ORIGINAL ARTICLE / ОРИГИНАЛНИ РАД

Prevalence and risk factors for Barrett's esophagus in patients with chronic gastroesophageal reflux disease

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SUMMARY

Introduction/Objective The most important complication of gastroesophageal reflux disease (GERD) is Barrett's esophagus (BE) and the development of esophageal adenocarcinoma. Prevalence of BE is 5–15% in patients with GERD symptoms.

The aim of the study was to investigate the prevalence and risk factors for BE in patients with chronic reflux symptoms. A prospective study was conducted in the Clinic of Gastroenterology, Niš Clinical Center. **Methods** We included 676 patients with chronic reflux symptoms, who underwent esophagogastroduodenoscopy. The biopsy specimens were obtained in a four-quadrant fashion at intervals of 2 cm from the circumferential endoscopic Barrett's epithelium in the distal esophagus. BE was diagnosed by pathological examination.

Results Out of the total number patients with GERD, 92 were diagnosed with columnar-lined esophagus (CLE), the prevalence being 13.6%. Histological examination of biopsy from 92 patients with CLE revealed specialized intestinal metaplasia in 15 patients, with the prevalence of 2.22%. Compared to patients without BE, patients with BE were older and more commonly male. Univariable analyses showed that hiatal hernia and *Helicobacter pylori* infection were two significant risk factors for the onset of esophagitis. The age and the presence of reflux symptoms were associated with the presence of BE. Older age could be considered a significant risk factor for the development of BE and GERD.

Conclusion Prevalence of biopsy-proven BE and CLE in Serbia was 2.22% and 13.6%, respectively, in patients with GERD symptoms.

Keywords: Barrett's esophagus; gastroesophageal reflux disease; chronic reflux symptoms

INTRODUCTION

Gastroesophageal reflux disease (GERD) is a long-term condition where stomach contents come back up into the esophagus resulting in either symptoms or complications. GERD is mild acid reflux that occurs at least twice a month, or moderate to severe acid reflux that occurs at least once a week. In 20% of the population, symptoms last longer than one week. The prevalence of GERD significantly varies among different populations. The prevalence of all forms of GERD is 40%, the weekly symptoms have 14% of the population, and the daily symptoms range 4–7% [1]. Peptic esophagitis, reflux esophagitis and erosive esophagitis, erosive reflux disease (ERD) are synonyms for the subgroup of GERD patients with histopathological changes of esophageal mucosa that usually correlate with the symptoms of acid reflux content. Non-erosive reflux disease (NERD) includes the group of patients with symptomatic GERD who have no macroscopic mucosal changes noticed on the esophagogastroduodenoscopy. It is estimated that 50-70% of patients with GERD have NERD. Symptoms and signs of esophageal reflux disease can be varying intensity and are not always in correlation with the severity of esophageal damage [2].

Barrett's esophagus (BE) is a consequence of chronic GERD, that predisposes the development of esophageal adenocarcinoma (EAC) [3]. Endoscopically, the prevalence of BE has been estimated at 1-2% in all patients who underwent upper endoscopy for any indication, and anywhere from 5% to 15% in patients with symptoms of GERD. Among the malignant tumors of the esophagus, the incidence of Barrett's adenocarcinoma is increasing. The incidence of EAC has been three to four times higher in the last two decades. It is believed that the main reason for this high percentage of Barrett's adenocarcinoma is related to an increased incidence of BE, which shows a close causal relationship with GERD [4]. However, not all patients with gastroesophageal reflux and erosive esophagitis will develop BE, and all patients with BE do not have a history of gastroesophageal reflux. At least 25% of patients with BE do not have a history of GERD. In many patients with reflux esophagitis, treatment leads to the regeneration of the mucosa. Some patients will develop BE with an increased risk of developing EAC. There are many risk factors that can

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contribute to the development of BE, which is the subject of many studies in the world [5, 6].

The esophagus lined with columnar epithelium (CLE) and BE are the conditions in which stratified squamous epithelium is continuously replaced by a cylindrical epithelium from an esophageal-gastric junction. BE is characterized by the presence of specialized intestinal metaplasia (SIM). As SIM is part of the definition and is the epithelial type associated with cancer, obtaining biopsies from the columnar lined distal esophagus is mandatory. The sensitivity and positive predictive values of standard upper endoscopy for diagnosing BE have been reported as 82% and 34%, respectively [7]. Guidelines of the American College of Gastroenterology state that every patient with gastroesophageal reflux symptoms should at least once in his/her lifetime be referred for BE screening endoscopy. Patients with SIM in CLE are currently advised to undergo a periodic endoscopic surveillance to detect progression to dysplasia at an early, potentially curable stage. New techniques such as chromoendoscopy and magnification endoscopy have been tried to improve recognition of SIM [4].

