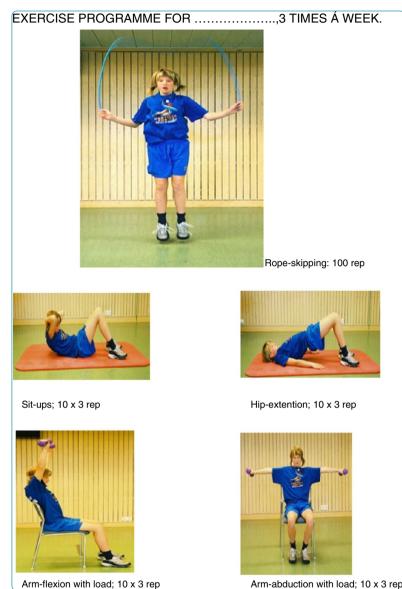


PEDIATRIC RHEUMATOLOGY



Arm-abduction with load; 10 x 3 rep

Muscle strength, physical fitness and well-being in children and adolescents with juvenile idiopathic arthritis and the effect of an exercise programme: a randomized controlled trial

Sandstedt et al.



RESEARCH



Open Access

Muscle strength, physical fitness and well-being in children and adolescents with juvenile idiopathic arthritis and the effect of an exercise programme: a randomized controlled trial

Eva Sandstedt^{1*†}, Anders Fasth^{1†}, Meta Nyström Eek^{1†} and Eva Beckung^{2†}

Abstract

Background: Decreased muscle strength, fitness and well-being are common in children and adolescents with juvenile idiopathic arthritis (JIA) compared to healthy peers. Biological drugs have improved health in children with JIA, but despite this pain is still a major symptom and bone health is reported as decreased in the group. The improvement made by the biological drugs makes it possible to more demanding exercises. To jump is an exercise that can improve bone heath, fitness and muscle strength. The aim of the study was to see if an exercise programme with jumps had an effect on muscle strength, physical fitness and well-being and how it was tolerated.

Methods: Muscle strength and well-being were studied before and after a 12-week exercise programme in 54 children and adolescents with JIA, 9–21 years old. The participants were randomized into an exercise and a control group. Muscle strength, fitness and well-being were documented before and after the training period and at follow-up after 6 months. Physical activity in leisure time was documented in diaries. The fitness/exercise programme was performed at home three times a week and included rope skipping and muscle strength training exercises.

Assessment included measurement of muscle strength with a handheld device, and with Grip-it, step-test for fitness with documentation of heart rate and pain perception and two questionnaires (CHAQ, CHQ) on well-being.

Results: There were no differences between exercise and control group regarding muscle strength, grip strength, fitness or well-being at base line. Muscle weakness was present in hip extensors, hip abductors and handgrip. For the exercise group muscle strength in hip and knee extensors increased after the 12-week exercise programme and was maintained in knee extensors at follow-up. There was no change in fitness tested with the individually adapted step-test. The CHQ questionnaire showed that pain was common in the exercise group and in the control group. There were only small changes in the CHAQ and CHQ after the training period. The fitness/exercise programme was well tolerated and pain did not increase during the study.

Conclusions: A weight bearing exercise programme, with muscle strength training with free weights and rope skipping was well tolerated without negative consequences on pain. It also improved muscle strength in the legs and can be recommended for children and adolescents with JIA.

Keywords: Muscle strength, Grip strength, Physical fitness, Well-being, JIA, Exercise programme

* Correspondence: eva.sandstedt@vgregion.se

[†]Equal contributors

¹Department of Paediatrics, Institute of Clinical Sciences, University of Gothenburg, Gothenburg, Sweden

Full list of author information is available at the end of the article



© 2013 Sandstedt et al.; licensee BioMed Central Ltd. This is an Open Access article distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/2.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Background

Children and adolescents with juvenile idiopathic arthritis (JIA) in most parts of the world have decreased muscle strength, bone health and well-being compared to healthy peers [1-8]. The disease can affect school performance, physical training, family life, and activities in leisure time with peers [9-11]. Kimura et al. declare in a study from 2008 that pain is one of the major symptoms and limits the activities, disrupts school attendance and contributes to psychosocial distress [12].

The last decade has seen the introduction of biological drugs, e.g. anti-tumor necrosis factor alpha (anti-TNF α) also for paediatric rheumatic disorders [13,14]. The medical effect of anti-TNF is especially high in children with polyarticular onset of JIA [13,14]. In subjects with JIA the effects are described as improvement in functional ability, health-related quality of life, pain, sleep quality and daily participation and in terms of less flares or inflammatory active joints [15-17]. Anti-TNF α drugs are effective, safe and well tolerated in children with JIA [14-19].

Despite the use of biological agents, pain is reported as the major symptom of the disease, and joint pain is the leading cause of disability in this disease [12]. The authors describe pain perception as multifactorial and therefore require "a bio-psychosocial model that includes the individual's age, developmental status, coping ability, mood, stress levels, and environmental and family factors, in addition to disease status and severity" [12].

Physical activity is important from a health perspective, especially in the subgroups with the polyarticular and extended oligoarticular categories [1,2,12,17,20]. Different physical activities have been studied such as jumping with a rope (rope-skipping) and exercise programs in water [4,6,7,21-24]. Jumping has influence on bone health and foot orthotics can significantly improve pain, speed of ambulation, and self-rated activity and functional ability [23,25].

Exercise programs with weight bearing exercises have been shown to improve both muscle strength and bone mass [6,22,23]. The exercise programmes in these studies were at different intensity levels and of different duration and physical activity in leisure time was not fully documented [4,6,22]. Takken showed that also cardiovascular fitness was decreased in children with JIA compared to healthy peers and point out the importance of cardiovascular fitness and motor performance as a part of total well-being [24]. Muscle strength is an important part in a fitness programme; muscle weakness in children with JIA is reported in many studies since the 1990s [2,8,9]. There is, however, a lack of knowledge about physical exercise levels and the impact on pain and well-being.

