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Dietary inflammatory index and prevalence of overweight and obesity in Brazilian graduates from the Cohort of Universities of Minas Gerais (CUME project)



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ABSTRACT

Objectives: The aim of this study was to evaluate the independent association of the dietary inflammatory index (DII[®]) score with overweight and obesity in Brazilian participants of the Cohort of Universities of Minas Gerais (CUME project).

Methods: This was a cross-sectional study consisting of 3,151 graduates and postgraduates (2197 women) with a mean (SD) age of 36.3 y (\pm 9.4 y). Sociodemographic characteristics, lifestyle, and anthropometric data were assessed via online self-reported questionnaire. Additionally, a validated food frequency questionnaire with 144 food items was used to generate energy-adjusted DII (E-DIITM) scores, which evaluated the inflammatory potential of the diet.

Results: The prevalence of overweight and obesity were 28.2% and 11%, respectively. Participants in the highest E-DII quartile (most proinflammatory diet) were more likely to be smokers/former smokers; sedentary; and consumers of red and ultra-processed meats, fats and oils (excluding olive oil), bottled fruit juices and soft drinks, sugars, sweets, and higher overall caloric intake, compared with the first quartile of E-DII. Both men and women in the fourth E-DII quartile had the highest prevalence of overweight and obesity (prevalence ratio [PR], 1.35; 95% confidence interval [CI], 1.14–1.59 and PR, 1.97; 95% CI, 1.20–3.22, respectively, in men; PR, 1.38; 95% CI, 1.17 to 1.65 and PR, 1.95; 95% CI, 1.31–2.90, respectively, in women).

Conclusion: The most proinflammatory dietary pattern was associated with a higher prevalence of overweight and obesity and other unhealthy lifestyles including being sedentary, smoking, and consuming a obesogenic diet.

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Introduction

Obesity, defined as excessive accumulation of body fat, is a chronic multifactorial disease considered a public health problem in Brazil [1] and worldwide [2]. In Brazil, the percentage of overweight individuals increased from 42.6% in 2006 to 53.8% in 2016, and obesity rose from 11.8% to 18.9% in the same time [1]. By 2030, the number of overweight or obese people worldwide could reach 3.3 billion [3].

In this sense, low-grade inflammation is a recognized link between obesity and other chronic diseases (e.g., diabetes, metabolic syndrome, and cardiovascular diseases) [4–6]. At the same time, healthy dietary patterns and specific dietary components (e.g., folate, unsaturated fatty acids, and polyphenols) have been associated with inflammation [7–9] as well as with comorbidity-related obesity [10–13].

Additionally, global dietary indexes allow a more comprehensive evaluation of dietary patterns. Among them, the dietary inflammatory index (DII[®]) has emerged as a new tool to assess the inflammatory potential of the diet and its relationship with cardio-metabolic risk, such as excess body weight, in different populations [14]. In fact, most proinflammatory diets (higher DII values) have been associated with higher occurrences of metabolic disorders, cardiovascular diseases, and overall and cause-specific mortality [15–18]. The relationship between DII and chronic disease outcomes has not been reported in the Brazilian population in general, but was documented in specific groups, such as adolescents [19], young adults [20], or individuals undergoing bariatric surgery [21].

In this cross-sectional study, we aimed to evaluate the independent association of DII scores with overweight and obesity at baseline in participants of the Cohort of Universities of Minas Gerais (CUME project) [22]. Our hypothesis is that higher DII score (more proinflammatory diet) would be associated with a higher prevalence of overweight or obesity, after accounting for several confounders.

Materials, participants, and methods

CUME project

The CUME project is a concurrent open cohort whose objective is to evaluate the effects of the Brazilian dietary pattern and the nutritional transition on non-communicable diseases (NCD) in adults ≥ 18 y of age, who are graduates and post-graduates at the Universidade Federal de Viçosa (UFV) or the Universidade Federal de Minas Gerais (UFMG), institutions located in the state of Minas Gerais, Brazil [22].

The cohort methodology has been previously described [22]. Briefly, potential volunteers were invited by e-mail and directed to the CUME's virtual page. A self-reported, online questionnaire consisted of two parts and were sent separately within 1 wk of each other. The first stage consisted of questions related to sociodemographic, anthropometric, lifestyle, and health-related data. In the second stage, participants completed the food frequency questionnaire (FFQ) and additional questions related to dietary practices and consumption of specialty products [22].

The CUME project was designed in accordance with the experience of the team as researchers of the study Seguimiento University of Navarra (SUN project), which was developed in Spain, and whose objective is to analyze the influence of the Mediterranean diet on health outcomes [23].

Study population

This was a cross-sectional analysis of baseline data from the CUME project [22]. Of the 4986 volunteers who answered the baseline CUME questionnaire from March to August 2016, 3151 were included in the present study. Inclusion criteria were residency in Brazil over the previous year and completion of the baseline questionnaire, including the FFQ.

The study was approved by the Human Research Ethics Committees of UFMG and UFV. All participants read the informed consent form and indicated online agreement (with an online command) before responding to the questionnaire [22].

Food consumption and DII computation

In order to evaluate food consumption and calculate the energy-adjusted DII (E-DIITM) score, a 144-item FFQ was applied, which was previously validated for the Brazilian population [24]. Each participant reported the frequency of consumption of a food (daily, weekly, monthly, or annually), the number of times it was consumed (0 to ≥ 9 times) and the portion size appropriate to each food. Moreover, they answered questions related to daily dietary practices, such as number of meals per day; intake of visible fat of meats; and addition of salt and sugar in ready-made meals. To enhance self-completion of the FFQ, the team developed a photographic album with 96 images of foods and serving utensils to facilitate the visualization of portions of food to enhance a reliable response regarding consumption.

Nutrient intakes were calculated using daily intake of each food item and its nutrient composition, according to the Brazilian Food Composition Table [25]. Where data were lacking in the Brazilian tool, the US Department of Agriculture table [26] was used. The Brazilian Carotenoid Composition Table in Foods [27] and the Phenol-Explorer database [28] were used for estimation of β -carotene and flavonoids, respectively.

