Cross-sectional associations between body mass index, waist circumference, and multimorbidity: *Pró-Saúde* study

Fernanda Muniz de Macedo Stumpf ^{Corresp., 1}, Alessandra Silva Dias de Oliveira ¹, Eduardo Faerstein ², Cintia Chaves Curioni ¹

¹ Instituto de Nutrição, Universidade do Estado do Rio de Janeiro, Rio de Janeiro, Rio de Janeiro, Brazil

² Instituto de Medicina Social, Universidade do Estado do Rio de Janeiro, Rio de Janeiro, Rio de Janeiro, Brazil

Corresponding Author: Fernanda Muniz de Macedo Stumpf Email address: fermuniz@hotmail.com

Background: This study aimed 1) To investigate the association between the Body Mass Index (BMI) and waist circumference (WC) with multimorbidity (MM), and, 2) To identify patterns of MM and investigate the relationship between BMI and WC with specific combinations of MM (patterns of MM). Methods: A cross-sectional study was conducted with 2,698 participants of the fourth phase of the Brazilian Longitudinal Study of Adult Health (Pró-Saúde Study). Body mass index (BMI) and waist circumference (WC) and MM was defined by the presence of two or more morbidities. MM patterns were identified by exploratory factor analysis based on tetrachoric correlations. Logistic regression models were used to assess associations (odds ratios [OR] with the respective confidence intervals [CI]). **Results:** Of the total number of participants, 39.5 % were overweight and 30.0% were obese; 89.0 % (n=1468) of women and 77.0 % (n=952) of men were abdominally obese. Indeed, 60.7% (n=1635) was identified with MM. For the category four or more morbidities, OR values of 5.98 (95% CI 4.84-7.13) and 7.48 (95% CI 6.14-8.18) were found for each point of increase in BMI, and 6.74 (95% CI 5.48–7.99) and 8.48 (95% CI 7.64–9.29) for each additional centimeter in the WC, for female and male, respectively. Five patterns of MM were identified: respiratory, osteoarticular, cardiometabolic, gastric, and thyroid diseases (56.4% of the total variance). Positive associations were found between BMI and patterns of cardiometabolic, osteoarticular, thyroid and gastric diseases (higher OR of 1.09 [95% CI 1.04–1.14]) and WC and patterns of cardiometabolic and osteoarticular (higher OR of 1,04 [95% CI 1.03-1.04]). **Conclusions:** The results showed that an increase of both BMI and WC was associated with a higher number of morbidities and with patterns of cardiometabolic and osteoarticular diseases.

Cross-sectional associations between body mass index, waist circumference, and multimorbidity: Pró-Saúde Study. Fernanda Muniz de Macedo Stumpf¹; Alessandra Silva Dias de Oliveira²; Eduardo Faerstein³; Cintia Chaves Curioni⁴ ¹ Instituto de Nutrição, Universidade do Estado do Rio de Janeiro, Rio de Janeiro, Brazil. ORCID 0000- 0002-1066-1932. ² Instituto de Nutrição, Universidade do Estado do Rio de Janeiro, Rio de Janeiro, Brazil. ORCID 0000- 0002-3232-5868 ³ Instituto de Medicina Social, Universidade do Estado do Rio de Janeiro, Rio de Janeiro, Brazil. ORCID 0000-0002-4027-4896 ⁴ Instituto de Nutrição, Universidade do Estado do Rio de Janeiro, Rio de Janeiro, Brazil. ORCID 0000- 0002-5160-9567. Corresponding Author: Fernanda Muniz de Macedo Stumpf – Instituto de Nutrição, Universidade do Estado do Rio de Janeiro - Rua São Francisco Xavier, 524, sala 12144 F - Rio de Janeiro, Brazil - Zipcode: 20550-900 - Telephone: +55 21 2334.0150/ Fax: +55 21 2334.0722 - Email:fermuniz@hotmail.com

40 Abstract

- 41 Background: This study aimed 1) To investigate the association between the body mass index
- 42 (BMI) and waist circumference (WC) with multimorbidity (MM) and, 2) To identify patterns of
- 43 MM and investigate the relationship between BMI and WC with specific combinations of MM
- 44 (patterns of MM).
- 45 Methods: A cross-sectional study was conducted with 2,698 participants of the fourth phase of
- 46 the Brazilian Longitudinal Study of Adult Health (*Pró-Saúde* Study). MM was defined by the
- 47 presence of two or more morbidities. MM patterns were identified by exploratory factor analysis
- 48 based on tetrachoric correlations. Logistic regression models were used to assess associations
- 49 (odds ratios [OR] with the respective confidence intervals [CI]).
- 50 **Results:** Of the total number of participants, 39.5 % were overweight and 30.0% were obese;
- 51 89.0 % of women and 77.0 % of men were abdominally obese. Indeed, 60.7% was identified
- 52 with MM. For the category four or more morbidities, OR values of 5.98 (95% CI 4.84–7.13) and
- 53 7.48 (95% CI 6.14–8.18) were found for each point of increase in BMI, and 6.74 (95% CI 5.48–
- 54 7.99) and 8.48 (95% CI 7.64–9.29) for each additional centimeter in the WC, for female and
- 55 male, respectively. Five patterns of MM were identified: respiratory, osteoarticular,
- 56 cardiometabolic, gastric, and thyroid diseases (56.4% of the total variance). Positive associations
- 57 were found between BMI and patterns of cardiometabolic, osteoarticular, thyroid, and gastric
- diseases (higher OR of 1.09 [95% CI 1.04–1.14]) and less pronounced between WC and patterns
- 59 of cardiometabolic and osteoarticular (higher OR of 1,04 [95% CI 1.03-1.04]).
- 60 **Conclusions:** The results showed that an increase of both BMI and WC was associated with a 61 higher number of morbidities and BMI with patterns of cardiometabolic and osteoarticular 62 diseases.
- 63 Keywords: Overweight; Obesity; Waist circumference; Multimorbidity; Multimorbidity patterns.
- 64
- 65
- 66
- 67
- 68
- 69
- 70

72 Introduction

73 Multimorbidity (MM) refers to the coexistence of multiple health conditions in a single individual (Barnett et al., 2012; MacMahon & The Academy of Medical Sciences, 2018). MM can 74 75 be measured in different ways, but assessing the number of morbidities involved, effects, and associations is still challenging (Zheng et al., 2021). Measuring MM using the number of reported 76 77 medical conditions is simple and useful, and it can assess the impact of different conditions on people's health (Diane Zheng et al., 2021). Studies indicate that diseases are grouped according to 78 79 specific patterns (Vetrano et al., 2018, 2019, 2020). However, the studies that assess such patterns are still scarce, especially in Brazilian context, and discrepancies exist regarding the number and 80 81 characteristics of the diseases considered, their severity levels, statistical analyses used, and other 82 methodological aspects (Prados-Torres et al., 2012, 2014). For example, it is known that one disease can be a risk factor for others or be involved in their pathophysiological mechanisms. 83 84 Furthermore, these MM patterns may be affected by socioeconomic factors and lifestyles, as well as eating habits and excess of weight (Van Den Akker, Buntinx & Knottnerus, 1996; Prados-Torres 85 86 et al., 2012; De Carvalho et al., 2018). This excessive fat accumulation, especially abdominal fat 87 is associated with important metabolic, hormonal, and inflammatory changes (Flor et al., 2015; 88 Hall et al., 2015; Catrysse & van Loo, 2017; Dragano, Haddad-Tovolli & Velloso, 2017; Nájera Medina et al., 2019). Therefore, excess of weight, particularly obesity, could be involved in the 89 90 development of multiple non-communicable chronic diseases (NCDs) (Samper-Ternent & Al Snih, 2012; Ezzati, 2017; Zhang et al., 2020). 91

