The importance of obesity and carbohydrate metabolism disorders on the course of gastroesophageal reflux disease – a pilot study

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Abstract

Introduction: Carbohydrate metabolism disorders, obesity and a severe course of gastroesophageal reflux correlate with more frequent development of esophageal complications. The aim of this study was to assess the influence of obesity and carbohydrate disorders on the characteristics of gastroesophageal reflux disease (GERD).

Methods: The study included 58 patients with excess weight. Anthropometric parameters (including the body mass index, BMI), data regarding GERD (severity of symptoms, gastroscopy and esophageal pH monitoring results) were included in the study. Correlations between obesity and GERD parameters were analyzed. Subjects were divided into a diabetic and a control group and the severity of GERD was compared. Results: GERD was diagnosed in 40 patients and occurred more frequently in the obese group (73%) than in the overweight group (57%). Increased GERD severity was associated with increased BMI only for postprandial parameters. GERD was diagnosed in most of the group with carbohydrate disorders (78% vs 63% in the non-diabetic group). No differences in the severity of GERD were observed between groups depending on carbohydrate disorders. Conclusions: In our study, GERD was common in obesity and in diabetic disorders. Increased severity of postprandial reflux was associated with an increased BMI. Diabetic disorders were not associated with more severe GERD.

Keywords: gastroesophageal reflux disease · obesity · diabetes mellitus

Citation

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Factors linked to modern lifestyle, e.g. physical inactivity, overnutrition and poor sleep quality have led to the widespread incidence of obesity, type 2 diabetes (T2DM) and gastroesophageal reflux disease (GERD) [1-5]. In many countries obesity is an important public health problem and its prevalence results in an increase in the incidence of GERD and type 2 diabetes [6-8]. Gastroesophageal reflux disease is a common health problem which still requires research. Depending on the region, the prevalence of GERD is estimated at 10-20% of the population [2, 5]. Overweight, especially abdominal obesity, correlates with the severity of GERD. The main causes of GERD are impaired function of the lower esophageal sphincter, hiatal hernia, impaired esophageal motility and increased intra-abdominal pressure [9-18]. Gastroesophageal reflux disease leads to the development of Barrett’s esophagus (BE) or metaplasia, which is a precancerous condition leading to the development of dysplasia and subsequently of esophageal adenocarcinoma (EAC) [4, 19-20]. Numerous studies have noted that not only the severity of GERD, but the presence of abdominal obesity and T2DM, increase the risk of developing BE and EAC [21-25].

In abdominal obesity, the development of BE and EAC plays a major role in the increased amount of adipokines and inflammatory cytokines produced by visceral adipose tissue, which leads to a chronic inflammatory process, and thus, promotes cancerous transformation [26-28]. Patients with T2DM are often obese, making them more prone to GERD. In T2DM, as in high-grade obesity, the clinical course of GERD is more often atypical. The differences between patients with T2DM result from additional factors contributing to GERD, including esophageal peristalsis disorders, gastroparesis, hyperglycemia, neuropathy and usage of diabetic medications (e.g. GLP-1 receptor agonists and metformin) [29]. The atypical, often mild or asymptomatic course of GERD in T2DM may delay the decision to perform diagnostics, leading to delayed diagnosis of complications, particularly BE and EAC. Therefore, it is important to study the natural course of GERD in obese and T2DM patients in order to reduce the risk of complications through early diagnosis and treatment.

Aim

The aim of our study was to characterize GERD in overweight patients and to assess the correlation between obesity parameters and the severity of GERD. The second aim of this study was to assess the differences in the clinical characteristics of GERD, depending on the diagnosis of carbohydrate disorders, which could explain the increased risk of complications in this group.