The aim of this study was to determine the prevalence and possible risk factors of BE in patients with chronic reflux symptoms.

METHODS

A prospective study conducted at the Clinic of Gastroenterology, Niš Clinical Center, included 676 patients with chronic reflux symptoms and all underwent esophagogastroduodenoscopy. The symptoms are defined as the presence of heartburn and regurgitation at least three times a week for one year. The questionnaire was completed by every patient; the questionnaire included information on age, sex, occupation, as well as the following criteria: primary referral symptoms, frequency of GERD symptoms, acid test, extraesophageal symptoms. Patients with a history of documented peptic disease, gastric or esophageal surgery, and those with motor disorders such as achalasia, diffuse esophageal spasm, or scleroderma, were excluded. Gastroesophageal junction (GEJ) is defined as the beginning of the proximal limit of gastric mucosal folds (Figure 1). CLE was identified as a columnar epithelium over 1 cm from the GEJ which had a reddish color and a velvety texture that could be easily distinguished from the normal pale and glossy esophageal squamous epithelium. The length of the CLE was estimated by subtracting the distance from the incisors to the squamocolumnar junction (Z-line) from the distance between the incisors and the GEJ (Figure 2). The patients were classified as short-segment BE (SSBE) if the length of the columnar appearing mucosa was less than 3 cm above the GEJ, and long-segment BE (LSBE) if the length of the columnar mucosa was equal to or greater than 3 cm. The diagnosis of BE is based on the presence of endoscopic findings compatible with columnar epithelium in the distal esophagus and confirmed by the presence of SIM on biopsies (Figure 3).



Figure 1. Endoscopic appearance of normal gastroesophagel junction; note that the squamocolumnar line corresponds with proximal extent of the gastric folds



Figure 2. Salmon-colored mucosa is seen extending proximal to the gastroesophagel junction consistent with Barrett's esophagus



Figure 3. Histological appearance of Barrett's epithelium; intestinalized mucosa with branching pits and goblet cells (H&E, ×20)

The study protocol was approved by the local ethics committee and all patients gave their informed consent to be included. All the patients were fully informed of the study protocol and agreed to undergo upper gastrointestinal endoscopy.

All upper endoscopies were performed using a GIF100 or GIF130 video endoscope (Olympus, Lake Success, NY, USA). Macroscopic mucosal changes of the distal esophagus were measured on the basis of the distance from the Z line, and mucosal damage was classified according to the Los Angeles classification of reflux esophagitis [8].

The presence of a hiatal hernia and its size was determined in all the patients during the withdrawal of the endoscope and was measured in centimeters. We investigated the presence of *Helicobacter pylori* infection in all the patients by using pathology and rapid urease test.

The biopsy specimens were obtained in a four-quadrant fashion at intervals of 2 cm from the circumferential endoscopic Barrett's epithelium in the distal esophagus. In patients with small islands or irregular tongues of columnar appearing mucosa, at least two specimens were obtained within the abnormal-appearing mucosa at intervals of 1cm from the GEJ to the proximal extent of the abnormality. All biopsy specimens were stained with hematoxylin and eosin and with Alcian blue (pH 2.5) stain.

Statistical analysis

The processing of the obtained data was made using SPSS for Windows, version 16.0 (SPSS Inc., Chicago, IL, USA). The data was processed using standard descriptive statistical methods (mean value, standard deviation, and percentage representation). The results were analyzed using the appropriate tests depending on the size of the group, type of mark, and type of distribution. We used the Student's t-test for continuous variables and χ^2 test for categorical variables, in comparative analyses. A univariate analysis was performed to determine the variables independently associated with the risk of BE. A p-value < 0.05 was considered statistically significant.

