

Depression And Anxiety In Patients With Juvenile Idiopathic Arthritis: Current Insights And Impact On Quality Of Life, A Systematic Review

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Abstract: Depression and anxiety are prevalent in children with rheumatologic diseases, including juvenile idiopathic arthritis (JIA). However, prevalence rates and the relationship with disease outcomes, including quality of life are conflicting in the early literature. To review the current literature, determine gaps in our knowledge, and identify areas in need of further investigation, we conducted a systematic review of studies examining depression and anxiety symptoms among children with JIA and the impact these symptoms may have on disease outcomes and quality of life. Six electronic databases were searched up until January 2019. Of 799 potential articles, 60 articles were included with the main focus on 28 articles from 2009 to 2019, to concentrate on the most current evidence. We found that JIA patients experience symptoms of depression and anxiety similar to other childhood chronic diseases and at higher rates than in healthy children. Patients who experience these symptoms have worse quality of life, with some evidence pointing to depression and anxiety symptoms having a greater impact on quality of life than other disease features, such as active joint count. Family members of JIA patients experience high rates of anxiety and depression symptoms which may impact their child's mental health and pain symptoms related to JIA. Conflicting reports of associations between depression/anxiety symptoms and disease features/disease outcomes and a paucity of longitudinal studies investigating the impact of treatment on mental health symptoms indicate areas in need of further research to effectively identify patients at greatest risk of depression and anxiety and to better understand how to treat and prevent these symptoms in youth with JIA. Family mental health should also be considered in investigations concerning mental health and disease outcomes of children with JIA.

Keywords: pediatric rheumatology, mental health, mood disorder, arthritis, autoimmune disease

Introduction

Juvenile idiopathic arthritis (JIA), diagnosed in children under 16, affects an estimated 1.7 to 8.4 million children worldwide.¹ Incidence rates are estimated to be around 1.6–23 per 100,000.² All races, ethnicities, and ages can be affected. However, females tend to be more frequently affected than males. Children with any of the six different classifications of JIA: oligoarticular (<5 joints), polyarticular (>4 joints), systemic, psoriatic, enthesitis-related (ERA), and undifferentiated can experience musculoskeletal pain and disability, fatigue, and symptoms of psychological distress such as depression and anxiety.

Currently, the incidence of mental health disorders is on the rise within the general United States (US) population. The National Survey on Drug Use and

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Health (NSDUH) found that from 2011 to 2016 the prevalence of depression in adolescents increased from 8.3% to 12.9%.³ Children who suffer from chronic disease have higher rates of anxiety and depression than the general population.^{4–6} In addition, children with chronic disease have less favorable courses regarding their mental health symptoms over time.⁷

Reviews of the earlier literature on mental health in children with juvenile arthritis indicated that symptoms of depression and anxiety are prevalent in this population and established the inter-connectedness of parental and child mental health in families affected by JIA.^{8–10} However, studies have reported a wide range of rates for both symptoms and diagnoses of mental health disorders and have conflicted regarding the relationship between these symptoms, disease features, and disease outcomes. Many were limited by small sample sizes, with some studies including fewer than 30 patients,^{11–13} few included significant proportions of racial and ethnic minorities, and few were conducted outside the US and Europe.⁸ Many studies examined children with JIA as part of a general cohort of children with other rheumatologic diseases,^{11,14–18} restricting our understanding of the relationship of psychological symptoms to disease-specific outcomes. In the interim, our treatments and even our classifications of chronic juvenile inflammatory arthritis have changed. The past decade of research in pediatric rheumatology has benefited from the development of longitudinal cohorts (institutional, multicenter, and even international) of both children and adults who suffer from diseases such as JIA.^{19–21} These studies have enhanced our understanding of the epidemiology, risk factors, and outcomes associated with rheumatologic diseases. Given the changes in the field and the general rising prevalence of mental health disorders, we aimed to conduct a systematic review of the recent literature of depression and anxiety in JIA to summarize our knowledge to date, determine gaps in our understanding of depression and anxiety in this population, and identify areas in need of further investigation that will improve care of youth with JIA.

Methods

Eligibility Criteria

Studies were included in the review if they (1) involved children, adolescents, and/or adults with JIA, (2) focused on depression and/or anxiety symptoms, and (3) were an original, primary source of research available in English.

Data Sources

The following databases were searched to identify relevant studies for this systematic review: Ovid MEDLINE, PsycINFO, Web of Science Core Collection, Scopus, the Cumulative Index of Nursing and Allied Health Literature (CINAHL), and Cochrane Database of Systematic Reviews and Cochrane Register of Controlled Trials. The search included literature up until January 2019 and was conducted by a medical librarian in February 2019.

Search Strategy And Study Selection

The published literature was searched using strategies designed by a medical librarian for the concepts of juvenile idiopathic arthritis or juvenile rheumatoid arthritis and depression or anxiety. Search terms related to juvenile rheumatoid arthritis included the MeSH term “arthritis, juvenile” and keywords related to childhood, childhood-onset or juvenile combined with terms for arthritis including arthrosis, rheumatism, and Still’s disease. Search terms related to mental health included the MeSH terms “depression”, “depressive disorder, major”, “anxiety” and “anxiety disorder”, and keywords such as depressive, depressed, anxious, or anxiety.

The initial search was done in Ovid MEDLINE, and the search strategy was peer reviewed by another librarian prior to being translated into subsequent databases. A total of 799 unique references were found. A flow-chart of the study identification process is provided in [Figure 1](#). Whereas, full search strategies are provided in the [supplementary appendix Table S1](#).