DII scores were calculated using a scoring algorithm based on a review of 1943 articles linking 45 food parameters and six inflammatory biomarkers (interleukin [IL]-1 β , IL-4, IL-6, IL-10, tumor necrosis factor [TNF]- α , and C-reactive protein [CRP]). Then, food parameters were assigned to a positive score (+1) if the effect was proinflammatory or a negative score (–1) if the effect was anti-inflammatory. A score of zero was assigned if the parameter did not generate significant changes in the inflammatory biomarkers, as previously detailed [14]. DII density scores, which adjust for energy intake, also were calculated; before standardizing to ation with the energy-adjusted global reference database, the parameters were each converted to 1000 kcal of energy intake, and energy intake as food parameters was excluded from the actual E-DII calculation. These scores were based on a similar algorithm that used an energy-adjusted global database [29,30]. Positive DII and E-DII scores represent food patterns with the most proinflammatory potential, whereas negative DII and E-DII values represent food patterns with the most anti-inflammatory potential. The DII has been construct validated with several inflammatory markers in various studies [14,31].

The E-DII scores in the present study were based on 35 food parameters available from the FFQ: energy intake (used for adjustment); carbohydrate; protein; total fat; cholesterol; saturated fat; trans fat; monounsaturated and polyunsaturated fat; ω -3 and ω -6; fiber; alcohol; niacin; thiamine; riboflavin; vitamins B₁₂ and B₆; folic acid; vitamins A, C, D, and E; iron; selenium; magnesium; zinc; caffeine; β -carotene; flavonol-3-ol; flavones, flavonols, and flavonoids; anthocyanidins; and isoflavones.

Determination of overweight and obesity

The participants reported their current weight and height by completing the online questionnaire. Body mass index (BMI; kg/m²) was calculated as body weight (in kg) divided by height (in m) squared. The weight status of the adults was classified according to the criteria defined by the World Health Organization [32] as being normal weight (BMI 18.5 to <25 kg/m²), overweight (BMI 25 to <30 kg/m²), or obese (BMI ≥ 30 kg/m²). For participants ≥ 60 y of age, weight status was classified according to the Pan American Health Organization [33]: normal weight (BMI 23 to <28 kg/m²), overweight (BMI 28 kg/m²), or obese (BMI ≥ 30 kg/m²). A validation study of a subsample of 172 CUME participants at baseline compared self-reported height and weight with measured height and weight and obtained adequate correlation coefficients (intraclass correlation coefficients [ICC], 0.989; 95% confidence interval [CI], 0.985–0.992 and ICC, 0.995; 95% CI, 0.993–0.996 for height and weight, respectively, and kappa coefficient of 0.882 between obesity diagnosis based on self-reported and measured data, indicates almost perfect agreement) [22,34].

Assessment of other variables

Information on the practice of physical activity (yes or no), smoking (never, former or current smoker), age (y), sex (male or female), and subject area of concentration (health sciences, humanities, physical sciences and agrarian sciences) also was collected.

Statistical analysis

Based on a previous study that investigated the relationship between DII and overweight/obesity cases [28], with a similar population to the CUME project, a minimal sample of 1426 participants was calculated for this study using a 95% CI (type 1 error rate of 0.05, two-tailed); 80% statistical power; ratio exposed/unexposed of 1; prevalence of overweight/obesity of 20% in individuals in the first quartile of the inflammatory diet index (most anti-inflammatory); and obesity prevalence ratio of 1.32.

The database was written in SPSS[®] version 20 (SPSS, Chicago, IL, USA). All analyses were performed with Stata[®] version 13 (StataCorp, College Station, TX, USA) and statistical significance was set at $\alpha = 5\%$ ($P < 0.05$). To control for the effect of

caloric intake on nutrients and food groups evaluated, models were adjusted by the residual nutrient method [35]. To evaluate the associations between E-DII score and nutrient intake, food groups, BMI, and prevalence of overweight and obesity, participants were categorized by E-DII quartile.

The comparisons of sociodemographic characteristics and BMI among E-DII quartiles were made with Pearson's χ^2 test for categorical variables and analysis of variance for continuous variables. When a significant difference was detected, the Bonferroni *post hoc* test was applied to correct multiple comparisons.

Multiple linear regression analyses were performed with adjustment for sex, age, BMI, smoking, and physical activity to compare the consumption of macro- and micro-nutrients, food groups, and feeding practices among the E-DII quartiles.

Finally, Poisson regression multivariate models, adjusted for age, smoking, physical activity, and graduation area, for male and female groups, and additionally adjusted for sex for total sample, were fitted to evaluate the association of a proinflammatory diet with overweight and obesity. Prevalence ratio (PR) and 95% CIs were estimated using the first E-DII (anti-inflammatory) quartile as the reference category.

Results

Of the 3151 participants in the CUME project, 30.3% were men and 69.7% women, with a mean age (SD) of 36.3 y (\pm 9.4 y), as only 3% of the population were >60 y of age. The E-DII score had a mean (SD) of -0.12 (1.63) and ranged from -5.48 to $+4.55$. The prevalence of overweight was 28.2% and obesity was 11%. Among overweight participants, 42.6% were men and 57.4% were women ($P < 0.001$).

The total energy intake of participants was ~ 3000 kcal/d across all E-DII quartiles ($P < 0.001$).

Table 1 shows the main sociodemographic characteristics of the participants according to E-DII quartile. The individuals included in the last quartile (most proinflammatory) were more likely to be former or current smokers, sedentary, and to consume more alcohol compared with those included in the first quartile (most anti-inflammatory). Regarding the graduation area, participants included in the first quartile were more likely to have concentrated in the health sciences.

Women were represented in higher proportions than men in all quartiles. Unlike men, however, their frequency decreased as E-DII

scores became most proinflammatory (Table 1). Furthermore, BMI values were significantly higher in the highest E-DII quartile, compared with the first quartile, for both men and women (Fig. 1).

In relation to food consumption according to E-DII scores, the intake of dairy, white and lean meats, fish or shellfish and eggs, whole grains, legumes, olive oil and oilseeds, fruits, vegetables, and natural fruit juices was significantly higher in the first quartile (most anti-inflammatory) and the consumption of red, fat, and ultra-processed meats, fats, and oils with the exception of olive oil, industrial juices, soft drinks, sugars and sweets was higher in the last quartile (most pro-inflammatory; Table 2).

