

Research

Body composition in children with juvenile idiopathic arthritis: effect of dietary intake of macronutrient: results from a cross sectional study

Asmae Hari^{1,8}, Samira Rostom¹, Asmae Hassani¹, Dalal El Badri¹, Ilham Bouaadi¹, Amina Barakat², Bouchra Chkirat², Khalid Elkari³, Bouchra Amine¹, Najia Hajjaj-Hassouni¹

¹Department of Rheumatology, El Ayachi Hospital, University Hospital of Rabat-Sale, Sale, Morocco, ²Department of Pediatrics, Hospital of Children, University Hospital of Rabat-Sale, Rabat, Morocco, ³Department of Nutrition, University Ibn Tofaïl, Faculty of Science of Kenitra, Kenitra, Morocco

Corresponding author: Asmae Hari, Department of Rheumatology, El Ayachi Hospital, University Hospital of Rabat-Sale, Sale, Morocco

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Abstract

Introduction: The aim of this study was to evaluate the relationship between macronutrient intake, body composition (lean body mass and fat mass) and bone mineral content in Moroccan children with juvenile idiopathic arthritis (JIA). **Methods:** A cross-sectional study, conducted between May 2010 and June 2011, covering out patient with JIA. The characteristics of patients were collected. The nutritional status was assessed by a food questionnaire including data of food intake during 7 consecutive days using 24-hour dietary recall. Food intake was quantified using the software Bilnut (Bilnut version 2.01, 1991). Dietary intake of macronutrients was expressed as percentage contribution to total energy. Body composition was evaluated with DXA total-body measurements (bone mineral content BMC expressed in g, lean body mass LBM and fat mass FM expressed in kg). **Results:** 33 patients were included. The mean age was 10.4 ± 4.3 years. The median disease duration was 2 (1-4.5) years. The median of LBM, FM and BMC were 19 kg (13.82-33.14), 5 kg (3.38-9.14) and 1044.90 g (630.40-1808.90) respectively. We found a positive correlation between LBM and dietary intake of carbohydrate (r = 0.4; p = 0.03). There were no significant association between LBM and intake of lipids, or protein. Moreover, no association was found between FM, BMC and intake of carbohydrates, lipids and proteins. **Conclusion:** This study suggests that there is a positive correlation between carbohydrates intake and LBM; however, dietary intake does not influence FM and BMC. Prospective studies with larger numbers of patients appear to be needed to confirm our findings.

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Introduction

In patients with juvenile idiopathic arthritis (JIA), growth impairment and altered body composition, including disturbed skeletal development, are well-known long-term complications [1]. Although, few studies have specifically addressed body composition in children with JIA.

Reduced bone mineral density (BMD) is also well recognized features [2]. Deficits of muscle mass have been described as a central factor in (secondary) bone loss [3]. The reasons for deficits in lean body mass are thought to be the disease activity itself, medication (corticosteroids), reduced physical activity, and malnutrition [4]. Apart from low BMD and lean mass, a higher fat mass was reported in children with rheumatic diseases [5].

Despite the considerable hereditary influence on body composition, the environmental factors, namely nutrition, play an important role as well [6]. Among the nutritional factors, macronutrients (proteins, carbohydrates and lipids), and their effects on bones receives most attention [7]. Inadequate dietary intake in childhood and adolescence might alter body composition with adverse implications later in life (obesity and osteoporosis). Therefore, understanding the effects of diet on body composition can guide the implementation of social policies and interventions that can help youths to develop healthy bodies.

In the literature, there was a very few studies assessing the relationship between dietary intake and body composition in children with JIA. In addition, there are no studies in a Moroccan population that evaluate the same subject. An abnormal body composition with low muscle mass and increased fat mass in children with JIA [4,5] prompted us to conduct this study which aims to evaluate the relationship between dietary intake of macronutrients, body composition (lean body mass and fat mass) and bone mineral content in Moroccan children with JIA.

Methods

Data collection

A cross sectional study of children with JIA was conducted between May 2010 and June 2011 at the department of rheumatology of El

Ayachi university hospital and department of pediatrics of university hospital of children of Rabat-Sale. Informed consent was obtained by parents from all subjects and the study was approved by ethics committee of our university hospital.