The remaining 799 articles were organized in Rayyan Qatar Computing Research Institute (QCRI), a web-based app for systematic reviews.²² All decisions for study inclusions and exclusions were taken by consensus based on how well the study fit the inclusion criteria. Each abstract was screened by two of three authors (D.F., M.R., and T.R.) in a blinded fashion. Discrepancies were resolved by the third author who had not yet screened the article. This led to a total of 158 articles for potential inclusion in the review. Finally, the full texts of all the potentially included articles were independently reviewed by three authors (D.F., M.R., and T.R.) to determine the final selection of articles. All initially included articles were reviewed, but focus was placed on those written from 2009 to 2019 in order to concentrate on the most current evidence. Full-text articles were excluded for the following reasons: (1) a duplicate article missed in the initial screening, (2) an article was found to be unavailable in

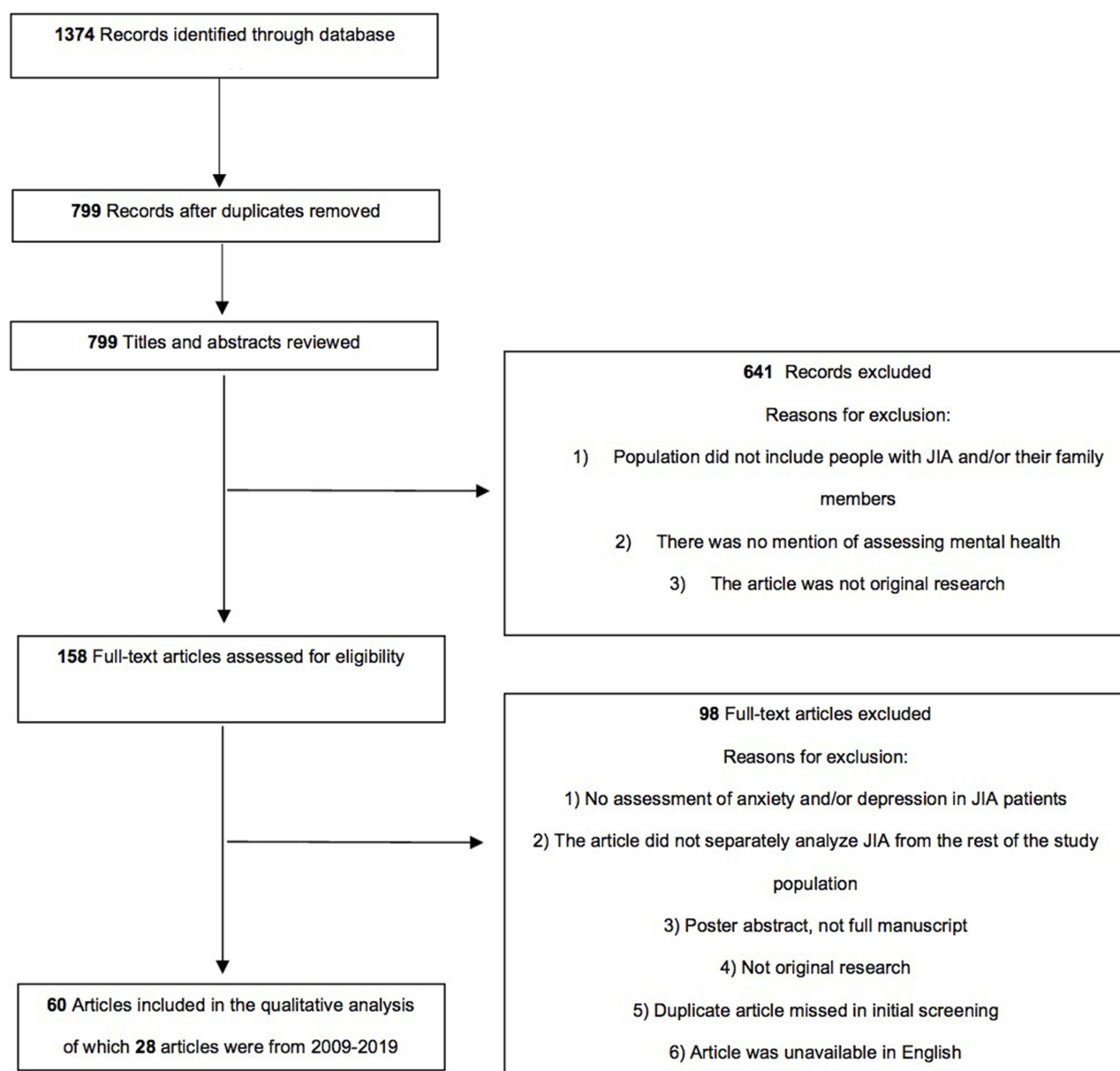


Figure 1 Flowchart of selection process for study inclusion in the systematic review.

the English language, (3) there was no assessment of anxiety and/or depression in JIA patients, (4) an article did not analyze JIA separately from the rest of the study population, (5) a poster abstract and not a full manuscript, and/or (6) an article was not original research. At the conclusion of the search and screening process, 60 articles were selected for inclusion of which 28 articles were written between 2009 and 2019.

Data Extraction

The data extracted from the final articles included the title, author(s), year of publication, country in which the

study was conducted, patient characteristics (age, gender, sample size, race, ethnicity, disease(s), subtype of JIA, disease duration, disease activity, and medication(s)), family socioeconomic characteristics (parent's educational level and family income), and findings related to depression and anxiety. Reviewers recorded the assessments that were used to screen for anxiety and/or depression and measure the quality of life in both patients and families along with the measures of JIA disease characteristics such as disease activity, mobility, pain, and global health scores. Lastly, reasons for exclusions and limitations were included as well.

Results

Study Characteristics

Overall, 28 articles from 2009 to 2019 were included (Table 1): three-fourths addressed the prevalence of depression, half addressed the prevalence of anxiety, and fewer than one-third addressed the relationship of mental health to JIA disease characteristics or the impact of mental health on patient quality of life. Of the 28 current articles, 24 were conducted in countries outside of the US, including countries outside of North America and Europe (Iran, Turkey, and Egypt).

