

## Review Article

# Anthropometric Parameters in Celiac Disease: A Review on the Different Evaluation Methods and Disease Effects

Allysson Costa  and Gleisson A. P. Brito

*Laboratory of Physiology and Developmental Biology, Federal University of Latin American Integration—UNILA, Foz do Iguaçu, Paraná, Brazil*

Correspondence should be addressed to Allysson Costa; [allyssoncostaa@gmail.com](mailto:allyssoncostaa@gmail.com)

Received 17 April 2019; Revised 3 August 2019; Accepted 20 August 2019; Published 9 September 2019

Academic Editor: Pedro Moreira

Copyright © 2019 Allysson Costa and Gleisson A. P. Brito. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

This review compiled anthropometric data from 29 original articles, published between 1995 and 2015, corresponding to a total sample of 6368 celiac disease subjects. Body mass index was the main parameter for measuring anthropometry (82.1%), followed by body mass (78.6%), body fat (51.7%), bone mineral density and bone mineral content (46.4%), and fat-free mass (44.8%). The main evaluation method was dual x-ray absorptiometry (83.3%), followed by bioimpedance (16.6%), skinfold thickness (16.6%), and isotope dilution (5.5%). This compilation suggests that celiac disease patients without a gluten-free diet (WGFD) and celiac disease patients with a gluten-free diet (GFD) show a lower body mass than the control group, with inconclusive data about WGFD versus GFD. Body mass index is lower in WGFD and GFD compared to control group, and is lower in WGFD compared to GFD. We observed lower values of FM and FFM in WGFD and GFD versus the control group. No difference was found between WGFD versus GFD. BMD and BMC are lower in WGFD versus GFD and GFD versus the control group, with inconclusive data about WGFD versus GFD. The findings of this review suggest that celiac disease patients must be periodically evaluated through anthropometric parameters, since the pathology has the potential to modulate such values even in a gluten-free diet, with these variables reflecting their healthy status. In parallel, the screening of different anthropometric assessment methodologies can provide support for more accurate evaluations by scientists and clinical professionals who work with celiac disease patients.

## 1. Introduction

**1.1. Celiac Disease.** Celiac disease is a chronic autoimmune illness that manifests itself in individuals according to a genetic predisposition with environmental interaction [1]. It is characterized by an inflammatory condition as a consequence of the body's difficulty to process gluten proteins from wheat, barley, and rye [2–4]. Epidemiological research reveals a prevalence of 1:100 (1%) in the United States population, with a variation between 1:80 (1.25%) and 1:140 (0.71%) [3]. A previous review by Fasano et al. [5] estimated that celiac disease is one of the most frequent genetic disorders, affecting 0.5% to 1% of the world population. However, its diagnosis can be outdated, since its clinical presentation overlaps with other more common conditions.

Celiac disease manifests itself clinically in five ways: (1) classic: small bowel mucosal malabsorption, chronic diarrhea, abdominal distension, abdominal pain, weight loss, and flatulence; (2) atypical: the most common display of the disease, in which there is an absence of or few gastrointestinal symptoms, iron deficiency anemia, osteoporosis or osteopenia, infertility, and short stature; (3) silent: asymptomatic, with a casual, histological, or serological diagnosis; (4) latent: (A) individuals who are responsive to a gluten-free diet with a normal histology and elevated intraepithelial lymphocytes; (B) normal small bowel mucosa, without restriction to gluten, with subsequent development of celiac disease; (5) refractory: patients with celiac disease who do not respond to a gluten-free diet [4].

Each manifestation has its own characteristics, from gastrointestinal symptoms [6] to metabolic alterations [7, 8]

and anthropometric changes, [9–11] mostly due to the unsatisfactory absorption of nutrients as a consequence of small bowel inflammation [6, 12]. The diagnosis of celiac disease is based on clinical manifestations and serological and histological laboratory tests from small bowel biopsies [13]. It is accepted that serological markers from tissue antitransglutaminase antibodies (TtG), immunoglobulin A (IgA), and antiendomysium are sensitive and specific to the initial celiac disease diagnosis [3, 14]. There is good evidence of a relationship between mucosal villi atrophies in the small intestine and the histopathological characteristics of the disease, and for this reason, a duodenal biopsy is recommended for diagnosis confirmation [14]. The only treatment for celiac disease is a gluten-free diet [2, 5], and patients with good adherence to it may present anthropometric values similar to healthy subjects [15]. However, other studies suggest that, compared to the control group, celiac patients with adherence to a gluten-free diet may still present decreased values in anthropometric parameters [9, 11, 16–18].

Considering the impact of celiac disease on metabolism and body composition, we now proceed to review these subjects and analyze data from experimental and epidemiological research.

**1.2. Metabolic and Anthropometric Alterations in Celiac Disease Patients.** The immunological process of celiac disease, triggered by gluten, leads to a chronic inflammatory response, resulting in lesions associated with atrophy in the small bowel mucosal villi [2], that results in unsatisfactory nutrient absorption, including fatty acids, iron, transferrin, glucose, electrolytes, vitamins, folic acid, and calcium [17, 19, 20]. As a consequence, short stature, muscular fatigue, iron deficiency anemia, hypovitaminosis (A, B12, D, K), muscle fatigue [6, 9, 17], bone mineral content reduction [10, 21], and osteoporosis, [22] among other intestinal symptoms [6], may occur. Furthermore, celiac disease patients oxidize more carbohydrates as an energy source, probably due to the insufficient absorption of nutrients by the small bowel mucosa [9]. In Bode et al.' study [16], celiac disease patients with good adherence to gluten-free diet, median age 42, compared to healthy subjects, had a lower body mass index, a lower body fat percentage, and a lower bone mineral content in the spine and forearms. González et al. [17] analyzed women with celiac disease, between 20 and 60 years, with or without a gluten-free diet, and compared them to a control group of women of the same age. Women with celiac disease, in both groups, had a lower body mass index and height compared to the control group, and women with celiac disease without a gluten-free diet presented a lower body mass index than those on a gluten-free diet, as well as a lower body fat and fat-free mass than the control group. Capristo et al. [9], when analyzing adult patients with celiac disease, with a mean age of  $29.9 \pm 7.6$  years, found that those on a gluten-free diet presented a lower body mass, fat, and fat-free mass than the control group, with such parameters even lower in individuals without a gluten-free diet when compared to the control group. The research of Bardella et al. [18], with 71