Participants in the last quartile (most pro-inflammatory) were more likely to be overweight (Table 3) or obese (Table 4), regardless of confounding factors, including sex, age, smoking status, physical activity, and graduation area. When the sample was stratified by sex, the results were similar for both outcomes (Tables 3 and 4).

Discussion

In the present study, participants of the CUME project in the most proinflammatory E-DII quartile had higher BMI values and a higher prevalences of overweight and obesity than those in the lower (most anti-inflammatory) E-DII quartiles.

Results from the present study are in agreement with those obtained in studies conducted with other populations. In the Spanish SUN cohort, with 7027 individuals having a mean age of 37.4 y, the most proinflammatory DII was also associated with a higher risk for weight gain and a greater risk for overweight and obesity, independently of total caloric intake, physical activity, history of obesity, and baseline weight [31]. In the PREDIMED (PREvención con Dieta MEDiterránea) trial (also in Spain), with 7236 individuals with cardiometabolic risk, the authors also observed that participants with a most proinflammatory diet (higher DII scores) presented higher values of BMI independent of total caloric intake, age, diabetes, hypertension, smoking

Table 1
Sociodemographic characteristics according to E-DII of the participants of the CUME project (N = 3151), 2016

Characteristics	E-DII				P-value
	Q1 (-5.48 to -1.37) (most anti-inflammatory)	Q2 (-1.37 to -0.26)	Q3 (-0.26 to 1.08)	Q4 (1.08 to 4.55) (most proinflammatory)	
	n = 787	n = 788	n = 788	n = 788	
Sex, n (%)					
Male	205 (26.1)	215 (27.3)	247 (31.4)	287 (36.4)	<0.001
Female	582 (73.9)	573 (72.7)	541 (68.6)	501 (63.6)	
Age, y, mean (SD)	36.5 \pm 9.9	36.8 \pm 10	35.7 \pm 8.8	36.1 \pm 8.9	0.04
Marital status, n (%)					
Married or stable union	413 (52.7)	410 (52.6)	403 (51.8)	392 (50.5)	0.84
Single, separated/divorced, widower	370 (47.2)	371 (47.4)	377 (48.2)	390 (49.5)	
Smoking, n (%)					
Never smoked	660 (83.8)	626 (79.4)	610 (77.4)	603 (76.5)	0.001
Former and current smokers	127 (16.2)	162 (20.6)	178 (22.6)	185 (23.5)	
Alcoholic beverage consumption, n (%)					
No	253 (32.2)	229 (29.1)	172 (21.8)	202 (25.7)	<0.001
Yes	534 (67.8)	559 (70.9)	616 (78.2)	586 (74.3)	
Physical activity practice, n (%)					
Yes	642 (81.6)	625 (79.3)	570 (72.4)	538 (68.3)	<0.001
No	145 (18.4)	163 (20.7)	218 (27.6)	250 (31.7)	
Graduation area, n (%)					
Health sciences	305 (38.7)	297 (37.7)	267 (33.9)	255 (32.3)	0.02
Humanities	271 (34.4)	250 (31.7)	262 (33.2)	288 (36.5)	
Physical sciences	141 (17.9)	156 (19.8)	163 (20.7)	139 (17.6)	
Agrarian sciences	70 (8.9)	85 (10.8)	96 (12.2)	106 (13.4)	

CUME, Cohort of Universities of Minas Gerais; E-DII, energy-adjusted dietary inflammatory index. P-value of analysis of variance or χ^2 test, for continuous or categorical variables, respectively.

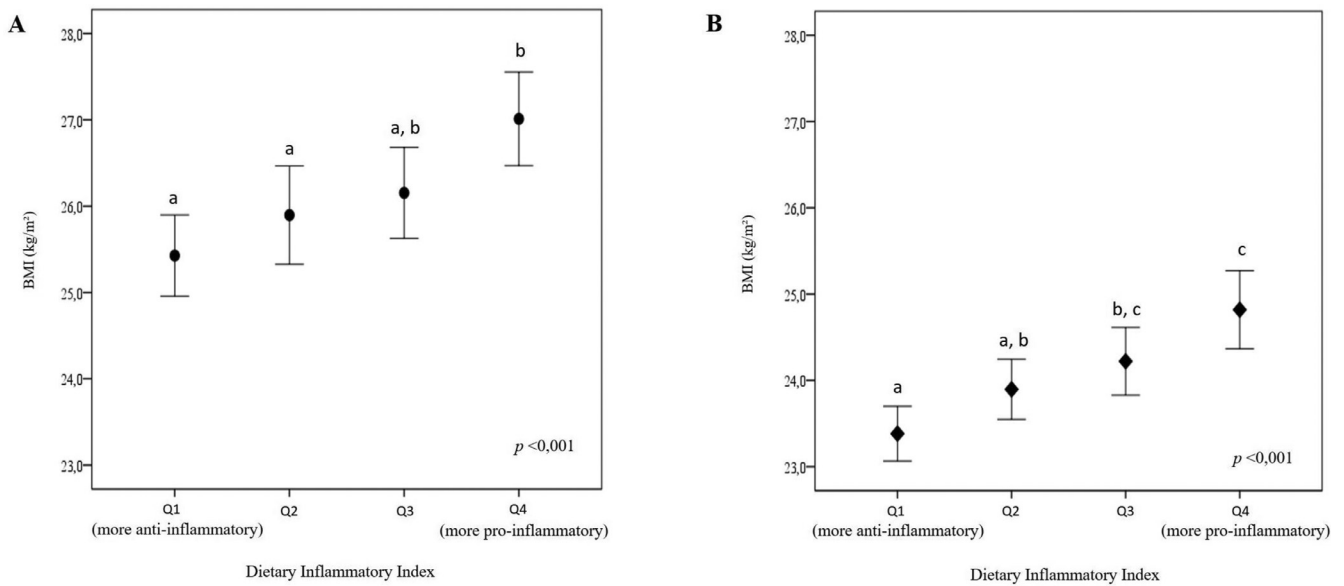


Fig. 1. Body mass index values according to E-DII for male (A) and female (B) participants in the CUME project (n = 3151), 2016.