Materials and Methods

Study population

Our analysis covers data collected from 58 patients who were diagnosed for sleep breathing disorders at the Department of Internal Diseases of the Institute of Rural Medicine in Lublin. The exclusion criteria were: chronic use of drugs which may interfere with the assessment of GERD parameters (proton pump inhibitors, H2 blockers, alkali, nitrates) or previous significant gastrointestinal surgery (gastrectomy, bowel resection). This patient group also served as study participants in our previous work [30]. We collected anthropometric measurements and the responses from GERD-complaint questionnaires. If GERD was suspected, patients were referred for additional tests (gastroscopy and esophageal pH-measurement). The results from these additional tests were included in the study.

Criteria for the diagnosis of GERD

The diagnosis of GERD was established according to the definitions of the Lyon consensus [4]. Therefore, the diagnosis was based on the combined assessment of clinical symptoms, endoscopic evaluation of the esophageal mucosa, esophageal pH monitoring and response to therapeutic intervention. Clinical diagnosis was made when the patient had persistent symptoms characteristic of GERD which include heartburn and acid regurgitation. Persistence of symptoms was recognized when symptoms of mild intensity occurred at least 2 days a week or when they were more severe and caused deterioration in general well-being and occurred at least 1 day a week. Moreover, the diagnosis of GERD confirmed the presence of inflammatory changes in the esophagus (LA grade C) and the result of pH-measurement with esophageal acid exposure time > 6%, DeMeester Score > 14.72 or > 80 reflux episodes per 24 hours. Additionally, the diagnosis of GERD was confirmed by the reduction of symptoms after starting treatment with a proton pump inhibitor [4].

Anthropometric data

All participants underwent a physical examination. The body mass index (BMI) was calculated as the body weight in kilograms divided by the height in meters squared (kg/m²). The waist circumference (the circumference at midpoint between the lower border of the rib cage and the iliac crest) was measured in the standing position. Overweight was diagnosed if BMI was in the range of 25-29.9 kg/m², class I obesity when BMI was 30-34.9 kg/m², class II obesity when BMI was 35-39.9 kg/m², and class III obesity when BMI was at ≥ 40 kg/m² [8].
Survey data regarding the severity of GERD complaints

The severity of GERD clinical symptoms was assessed using a questionnaire about the overall intensity of complaints (within a score range of 0-10) and the intensity and frequency of symptoms considered typical for GERD (within a score range of 0-52). GERD symptoms included the feeling of heartburn and presence of regurgitation typical for GERD situations, acid reflux and dysphagia. In addition, the presence of GERD symptoms was assessed at night, during sleep and after an overnight sleeping period. Patients were instructed to describe the symptoms occurring in the month prior to completing the questionnaire. Survey data from all patients were collected by the primary investigator. The full version of the questionnaire is available in the Supplementary Materials [in Polish]: https://ejtc.mumed.edu.pl/files/67.

Gastroscopy and esophageal pH monitoring

All procedures were undertaken by experienced physicians before the introduction of GERD treatment. All gastroscopic examinations were done using the Fujifilm (Japan or Pentax) (Japan). Whereas the esophageal pH monitoring was done using the ComforTEC Plus PHNS single-channel probe and a recorder made by Sandhill Scientific (USA, REF: Z07-2000-A, SN: H109007C). The degree of esophagitis was determined according to the Los Angeles classification, and numerical values with a range of 1 to 4 were given for subsequent grades (grade A-D); 0 was designated as no inflammatory lesions [4, 10].

Diagnosis of pre-diabetes and type 2 diabetes

In order to assess the importance of carbohydrate disorders on the course of GERD, the subjects were divided into a group with carbohydrate disorders (23 patients) and a control group (35 patients). The group with carbohydrate disorders included patients diagnosed with T2DM, impaired glycemic tolerance (IGT) or impaired fasting glycemia (IFG). In all participants with T2DM, diabetes was well-controlled. Most of the gastroscopic examinations revealed esophageal inflammatory changes with grade A, B and C of esophagitis recognized in 22, 7, and 1 subject, respectively. Grade D lesions or peptic stricture were not observed in any of the patients. Moreover, neither BE nor EAC were diagnosed. In 23 patients, pH-metry was performed. Table 1 presents the basic characteristics of the study population.