Population Characteristics

Twenty-five articles included youth with JIA while 3 articles studied adults. Four studies compared JIA to other chronic childhood diseases. As expected based on the general demographic patterns associated with JIA, study populations included a majority of female patients (range 52–81%) with the exception of one study.²³ Among the included pediatric studies, the mean age was 6.9 to 15.5 years and the age range in the adult studies was 18 to 73 years. Most studies did not report race and/or ethnicities. In the 6 studies that did report race, $\geq 70\%$ of the subjects were Caucasian or white.^{24–29} JIA disease duration ranged between 3 months to 15 years in children and 10 to 27 years in adults.^{24–28,30–44}

JIA subtype was described in 14 studies: 13 included polyarticular,^{24–26,30,31,35–40,45,46} 11 included oligoarticular,^{24,26,30,31,35–40,46} 10 included systemic,^{25,26,30,31,35–40} and fewer included ERA, psoriatic, undifferentiated, spondyloarthritis, and other forms of arthritis.^{25,26,30,31,34–37,40,46} JIA subtypes were described by all 3 of the adult studies.^{42–44}

Mental Health Measures

Recent studies mostly relied on patient-reported symptom scales, though parent-reported tools were often used in conjunction (Table 2). Depression was evaluated by a variety of questionnaires, most commonly the Children's Depression Inventory (CDI; in 9/22 depression studies). Other measures used were the Mood and Feelings Questionnaire (MFQ), Beck Depression Inventory-2 (BDI-II), and the Center for Epidemiological Studies Depression Scale for Children (CES-DC). Anxiety was evaluated most commonly by the Screen for Child Anxiety Related Disorders (SCARED; in 3/14 anxiety studies). Other anxiety measures used were the State Trait Anxiety Inventory (STAI), Anxiety Checklist for

Children and Adolescents, and the Self-Administered Psychiatric Scales for Children and Adolescents-Anxiety questionnaire (SAFA-A). To assess symptoms of either depression or anxiety, the Achenbach Child Behavioral Checklist (CBCL), Mini-International Neuropsychiatric Interview for Children (MINI-KID), Schedule for Affective Disorders and Schizophrenia for School-Age Children-Present and Lifetime Version (KSADS-PL), self-report, and psychiatric interviews were used.

Prevalence Of Depressive Symptoms In JIA

Most studies that used validated screens for depressive symptoms reported prevalences of clinically significant symptoms in the range of 7–36% in children with JIA.^{23–26,28,29,35,38,39,41,46,47} Reda et al used the MINI-KID psychiatric interview and reported that 93% of the patients had dysthymic disorder while 3% had major depressive disorder.³⁰ El Najjar et al found that all JIA patients scored above the cut-off for clinically significant symptoms on the CES-DC questionnaire.³⁷ There were 4 studies that reported no clinically significant depressive symptoms within their study population.^{27,31–33} These studies all used the CDI or BDI, screens that were used to detect prevalence rates of clinically significant symptoms in other studies from 9% to 36%,^{24,25,39} but were notable for patient populations from reportedly high socioeconomic levels^{27,33} and/or restricted to either non-acute patients^{32,33} or patients without comorbid psychiatric diagnoses.³¹

Three studies compared JIA patients to healthy controls, of which two found that JIA patients had worse depressive symptoms.^{23,35} Two additional studies reported that youth with JIA had worse anxiety/depressive symptoms than healthy controls; however, these symptoms were not analyzed separately.^{41,48} The exception, Tarakci et al, did not find a significant difference between JIA patients and healthy controls in regards to depression and/or anxiety symptoms; however, they excluded JIA patients with known comorbid psychiatric diagnoses.³¹ Notably, Kayan Ockagolu et al reported that 28% of their JIA cohort had a known diagnosis of major depressive disorder.²³

From the 3 studies that evaluated adults with JIA, two noted a particularly high prevalence of either depression or anxiety symptoms. Barth et al found significantly higher rates of anxiety/depression in JIA adults (18% in men, 37% in women) compared to healthy adults, and Rebane

Table 1 Demographic And Disease Characteristics Of Studies Of Depression And Anxiety In Youth With JIA (2009–2019)

Author (Year)	Country	Study Design	Sample Size		Female %	Mean Age Years (SD)*	Disease Duration (Months)	Percentage By JIA Subtype %							
			Subjects	Control				Oligo-Articular	Poly-Articular	RF – Poly	RF +Poly	ERA	PSA	Systemic	Undifferentiated
Banasiak et al (2010) ³⁰	Poland	Cross-Sectional	30	0	63	15.4	NS	NS	NS	NS	NS	NS	NS	NS	NS
Reda et al (2011) ³⁰	Egypt	Cross-Sectional	30	30	53	8-16	6-144	7	76	53	23	NS	NS	17	0
Tarakci et al (2011) ³¹	Turkey	Cross-Sectional	52	51	63	12.1 (2.9)	5-180	28	52	35	17	8	6	4	2
Vuorimaa et al (2011) ²⁴	Finland	Cross-Sectional	142 ^a	0	73	11.9 (3)	12-168	50	50	NS	NS	NS	NS	NS	NS
Anthony et al (2011) ²⁵	USA	Cross-Sectional	51 ^a	0	61	12.4 (2.8)	3-144	0	63	NS	NS	8	6	23	NS
Lal et al (2011) ²⁶	UK	Longitudinal	204 ^a	0	62	13.4	4-27	46	23	18	5	13	7	6	5
Barsdorf (2011) ²⁷	USA	Cross-Sectional	36	0	75	15.5	12-180	NS	NS	NS	NS	NS	NS	NS	NS
Russo et al (2012) ³²	Italy	Cross-Sectional	33	0	70	10.5 (4)	At least 6	NS	NS	NS	NS	NS	NS	NS	NS
Hrelc et al (2013) ²⁸	USA	Cross -Sectional	49 ^a	0	71	13	NA	NA	NA	NA	NA	NA	NA	NA	NA
Bromberg et al (2012) ⁴⁵	USA	Longitudinal	51	0	65	12.4	At least 1.5	0	100	NS	NS	0	0	0	0
Castaneda et al (2013) ³³	Finland	Cross -Sectional	34 ^b	23 ^f	61	15.5 (1.2)	<12: 9% 12-24: 4% >24 87%	NS	NS	NS	NS	NS	NS	NS	NS
Sevanovic et al (2013) ³⁴	Serbia	Cross -Sectional	67 ^a	0	71	13.6	6-168	NS	NS	NS	NS	NS	NS	NS	NS
Bomba et al (2013) ³⁵	Italy	Cross -Sectional	39	80	69	11.4 (2.1)	5-70.24	62	28	NS	NS	3	3	4	0
Mulligan et al (2013) ³⁶	UK	Cross -Sectional	171 ^c	0	72	9 (4.1)	8.4-87.6	32	43	37	5	6	4	15	0
El-Naijar et al (2014) ³⁷	Egypt	Cross-Sectional	54	0	67	11 (2.7)	12-120	39	28	NS	NS	11	0	22	0
Abdul-Sattar, Magd et al (2014) ³⁸	Egypt	Cross-Sectional	52	0	67	9.5	12-132	46	31	NS	NS	0	0	23	0

(Continued)

Table 1 (Continued).