P value < 0001 from ANOVA tests. Different letters (a, b and c) indicate mean statistical differences according to Bonferroni post hoc to correct multiple comparisons. Q, quartile.

status, physical activity, educational level, and marital status [36]. A positive correlation between BMI and DII values was reported by Ruiz-Canela et al. [37] in a review that included data from the following:

- PREDIMED;
- The SUN project;
- The Geelong Osteoporosis Study with 1363 Australian men ≥ 18 y of age;

Table 2

Daily food consumption according to E-DII of the participants of the CUME project (n = 3151), 2016

	Daily consumption E-DII				<i>P</i> _{trend} *
	Q1	Q2	Q3	Q4	
	(−5.48 to −1.37) (most anti-inflammatory) n = 787	(−1.37 to −0.26) n = 788	(−0.26 to 1.08) n = 788	(1.08 to 4.55) (most proinflammatory) n = 788	
Energy intake, kcal	2899 ± 1483	3072 ± 1735	3155 ± 1471	3330 ± 1573	<0.001
Carbohydrate, %EI	50.7 ± 7.8	50.1 ± 9.5	49.5 ± 9.3	49.9 ± 12.8	0.13
Protein, %EI	19 ± 4.5	18.8 ± 5.9	18.7 ± 4.6	18.3 ± 5.7	0.002
Lipid, %EI	30.2 ± 6.3	31 ± 7.3	31.7 ± 7.5	31.9 ± 10.3	<0.001
MUFA, %EI	14.3 ± 4.3	13.8 ± 4.8	13.5 ± 4.8	12.6 ± 5.7	<0.001
PUFA, %EI	5.8 ± 2	5.4 ± 2.1	5.4 ± 2.4	4.6 ± 2.4	<0.001
SFA, %EI	7.9 ± 2.2	8.6 ± 3	9 ± 3.2	9.4 ± 4.6	<0.001
Food groups (g/d)					
Dairy products	274.5 ± 207.5	266.2 ± 207.7	258.1 ± 199.8	253.4 ± 232.2	0.01
Red, fat, and ultra-processed meats	115.8 ± 82	125.5 ± 99.0	142.4 ± 110.7	156.4 ± 153.4	<0.001
White and lean meats, fish/shellfish, and eggs	133.7 ± 117.6	121.6 ± 159.3	107.9 ± 116.5	91.5 ± 128	<0.001
Refined cereals (breads and noodles)	131.0 ± 89.6	142.9 ± 94.3	147.8 ± 99.5	149.2 ± 117.6	0.05
Whole grains (breads, oats, and rice)	63.3 ± 58.8	53.7 ± 55.1	46.4 ± 53.5	34.9 ± 51.5	<0.001
Legumes	115.5 ± 122.3	72.7 ± 96.5	66.8 ± 73.2	42.1 ± 46.1	<0.001
Fats and oils, excluding olive oil	14.1 ± 12.3	16.3 ± 14.9	18 ± 15.5	18.6 ± 17.8	<0.001
Olive oil	4.6 ± 3.6	4.2 ± 3.3	4.2 ± 4.3	3.9 ± 4.4	0.003
Fruits	766.6 ± 467.5	569.2 ± 333.4	397.7 ± 298.3	224.4 ± 241.1	<0.001
Vegetables, excluding potatoes	261.4 ± 169.7	199.5 ± 110	154.7 ± 93.4	109.1 ± 84.9	<0.001
Natural fruit juice	153.9 ± 143.4	130.2 ± 124.6	106.7 ± 118.2	85.8 ± 126.8	<0.001
Industrial juices and soft drinks	100.5 ± 142.1	110.7 ± 152.8	133.4 ± 165.9	138.8 ± 192.9	0.002
Alcoholic beverage	93.0 ± 136.7	95.6 ± 173.5	108.2 ± 193.2	92.7 ± 145.9	0.15
Oilseeds	18.2 ± 23.2	15.7 ± 29.9	13.5 ± 29.6	9.6 ± 22.5	<0.001
Sugars and sweets	33.2 ± 33.7	34.9 ± 37.7	39.4 ± 39	40.5 ± 51.1	0.001
Preparation of the refined cereal base and fried foods	61.9 ± 59	69.8 ± 72.2	75.6 ± 79.8	69.4 ± 102	0.49

BMI, body mass index; CUME, Cohort of Universities of Minas Gerais; E-DII, energy-adjusted dietary inflammatory index; MUFA, monounsaturated fatty acid; PUFA, polyunsaturated fatty acid; SFA, saturated fatty acid; EI, energy intake.

Data are expressed as mean ± SD.

*P-values < 0.05 are in bold style by the multiple linear regression model, adjusted for sex (male or female), age (y), BMI (kg/m²), smoking (never smoked or former or current smokers).

Table 3Prevalence ratio for overweight^{*†} according to E-DII of the participants in CUME project (N = 3151), 2016

Total	Simple Poisson regression		Adjusted Poisson regression ^{‡§}	
	PR (95% CI)	P-value	PR (95% CI)	P-value
E-DII (quartile)				
1 (most anti-inflammatory)	1 (ref.)		1 (ref.)	
2	1.15 (1.01–1.33)	0.036	1.13 (0.98–1.29)	0.08
3	1.26 (1.11–1.44)	0.001	1.22 (1.07–1.39)	0.002
4 (most proinflammatory)	1.50 (1.32–1.70)	<0.001	1.39 (1.23–1.57)	<0.001
Male				
E-DII (quartile)				
1 (most anti-inflammatory)	1 (ref.)		1 (ref.)	
2	1.08 (0.90–1.32)	0.381	1.08 (0.89–1.31)	0.40
3	1.10 (0.91–1.32)	0.310	1.10 (0.91–1.32)	0.31
4 (most pro-inflammatory)	1.35 (1.14–1.59)	<0.001	1.35 (1.14–1.59)	<0.001
Female				
E-DII (quartile)				
1 (most anti-inflammatory)	1 (ref.)		1 (ref.)	
2	1.19 (0.99–1.43)	0.064	1.14 (0.95–1.36)	0.14
3	1.32 (1.10–1.58)	0.002	1.30 (1.10–1.55)	0.003
4 (most proinflammatory)	1.46 (1.22–1.74)	<0.001	1.38 (1.17–1.65)	<0.001

BMI, body mass index; CUME, Cohort of Universities of Minas Gerais; E-DII, energy-adjusted dietary inflammatory index; PR, prevalence ratio.