Physical fitness is described as a state of well-being with energy to participate in a variety of physical activities [26]. Frankala-Pinkham et al. stress the importance to incorporate more strategies to increase fitness, physical activity, and participation in the rehabilitation programme to improve quality of life (QoL) [27]. Questions concerning well-being and the impact of social and psychological functioning are well covered in the Child Health Questionnaire (CHQ) [28-30]. A couple of studies have reported that children with JIA after exercise interventions have less physical impairment or discomfort but report low levels of psychosocial abilities such as self esteem, psychosocial functioning and high levels of pain [21,22,25]. This seems to be a pattern for children with different chronic diseases or disabilities [31,32].

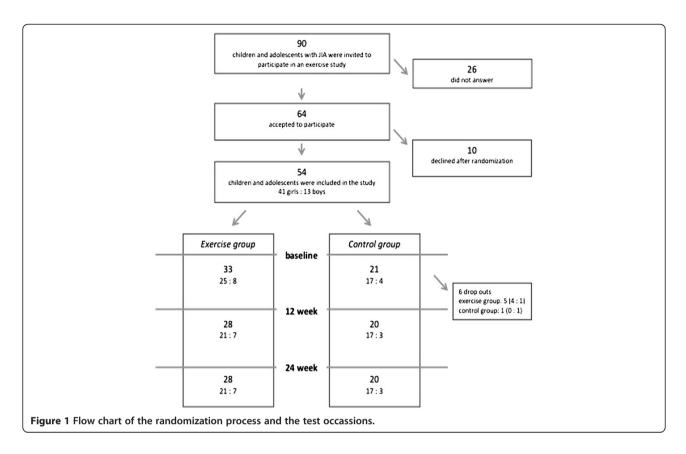
At our hospital the children with JIA attend the hospital regularly for physical training, which is time-consuming and costly both for the families and the health care system. An easy-to-handle home-based exercise programme making the patient less dependent on the physical therapist was needed, which was the impetus for this study. The aim of the study was to evaluate muscle strength, grip strength, physical fitness and well-being in a cohort of children and adolescents with JIA and the effects of a home-based exercise programme.

Methods

Subjects

The study is the second part of a randomized controlled trial of 54 children and adolescents with JIA that studied the effects of an exercise programme on bone health, muscle strength, fitness and well-being. Bone health and leisure time activities have been reported earlier and the randomization process has been described in detail [23]. The inclusion criteria were polyarticular or extended oligoarticular arthritis, treated with methotrexate, TNFblockers and/or prednisone, and in need of repeated corticosteroid injections of joints in the lower extremities. Medical records were obtained and three participants were found to be diagnosed with enthesitis related and psoriatic arthritis. After written consent by the parents and assent by the children, the subjects were randomized into an exercise or a control group. The person carrying out the group allocation was blinded. A flow chart of randomization and training is presented in Figure 1. There were 10 dropouts from the control group after the randomization, as they had preferred to belong to the exercise group. There were another six dropouts after the first test occasion due to the families' lack of time.

Muscle strength, range of motion, balance, fitness and well-being were studied before and after a 12-week exercise programme. The participants were evaluated three times; at base line, after 3 months at the end of the training period and at follow-up at 6 months. The same physiotherapist, who was blinded to the previous measurement, performed all measurements.



Range of motion and balance

Range of motion (ROM) was measured with a plastic goniometer in dorsal and plantar flexion of the ankle; flexion, abduction and rotation of the hip; and rotation, abduction and flexion of the shoulder.

The Balance Reach Test for children [33] was performed before and after the exercise programme. It has a good test-retest and inter-rater reliability with intraclass correlation coefficients between 0.54 and 0.88 and 0.54 -0.93 respectively [33].

Muscle strength

Muscle strength in arms and legs was tested with a handheld device (adapted Chatillon[®], dynamometer; Axel Ericson Medical AB, Gothenburg Sweden), in eight muscle groups (shoulder abduction, elbow: extension, flexion, hip: extension, flexion, abduction, knee: extension, ankle: dorsal flexors) using the "make" technique and with standardized positions. After instruction and familiarisation with the procedure, three attempts were made and the maximum recording was used for data analysis. Lever arm for each muscle group was measured with a tape measure and torque was calculated (Nm). For six muscle groups the positions used in this study were similar to the one in a normative study [34], presenting equations for a predicted value for every muscle group based on age, sex and body weight. Muscle strength was compared with the

normative material and calculated as a percentage of the predicted value. Thus it was possible to evaluate the whole group despite their different ages.

Grip strength was measured with Grippit (Detektor AB, Goteborg, Sweden) [35]. The instrument estimates peak strength over a 10 s period and the test was performed three times for each hand. The maximum recording was used for statistical analysis. Measurements were compared to normative values obtained with the same device [35], with data presented as mean ± 1 SD and grouped according to age and sex. The data in this study was classified in a three level ordinal scale: *strong* = outside +1SD from mean, *average* = within mean ± 1 SD and *weak* = outside -1 SD from mean.

Physical fitness

Fitness was tested with a step-test. In the Harward step test from 1956 the step board was 45 cm high and the speed 30 steps per minute for 5 minutes or until exhaustion [36]. In this study the test was adapted, by using a lower step board (20 cm high), in order not to provoke pain. The participants were stepping on and off the step board for six minutes and a metronome was used for keeping an individually chosen speed. Heart rate was documented once a minute during the test and exertion was documented with the Borg Scale 6–20 [37]. The power in Watt was calculated taking into account body weight, gravity, the height of the step board and speed ($P = m \times g \times v$) and was normalised to body weight (W/Kg). The test was considered as an individual sub maximal test.

