

Visual Function And Quality Of Life In A Cohort Of Swedish Children With Juvenile Idiopathic Arthritis

This article was published in the following Dove Press journal:
Clinical Ophthalmology

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Purpose: To evaluate quality of life (QoL) in children with juvenile idiopathic arthritis (JIA).

Methods: Forty children with a mean age of 7.9 years were included. The children underwent an ophthalmological examination and completed questionnaires on physical function (CHAQ) and vision-related (VR) QoL (EYE-Q).

Results: No differences regarding visual acuity (VA), refraction, intraocular pressure or physical or VRQoL were found between those with JIA without (n=33) and those with JIA-associated uveitis (n=7). When comparing physical function measured by CHAQ disability index and JIA subtype, a difference was found; children with polyarthritis scored the worst (p=0.0098). Children with subnormal VA scored worse on EYE-Q compared with those with normal VA (p=0.013). We found correlations between duration of JIA and CHAQ disability index (r=-0.42, p=0.0007) and CHAQ well-being (r=-0.34, p=0.022).

Conclusion: This study indicates the importance of measuring not only physical function but also VRQoL in children with JIA and JIA-associated uveitis.

Keywords: child arthritis, juvenile idiopathic arthritis, PROM, uveitis, quality of life

Introduction

Juvenile idiopathic arthritis (JIA) consists of several subtypes of chronic arthritis with its onset of diagnosis before 16 years of age, and is defined as swelling or limitation of motion of the joints accompanied by heat, pain or tenderness of at least six weeks duration with no other identifiable causes of arthritis.¹ Juvenile idiopathic arthritis is the most common chronic rheumatic disease in children in the Western world, and it may be a significant cause of chronic pain, disability and reduced quality of life (QoL).^{2,3}

The incidence and prevalence in European and North American populations range from 2 to 20 and from 16 to 150 per 100,000, respectively.⁴ The International League of Associations for Rheumatology (ILAR) has provided classification of JIA consisting of seven subtypes, including systemic arthritis, oligoarthritis (OA) extended and persistent, polyarthritis (PA), rheumatoid factor (RF) positive and RF negative, enthesitis-related arthritis, psoriatic arthritis and undifferentiated arthritis or "other" JIA.^{5,6}

Vision-threatening uveitis is the most common extra-articular manifestation of JIA, with a cumulative incidence of approximately 9–21% in these patients.^{7–10}

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JIA-associated uveitis (JIA-U) may develop before, at the same time as, or after the arthritis onset and is often asymptomatic at the time of onset. JIA-U can lead to severe visual impairment and is a relevant cause of ocular morbidity in children.^{11,12} The JIA-U is typically anterior and chronic, and frequently affects both eyes. Serious ocular complications can occur (e.g. cataract, glaucoma, band keratopathy and posterior synechiae), probably causing visual impairment in affected children.¹³ Risk factors for developing uveitis include antinuclear antibody (ANA) positivity, early onset of JIA, JIA duration, certain human leukocyte antigen (HLA) markers and active disease state and elevated erythrocyte sedimentation rate (ESR).^{8–10,13–15}

Our primary objective was to evaluate and compare visual and physical function and vision-related (VR)QoL in children with JIA and JIA-U who were screened and/or treated for uveitis at the Department of Pediatric Ophthalmology at the Queen Silvia Children's Hospital in Gothenburg, Sweden.

Materials And Methods

A prospective cohort study was performed on 40 Caucasian children (31 girls, 9 boys) with a mean age of 7.9 years (range 3.1–11.8 years), diagnosed with JIA according to ILAR criteria. Patients having JIA with ongoing uveitis or children followed up for previous uveitis as well as JIA children screening for uveitis during 2013–2014 were included. All children underwent a detailed ophthalmological examination (by one of the authors, MP) including best corrected visual acuity (BCVA), refraction, intraocular pressure (IOP), slit-lamp inspection, ophthalmoscopy of the fundus and optic coherence tomography (OCT). We also registered any current and/or previous ocular complications secondary to uveitis, previous eye surgery and age at onset of uveitis. Ocular inflammation was defined according to the Standardization of Uveitis localisation of the inflammation. The classification includes 1) location of uveitis, 2) onset, duration and course of uveitis and 3) severity and activity of uveitis.¹⁶