Author (Year)	Country	Study Design	Sample Size		Female %	Mean Age Years (SD)*	Disease Duration (Months)	Percentage By JIA Subtype %							
			Subjects	Control				Oligo-Articular	Poly-Articular	RF – Poly	RF +Poly	ERA	PSA	Systemic	Undifferentiated
Abdul-Sattar, Elewa et al (2014) ³⁹	Egypt	Cross-Sectional	58	0	71	7-17	12-120	48	30	9	21	0	0	22	0
Haverman et al (2014) ⁴⁰	Netherlands	Cross-Sectional	155 ^c	0	67	11.5 (4.6)	2-168	36	42	39	3	11	0	4	5
Kuburovic et al (2014) ⁴⁷	Serbia	Cross-Sectional	25 ^d	50 JIA 89 Healthy	52	NS	NS	NS	NS	NS	NS	NS	NS	NS	NS
Memari et al (2016) ⁴¹	Iran	Cross-Sectional	51	75	55	11.2 (3.5)	27.6-70.6	NS	NS	NS	NS	NS	NS	NS	NS
Graziano et al (2016) ⁴⁸	Italy	Cross-Sectional	15 ^e	15 ^f	60	6.6 (3.2)	NS	NS	NS	NS	NS	NS	NS	NS	NS
Hanns et al (2018) ⁴⁶	UK	Longitudinal	102	0	57	13.2	NS	52	30	NS	NS	18	NS	NS	NS
Butler et al (2018) ³⁹	Canada	Longitudinal	28% asthma 18% diabetes 16% epilepsy 16% food allergy 22% JIA	0	48	11.3	NS	NS	NS	NS	NS	NS	NS	NS	NS
Kayan Okacoglu et al (2018) ²³	Turkey	Cross-Sectional	48 PID 32 JIA	30	28	NS	NS	NS	NS	NS	NS	NS	NS	NS	NS
Korte-Bouws et al (2019) ⁵³	Netherlands	Cross-Sectional	24	0	63	NS	NS	NS	NS	NS	NS	NS	NS	NS	NS

Notes: ^aProxy also included as part of the study. ^bStudy population was inflammatory bowel disease. ^cStudy population was juvenile idiopathic arthritis. ^dStudy population was patients with primary immunodeficiencies. ^eStudy population was patients with malignant solid tumors. ^fControl population was juvenile idiopathic arthritis. *When mean age was not provided, reported age range for the study is given.

Abbreviations: JIA, juvenile idiopathic arthritis; PID, primary immunodeficiency; NS, not stated; NA, not applicable.

Table 2 Depression And Anxiety Screening Measures Used For Youth With JIA (2009–2019)

Author	N (JIA)	Screening Measure For Anxiety	Anxiety Screen Interpretation	Prevalence Of Anxiety Symptoms/ Or Mean Score (SD) ^a	Screening Measure For Depression	Depression Screen Interpretation	Prevalence Of Depressive Symptoms/ Or Mean Score (SD) ^a
Banasiak et al (2010) ⁵⁰	30	STAI (s)	Low: 1-4 Moderate: 5-6 High: 7-10	Anxiety: 47% moderate, 10% high anxiety	NA	NA	NA
Tarakci et al (2011) ³¹	52 ^b	SCARED (s)	≥ 25	24.2 (9.7)	CDI (s)	≥19	8.1 (4.8)
Anthony et al (2011) ²⁵	51	RCMAS (s)	T-score > 60	8%	CDI (s)	T-score > 60	12%
Reda et al (2011) ³⁰	30	MINI-KID (s)	Diagnostic Criteria for Specific Phobia, Social Phobia, Obsessive Compulsive Disorder	40%	MINI-KID	Diagnostic Criteria for Dysthymia, Major Depressive Disorder	97%
Vuorimaa et al (2011) ²⁴	142	STAI (s)	T-scale	7%	CDI (s)	≥13	12%
Barsdorf (2011) ²⁷	36	NA	NA	NA	CDI (s)	≥19	7.6 (4.6)
Lal et al (2011) ²⁶	204	NA	NA	NA	MFQ (s)	≥27	7%
Russo et al (2012) ³²	33	Anxiety Checklist for Children and Adolescents (s)	z scores	z-score range: -1.7-0.04	CDI (s)	z scores	z-score range: -1.7-0.2
Hrelc et al (2013) ²⁸	49	RCMAS (s)	NS	18%	CES-DC (s)	≥15	33%
Bomba et al (2013) ³⁵	39	SAFA-A (s)	≥43	53%	CDI (s)	≥19	23%
Castaneda et al (2013) ³³	23 ^c	NA	NA	NA	BDI (s)	0-9=no symptoms 10-18=mild 19-29=moderate 30-63=severe	Depression: 9% mild 0% moderate-severe
Kuburovic et al (2014) ⁴⁷	50	SCARED (s, p)	≥25	14%	MFQ (s)	≥27	10%
El-Najjar et al (2014) ³⁷	54	NA	NA	NA	CES-DC (s)	≥15	19-60
Abdul-Sattar, Magd, et al (2014) ³⁸	52	NA	NA	NA	CDI (s)	>35 (Median)	36%
Abdul-Sattar, Elewa et al (2014) ³⁹	58	NA	NA	NA	CDI (s)	T-score ≥ 70	31%

(Continued)

Table 2 (Continued).

Author	N (JIA)	Screening Measure For Anxiety	Anxiety Screen Interpretation	Prevalence Of Anxiety Symptoms/ Or Mean Score (SD) ^a	Screening Measure For Depression	Depression Screen Interpretation	Prevalence Of Depressive Symptoms/ Or Mean Score (SD) ^a
Graziano et al (2016) ⁴⁸	15	CBCL (p)	NS	NS	CBCL (p)	NS	NS
Memari et al (2016) ⁴¹	51 ^c	CBCL (p)	≥98th percentile	13%	CBCL (p)	≥98th percentile	27%
Butler et al (2018) ²⁹	22	MINI-KID	Diagnostic Criteria for Phobia, Generalized Anxiety, Separation Anxiety	64% (baseline) 56% (6 months)	MINI-KID	Major Depressive Disorder	36% (baseline) 22% (6 months)
Hanns et al (2018) ⁴⁶	102	NA	NA	NA	MFQ	≥27	15%
Kayan Ocakoglu et al (2018) ²³	24	1. KSADS-PL 2. SCARED	SCARED ≥25	25%	1. KSADS-PL 2. CDI	CDI ≥19	31%

Notes: ^aInterquartiles ranges given when medians reported. ^bExcluded patients with known psychiatric disorders. ^cExcluded patients with active arthritis.