Quality of life

The Child Health Assessment Questionnaire (CHAQ) is a questionnaire that is diagnose-specific for JIA, and is translated into and validated in Swedish [29]. The instrument refers to the last 14 days and includes eight different categories of activities (dressing, eating, walking, getting up, reaching, gripping, hygiene and activity). Each question is scored from 0 to 3 (0= no difficulty, 1= some difficulty, 2= much difficulty, 3= unable to complete task). The total score varies from 0 (no limitation) to 3 (extensive limitation). The instrument is recommended for children with JIA by the International League of Associations for Rheumatology (ILAR) and the Paediatric Rheumatology International Trials Organization (PRINTO) [29].

The Child Health Questionnaire (CHQ-C87) is a survey of the physical and psychosocial health of children 5 years of age and older. The questionnaire refers to the wellbeing status for the last four weeks. It was developed for children in the general population (for which normative data are available), and for children with chronic conditions [30]. The instrument has been validated for Swedish children, 9–16 years old, with epilepsy, diabetes and JIA [28]. It has a multidimensional profile consisting of 87 questions in twelve different domains (see Table 1). Scoring algorithms are provided for the different domains [30]. The manual consists of a Scale Scoring for a clinical sample of children with JIA, epilepsy, asthma and with psychiatric disorders.

Pain

The children were asked to report if pain occurred during the test occasions. Presence of pain was documented with a 10 centimetres visual analogue scale (VAS). Pain in a perspective of health/well-being was also reported within the questionnaires CHAQ and CHQ.

Fitness programme

The participants fulfilled a training programme three times a week for 12 weeks. The exercise programme consisted of rope skipping, muscle strength, core exercises and exercises with free weights for arms (Appendix). The programme has been described in detail earlier [23]. The number of repetitions performed was documented in an exercise diary. Physical activity in leisure time outside the programme was also documented in an activity diary.

Statistical methods

For comparison between groups, Mann Whitney *U*-test was used for grip strength and for the questionnaires. *T*-test was used for muscle strength with myometer. The repeated measures ANOVA method was used for comparison of muscle strength and for step-test at baseline, after training and at follow-up. Data was tested with Mauchlys test of sphericity, and if sphericity was not assumed the Greenhouse-Geisser method/procedure was used for analysis.

As the results from the questionnaires were not normally distributed the Friedman test was used for repeated measures followed by post hoc testing with Wilcoxon singed rank test.

P-values of 0.05 or less were considered evidence of statistically significant findings. In the post hoc analysis Bonferroni adjustment for multiple comparisons was used.

Software packages Statview, SPSS (version 17.0) and SPSS for Mac (version 19.0) were used for statistical analysis.

Ethics

This study was carried out in compliance with the Helsinki Declaration and was approved by The Regional

Table 1 CHAQ and CHQ, mean and median at baseline

				Exercise group n=32		Control group n=21	
				Mean	Median	Mean	Median
CHAQ				.55	.50	.50	.38
CHQ domain	Nr of items	Low Score	High Score				
Physical functioning	9	Child is limited a lot in performing all physical activities, including self-care, due to health.	Child performs all types of physical activities, including the most vigorous, without limitations, due to health	79.4	83.5	81.0	80.0
Role Emotional	3	Child is limited a lot in school work or activities with friends as a result of emotional problems	Child has no limitations in school work or activities with friends as a result of emotional problems	89.0	100.0	83.3	100.0
Role Behavioural	3	Child is limited a lot in school work or activities with friends as a result of behavioural problems	Child has no limitations in school work or activities with friends as a result of behavioural problems	89.9	100.0	94.6	100.0

Ethics Committee in Gothenburg. Written consent was obtained from the participants and from their parents.

Results

Fifty-four children and adolescents were included in the study, with a mean age of 13.9 years (range 8.8-21.6). There were 41 girls and 13 boys, randomized in an exercise and in a control group (Table 2). There was a difference in age between the exercise group and the control group, not reaching statistical significance (p=0.059), but there were a statistically significant difference in height (p=0.007) and weight (p=0.026).

Range of motion and balance

There were no differences between groups at baseline for measurement of ROM in the Balance Reach Test and there were no significant changes during the study.

Muscle strength

All children did not fulfil the whole protocol at all test occasions and only muscle groups with complete measurements were analysed. Muscle strength measurements taken at baseline were compared with the normative material for six muscle groups (see Table 3 and Figure 2). Values for hip abductors (33-38%) and hip extensors (52-55%) were below the limits of the 95% prediction interval. There were no significant differences between control and exercise group (Table 3). Values were also compared in order to see if age had influence on muscle strength. No significant differences were found when younger children (8–12 years) were compared with older children (13–16 years) (Figure 2).

Forty-five children had measurement of grip strength that could be compared to normative values [33], 17 in control and 28 in exercise group. The comparison showed weakness, with 28 children showing values below the normative mean -1 SD. Sixteen children were average

Table 2 Distribution	of gender,	age,	height,	weight,
disease onset and ty	pe			

	Exercise group n=33	Control group n=21
Girls (n=)	25 (76%)	17 (81%)
Age (years)	13.3 (8.8-19.9)	14.9 (8.8-20.6)
Height (cm)	153.4 (128.6-175.8)	163.8 (129–185.8) *
Weight (kg)	47.9 (22.3-78)	56 (35.8-82.9) *
Disease onset (years)	6.1 (1.2-16.5)	4.8 (1–13.4)
JIA categories at onset		
Polyarticular	20	9
Oligoarticular	7	8
Entesites related or psoriatic	1	3

* = statistically significant difference p<0.05.

