

High rates of obesity and greater associated disability among people with rheumatoid arthritis in Canada

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Abstract Obesity in rheumatoid arthritis has been associated with increased risk of comorbidities, larger medical costs, decreased quality of life, higher disease activity, and reduced therapeutic responses. We assessed the burden of obesity among rheumatoid arthritis patients and its impact on patient-reported outcomes. Patients receiving care at two Canadian University Centers were included. Height and weight were measured and selected sociodemographic and rheumatoid arthritis (RA) characteristics as well as patient-reported outcomes were obtained. Patients were classified according to WHO criteria and proposed RA cut points, and results were compared with national data. Using WHO criteria, 68 (34 %) RA patients were classified as obese (vs. ~25 % of Canadians). Using RA cut points, 112 (55 %) RA patients were classified as obese. With both classification methods, obese individuals had significantly higher mean HAQ scores and a higher odds of significant disability (HAQ \geq 1: WHO OR 2.3;

95 % CI 1.2, 4.2 and RA-specific OR 1.8; 95 % CI 1.0, 3.2). Independent of the classification method use, RA patients have significantly higher rates of obesity than national prevalence estimates. Obese RA patients had about twice the odds of reporting moderate to severe disability.

Keywords Cardiovascular disease risk · Disability · Obesity · Rheumatoid arthritis

Introduction

Rheumatoid arthritis (RA) patients have a reduced life expectancy compared with the general population, mainly due to an increased prevalence of, and worse outcomes from, cardiovascular (CV) disease (CVD) [1]. Genetic predisposition, traditional CV risk (CVR) factors, and the effects of systemic inflammation on the vasculature contribute to the excess CV morbidity and mortality in RA [2, 3]. Among traditional CVR factors, obesity is associated with increased risk of comorbidities, the need for total joint replacement, increased medical costs, and decreased quality of life [4]. Excess weight is also associated with higher RA disease activity and reduced therapeutic benefits of disease-modifying anti-rheumatic drugs [5].

Body mass index (BMI) provides a relative estimate of body fatness, and excess fat is a potent predictor of CVD and overall mortality [6]. The World Health Organization (WHO) defines overweight as BMI 25–29.9 kg/m² and obesity as BMI \geq 30 kg/m² [7]. In RA, WHO cut points appear to underestimate body fat, and lower sex-specific thresholds to define overweight and obesity based on DEXA-derived estimates of fatness have been proposed

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[8, 9]. The goal of this study was to estimate the burden of obesity in RA patients using both classification systems.

Materials and methods

Consecutive RA patients ≥ 18 years of age treated at arthritis clinics of McGill University ($n=94$) and the University of Manitoba ($n=106$) between 2012 and 2013 were included. All met the 2010 ACR/EULAR classification criteria for RA. This study was approved by the ethics committees at each center.

Demographics, smoking status, and RA characteristics (disease duration, RF and anti-CCP antibodies, ESR, CRP, tender and swollen joint counts) were abstracted from medical records. BMI and patient-reported outcomes (PROs) were obtained at the time of evaluation.

Patients were classified using WHO and RA sex-specific (BMI ≥ 24.7 kg/m² in men; ≥ 26.1 kg/m² in women) BMI categories [9]. We used 2008 Statistics Canada data to compare the obesity rates with the general Canadian population. Obese vs. non-obese groups were compared with *t* tests and chi-square for continuous and categorical variables. Statistical analyses were performed with SPSS V21 and $p < 0.05$ was considered as statistically significant.

Results

Data from 94 patients from Montreal and 106 from Winnipeg were included. Demographics and RA characteristics are presented in Table 1.

Prevalence of obesity

Using WHO criteria, 68 (34 %) RA patients were classified as obese (Table 2). Using RA thresholds, 112 (55 %) were classified as obese. With both classification methods, sociodemographic characteristics were similar between obese and non-obese groups, and the proportion of people classified as obese was much higher than in the adult Canadian population (25 %) [10]. Using WHO criteria, the risk ratio (RR) for obesity was 1.4 (95 % confidence interval [CI] 0.9, 2.2); more women than men were classified as obese (36 vs. 29 %, respectively; Fig. 1). Using RA thresholds, 55 % were classified as obese (RR 2.3; 95 % CI 1.5, 3.4); however, more men than women were obese (67 vs. 53 %, respectively).