RESULTS

Patient with gastroesophageal reflux disease

The average age of subjects with the symptoms of reflux disease was 50 ± 13 years. There were 381 men (56.36%) and 295 women (43.64%). Based on endoscopic findings, patients were divided into two groups: the NERD group included 403 (59.61%) patients, and the ERD group included 273 patients (40.39%). Esophagitis A grade was found in 64.44%, B grade in 26.66%, and C grade in 8.88% of the ERD group patients. Esophagitis D grade was not found. The mean age of patients in both groups did not differ significantly (p = 0.07). The percentage of respondents by sex was approximately the same. Of the clinical manifestations of reflux disease, the heartburn symptom significantly correlates with ERD (p = 0.013). Heartburn was

equally represented in the groups compared to daytime. In both groups of patients, heartburn was more frequent during the day (ERD, p = 0.00001; NERD, p = 0.00001), while fewer patients in both groups had heartburn at night. The symptom of regurgitation was more frequent in the NERD group in 222 (55.08%) patients, but without statistical significance. Hiatal hernia was more frequent in the ERD group, with a statistically significant (p = 0.001). *H. pylori* infection was significantly higher in NERD patients, 24.81% (n = 100). There was no correlation between the presence of *H. pylori* infection and the existence of reflux symptoms (Table 1).

Characteristics	NERD (n = 403)	ERD (n = 273)	p-value
Age	49 ± 15	52 ± 17	0.07
Sex Male Female	220 (54.59%) 183 (45.41%)	161 (58.97%) 112 (41.03%)	0.30
Hiatal hernia Yes No	91 (22.58%) 312 (77.42%)	160 (58.61%) 113 (41.39%)	0.001
RUT Yes No	100 (24.81%) 303 (75.19%)	86 (31.5%) 187 (68.5%)	0.05
Heartburn	239 (59.3%)	190 (69.58%)	0.013
Regurgitation	222 (55.09%)	158 (57.87%)	0.54

Table 1. Background characteristics of the study groups

 $\mathsf{NERD}-\mathsf{non}\mathsf{-}\mathsf{erosive}$ reflux disease; $\mathsf{ERD}-\mathsf{erosive}$ reflux disease; $\mathsf{RUT}-\mathsf{rapid}$ urease test

Prevalence of columnar-lined esophagus

Of all the patients with GERD, 92 had CLE, with the prevalence of 13.6%. Sixty-five patients were found to have normal endoscopy results, and 27 had erosive esophagitis ($\chi^2 = 27.39$; p = 0.001). On endoscopic examination of all 92 patients, 35% had circumferential CLE, 34% had tongue-like extensions, and 31% isolated islands. A short CLE segment was found in 56% of the patients, and a long CLE segment was found in 13% of the patients.

Prevalence of Barrett's esophagus

Histological examination of biopsy from 92 patients with CLE revealed SIM in 15 patients, with the prevalence of 2.22% in our study. Of the 15 patients with BE, nine patients were found to have a long BE segment and six had a short BE segment. The average age of patients with BE was 59 ± 15 years, and 12 of them (80%) were male. The percentage of patients with CLE who had SIM was 16.3%; this was more frequent with a long CLE segment. The largest number of patients did not have erosive changes in the esophagus during endoscopy (87%), and the hiatal hernia was noticed in 80% of patients with BE (Table2).

Prevalence of Barrett's esophagus in gastroesophageal reflux disease

Compared to patients without BE, patients with BE were older and more commonly men, with statistical significance

No metaplasia Metaplasia Characteristics p-value (n = 77)(n = 15)Age 49 ± 12 59 ± 15 0.001 59 (76.62%) 12 (80%) Male 0.61 Female 18 (23.38%) 3 (20%) 0.58 Heartburn 0.004 53 (68.83%) 2 (13.33%) Regurgitation 19 (24.68%) 10 (66.67%) 0.12 NERD 52 (67.53%) 13 (86.67%) 0.34 FRD 25 (32.47%) 2 (13.34%) 0.25 Hiatal hernia 40 (51.95%) 12 (80%) 0.17 CLE Short segment 47 (61.04%) 6 (33.34%) 0.29 9 (53.34%) 0.005 Long segment 3 (3.89%)

Table 2. Predictors of specialized intestinal metaplasia or Barrett's esophagus

NERD – non erosive reflux disease; ERD – erosive reflux disease; CLE – the esophagus lined with columnar epithelium

Table 3. Background characteristics of the study groups

Characteristics	BE (n = 15)	Without BE (n = 661)	p-value
Age	59 ± 15	49 ± 15	0.001
Male Female	12 (80%) 3 (20%)	372 (56.28%) 289 (43.72%)	0.06
Heartburn	2 (13.33%)	414 (62.63%)	0.04
Hiatal hernia Yes No	12 (80%) 3 (20%)	244 (36.91%) 417 (63.09%)	< 0.05
RUT Yes No	4 (26.66%) 11 (73.34%)	182 (27.53%) 479 (73.47%)	0.43

BE – Barrett's esophagus; RUT – rapid urease test

(p = 0.001). The symptom of heartburn was the dominant symptom, statistically occurring more frequently in patients with BE (p = 0.04). The univariate analyses showed that hiatal hernia and *H. pylori* infection were the most significant risk factors for the onset of esophagitis. The age and the presence of reflux symptoms are associated with the presence of BE (Table3).