Ophthalmological Examination

VA was tested with a linear KM-Boks chart.¹⁷ If a child could not manage to read the KM-Boks chart, an HOTV chart was used. Distance VA was tested monocularly and binocularly at a distance of 3 m. Values were noted in decimal and transformed to logMAR units. Visual acuity of children with JIA was compared with an age- and sex-matched control group consisting of 55 healthy

children (13 boys, 42 girls) with a mean age of 7.9 years (range 4.1–12.1 years). Refraction was performed with an autorefractor (Topcon A6300; Topcon Corporation, Tokyo, Japan), undilated as well as dilated after a single instillation of a mixture of cyclopentolate (0.85%) and phenylephrine (1.5%). Refractive errors were defined as the spherical equivalent (SE) of myopia ≥ 0.5 dioptres (D), hyperopia ≥ 2.0 D or anisometropia ≥ 1.0 D. Astigmatism was considered significant at a level of ≥ 1.0 D.

IOP was measured by a handheld tonometer (TA0li, I care, Finland Oy, Espoo, Finland). Readings with high deviations were discarded and repeated. Findings such as band keratopathy, flare and cells, posterior synechiae, cataract, vitritis, and macular and/or papilledema were especially noted. The morphology of the fundus was further examined using an OCT (3D OCT-2000, Topcon, Corporation, Japan, 2012), e.g. for detection of macular edema.

Paediatric Data

The following variables were collected and documented from medical records: gender, age, ethnicity, arthritis subtype, age at onset and duration of arthritis and presence of ANA. All topical and systemic anti-inflammatory and immunosuppressive medications were recorded.

Questionnaires On Quality Of Life Visual Function Assessment And Vision-Related Quality Of Life

Visual function and VRQoL were measured by the Effects of Youngsters' Eyesight on Quality of Life instrument (EYE-Q). The EYE-Q instrument consists of 13 self-reported items for individuals ≥ 8 years of age that evaluate competence in performing daily tasks that rely on vision in the school and at home.^{2,18} It requires 10 mins to complete. Those children younger than 8 years old completed the questionnaire together with the parents. This questionnaire uses a 5-point Likert scale in 12 of the items (items which assess the difficulties to see the colors on a stoplight, the words in the school book, the food on the plate, etc.). The following response format is used to assess difficulty in performing tasks, with response options as follows: 1 (not hard/never); 2 (a little hard/rarely); 3 (hard/sometimes); 4 (very hard/often) and 5 (cannot do). The items were rescaled, with the scores ranging from 0 to 4² higher scores indicated worse QoL.^{2,18–20} The 13th item, which is about whether there are other things that are difficult to do related to the child's vision, uses yes or no alternatives.

Physical Function Assessment

Physical function was measured by the Child Health Assessment Questionnaire (CHAQ), which is a disease-specific instrument for JIA and a valid measurement of functional disability.^{18,21} It is adapted from the Stanford Health Assessment Questionnaire and has well-documented reliability and validity.^{2,22} We evaluated two parallel questionnaires, one for parents (n=25) and another for children older than 8 years of age (n=15).

It consists of a “disability index” of 30 questions within eight functional components: 1) dressing and grooming, 2) arising, 3) eating, 4) walking, 5) hygiene, 6) reach, 7) grip and (8) activities. There are three components within each area: 1) difficulty in performing daily functions, 2) use of special aids or devices and 3) activities that require assistance from another person. Each question has a 4-point response scale (0=no difficulty; 1=some difficulty; 2=much difficulty; 3=unable to do). Scores range from 0 (no disability) to 3 (maximum disability), where higher scores indicate worse physical function.^{2,18,22}

CHAQ also includes two other components: 1) “Assessment of discomfort”, using a 10-cm visual analogue scale (VAS) for the evaluation of pain and its severity in the past week, whereby a score of 0 indicates “no pain” and 10 indicates “extreme pain”²³ 2) “Overall well-being” (global assessment), also using a 10-cm VAS, ranging from “very well” (score of 0) to “very bad” (score of 10).²⁴

In the Swedish version of CHAQ, two other parts are also included, one consisting of information about whether the children have been absent from school due to arthritis during the last two months, and another component providing information about the children’s participation in school gymnastics.

Correlations

Age at onset and duration of both JIA and uveitis as well as VA in the better eye were analysed to establish any correlation with the following variables: physical function (CHAQ disability index; CHAQ discomfort; CHAQ well-being) and visual function and VRQoL measured by EYE-Q.