92 Most studies on MM focused on evaluating its definition and prevalence (Willadsen et 93 al., 2016; Xu, Mishra & Jones, 2017; Araujo et al., 2018; Nguyen et al., 2019). Some of them 94 sought to identify risk factors for MM. Among the risk factors considered, many of them included 95 body mass index (BMI), and only a few considered waist circumference (WC) (Jovic, Marinkovic 96 & Vukovic, 2016; Leal Neto, Barbosa & Meneghini, 2016; Kivimäki et al., 2017; De Carvalho et al., 2018; Jawed et al., 2020; Zhang et al., 2020; Flores et al., 2021). Zhang et al. (2020) observed 97 98 an association between BMI, WC, and waist/hip ratio with MM in older adults. In Brazil, a recent 99 study with a representative sample of the population found an association between increased BMI 100 and MM but did not assess WC (Flores et al., 2021). Petarli et al. (2019) identified an increased 101 prevalence of MM associated with increased WC. Carvalho et al. (2018) found an association of 102 BMI and WC with MM and BMI with the following patterns: cardiometabolic, oncological,

mental/occupational, musculoskeletal, and respiratory diseases (De Carvalho et al., 2018; Petarli
et al., 2019; Zhang et al., 2020). Thus, a Brazilian study that simultaneously assesses the impact
of increased BMI and WC on MM, as the main exposure, considering the number of diseases
involved and the patterns in which the diseases are grouped, can contribute to this field of research,
especially in similar contexts.

The studies have suggested a positive association between the overweight and MM. This Brazilian study hypothesizes that excess weight and accumulation of fat in the abdominal area, represented by the BMI and WC, is associated with a greater number of morbidities grouped. Moreover, it is possible that both BMI and WC are related to specific diseases patterns. The objective of this study was to investigate the relation between BMI and WC with MM, and to identify patterns of MM and investigate the relationship between BMI and WC with specific combinations of MM (patterns of MM).

115

116 Materials & Methods

- 117
- 118 Study design and population

This was a cross-sectional study inserted in a prospective cohort study conducted at several university campuses in the State of Rio de Janeiro, Brazil, in 2011 and 2012, involving non-faculty civil servants, *Pro-Saúde* Study. All technical and administrative permanent staff members were invited to the study. The exclusion criteria were current non-medical leave of absence and working relocation to another institution (Faerstein et al., 2005).

124 The population of the present study included 2,698 participants of the fourth phase of the 125 *Pró-Saúde* Study. The protocols of the 2011–2012 *Pró-Saúde* Study were approved by the 126 Research Ethics Committee of the Institute of Social Medicine, State University of Rio de Janeiro 127 (CAAE: 0041.0.259.000-11). All participants signed an informed consent form.

128

129 *Study variables*

130 MM was evaluated in a self-reported way, through the following question: *Have you ever* 131 been told by any doctors you have had...? For each condition or disease, check Yes or No. The 18 chronic conditions: 132 options involved medical hypertension, diabetes (DM), hypercholesterolemia, acute myocardial infarction, angina, stroke, asthma, pulmonary 133

emphysema, cholecystitis, peptic ulcer, gastritis, nephrolithiasis, arthrosis, slipped disc, repetitive strain injury (RSI), hyperthyroidism, hypothyroidism, and tuberculosis. All affirmative answers (Yes) for each condition/morbidity were considered for the creation of the MM variable. The original list of the study was maintained due to the lack of consensus as for the number of diseases to be considered for assessing the extent of MM, and according to the results of a systematic review, the prevalence was severely underestimated when studies considered a list with less than twelve conditions/diseases (Chua et al., 2021).

MM was evaluated from the simple count of self-reported diseases in the questionnaire applied in phase 4 of the *Pró-Saúde Study*, thus, the presence of two or more chronic health conditions was considered MM (Hudon, Fortin & Vanasse, 2005; Valderas, Starfi & Sibbald, 2009). From the morbidities count, participants were grouped as follows: with no MM (zero or one morbidity), with two morbidities, with three morbidities, and with four or more morbidities.

BMI and WC are the exposure variables of central interest. The following anthropometric measures were performed by trained personnel using portable digital scale with precision of 50 grams, stadiometer and anthropometric tape used: weight (kg), height (m), and WC (cm), measured according to Lohman's protocols, that recommend to use the midpoint between the lower margin of the last palpable rib and the top of the iliac crest (Lohman TG, Roche AF, 1988). BMI (weight/height²) was calculated based on the weight and height data.

In the analyses, BMI and WC were used continuously, but to characterize the study 152 153 population, the following categories were used for BMI: underweight/eutrophic, BMI up to 24.9 kg/m²; overweight, BMI between 25 and 29.9 kg/m²; and obesity, BMI \geq 30 kg/m² (WHO, 2000). 154 155 The cut-off points recommended by the World Health Organization for South Americans were used for the WC, an abdominal adiposity indicator, considering abdominal obesity and increased 156 157 cardiovascular risk as measured by WC > 90 cm in men and > 80 cm in women (WHO, 2000). The investigated covariates were gender (female, male), age (in years), education (university or 158 more, high school, elementary school), marital status (married/stable union, separated/divorced, 159 widowed, single), race (black/brown, white, yellow/indigenous), and modified equivalent income 160 161 proposed by the Organization for Economic Cooperation and Development (OECD) (Celeste & 162 Bastos, 2014), which for the descriptive analyses was categorized into up to three minimum wages (MW), from three to six MW and more than six MW, and for the ordinal and binary logistic 163 164 regression analyses it was used continuously, smoking (smoker, ex-smoker, non-smoker), physical

activity in the last two weeks (yes or no), and consumption of fruits and vegetables (< or \ge 4 times a week).

167

168 *Statistical analyses*

A descriptive analysis was performed for the following variables: gender, age, education, marital status, race, income, BMI, abdominal obesity, smoking, physical activity, number of morbidities and, fruit and vegetable consumption, using the mean and standard deviations of continuous variables and the absolute and relative frequency of categorical variables, since all of them follow a normal distribution according to the Shapiro Wilk test.

Associations of BMI and WC and MM categories (none or one; two, three, and four or more morbidities) were estimated using ordinal logistic regression with the unadjusted and adjusted model, providing the OR and 95% CI. Univariate analyses of the covariates (gender, age, education, marital status, race, income, smoking, physical activity, number of morbidities and, fruit and vegetable consumption), were performed and those with a p-value < 0.2 were included in the model.

180 MM patterns were identified after submitting data to an exploratory factor analysis, technique used to investigate patterns for a large number of variables into fewer numbers of factors, 181 182 to identify concurrent morbidities (Ferrando & Lorenzo-Seva, 2017). Since morbidity is a binary variable (presence or absence), factor analysis was implemented using a tetrachoric matrix, where 183 184 both the variables were binary and the extraction method Robust Diagonally Weighted Least Squares (RDWLS) (Asparouhov & Muthén, 2010). The adequacy of the model was assessed using 185 186 the Root Mean Square Error of Approximation (RMSEA), Comparative Fit Index (CFI), and Tucker-Lewis Index (TLI). According to the literature, the RMSEA values must be < 0.08, with a 187 188 < 0.10 CI, and the CFI and TLI values must be > 0.90 or preferably > 0.95 (Brown, 2006). It is suggested that composite trustworthiness values should be > 0.70. However, values between 0.6 189 and 0.7 are acceptable, if the other quality indices of the model are adequate (Hair, J. F., Black, 190 W. C., Babin, B. J., Anderson, R. E., & Tatham, 2009). Morbidities with factor loadings > 0.30 in 191 192 a given factor were considered to belong to that comorbidity pattern. The stability of the factors was evaluated by index H (Ferrando & Lorenzo-Seva, 2018). However, values between 0.6 and 193 0.7 are acceptable, if the other quality indices of the model (CFI, TLI, and RMSEA) are adequate 194

(Hair, J. F., Black, W. C., Babin, B. J., Anderson, R. E., & Tatham, 2009). Exploratory factoranalysis was presented without stratification by sex, since it is a better fit to the model.