DISCUSSION

In previous decades, the lower part of the esophagus and cardia have been in the focus of extensive research. The reason for this is a dramatic increase in the incidence of adenocarcinoma of the esophagogastric junction. In comparison, the incidence of GERD and BE as one of its complications was also noticed. Some data indicate a 10-fold increase in the incidence of Barrett's esophagus in Western European countries over the last few decades. Barrett's metaplasia is considered an intermediary event in the development of EAC [9].

In our study, the average age of subjects with symptoms of reflux disease was 50 ± 13 . Almost 60% of patients with GERD did not have endoscopic signs of esophagitis, which is similar to those of Western countries, which shows that 60-70% of patients with typical reflux symptoms do not have damage of esophageal mucosa during endoscopy. In both groups, there were more male than female patients,

though without statistical significance. Male sex has been reported to be an independent risk factor for esophagitis. Different parietal cell mass, lower esophageal function or body mass index between sexes have been proposed as possible causes to explain the sex effect [10]. Kumar et al. [11] show the prevalence of the male sex in patients with GERD.

Of the clinical manifestations of GERD, the heartburn symptom was statistically more frequent in the ERD group compared to the NERD group (p = 0.013), but there was no statistically significant association of heartburn symptoms with the degree of esophagitis. GERD symptoms have been inconsistently correlated with endoscopic findings of eosinophilic esophagitis in different studies, some of which favor such correlation, though not with all reflux symptoms, and some argue against it [12].

Hiatal hernia is present in 37.13% of patients with GERD. In the ERD group, the hiatal hernia is present in 58.61% of the patients. We found that the presence of hiatal hernia is a strong risk factor for esophagitis (p = 0,001) [13].

The relationship between *H. pylori* and GERD infection is relatively unclear. H. pylori gastritis can lead to acid hyposecretion and loss of symptoms of burning sensation [14]. In our study, *H. pylori* infection was statistically more common in the NERD than in the ERD group (p = 0.04). We did not find a statistically significant relationship between the presence of *H. pylori* infection and the presence of typical reflux symptoms.

Of all patients with GERD, suspected CLE was found in 92% of patients, representing prevalence of 13.6% of patients with GERD. Sixty-five patients were in the NERD group, and 27 in the ERD group. ($\chi^2 = 27.39$; p = 0.001). Of the 92 patients with suspected CLE revealed, SIM was present in 15 patients, with the prevalence of 2.22%. The prevalence of BE worldwide is different; it is assumed to be higher in the western than in the eastern countries of the world. Westhoff et al. [15] showed a prevalence of 13.2%. Ronkainen et al. [16] showed a prevalence of 2.3% in Sweden, while Kim et al. [17] show a prevalence of less than 1% in Korea. In our study, BE was more common in men (80%) than in patients without BE (56.02%). BE prevalence was statistically more common in men than in women (p < 0.05). Lin et al. [18] in their study showed that 14% of women had BE, compared to 23% of men with BE (p < 0.05). Male sex has been reported to be a risk factor for BE. Age has been also considered a risk factor for it. Edelstein et al. [19] noted that risk of BE increased with increased age. In our study, patients with BE were significantly older than those without BE (p = 0.001). In clinical manifestation, we found a significant difference for heartburn between patients with BE and those without BE, which was more evident in patients with BE. The symptoms of reflux in our study was a good predictor of the risk for BE (p = 0.04), which is in a line with another study. Hak et al. [20] in their study showed that the duration of reflux symptoms is longer in patients with BE than in those without it. In our study, we noticed a significant difference in the existence of hiatal hernia between the

groups – hiatus hernia was more common in patients with BE. Herrera et al. [21] in their study showed that hiatus hernia is independently associated with the presence of BE.