celiac disease individuals without a gluten-free diet, between 17 to 58 years, verified that height in men, body mass, and body mass index in men and women were significantly lower than the control group. Brambilla et al. [11] studied 150 celiac disease children without a gluten-free diet and found lower values for body mass and body mass index when compared to the control groups.

Celiac disease can be effectively treated through a gluten-free diet [2, 5]. In the research by Barera et al. [23], individuals diagnosed with celiac disease revealed a lower body mass, fat mass, fat-free mass, and bone mineral content when compared to the control group, but after one year on a gluten-free diet, anthropometric values were similar to healthy subjects. However, other evidence suggests that even with a gluten-free diet, patients with celiac disease can present lower anthropometric values, [9, 11, 16–18] in addition to nutritional deficiencies, such as lower levels of B6 and B12 vitamins, folic acid, iron, and transferrin [9, 24].

## 2. Methodology

For this review, original articles written in English were selected through the Google Scholar search engine, using the keywords: celiac disease, body composition, anthropometry, anthropometric, body mass, body mass index, lean mass, fat mass, fat-free mass, bone mineral content, and bone mineral density. The articles were published between 1995 and 2015, covering a period of 20 years.

The inclusion criterion used was the presence of anthropometric data in celiac disease patients. The search returned 29 original articles, corresponding to a total sample of 6368 subjects with celiac disease. All data were included for review and discussion.

Data were compiled according to evaluated anthropometric parameters, the sex and age of the sample, and the methodology undertaken. Data from the anthropometric parameters were distributed in 3 groups: celiac disease patients without a gluten-free diet (WGFD), celiac disease patients with a gluten-free diet (GFD), and the control group.

## 3. Results

**3.1. Measured Parameters and Sample Characteristics.** In the reviewed studies, an analysis of body mass index (BMI) was present in 82.1%, body mass (BM) in 78.6%, fat mass in 54.7%, fat-free mass in 44.8%, and bone mineral density (BMD) and bone mineral content (BMC) in 46.4% of the studies (Table 1).

Table 2 compiles the sample data of age and sex and the presence of control groups in the reviewed studies. A prevalence of adults, 3197, in relation to children and adolescents, 443, was verified in the proportion of 7:1, with more female adult (52.9%) than male adult (47.1%) patients. In children and adolescents, we are also found more females (52.2%) than males (47.8%). In the control groups adults were more prevalent, 70.5%, with children and adolescents representing 29.5%.

TABLE 1: Anthropometric parameters measured in studies with celiac patients.

Scientific publication	BM	BMI	BMC BMD	BF	FFM
Bode et al. [25]	X	X	X	X	
Bardella et al. [26]	X	X			
Gonzalez et al. [27]	X	X	X	X	X
Rea et al. [28]	X	X	X	X	X
Mautalen et al. [29]			X		
Smecuol et al. [30]	X		X	X	X
Dickey and Bodkin [31]	X	X			
De Lorenzo et al. [32]	X	X	X	X	X
Bardella et al. [18]	X	X	X	X	X
Barera et al. [23]	X	X	X	X	X
Capristo et al. [9]	X			X	X
Fabiani et al. [33]					
Carbone et al. [10]	X		X	X	X
Zipser et al. [34]		X			
West et al. [35]		X			
Capristo et al. [36]	X	X		X	X
Dickey and Kearney [37]	X	X	X		
West et al. [38]		X			
Capristo et al. [20]	X	X		X	X
Cheng et al. [39]	X	X			
Duerksen and Leslie [40]	X	X	X	X	
Reilly et al. [41]		X			
Kabanni et al. [42]		X			
Passananti et al. [21]	X	X	X		
Brambilla et al. [11]	X	X			
Valente [43]	X	X		X	X
Kurpaa et al. [44]	X	X	X		
Silva et al. [45]	X	X		X	X
Churruca et al. [46]	X			X	X
Totais de um mesmo parâmetro	78.6%	82.1%	46.4%	51.7%	44.8%

BM, Body mass; BMI, body mass index; BMD, bone mineral density; BMC, bone mineral content; FM, fat mass; FFM, fat-free mass.

**3.2. Anthropometric Measurement Methods.** In order to evaluate anthropometry, the studies used different methodologies, including digital scales and weight balances for body mass, wall, and infant stadiometers for height measurement, plicometer, bioimpedance, isotope dilution, and dual-energy x-ray absorptiometry (DXA) for fat mass and fat-free mass, and DXA for measuring bone mineral density and bone mineral content. The body mass index (BMI) was mathematically defined by the weight/height ratio, with the formula  $BMI = \text{weight (kg)} / \text{height}^2 \text{ (m)}$ .

In Table 3, the frequency of different anthropometric measurement methods is compiled. DXA is present in 83.3%, bioimpedance in 16.6%, skinfold thickness in 16.6%, and isotope dilution in 5.5% of the studies. Moreover, the most used DXA system was Lunar DPX, present in 9 studies, followed by Lunar Prodigy and Hologic Delphy, present in 2 studies each, and Lunar DPA was used in 1 study.