Statistical Analysis

Percentages, means and standard deviations (SDs), median and ranges were calculated for descriptive purposes. For comparison between groups, Fisher’s Exact test was used for dichotomous variables, and Mann–Whitney *U*-test and

Kruskal-Wallis test were used for continuous variables. Correlation analyses were performed using Pitman’s partial-rank correlation analysis. Test results were considered significant if $p < 0.05$.

The healthy controls were selected individually by minimizing the maximal *t*-values between the group of JIA children at assessment and a reference group of healthy Swedish school-aged children²⁵ over the variables age and sex. This method matches two populations by iteratively selecting the individuals with the minimum *t*-test score.²⁶

Ethical Approval

The study was approved by the Research Ethics Board at the University of Gothenburg, Sweden. All candidates and their guardians received detailed information about the nature of the investigation, after which written informed consent was collected. All procedures were performed in accordance with the tenets of the Declaration of Helsinki.

Results

The clinical characteristics of the 40 children with JIA, divided into those with JIA-U (n=7) or JIA alone (n=33) are summarized in Table 1. The children with JIA were categorized by subtype, as defined by ILAR. Characteristics of the seven children (17.5%), two boys and five girls, who had uveitis at assessment or have had uveitis in the past, are shown in Table 2. The mean age of arthritis onset in children with JIA-U was 3.3 years (range 1.5–5.3 years), and in children without uveitis it was 3.8 years (range 1.4–11.3 years). All children with JIA-U developed uveitis after being diagnosed with JIA. Several ocular complications were found such as bilateral band keratopathy, synechiae, cataract and ocular hypertension. Furthermore, ongoing inflammation (keratic precipitate, flare and cells) was noted in two cases at assessment.

Results of BCVA and refraction are shown in Table 3. There was no significant difference regarding IOP between the children with JIA alone (RE mean (SD) 14.9 ± 3.7 mm Hg; LE mean (SD) 15 ± 3.2 mm Hg) and those with JIA-U (RE mean (SD) 15.4 ± 4.3 mm Hg; LE mean (SD) 14.6 ± 2.6 mm Hg; $p=0.79$, $p=0.77$, respectively). Performed OCT on the children showed normal imaging, i.e. no sign of macular edema could be detected.

The results from the questionnaires (CHAQ and EYE-Q) are summarized in Table 4. When comparing the three major components of CHAQ between the self-reports (children ≥ 8 years, n=15) and the parent reports (children < 8 years, n=25), we could not find

Table 1 Demographic Data On 40 Children With Juvenile Idiopathic Arthritis (JIA) Divided Into Those With JIA-Associated Uveitis (n = 7) Or Not (n = 33)

Demographics	JIA children n = 40*	Uveitis n = 7*	No uveitis n = 33*
Age at assessment (yrs) Mean (range)	7.9 (3.1 – 11.8)	7.9 (4.2 – 11.0)	7.8 (3.1 – 11.8)
Gender, n (%) • Male • Female	9 (22.5%) 31 (77.5%)	2 (29%) 5 (71%)	7 (21%) 27 (79%)
Race, n (%) • Caucasian	40 (100%)	7 (100%)	33 (100%)
JIA disease characteristics • Age at JIA diagnosis (yrs) Mean (range) • Duration of JIA (yrs) Mean (Range)	3.7 (0.4 – 11.3) 4.2 (0.7 – 9)	3.3 (1.5 – 5.3) 4.9 (1.6 – 8.3)	3.8 (1.4 – 11.3) 4.1 (0.7 – 9)
JIA subtype, n (%) • Systemic • Oligoarthritis persistent • Oligoarthritis extended • Psoriatic • Polyarthritis	1 (2.5%) 25 (62.5%) 5 (12.5%) 1 (2.5%) 8 (20%)	0 4 (57%) 1 (14%) 0 2 (29%)	1 (3%) 21 (64%) 4 (12%) 1 (3%) 6 (18%)
Labs, n (%) • ANA (+) • RF (–) • HLA B27 (+)	26 (65%) 26 (100%) 3 (100%) n = 26 n = 3	5 (71%) 3 (100%) 0 n = 3	21 (64%) 23 (100%) 3 (100%) n = 23 n = 3

Notes: *Where numbers differ from the total number of children in the group, they are given separately for each category.