Table 1 Participant characteristics by site

	N	Winnipeg	N	Montreal
Sociodemographic characteristics				
Age (years)	106	57.5 (13.6)	94	56.3 (17.2)
Female sex; <i>n</i> (%)	106	78 (73.6)	94	74 (78.7)
BMI (kg/m ²)	106	28.1 (6.0)	94	28.1 (7.3)
WHO BMI categories, <i>n</i> (%)				
Normal (<25)		35 (33.0)		35 (37.2)
Overweight (25–29.9)		36 (34.0)		26 (27.7)
Obese (30–39.9)		30 (28.3)		28 (29.8)
Severe obesity (40+)		5 (4.7)		5 (5.3)
Ever smoked* <i>n</i> (%)	102	67 (65.7)	70	22 (31.4)
RA characteristics				
RA duration (years)	103	6.9 (4.3)	94	5.8 (7.6)
RA <1 year; <i>n</i> (%)	103	8 (7.8)	94	10 (10.6)
DAS 28-CRP*	106	2.6 (1.4)	67	3.4 (1.5)
Remission; <i>n</i> (%)				
Low; <i>n</i> (%)		64 (65)		18 (27)
Moderate; <i>n</i> (%)		17 (17)		4 (6)
High; <i>n</i> (%)		13 (13)		39 (58)
Tender joint count*	106	3.0 (6.4)	91	5.4 (6.0)
Swollen joint count*	106	1.6 (4.0)	83	2.9 (4.9)
CRP (mg/dL); mdn (IQR)*	99	3.8 (6.8)	87	2.7 (5.0)
ESR (mm/h); mdn (IQR)*	99	7.0 (16.0)	81	20.0 (22.0)
Anti-CCP+; <i>n</i> (%)	106	71 (67.0)	86	49 (57.0)
RF+; <i>n</i> (%)	99	86 (86.9)	90	71 (78.9)
Patient-reported outcomes				
AM stiffness (min)	105	42.5 (61.8)	89	30.7 (49.5)
Pain* (100 mm VAS)	99	27.5 (26.8)	50	42.4 (29.4)
Patient global* (100 mm VAS)	99	25.0 (25.8)	81	37.3 (27.6)
HAQ ^a			92	0.8 (8)

Values are the mean (SD) unless otherwise indicated

* $p < .04$, significantly different

^aHealth Assessment Questionnaire

Disease activity

Using WHO criteria, ESR and CRP levels were significantly higher in obese individuals; and while DAS28 and tender joint counts also were higher, differences were not statistically significant. Using RA thresholds, disease activity indicators were similar between groups.

Patient-reported outcomes

With both classification methods, obese individuals had significantly higher mean HAQ scores (Table 2) and a higher odds of significant disability (i.e., HAQ ≥ 1 : WHO OR 2.3; 95 % CI 1.2, 4.2 and RA-specific OR 1.8; 95 % CI

Table 2 Characteristics by obesity status using WHO and proposed RA criteria

	WHO criteria		Proposed RA criteria	
	Not obese (N=132)	Obese (N=68)	Not obese (N=88)	Obese (N=112)
Sociodemographic characteristics				
Age (years)	56.7 (15.4)	57.3 (15.3)	55.6 (16.3)	58.0 (14.5)
Female sex; n (%)	98 (74.2)	54 (79.4)	72 (81.8)	80 (71.4)
Ever smoked; n (%)	61 (53.5)	28 (48.3)	50 (51.9)	49 (51.6)
RA characteristics				
RA duration (years)	6.8 (6.7)	4.7 (5.5)	7.1 (7.6)	5.9 (4.6)
Early RA (<1 year); n (%)	10 (7.6)	8 (12.1)	7 (8.0)	11 (10.0)
DAS28-CRP	2.8 (1.5)	3.1 (1.6)	2.9 (1.6)	2.8 (1.3)
Tender joint count	3.8 (6.7)	4.7 (5.5)	4.5 (7.6)	3.8 (5.1)
Swollen joint count	2.2 (4.4)	2.1 (4.6)	2.7 (4.9)	1.8 (4.0)
CRP (mg/dL); mdn (IQR)	2.5 (4.0)	5.4 (9.8)*	2.4 (4.3)	4.2 (8.4)
ESR (mm/h); mdn (IQR)	10.0 (19.0)	20.0 (28.0)*	12.0 (20.0)	14.0 (22.0)
CRP (mg/dL); mdn (IQR)	2.5 (4.0)	5.4 (9.8)*	2.4 (4.3)	4.2 (8.4)
Patient-reported outcomes				
AM stiffness (min)	34.2 (53.7)	42.7 (62.2)	35.2 (57.4)	38.6 (56.3)
Pain (100 mm VAS)	30.5 (28.4)	36.7 (28.1)	30.3 (29.4)	34.1 (27.5)
Patient global (100 mm VAS)	29.3 (27.4)	32.3 (26.9)	28.5 (27.3)	31.8 (27.2)
HAQ ^a	0.6 (0.7)	1.2 (0.8)*	0.7 (0.7)	0.9 (0.8)**