**3.3. Body Mass in Patients with Celiac Disease.** The WGFD group presented a lower BM when compared to the GFD group, in 50% of the studies. This parameter was not different in the other 50% of studies. However, when compared

TABLE 2: Sample characteristics in studies with celiac patients.

Scientific publication	<18 years		>18 years		Control
	M	F	M	F	
Bode et al. [25]			7	15	No
Bardella et al. [26]			31	127	No
Gonzalez et al. [27]				32	85
Rea et al. [28]	8	15			No
Mautalen et al. [29]			5	9	No
Smecuol et al. [30]			1	24	No
Dickey and Bodkin [31]			35	15	No
Lorenzo et al. [32]	12	31			30
Bardella et al. [18]	14	15	8	15	52
Barera et al. [23]			20	51	142
Capristo et al. [9]			16	23	63
Fabiani et al. [33]			26	18	No
Carbone et al. [10]	15	33			41
Zipser et al. [34]			748	248	No
West et al. [35]		2649*	17925		
Capristo et al. [36]				18	22
Dickey and Kearney [37]			114	257	No
West et al. [38]			57	30	No
Capristo et al. [20]			17	9	No
Cheng et al. [39]			248	121	No
Duerksen and Leslie [40]			6	37	233
Reilly et al. [41]	60	82			No
Kabanni et al. [42]			166	513	No
Passananti et al. [21]				95	No
Brambilla et al. [11]	103	47			288
Valente [43]			8	12	39
Kurpaa et al. [44]			40*	No	
Silva et al. [45]	23	8			31
Churruca et al. [46]				54	No
Totais	212	231	1513	1725	18975

The data was separated in children and adolescents (<18 years) and adults (>18 years), subdivided by sex, male (M) and female (F). Control: the presence of a control groups. \*Sex data not informed. These values were not included in stratification by sex.

to the control group, WGFD presented a lower BM in 100% of the reviewed studies. When GFD was compared to the control group, 66% of the studies showed lower values, while 33.35% revealed no difference between these groups (Table 4). Seven studies were not included in the BM table because they did not discriminate between these data.

**3.4. Body Mass Index in Patients with Celiac Disease.** When WGFD versus GFD and WGFD versus the control group were compared, 71.4% of studies showed a lower BMI, while 28.6% presented no difference between the groups. However, GFD compared to the control group presented a lower BMI in 60% of the studies, and no difference in 40% of them (Table 5). Eight articles did not discriminate this parameter between the groups and were not included in Table 5. Passananti et al. compared WGFD to GFD in two time periods, at two and five years, and found different results, which were included separately in Table 5.

**3.5. Fat Mass and Fat-Free Mass in Patients with Celiac Disease.** FM data are compiled in Table 6. When

TABLE 3: Anthropometric measurement methods in celiac patients studies.

Scientific publication	ST	BIA	DXA	ID	DXA devices
Bode et al. [25]	X		X		Lunar DPA
Gonzalez et al. [27]			X		Lunar DPX
Rea et al. [28]	X		X		Lunar DPX
Smecuol et al. [30]			X		Lunar DPX
Lorenzo et al. [32]	X	X	X		Lunar DPX
Bardella et al. [18]			X		Hologic
Barera et al. [23]			X		Lunar DPX
Capristo et al. [9]				X	
Carbone et al. [10]			X		Lunar DPX
Capristo et al. [36]			X		Lunar DPX
Dickey and Kearney [37]			X		Not shown
Capristo et al. [20]			X		Lunar prodigy
Duerksen and Leslie [40]			X		Lunar DPX
Passananti et al. [21]			X		Hologic
Valente [43]			X		Lunar DPX
Kurpaa et al. [44]			X		Lunar prodigy
Silva et al. [45]		X			
Churrua et al. [46]		X			
Totais	16.6%	16.6%	83.3%	5.5%	

ST, skinfold thickness; BIA, bioimpedance; DXA, dual-energy X-ray absorptiometry; ID, isotope dilution.

TABLE 4: Body mass of celiac patients with and without gluten-free diet.

Scientific publication	WGFD versus GFD	WGFD versus control	GFD versus control
Gonzalez et al. [27]	Lower	Lower	Lower
Rea et al. [28]	Equal	Lower	Lower
Smecuol et al. [30]	Lower		
Lorenzo et al. [32]			Equal
Bardella et al. [18]			Lower
Barera et al. [23]		Lower	Equal
Capristo et al. [9]	Lower	Lower	Lower
Carbone et al. [10]		Lower	Lower
Capristo et al. [36]	Equal	Lower	Lower
Capristo et al. [20]		Lower	Lower
Duerksen and Leslie [40]		Lower	
Brambilla et al. [11]			Lower
Valente [43]			Equal
Kurpaa et al. [44]	Equal		
Silva et al. [45]			Equal

compared to GFD and to the control group, 100% of the studies presented a lower FM in the WGFD group. When GFD was compared to the control group, 50% of

TABLE 5: Body mass index of celiac patients with and without gluten-free diet.

Scientific publication	WGFD versus GFD	WGFD versus control	GFD versus control
Bode et al. [25]			Lower
Bardella et al. [26]		Lower	Lower
Gonzalez et al. [27]	Lower	Lower	Lower
Rea et al. [28]		Equal	Equal
Bardella et al. [18]			Lower
Barera et al. [23]		Equal	Equal
Capristo et al. [36]	Lower	Lower	Lower
Capristo et al. [20]	Lower		
Duerksen and Leslie [40]		Equal	
Kabanni et al. [42]	Lower	Lower	
Passananti et al. [21]	Equal		
Passananti et al. [21]	Lower		
Brambilla et al. [11]			Lower
Valente [43]			Equal
Kurpaa et al. [44]	Equal		
Silva et al. [45]			Equal

TABLE 6: Fat mass of celiac patients with and without gluten-free diet.