Abbreviations: ANA, antinuclear antibodies; HLA B27, human leukocyte antigen subtype B27; JIA, juvenile idiopathic arthritis; n, number of children; RF, rheumatic factor.

any differences regarding CHAQ disability index and CHAQ discomfort. However, the parents reported higher CHAQ well-being scores, lower QoL than the children ($p=0.012$). When comparing physical function measured by CHAQ disability index and JIA subtype, a difference was found ($p=0.0098$), with children with PA ($n=8$) having the highest score (0.46). We could not find any differences between JIA subtype and uveitis/no uveitis status.

All 40 children with JIA answered the 13th item of the EYE-Q. One of seven JIA-U children reported to have

difficulty with photosensitivity. No children without uveitis reported any difficulty on the 13th item of EYE-Q.

Parents reporting any need of aids or devices for the children, and/or assistance of another person because of the disease, are shown in Figure 1A and B, divided into those with and without JIA-U. The self-reporting children did not report any need of aids and/or devices. Regarding children being home from school/pre-school due to their arthritis, the parents reported a total number of 10 children having a mean of 2.1 (range 1–4) days of absence in the past 2 months; three of these children had JIA-U. Thirty-four of the 40 children with JIA went to school, and all of them participated in school gymnastics.

There were significant correlations between the duration of JIA and CHAQ disability index ($r=-0.42$; $p=0.0007$) and CHAQ well-being ($r=-0.34$; $p=0.022$). A correlation between BCVA LogMAR best eye and EYE-Q ($r=-0.36$; $p=0.042$) was also found. When we compared JIA children with BCVA in the better eye ≤ 0.7 decimal (0.2 LogMAR; $n=5$) versus children with normal BCVA ($n=35$), we noticed that those with BCVA ≤ 0.7 decimal had significantly worse EYE-Q scores ($p=0.013$). However, this was not true for the three components of CHAQ regarding BCVA ≤ 0.7 decimal versus normal BCVA (Table 5). Analysis at the individual level did not reveal any association between those with both JIA and uveitis and worse general QoL or worse EYE-O.

A total of 25 patients out of the 40 used some kind of disease-modifying antirheumatic drugs (DMARDs) and/or non-steroidal anti-inflammatory drugs, five with JIA-U and 20 without. Eleven of the 20 children with JIA alone and five children with JIA-U used DMARDs, either alone or in combination with both methotrexate and tumour necrosis factor (TNF)-alfa inhibitor. The TNFalpha drugs used in these cases were adalimumab[®] and etanercept[®]. One child who had had JIA-U was not using any current medication, while another child with JIA-U had only used NSAID before the study enrolment (Figure 2A and B). One of the children with JIA-U was currently being treated with a combination of both Methotrexate[®] and Adalimumab[®] as well as local steroids (Dexamethasone[®]) once a day, and antihypertensive eye drops (Betoptic S[®]) twice a day. This patient had several ocular manifestations, such as cataract, ocular hypertension, posterior synechiae, vitritis and band keratopathy. None of our patients was on oral steroids.

Table 2 Characteristics Of Children With Juvenile Idiopathic Arthritis (JIA)-Associated Uveitis At Study Enrolment

	Patient No 1	Patient No 2	Patient No 3	Patient No 4	Patient No 5	Patient No 6	Patient No 7
Age at assessment (yrs)	8	4.2	8.1	9.1	6.2	8.8	11
Sex m/f	f	f	f	f	m	m	f
JIA debut (yrs)	1.5	1.5	4.5	2.8	4.5	3	5.3
ANA (\pm)	+	-	+	+	+	-	+
JIA subtype	Oligoarthritis persistent	Polyarthritis	Polyarthritis	Oligoarthritis extended	Oligoarthritis persistent	Oligoarthritis persistent	Oligoarthritis persistent
Age at uveitis diagnosis (yrs)	3.7	4.2	4.9	5	4.7	6.8	5.4
Interval between uveitis and arthritis (yrs)	1.7	2.7	0.4	2.2	0.2	3.8	0.1
Bilateral involvement (Y/N)	Y	N	Y	Y	Y	N	N
Location uveitis	Anterior	Anterior	Anterior	Anterior	Anterior	Anterior	Anterior
BCVA best eye (LogMAR)	-0.1	0.1	0.4	0.0	0.0	0.0	-0.1
VA RE	1.0	0.8	0.4	1.0	1.0	1.0	1.25
VA LE	1.3	0.65	0.4	1.0	1.0	1.0	1.25
Ocular complication(s)	Synechiae, ocular hypertension		Bilateral band keratopathy and synechiae, cataract*, ocular hypertension				Unilateral subcapsular cataract
Current treatment	Methotrexate [®] , Adalimumab,	Erancept [®] , ocular local	Methotrexate [®] , Adalimumab [®] , Betoptic S [®] , Dexamethasone [®] , ocular local		Adalimumab	Methotrexate [®]	

Note: *This child had underwent cataract surgery.