Data Analyses and Statistical Methods

All statistical analyses were carried out with the Statistica software package (version 13, TIBCO Software Inc., USA). After confirming that all variables meet the criteria of a normal distribution, Spearman’s rank correlation coefficient was used for the analysis. Correlations between variables were calculated using Spearman’s rank correlation coefficient, while comparisons between two independent groups were performed using the Mann–Whitney U test. P values < 0.05 were considered statistically significant. Because the distribution of GERD measurements was characterized by a large asymmetry, the median should be taken as the key measurement (possibly including the range of variability in the form of IQR) while assessing its severity depending on the presence of diabetic disorders.

Results

Baseline data

The study population consisted of 58 patients (48 males and 10 females) aged 34-75 years (mean = 54.5 years; Me = 56 years; s = 11.2 years). In the study group, obesity was diagnosed in 44 subjects (75.9%). Of these, class I obesity was seen in 17 subjects, class II obesity was seen in 12 subjects, and class III obesity was seen in 15 subjects. Fourteen (24.1%) patients were overweight. The average patient weight was 104 kg (range 77-161 kg), while average waist circumference was 115.5 cm (range 96-147 cm). The mean BMI was 34.8 kg / m² (range 25.1-49.7 kg / m²). Carbohydrate disorders were reported by 23 subjects (~40%), of which 14 subjects (24.1%) had T2DM and 9 subjects (15.5%) had pre-diabetes. Table 1 presents the basic characteristics of the study population.

In our study, GERD was diagnosed in 40 patients (~69%). In the obese group, GERD was diagnosed in 32 subjects (73%), while in the overweight group, GERD was diagnosed in 8 subjects (57%). Gastroscopy was performed in 32 patients. In 2 cases, no esophageal inflammatory changes were observed. Most of the gastroscopic examinations revealed esophageal inflammatory changes with grade A, B and C of esophagitis recognized in 22, 7, and 1 subject, respectively. Grade D lesions or peptic stricture were not observed in any of the patients. Moreover, neither BE nor EAC were diagnosed. In 23 patients, pH-metry was performed. Table 1 presents the distribution of the variables.

Apart from obesity, carbohydrate disorders and GERD, the majority participants had comorbidities mainly related to the circulatory system. The most common were: arterial hypertension (47 patients, 81%); dyslipidemia (34 patients, 59%), coronary artery disease (16 patients, 28%); hyperuricemia (10 patients, 17%); chronic obstructive pulmonary disease (5 patients, 9%) and heart failure (4 patients; 7%).

Gastroesophageal reflux in obesity and in diabetic disorders

We investigated the relationship between obesity parameter values (body weight, abdominal circumference, and BMI) and GERD severity parameters. Since all considered
features were consistent with a normal distribution, Spear- 
man’s rank correlation coefficient was used for the analysis. 
Table 2 presents correlation coefficient values between indi-

gual features along with the assessment of their statistical 

significance. In the studied group, no statistically signifi-
cant relationships were found between the obesity parameters 
and the assessed reflux parameters. 

A tendency toward greater values with increasing obe-
sity parameters was seen only for the “feeling of heartburn 
after meals” parameter.

The relationships between GERD parameters from gas-
troscopy, pH measurement and obesity parameters were 
calculated. Only the correlation between the number of 
postprandial reflux episodes and BMI was near statistical 
significance (test probability values $p = 0.0084$), however its 
strength was rather small $r_S = 0.36$. In addition, there was 
a trend toward greater values for the postprandial De 
Meester index and for the duration of gastric acid exposure 
with increasing BMI (Table 3).