In our study, we did not find that eosinophilic esophagitis is a predictor for the appearance of BE. Different morphological types of BE are not a risk factor for BE. The CLE length is a risk factor for BE. The CLE length was 3 cm in patients with BE, compared to 1.8 cm in patients without BE (p = 0.001). Okita et al. [22], as well as others, also proved that the long segment of BE is a predictor of SIM in the histological examination [23, 24, 25]. In our study, we did not show the presence of dysplasia in any of the patients with BE.

In conclusion, the prevalence of endoscopically suspected CLE in GERD patients is 13.6%. The prevalence of histologically proven BE was 2.22% in patients with GERD in our area. The presence of hiatal hernia, reflux

REFERENCES

- Dent J, El-Serag HB, Wallande MA, Johansson S. Epidemiology of gastro-oesophageal reflux disease: a systematic review. Gut. 2005; 54(5):710–17.
- Armstrong D. Systematic review: persistence and severity in gastroesophageal reflux disease. Aliment Pharmacol Ther. 2008; 28(7):841–53.
- Labenz J, Koop H, Tannapfel A, Kiesslich R, Hölscher AH. The epidemiology, diagnosis and treatment of Barrett's carcinoma. Dtsch Arztebl Int. 2015; 112(13):224–34.
- Shaheen NJ, Falk GW, Iyer PG, Gerson LB. ACG clinical guideline: Diagnosis and management of Barrett's esophagus. Am J Gastroenterol. 2016; 111(1):30–51.
- Ronkainen J, Aro P, Storskrubb T, Johansson SE, Lind T, Bolling-Sternevald E, et al. Prevalence of Barrett's esophagus in the general population: An endoscopic study. Gastroenterology. 2005; 129(6):1825–31.
- Fan X, Snyder N. Prevalence of Barrett's esophagus in patients with or without GERD symptoms: role of race, age, and gender. Dig Dis Sci. 2009; 54(3):572–7.
- Spechler SJ, Souza RF. Barrett's esophagus. N Engl J Med. 2014; 371(9):836–45.
- Vakil N, van Zanten SV, Kahrilas P, Dent J, Jones R; Global Consensus Group. The Montreal definition and classification of gastroesophageal reflux disease (GERD) – a global evidencebased consensus. Am J Gastroenterol. 2006; 101(8):1900–20.
- Fitzgerald RC, di Pietro M, Ragunath K, Ang Y, Kang JY, Watson P, et al. British Society of Gastroenterology guidelines on the diagnosis and management of Barrett's oesophagus. Gut. 2014; 63(1):7–42.
- Rubenstein JH, Mattek N, Eisen G. Age and sex specific yield of Barrett's esophagus by endoscopy indication. Gastrointest Endosc. 2010; 71(1):21–7.
- Kumar S, Sharma S, Norboo T, Dolma D, Norboo A, Stobdan T, et al. Population based study to assess prevalence and risk factors of gastroesophageal reflux disease in a high altitude area. Indian J Gastroenterol. 2011; 30(3):135–43.
- Koek GH, Sifrim D, Lerut T, Janssens J, Tack J. Multivariate analysis of the association of acid and duodeno-gastro-oesophageal reflux exposure with the presence of oesophagitis, the severity of oesophagitis and Barrett's oesophagus. Gut. 2008; 57(8):1056–64.
- Jones MP, Sloan SS, Rabine JC, Ebert CC, Huang CF, Kahrilas PJ. Hiatal hernia size is the dominant determinant oesophagitis presence and severity in gastroesophageal reflux disease. Am J Gastroenterol. 2001; 96(6):1711–7.

symptoms, and long segment of CLE are independently associated with the presence of BE. Older age could be considered a significant risk factor for the development of BE and GERD.

CONCLUSION

A large number of studies have noted that most patients who have endoscopically suspected BE did not have SIM on histological samples. Multicenter studies are required for determining the epidemiology of BE more precisely, after which a cost-effective strategy for BE screening and surveillance can be developed. Studies should be carried out to determine endoscopic predictors, which can be used as surrogate markers for the histological BE, so that only patients with this predecessor are subjected to biopsy.