Abbreviations: ANA, antinuclear antibodies; BCVA, best corrected visual acuity; JIA, juvenile idiopathic arthritis; LogMAR, logarithm of the minimum angle of resolution; N, no; VA, visual acuity; Y, yes.

Table 3 Visual Acuity And Refraction In 40 Children With Juvenile Idiopathic Arthritis (JIA) Compared With An Age- And Sex-Matched Control Group (n = 55)

Visual Acuity (logMAR)	JIA Children, n = 40*	Uveitis, n = 7	No Uveitis, n = 33*	Uveitis Vs No Uveitis p-value	Controls, n = 55	JIA vs Control p-Value	Uveitis vs Control p-Value	No Uveitis vs Control p-Value
BCVA best eye Mean \pm SD	0.015 \pm 0.108	0.029 \pm 0.170	0.012 \pm 0.093	0.76	0.027 \pm 0.091	0.19	0.30	0.28
VA RE Mean \pm SD	0.023 \pm 0.111 n = 39	0.057 \pm 0.162	0.016 \pm 0.099 n = 32	0.61	0.037 \pm 0.096	0.16	0.79	0.14
VA LE Mean (SD)	0.036 \pm 0.116 n = 39	0.086 \pm 0.168	0.025 \pm 0.102 n = 32	0.41	0.038 \pm 0.088	0.43	0.74	0.30
Refraction** D SE								
RE Mean \pm SD	+0.063 \pm 1.479 n = 38	-0.411 \pm 1.980	+0.169 \pm 1.360 n = 31	0.56				
LE Mean \pm SD	+0.240 \pm 1.540 n = 37	-0.286 \pm 0.970	+0.363 \pm 1.633 n = 30	0.42				
Refractive error n (%)								
Myopia (≥ 0.5 D SE)	16 (42%) n = 38	4	12 (39%) n = 31	0.43				
Hyperopia (≥ 2.0 D SE)	5 (13%) n = 38	1	4 (13%) n = 31	1.00				
Astigmatism (≥ 0.75 D)	4 (11%) n = 38	2	3 (10%) n = 31	0.21				
Anisometropia (≥ 1.0 D SE)	5 (13%) n = 38	2	3 (10%) n = 31	0.21				

Notes: *Where numbers differ from the total number of children in the group, they are given separately for each category. **Undilated.

Abbreviations: BCVA, best corrected visual acuity; JIA, juvenile idiopathic arthritis; D, diopter; IOP, intraocular pressure; LE, left eye; LogMAR, logarithm of the minimum angle of resolution; n, number of children; RE, right eye; SD, standard deviation; SE, spherical equivalent; VA, visual acuity.

Table 4 Effects Of Youngsters' Eyesight On Quality Of Life Questionnaire (EYE-Q) And Child Health Assessment Questionnaire (CHAQ) Answered By Parents And Children at Assessment

Questionnaires On QoL	JIA Children, n = 40*	Uveitis, n = 7*	No Uveitis, n = 33*	P-Value
EYE-Q (0 – 4)^a Median outcome scores (range)**	0 (0 – 0.56)	0 (0 – 0.08)	0 (0 – 0.56)	0.78
CHAQ (0 – 3)^a disability index Median outcome scores (range)**	0 (0 – 1.32)	0 (0 – 0.38)	0 (0 – 1.32)	0.97
CHAQ (0 – 3)^a self report ≥ 8 yrs	0 (0 – 0.52) n = 15	0 (0 – 0) n = 2	0 (0 – 0.52) n = 13	0.44
CHAQ (0 – 3)^a parent report <8yrs	0.03 (0 – 1.32) n = 25	0.14 (0 – 0.38) n = 5	0.017 (0 – 1.32) n = 20	0.86
CHAQ VAS Median (range)				
Pain (0 – 10)^a	0 (0 – 4.7)	0 (0 – 3.5)	0 (0 – 4.7)	0.58
Pain (0 – 10)^a self report ≥ 8 yrs	0 (0 – 4.2) n = 15	0 (0 – 0) n = 2	0 (0 – 4.2) n = 13	0.44
Pain (0 – 10)^a parent report <8yrs	0.2 (0 – 4.7) n = 25	0 (0 – 3.5) n = 5	0.3 (0 – 4.7) n = 20	0.83
Overall well-being (0 – 10)^a	0.25 (0 – 6.4)	0 (0 – 5.4)	0.3 (0 – 6.4)	0.84
Overall well-being (0 – 10)^a self report ≥ 8 yrs	0 (0 – 2.5) n = 15	0 (0 – 0) n = 2	0 (0 – 2.5) n = 13	0.45
Overall well-being (0 – 10)^a parent report <8yrs	0.9 (0 – 6.4) n = 25	1. (0 – 5.4) n = 5	0.75 (0 – 6.4) n = 20	1.00