P-values in bold style are presented in significant regression models.

*Overweight for BMI ≥ 25 kg/m² [33] and BMI ≥ 28 kg/m² [32].

†Total sample shows 1233 overweight individuals: 525 men and 708 women.

‡Adjusted for age (y), smoking (never smoked or former or current smokers), physical activity (yes or no), and graduation area (health, human, exact, and agrarian sciences) for male and female groups.

§Additional adjustment for sex (male or female), for total sample.

- The SU.VI.MAX (Supplementation Study on Vitamins and Minerals Antioxidants) study with 7743 French women between 35 and 60 y of age and French men 45 to 60 y of age; and
- The NHANES (National Health and Nutrition Examination Survey) with 1734 Americans with previously diagnosed cardiovascular disease [38].

Together, these outcomes support the hypothesis that a more proinflammatory diet is related to the prevalence of overweight and obesity.

In this context, the origin of inflammation during obesity is not yet fully understood [39]. Inflammation is known to be associated

with adiposity, but this relationship could be bidirectional, creating a vicious cycle of positive feedback [40]. Additionally, excess of specific nutrients and some foods that have been associated with inflammation [41–44] may exacerbate the effects of adiposity. Furthermore, the most proinflammatory DII (highest quartile) in this study was associated with the most unhealthy eating habits (e.g., higher consumption of red, fat, and ultra-processed meats, and fats, soft drinks, sugar, and sweets), and the lowest DII quartile was associated with the healthiest diet (e.g., higher consumption of dairy products, white and lean meats, fish/shellfish and eggs, whole grains, legumes, olive oil, fruits, and vegetables). In fact, dietary patterns provide one of the best approaches to understanding

Table 4Prevalence ratio for obesity^{*†} according to E-DII of the participants of CUME project (N = 3151), 2016

Total	Simple Poisson regression		Adjusted Poisson regression ^{‡§}	
	PR (95% CI)	P-value	PR (95% CI)	P-value
E-DII (quartile)				
1 (most anti-inflammatory)	1 (ref.)		1 (ref.)	
2	1.63 (1.17–2.27)	0.004	1.56 (1.12–2.17)	0.008
3	1.78 (1.28–2.47)	<0.001	1.68 (1.21–2.31)	0.002
4 (most proinflammatory)	2.24 (1.64–3.06)	<0.001	1.99 (1.46–2.71)	<0.001
Male				
E-DII (quartile)				
1 (most anti-inflammatory)	1 (ref.)		1 (ref.)	
2	1.65 (0.97–2.81)	0.064	1.63 (0.96–2.77)	0.07
3	1.70 (1.01–2.85)	0.043	1.70 (1.01–2.84)	0.04
4 (most proinflammatory)	1.99 (1.22–3.26)	<0.001	1.97 (1.20–3.22)	<0.001
Female				
E-DII (quartile)				
1 (most anti-inflammatory)	1 (ref.)		1 (ref.)	
2	1.60 (1.05–2.44)	0.028	1.51 (0.99–2.29)	0.05
3	1.75 (1.16–2.66)	0.008	1.65 (1.09–2.50)	0.02
4 (most proinflammatory)	2.24 (1.50–3.36)	<0.001	1.95 (1.31–2.90)	0.001

BMI, body mass index; CUME, Cohort of Universities of Minas Gerais; E-DII, energy-adjusted dietary inflammatory index; PR, prevalence ratio.

P-values in bold style are presented in significant regression models.

*Obesity for BMI ≥ 30 kg/m² [32,33].

†Total sample presents 347 individuals with obesity, 144 men and 203 women.

‡Adjusted for age (y), smoking (never smoked or former or current smokers), physical activity (yes or no) and graduation area (health, human, exact, and agrarian sciences) for male and female groups.

the relationship between diet and disease because they allow for the evaluation of the inflammatory potential of diet, taking into account the potential synergies of different foods [45,46]. Given this, it is not surprising that diet quality tends to be better among those individuals with lower relative weight [47]. In turn, results from the present study also point to a possible solution for controlling chronic systemic inflammation. Although 8 of the parameters that comprise the DII are proinflammatory, 37 are anti-inflammatory. This latter group is characterized by being colorful, flavorful, nutrient dense, and calorie sparse [14]. So, consuming these constituents will not only reduce inflammation, but it is likely that it will lead to weight loss.

In the cross-sectional HELENA (Healthy Lifestyle in Europe by Nutrition in Adolescents) study, conducted with 3528 adolescents, Shivappa et al. [48] found that consumption of food items that tended to increase DII include bread and rolls; chocolate; margarine, butter, animal fats, and vegetable lipids; carbonated soft drinks; meat; and cakes, pies, biscuits, and sugar/honey/jam; and had lower consumption of vegetables (excluding potatoes), fruits, fruits and vegetable juices, and fish. Additionally, higher serum concentrations of IL-1, IL-2, and TNF- α were observed in adolescents with higher DII scores. Studies using DII scores for analysis also have reported positive associations of unhealthy diets (proinflammatory DII) with increased IL-6, IL-1, TNF- α , and CRP concentrations and lower concentrations of IL-10; whereas in healthy diets (anti-inflammatory DII) the concentrations of these markers were uniformly lower [31,35,49,50]. Thus, we can conclude that a diet considered healthy tends to be associated with lower DII (more anti-inflammatory) scores which, in turn, will reduce the occurrence of subclinical inflammatory processes.

The present study also found other attributes of healthier lifestyles among those who consumed a most anti-inflammatory diet: They were more likely to be nonsmokers, non-consumers of alcohol, and were more physically active, as has been reported by previous studies [31,51,52]. This combination of factors can result in a better quality of life and reduce the risk for developing future problems such as chronic inflammation, obesity, and other chronic NCDs [53–55].

On other hand, our sample was predominantly composed of young adults (only 3% of the population is ≥ 60 y of age) and all participants graduated from university. Other studies have found associations between healthier (most anti-inflammatory) diet and higher educational attainment (>4 years of college) among older people [56,57].