- Rubenstein JH, Inadomi JM, Scheiman J, Schoenfeld P, Appelman H, Zhang M, et al. Association between helicobacter pylori and Barrett's oesophagus, erosive esophagitis and gastroesophageal reflux symptoms. Clin Gastroenterol Hepatol. 2014; 12(2):239–45.
- Westhoff B, Brotze S, Weston A, McElhinney C, Cherian R, Mayo MS, et al. The frequency of Barrett's esophagus in high-risk patients with chronic GERD. Gastrointest Endosc. 2005; 61(2):226– 31.
- Ronkainen J, Aro P, Storskrubb T, Johansson SE, Lind T, Bolling-Sternevald E, et al. Prevalence of Barrett's esophagus in the general population: an endoscopic study. Gastroenterology. 2005; 129(6):1825–31.
- Kim JH, Rhee PL, Lee JH, Lee H, Choi YS, Son HJ, et al. Prevalence and risk factors of Barrett's esophagus in Korea. J Gastroenterol Hepatol. 2007; 22(6):908–12.
- Lin M, Gerson LB, Lascar R, Davila M, Triadafilopoulos G. Features of gastroesophageal reflux disease in women. Am J Gastroenterol. 2004; 99(8):1442–7.
- Edelstein ZR, Bronner MP, Rosen SN, Vaughan TL. Risk factors for Barrett's esophagus among patients with gastroesophageal reflux disease: a community clinic-based case-control study. Am J Gastroenterol. 2009; 104(4):834–42.
- Hak NG, Mostafa M, Salah T, El-Hemaly M, Haleem M, Abd El-Raouf A, et al. Acid and bile reflux in erosive reflux disease, non-erosive reflux disease and Barrett's esophagus. Hepatogastroenterology. 2008; 55(82-83):442–7.
- Herrera Elizondo JL, Monreal Robles R, García Compean D, González Moreno El, Borjas Almaguer OD, Maldonado Garza HJ, et al. Prevalence of Barrett's esophagus: An observational study from a gastroenterology clinic. Rev Gastroenterol Mex. 2017; 82(4):296–300.
- Okita K, Amano Y, Takahashi Y, Mishima Y, Moriyama N, Ishimura N, et al. Barrett's esophagus in Japanese patients: its prevalence, form, and elongation. J Gastroenterol. 2008; 43(12):928–34.
- 23. Spechler SJ. Barrett esophagus and risk of esophageal cancer: a clinical review. JAMA. 2013; 310(6):627–36.
- Bhat SK, McManus DT, Coleman HG, Johnston BT, Cardwell CR, McMenamin U, et al. Oesophageal adenocarcinoma and prior diagnosis of Barrett's oesophagus: A population-based study. Gut. 2015; 64(1):20–5.
- Pohl H, Sirovich B, Welch HG. Esophageal adenocarcinoma incidence: are we reaching the peak? Cancer Epidemiol Biomarkers Prev. 2010;19(6):1468–70.

Преваленца и фактори ризика за настанак Баретовог једњака код болесника са хроничном гастроезофагеалном рефлуксном болешћу

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САЖЕТАК

Увод/Циљ Најважнија компликација гастроезофагеалне рефлуксне болести (ГЕРБ) јесте појава Баретовог једњака (БЈ) и настанак аденокарцинома. Преваленца БЈ је од 5 до 15% код болесника са симптомима ГЕРБ-а. Циљ ове студије био је испитивање преваленце и ризичних фактора за настанак БЈ код болесника са хроничним симптомима рефлукса. Истраживање је спроведено у Клиници за гастроентерологију Клиничког центра у Нишу.

Методе Укључено је 676 болесника са хроничним рефлуксним симптомима, којима је урађена езофагогастродуоденоскопија. Биопсије су узимане из четири квадранта у дисталном делу једњака, на удаљености од 2 *ст* од ендоскопски суспектног БЈ. БЈ је дијагностикован патолошким прегледом. **Резултати** Од укупног броја болесника са ГЕРБ-ом, суспектан БЈ је нађен код 92 болесника, што чини преваленцу од 13,60% у нашој студији. Након хистолошког испитивања биопсије суспектног БЈ нађена је специјализована интестинална метаплазија код 15 болесника, са преваленцом од 2,22%. У поређењу са болесницима без БЈ, болесници са БЈ су старији, чешће мушкарци, у оба параметра са статистичком значајношћу. Хијатална хернија и инфекција бактеријом *Helicobacter pylori* су два значајна фактора ризика за настанак езофагитиса. Старост и присуство симптома рефлукса су повезани са присуством БЈ. Старији узраст може представљати значајан фактор ризика за развој БЈ и ГЕРБ-а.

Закључак Преваленца хистолошки доказаног БЈ и суспектног БЈ у Србији је била 2,22%, а 13,60% код болесника са симптомима ГЕРБ-а.

Кључне речи: Баретов једњак; гастроезофагеална рефлуксна болест; хронични рефлуксни симптоми