Notes: *Where numbers differ from the total number of children in the group, they are given separately for each category. **Median (range) is calculated on the mean of the scores for CHAQ disability index – parent and self-reports. ^aHigher scores indicate worse quality of life. Both pain and overall well-being were measured by a visual analog scale based on a Likert scale of 0-10 cm, where higher score indicates more pain/worse well-being. CHAQ disability index scores range from 0 to 3, at which 0 means without any difficulty and 3 means unable to do. We have excluded alternative 4 (not applicable) in our conclusion. EYE-Q scores range from 0 to 5, at which 0 means no difficulty and 4 means unable to do. We have excluded alternative 5 (not applicable) in our conclusion. Higher scores for both CHAQ and EYE-Q indicate worse quality of life.

Abbreviations: CHAQ, Childhood Health Assessment Questionnaire; EYE-Q, Effects of Youngsters' Eyesight on Quality of Life Questionnaire; JIA, juvenile idiopathic arthritis; n, number of children; QoL, quality of life; VAS, visual analogue scale.

Discussion

JIA is the most common chronic rheumatic disease in childhood. Children with JIA may have long-term disability and affected QoL. Therefore, there has been an increasing need to assess the daily physical function as well as visual function status and VRQoL in patients with JIA and JIA-U. CHAQ is a valid, culturally adapted, reliable and sensitive tool for the evaluation of functional outcomes in children with chronic arthritis.^{22,27} The aim of this study was to further evaluate visual and physical function and VRQoL in 40 children with JIA, of whom seven had JIA-U. We could not find any significant difference in QoL measured by CHAQ and EYE-Q between those with JIA alone and those with JIA-associated uveitis. However, this could be due to the fact that the patients were relatively stable in their JIA, but also due to the few cases with uveitis and/or other ocular problems. Furthermore, based

on our results, the longer the duration of the arthritis disease, the more the overall QoL is affected.

Our results are consistent with previous studies showing that children with JIA are primarily Caucasian, female and have positive ANA.^{7,28} Having positive ANA is the most important predictor for developing uveitis.⁹ The mean age of arthritis onset in children who developed uveitis was 3.3 years and the mean time interval between diagnosis of arthritis and the diagnosis of uveitis was 1.6 years, which is in accordance with Heiligenhaus et al⁷ and Papadopoulou et al⁹ reporting 3.8 years/21 months and 3.4 years/17 months, respectively. The median age of onset of uveitis after being diagnosed with arthritis was in the present study 1.7 years, which is in accordance with Papadopoulou et al⁹ presenting 12 months and Cassidy et al²⁹ reporting that uveitis can occur years after JIA onset. A recent Nordic-population-based

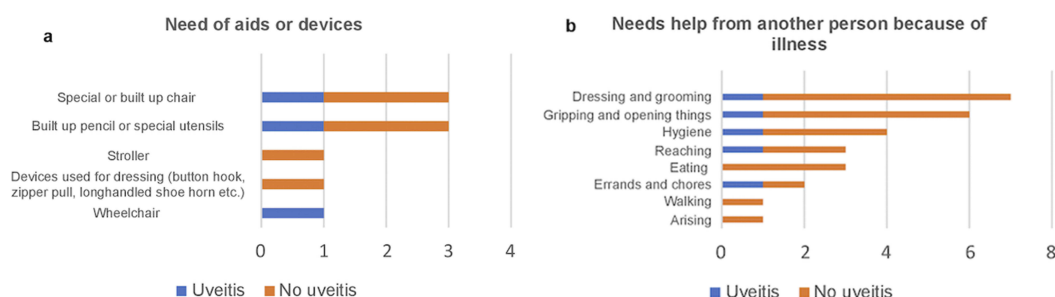


Figure 1 Number of children with juvenile idiopathic arthritis (JIA) needing special aids or devices documented by Child Health Assessment Questionnaire (CHAQ) (A) and number of children with JIA needing help from another person because of illness (B).