Table 1. Basic characteristics of the study population, severity of GERD symptoms and distribution of esophageal pH monitoring results

| Features                           | Mean  | Median | Std. dev. | Min  | Max  |
|------------------------------------|-------|--------|-----------|------|------|
| Age (yr)                           | 54.5  | 56     | 11.2      | 34   | 75   |
| BMI (kg/m²)                        | 34.8  | 34.6   | 6.7       | 25.1 | 49.7 |
| Waist circumference (cm)           | 115.5 | 115.0  | 12.8      | 96.0 | 147.0|
| GERD symptoms day                  | 11.9  | 12     | 9.2       | 0    | 28   |
| GERD symptoms night                | 2.8   | 3      | 2.9       | 0    | 12   |
| GERD symptoms overall              | 14.7  | 15     | 11.5      | 0    | 40   |
| De Meester index (n = 23)          | 27.8  | 15.1   | 33.2      | 1.7  | 136.4|
| De Meester index – post meal (n = 23) | 9.8   | 7.5    | 9.0       | 1.5  | 37   |
| Reflux episodes – recumbent (n = 22) | 34.5  | 10     | 57.7      | 0    | 250  |
| Reflux episodes – 24/h (n = 22)    | 136.1 | 107    | 127.9     | 7    | 570  |
| Mean pH night (n = 23)             | 6.4   | 6.5    | 0.8       | 4.3  | 8.1  |
| Mean pH – 24/h (n = 23)            | 6.2   | 6.2    | 0.5       | 5.0  | 7.2  |
| Esophageal clearance time (n = 22) | 37.0  | 35.5   | 21.2      | 8    | 91   |
| Esophageal clearance time – recumbent (n = 22) | 31.7  | 22.5   | 33.1      | 0    | 138  |
| Longest reflux episode (n = 23)    | 13.7  | 4.9    | 17.9      | 0.3  | 59.3 |
Table 2. Distribution of correlations between obesity parameters and the severity of GERD complaints

| GERD symptoms                          | Obesity parameters (n = 58) |   |   |   |
|----------------------------------------|----------------------------|---|---|---|
|                                        | Weight                     | Waist circumference | BMI |
| GERD symptoms day                      | 0.05 (p = 0.7338)          | 0.07 (p = 0.6178)   | 0.03 (p = 0.7973) |
| GERD symptoms night                    | 0.07 (p = 0.5866)          | 0.06 (p = 0.6513)   | 0.04 (p = 0.7892) |
| GERD symptoms day and night            | 0.05 (p = 0.7039)          | 0.07 (p = 0.5779)   | 0.04 (p = 0.7681) |
| Overall intensity of GERD-related complaints | -0.01 (p = 0.9461)      | 0.07 (p = 0.5988)   | 0.10 (p = 0.4678) |
| Burning sensation in the chest (Post Meal) | 0.16 (p = 0.2302)      | 0.16 (p = 0.2440)   | 0.19 (p = 0.1605) |

Table 3. Correlations between GERD parameters from gastroscopy, pH measurement and obesity parameters

| GERD severity parameters               | Obesity parameters (n = 58) |   |   |   |
|----------------------------------------|----------------------------|---|---|---|
|                                        | Weight                     | Waist circumference | BMI |
| Severity of inflammatory changes in the esophagus (gastroscopy assessment) | -0.08 (p = 0.6753) | -0.04 (p = 0.8218) | -0.09 (p = 0.6203) |
| De Meester Index (Post Meal)           | 0.02 (p = 0.9340)          | 0.10 (p = 0.6415)   | 0.24 (p = 0.2636) |
| De Meester Index                       | -0.10 (p = 0.6608)         | 0.00 (p = 0.9964)   | 0.11 (p = 0.6198) |
| Reflux Episodes (Post Meal)            | 0.16 (p = 0.4610)          | 0.22 (p = 0.3198)   | **0.36 (p = 0.0884)** |
| Reflux Episodes (24 h)                 | -0.04 (p = 0.8554)         | -0.06 (p = 0.7859)  | 0.06 (p = 0.8029) |
| Exposure to gastric acid following a meal | 0.03 (p = 0.9055)    | 0.12 (p = 0.5803)   | 0.22 (p = 0.3044) |
| Exposure to gastric acid (24 h)        | -0.05 (p = 0.8085)         | -0.05 (p = 0.8305)  | 0.03 (p = 0.8888) |
| Mean pH (Post Meal)                    | 0.12 (p = 0.5836)          | 0.00 (p = 0.9839)   | -0.11 (p = 0.6025) |
| Mean pH (24 h)                         | 0.15 (p = 0.4928)          | -0.06 (p = 0.7962)  | -0.13 (p = 0.5468) |
| Esophageal Clearance (Post Meal)       | -0.13 (p = 0.5452)         | 0.04 (p = 0.8540)   | 0.08 (p = 0.7216) |
| Esophageal Clearance (24 h)            | -0.05 (p = 0.8308)         | 0.07 (p = 0.7636)   | 0.02 (p = 0.9284) |
Next, we analyzed the differences between the parameters of GERD severity depending on the diagnosis of carbohydrate disorders. Among the participants with carbohydrate disorders, GERD was diagnosed in 18 patients (78%), of which 12 (86%) had T2DM, and 6 (67%) had pre-diabetes. Whereas in the group without carbohydrate disorders (n = 35) GERD was diagnosed in 22 participants (63%). No differences in the severity of GERD were observed between the groups with and without carbohydrate disorders. In our study, the GERD parameters in both groups were very similar (Table 4).