197 Adjusted binary logistic regressions were conducted to assess the association between BMI 198 and WC and each MM pattern. It should be noted that each subject could belong to more than one 199 pattern, as long as they accumulated more than two morbidities in each of the patterns to which 200 they belonged. A univariate model was used to assess the associations of covariates with MM and 201 its patterns. Variables with a p-value < 0.2 were included in the multivariate model. Regression 202 models were stratified by sex.

- 203 Descriptive and regression analyses were performed using the SPSS V.20 (IBM) software 204 and exploratory factor analysis was performed using the FACTOR 10.10 software.
- 205
- 206

207 **Results**

In the studied population, 56.3% were female with a mean age of 51.5 years (SD=9.0). 208 Regarding BMI, 39.5% were overweight and 30.0% were obese. For WC, 89.0% of women and 209 77% of men exhibited abdominal obesity. As for the number of morbidities, the median showed 210 211 a value of 2.0 (interquartile range: 2.0-3.0), and the total prevalence of MM was 60.7%. 212 Approximately half of the participants had higher education or more (53.3%), were married 213 (61.3%), white (50.1%), and with income above three minimum wages (73.2%) (Table 1). As for 214 lifestyle, most were non-smokers (60.3%), did not perform any physical activity in recent weeks 215 (57.0%), and consumed fruits and vegetables more than four times a week (n=1611, 54.9% and 216 n=1757, 60.7%, respectively).

217 Table 2 shows the results of the ordinal logistic regressions for each MM category. The results showed measures of a positive association in the adjusted models, revealing the impact of 218 219 covariates (age, gender, race, income, education, marital status, and fruit and vegetable 220 consumption) on the outcome, contrasting with the unadjusted model. There was an increased odds of MM with BMI and WC for all categories of MM, with a progressive increase in the OR (dose-221 222 response gradient). Considering the category four or more morbidities, we found in the adjusted 223 model to age, race, income, marital status, and consumption of fruits and vegetables, for each point 224 of increase in BMI, OR values of 7 (95% CI 5.75-8.15) for the total population, 5.98 (95% CI 4.84-7.13) for female, and 7.48 (95% CI 6.14-8.18) for male. For each additional centimeter in 225 the WC, OR values of 7.48 (95% CI 6.16-8.48) for the total population, 6.74 (95% CI 5.48–7.99) 226

for female, and 8.48 (95% CI 7.64–9.29) for male. The results of the unadjusted model are shownin the supplementary material (Supplemental files).

229 Five patterns of MM were identified, using an exploratory factor analysis, as described in the materials and methods section: respiratory diseases (asthma and emphysema), osteoarticular 230 diseases (RSI, arthrosis, and slipped disc), thyroid diseases (hypo- and hyperthyroidism), 231 cardiometabolic diseases (hypertension, DM, hypercholesterolemia, myocardial infarction, 232 angina, stroke), and gastric diseases (ulcer and gastritis), which explained 56.4% of the total 233 variance. This factorial solution showed adequate fit indices (RMSEA = 0.019; CFI = 0.967; TLI 234 = 0.930). The composite reliability of the factors was also acceptable (> 0.70) for almost all factors, 235 236 except for the pattern of osteoarticular diseases (CR = 0.565). However, this pattern was maintained in this study, since the adjustment of the model, evaluated by CFI, TLI, and RMSEA 237 was adequate, and the morbidities of this factor also presented adequate factor loadings. Table 3 238 239 shows the factor loadings of the morbidities, as well as the composite reliability and the H index.

In the adjusted model, positive associations were identified between BMI and WC and patterns of cardiometabolic diseases for BMI and WC, respectively; and less pronounced between osteoarticular diseases and BMI and WC; and thyroid disease and BMI, OR of 1.09 (95% CI 1.04– 1.14) for BMI (Table 4).

244

245 **Discussion**

Our study confirmed the hypothesis that an increase in BMI and WC is associated with MM emphasizing the impact of the number of diseases involved and the patterns of MM. We took a more sensitive look at variables such as BMI and WC, continuously evaluating them and moving away from BMI categories.

Studies have shown that the risk of MM increases with increasing BMI, being twice as high in overweight individuals and ten times higher in individuals with class III obesity compared to eutrophic individuals (Kivimäki et al., 2017).

In Brazil, few studies have analyzed the association of BMI and WC with MM patterns. MM was evaluated in different ways, considering categories (number of diseases involved), and respecting disease patterns. It might be a differential, since other studies separately assessed the impact of body weight, through BMI, and few evaluated the accumulation of abdominal fat, by

waist circumference, which, as already described, plays a fundamental role in the development ofchronic diseases.

259 Results of previous studies corroborate the association found between BMI, WC, and MM, although the vast majority only assess BMI and not indicators of abdominal adiposity, such as WC 260 (Agborsangava et al., 2013; Jovic, Marinkovic & Vukovic, 2016; Christofoletti, Streb & Del Duca, 261 262 2018; Zhang et al., 2020). The study by Petarli et al. (2019) investigated the association of WC in MM in adults (Petarli et al., 2019), and Zhang et al. (2020) assessed the WC among older adults 263 (Zhang et al., 2020) and found an association between increased abdominal fat and MM. WC is 264 considered relevant due to the role that abdominal adiposity plays in the development of NCDs 265 (Melo, 2011; Nájera Medina et al., 2019). 266

The *Pró-Saúde Study* participants showed a very high WC (96.33 cm) and considering the cut-off points for cardiometabolic risk of 80 cm for female and 90 for male, it shows a greater predisposition to a higher risk of diseases, both in male and in female (ABESO, 2016). Excess weight, especially when associated with increased abdominal adiposity, increases the risk of developing hypertension between 65% and 75% (Hall et al., 2015; Seravalle & Grassi, 2017). It is estimated that 85% of adult type 2 diabetics are also obese (Ezzati, 2017; Chait & Hartigh, 2020).

Studies suggested that female sex, aging, and the presence of obesity are some of the factors 273 274 associated with MM (Leal Neto, Barbosa & Meneghini, 2016; Kivimäki et al., 2017; De Carvalho et al., 2018; Jantsch, Alves & Faerstein, 2018; Nguyen et al., 2019; Song et al., 2019). The present 275 276 analyses, stratified by sex, showed associations of BMI and WC in both genders, however with greater association in male, which contrasts with the findings of previous studies. This result can 277 278 be justified by the difference in body composition between male and female. Although female have proportionally more fat mass than male, the latter tend to have greater accumulation of 279 280 abdominal fat (Song et al., 2014; Bredella, 2017; Schorr et al., 2018).

A systematic review found more than 97 patterns with two or more diseases; however, three groups prevailed in all analyzed studies, those of cardiometabolic, musculoskeletal, and mental health diseases (Prados-Torres et al., 2014). In addition, a Brazilian study conducted with the objective of defining the grouping patterns of diseases and relating them to socioeconomic and lifestyle factors found four groups of diseases (cardiometabolic/cancer, mental/occupational, musculoskeletal, and respiratory) and an association with high BMI values was also identified (De Carvalho et al., 2018).