Abbreviations: STAI, State Trait Anxiety Inventory; SCARED, Screen for Child Anxiety Related Disorders; CDI, Children's Depression Inventory; RCMAS, Revised Children's Manifest Anxiety Scale; MINI-KID, Mini-International Neuropsychiatric Interview for Children and Adolescents; MFQ, Mood and Feeling Questionnaire; CES-DC, Center for Epidemiological Studies Depression Scale for Children; SAFA-A, Self-Administered Psychiatric Scales for Children and Adolescents-Anxiety Questionnaire; BDI, Beck Depression Inventory; CBCL, Child Behavior Checklist; KSADS-PL, Schedule for Affective Disorders and Schizophrenia for School-Age Children-Present and Lifetime Version; S, self-report; p, parent-proxy report; NS, not stated; N/A, not applicable.

et al reported a depression prevalence of 34%.^{43,44} In contrast, Raab et al found that only 4.9% of the adults with JIA self-reported depression, significantly fewer compared to the general population.⁴²

While most studies reviewed did not analyze a differential prevalence of depressive symptoms based on age or gender, those that did agree with trends in the general population with older adolescents and females showing worse scores.^{37,46,49}

Prevalence Of Anxiety Symptoms In JIA

The prevalence of clinically significant symptoms of anxiety in youth with JIA ranged from 7% to 64%.^{23,25,28,29,35,41,47,50} While most studies reported higher rates of depression than anxiety, a few notable studies found the opposite.^{29,35,47} Most studies comparing to healthy controls found higher rates among youth with JIA.^{35,41,47} Kayan Ocakoglu et al found that JIA patients scored worse on the SCARED and were more likely to have diagnoses of anxiety disorders compared to healthy controls, but the difference did not reach statistical significance, likely due to a small sample size.²³ Tarakci et al, another exception, had a cohort of JIA patients that excluded those with comorbid psychiatric disease.³¹ Bomba et al looked at specific subscales for different types of anxiety (generalized, social, separation, and school anxiety) and found that with each, youth with JIA scored worse than controls.³⁵

Butler et al was the only longitudinal study reviewed that examined anxiety and theirs was a pilot study of children with multiple chronic diseases, including a very small number of JIA patients. They found both overall and in JIA patients, high rates of anxiety (up to 36%) at baseline and 6-month measurements.²⁹

Few studies analyzed the prevalence of anxiety symptoms by age or gender. Russo et al found that children with JIA, ages 6–10 years, scored slightly higher on an anxiety questionnaire than adolescents with JIA, but this finding was not statistically significant. They also noted that psychological unease was reported in all JIA children, ages 6–10 years, compared to 42% of the adolescents.³² Raab et al was the only study to evaluate anxiety in adult patients with JIA and reported that 2% of the adults with JIA self-reported anxiety disorders in a comorbidity questionnaire.⁴²

Pathophysiology Of Mental Health In JIA

In 2011, Capuron et al found an association between chronic low-grade inflammation in elderly individuals with altered neurotransmitter metabolism and disturbed mental well-being.⁵¹ Then, in 2018, a review was published discussing the association between inflammation in rheumatoid arthritis and depression.⁵² Therefore, Korte-Bouws et al decided to evaluate this relationship in JIA by measuring serum samples of a panel of neurotransmitters in children with active JIA. They discovered that ratios between specific neurotransmitters indicated likely elevated activity of enzymes in dopamine and serotonin pathways. This may explain a pathway between active inflammation in JIA and symptoms of depression, anxiety, fatigue, and even cognitive impairment.⁵³

Reda et al found that the patterns of cerebral perfusion deficit in patients with JIA are similar to perfusion patterns observed in patients with diagnosed depression and/or anxiety. With single-photon emission computed tomography (SPECT) imaging, they found that children with JIA had a significant decrease in perfusion of the right frontal and right parietal lobes and a significant increase in perfusion of the left parietal and left occipital lobes when compared to healthy controls. These regions of the brain are known to be involved with mood regulation, attention span, and anxiety.³⁰

Risk Factors For Depression And Anxiety In Youth With JIA

Patient Characteristics

Hanns et al found that polyarticular patients scored significantly higher than oligoarticular and ERA patients on depressive symptoms. There was no difference in scores for depressive symptoms between the oligoarticular and ERA patients.⁴⁶ Conversely, El-Najjar et al did not find any statistically significant difference between the depressive scores of polyarticular, oligoarticular, systemic, and ERA subtypes. However, all patients scored above the clinically significant cut off for depressive symptoms by the CES-DC.³⁷

Disease Activity

Many of the articles found a correlation between disease activity markers and depression symptoms. Specifically, Hanns et al noted that both a higher active joint count and higher limited joint count correlated with higher depressive scores in the first year, but this relationship was no longer significant after follow up four years later.⁴⁶

Similarly, another study found a statistically significant correlation between depression scores and physician global scores, number of tender joints, number of swollen joints, number of limited joints, and the Juvenile Arthritis Disease Activity Score (JADAS-27).³⁷ In Barth et al's study of adult JIA patients, they found that anxiety/depression symptoms on a health-related quality of life questionnaire were significantly worse in patients with active versus inactive disease. Active disease in this study was also associated with worse mobility, self-care, usual activities, and pain/discomfort.⁴³ Conversely, Tarakci et al did not find any significant correlation between depression or anxiety and total active joint count.³¹

Two studies evaluated the relationship between disease duration and anxiety or depression with opposing findings regarding a correlation between duration and symptoms. El-Najjar et al found a strong correlation between worse depression symptoms by CES-DC,³⁷ while Tarakci et al found no correlation between disease duration and the CDI.³¹ In part, this may be due to the differing populations of JIA subtypes in these two studies (28% vs 52% polyarticular, respectively).