The strengths of the present study are its large sample size, which guarantees a high statistical power; use of a validated instrument to measure the inflammatory potential of the diet; control of important confounding factors in the analysis; the high level of education of the participants, which would tend to increase reliability of the self-reported information and to ensure a degree of homogeneity, increasing the internal validity of our study and reducing confounding related to education. Moreover, we validated self-reported weight, height, and BMI in a subsample of our cohort [22,34]. Our study is only the second study of the DII in relation to food intakes and cardiometabolic outcome in the Brazilian population and the third study in South American. The other two studies were in Argentina, where higher DII scores were associated with increased incidence of colorectal [58] and prostate [59] cancers. The other studies included one in northeastern Brazil, where inflammatory bowel disease was positively correlated with BMI in people with progressive multiple sclerosis [60] and a cross-sectional study (nested within a cohort) in Ribeirão Preto, São Paulo, where the DII was evaluated in young adults and had no association with insulin resistance or metabolic syndrome in either sex [20].

Some potential limitations of the study need to be recognized: The use of BMI as an indicator of adiposity did not allow us to differentiate lean mass from fat mass, or even total adiposity versus

central adiposity, despite being a tool widely used in epidemiologic studies, owing to its easy measurement and good correlation with body fat [61]. We did not measure inflammatory markers in plasma to assess their association with E-DII scores in the sample studied, due to the nature of online data collection. However, such associations had previously been reported [62–64]. In this context, Garcia-Arellano et al. [9] point out that the use of DII could help in the study of associations between food exposures and clinical events, with lower costs and avoidance of blood collection. Finally, the analysis of physical activity was classified only as whether or not it was practiced, as the study does not yet present the evaluation by the Brazilian version of the International Physical Activity Questionnaire [65] or the Global Physical Activity Questionnaire [66]. However, even using this crude measure we showed that the non-practice of physical activity contributes to the presence of obesity.

Conclusion

The most proinflammatory dietary pattern was associated with a higher prevalence of overweight and obesity in men and women participating in the baseline of the CUME study. E-DII scores also were associated with other unhealthy lifestyle characteristics, including sedentariness, smoking, and consumption of an obesogenic diet.

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References

- [1] VIGITEL. Vigitel Brasil 2017 - Vigilância De Fatores De Risco E Proteção Para Doenças Crônicas Por Inquérito Telefônico. 2018. Available at: <https://portal-arquivos2.saude.gov.br/images/pdf/2019/julho/25/vigitel-brasil-2018.pdf>. Accessed November 7, 2019
- [2] World Health Organization. World Health statistics 2014. Geneva, Switzerland: Author; 2014.
- [3] Magueresse-Battistoni BE, Labaronne E, Vidal H, Naville D. Endocrine disrupting chemicals in mixture and obesity, diabetes and related metabolic disorders. *World J Biol Chem* 2017;8:108.
- [4] Hermsdorff HHM, Volp ACP, Puchau B, Barbosa KBF, Zulet MÁ, Bressan J, et al. Contribution of gender and body fat distribution to inflammatory marker concentrations in apparently healthy young adults. *Inflamm Res* 2012;61:427–35.
- [5] Hermsdorff HHM, Zulet MÁ, Puchau B, Martínez JA. Central adiposity rather than total adiposity measurements are specifically involved in the inflammatory status from healthy young adults. *Inflammation* 2011;34:161–70.
- [6] Lavie CJ, Laddu D, Arena R, Ortega FB, Alpert MA, Kushner RF. Healthy weight and obesity prevention: JACC health promotion series. *J Am Coll Cardiol* 2018;72:1506–31.
- [7] Esmailzadeh A, Kimiagar M, Mehrabi Y, Azadbakht L, Hu FB, Willett WC. Dietary patterns and markers of systemic inflammation among Iranian women. *J Nutr* 2007;137:992–8.
- [8] Bressan J, Hermsdorff HHM, Zulet MÁ, Martínez JA. Hormonal and inflammatory impact of different dietetic composition: emphasis on dietary patterns and specific dietary factors. *Arg Braz Endocrinol Metabol* 2009;53:572–81.
- [9] Garcia-Arellano A, Ramallal R, Ruiz-Canela M, Salas-Salvadó J, Corella D, Shivappa N, et al. Dietary inflammatory index and incidence of cardiovascular disease in the PREDIMED study. *Nutrients* 2015;7:4124–38.
- [10] Ribeiro PV, de M, Andrade PA, Hermsdorff HHM, dos Santos CA, Cotta RMM, Estanislau J, de ASG, et al. Dietary non-nutrients in the prevention of non-communicable diseases: potentially related mechanisms. *Nutrition* 2019;66:22–8.
- [11] Pereira GA, Bressan J, Oliveira FLP, Sant'ana HMP, Pimenta AM, Lopes LL, et al. Dietary folate intake is negatively associated with excess body weight in Brazilian graduates and postgraduates (CUME project). *Nutrients* 2019;11:518.
- [12] Silveira BKS, De Novaes JF, Reis NDA, Lourenço LP, Capobianco AHM, Vieira SA, et al. "Traditional" and "healthy" dietary patterns are associated with low cardiometabolic risk in Brazilian subjects. *Cardiol Res Pract* 2018;2018.
- [13] Rocha DMUP, Lopes LL, Da Silva A, Oliveira LL, Bressan J, Hermsdorff HHM. Orange juice modulates proinflammatory cytokines after high-fat saturated meal consumption. *Food Funct* 2017;8:4396–403.
- [14] Shivappa N, Steck SE, Hurley TG, Hussey JR, Hébert JR. Designing and developing a literature-derived, population-based dietary inflammatory index. *Public Health Nutr* 2014;17:1689–96.