Table 5 Comparison Of Effects Of Youngsters' Eyesight On Quality Of Life Questionnaire (EYE-Q) And Child Health Assessment Questionnaire (CHAQ) In Children With Juvenile Idiopathic Arthritis Having Visual Acuity \leq or >0.7 Decimal (0.2 Logarithm Of The Minimum Angle Of Resolution)

Questionnaires On VR And HR QoL	JIA Children, n = 40*	VA Best Eye ≤ 0.7 Decimal (0.2 LogMAR), n = 5	VA Best Eye > 0.7 Decimal (0.2 LogMAR), n = 35*	P-Value
EYE-Q (0 – 4) ^a Median outcome scores (range)**	0.0 (0.0 – 0.56)	0.0 (0.0 – 0.56) n = 5	0 (0.0 – 0.11) = 35	0.013
CHAQ (0 – 3) ^a disability index Median outcome scores (range)**	0.0 (0.0 – 1.32) n = 40	0.143 (0.0 – 1.321) n = 5	0.0 (0.0 – 1.318) n = 35	0.066
CHAQ (0 – 3) ^a self report ≥ 8 yrs	0.0 (0.0 – 0.517) n = 15		0.0 (0.0 – 0.517) n = 15	
CHAQ (0 – 3) ^a parent report < 8 yrs	0.033 (0.0 – 1.321) n = 25	0.143 (0.0 – 1.321) n = 5	0.0 (0.0 – 1.318) n = 20	0.21
CHAQ VAS Median (range)				
Pain (0 – 10) ^a	0.0 (0.0 – 4.700) n = 40	0.40 (0.0 – 2.200) n = 5	0.0 (0.0 – 4.700) n = 35	0.63
Pain (0 – 10) ^a self report ≥ 8 yrs	0.0 (0.0 – 4.200) n = 15		0.0 (0.0 – 4.200) n = 15	
Pain (0 – 10) ^a parent report < 8 yrs	0.20 (0.0 – 4.700) n = 25	0.40 (0.0 – 2.200) n = 5	0.10 (0.0 – 4.700) n = 20	1.00
Overall well-being (0 – 10) ^a	0.0 (0.0 – 2.500) n = 15	1.70 (0.0 – 3.80) n = 5	0.0 (0.0 – 6.400) n = 35	0.12
Overall well-being (0 – 10) ^a self report ≥ 8 yrs	0.0 (0.0 – 2.500) n = 15		0.0 (0.0 – 2.500) n = 15	
Overall well-being (0 – 10) ^a parent report < 8 yrs	0.90 (0.0 – 6.40) n = 25	1.70 (0.0 – 3.80) n = 5	0.75 (0.0 – 6.40) n = 20	0.39

Notes: *Where numbers differ from the total number of children in the group, they are given separately for each category. **Median (range) is calculated on the mean of the scores for CHAQ disability index – parent and self-reports. ^aHigher scores indicate worse quality of life.

Abbreviations: CHAQ, Childhood Health Assessment Questionnaire; EYE-Q, Effects of Youngsters' Eyesight on Quality of Life Questionnaire; HR, health-related; JIA, juvenile idiopathic arthritis; LogMAR, logarithm of the minimum angle of resolution; n, number of children; QoL, quality of life; VA, visual acuity; VAS, visual analogue scale; VR, visual-related.

cohort study of 435 JIA children reported that uveitis occurred in 20.5% of these children, whereas in 53.9%, uveitis developed within the first year after the onset of arthritis.¹⁰

According to other studies, uveitis is more common in the OA subtype, whereas the extended OA subtype is thought to have the highest prevalence of uveitis (25%).^{28,30} In our