Scientific publication	WGFD versus GFD	WGFD versus control	GFD versus control
Bode et al. [25]			Lower
Gonzalez et al. [27]	Lower	Lower	Lower
Rea et al. [28]		Lower	Equal
Smecuol et al. [30]	Lower	Lower	Equal
Lorenzo et al. [32]			Lower
Bardella et al. [18]		Lower	Lower
Barera et al. [23]		Lower	Equal
Capristo et al. [9]			Higher
Carbone et al. [10]			Lower
Capristo et al. [36]		Lower	Equal
Capristo et al. [20]	Lower		
Duerksen and Leslie [40]		Lower	
Valente [43]			Equal
Silva et al. [45]			Equal

studies found no difference, 41.7% showed a lower FM, and 8.3% a higher FM.

Table 7 presents the compiled FFM data. Compared to GFD, WGFD showed a lower FFM in 66% of studies, and

TABLE 7: Free fat mass of celiac patients with and without gluten-free diet.

Scientific publication	WGFD versus GFD	WGFD versus control	GFD versus control
Gonzalez et al. [27]	Lower	Lower	Lower
Rea et al. [28]		Lower	Equal
Smecuol et al. [30]		Lower	Lower
Lorenzo et al. [32]			Lower
Bardella et al. [18]		Lower	Lower
Barera et al. [23]		Lower	Equal
Capristo et al. [9]	Equal	Lower	Lower
Carbone et al. [10]			Lower
Capristo et al. [36]		Equal	Lower
Capristo et al. [20]	Lower		
Valente [43]			Equal
Silva et al. [45]			Equal

there was no difference in 33.3% of them. However, WGFD versus control subjects presented lower FFM values in 100% of the studies. When compared to the control group, 36.4% of studies showed a lower FFM in GFD, while 63.6% found no difference.

Four articles, one for FM and three for FFM, did not discriminate data relative to this parameter and were not included in Tables 6 and 7.

### 3.6. Bone Mineral Density and Bone Mineral Content.

When WGFD was compared to the control group, it was found to have a lower BMD and BMC in 90% of the studies. In 66.5% of the studies, GFD presented a lower BMD and BMC versus control patients, and there was no difference in this comparison in 33.3% of the studies. There were not enough data to compare WGFD and GFD (Table 8) in relation to BMD and BMC.

## 4. Discussion

The anthropometric assessment is an important variable to understand human metabolism in different health conditions, [47, 48] including celiac disease. The data presented in our sample of studies, there is a greater level of verification in the body mass index (BMI) 82.1%, and body mass (BM) 78.6%. These techniques are complementary, can be easily accessed, and have a low cost. More complex and expensive variables were used in smaller proportions in anthropometric studies involving patients with celiac disease: fat mass (FM) 53.6%, fat-free mass (FFM) 44.8%, bone mineral density (BMD) 46.4%, and bone mineral content (BMC) 9%. Despite the usefulness of BMI in the classification of subjects, [49] other variables like fat mass, fat-free mass, bone

TABLE 8: Bone mineral density and bone mineral content of celiac patients with and without gluten-free diet results.

Scientific publication	WGFD versus GFD	WGFD versus control	GFD versus control
Bode et al. [25]			Lower
Gonzalez et al. [27]	Lower	Lower	Lower
Mautalen et al. [29]		Lower	Lower
Smecuol et al. [30]		Lower	Lower
De Lorenzo et al. [32]		Lower	
Bardella et al. [18]		Lower	
Barera et al. [23]		Lower	Equal
Carbone et al. [10]		Lower	Equal
Dickey and Kearney [37]		Lower	
Duerksen and Leslie [40]		Lower	Lower
Passananti et al. [21]		Lower	Lower
Kurpaa et al. [44]		Equal	Equal

mineral density, and bone mineral content enable a more comprehensive analysis of anthropometry. Future studies on these topics needs to consider the complexity effects of celiac disease in anthropometric parameters to provide for an appropriate selection of assessment methods.

The reviewed studies presented a higher proportion of adult patients with celiac disease than children and adolescents, being 7 : 1, respectively, and a greater participation of females (53.3%) than males (46.7%). A recent review of ninety-six studies published between 1991 and 2016, indicated a higher prevalence of celiac disease in females than males (0.6% versus 0.4%) and in children than adults (0.9% versus 0.5%). [50] Our data suggests that more research with children and adolescents is necessary to establish a better understanding of the anthropometric impacts as well as the growth and development of celiac disease.

Regarding the anthropometry measurement methods, a higher prevalence of DXA was observed, at 83.3% which, despite its high cost, has the main advantage of simultaneously analyzing fat mass, fat-free mass, and bone mineral content; moreover, it is considered the gold standard for body composition measurements. [47, 48, 51, 52] In this review, Lunar DPX was the most used DXA device, followed by Lunar Prodigy, Hologic Delphy, and Lunar DPA. A previous review from Marinangeli and Kassis [53] about DXA comparative studies indicate that there are inter- and intradevices differences, with under or overestimated values, which may be explain by factors like beam type, gender sample, and scan speed. However, quality-control procedures may help the correct identification of changes in body composition parameters [53].

