222
- 9. Waizbard-Bartov E, Ferrer E, Young GS, Heath B, Rogers S, Wu Nordahl C, et al. Trajectories of autism symptom severity change during early childhood. *J Autism Dev Disord*. (2021) 51:227–42. doi: 10.1007/s10803-020-04526-z
- 10. Samadi H, Samadi SA. Understanding different aspects of caregiving for individuals with autism spectrum disorders (ASDs) a narrative review of the literature. *Brain Sci.* (2020) 10:557. doi: 10.3390/brainsci10080557
- 11. Shepherd D, Buchwald K, Siegert RJ, Vignes M. Using network analysis to identify factors influencing the heath-related quality of life of parents caring for an autistic child. *Res Dev Disabilities*. (2024) 152:104808. doi: 10.1016/j.ridd.2024.104808
- 12. Mahjoob M, Paul T, Carbone J, Bokadia H, Cardy RE, Kassam S, et al. Predictors of health-related quality of life in neurodivergent children: A systematic review. *Clin Child Family Psychol Review*. (2024) 27:91–129. doi: 10.1007/s10567-023-00462-3
- 13. Simpson K, Paynter J, Westerveld M, van der Meer L, Patrick L, Hogg G, et al. Time to change how we measure quality of life and well-being in autism: A systematic review. *Rev J Autism Dev Disord*. (2024) 2024:1–16. doi: 10.1007/s40489-024-00440-7
- 14. Chezan LC, Liu J, Gao R, Drasgow E. Investigating the quality of life for children with autism spectrum disorder scale using Rasch methodology. *Autism Res.* (2023) 16:2172–83. doi: 10.1002/aur.v16.11
- 15. Rice C, Carpenter L, Morrier M, Lord C, DiRienzo M, Boan A, et al. Defining in detail and evaluating reliability of DSM-5 criteria for autism spectrum disorder (ASD) among children. *J Autism Dev Disord*. (2022) 52:5308–20. doi: 10.1007/s10803-021-05377-y
- 16. Katusic MZ, Myers SM, Weaver AL, Voigt RG. IQ in autism spectrum disorder: a population-based birth cohort study. *Pediatrics*. (2021) 148:1–9. doi: 10.1542/peds.2020-049899
- 17. Coxe S, Sibley MH. Harmonizing DSM-IV and DSM-5 versions of ADHD "A Criteria": An item response theory analysis. *Assessment*. (2023) 30:606–17. doi: 10.1177/10731911211061299
- 18. Menezes M, Mazurek MO. Associations between domains of health-related quality of life and co-occurring emotional and behavioral problems in youth with autism spectrum disorder. *Res Autism Spectr Disord.* (2021) 82:101740. doi: 10.1016/j.rasd.2021.101740
- 19. Donaghy B, Moore D, Green J. Co-occurring physical health challenges in neurodivergent children and young people: A topical review and recommendation. *Child Care Practice.* (2023) 29:3–21. doi: 10.1080/13575279.2022.2149471
- 20. Dunn K, Rydzewska E, Fleming M, Cooper S-A. Prevalence of mental health conditions, sensory impairments and physical disability in people with co-occurring intellectual disabilities and autism compared with other people: a cross-sectional total population study in Scotland. *BMJ Open.* (2020) 10:e035280. doi: 10.1136/bmjopen-2019-035280