Table 3 Muscle strength, expressed as a percentage of predicted value, mean (SD)

Muscle –group	n	Exercise	n	Control	<i>p</i> -value ^a
Shoulder abductors	28	70.5 (17.6)	15	73.4 (21.6)	0.636
Elbow extensors	28	84.1 (19.6)	15	75.7 (12.7)	0.144
Hip extensors	27	52.1 (18.9)	15	55.9 (13.1)	0.496
Hip flexors	28	91.8 (17.8)	15	92.9 (25.5)	0.870
Hip abductors	28	33.9 (10.2)	15	38.8 (10.4)	0.138
Knee extensors	28	80.9 (21.5)	15	84.4 (15.4)	0.575
a					

unpaired *t*-test.

(within 1 SD from mean) and one was strong with values above mean +1SD. There were no significant differences between groups when corrected for/compared to age (Table 4).

Changes after fitness programme

There were no changes in grip strength during or after the training period (see Table 4).

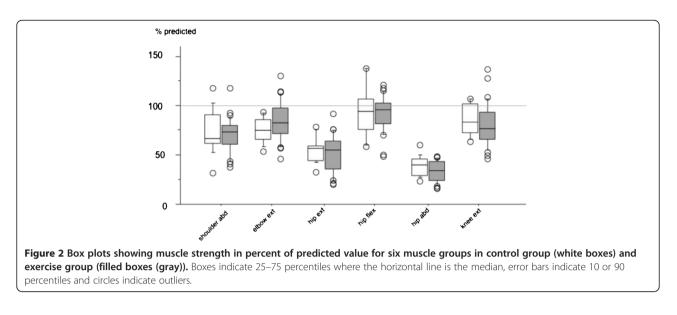
Measurements of muscle strength of the legs are presented in Table 4. Statistically significant changes were found in the exercise group after training, with an increase in hip extensors and knee extensors, compared to baseline. Knee extensor strength was maintained at follow-up.

Physical fitness

There were nine dropouts in the step-test due to pain; five experienced pain in the knee, one in the hip and three in the foot. The power in W/Kg and heart rate in the step-test are shown in Table 5. There were no differences between groups before training started regarding power and heart rate. There were no changes in heart rate or perceived exertion after training (Table 6).

QoL and well-being

Results of CHAQ and CHQ are shown in Tables 1 and 7. Fifty-three children completed the CHAQ and CHQ at baseline. There were no differences between the exercise and the control group. The CHQ was only used in scientific studies such as the study of Norrby [38]. The CHQ was used for all the participants in this study. The adolescents even the one 20.6 years of age and those older than 16 years were considered adolescents as the participants still were patients at the Children's hospital. Our subjects showed low values in the domain "bodily pain" and also in the domains "general health" and "mental health" at baseline. 35 children fulfilled the CHAQ and 39 the CHQ at all test occasions. There was no increase in pain during the study. There were only small changes in both of the questionnaires. In the control group there was a statistically significant increase in CHQ domain "role physical" at the end of the study period. There was a tendency to improved "mental



health" in the exercise group and deterioration in the control group in "general health".

Exercise programme

The participants in the exercise group fulfilled about 70% of expected numbers of exercises (Figure 3).

Discussion

The study revealed muscle weakness and the presence of pain in children and adolescents with JIA. Pain is not an obstacle for performing the physical fitness programme, but there were ten dropouts mainly due to pain in the testing procedure.

Table 4 Grip and muscle strength before, after training and at follow-up

			Exercise group n=2	6	Control group n=19			
		Before	3 months	6 months	Before	3 months	6 months	
Grip strength	Right	157.9 (73.0)	173.4 (76.1)	174.4 (68.5)	197.1 (66.1)	208.8 (75.1)	216.9 (72.5)	
	Left	150.5 (74.1)	149.1 (62.4)	155.1 (62.5)	185.4 (60.2)	185.1 (71.8)	194.0 (72.8)	
Muscle group								
Shoulder abd	Right	21.8 (10.3)	23.3 (10.9)	24.0 (11.8)	26.5 (10.9)	25.5 (8.8)	30.2 (8.6) †	
	Left	21.9 (10.2)	24.0 (11.3) *	23.9 (11.1)	25.0 (8.7)	25.5 (7.7)	27.0 (7.7)	
Elbow ext	Right	21.6 (12.7)	21.9 (9.3)	23.4 (9.8)	26.0 (11.5)	25.4 (10.6)	26.7 (9.7)	
	Left	21.1 (9.0)	22.2 (8.8)	23.0 (8.0)	24.9 (8.0)	25.8 (8.5)	26.3 (8.8)	
Elbow flex	Right	22.2 (12.9)	24.8 (10.8)	26.5 (12.6) *	31.0 (14.1)	31.8 (12.0)	33.7 (13.2)	
	Left	21.1 (11.6)	23.5 (10.9)	24.6 (10.9) *	30.0 (10.9)	31.5 (12.2)	31.1 (10.8)	
Hip ext	Right	47.5 (21.4)	54.3 (22.4) *	49.1 (19.6)	66.7 (24.4)	64.0 (27.3)	60.7 (25.1)	
	Left	47.3 (19.3)	56.1 (22.5) *	53.3 (20.4)	64.1 (20.1)	64.9 (25.0)	61.7 (22.9)	
Hip flex	Right	53.5 (20.7)	56.0 (21.7)	54.1 (21.9)	68.8 (25.5)	64.3 (22.4)	66.2 (21.6)	
	Left	49.9 (17.9)	53.9 (19.4)	52.4 (20.1)	62.0 (21.4)	60.8 (17.0)	63.4 (18.9)	
Hip abd	Right	26.5 (13.6)	28.9 (15.1)	29.6 (10.6)	40.6 (18.3)	40.2 (15.5)	33.3 (9.7)	
	Left	24.6 (12.3)	30.7 (15.3) *	32.9 (14.1) *	35.9 (12.3)	37.0 (14.1)	38.0 (12.8)	
Knee ext	Right	57.3 (21.5)	65.8 (22.3) *	67.5 (22.3) *	79.3 (19.2)	78.7 (21.9)	77.1 (20.9)	
	Left	55.8 (22.8)	61.7 (18.0) *	65.9 (26.4) *	78.8 (18.3)	79.6 (20.3)	78.8 (18.1)	
Dorsiflex	Right	19.3 (5.2)	20.1 (6.3)	21.8 (6.1) *	23.8 (6.8)	22.1 (6.5)	25.2 (5.6) †	
	Left	20.3 (5.5)	20.7 (6.7)	21.8 (5.8)	23.9 (6.5)	21.9 (4.8)	23.8 (5.0)	