Bioimpedance, with a frequency of 16.6%, depends on many factors to obtain validity and precision, such as the type of apparatus, researcher handling, room temperature,



and the preparation of subjects in relation to feeding, hydration, physical exercise routine, and the consumption of alcohol and medicine [54]. These limitations may explain the lower utilization of bioimpedance in these studies. The skinfold thickness evaluation is one of the most popular research methods because of its low cost and simplicity, and there is a good relation between subcutaneous fat and total fat mass [47, 55]. However, its frequency of use in these studies was low, a factor that can be explained by its limitation to predict bone mineral density and bone mineral content, being variables present in a larger proportion of the reviewed studies. Isotope dilution was the least utilized method, at 5.5%. This method verifies the deuterium, oxygen eighteen, and tritium concentration, thus determining the total water, fat mass, and fat-free mass for the whole body [56]. Despite having good precision, the technique is very difficult to analyze, [47] being a possible cause for its restricted use. We credit the greater preference for the DEXA method among others to two factors: (1) the possibility of accessing many variables simultaneously, allowing for more complete and faster analyses; (2) the higher incidence of bone mineral density and bone mineral content in the reviewed methodologies, which are variables easily measured by DEXA devices.

Considering the lack of studies with two or more body composition assessment methodologies, we cannot do a comparative evaluation between them, since it is not possible to claim that the result is a consequence of the selected method or of group characteristics. However, previous research suggests that skinfold thickness and bioimpedance seems to provide underestimated results for body fat when compared to DXA devices [57, 58].

Body mass was evaluated in 80.7% of the reviewed studies. All studies that evaluated body mass in WGFD patients and most of the GFD studies revealed significantly lower levels of this parameter when compared to the control group, corroborating previous findings [9–11, 18, 23, 27, 28, 32, 36]. Although many publications present a gluten-free diet as a body mass promoter [11, 20, 23, 30, 40], our review found that in half of the studies, there was no significant difference between WGFD and GFD groups.

The majority of the studies indicated that WGFD presented lower values of BMI than control subjects, in accordance with previous observations [27, 36, 40, 42, 59]. However, the majority of studies showed higher BMI values in WGFD compared to GFD, suggesting that BMI seems to be influenced by a gluten-free diet [11, 20, 21, 30, 41]. However, most studies also demonstrated that WGFD have lower values of BMI when compared to the control group, suggesting that the gluten-free diet may not be able to normalize this parameter [11, 16, 32, 39]. In the work of West et al. [35], which proposed a categorization of samples thorough BMI, a greater prevalence of WGFDs in the underweight classification was verified. In addition, Kabbani et al. [42] observed a significant probability of WGFD and GFD patients to be in the underweight classification in relation to the overweight and obesity classifications.

The mensuration of fat mass and fat-free mass in patients with celiac disease constitutes an important parameter to

investigate the effects of this disease on anthropometric and metabolic functioning. Moreover, the increased body mass found in GFD subjects seems to be essentially related to an increase in fat mass [9, 20, 30, 36]. The studies compiled indicated lower values of FM in WGFD compared to GFD and the control group, corroborating with previous publications [9, 10, 16, 18, 23, 27, 28, 30, 32, 36, 40]. Although all the publications revealed a rise of FM in GFD patients, only half of these studies showed the same values when GFD patients are compared to the control group [23, 28, 30, 36, 43, 60].

In the majority of publications, GFD presented lower values of FFM than the control group, suggesting that a gluten-free diet may not be able to normalize FFM values in relation to healthy people [9, 10, 17, 18, 30, 32, 36].

The analysis of bone mineral density and bone mineral content revealed that the majority of publications found significantly lower levels of these variables in WGFD than in control groups [10, 17, 18, 21, 23, 29, 30, 32, 37, 40, 44]. A few publications suggest that a gluten-free diet significantly promotes bone mineral density and bone mineral content [21, 29, 30] with some articles indicating that celiac disease patients submitted to a one year gluten-free diet can present normal values of these parameters [10, 23, 44]. However, most of the reviewed studies showed lower mineral density and bone mineral content values in GFD patients compared to control groups [17, 21, 25, 29, 30, 40, 44].

Our limitations are similar to those observed by Bardella et al., which includes the difficulty in comparing previously published anthropometry and nutritional data in patients with celiac disease. Data ambiguity and the dependence on other variables such as age at diagnosis, symptom duration before diagnosis, and the presence or absence of unsatisfactory nutrient absorption constitutes important factors of interference.

## 5. Conclusion

Celiac disease significantly changes anthropometric parameters, which can be improved by a gluten-free diet. Despite this, celiac disease patients may not improve anthropometric variables to values similar in healthy people.

This review demonstrated that WGFD compared to the control group showed lower values in all anthropometric variables. GFD compared to WGFD presented higher values in BMI, FM, and FFM, and there was inconclusive data about BM, BMD, and BMC. GFD patients did not present different FFM values from control.

Anthropometric parameters that were more utilized included body mass index and body mass, followed by fat mass, bone mineral density and bone mineral content, and fat-free mass, which were most measured by dual-energy x-ray absorptiometry (DXA), followed by bioimpedance, skinfold thickness, and isotope dilution.

The findings of this review suggest that celiac disease patients must be periodically evaluated through anthropometric parameters, since the pathology has the potential to modulate such values even in a gluten-free diet, with these variables reflecting their healthy status. In parallel, the

screening of different anthropometric assessment methodologies can provide support for more accurate evaluations by scientists and clinical professionals who work with celiac disease patients.

## Abbreviations

BMI:	Body mass index
BM:	Body mass
BF:	Body fat
BMD:	Bone mineral density
BMC:	Bone mineral content
FFM:	Fat-free mass
WGFD:	Celiac disease patients without a gluten-free diet
GFD:	Celiac disease patients with a gluten-free diet
TtG:	Antitransglutaminase antibodies
IgA:	Immunoglobulin A
DXA:	Dual-energy X-ray
ST:	Skinfold thickness
BIA:	Bioimpedance
ID:	Isotope dilution.