The diagnosis of JIA was based on the criteria of the International League of Association for Rheumatology (ILAR) [8]. Patients were recruited in consultation or during hospitalization. We excluded patients with any other chronic disease (endocrinal, neurological, cardiac, and renal) that affect bone metabolism. The disease and patients characteristics considered as explanatory measures were: age (year), gender, diagnosis (JIA subtype), disease duration (years). Disease activity was assessed using a visual analogical scale (VAS), number of tender joints, number of swollen joints, erythrocyte sedimentation rate (ESR), Disease activity score (DAS 28) for polyarticular and oligoarticular JIA [9]. The Maastricht AS Enthesitis Score and Bath AS Disease Activity Index (BASDAI) were used for juvenile spondylarthropathy [10]. Functional disability was determined by using the Moroccan version of Childhood Health Assessment Questionnaire (CHAQ) [11]. Treatment with NSAIDs, steroids and disease modifying anti-rheumatic drugs (DMARDs) was determined.

Body composition

Body composition was evaluated with DXA total-body measurements (whole body bone mineral content BMC expressed in g, lean body mass LBM and fat mass FM expressed in kg) using the same DXA instrument (Lunar Prodigy; GE Lunar, Madison, WI). According to international consensus, DXA measurements without head were used. The instrument automatically alters scan depth depending on the thickness of the subject, as estimated from age, height, and weight. All scans were performed while the subjects were wearing light indoor clothing and no removable metal objects.

Medical history for bone fractures was negative in all patients. All subjects underwent plain Vertebral Fracture Assessment (VFA) to exclude unknown vertebral fractures [12].

Dietary evaluation

Nutrient intake was determined using the 24 hour diet recall during 7 consecutive days [13]. The food questionnaire identified all foods consumed during the day previous to the interview. Two nutritionists analyzed the food dietary to quantify the food

consumed from the recorded information. Nutrient intake was analyzed by software bilnut (Bilnut version 2.01, 1991), validated and standardized. Dietary intake of macronutrients was expressed as percentage contribution to total energy. None of the children was taking vitamin or mineral supplements at the time of recruitment

Anthropometric measures

Weight (kg) and height (m) were measured according to the recommendation of the World Health Organization (WHO). The results of the BMI (Kg/m^2) were compared with reference values of Hammer and al [14].

Statistics

Analysis was carried out using the statistical package for the social sciences (SPSS) version 16.0. Data for patients were presented as mean ± standard deviation or median and quartile for continuous variables and as frequencies and percentage for categorical variables. The two-sample t-test was used for comparisons of scores within categorical variable subgroups. Pearson's linear correlation was used for the quantification between the observed numerical variables. Significance level was p value less than 0.05.

Results

Thirty three patients were included in this study. The mean age of our patients was 10.4 ± 4.35 . 54.5% of our patients were males. The median disease duration was equal to 2(1-4.5) years. Demographic and clinical characteristics of patients are shown in **Table 1**. The median of lean body mass (LBM), total body fat mass (FM) and bone mineral content (BMC) were 19 kg (13.82-33.14), 5 kg (3.38-9.14) and 1044.90 g (630.40-1808.90) respectively (**Table 1**).

The dietary intake data are shown in **Table 2**. Mean values for diet composition indicated that carbohydrates provided 53% of energy, lipids provided 32.5%, and proteins provided 16.1%.

We found a positive correlation between LBM and carbohydrates intake (r=0.4; p=0.03). There were no significant association between LBM and intake of lipids or proteins. Also, no association was found between FM, BMC and dietary intake of carbohydrates, lipids and proteins (**Table 3**).

Discussion

Our data show that there is a positive correlation between LBM and carbohydrates ingested (expressed as a proportion of energy intake) but not between LBM, lipids and proteins intake. In the study of Volek JS et al, they found that a carbohydrate-restricted diet resulted in a significant reduction in fat mass and increase in lean body mass, which may be partially mediated by the reduction in circulating insulin concentrations [15]. Chandler-Laney PC et al showed that maternal glucose concentration during pregnancy was positively associated with children's lean mass and fat mass [16]. In the other study, Lean muscle mass was not associated with physical activity or dietary intakes [17].