* = statistically significant difference to measurement 1 (baseline) p<0.03.

 \dagger = statistically significant difference to measurement 2 (after 12 week training period) p<0.03.

Data in Nm, mean (SD).

Tab	le 5	Result	s from	the	e step	-test:	power	and	heart	rate
-----	------	--------	--------	-----	--------	--------	-------	-----	-------	------

	Exercise group n = 33	Control group n = 19	<i>p</i> -value ^a				
W/Kg	1.20 (1.43)	1.23 (0.17)	0.508				
Heart rate	170.9 (16.14)	176.8 (19.8)	0.246				
Companies hot was an ended hot training many (CD)							

Comparison between groups before training, mean (SD).

Muscle strength in the leg

Muscle weakness in knee extensors, elbow flexors and in ankle dorsal flexors has since 1995 been reported in children with JIA compared to healthy peers [2,3,7,8]. In our study muscle weakness was found in hip extensors, hip abductors and in handgrip strength. The other muscle groups were also below predicted values but within the normative range. In contrast to other studies our group was within normal values in knee extensors [3,7,8] Saarinen, Lindeman, and Broström all found that muscle weakness was present in children and adolescents with JIA, [3,7,8].

Grip strength

More than 50% of the children in this study had a lower grip strength compared to peers and many of the children in the group had received corticosteroids in hands and fingers. No increase was found in hand grip strength after the training, which was not to be expected, as there were no specific exercises for grip strength in the programme. Earlier studies with children and adolescents report lower grip strength and that performance in writing and drawing in school give negative consequences compared to peers [39,40]. A pilot study shows that children with JIA are suffering from handwriting difficulties and are limited mainly due to pain and the inability to sustain handwriting for a longer period of time [40].

Exercise programme

The increased muscle strength in hip and knee extensors found after 12 weeks exercise correlates well with the training programme that included of exercises for both hip and knee extensors. Rope skipping seemed to be an effective exercise to improve muscle strength in these muscle groups. Improvement in knee extensors is especially important, as the knee joint is the most affected joint in children and adolescents with JIA [10]. Earlier studies on groups of children and adolescents with JIA [1-4,20-25,38,41-43] show that this group is in inferior regarding functional ability, physical fitness and cardiovascular capacity compared to peers.

Earlier studies with rope skipping also reported significant improvement in bone health and muscle strength both for children with JIA and for healthy children [6,23,44]. The physical fitness programme also covered items for muscle strength in core muscles and muscles around the shoulders. The failure to increase muscle strength in these muscle groups may have been due to that the weights used were not heavy enough.

No changes in fitness, in terms of heart rate and exertion, were found after training. This may be due to the tests not being sensitive enough or the exercises were not sufficiently challenging regarding fitness. The steptest was performed on a 20 cm step board in order not to provoke knee pain; despite this ten children did not fulfil the test mainly due to pain in knee, hip or foot. A higher step board had been more demanding and could perhaps have shown a difference in the measurement. Measuring physical fitness with a cycle ergometer may be less painful for children with JIA. On the other hand the step-test with weight bearing exercise is a test close to functions that are important in daily life. Functional ability, physical fitness and cardiovascular capacity have earlier been studied in groups of children with JIA [20-24,40] who found that they were inferior regarding physical fitness compared to peers. There was a lack of knowledge regarding to the participants' ability to perform the programme and how well they adhered to it. The focus had been on daily activity and of participating in physics at school. The focus had not been on progressive cardiovascular fitness.

With the improvement in medical treatment it is important to keep addressing physical fitness, as it is a prerequisite for good health [45,46].

The exercise programme in our study was designed to meet the need for physical training, in accordance to the ILAR recommendation. The frequency of three times a week and the level of cardiovascular effort and weights were well tolerated. In this study we did not individualize the programme or increase the number of repetitions or the load during the training period, which might have given another outcome.

Noteworthy, our protocol did not render any increase in pain during the training period. By the introduction of an exercise programme we also hoped to encourage

Table 6 Step-test; heart rate and exertion at baseline and after 3 and 6 months, mean (SD)

		Exercise group	n=25		Control group r	า=17
	Before	3 months	6 months months	Before	3 months	6 months months
Heart rate	168.4 (16.4)	169.1 (15.2)	168.0 (16.9)	177.6 (20.9)	173.6 (23.6)	177.0 (18.4)
Borg scale	14.6 (2.86)	14.0 (3.03)	14.2 (3.58)	15.5 (1.97)	15.0 (3.00)	14.9 (2.89)