Values are the mean (SD) unless otherwise indicated. Numbers in italics are the groups that are being compared

*Groups significantly ($p < 0.05$) different ** $p = 0.08$

^a Health Assessment Questionnaire: $N = 60$ for not obese and $N = 32$ for obese

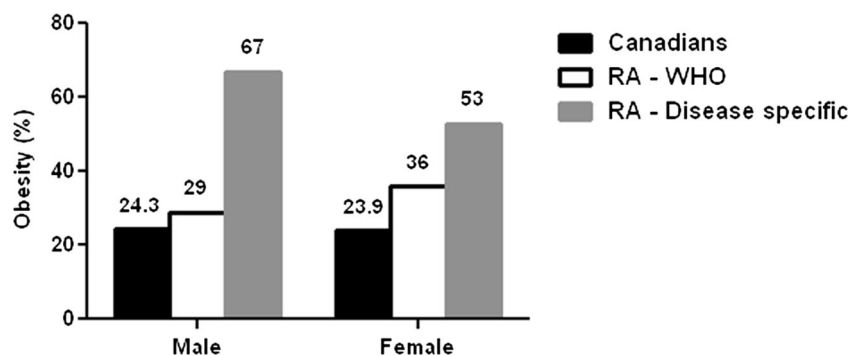
1.0, 3.2). Although morning stiffness, pain, and patient global scores were higher in obese patients with both classification methods, differences were not statistically significant.

Discussion

Our results contribute to the growing body of evidence suggesting rates of obesity are higher in people with RA [3, 11]. Similar to others, one third of our sample (vs. ~25 % of Canadians) were obese using WHO criteria [3, 8, 9]. Using

RA-specific criteria, two thirds of men and more than half of women in our sample were obese. With either weight classification system, obesity was associated with greater disability, and obese RA patients had about twice the odds of reporting moderate to severe disability (i.e., $HAQ \geq 1$). A 2015 meta-analysis of BMI, disease activity, and selected RA outcomes, and a study of inflammatory polyarthritis, also reported greater disability among obese patients [4, 12]. Similarly, the increased disability we observed could be due to the higher prevalence of obesity in our sample. The complex interactions between patients’ health-related thoughts about arthritis and weight have been recently

Fig. 1 Prevalence of obesity by sex using World Health Organization and proposed RA-specific cut points



explored in a study by Sommers et al. showing that RA patients with higher levels of pain catastrophizing (i.e., tendency to focus on and magnify pain sensations and to feel helpless in the face of pain) and lower levels of confidence for managing (i.e., self-efficacy) both arthritis and weight were more likely to report higher levels of pain, poorer physical function, and more overeating [13]. Inflammatory indicators also were higher in obese persons using WHO (but not RA-specific) cut points. Several, but not all studies, have reported higher disease activity and greater pain and patient global scores in RA obese individuals [4, 14, 15]. Although DAS and tender (but not swollen) joint counts and PROs were higher in our sample, differences were not statistically significant in part due to smaller sample sizes.

Strengths of our study include the diverse sample of people with a wide range of BMIs (17–59). Participants were similar in age, pain, and disability to the sample on which the new RA cut points were derived, although our sample had more females, with shorter RA duration, and lower CRP. Our study has several limitations including the cross-sectional nature of the described associations and the lack of assessment of the impact of steroids, and other obesogenic medications. We did not specifically assess abdominal obesity, which is more strongly associated with negative health outcomes [16].

Given the deleterious effects of excess weight on health and quality of life [17], our results emphasize the importance of routinely identifying and addressing obesity in RA patients, and preventing weight gain in overweight individuals. It remains unclear whether weight loss can attenuate the excess CVD morbidity and mortality associated with RA, optimize response to treatment, and enhance physical function.

In summary, in a sample of Canadian RA patients, we found significantly higher rates of obesity and disparities in the prevalence of obesity by sex. Using traditional BMI cut points, one third of the RA patients were obese, and they had significantly higher levels of inflammatory markers and greater levels of disability. Using recently proposed RA cut points, two thirds of men and more than half of women with RA would be classified as obese. These results underscore the need to consider weight as one component of optimal disease management.

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Compliance with ethical standards

Disclosures None.

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