- 21. Øzerk K, Özerk G, Silveira-Zaldivar T. Developing social skills and social competence in children with autism. *Int Electronic J Elementary Education*. (2021) 13:341–63. doi: 10.26822/iejee.2021.195
- 22. Pascoe MI, Forbes K, de la Roche L, Derby B, Psaradellis E, Anagnostou E, et al. Exploring the association between social skills struggles and social communication difficulties and depression in youth with autism spectrum disorder. *Autism Res.* (2023) 16:2160–71. doi: 10.1002/aur.v16.11
- 23. 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) 40:721–9. doi: 10.1007/s10803-009-0921-2
- 24. Eapen V, Snedden C, Črnčec R, Pick A, Sachdev P. Tourette syndrome, comorbidities and quality of life. *Aust New Z J Psychiatry*. (2016) 50:82–93. doi: 10.1177/0004867415594429
- 25. Fong VC, Gardiner E, Iarocci G. Can a combination of mental health services and ADL therapies improve quality of life in families of children with autism spectrum disorder? *Qual Life Res.* (2020) 29:2161–70. doi: 10.1007/s11136-020-02440-6
- 26. Sun X, Allison C, Auyeung B, Baron-Cohen S, Brayne C. Parental concerns, socioeconomic status, and the risk of autism spectrum conditions in a population-based study. *Res Dev Disabilities*. (2014) 35:3678–88. doi: 10.1016/j.ridd.2014.07.037
- 27. Hrdlicka M, Vacova M, Oslejskova H, Gondzova V, Vadlejchova I, Kocourkova J, et al. Age at diagnosis of autism spectrum disorders: is there an association with socioeconomic status and family self-education about autism? *Neuropsychiatr Dis Treat.* (2016) 12:1639–44. doi: 10.2147/NDT.S107239
- 28. Katzman MA, Bilkey TS, Chokka PR, Fallu A, Klassen LJ. Adult ADHD and comorbid disorders: clinical implications of a dimensional approach. *BMC Psychiatry*. (2017) 17:1–15. doi: 10.1186/s12888-017-1463-3
- 29. Coghill D. The impact of medications on quality of life in attention-deficit hyperactivity disorder: a systematic review. *CNS Drugs*. (2010) 24:843–66. doi: 10.2165/11537450-00000000-00000
- 30. Hsiao Y-J. Pathways to mental health-related quality of life for parents of children with autism spectrum disorder: roles of parental stress, children's performance, medical support, and neighbor support. *Res Autism Spectr Disord*. (2016) 23:122–30. doi: 10.1016/j.rasd.2015.10.008
- 31. Mathew NE, Burton KL, Schierbeek A, Črnčec R, Walter A, Eapen V. Parenting preschoolers with autism: Socioeconomic influences on wellbeing and sense of competence. *World J Psychiatry*. (2019) 9:30. doi: 10.5498/wjp.v9.i2.30
- 32. Koukouriki E, Soulis S-G. Self-reported Health-related quality of life (HRQOL) and anxiety among greek school-age siblings of individuals with autism spectrum disorders (ASD) in relation to parental mental health and social support. *J Autism Dev Disord.* (2020) 50:2913–30. doi: 10.1007/s10803-020-04395-6
- 33. D'Agati E, Curatolo P, Mazzone L. Comorbidity between ADHD and anxiety disorders across the lifespan. *Int J Psychiatry Clin Practice*. (2019) 23:238–44. doi: 10.1080/13651501.2019.1628277
- 34. Blackstock PM. The Perceptions of Single Parents Raising a Child Diagnosed With Attention Deficit Hyperactivity Disorder With Behavioral and Emotional Concerns. Minneapolis, Minnesota, USA: Walden University (2024).
- 35. Martin CA, Papadopoulos N, Chellew T, Rinehart NJ, Sciberras E. Associations between parenting stress, parent mental health and child sleep problems for children with ADHD and ASD: Systematic review. *Res Dev disabilities.* (2019) 93:103463. doi: 10.1016/j.ridd.2019.103463
- 36. Yerys BE, McQuaid GA, Lee NR, Wallace GL. Co-occurring ADHD symptoms in autistic adults are associated with less independence in daily living activities and lower subjective quality of life. *Autism.* (2022) 26:2188–95. doi: 10.1177/13623613221112198
- 37. Davoody S, Goeschl S, Dolatshahi M, Davari-Ashtiani R, Saffarpour R, Sodeifian F, et al. Relation between ADHD and COVID-19: a narrative review to guide advancing clinical research and therapy. *Iranian J Psychiatry.* (2022) 17:110. doi: 10.18502/ijps.v17i1.8055
- 38. Arildskov TW, Thomsen PH, Sonuga-Barke EJ, Lambek R, Østergaard SD, Virring A. Is attention-deficit/hyperactivity disorder (ADHD) a dimension or a category? What does the relationship between ADHD traits and psychosocial quality of life tell us? *J Attention Disord*. (2024) 28:1035–44. doi: 10.1177/10870547231222228
- 39. Stolte M, Trindade-Pons V, Vlaming P, Jakobi B, Franke B, Kroesbergen EH, et al. Characterizing creative thinking and creative achievements in relation to

symptoms of attention-deficit/hyperactivity disorder and autism spectrum disorder. Front Psychiatry. (2022) 13:909202. doi: 10.3389/fpsyt.2022.909202

- 40. Young S, Hollingdale J, Absoud M, Bolton P, Branney P, Colley W, et al. Guidance for identification and treatment of individuals with attention deficit/ hyperactivity disorder and autism spectrum disorder based upon expert consensus. *BMC Med.* (2020) 18:1–29. doi: 10.1186/s12916-020-01585-y
- 41. Young S, Adamo N, Ásgeirsdóttir BB, Branney P, Beckett M, Colley W, et al. Females with ADHD: An expert consensus statement taking a lifespan approach providing guidance for the identification and treatment of attention-deficit/
- hyperactivity disorder in girls and women. BMC Psychiatry. (2020) 20:1–27. doi: $10.1186/\mathrm{s}12888-020-02707-9$
- 42. Eapen V, Islam R, Azim SI, Masi A, Klein L, Karlov L. Factors impacting parental quality of life in preschool children on the autism spectrum. *J Autism Dev Disord*. (2024) 54:943–54. doi: 10.1007/s10803-022-05848-w
- 43. Mavroeidi N, Sifnaios C, Ntinou A, Iatrou G, Konstantakopoulou O, Merino Martinez M, et al. Exploring the potential impact of training on short-term quality of life and stress of parents of children with autism: the integrative parents' Autism training module. *Int J Environ Res Public Health*. (2024) 21:474. doi: 10.3390/ijerph21040474