Table 4. Comparison of GERD severity parameters depending on the presence of diabetic disorders (p – test probability values were calculated using the Mann-Whitney test)

| GERD severity parameters                                      | Diabetic disorders |   |   |   |   |   |   |   |
|--------------------------------------------------------------|-------------------|---|---|---|---|---|---|---|
|                                                             | Yes               | No | p  |
|                                                             | n | Mean | Me | IQR | n | Mean | Me | IQR |
| GERD symptoms day and night                                  | 23 | 16.5 | 18 | 21.0 | 35 | 13.8 | 14 | 22.0 | 0.4206 |
| Severity of inflammatory changes in the esophagus (gastroscopy Assessment) | 12 | 1.1 | 1 | 0.0 | 20 | 1.3 | 1 | 1.0 | 0.2551 |
| Reflux Episodes (24 h)                                       | 10 | 111.8 | 66.5 | 165.0 | 12 | 156.4 | 146.5 | 151.5 | 0.4562 |
| Reflux Episodes (Post Meal)                                  | 10 | 45.9 | 28 | 49.0 | 13 | 69.2 | 34 | 119.0 | 0.7381 |
| Exposure to gastric acid following a meal (%)                | 10 | 5.6 | 2.5 | 8.5 | 12 | 8.1 | 4.5 | 9.5 | 0.8718 |
| Mean pH (24 h)                                               | 10 | 6.1 | 5.9 | 0.6 | 13 | 6.3 | 6.4 | 0.6 | 0.2316 |
| Mean pH (day)                                                | 10 | 6.1 | 6.0 | 0.8 | 13 | 6.1 | 6.3 | 1.1 | 0.7844 |
| Mean pH (night)                                              | 10 | 6.1 | 6.1 | 0.8 | 13 | 6.6 | 6.7 | 0.3 | 0.0666 |
| De Meester Index                                             | 10 | 24.7 | 10.6 | 27.9 | 13 | 30.2 | 18.1 | 29.0 | 1.0000 |
| De Meester Index (Post Meal)                                 | 10 | 7.8 | 6.4 | 9.8 | 13 | 11.3 | 8.4 | 8.8 | 0.7844 |
| Esophageal Clearance – 24 h (sec)                            | 10 | 40.7 | 38 | 20.0 | 12 | 33.8 | 33 | 24.5 | 0.3136 |
| Overall intensity of GERD-related complaints                  | 23 | 3.8 | 2.0 | 8.0 | 35 | 2.5 | 2.0 | 5.0 | 0.3271 |
| GERD symptoms day                                            | 23 | 13.7 | 15 | 17.0 | 35 | 11.0 | 10 | 20.0 | 0.3041 |
| GERD symptoms night                                          | 23 | 2.8 | 3 | 4.0 | 35 | 2.8 | 2 | 5.0 | 0.9497 |
| Burning sensation in the chest (Post Meal)                   | 23 | 2.9 | 3.0 | 5.0 | 35 | 2.5 | 3.0 | 4.0 | 0.4973 |
The analysis of the probability of GERD depending on the selected variables (BMI, waist circumference, diagnosis of carbohydrate disorders and diagnosis of diabetes) was performed using the logistic regression model. Based on the analyzes performed, no statistical evidence was found that any of the proposed factors had a significant influence on the diagnosis of GERD (Table 5). Moreover, an attempt was made to search for a model containing statistically significant variables. The best model that included only the diagnosis of diabetes was still not statistically significant: OR (95% CI) = 3.429; p = 0.14.