Previous observational studies of dietary protein intake and body composition have shown mixed results. Protein intake was not associated with LM in cross-sectional studies [17, 18]. In the Health ABC Study cohort, they found an association between protein intake and changes in LM over 3 years of follow-up [19]. However, few studies have examined the effect of protein intake on LBM in children.

The problem of overweight and obesity is a growing public health concern affecting numerous countries. The amount of fat ingested has been implicated as a causal or facilitating factor in the deposition of body fat. Thus, it is proposed that dietary fat can directly or indirectly manipulate human adipose tissue. There is suggestion in the literature that protein intake, not fat intake, may be associated with the development of adiposity in childhood [20]. It has been proposed that a high protein intake during early childhood stimulates insulin-like growth factor I production, thereby triggering precocious adipocyte multiplication [21]. In a prospective case-control study, they found that fat intake predicted gain in percentage of body fat in both adolescent girls with type 1 diabetes and healthy control girls [22]. In the other study, they did not find any association between dietary protein intake and percentage body fat [23]. Dietary carbohydrate expressed as a percentage of energy intake is often inversely related to body fat, including in childhood obesity [24]. In this study of children with JIA, there was no association between FM and intake of macronutrients.

Despite the methodological limitations of the present study; sample size, cross-sectional and non-controlled design, this study was the first Moroccan study that assesses the relationship between dietary intake and body composition in children with JIA, but the causeeffect relationship remains to be determined.

Conclusion

This study suggests that there is a positive correlation between carbohydrates intake and LBM; however, dietary intake does not influence FM and BMC. Furthermore, Controlled feeding studies will be required to clarify possible roles of dietary intake in body composition of children with JIA.

Competing interests

The authors declare no competing interests.

Authors' contributions

SR, BA and NHH conceived the study and supervised its design, execution, and analysis and participated in the drafting and critical review of the manuscript. AH and SR did data management and statistical analyses. AB and KEK analyzed the food dietary. All authors participated in critical revision of the manuscript. AH wrote the paper with input from all investigators. All authors read and approved the final manuscript.

Tables

Table 1: demographic and clinical characteristics of patients

 Table 2: energy, macronutrient intakes

Table 3: pearson's correlation coefficients between dietary intake and body composition variables

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Table 1: demographic and clinical characteristics of patients			
Characteristics of patients	n=33		
Age (years), mean ± DS	10.4 ± 4.3		
Female sex, n (%)	15(45.5)		
Duration of disease (year), mediane (IQ)	2(1-4.5)		
BMI, n (%)			
Obesity	5(15.2)		
Normal	19(57.6)		
Underweight	9(27.3)		
Subtype of JIA, n (%)			
Systemic	8(24.2)		
Oligoarticular	9(27.3)		
Polyarticular	16(48.5)		
Medications used, n (%)			
NSAID	26 (79)		
Corticosteroids	19(58)		
DMARDs	17(51.5)		
LBM (kg), mediane (IQ)	19 (13.82-33.14)		
FM (kg), mediane (IQ)	5 (3.38-9.14)		
BMC (g), mediane (IQ)	1044.90 (630.40-1808.90)		

BMI: body mass index; JIA: juvenile idiopathic arthritis; NSAIDs: non-steroidal antiinflammatory drugs; DMARDs: disease modifying anti-rheumatic drugs; LBM: lean body mass index; FM: fat mass, BMC: bone mineral content

Table 2: energy, macronutrient intakes							
	Total sample (33)	Boys (18)	Girls (15)	p			
Carbohydrates (% of energy)	53±17.3	53.18±14.7	52.5±20	0.9			
Lipids (% of energy)	32,5±13.1	30.8±13.4	34.6±12,8	0,4			
Protein (% of energy)	16,08±4.3	16±3.8	16.2±5	0.9			

Results are presented in mean±SD; Differences were tested using t test

Table 3: pearson's correlation coefficients between dietary intake and body composition variables

	LBM	FM	ВМС			
Carbohydrates (%	0.4+	0.3	0.3			
of energy)	0.4	0.5	0.5			
Lipids (% of energy)	-0.2	-0.09	0.2			
	-0.2	-0.09	-0.2			
Proteins (% of	0.1	0.04	0.1			
energy)	-0.1	-0.04	-0.1			
		I	I			

*p<0.05, LBM: lean body mass, FM: fat mass, BMC: bone mineral content