- [15] Phillips CM, Chen L-W, Heude B, Bernard JY, Harvey NC, Duijts L, et al. Dietary inflammatory index and non-communicable disease risk: a narrative review. *Nutrients* 2019;11:1873.
- [16] Okada E, Shirakawa T, Shivappa N, Wakai K, Suzuki K, Date C, et al. Dietary inflammatory index is associated with risk of all-cause and cardiovascular disease mortality but not with cancer mortality in middle-aged and older Japanese adults. *J Nutr* 2019;149:1451–9.
- [17] Garcia-Arellano A, Martínez-González MA, Ramallal R, Salas-Salvadó J, Hébert JR, Corella D, et al. Dietary inflammatory index and all-cause mortality in large cohorts: the SUN and PREDIMED studies. *Clin Nutr* 2019;38:1221–31.
- [18] Park SY, Kang M, Wilkens LR, Shvetsov YB, Harmon BE, Shivappa N, et al. The dietary inflammatory index and all-cause, cardiovascular disease, and cancer mortality in the multiethnic cohort study. *Nutrients* 2018;10:1–12.
- [19] Ferreira YAM, Kravchychyn ACP, Vicente S, de CF, Campos RM, Tock L, Oyama LM, et al. An interdisciplinary weight loss program improves body composition and metabolic profile in adolescents with obesity: associations with the dietary inflammatory index. *Front Nutr* 2019;6:77.
- [20] Carvalho CA, Silva AAM, Assunção MCF, Fonseca PCA, Barbieri MA, Bettiol H, et al. The dietary inflammatory index and insulin resistance or metabolic syndrome in young adults. *Nutrition* 2019;58:187–93.
- [21] Andrade PA, Hermsdorff HHM, Leite JJA, Shivappa N, Hébert JR, Henriques HKF, et al. Baseline pro-inflammatory diet is inversely associated with change in weight and body fat 6 months following-up to bariatric surgery. *Obes Surg* 2019;29:457–63.
- [22] Gomes Domingos A, Miranda AES, Pimenta AM, Hermsdorff HHM, Oliveira FLP de, dos Santos LC, et al. Cohort profile: the cohort of Universities of Minas Gerais (CUME). *Int J Epidemiol* 2018;47(6):1743–4.
- [23] Seguí-Gómez M, de la Fuente C, Vázquez Z, de Irala J, Martínez-González MA. Cohort profile: the "Seguimiento Universidad de Navarra" (SUN) study. *Int J Epidemiol* 2006;35:1417–22.
- [24] Henn RL, Fuchs SC, Moreira LB, Fuchs FD. Development and validation of a food frequency questionnaire (FFQ-Porto Alegre) for adolescent, adult and elderly populations from Southern Brazil. *Cad Saude Publica* 2010;26(11):2068–79.
- [25] TACO. Brazilian Food Composition Table vol. 1. 2011. Available at: http://189.28.128.100/nutricao/docs/taco/tab_bras_de_comp_de_alim_doc.pdf. Accessed November 7, 2019.
- [26] United States Department of Agriculture. USDA Food Composition Databases 2018:9894. Available at: <https://ndb.nal.usda.gov/ndb/search/list>. Accessed 7 November 2019.
- [27] Rodrigues-Amaya DB. [Brazilian sources of carotenoides: table of composition of carotenoids in foods], 2008. Available at: https://www.mma.gov.br/estruturas/sbf_agrobio/_publicacao/89_publicacao09032009113306.pdf. Accessed 7 November 2018.
- [28] Neveu V, Perez-Jiménez J, Vos F, Crespy V, du Chaffaut L, Mennen L, et al. Phenol-Explorer: an online comprehensive database on polyphenol contents in foods. *Database (Oxford)* 2010;2010:1–9.
- [29] Shivappa N, Hebert JR, Anderson LA, Shrubsole MJ, Murray LJ, Getty LB, et al. Dietary inflammatory index and risk of reflux oesophagitis, Barrett's oesophagus and oesophageal adenocarcinoma: a population-based case-control study. *Br J Nutr* 2017;117:1323–31.
- [30] Shivappa N, Hébert JR, Rashidkhani B. Dietary Inflammatory index and risk of esophageal squamous cell cancer in a case-control study from Iran. *Nutr Cancer* 2015;67:1253–9.
- [31] Ramallal R, Toledo E, Martínez JA, Shivappa N, Hébert JR, Martínez-González MA, et al. Inflammatory potential of diet, weight gain, and incidence of overweight/obesity: the SUN cohort. *Obesity* 2017;25:997–1005.
- [32] Bjorntorp P, Bray GA, Carroll KK, Chuchalin A, Dietz WH, Ehrlich GE, et al. Obesity: preventing and managing the global epidemic. WHO Tech Rep Ser 2000. Geneva, Switzerland: Author; 2000.
- [33] Pan American Health Organization. XXXVI Reunión del Comité Asesor de Investigaciones en Salud – Encuesta Multicéntrica – Salud Bienestar y Envejecimiento (SABE) en América Latina e el Caribe 2009. Available at: https://www.paho.org/hq/index.php?option=com_content&view=article&id=1734:2009-documentos-cais&Itemid=1654&lang=es. Accessed November 7, 2019.
- [34] Miranda AES, da S, Ferreira AVM, Oliveira FLP, Hermsdorff HHM, Bressan J, et al. Validation of Metabolic syndrome and its self reported components in the CUME study. *REME Rev Min Enferm* 2017;21:1–7.
- [35] Willett WC. *Nutritional epidemiology*. 2nd ed. New York: Oxford University Press; 1998.
- [36] Protein C, Cavicchia PP, Steck SE, Hurley TG, Hussey JR, Ma Y, et al. A new dietary inflammatory index predicts interval changes in serum high-sensitivity. *J Nutr* 2009;139:2365–72.
- [37] Ruiz-Canela M, Zazpe I, Shivappa N, Hébert JR, Sánchez-Tainta A, Corella D, et al. Dietary inflammatory index and anthropometric measures of obesity in a population sample at high cardiovascular risk from the PREDIMED (PREVENCIÓN con Dieta MEDiterránea) trial. *Br J Nutr* 2015;113:984–95.
- [38] Ruiz-Canela M, Bes-Rastrollo M, Martínez-González MA. The role of dietary inflammatory index in cardiovascular disease, metabolic syndrome and mortality. *Int J Mol Sci* 2016;17.
- [39] Archer E, Lavie CJ, Hill JO. The contributions of 'diet' genes and physical activity to the etiology of obesity: contrary evidence and consensus. *Prog Cardiovasc Dis* 2018;61:89–102.