## Conflicts of Interest

The authors declare no conflicts of interest.

## Acknowledgments

The authors thank the Federal University of Latin American Integration—UNILA, Paraná, Brasil.

## References

- [1] F. Clot and M.-C. Babron, "Genetics of celiac disease," *Molecular Genetics and Metabolism*, vol. 71, no. 1-2, 2000.
- [2] L. M. Sollid, "Coeliac disease: dissecting a complex inflammatory disorder," *Nature Reviews Immunology*, vol. 2, no. 9, pp. 647–655, 2002.
- [3] M. F. Kagnoff, "AGA institute medical position statement on the diagnosis and management of celiac disease," *Gastroenterology*, vol. 131, no. 6, pp. 1977–1980, 2006.
- [4] T. Sudbrack Da Gama E Silva and W. Furlanetto, "Artigo de revisão diagnóstico de doença celíaca em adultos," *Revista da Associação Médica Brasileira*, vol. 56, no. 1, pp. 122–126, 2010.
- [5] A. Fasano, M. Araya, S. Bhatnagar et al., "Federation of international societies of pediatric gastroenterology, hepatology, and nutrition consensus report on celiac disease," *Journal of Pediatric Gastroenterology and Nutrition*, vol. 47, no. 2, pp. 214–219, 2008.
- [6] C. Catassi, I. M. Räscher, E. Fabiani et al., "Coeliac disease in the year 2000: exploring the iceberg," *The Lancet*, vol. 343, no. 8891, pp. 200–203, 1994.
- [7] C.-H. Florén and P. Alm, "Defective synthesis of apolipoprotein A-I in jejunal mucosa in coeliac disease," *Scandinavian Journal of Gastroenterology*, vol. 23, no. 7, pp. 856–860, 1988.
- [8] N. Malandrino, E. Capristo, S. Farnetti et al., "Metabolic and nutritional features in adult celiac patients," *Digestive Diseases*, vol. 26, no. 2, pp. 128–133, 2008.
- [9] E. Capristo, G. Addolorato, G. Mingrone et al., "Changes in body composition, substrate oxidation, and resting metabolic rate in adult celiac disease patients after a 1-y gluten-free diet treatment," *The American Journal of Clinical Nutrition*, vol. 72, no. 1, pp. 76–81, 2000.
- [10] M. C. Carbone, G. Pitzalis, M. Ferri et al., "Body composition in coeliac disease adolescents on a gluten-free diet: a longitudinal study," *Acta Diabetologica*, vol. 40, no. 1, pp. S171–S173, 2003.
- [11] P. Brambilla, M. Picca, D. Dilillo et al., "Changes of body mass index in celiac children on a gluten-free diet," *Nutrition, Metabolism and Cardiovascular Diseases*, vol. 23, no. 3, pp. 177–182, 2013.
- [12] D. Schuppan, Y. Junker, and D. Barisani, "Celiac disease: from pathogenesis to novel therapies," *Gastroenterology*, vol. 137, no. 6, pp. 1912–1933, 2009.
- [13] A. P. A. Santos, C. de S. Monteiro, H. F. Arantes, M. S. Costa, R. M. Rosa, and G. M. Ribeiro, "Celiac disease: pathways for diagnosis," *Revista Médica de Minas Gerais*, vol. 24, no. 3, pp. 381–387, 2014.
- [14] I. D. Hill, M. H. Dirks, G. S. Liptak et al., "Guideline for the diagnosis and treatment of celiac disease in children: recommendations of the north American society for pediatric gastroenterology, hepatology and nutrition," *Journal of Pediatric Gastroenterology and Nutrition*, vol. 40, no. 1, pp. 1–19, 2005, <http://www.ncbi.nlm.nih.gov/pubmed/15625418>.
- [15] S. Mora, G. Barera, A. Ricotti, G. Weber, C. Bianchi, and G. Chiumello, "Reversal of low bone density with a gluten-free diet in children and adolescents with celiac disease," *The American Journal of Clinical Nutrition*, vol. 67, no. 3, pp. 477–481, 1998.
- [16] S. Bodé, C. Hassager, E. Gudmand-Hoyer, and C. Christiansen, "Body composition and calcium metabolism in adult treated coeliac disease," *Gut*, vol. 32, no. 11, pp. 1342–1345, 1991.
- [17] D. González, R. Mazure, C. Mautalen, H. Vazquez, and J. Bai, "Body composition and bone mineral density in untreated and treated patients with celiac disease," *Bone*, vol. 16, no. 2, pp. 231–234, 1995.
- [18] M. T. Bardella, C. Fredella, L. Prampolini, N. Molteni, A. M. Giunta, and P. A. Bianchi, "Body composition and dietary intakes in adult celiac disease patients consuming a strict gluten-free diet," *The American Journal of Clinical Nutrition*, vol. 72, no. 4, pp. 937–939, 2000.
- [19] C. Catassi, I. M. Räscher, E. Fabiani et al., "Coeliac disease in the year 2000: exploring the iceberg," *The Lancet*, vol. 343, no. 8891, pp. 200–203, 1994, <http://www.ncbi.nlm.nih.gov/pubmed/7904667>.
- [20] E. Capristo, N. Malandrino, S. Farnetti et al., "Increased serum high-density lipoprotein-cholesterol concentration in celiac disease after gluten-free diet treatment correlates with body fat stores," *Journal of Clinical Gastroenterology*, vol. 43, no. 10, pp. 946–949, 2009.
- [21] V. Passananti, A. Santonicola, C. Bucci et al., "Bone mass in women with celiac disease: role of exercise and gluten-free diet," *Digestive and Liver Disease*, vol. 44, no. 5, pp. 379–383, 2012.
- [22] T. Kemppainen, H. Kröger, E. Janatuinen et al., "Osteoporosis in adult patients with celiac disease," *Bone*, vol. 24, no. 3, pp. 249–255, 1999, <http://www.ncbi.nlm.nih.gov/pubmed/10071918>.
- [23] G. Barera, S. Mora, P. Brambilla et al., "Body composition in children with celiac disease and the effects of a gluten-free diet: a prospective case-control study," *The American Journal of Clinical Nutrition*, vol. 72, no. 1, pp. 71–75, 2000.
- [24] C. Hallert, C. Grant, S. Grehn et al., "Evidence of poor vitamin status in coeliac patients on a gluten-free diet for 10 years,"