288 It is noteworthy that the patterns found grouped two or more diseases, with a well-related physiological mechanism. The cardiometabolic pattern, in addition to grouping the largest number 289 290 of diseases, was also associated with BMI and WC, corroborating the literature that shows the impact of excess weight on the pattern of diseases (Flor et al., 2015; Murray et al., 2015; Tang, 291 Liebeskind & Towfighi, 2017; Nájera Medina et al., 2019). The pattern of osteoarticular diseases 292 293 was also associated with adiposity. A population-based cohort study with 1,764,061 participants found a positive association of being overweight and the development of knee, hip, and hand 294 295 osteoarthritis. In obese class II individuals, the risk of developing knee osteoarthritis was 4.7 times higher compared to eutrophic individuals (Reves et al., 2016). 296

In the present study, we identified five patterns of MM: respiratory, osteoarticular, thyroid, cardiometabolic, and gastric diseases. It was similar to other studies, except for mental health diseases, which was not included in the list of diseases in the baseline study questionnaire. Furthermore, we observed positive associations between BMI and patterns of cardiometabolic, osteoarticular, thyroid, and gastric diseases. And less pronounced between WC and patterns of cardiometabolic and osteoarticular diseases.

In view of the above, high BMI and WC values impact the number of associated morbidities in the same individual. Therefore, in practice, monitoring the variables can contribute to the prevention of MM. In addition, the knowledge of the main disease patterns related to this increase can help public health policies and actions. The development of longitudinal studies on the subject is encouraged, aiming to contribute even more significantly to this field of research.

308

309 Limitations

A limitation of this study was the use of the MM variable with closed questions, yes or no, and not with medical data or medical records, leaving the perception of their health and/or diseases to the participants, which often may not mirror reality, due to the omission of certain conditions of morbidity. Although MM has been self-reported, it was worked with a preestablished list of morbidities, which does not allow considering other conditions related to mental health, very prevalent in MM studies (Prados-Torres et al., 2014; Violan et al., 2014; Vetrano et al., 2020) and other important chronic morbidities.

317 It is worth mentioning that the sample is not a true representation of the Brazilian318 population. Most participants were white, with incomes higher than the average for Brazilians,

319	and had a higher proportion of obesity and higher values for WC. In addition, the cross-sectional
320	design does not allow for affirming the causality of the associations.
321	
322	Conclusion
323	Both BMI and WC showed a positive association with MM in all MM categories and
324	were more expressive in men. The increase in BMI or WC was positively associated with the
325	accumulation of two or more morbidities. Of the five MM patterns found, the ones that showed
326	an association with BMI and WC were those of cardiometabolic and osteoarticular diseases.in

addition to these, those of gastric and thyroid diseases were associated with BMI. It would be

328 required additional longitudinal research that assesses the association of adiposity indicators and

329 MM patterns in different contexts and populations, especially when considering the diversity of

definitions and methods of MM evaluation and its patterns.

Therefore, this study highlights how the increase in BMI and WC impacts MM, and MM patterns, opening the way for a more careful look at BMI and WC, which are simple tools that can be used in the prevention and /or worsening of MM.

334 335

336 Acknowledgements

337 None.338

- 339
- 340
- 341
- 342
- 343
- 344

345 346

347

- 348
- 349
- 350

351 **References**

- ABESO. 2016. Diretrizes brasileiras de obesidade 2016/ABESO. DOI: 10.1590/S1415 52732000000100003.
- Agborsangaya CB, Ngwakongnwi E, Lahtinen M, Cooke T, Johnson JA. 2013. Multimorbidity
 prevalence in the general population: The role of obesity in chronic disease clustering. *BMC Public Health* 13. DOI: 10.1186/1471-2458-13-1161.
- 357 Van Den Akker M, Buntinx F, Knottnerus JA. 1996. Comorbidity or multimorbidity: What's in a
 358 name? A review of literature. *European Journal of General Practice* 2:65–70. DOI:
 359 10.3109/13814789609162146.
- Araujo MEA, Silva MT, Galvao TF, Nunes BP, Pereira MG. 2018. Prevalence and patterns of
 multimorbidity in Amazon Region of Brazil and associated determinants: A cross-sectional
 study. *BMJ Open* 8. DOI: 10.1136/bmjopen-2018-023398.
- 363 Asparouhov T, Muthén BO. 2010. Simple second order chi-square correction. :1–8.
- Barnett K, Mercer SW, Norbury M, Watt G, Wyke S, Guthrie B. 2012. Epidemiology of
 multimorbidity and implications for health care, research, and medical education: A crosssectional study. *The Lancet* 380:37–43. DOI: 10.1016/S0140-6736(12)60240-2.
- Bredella MA. 2017. Sex Differences in Body Composition. In: Mauvais-Jarvis F. (eds) Sex and
 Gender Factors Affecting Metabolic Homeostasis, Diabetes and Obesity. Advances in
 Experimental Medicine and Biology 1043:9–27.
- Brown T. 2006. Confirmatory Factor Analysis for Applied Research. *The American Statistician* 62:91–92. DOI: 10.1198/tas.2008.s98.
- De Carvalho JN, De Camargo Cancela M, Leandro D, De Souza B. 2018. Lifestyle factors and
 high body mass index are associated with different multimorbidity clusters in the Brazilian
 population. *PLoS ONE* 13:1–15. DOI: 10.1371/journal.pone.0207649.
- Catrysse L, van Loo G. 2017. Inflammation and the Metabolic Syndrome: The Tissue-Specific
 Functions of NF-κB. *Trends in Cell Biology* 27:417–429. DOI: 10.1016/j.tcb.2017.01.006.
- Celeste RK, Bastos JL. 2014. Mid-point for open-ended income category and the effect of
 equivalence scales on the income-health relationship. *Revista de Saude Publica* 47:168–
 171. DOI: 10.1590/S0034-8910.2013047004388.
- Chait A, Hartigh LJ den. 2020. Adipose Tissue Distribution, Inflammation and Its Metabolic
 Consequences, Including Diabetes and Cardiovascular Disease. *Frontiers in Cardiovascular Medicine* 7:1–41. DOI: 10.3389/fcvm.2020.00022.
- Christofoletti M, Streb AR, Del Duca GF. 2018. Body mass index as a predictor of
 multimorbidity in the Brazilian population. *Revista Brasileira de Cineantropometria e*
- 385 *Desempenho Humano* 20:555–565. DOI: 10.5007/1980-0037.2018v20n6p555.
- Chua YP, Xie Y, Lee PSS, Lee ES. 2021. Definitions and prevalence of multimorbidity in large
 database studies: A scoping review. *International Journal of Environmental Research and Public Health* 18:1–12. DOI: 10.3390/ijerph18041673.
- Diane Zheng D, Loewenstein DA, Christ SL, Feaster DJ, Lam BL, McCollister KE, Curiel-Cid
 RE, Lee DJ. 2021. Multimorbidity patterns and their relationship to mortality in the US
 older adult population. *PLoS ONE* 16:1–15. DOI: 10.1371/journal.pone.0245053.
- 392 Dragano NRV, Haddad-Tovolli R, Velloso LA. 2017. Leptin, Neuroinflammation and Obesity.
 393 Frontiers of Hormone Research 48:84–96. DOI: 10.1159/000452908.
- 394 Ezzati M. 2017. Excess weight and multimorbidity: putting people's health experience in risk
- factor epidemiology. *The Lancet Public Health* 2:e252–e253. DOI: 10.1016/S2468 2667(17)30093-2.