- [40] Gregor MF, Hotamisligil GS. Inflammatory mechanisms in obesity. *Annu Rev Immunol* 2011;29:415–45.
- [41] Rocha DM, Bressan J, Hermsdorff HH. The role of dietary fatty acid intake in inflammatory gene expression: a critical review. *Sao Paulo Med J* 2017;135:157–68.
- [42] Hermsdorff HHM, Zulet MÁ, Puchau B, Martínez JA. Fruit and vegetable consumption and proinflammatory gene expression from peripheral blood mononuclear cells in young adults: a translational study. *Nutr Metab* 2010;7:1–11.
- [43] Hermsdorff HHM, Zulet MÁ, Abete I, Martínez JA. A legume-based hypocaloric diet reduces proinflammatory status and improves metabolic features in overweight/obese subjects. *Eur J Nutr* 2011;50:61–9.
- [44] Khan N, Khymenets O, Urpi-Sardà M, et al. Cocoa polyphenols and inflammatory markers of cardiovascular disease. *Nutrients* 2014;6(2):844–80.
- [45] Jacobs DR, Steffen LM. Nutrients, foods, and dietary patterns as exposures in research. *Am J Clin Nutr* 2003;78:508–13.
- [46] DiNicolantonio JJ, Mehta V, Onkaramurthy N, O'Keefe JH. Fructose-induced inflammation and increased cortisol: a new mechanism for how sugar induces visceral adiposity. *Prog Cardiovasc Dis* 2018;61:3–9.
- [47] Pate RR, Taverno Ross SE, Liese AD, Dowda M. Associations among physical activity, diet quality, and weight status in US adults. *Med Sci Sports Exerc* 2015;47:743–50.
- [48] Shivappa N, Hebert JR, Marcos A, Diaz LE, Gomez S, Nova E, et al. Association between dietary inflammatory index and inflammatory markers in the HEL-ENA study. *Mol Nutr Food Res* 2017;61:1–10.
- [49] Nettleton JA, Steffen LM, Mayer-davis EJ, Jenny NS, Jiang R, Herrington DM, et al. Dietary patterns are associated with biochemical markers of inflammation and endothelial activation 2006:1–3.
- [50] Julia C, Meunier N, Touvier M, Ahluwalia N, Sapin V, Papet I, et al. Dietary patterns and risk of elevated C-reactive protein concentrations 12 years later. *Br J Nutr* 2013;110:747–54.
- [51] Pimenta AM, Toledo E, Rodriguez-Diez MC, Gea A, Lopez-Iracheta R, Shivappa N, et al. Dietary indexes, food patterns and incidence of metabolic syndrome in a Mediterranean cohort: the SUN project. *Clin Nutr* 2015;34:508–14.
- [52] Shivappa N, Steck SE, Hussey JR, Ma Y, Hebert JR. Inflammatory potential of diet and all-cause, cardiovascular, and cancer mortality in National Health and Nutrition Examination Survey III study. *Eur J Nutr* 2017;56:683–92.
- [53] Fletcher GF, Landolfo C, Niebauer J, Ozemek C, Arena R, Lavie CJ. Promoting Physical activity and exercise: JACC health promotion series. *J Am Coll Cardiol* 2018;72:1622–39.
- [54] Ozemek C, Laddu DR, Lavie CJ, Claeys H, Kaminsky LA, Ross R, et al. An update on the role of cardiorespiratory fitness, structured exercise and lifestyle physical activity in preventing cardiovascular disease and health risk. *Prog Cardiovasc Dis* 2018;61:484–90.
- [55] Harber MP, Kaminsky LA, Arena R, Blair SN, Franklin BA, Myers J, et al. Impact of cardiorespiratory fitness on all-cause and disease-specific mortality: advances since 2009. *Prog Cardiovasc Dis* 2017;60:11–20.
- [56] Hiza HAB, Casavale KO, Guenther PM, Davis CA. Diet quality of Americans differs by age, sex, race/ethnicity, income, and education level. *J Acad Nutr Diet* 2013;113:297–306.
- [57] Hsiao PY, Mitchell DC, Coffman DL, et al. Dietary patterns and diet quality among diverse older adults: the University of Alabama at Birmingham Study of Aging. *J Nutr Health Aging* 2013;17(1):19–25.
- [58] Niclis C, Pou SA, Shivappa N, Hébert JR, Steck SE, Diaz M, del P. Proinflammatory dietary intake is associated with increased risk of colorectal cancer: results of a case-control study in Argentina using a multilevel modeling approach. *Nutr Cancer* 2018;70:61–8.
- [59] Shivappa N, Niclis C, Coquet JB, Román MD, Hébert JR, Diaz M, del P. Increased inflammatory potential of diet is associated with increased odds of prostate cancer in Argentinian men. *Cancer Causes Control* 2018;29:803–13.
- [60] Da Costa Silva BY, De Carvalho Sampaio HA, Shivappa N, Hébert J, Silva Albuquerque LDA, Ferreira Carioca AA, et al. Interactions between dietary inflammatory index, nutritional state and multiple sclerosis clinical condition. *Clin Nutr ESPEN* 2018;26:35–41.
- [61] Lear SA, Humphries KH, Kohli S, Birmingham CL. The use of BMI and waist circumference as surrogates of body fat differs by ethnicity. *Obesity* 2007;15:2817–24.
- [62] Shivappa N, Steck SE, Hurley TG, Hussey JR, Ma Y, Ockene IS, et al. A population-based dietary inflammatory index predicts levels of C-reactive protein in the Seasonal Variation of Blood Cholesterol Study (SEASONS). *Public Health Nutr* 2014;17:1825–33.
- [63] Wirth MD, Shivappa N, Davis L, Hurley TG, Ortaglia A, Drayton R, et al. Construct validation of the dietary inflammatory index among African Americans. *J Nutr Heal Aging* 2017;21:487–91.
- [64] Bodén S, Wennberg M, Van Guelpen B, Johansson I, Lindahl B, Andersson J, et al. Dietary inflammatory index and risk of first myocardial infarction; a prospective population-based study. *Nutr J* 2017;16:1–10.
- [65] IPAQ Group. International Physical Activity Questionnaire. Available at: <https://sites.google.com/site/theipaq/>. Accessed November 7, 2019.
- [66] World Health Organization. Global Physical Activity Questionnaire (GPAQ) WHO STEPwise approach to NCD risk factor surveillance. *Surveill Popul Prev Noncommunicable Dis Dep* 2008;1–3.