Table 7 CHAQ and CHQ

		Exei	cise group			Control group			
		Baseline	3 months	6 months		Baseline	3 months	6 months	
		Mean	Mean	Mean		Mean	Mean	Mean	
CHAQ	n=20	0.50	0.41	0.50	n=15	0.44	0.46	0.37	
CHQ domain	n=23				n=16				
Physical functioning		83.4	88.4	87.6		80.4	86.9	84.63	
Role Emotional		91.7	91.9	87.2		81.6	86.8	90.5	
Role Behavioural		88.9	93.7	90.3		92.9	97.8	97.1	
Role physical		91.2	94.2	89.5		80.2	89.8	96.3*	
Bodily pain	n=22	59.3	71.8	64.1	n=15	56.7	56.7	64.7	
Behaviour	n=18	82.7	82.5	83.6	n=15	84.5	82.4	86.5	
Mental health		71.0	74.3	77.2		70.1	70.3	71.6	
Self esteem		78.7	82.0	80.0		80.3	78.6	81.9	
General health	n=22	67.2	70.6	70.6	n=14	66.7	59.2	69.7	

* difference compared to baseline, p<0.017. Bonferroni adjustment for multiple comparisons.

Outcome in mean at baseline, after 3 and 6 months.

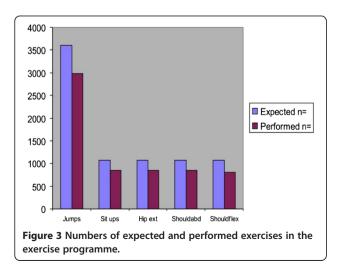
the children to change from a sedentary to an active life style. As reported earlier, physical activity increased in the group during the study and at follow-up [23].

Out of the 54 participants 48 completed the training, but there were dropouts in some of the measurements. This was explained by; lack of time, pain, and different psychosocial reasons.

Well-being

The questionnaires showed that the children in the study did not have any difficulties to carry out daily activities.

Compared to an earlier study in the same region and compared to the results from normative data [38] our group scored higher on the domain "general health" but were at about the same levels on the other domains. There were only a few changes during the 6-month period. The CHAQ has been debated because of lack of



sensitivity in view of changes in rehabilitation [29]. Dempster et al. in a study from 2001 showed that a minimal clinically important improvement is represented by a median change of -0, 13 in the CHAQ [47]. They also found a minimal clinically important deterioration by a median change of 0.75. In our study no improvement or deterioration was found.

Our results confirm that it is not an optimal questionnaire for children with a moderate impairment and being in an inactive phase of the disease, as it did not capture the spontaneous positive comments from participants during the study.

In our study the children reported low levels in the domain "bodily pain" and also in the domains "general health" and "mental health". The pathogenesis of pain in children with rheumatic diseases is multifactorial, and disease treatment alone is often not enough to alleviate it. Many researchers stress that children with chronic conditions can have "hidden" consequences on self esteem and well-being why pain treatment should include non-pharmacological interventions, for example exercise and cognitive-behavioural therapy, for better outcome on their general health [12,17,45,46].

Conclusions

Muscle weakness was present in hip extensors, hip abductors and in the handgrip. Muscle strength in hip and knee extensors increased after the 12-week exercise programme and was maintained in knee extensors at follow-up. Pain was common in the group. The exercise programme was well tolerated, there was a compliance of 70% to the programme and pain did not increase during the study. The study shows that a weight bearing fitness programme with muscle strength training including free weights and rope skipping can be recommended for children and adolescents with JIA.

Abbreviations

anti-TNFa: anti Tumor Necrosis Factor alpha; CHAQ: Child Health Assessment Questionnaire; CHQ: Child Health Questionnaire; ILAR: International League of Association for Rheumatology; JIA: Juvenile Idiopathic Arthritis; Nm: Newton metre; PRINTO: Peadiatric Rheumatology International Trials Organization; QoL: Quality of Life; ROM: Range of movement; VAS: Visual Analogue Scale; W/kg: Weight/kilogram.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

ES carried out the conception and design, all measurements (except DXA), performed analyses, supervised the training period and drafted the manuscript. AF participated in the design of the study and was involved in drafting the manuscript. MNE performed statistical analyses, provided reference data and was involved in drafting the manuscript. EB participated in the design of the study and drafted the manuscript. All authors have read and approved the final manuscript.

Acknowledgements

We would like to thank all children and parents who participated and made this study possible. We thank the Norrbacka Eugenia Foundation, The Research and Development Foundation of Göteborg and Bohuslän for funding this research, Västra Götaland County Council (agreement between government and university hospital regarding education of medical students and medical research).

Author details

¹Department of Paediatrics, Institute of Clinical Sciences, University of Gothenburg, Gothenburg, Sweden. ²Department of Neuroscience and Physiotherapy, Sahlgrenska Academy, University of Gothenburg, Gothenburg, Sweden.

Received: 16 November 2012 Accepted: 9 February 2013 Published: 22 February 2013

References

- Maggio AB, Hofer MF, Martin XE, Marchand LM, Beghetti M, Farpour-Lambert NJ: Reduced physical activity level and cardiorespiratory fitness in children with chronic diseases. *Eur J Pediatr* 2010, 169:1187–1193.
- Burnham JM, Shults J, Dubner SE, Sembhi H, Zemel BS, Leonard MB: Bone density, structure, and strength in juvenile idiopathic arthritis: importance of disease severity and muscle deficits. *Arthritis Rheum* 2008, 58:2518–2527.
- Saarinen J, Lehtonen K, Malkia E, Lahdenne P: Lower extremity isometric strength in children with juvenile idiopathic arthritis. *Clin Exp Rheumatol* 2008, 26:947–953.
- 4. Takken T, van der Net J, Kuis W, Helders PJ: Physical activity and health related physical fitness in children with juvenile idiopathic arthritis. *Ann Rheum Dis* 2003, **62**:885–889.
- Ganotti ME, Nahorniak M, Gorton GE: Can exercise influence low bone mineral density in children with juvenile rheumatoid arthritis? *Pediatr Phys Ther* 2007, 19:128–139.
- Klepper SE: Exercise in pediatric rheumatic diseases. Curr Opin Rheumatol 2008, 20:619–624.
- Hedengren E, Knutson LM, Haglund-Akerlind Y, Hagelberg S: Lower extremity isometric joint torque in children with juvenile chronic arthritis. Scand J Rheumatol 2001, 30:69–76.
- Lindehammar H, Sandstedt P: Measurement of quadriceps muscle strength and bulk in juvenile chronic arthritis. A prospective, longitudinal, 2 year survey. J Rheumatol 1998, 25:2240–2248.
- 9. Andersson Gäre B, Fasth A, Wiklund I: Measurement of functional status in juvenile chronic arthritis; evaluation of a Swedish version of the