- Alimentary Pharmacology and Therapeutics*, vol. 16, no. 7, pp. 1333–1339, 2002.
- [25] S. H. Bode, E. H. Bachmann, E. Gudmand-Hoyer, and G. B. Jensen, "Stature of adult coeliac patients: no evidence for decreased attained height," *European Journal of Clinical Nutrition*, vol. 45, no. 3, pp. 145–149, 1991.
  - [26] M. T. Bardella, M. Fraquelli, M. Quatrini, N. Molteni, P. Bianchi, and D. Conte, "Prevalence of hypertransaminasemia in adult celiac patients and effect of gluten-free diet," *Hepatology*, vol. 22, no. 3, pp. 833–836, 1995.
  - [27] D. González, R. Mazure, C. Mautalen, H. Vazquez, and J. Bai, "Body composition and bone mineral density in untreated and treated patients with celiac disease," *Bone*, vol. 16, no. 2, pp. 231–234, 1995, <http://www.ncbi.nlm.nih.gov/pubmed/7756052>.
  - [28] F. Rea, C. Polito, A. Marotta et al., "Restoration of body composition in celiac children after one year of gluten-free diet," *Journal of Pediatric Gastroenterology & Nutrition*, vol. 23, no. 4, pp. 408–412, 1996.
  - [29] C. Mautalen, D. Gonzalez, R. Mazure et al., "Effect of treatment on bone mass, mineral metabolism, and body composition in untreated celiac disease patients," *American Journal of Gastroenterology*, vol. 92, no. 2, pp. 313–318, 1997.
  - [30] E. Smecuol, D. Gonzalez, C. Mautalen et al., "Longitudinal study on the effect of treatment on body composition and anthropometry of celiac disease patients," *American Journal of Gastroenterology*, vol. 92, no. 4, pp. 639–643, 1997.
  - [31] W. Dickey and S. Bodkin, "Prospective study of body mass index in patients with coeliac disease," *BMJ*, vol. 317, no. 7168, p. 1290, 1998, <http://www.ncbi.nlm.nih.gov/pubmed/9804717>.
  - [32] A. De Lorenzo, C. Di Campli, A. Andreoli, G. F. Sasso, M. Bonamico, and A. Gasbarrini, "Assessment of body composition by bioelectrical impedance in adolescent patients with celiac disease," *American Journal of Gastroenterology*, vol. 94, no. 10, pp. 2951–2955, 1999.
  - [33] E. Fabiani, L. M. Taccari, I. M. Rättsch, S. Di Giuseppe, G. V. Coppa, and C. Catassi, "Compliance with gluten-free diet in adolescents with screening-detected celiac disease: a 5-year follow-up study," *Journal of Pediatrics*, vol. 136, no. 6, pp. 841–843, 2000, <http://www.ncbi.nlm.nih.gov/pubmed/10839888>.
  - [34] R. D. Zipser, S. Patel, K. Z. Yahya, D. W. Baisch, and E. Monarch, "Presentations of adult celiac disease in a nationwide patient support group," *Digestive Diseases and Sciences*, vol. 48, no. 4, pp. 761–764, 2003, <http://www.ncbi.nlm.nih.gov/pubmed/12741468>.
  - [35] J. West, R. F. A. Logan, T. R. Card, C. Smith, and R. Hubbard, "Risk of vascular disease in adults with diagnosed coeliac disease: a population-based study," *Alimentary Pharmacology and Therapeutics*, vol. 20, no. 1, pp. 73–79, 2004.
  - [36] E. Capristo, S. Farnetti, G. Mingrone et al., "Reduced plasma ghrelin concentration in celiac disease after gluten-free diet treatment," *Scandinavian Journal of Gastroenterology*, vol. 40, no. 4, pp. 430–436, 2005.
  - [37] W. Dickey and N. Kearney, "Overweight in celiac disease: prevalence, clinical characteristics, and effect of a gluten-free diet," *American Journal of Gastroenterology*, vol. 101, no. 10, pp. 2356–2359, 2006.
  - [38] J. West, R. F. A. Logan, P. G. Hill, and K. Khaw, "The iceberg of celiac disease: what is below the waterline?," *Clinical Gastroenterology and Hepatology*, vol. 5, no. 1, pp. 59–62, 2007.
  - [39] J. Cheng, P. S. Brar, A. R. Lee, and P. H. R. Green, "Body mass index in celiac disease: beneficial effect of a gluten-free diet," *Journal of Clinical Gastroenterology*, vol. 44, no. 4, pp. 267–271, 2010.
  - [40] D. R. Duerksen and W. D. Leslie, "Longitudinal evaluation of bone mineral density and body composition in patients with positive celiac serology," *Journal of Clinical Densitometry*, vol. 14, no. 4, pp. 478–483, 2011.
  - [41] N. R. Reilly, K. Aguilar, B. G. Hassid et al., "Celiac disease in normal-weight and overweight children: clinical features and growth outcomes following a gluten-free diet," *Journal of Pediatric Gastroenterology and Nutrition*, vol. 53, no. 5, pp. 528–531, 2011.
  - [42] T. A. Kabbani, A. Goldberg, C. P. Kelly et al., "Body mass index and the risk of obesity in coeliac disease treated with the gluten-free diet," *Alimentary Pharmacology & Therapeutics*, vol. 35, no. 6, pp. 723–729, 2012.
  - [43] F. X. Valente, "Avaliação dos fatores de risco para doenças cardiovasculares em portadores de doença celiaca," *Dissertação, Universidade Federal de Viçosa, Viçosa, Brazil*, 2013, <https://www.locus.ufv.br/bitstream/handle/123456789/7271/texto%20completo.pdf?sequence=1&isAllowed=y>.
  - [44] K. Kurppa, A. Paaola, P. Collin et al., "Benefits of a gluten-free diet for asymptomatic patients with serologic markers of celiac disease," *Gastroenterology*, vol. 147, no. 3, pp. 610–617.e1, 2014.
  - [45] M. M. D. S. E. Silva, M. Bahia, F. J. Penna, and L. Gandra, "Anthropometric profile of patients with celiac disease tended at the pediatric gastroenterology clinic of UFMG, Belo Horizonte," *Revista Médica de Minas Gerais*, vol. 24, no. 4, pp. 457–463, 2014.
  - [46] I. Churrua, J. Miranda, A. Lasa, M. A. Bustamante, I. Larretxi, and E. Simon, "Analysis of body composition and food habits of Spanish celiac women," *Nutrients*, vol. 7, no. 7, pp. 5515–5531, 2015.
  - [47] D. R. Wagner and V. H. Heyward, "Techniques of body composition assessment: a review of laboratory and field methods," *Research Quarterly for Exercise and Sport*, vol. 70, no. 2, pp. 135–149, 1999.
  - [48] C. V. Albanese, E. Diessel, and H. K. Genant, "Clinical applications of body composition measurements using DXA," *Journal of Clinical Densitometry*, vol. 6, no. 2, pp. 75–85, 2003, <http://www.ncbi.nlm.nih.gov/pubmed/12794229>.
  - [49] World Health Organization, "Consultation on obesity. Obesity: preventing a managing the global epidemic. Geneva. Switzerland: division of non communicable diseases, program of nutrition, family and reproductive," *Journal of Clinical Endocrinology and Metabolism*, vol. 7, no. 1, pp. 1–178, 1998.
  - [50] P. Singh, A. Arora, T. A. Strand et al., "Global prevalence of celiac disease: systematic review and meta-analysis," *Clinical Gastroenterology and Hepatology*, vol. 16, no. 6, pp. 823–836.e2, 2018.
  - [51] T. Erselcan, F. Candan, S. Saruhan, and T. Ayca, "Comparison of body composition analysis methods in clinical routine," *Annals of Nutrition and Metabolism*, vol. 44, no. 5-6, pp. 243–248, 2000.
  - [52] A. Scafoglieri and J. P. Clarys, "Dual energy X-ray absorptiometry: gold standard for muscle mass?," *Journal of Cachexia, Sarcopenia and Muscle*, vol. 9, no. 4, pp. 786–787, 2018.
  - [53] C. P. Marinangeli and A. N. Kassis, "Use of dual X-ray absorptiometry to measure body mass during short- to medium-term trials of nutrition and exercise interventions," *Nutrition Reviews*, vol. 71, no. 6, pp. 332–342, 2013.
  - [54] V. H. Heyward and L. M. Stolarczyk, "Avaliação da composição corporal aplicada," *SP: Manole*, vol. 2015, pp. 2-3, 2000.