Discussion

All participants in our study were overweight and in this group the occurrence of GERD was much more frequent than in the general population. Obesity is an important risk factor for GERD and numerous studies demonstrated its more frequent occurrence in people with excess weight [11-17]. In our study, no statistically significant correlation was found between individual obesity measures and GERD parameters. However, there was a trend towards greater severity of postprandial GERD clinical symptoms and worsening of postprandial GERD pH parameters with increasing BMI. Most of the published data demonstrate a more severe course of GERD in people with excess weight.

Obesity, especially the visceral type, causes GERD due to changes in the anatomy and physiology of the gastroesophageal junction (GEJ) [11, 18]. Additionally, it is believed that the pro-inflammatory effects of cytokines synthesized in visceral adipose tissue plays an important role [12, 26]. Akyuz et al. showed a significant correlation between BMI and the severity of GERD in pH measurements. They also found that the severity of esophageal inflammatory changes seen in gastroscopy did not differ significantly in the obese group, however the authors did not assess the severity of GERD in the context of its association with abdominal obesity [11]. In our study, we also did not observe any relationship between obesity and the severity of esophageal inflammatory changes in gastroscopy. The correlation between BMI and GERD severity in pH measurements was also present, but did not reach the level of statistical significance.

We did not observe a correlation between abdominal circumference and the severity of GERD. However, most studies indicate that it does have greater a role than BMI in terms of GERD severity. A study by Wu et al. investigated the correlations of obesity parameters with GERD symptoms and esophageal inflammatory activity via the measurement of glucose metabolism in 18F-Fluorodeoxyglucose positron emission tomography (PET-CT). There was a significant correlation between GERD symptoms and esophageal inflammatory activity in PET-CT with all obesity parameters (BMI, abdominal circumference, and the amount of subcutaneous and visceral adipose tissue) [13]. Nam et al. described the correlations between the amount of visceral fat, concentrations of inflammatory cytokines synthesized within it, and the intensity of esophageal inflammatory changes [12]. Similar results were obtained in large-scale studies in Japan and South Korea. An increased incidence of reflux esophagitis has been observed in obesity and in the metabolic syndrome. Hyperglycemia, high BMI, and in particular, greater abdominal circumference and increased visceral fat, correlated with an increased risk for GERD [14-15].

Gastroesophageal reflux disease is linked to a higher risk of BE [4, 20]. Population-based studies have indicated that the risk of BE and EAC is also significantly increased in obesity, especially in the abdominal type. This risk is increased regardless of the presence of GERD symptoms [21, 23-25]. In a study by Nelsen et al., the risk of developing BE and dysplasia correlated with the amount of visceral adipose tissue and adipose tissue in the GEJ fat area; however, it was independent of the BMI value and the presence of GERD symptoms [24]. Similar conclusions were obtained by El-Seraq et al. [19]. Moreover, in an investigation by Corley et al., abdominal circumference and abdominal obesity (but not BMI) correlated with a greater risk for BE [25].