Child Health Assessment Questionnaire. Clin Exp Rheumatol 1993, 11:569–576.

- Berntsson L, Berg M, Brydolf M, Hellström AL: Adolescents' experiences of well-being when living with a long-term illness or disability. Scand J Caring Sci 2007, 21:419–425.
- Berntson L, Andersson Gäre B, Fasth A, Herlin T, Kristinsson J, Lahdenne P, Marhaug G, Nielsen S, Pelkonen P, Rygg M, Nordic Study Group: Incidence of juvenile idiopathic arthritis in the Nordic countries. A population based study with special reference to the validity of the ILAR and EULAR criteria. J Rheumatol 2003, 30:2275–2282.
- 12. Kimura Y, Walco GA: Treatment of chronic pain in pediatric chronic disease. *Nat Clin Pract Rheumatol* 2007, **3:**210–218.
- 13. Wallace CA, Giannini EH, Spalding SJ, Hashkes PJ, O'Neil KM, Zeft AS, Szer IS, Ringold S, Brunner HI, Scjanberg LE, Sundel RP, Milojevic D, Punaro MG, Chira P, Gottlieb BS, Higgins GC, Ilowite NT, Kimura Y, Hamilton S, Johnson A, Haug B, Lovell DJ, Childhood Arthritis and Rheumatology research Alliance: Trial of early aggressive therapy in polyarticular juvenile idiopathic arthritis. Arthritis Rheum 2012, 64:2012–2021.
- 14. Murray KJ, Lovell DJ: Advanced therapy for juvenile arthritis. *Best Practice Res Clin Rheumatol* 2002, **16:**361–378.
- 15. Ruperto N, Lovell DJ, Li T, Sztajnbok F, Goldenstein-Schainberg C, Scheinberg M, Penades IC, Fischbach M, Alcala JO, Hashkes PJ, Hom C, Jung L, Lepore L, Oliveira S, Wallace C, Alessio M, Quartier P, Cortis E, Eberhard A, Simonini G, Lemelle I, Chalom EC, Sigal LH, Block A, Covucci A, Nys M, Martini A, Giannini EH: Paediatric Rheumatology International Trials Organisation (PRINTO); Pediatric Rheumatology Collaborative Study Group (PRCSG). Abatacept improves health-related quality of life, pain, sleep quality, and daily participation in subjects with juvenile idiopathic arthritis. Arthritis Care Res (Hoboken) 2010, 62:1542–1551.
- 16. Prince FHM, Geerdink LM, Borsboom GJJM, Twilt M, van Rossum MAJ, Hoppenreijs EPAH, ten Cate R, Koopman-Keemink Y, van Santen-Hoeufft M, Raat H, van Suijlekom-Smit LWA: Major improvements in health-related quality of life during the use of etanercept in patients with previously refractory juvenile idiopathic arthritis. Ann Rheum Dis 2010, 69:138–142.
- Perandino LA, de Sá-Pinto AL, Roschel H, Benatti FB, Lima FR, Bonfá E, Gualano B: Exercise as a therapeutic tool to counteract inflammation and clinical symptoms in autoimmune rheumatic diseases. *Autoimmun Rev* 2012, 12:218–224.
- Halbig M, Horneff G: Improvement of functional ability in children with juvenile idiopathic arthritis by treatment with etanercept. *Rheumatol Int* 2009, 30:229–238.
- Sevcic K, Orban I, Brodszky V, Bazso A, Balogh Z, Poor G, Kiss E: Experiences with tumor necrosis factor –(alpha) inhibitors in patients with juvenile idiopathic arthritis: Hungarian data from the National Institute of Rheumatology and Physiotherapy Registry. *Rheumatology (Oxford)* 2011, 50:1337–1340.
- Stephens S, Singh-Grewald D, Bar-Or O, Beyene J, Cameron B, Leblanc C, Schneider R, Scneiderman-Walker J, Selvadurai H, Silverman ED, Spiegel L, Tse S, Wright V, Laxer RM, Feldman BM: Reliability of Testing and Functional Activity Questionnaires in Children With Juvenile Arthritis. Arthritis Rheum 2007, 57:1446–1452.
- 21. Klepper SE: Effects of an eight-week physical conditioning programme on disease signs and symptoms in children with chronic arthritis. *Arthritis Care Res* 1999, **12**:52–60.
- Farpour-Lambert NJ, Keller-Marchand L, Rizzoli R, Schwitzgebel V, Dubuis JM, Hans D, Hofer MF, Suter S: Physical exercise and bone development in chronically ill children. *Rev Med Suisse Romande* 2004, 124:73–75.
- Sandstedt E, Fasth A, Fors H, Beckung E: Bone health in children and adolescents with juvenile idiopathic arthritis and the influence of shortterm physical exercise. *Pediatr Phys Ther* 2012, 24:155–161.
- 24. Takken T, Van Der Net J, Kuis W, Helders PJ: Aquatic fitness training for children with juvenile idiopathic arthritis. *Rheumatology (Oxford)* 2003, 42:1408–1414.
- Powell M, Seid M, Szer IS: Efficacy of custom foot orthotics in improving pain and functional status in children with juvenile idiopathic arthritis: a randomized trial. J Rheumatol 2005, 32:943–950.
- 26. Howely ET, Franks BD: *Health Fitness Instructor's Handbook*. Champaign, Illinois: Human Kinetics Pub; 1997.
- Fragala-Pinkham M, O'Neil M, Bjornson K, Boyd R: Fitness and physical activity in children and youth with disabilities. *Int J Pediatr* 2012, 2012:162648. doi:10.1155/2012/162648. Epub 2012 Dec 30.