- [55] J. V. G. A. Durnin and J. Womersley, "Body fat assessed from total body density and its estimation from skinfold thickness: measurements on 481 men and women aged from 16 to 72 years," *British Journal of Nutrition*, vol. 32, no. 1, pp. 77–97, 1974.
- [56] D. Halliday and A. G. Miller, "Precise measurement of total body water using trace quantities of deuterium oxide," *Biological Mass Spectrometry*, vol. 4, no. 2, pp. 82–87, 1977, <http://www.ncbi.nlm.nih.gov/pubmed/884210>.
- [57] J. Grove and Y.-J. Hung, "Body fat prediction equations for skinfold and bioelectrical impedance analysis using dual-energy x-ray absorptiometry data as the criterion," *Journal of Physical Therapy and Sports Medicine*, vol. 1, no. 1, 2017.
- [58] K. González-Ruíz, M. Medrano, J. E. Correa-Bautista et al., "Comparison of bioelectrical impedance analysis, slaughter skinfold-thickness equations, and dual-energy X-ray absorptiometry for estimating body fat percentage in Colombian children and adolescents with excess of adiposity," *Nutrients*, vol. 10, no. 8, p. 1086, 2018.
- [59] M. T. Bardella, M. Fraquelli, M. Quatrini, N. Molteni, P. Bianchi, and D. Conte, "Prevalence of hypertransaminasemia in adult celiac patients and effect of gluten-free diet," *Hepatology*, vol. 22, no. 3, pp. 833–836, 1995, <http://www.ncbi.nlm.nih.gov/pubmed/7657290>.
- [60] M. M. de S. E Silva, M. Bahia, F. J. Penna, and L. Gandra, "Perfil antropométrico de pacientes com doença celíaca atendidos pelo ambulatório de gastroenterologia pediátrica da UFMG, Belo Horizonte, MG—Brasil," *Revista Médica de Minas Gerais*, vol. 24, no. 4, 2014, <http://www.rmmg.org/artigo/detalhes/1704>.