Disease Burden

Three studies found a statistically significant relationship between the Childhood Health Assessment Questionnaire (CHAQ), a patient-reported outcome of physical function, and depression scores.^{31,37,46} Additionally, Hanns et al found that a higher baseline depression score predicted worse disability in patients one and four years later.⁴⁶ On the other hand, Tarakci et al did not find a significant relationship between functional ability level, as measured by physical activities and energy expenditure, and anxiety or depression.³¹

Though most studies supported an association between active disease and worse mental health symptoms, managing disease with medications and blood test monitoring can be a source of psychological burden, as well. Mulligan et al assessed anxiety specific to injections and blood tests in children with JIA on methotrexate for ≥ 6 months. Over half reported feeling anxious about either. Anxiety about injections was associated with younger age, higher current disease activity, and subcutaneous administration, while anxiety about blood tests was significantly related to a younger age and a shorter duration of time on methotrexate. Importantly, blood test anxiety was an independent predictor for poorer scores in psychosocial quality of life.³⁶ Barth

et al's study of adults found an association between treatment for JIA as an adult and overall lower quality of life.⁴³ Notably, treatment for JIA in adulthood may be a marker for progressive, more severe disease.

Pain

Many studies evaluated the relationship between mental health symptoms and pain and most found an association between pain and either symptoms of anxiety or depression.^{24,37,46,50} The experience of pain may be influenced by particular stress-coping strategies reflective of a child or adolescent's level of anxiety.⁵⁰ Hanns et al noted that a higher baseline depression score predicted worse pain at one and four years later.⁴⁶ Bromberg et al found that daily reported mood was associated with daily measures of pain.⁴⁵ Tarakci et al was the outlier, finding no significant correlation between pain and depression or anxiety.³¹

Impact On Quality Of Life

Throughout the literature, depression symptoms consistently correlated with impaired overall quality of life.^{31,34,35,39} Abdul-Sattar et al found that over three-fourths of patients with significant depression symptoms had poor quality of life.³⁹ As expected, studies found that depression symptoms inversely correlated to emotional and social function on measures of quality of life.³⁵ Proxy parent reports of symptoms of their children's depression significantly predicted worse quality of life.³⁴

Anxiety also appears linked to quality of life in the reviewed literature. Children with JIA with a negative perception of their own life have higher anxiety scores.³⁴ However, Stevanovic et al found that after accounting for disease features, pain, and demographics, the association between anxiety and quality of life was lost.³⁵

A few studies looked specifically at quality of sleep and school functioning as markers of quality of life. Hrelc et al found that 80% of the children with JIA reported sleep disturbances of which night waking and parasomnias were significantly correlated with both anxiety and depression symptoms.²⁸ Bromberg et al reported that mood moderated the relationship between quality of sleep and pain.⁴⁵ Higher depression scores are strongly associated with poorer school functioning and school absenteeism in youth with JIA. Among JIA patients with significant depression, 88% had school absences and 69% had poor school functioning.³⁸

A study of adults with JIA evaluated patients on whether they were socially restricted versus non-restricted. The restricted patients were unemployed or on disability while the non-restricted were studying, working, on maternity leave, or in the military. They found that 83% of the patients were socially non-restricted compared to 16% restricted. Of the 16% that were restricted, the majority (85%) were unemployed. The socially restricted JIA patients spent less time in leisure non-physical activities, felt more disturbed during leisure time, had lower self-esteem scores, and higher disability scores. They also had statistically significant lower health-related quality of life scores in physical and social functioning as well as emotional well-being compared to the non-restricted. However, they did not find any difference between the depressive scores between the two groups. Of note, 59% of the restricted patients had polyarticular JIA compared to 47% of the non-restricted patients; therefore, it could be hypothesized that some differences between the two groups were secondary to worse disease activity in the restricted patients. In contrast, there was no statistically significant difference in disease duration between the two groups.⁴⁴

Impact On Parents And Family

A few studies addressed the mental health of caregivers of children with JIA. Vuorimaa et al found that 12% of the caregivers had clinically significant depressive symptoms while 7% had anxiety. Children's pain was significantly associated with their parents' depressive symptoms and children's depressive and anxiety symptoms moderately correlated with those of their parents.²⁴ Russo et al found that 17% of the mothers of children with JIA and a third of their siblings sought psychological support, while a quarter of mothers endorsed being on an anti-depressant.³² Through a health-related quality of life questionnaire, parents of children with active disease scored significantly worse in depressive symptoms than the parents of children with inactive disease.⁴⁰

Haverman et al report that parents of children with JIA perceived their child as more vulnerable compared to parents of healthy children and parents of children with other chronic diseases.⁴⁰ Anthony et al found that higher parent perception of child vulnerability correlated with child depressive symptoms, and children perceived as more vulnerable were also more anxious. In addition, parent daily stress, specifically more intense daily hassles and fewer daily uplifts, predicted higher depressive

symptoms in their children with JIA. They concluded that children of parents with more daily stress and less stress relief were more susceptible to depressive symptoms, and that parent experiences of daily uplifts or pleasurable events may be protective to the development of depressive symptoms in their children with JIA.²⁵ In contrast to the psychopathology described among JIA caregivers in these studies, Graziano et al found that with regards to emotional complaints and psychosocial distress, JIA mothers appeared to have more in common with mothers of healthy controls than with mothers of cancer patients.⁴⁸

Comparison Of Mental Health In JIA And Other Chronic Diseases

Four articles analyzed mental health in JIA compared to other chronic childhood diseases including inflammatory bowel disease (IBD), primary immunodeficiencies (PIDs), and cancer.^{23,33,47,48} Overall, JIA patients had similar depression/anxiety symptom levels compared to the other chronic diseases, with the exception of IBD patients who were found to be more depressed.³³ Kayan Ocakoglu et al had children evaluated through a psychiatric interview and found that 63% of the JIA children, 70% of the PIDs, and 23% of the healthy children had some form of a psychiatric disorder (this included disorders other than mood and anxiety such as psychotic, elimination, disruptive, substance abuse, eating disorders, and tic disorders).²³