397 Faerstein E, Chor D, Lopes C de S, Werneck GL. 2005. Estudo Pró-Saúde: características gerais 398 e aspectos metodológicos. Revista Brasileira de Epidemiologia 8:454-466. DOI: 399 10.1590/s1415-790x2005000400014. 400 Ferrando PJ, Lorenzo-Seva U. 2017. 10 años del programa FACTOR: Una revisión crítica de sus 401 orígenes, desarrollo y líneas futuras. Psicothema 29:236-240. DOI: 10.7334/psicothema2016.304. 402 403 Ferrando PJ, Lorenzo-Seva U. 2018. Assessing the Quality and Appropriateness of Factor 404 Solutions and Factor Score Estimates in Exploratory Item Factor Analysis. Educational and 405 Psychological Measurement 78:762–780. DOI: 10.1177/0013164417719308. Flor LS, Campos MR, de Oliveira AF, Schramm JM de A. 2015. Diabetes burden in Brazil: 406 407 Fraction attributable to overweight, obesity, and excess weight. Revista de Saude Publica 49. DOI: 10.1590/S0034-8910.2015049005571. 408 409 Flores TR, Dos Santos Rodrigues AP, Neves RG, Batista SR, Da Cruz Teixeira DS, Da Silveira EA, Malta DC, Nunes BP. 2021. The risk of multimorbidity associated with overweight and 410 obesity: Data from the brazilian national health survey 2013. Journal of Obesity and 411 412 Metabolic Syndrome 30:155–162. DOI: 10.7570/JOMES20110. 413 Hair, J. F., Black, W. C., Babin, B. J., Anderson, R. E., & Tatham RL. 2009. Análise 414 multivariada de dados. Hall JE, Do Carmo JM, Da Silva AA, Wang Z, Hall ME. 2015. Obesity-Induced Hypertension: 415 416 Interaction of Neurohumoral and Renal Mechanisms. Circulation Research 116:991–1006. 417 DOI: 10.1161/CIRCRESAHA.116.305697. Hudon C, Fortin M, Vanasse A. 2005. Cumulative Illness Rating Scale was a reliable and valid 418 419 index in a family practice context. Journal of Clinical Epidemiology 58:603-608. DOI: 420 10.1016/j.jclinepi.2004.10.017. 421 Jantsch AG, Alves RFS, Faerstein E. 2018. Educational inequality in rio de janeiro and its 422 impact on multimorbidity: Evidence from the pró-saúde study. a cross-sectional analysis. Sao Paulo Medical Journal 136:51-58. DOI: 10.1590/1516-3180.2017.0209100917. 423 424 Jawed M, Inam S, Shah N, Shafique K. 2020. Association of obesity measures and 425 multimorbidity in Pakistan: findings from the IMPACT study. Public Health 180:51–56. 426 DOI: 10.1016/j.puhe.2019.10.017. 427 Jovic D, Marinkovic J, Vukovic D. 2016. Association between body mass index and prevalence 428 of multimorbidity: a cross-sectional study. Public Health 139:103-111. DOI: 429 10.1016/j.puhe.2016.05.014. Kivimäki M, Kuosma E, Ferrie JE, Luukkonen R, Nyberg ST, Alfredsson L, Batty GD, Brunner 430 EJ, Fransson E, Goldberg M, Knutsson A, Koskenvuo M, Nordin M, Oksanen T, Pentti J, 431 Rugulies R, Shipley MJ, Singh-Manoux A, Steptoe A, Suominen SB, Theorell T, Vahtera J, 432 Virtanen M, Westerholm P, Westerlund H, Zins M, Hamer M, Bell JA, Tabak AG, Jokela 433 434 M. 2017. Overweight, obesity, and risk of cardiometabolic multimorbidity: pooled analysis 435 of individual-level data for 120 813 adults from 16 cohort studies from the USA and Europe. The Lancet Public Health 2:e277–e285. DOI: 10.1016/S2468-2667(17)30074-9. 436 Leal Neto J de S, Barbosa AR, Meneghini V. 2016. Diseases and chronic health conditions, 437 438 multimorbidity and body mass index in older adults. Revista Brasileira de Cineantropometria e Desempenho Humano 18:509-519. DOI: 10.5007/1980-439 440 0037.2016v18n5p509. 441 Lohman TG, Roche AF MR. 1988. Anthropometric standardization reference manual. 442 Champaign: Human Kinetics Book.

443 MacMahon S, The Academy of Medical Sciences. 2018. Multimorbidity: a priority for global 444 health research. Academy of Medical Sciences. Melo ME de. 2011. Doenças Desencadeadas ou Agravadas pela Obesidade. Associação 445 446 Brasileira para o Estudo da Obesidade e da Síndrome Metabólica (ABESO):1–10. 447 Murray CJL, Barber RM, Foreman KJ, Ozgoren AA, Abd-Allah F, Abera SF, Aboyans V, Abraham JP, Abubakar I, Abu-Raddad LJ, Abu-Rmeileh NM, Achoki T, Ackerman IN, 448 449 Ademi Z, Adou AK, Adsuar JC, Afshin A, Agardh EE, Alam SS, Alasfoor D, Albittar MI, 450 Alegretti MA, Alemu ZA, Alfonso-Cristancho R, Alhabib S, Ali R, Alla F, Allebeck P, Almazroa MA, Alsharif U, Alvarez E, Alvis-Guzman N, Amare AT, Ameh EA, Amini H, 451 452 Ammar W, Anderson HR, Anderson BO, Antonio CAT, Anwari P, Arnlöv J, Arsenijevic 453 VSA, Artaman A, Asghar RJ, Assadi R, Atkins LS, Avila MA, Awuah B, Bachman VF, Badawi A, Bahit MC, Balakrishnan K, Banerjee A, Barker-Collo SL, Barquera S, Barregard 454 L, Barrero LH, Basu A, Basu S, Basulaiman MO, Beardsley J, Bedi N, Beghi E, Bekele T, 455 Bell ML, Benjet C, Bennett DA, Bensenor IM, Benzian H, Bernabé E, Bertozzi-Villa A, 456 Beyene TJ, Bhala N, Bhalla A, Bhutta ZA, Bienhoff K, Bikbov B, Biryukov S, Blore JD, 457 Blosser CD, Blyth FM, Bohensky MA, Bolliger IW, Basara BB, Bornstein NM, Bose D, 458 459 Boufous S, Bourne RRA, Boyers LN, Brainin M, Brayne CE, Brazinova A, Breitborde 460 NJK, Brenner H, Briggs AD, Brooks PM, Brown JC, Brugha TS, Buchbinder R, Buckle GC, Budke CM, Bulchis A, Bulloch AG, Campos-Nonato IR, Carabin H, Carapetis JR, 461 462 Cárdenas R, Carpenter DO, Caso V, Castañeda-Orjuela CA, Castro RE, Catalá-López F, 463 Cavalleri F, Cavlin A, Chadha VK, Chang JC, Charlson FJ, Chen H, Chen W, Chiang PP, Chimed-Ochir O, Chowdhury R, Christensen H, Christophi CA, Cirillo M, Coates MM, 464 465 Coffeng LE, Coggeshall MS, Colistro V, Colquhoun SM, Cooke GS, Cooper C, Cooper LT, Coppola LM, Cortinovis M, Criqui MH, Crump JA, Cuevas-Nasu L, Danawi H, Dandona 466 L, Dandona R, Dansereau E, Dargan PI, Davey G, Davis A, Davitoiu D V., Dayama A, De 467 468 Leo D, Degenhardt L, Del Pozo-Cruz B, Dellavalle RP, Deribe K, Derrett S, Des Jarlais DC, Dessalegn M, Dharmaratne SD, Dherani MK, Diaz-Torné C, Dicker D, Ding EL, 469 470 Dokova K, Dorsey ER, Driscoll TR, Duan L, Duber HC, Ebel BE, Edmond KM, Elshrek 471 YM, Endres M, Ermakov SP, Erskine HE, Eshrati B, Esteghamati A, Estep K, Faraon EJA, Farzadfar F, Fav DF, Feigin VL, Felson DT, Fereshtehnejad SM, Fernandes JG, Ferrari AJ, 472 Fitzmaurice C, Flaxman AD, Fleming TD, Foigt N, Forouzanfar MH, Fowkes FGR, Paleo 473 474 UF, Franklin RC, Fürst T, Gabbe B, Gaffikin L, Gankpé FG, Geleijnse JM, Gessner BD, 475 Gething P, Gibney KB, Giroud M, Giussani G, Dantes HG, Gona P, González-Medina D, Gosselin RA, Gotay CC, Goto A, Gouda HN, Graetz N, Gugnani HC, Gupta R, Gupta R, 476 Gutiérrez RA, Haagsma J, Hafezi-Nejad N, Hagan H, Halasa YA, Hamadeh RR, Hamavid 477 H. Hammami M. Hancock J. Hankey GJ. Hansen GM. Hao Y. Harb HL. Haro JM. 478 Havmoeller R, Hay SI, Hay RJ, Heredia-Pi IB, Heuton KR, Heydarpour P, Higashi H, Hijar 479 480 M, Hoek HW, Hoffman HJ, Hosgood HD, Hossain M, Hotez PJ, Hov DG, Hsairi M, Hu G, 481 Huang C, Huang JJ, Husseini A, Huynh C, Iannarone ML, Iburg KM, Innos K, Inoue M, Islami F, Jacobsen KH, Jarvis DL, Jassal SK, Jee SH, Jeemon P, Jensen PN, Jha V, Jiang G, 482 Jiang Y, Jonas JB, Juel K, Kan H, Karch A, Karema CK, Karimkhani C, Karthikeyan G, 483 484 Kassebaum NJ, Kaul A, Kawakami N, Kazanjan K, Kemp AH, Kengne AP, Keren A, Khader YS, Khalifa SEA, Khan EA, Khan G, Khang YH, Kieling C, Kim D, Kim S, Kim 485 Y. Kinfu Y. Kinge JM, Kivipelto M, Knibbs LD, Knudsen AK, Kokubo Y, Kosen S. 486 487 Krishnaswami S, Defo BK, Bicer BK, Kuipers EJ, Kulkarni C, Kulkarni VS, Kumar GA, 488 Kyu HH, Lai T, Lalloo R, Lallukka T, Lam H, Lan Q, Lansingh VC, Larsson A,