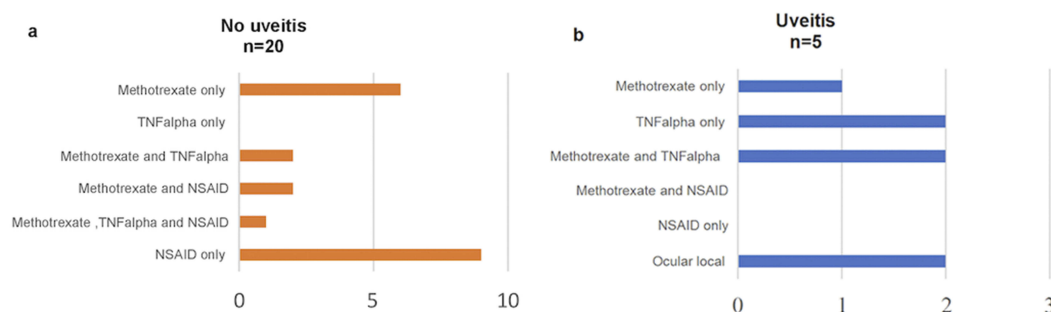


Figure 2 Medical treatment in 40 children without (A) and with (B) juvenile idiopathic arthritis (JIA)-associated uveitis at the time of assessment.

study, OA persistent was most common (57%), which is in accordance with Papadopoulou et al⁹ reporting 50% of persistent OA subtype among the observed uveitis population.

Fledelius et al³¹ found, in a long-term follow-up, that 40% of 65 patients with juvenile chronic arthritis were myopic, and the mean refraction in these patients was significantly more towards myopia than that of a control group. A possible explanation, according to them, is weakness of scleral connective tissue in the early active phases of eye growth due to chronic inflammation. However, it is unclear whether the patients in the previous mentioned study have/have had uveitis. Notably, 40% of the children in our study was myopic, of which four of these were children with JIA-U.

Angeles-Han and colleagues found, when comparing EYE-Q with logMAR VA in the best eye/unaffected eye, moderate correlations between child and parent EYE-Q scores.¹⁸ We found a weak correlation between BCVA LogMAR best eye and EYE-Q. However, worse EYE-Q scores were not necessarily associated with having uveitis; in fact, there was no sign of uveitis in two patients with the highest scores in EYE-Q. These patients may have/had other manifestations affecting their VR-QoL of yet unknown cause other than age. In several patients with the lowest scores possible in both CHAQ and EYE-Q, there was no need of medication. We also noted that the parent reports regarding need of aids/devices or need of assistance differed from the self-reports; the children older than 8 years of age did not need any of these.

The largest limitation of this study is the relatively small number of children with JIA and especially children with JIA-U which limits meaningful statistical analysis regarding EYE-Q and health-related OoL; however, our main focus was on the entire group and not only on those children with JIA-associated uveitis, especially when measuring with QoL in general. Most of the epidemiological data found in our study are in accordance with previous studies, despite

our small study population. Other studies such as Singh et al²¹ have found a strong correlation between the parents' and children's responses when comparing the disability index of CHAQ. In our study, we found a difference in the responses of CHAQ well-being between the self-reports and parent reports, which indicates that the parents of 25 patients younger than 8 years experienced lower QoL compared to the older children, but for better comparable analysis we would like to have the questionnaires completed by the children of the 25 parents as well. However, we should consider whether parents can serve as reliable proxy reporters of their child's capabilities. According to Eiser et al,³² relying on either the child's self-report or the parent's proxy report alone will not give a complete picture in the clinical setting; parents' reports of their children's health may be influenced by their own well-being, their involvement in treatment and their responsibility for the child's daily care as well as them having information of diagnosis, prognosis, etc. However, regardless of the reason for the differences between the self-reports and parents-reports, it is important to be aware of these differences as they may contribute to a great clinical significance.

In conclusion, children with PA JIA subtype in our study had the highest scores of physical function assessment, which indicates worst QoL. Worse VA is indeed in accordance with worse VR function. Both the CHAQ and EYE-Q instruments scored low in general, which we believe is due to effective management of the arthritis disease as well as good screening programmes, strong collaboration between the ophthalmic and the paediatric clinics.

Acknowledgments

The authors would like to thank all the children and their families for participating in the study, and Anders Pehrsson, Senior Biostatistician of Statistiska Konsultgruppen, Gothenburg, for the statistical analyses. This work was

supported by the Gothenburg Medical Society; the Aina Wallström and Mary-Ann Sjöblom Foundation for Medical Research; and the Wilhelm and Martina Lundgren Science Foundation.

Disclosure

The authors report no conflicts of interest in this work.

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