Discussion

Studies of depression and anxiety in youth with JIA conducted in the last 10 years reported that between 7% and 64% of the JIA patients have symptoms of anxiety or depression. These prevalence rates are at least as high as the general pediatric population. Most studies found that children with JIA had significantly higher levels of anxiety and/or depressive symptoms than healthy children.^{23,33,47,48}

The wide range in symptom prevalence among studies may be attributed to differences in populations, differences in definition or measurement of symptomatology, and differences in methods used to obtain information about patients' anxiety and depression symptoms. In contrast to the earlier literature that depended heavily on parent-reported scales for depression and anxiety,⁸ the reviewed studies mostly relied on patient self-reported symptom scales. Lal et al assessed the level of agreement between adolescents with JIA and their caregivers with regards to mental health and well-being. Adolescents who reported more depressive

symptoms were more likely to disagree with their parents about depressive symptoms.²⁶ Other reviewed studies also found discrepancies between patient self-reported and parent-reported symptoms of depression/anxiety through screening tools such as the CBCL.^{54,55} In pre-adolescent children, there is likely a role for both patient self-report and parent-proxy reports to build a complete picture of a child's mental health.

Likewise, it is still unclear how disease duration may impact the risk of anxiety and depression. Butler et al hypothesized that symptoms would improve after the initial adjustment to illness and found a slight decrease in symptoms at 6 months of follow-up.²⁹ This is supported by an earlier study that found that adolescents with lower symptoms of anxiety and depression tended to have longer duration of disease.⁵⁶ El-Najjar et al found the opposite correlation between disease duration and depressive symptoms.³⁷ It is possible that longer disease durations are associated with increased damage and changes to function that may increase the risk of developing symptoms of depression. This relationship has been seen with depression in diabetes.^{57,58}

Unfortunately, most studies did not perform a separate analysis of mental health symptoms by JIA subtype due to small sample sizes. In one study, polyarticular JIA patients were found to have significantly higher depressive scores than both oligoarticular and ERA patients.⁴⁶ Similarly, an older study reported that adults with RF-negative polyarticular JIA have particularly high rates of depressive symptoms in comparison to adults with other subtypes.⁵⁹ It is possible that polyarticular JIA patients are at an increased risk for depression compared to other subtypes because they commonly suffer from higher active joint counts and higher disease activity which may impair overall function and lead to lower quality of life. On the other hand, El-Najjar et al did not find any significant differences in depressive scores between oligoarticular, polyarticular, systemic, and ERA patients; though in this study, all of the patients scored above the cutoff for significant depressive symptoms.³⁷ No included studies examined anxiety symptoms and JIA subtype.

Only a quarter of the studies investigated relationships between other JIA disease characteristics and mental health symptoms. Disease characteristics were assessed in various ways, making comparisons between studies difficult. Most, but not all, of these studies found a positive correlation between higher disease activity and depression/

anxiety symptoms.^{37,43,46} Of note, some of the earlier literature conflicts with these findings.⁵⁵

The impact of JIA treatment on symptoms of depression and anxiety was rarely approached in the reviewed studies; however, our search approach was not wide enough to include studies of treatments that reported symptoms of depression and anxiety outside of the main findings highlighted in their abstracts. This is an area deserving of study, especially considering that systemic and brain inflammation have been implicated in psychiatric conditions⁶⁰ such as depression, and anti-tumor necrosis factor (anti-TNF) therapy was recently studied as a potential treatment for refractory depression with mixed results.⁶¹ In a randomized controlled trial of infliximab for refractory depression, infliximab did not improve depression scores compared to placebo in the overall study population; however, in an exploratory analysis of patients with elevated baseline inflammatory markers, there was a decrease in depression symptoms with infliximab compared to placebo.⁶¹ In contrast, depression and anxiety have been reported in small numbers in clinical trials of TNF-inhibitors and other treatments used for arthritis in patients with JIA.^{62–64} Hider et al specifically evaluated depression in adults with rheumatoid arthritis on anti-TNF therapy. Although 48% of the patients had depression at baseline, after a year of anti-TNF therapy the proportion of patients with persistent depression fell to 15% and only 6% of the patients on anti-TNF therapy developed new-onset depression. Therefore, they concluded that anti-TNF therapy is not likely to adversely affect mood.⁶⁵

Because the vast majority of studies reviewed were cross-sectional, the direction of the relationship between disease activity and mental health symptoms is still poorly described. It is plausible that depressive symptoms may lead to worse disease activity, especially if patients become non-adherent with medical care as a result of their mental health symptoms. In other disease processes, including childhood-onset lupus, medication adherence has been linked to depressive symptoms.⁶⁶ But it is equally plausible that worse disease activity is leading to the development of depressive symptoms. Although most studies were cross-sectional, three studies in adult JIA patients were reviewed to address long-term consequences of JIA. Two were designed to specifically answer the question of how JIA patients fared emotionally as adults and found that 30–40% of the adults with JIA had symptoms of anxiety or depression.^{43,44} The other study by Raab et al found that adults with JIA had a lower rate of

lifetime prevalence of depression compared to a community sample of age-/gender-matched controls.⁴² However, the prior literature also indicates that adults with JIA may have worse anxiety and depression than their healthy peers.^{59,67,68} Therefore, significant psychopathology can impact patients into adulthood and may influence their emotional, social, and overall quality of life. This is similar to what has been reported in children with other chronic diseases such as chronic kidney disease and epilepsy along with many others.^{4–6,69,70} However, in contrast to many other chronic conditions, JIA is characterized by unpredictable flares amidst periods of quiescence and often preceded by significant delays in diagnosis, all of which may further contribute to the development of anxiety and depression.

Although there were two articles that did not find any significant relationship between pain and depression or anxiety symptoms, most of the articles did find that pain was significantly correlated with symptoms.^{24,31,37,45,46,50} This is consistent with earlier literature.¹⁸ Older literature in JIA reported mixed results with many studies finding a correlation between pain and depression/anxiety symptoms, while some reported weak or no significant association.^{12,71} Hanns et al even found that baseline depressive symptoms predicted worse pain and disability up to 4 years later.⁴⁶ Overall, most of the literature points to an association between pain and psychological symptoms, though much of it does not account for all the potential confounders such as disease severity, activity, disability, and patient demographics.