489 Lawrynowicz AEB, Leasher JL, Leigh J, Leung R, Levitz CE, Li B, Li Y, Li Y, Lim SS, 490 Lind M, Lipshultz SE, Liu S, Liu Y, Lloyd BK, Lofgren KT, Logroscino G, Looker KJ, Lortet-Tieulent J, Lotufo PA, Lozano R, Lucas RM, Lunevicius R, Lyons RA, Ma S, 491 492 Macintyre MF, Mackay MT, Majdan M, Malekzadeh R, Marcenes W, Margolis DJ, 493 Margono C, Marzan MB, Masci JR, Mashal MT, Matzopoulos R, Mayosi BM, Mazorodze 494 TT, McGill NW, McGrath JJ, McKee M, McLain A, Meaney PA, Medina C, Mehndiratta 495 MM, Mekonnen W, Melaku YA, Meltzer M, Memish ZA, Mensah GA, Meretoja A, 496 Mhimbira FA, Micha R, Miller TR, Mills EJ, Mitchell PB, Mock CN, Ibrahim NM, Mohammad KA, Mokdad AH, Mola GLD, Monasta L, Hernandez JCM, Montico M, 497 498 Montine TJ, Mooney MD, Moore AR, Moradi-Lakeh M, Moran AE, Mori R, Moschandreas 499 J, Moturi WN, Moyer ML, Mozaffarian D, Msemburi WT, Mueller UO, Mukaigawara M, Mullany EC, Murdoch ME, Murray J, Murthy KS, Naghavi M, Naheed A, Naidoo KS, 500 Naldi L, Nand D, Nangia V, Narayan KMV, Nejjari C, Neupane SP, Newton CR, Ng M, 501 502 Ngalesoni FN, Nguyen G, Nisar MI, Nolte S, Norheim OF, Norman RE, Norrving B, Nyakarahuka L, Oh IH, Ohkubo T, Ohno SL, Olusanya BO, Opio JN, Ortblad K, Ortiz A, 503 504 Pain AW, Pandian JD, Panelo CIA, Papachristou C, Park EK, Park JH, Patten SB, Patton 505 GC, Paul VK, Pavlin BI, Pearce N, Pereira DM, Perez-Padilla R, Perez-Ruiz F, Perico N, 506 Pervaiz A, Pesudovs K, Peterson CB, Petzold M, Phillips MR, Phillips BK, Phillips DE, Piel FB, Plass D, Poenaru D, Polinder S, Pope D, Popova S, Poulton RG, Pourmalek F, 507 508 Prabhakaran D, Prasad NM, Pullan RL, Qato DM, Quistberg DA, Rafay A, Rahimi K, 509 Rahman SU, Raju M, Rana SM, Razavi H, Reddy KS, Refaat A, Remuzzi G, Resnikoff S, Ribeiro AL, Richardson L, Richardus JH, Roberts DA, Rojas-Rueda D, Ronfani L, Roth 510 511 GA, Rothenbacher D, Rothstein DH, Rowley JT, Roy N, Ruhago GM, Saeedi MY, Saha S, Sahraian MA, Sampson UKA, Sanabria JR, Sandar L, Santos IS, Satpathy M, Sawhney M, 512 Scarborough P, Schneider IJ, Schöttker B, Schumacher AE, Schwebel DC, Scott JG, Seedat 513 514 S, Sepanlou SG, Serina PT, Servan-Mori EE, Shackelford KA, Shaheen A, Shahraz S, Levy TS, Shangguan S, She J, Sheikhbahaei S, Shi P, Shibuya K, Shinohara Y, Shiri R, Shishani 515 516 K, Shiue I, Shrime MG, Sigfusdottir ID, Silberberg DH, Simard EP, Sindi S, Singh A, 517 Singh JA, Singh L, Skirbekk V, Slepak EL, Sliwa K, Soneji S, Søreide K, Soshnikov S, Sposato LA, Sreeramareddy CT, Stanaway JD, Stathopoulou V, Stein DJ, Stein MB, Steiner 518 C, Steiner TJ, Stevens A, Stewart A, Stovner LJ, Stroumpoulis K, Sunguya BF, 519 520 Swaminathan S, Swaroop M, Sykes BL, Tabb KM, Takahashi K, Tandon N, Tanne D, 521 Tanner M, Tavakkoli M, Taylor HR, Te Ao BJ, Tediosi F, Temesgen AM, Templin T, Ten Have M, Tenkorang EY, Terkawi AS, Thomson B, Thorne-Lyman AL, Thrift AG, Thurston 522 GD, Tillmann T, Tonelli M, Topouzis F, Toyoshima H, Traebert J, Tran BX, Trillini M, 523 524 Truelsen T, Tsilimbaris M, Tuzcu EM, Uchendu US, Ukwaja KN, Undurraga EA, Uzun SB, Van Brakel WH, Van De Vijver S, Van Gool CH, Van Os J, Vasankari TJ, 525 526 Venketasubramanian N, Violante FS, Vlassov V V., Vollset SE, Wagner GR, Wagner J, 527 Waller SG, Wan X, Wang H, Wang J, Wang L, Warouw TS, Weichenthal S, Weiderpass E, Weintraub RG, Wenzhi W, Werdecker A, Westerman R, Whiteford HA, Wilkinson JD, 528 529 Williams TN, Wolfe CD, Wolock TM, Woolf AD, Wulf S, Wurtz B, Xu G, Yan LL, Yano 530 Y, Ye P, Yentür GK, Yip P, Yonemoto N, Yoon SJ, Younis MZ, Yu C, Zaki ME, Zhao Y, Zheng Y, Zonies D, Zou X, Salomon JA, Lopez AD, Vos T. 2015. Global, regional, and 531 national disability-adjusted life years (DALYs) for 306 diseases and injuries and healthy life 532 533 expectancy (HALE) for 188 countries, 1990-2013: Quantifying the epidemiological 534 transition. The Lancet 386:2145–2191. DOI: 10.1016/S0140-6736(15)61340-X.