- Norrby U, Nordholm L, Fasth A: Reliability and validity of the Swedish version of child health questionnaire. Scand J Rheumatol 2003, 32:101–107.
- Andersson-Gäre B, Ruperto N, Berg S, Hagelberg S, Jonsson NO, Magnusson B, Martinell J, Erling A, Landgraf JM, Peadiatric Rheumatology International Trials Organisation: The Swedish version of the child health assessment questionnaire (CHAQ) and the child health questionnaire (CHQ). *Clin Exp Rheumatol* 2001, 19(suppl 23):S146–S150.
- Landgraf JM, Maunsell E, Speechley KN, Bullinger M, Campbell S, Abetz L, Ware JE: Canadian-French, German and UK versions of Child Health Questionnaire: methodology and preliminary items scaling results. *Qual Life Res* 1998, 7:433–445.
- Carlsson M, Olsson I, Hagberg G, Beckung E: Behaviour in children with cerebral palsy with and without epilepsy. *Dev Med Child Neurol* 2008, 50:784–789.
- Beckung E, White-Koning M, Sarcelli M, McManus V, Michelsen S, Parkes J, Parkinson K, Thyen U, Arnaud C, Fauconnier J, Colver A: Health status of children with cerebral palsy living in Europe: a multi-centre study. *Child Care Health Dev* 2008, 34:806–814.
- Bartlett D, Birmingham T: Validity and reliability of a pediatric reach test. Pediatr Phys Ther 2003, 15:84–92.
- Eek MN, Kroksmark AK, Beckung E: Isometric muscle torque in children 5 to 15 years of age: normative data. Arch Phys Med Rehabil 2006, 87:1091–1099.
- 35. Häger-Ross C, Rösblad B: Norms for grip strength in children aged 4–16 years. Acta Paediatr 2002, 91:617–625.
- Keen E, Sloan AW: Observation on the Harvard step test. J Appl Physiol 1958, 13:241–243.
- Borg GA: Phychophysical bases of perceived exertion. Med Sci Sports Exerc 1982, 14:377–381.
- Norrby U, Nordholm E, Andersson-Gäre B, Fasth A: Health-related quality of life in children diagnosed with asthma, diabetes, juvenile chronic arthritis or short stature. Acta Paediatr 2006, 95:450–456.
- Lindehammar H: Hand strength in juvenile chronic arthritis: a two-year follow-up. Acta Paediatr 2003, 92:1291–1296.
- Haberfehlner H, Visser B, Daffertshofer A, van Rossum MA, Roorda LD, van der Leeden M: Handwriting difficulties in juvenile idiopathic arthritis: a pilot study. *Clin Exp Rheumatol* 2011, 29:887–893.
- Henderson CJ, Lovell DJ, Specker BL, Campaigne BN: Physical activity in children with juvenile rheumatoid arthritis: quantification and evaluation. Arthritis Care Res 1995, 8:114–119.
- Takken T, Van Brussel M, Engelbert RH, Van Der Net J, Kuis W, Helders PJ: Exercise therapy in juvenile idiopathic arthritis: a Cochrane Review. Eur J Phys Rehabil Med 2008, 44:287–297.
- 43. Gutiérrez-Suárez R, Pistorio A, Cespedes Cruz A, Norambuena X, Flato B, Rumba I, Harjacek M, Nielsen S, Susic G, Mihaylova D, Huemer C, Melo-Gomes J, Andersson-Gare B, Balogh Z, De Cunto C, Vesely R, Pagava K, Romicka AM, Burgos-Vargas R, Martini A, Ruperto N, Pediatric Rheumatology International Trials Organisation (PRINTO): Health-related quality of life of patients with juvenile idiopathic arthritis coming from 3 different geographic areas. The PRINTO multinational quality of life cohort study. *Rheumatology (Oxford)* 2007, 46:314–320.
- Jürimäe T, Hurbo T, Jürimäe J: Relationship between legs bone mineral density, anthropometry and jumping height in prepuberal children. *Coll Antropol* 2008, 32:61–66.
- Gualano B, Sa Pinto AL, Perondi B, Leite Prado DM, Omori C, Almeida RT: Evidence for prescribing exercise as treatment in pediatric rheumatic diseases. Autoimmun Rev 2010, 9:569–573.
- Philpott JF, Houghton K, Luke A: Physical activity recommendations for children with specific chronic health conditions: Juvenile idiopathic arthritis, hemophilia, asthma and cystic fibrosis. *Clin J Sport Med* 2010, 20:167–172.
- Dempster H, Porepa M, Young N, Feldman MB: The clinical meaning of functional outcome scores in children with juvenile arthritis. Arthritis Rheum 2001, 44:1768–1774.

doi:10.1186/1546-0096-11-7

Cite this article as: Sandstedt *et al.*: Muscle strength, physical fitness and well-being in children and adolescents with juvenile idiopathic arthritis and the effect of an exercise programme: a randomized controlled trial. *Pediatric Rheumatology* 2013 11:7.

Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

) BioMed Central

Submit your manuscript at www.biomedcentral.com/submit