Physical disability appears to be correlated to mental health symptoms, as well. In two studies, the CHAQ was found to correlate with depressive symptom scores.^{31,37} An earlier cluster analysis of JIA patients by age, anxiety, and depression symptoms found that adolescents with JIA who were more symptomatic of both depression and anxiety also had higher rates of pain, disability, and active joint counts.⁵⁶ Other older literature has reported moderate associations between disability and depression/anxiety symptoms, both in children and adults with JIA.^{59,72} Difficulties in functioning and overall ability to complete activities of daily living may cause children with JIA to feel different than their healthy peers, which may ultimately affect their mood. Additionally, any physical changes or difficulties that are apparent to their peers could lead to bullying which is tied to the development of mental health symptoms.^{73,74}

The overall quality of life was worse or impaired in JIA patients with higher depressive scores.^{31,34,35,39} Sleep quality of life was also impacted with most JIA patients reporting some sleep disturbance which correlated with anxiety and depressive symptoms, and mood moderated the relationship between poor sleep and pain.^{28,45} This may create a vicious cycle. Pain can interfere with sleep, sleep disturbances can worsen pain, worse pain can lead to feelings of depression, depression can impact sleep and exacerbate pain, and poor sleep can worsen depression further. School quality of life was also impacted by depression, an association that can be explained by how people with depression struggle with participating and performing daily activities.³⁸ Prior studies have documented the relationship between depressive symptoms and school impairments in other chronic patient populations such as chronic pain.⁷⁵

Building off the previous literature that indicated that family and parental characteristics and adjustment are associated with child adaptation to illness,⁷⁶ several of the recent studies we reviewed addressed the interconnectivity between family and patient mental health in families affected by JIA. Many of these studies found that parents of children with JIA had clinically significant depressive or anxious symptoms, and that symptoms of depression and anxiety may be worse in family members of children with active disease.^{24,32,40} In addition, more parent daily stress with little stress relief also predicted higher depressive symptoms in their children with JIA.²⁵ Vuroimaa et al found that child and parent depressive and anxious symptoms correlated with each other and the child's pain was associated with the parent's depressive symptoms.²⁴ This is likely because of shared environmental and genetic predispositions as well as the possibility that one family member's mental health symptoms are influencing the other's.⁷⁷⁻⁷⁹ With evidence of significant family psychopathology and psychological distress, it is likely beneficial for the child's mental health to address the parent's mental health, too.

Study Limitations

Limitations to this review include the challenges of developing a screening process for a broad topic that is discussed with multiple different terms. We may have missed important studies that had keywords related to emotional or mental health and addressed internalized symptoms such as depression and anxiety, without specifically listing them as keywords. Furthermore, we missed articles by limiting our scope to English-language publications.

Knowledge gained by our review was also limited by the limitations of the individual studies. Most of the studies that were included were cross-sectional studies, so symptoms of depression/anxiety were evaluated only at one particular time point. Therefore, trends in symptoms over time could not be evaluated to see if they persisted, worsened, or improved. Consequently, patients who were experiencing symptoms of anxiety/depression outside of the time frame they participated in the study would have been missed. Studies, especially cross-sectional, may introduce non-response bias in which the responses of the people who chose to participate may differ from those who chose to not participate. This may be particularly true for studies involving mental health, where stigma still exists around the topic in some cultures. Lastly, cross-sectional studies allow you to make conclusions about possible associations, but not about causation.

Many of the studies had small sample sizes which can impact the significance and generalizability of the results. Although, most studies were still able to report on statistically significant differences comparing JIA to controls. However, small sample sizes did not provide enough power for most of the studies to complete any analysis about how JIA disease characteristics, medications, or subtypes may be correlated with mental health symptoms. Disease activity was assessed and analyzed with regards to mental health symptoms by varying measures of disease activity making it hard to make comparisons and overall conclusions. Furthermore, some studies only included patients who were in remission or had low/mild disease activity and others excluded patients with known psychiatric diseases. This can lead to potential selection bias, challenging us to compare these and synthesize the disparate observations into a singular understanding of how anxiety and depression impact youth with JIA.

Additionally, few studies reported patient sociodemographics, and US and European countries that reported race/ethnicity of patients still included few racial and ethnic minorities again limiting generalizability.²⁴⁻²⁹

Conclusion

Recent literature indicates that youth with JIA experience clinically significant symptoms of anxiety and depression similar to youth with other chronic diseases and likely at higher rates than healthy children. These studies have added to the body of literature that address JIA specifically, as opposed to general juvenile rheumatic diseases and help to reinforce the need and importance of mental

health screening and surveillance specifically in children with JIA. The reviewed studies show an increasing inclusion of work worldwide, even with our limitation in scope to English-language journals. Across diverse geo-cultural backgrounds, it is clear that mental health symptoms are an important part of the experience of JIA and something to which pediatric rheumatology providers must attend.

However, many of the challenges of the early literature still beleaguer the literature of the last ten years. It is likely that large populations of youth with JIA are still not well represented in the literature and this inhibits our understanding of how mental health symptoms impact all patients and which patients are at greatest risk. Disease activity and pain were found in a few articles to correlate with worse depressive and anxiety symptoms; however, the majority of the articles did not perform an analysis of the associations between JIA disease characteristics or subtype with mental health symptoms. More studies are needed to better understand the complex nature of how parents' and other family members' emotional health impacts the affected child's and vice versa. Furthermore, the studies are almost uniformly cross-sectional studies that still leave unanswered questions about how patients progress over time and how the unpredictable and undulating course of JIA disease activity impacts mood and mental health.

As the importance of mental health symptoms to patients and families is continually raised by the arthritis patient community, measurements of these symptoms should be incorporated in the large international longitudinal pediatric rheumatology registries. These studies, which may benefit from a coupling of biospecimen banks, will enhance our understanding of how JIA disease and mental health symptoms interplay over time, how they each impact one another, and may help us understand whether mental health symptoms influence the pathophysiology of inflammatory arthritis or vice versa. This information could provide evidence to target mental health screening efforts, identify novel and effective treatment pathways, and develop individualized therapy plans for youth with JIA.

Though gaps persist in our knowledge of how depression and anxiety influence specific disease factors in JIA, the recent literature clearly points to the role that depression and anxiety symptoms have on quality of life, both for the child with JIA and their family. Pediatric rheumatologists need to consider addressing these symptoms if they wish to promote improved quality of life for their patients.

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