Peer.

Nájera Medina O, Paulina Rodríguez-López C, Cristina González-Torres M, Cruz-Bautista I,
Nájera-Medina O. 2019. Nutrición Hospitalaria Trabajo Original Obesidad y síndrome
metabólico Visceral obesity, skeletal muscle mass and resistin in metabolic syndrome
development Obesidad visceral, masa musculoesquelética y resistina en el desarrollo de

539 síndrome metabólico. *Nutr Hosp* 36:43–50. DOI: 10.20960/nh.1889.

- Nguyen H, Manolova G, Daskalopoulou C, Vitoratou S, Prince M, Prina AM. 2019. Prevalence
 of multimorbidity in community settings: A systematic review and meta-analysis of
 observational studies. *Journal of Comorbidity* 9:2235042X1987093. DOI:
- 54310.1177/2235042x19870934.
- Petarli GB, Cattafesta M, Sant'Anna MM, de Paula Alves Bezerra OM, Zandonade E, Salaroli
 LB. 2019. Multimorbidity and complex multimorbidity in Brazilian rural workers. *PLoS ONE* 14:1–17. DOI: 10.1371/journal.pone.0225416.
- 547 Prados-Torres A, Calderón-Larrañaga A, Hancco-Saavedra J, Poblador-Plou B, Van Den Akker
 548 M. 2014. Multimorbidity patterns: A systematic review. *Journal of Clinical Epidemiology*549 67:254–266. DOI: 10.1016/j.jclinepi.2013.09.021.
- Prados-Torres A, Poblador-Plou B, Calderón-Larrañaga A, Gimeno-Feliu LA, González-Rubio
 F, Poncel-Falcó A, Sicras-Mainar A, Alcalá-Nalvaiz JT. 2012. Multimorbidity patterns in
 primary care: Interactions among chronic diseases using factor analysis. *PLoS ONE* 7. DOI: 10.1371/journal.pone.0032190.
- Reyes CM, M Leyland, K, Peat G, Cooper C, K Arden N and, Prieto-Alhambra D. 2016.
 Association between overweight and obesity and risk of clinically diagnosed knee, hip, and hand osteoarthritis: a population-based cohort study. *Arthritis & Rheumatology* 68:1869– 1875. DOI: https://dx.doi.org/10.1002%2Fart.39707.
- Samper-Ternent R, Al Snih S. 2012. Obesity in older adults: Epidemiology and implications for
 disability and disease. *Reviews in Clinical Gerontology* 22:10–34. DOI:
 10.1017/S0959259811000190.
- Schorr M, Dichtel LE, Gerweck A V., Valera RD, Torriani M, Miller KK, Bredella MA. 2018.
 Sex differences in body composition and association with cardiometabolic risk. *Biology of* Sex Differences 9. DOI: 10.1186/s13293-018-0189-3.
- Seravalle G, Grassi G. 2017. Obesity and hypertension. *Pharmacological Research* 122:1–7.
 DOI: 10.1016/j.phrs.2017.05.013.
- Song X, Tabák AG, Zethelius B, Yudkin JS, Söderberg S, Laatikainen T, Stehouwer CDA,
 Dankner R, Jousilahti P, Onat A, Nilsson PM, Satman I, Vaccaro O, Tuomilehto J, Qiao Q.
 2014. Obesity attenuates gender differences in cardiovascular mortality. *Cardiovascular*
- 569 *Diabetology* 13:1–10. DOI: 10.1186/s12933-014-0144-5.
- Song RH, Wang B, Yao QM, Li Q, Jia X, Zhang JA. 2019. The Impact of Obesity on Thyroid
 Autoimmunity and Dysfunction: A Systematic Review and Meta-Analysis. *Frontiers in Immunology* 10:1–11. DOI: 10.3389/fimmu.2019.02349.
- Tang XN, Liebeskind DS, Towfighi A. 2017. The Role of Diabetes, Obesity, and Metabolic
 Syndrome in Stroke. *Seminars in Neurology* 37:267–273. DOI: 10.1055/s-0037-1603753.
- Valderas JM, Starfi B, Sibbald B. 2009. Understanding Health and Health Services. *Annals Of Family Medicine*:357–363. DOI: 10.1370/afm.983.Martin.
- 577 Vetrano DL, Rizzuto D, Calderón-Larrañaga A, Onder G, Welmer AK, Bernabei R, Marengoni
 578 A, Fratiglioni L. 2018. Trajectories of functional decline in older adults with
- 579 neuropsychiatric and cardiovascular multimorbidity: A Swedish cohort study. *PLoS*
- 580 *Medicine* 15:1–15. DOI: 10.1371/journal.pmed.1002503.

- Vetrano DL, Rizzuto D, Calderón-Larrañaga A, Onder G, Welmer AK, Qiu C, Bernabei R,
 Marengoni A, Fratiglioni L. 2019. Walking Speed Drives the Prognosis of Older Adults
 with Cardiovascular and Neuropsychiatric Multimorbidity. *American Journal of Medicine*122:1207-1215 c (DOI: 10.1016/j.amira.d.2010.05.005)
- 584 132:1207-1215.e6. DOI: 10.1016/j.amjmed.2019.05.005.
- Vetrano DL, Roso-Llorach A, Fernández S, Guisado-Clavero M, Violán C, Onder G, Fratiglioni
 L, Calderón-Larrañaga A, Marengoni A. 2020. Twelve-year clinical trajectories of
 multimorbidity in a population of older adults. *Nature Communications* 11. DOI:
- 588 10.1038/s41467-020-16780-x.
- Violan C, Foguet-Boreu Q, Flores-Mateo G, Salisbury C, Blom J, Freitag M, Glynn L, Muth C,
 Valderas JM. 2014. Prevalence, determinants and patterns of multimorbidity in primary
 care: A systematic review of observational studies. *PLoS ONE* 9:3–11. DOI:
 10.1371/journal.pone.0102149.
- 593 WHO. 2000. Obesity: preventing and managing the global epidemic. Report of a WHO
 594 consultation. *World Health Organization Technical Report Series*.
- Willadsen TG, Bebe A, Køster-Rasmussen R, Jarbøl DE, Guassora AD, Waldorff FB, Reventlow
 S, Olivarius N de F. 2016. The role of diseases, risk factors and symptoms in the definition
 of multimorbidity a systematic review. *Scandinavian Journal of Primary Health Care*34:112–121. DOI: 10.3109/02813432.2016.1153242.
- Xu X, Mishra GD, Jones M. 2017. Evidence on multimorbidity from definition to intervention:
 An overview of systematic reviews. *Ageing Research Reviews* 37:53–68. DOI:
 10.1016/j.arr.2017.05.003.
- Zhang J, Xu L, Li J, Sun L, Qin W. 2020. Association between obesity-related anthropometric
 indices and multimorbidity among older adults in Shandong, China: a cross-sectional study.
 BMJ open 10:e036664. DOI: 10.1136/bmjopen-2019-036664.