Table 5. Logistic regression model of the probability of GERD diagnosis depending on selected variables

| Independent variables          | GERD diagnosis |        | p          |
|-------------------------------|----------------|-------|------------|
|                               | OR (95% CI)    |       |            |
| BMI                           | 1.060 (0.867-1.295) | 0.5704 |
| WC                            | 0.975 (0.880-1.079) | 0.6207 |
| Diagnosis of carbohydrate disorders | 1.158 (0.244-5.496) | 0.8532 |
| Diagnosis of diabetes         | 3.195 (0.392-26.073) | 0.2781 |

OR – odds ratio; CI – confidence interval; GERD – gastroesophageal reflux disease; BMI – body mass index (kg/m2); waist circumference (cm)
Gastroesophageal reflux disease is common in T2DM and is more likely to be atypical or present with mild symptoms [29]. In addition, these patients are at an increased risk for developing metaplasia [22]. In our study, no differences in the severity of GERD were found between subjects with T2DM or pre-diabetes and the group without these disorders. Notably, the two groups were very similar in terms of GERD characteristics. Lorentzen et al. compared the features of GERD in patients with a high degree of obesity, depending on the diagnosis of T2DM. As in our study, GERD was more common in the obese group than in the general population. However, a large proportion of the respondents had asymptomatic GERD, regardless of whether or not they suffered from T2DM. In this study, clinical symptoms were reported by approximately 29% of the respondents, but esophagitis in gastroscopy was seen in 58% of patients in the T2DM group and in 47% of patients in the non-T2DM group. Among subjects with inflammatory changes in the esophagus, 68-80% did not report symptoms of GERD. In the T2DM group (only T2DM patients underwent pH-metry), 55% of subjects had pathologic acid reflux, whereas 67% of subjects were asymptomatic [16]. In our study, the severity of GERD clinical symptoms was similar in both groups, which may be due to the lower number of severely obese patients when compared to the cited study.

As reported by Ortiz et al., the asymptomatic course of GERD in obese patients may be related to the decreased esophageal sensitivity to acid content observed in this group. The authors indicated that the absence of typical GERD symptoms in these patients may delay the diagnosis of GERD complications, especially BE [32]. Promberger et al. also observed the frequent occurrence of atypical GERD symptoms in T2DM [33]. Furthermore, Lluch et al. found that GERD was common in diabetic patients, but it was more often asymptomatic [34]. The above conclusions are of clinical significance in the context of a report by Leggett et al., which found an increased risk of BE in patients with metabolic syndrome, regardless of GERD symptoms [23]. The role of carbohydrate disorders in the pathogenesis of GERD is unclear. Gokturk et al. observed a more severe course of GERD in subjects with T2DM, but the presence of reflux episodes was associated with obesity rather than hyperglycemia [17]. In the study by Wang et al., the occurrence of GERD symptoms in diabetes was observed more frequently, and their severity clearly increased in the group of patients with diabetic neuropathy [35].

In our study group, we did not observe an atypical course of GERD in T2DM, but in all cases, these were patients with pre-diabetes or well-controlled diabetes without complications. Although the patients with carbohydrate disorders more often suffer from GERD and its complications, the clinical course does not correlate with the risk of complications. Because of this, they may benefit from early gastroscopic evaluation.

The limitations of our study include the small number of participants and the lack of assessment of other obesity parameters, such as the waist-hip ratio. Moreover, not all patients underwent endoscopic examinations or pH-metry. Another limitation was the inclusion of only hospitalized patients with suspected sleep apnea who were mostly obese, which makes it difficult to transfer the obtained conclusions to the general population. Despite these limitations, the collected results allowed us to demonstrate a greater incidence of GERD in obesity and to show that the presence of carbohydrate disorders was not associated with a more severe clinical course of GERD (in the context of clinical symptoms, changes in gastroscopy, and pH measurement).

Conclusions

In our study group, we observed that GERD is more common in obesity and in T2DM; however, the diagnosis of diabetic disorders was not associated with more severe GERD. Our results and a review of the current literature indicate that due to a mild or atypical course, GERD may be underdiagnosed in the group of severely obese and T2DM patients. Finally, although patients with carbohydrate disorders more often suffer from GERD and its complications, the clinical course does not correlate with the risk of complications, and because of this, these patients may benefit from early gastroscopic evaluation.

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Conflicts of interest

There are no conflicts of interest to report for any of the authors.
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