605



Table 1(on next page)

Table 1 – Characteristics of the participants (2,698) of the *Pró-Saúde* Study, Rio de Janeiro - Brazil (2012–2013).

Variables			Total	
variables			%	
$C_{\text{ondor}}(n-2608)$	Male	1178	43.7	
Gender (n=2070)	Female	1520	56.3	
	<40 years	272	10.1	
	40 a 49 years	901	33.4	
Age (years) (n=2698)	50 a 59 years	1056	39.1	
	60 a 69 years	401	14.8	
	70 a 79 years	68	2.5	
	University or more	1429	53.3	
Education (2680)	High School	940	35.1	
	Elementary School	311	11.6	
	Married/stable union	1645	61.3	
Marital status (n=7683)	Separated/ divorced	465	17.3	
Maritar status (n=2003)	Widowed	151	5.6	
	Single	422	15.7	
	Black/ Brown	1302	48.7	
R ace $(n=2674)$	White	1340	50.1	
Race (11-20/4)	Yellow	16	0.6	
	Indigenous	16	0.6	
	Into up to 3	718	26.8	
Modified equivalent income by OCDE (MW) (n=2673)	From the 3 to 6	1039	38.9	
	More than 6	916	34.3	
	Underweight/eutrophic	791	30.3	
BMI (Kg/m ²) (n=2608)	Overweight	1031	39.5	
	Obesity	786	30.2	
Abdominal obseits $(n-2)(1)$	Yes	2224	83.6	
Addominal obesity (n=2001)	No	437	16.4	
	Smoker	341	12.8	
Smoking (n=2678)	ex-smoker	721	26.9	
	Non-smoker	1616	60.3	
Deviced estimates in the last 2 modes $(n-2)(90)$	yes	1149	42.9	
Physical activity in the last 2 weeks (n=2080)	No	1531	57.1	
	0 to 1 morbidity	1063	39.4	
	2 morbidities	614	22.8	
Number of morbidities (n=2698)	3 morbidities	463	17.2	
	More than 4 morbidities	558	20.7	
	Yes	1057	39.4	
Consumption of truits (≥ 4 times a week) (n=2684)	No	1627	60.6	
	Yes	1613	60.1	
Consumption of vegetables (≥ 4 times a week) (n=2685)	No	1072	39.9	

*Mean (SD): age 51.5 (8.96); BMI 27.92 (5.20); WC 96.33 (12.88)

**Median of morbidities 2.0 (interquartile range: 2.0-3.0)

1



Table 2(on next page)

Odds ratio adjusted and 95% confidence interval (CI) of the association between BMI and WC and the multimorbidity categories.

	BMI			WC		
Number of	OR adjusted (IC95%)			OR adjusted (IC95%)		
morbidities	Total	Female	Male	Total	Female	Male
0-1	-	-	-	-	-	-
	5,05	3.99	5.43	5.54	4.75	6.49
2	(3.82-6,29)	(2.87-5.12)	(4.98-6.79)	(4.25-6.84)	(3.52-5.98)	(5.08-7.89)
	6.07	5.00	6.56	6.55	5.76	7.57
3	(4.83-7.31)	(3.87-6.13)	(5.24-7.88)	(5.25-7.85)	(4.52-7.00)	(6.14-8.99)
Marrie theory 4	7	5.98	7.48	7.48	6.74	8.48
Nore than 4	(5.75-8.25)	(4.84-7.13)	(6.14-8.18)	(6.16-8.78)	(5.48-7.99)	(7.64-9.29)

Table 2 – Odds ratio adjusted and 95% confidence interval (CI) of the association between BMI and WC and the multimorbidity categories.

p value<0,001 for all variables; * adjusted model to age, race, income, education, marital status and consumption of fruit and vegetables; number of morbities 0-1= no MM, the reference category.

Table 3(on next page)

Factor loads of morbidities.

VARIABLE	F1	F2	F3	F4	F5
Asthma	0.536	0.198	0.141	-0.071	-0.039
Emphysema	1.048	-0.019	0.000	0.018	0.021
RSI	0.016	0.562	-0.003	-0.025	0.006
Arthrosis	-0.011	0.542	-0.071	0.148	0.026
Slipped disc	-0.001	0.505	0.020	0.041	0.029
Hyperthyroidism	-0.072	-0.043	0.655	0.163	-0.172
Hipothyroidism	0.068	0.008	0.839	-0.050	0.069
Hypertension	0.128	0.059	-0.119	0.660	-0.098
DM	0.211	-0.039	-0.014	0.616	-0.079
Hypercholesterolemia Acute myocardial	0.179	0.118	0.132	0.346	0.047
infarction	-0.189	0.026	0.177	0.715	0.112
Angina	-0.022	0.042	0.012	0.711	0.175
Stroke	-0.017	0.088	-0.030	0.526	-0.168
Peptic ulcer	-0.008	-0.035	-0.018	0.071	0.914
Gastritis	0.073	0.155	0.028	-0.095	0.661
Nephrolithiasis	0.014	0.164	-0.161	0.104	0.006
Cholecystitis	0.046	0.200	0.021	0.151	0.078
Tuberculosis	0.176	-0.257	-0.277	0.020	0.079
Composite reliability	0.794	0.565	0.720	0.772	0.773
H index	1.100	0.641	0.783	0.840	0.867



Table 4(on next page)

Odds ratio and 95% confidence interval (CI) of the association of multimorbidity patterns with BMI and WC.

1	Table 4 - Odds ratio and 95% confidence interval (CI) of the association of multimorbidity patterns with BMI and
2	WC.

	BM	II	WC		
Detterns of MMI	Unadjusted OR	Adjusted OR	Unadjusted	Adjusted OR	
	(IC 95%)	(IC 95%)	OR (IC 95%)	(IC 95%)	
Pattern 1 – Respiratory	1.04	1.04	1.01	1.01	
diseases ²	(0.98-1.09)	(0.98-1.10)	(0.99-1.04)	(0.99-1.04)	
Pattern 2 – Osteoarticular	1.06	1.03	1.02	1.02	
diseases ³	(1.05 - 1.09)	(1.02 - 1.04)	(1.01 - 1.02)	(1.01 - 1.03)	
Dattorn 2 Thyraid diagonast	0.96	1.09	0.98	0.98	
Pattern 5 – Thyloid diseases	(0.88-1.04)	(1.04 - 1.14)	(0.95 - 1.01)	(0.95 - 1.01)	
Pattern 4 – Cardiometabolic	1.10	1.07	1.04	1.04	
diseases ⁵	(1.08-1.12)	(1.06-1.09)	(1.03 - 1.05)	(1.03-1.04)	
Dettern 5 Costria disassas	1.00	1.03	1.00	0.99	
Pattern 5 – Gastric diseases	(0.96-1.03)	(1.01 - 1.05)	(0.98-1.01)	(0.98-1.01)	

3 1 Total of participants who had MM in one of the patterns 1264 (43%); 2. Pattern 1 (n=36) adjusted model to consumption of fruits and vegetables and physical activity; 3. Pattern

4 5 2 (n=384) adjusted model to gender, age, income and race; 4. Pattern 3 (n=25) adjusted model to gender, age, consumption of fruits and vegetables and physical activity; 5. Pattern

4 (n=690) adjusted model to income, marital status, race and education.; 6. Pattern 5 (n=129) adjusted model to gender, age, consumption of fruits and physical activity.

6