FRONTIER
4889  Treatment repurposing for inflammatory bowel disease using literature-related discovery and innovation
Kostoff RN, Briggs MB, Shores DR

REVIEW
4900  Tumor microenvironment in primary liver tumors: A challenging role of natural killer cells
Polidoro MA, Mikulak J, Cazzetta V, Lleo A, Mavilio D, Torzilli G, Donadon M

MINIREVIEWS
4919  Exploring the food-gut axis in immunotherapy response of cancer patients
Russo E, Nannini G, Dinu M, Pagliai G, Sofi F, Amedei A

ORIGINAL ARTICLE
Basic Study
4933  Tumor necrosis factor alpha receptor 1 deficiency in hepatocytes does not protect from non-alcoholic steatohepatitis, but attenuates insulin resistance in mice

4945  Resveratrol alleviates intestinal mucosal barrier dysfunction in dextran sulfate sodium-induced colitis mice by enhancing autophagy
Pan HH, Zhou XX, Ma YY, Pan WS, Zhao F, Yu MS, Liu JQ

Retrospective Study
4960  Effects of denosumab treatment in chronic liver disease patients with osteoporosis
Saeki C, Saito M, Oikawa T, Nakano M, Torisu Y, Saruta M, Tsubota A

4972  Bowel function and quality of life after minimally invasive colectomy with D3 lymphadenectomy for right-sided colon adenocarcinoma
Lee KM, Baek SJ, Kwak JM, Kim J, Kim SH

4983  Acute liver failure and death predictors in patients with dengue-induced severe hepatitis
Teerasarntipan T, Chaiteerakij R, Komolmit P, Tangkijvanich P, Treeprasertsuk S

4996  Liver fat accumulation measured by high-speed T2-corrected multi-echo magnetic resonance spectroscopy can predict risk of cholelithiasis
Chen H, Zeng WK, Shi GZ, Gao M, Wang MZ, Shen J

5008  Radiomics of rectal cancer for predicting distant metastasis and overall survival
Li M, Zhu YZ, Zhang YC, Yue YF, Yu HP, Song B
SYSTEMATIC REVIEWS

5022 Neutrophil to lymphocyte ratio and albumin bilirubin grade in hepatocellular carcinoma: A systematic review
Bannaga A, Arasaradnam RP

CASE REPORT

5050 Surveilling Russell body Helicobacter pylori-negative gastritis: A case report and review of literature
Peruhova M, Peshevska-Sekalovska M, Georgieva V, Panayotova G, Dikov D
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Editorial Board of *World Journal of Gastroenterology*, Dr. Dario Sorrentino is a Professor of Medicine at Virginia Tech – Carilion School of Medicine and Research Institute (since 2013). His career research experience has ranged from the bench to the bedside focusing on IBDs, and carried out on three different continents. Fifteen years ago, he and his professional colleagues proposed a groundbreaking strategy to prevent post-surgical recurrence of Crohn’s disease that has evolved into today’s standard-of-care. More recently, he and his team developed a novel approach for diagnosing and treating pre-clinical Crohn’s disease, representing a revolutionary approach to IBD management and research. Dr. Sorrentino has published > 150 high-quality publications and delivered speeches on his own research worldwide. His recent work in the United States has garnered awards of research funds exceeding 2 million dollars from major foundations and private sources. (L-Editor: Filipodia)

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Surveilling Russell body *Helicobacter pylori*-negative gastritis: A case report and review of literature

Milena Peruhova, Monika Peshevska-Sekulovska, Viktoriya Georgieva, Gabriela Panayotova, Dorian Dikov

**Abstract**

**BACKGROUND**

Russell body gastritis (RBG) is very rare type of chronic inflammation of gastric mucosa. The pathologic hallmark of the disease is Russell bodies (RB) which represent accumulation of eosinophilic cytoplasmic inclusions in endoplasmic reticulum of mature plasma cells (Mott cells). Most published cases are associated with *Helicobacter pylori* (*H. pylori*) infection because of correlation between plasma cell activation and antigenic stimulation. There are insufficient data about *H. pylori*-negative RBG and very little is known about the natural course of the disease.

**CASE SUMMARY**

A 51-year-old male patient underwent endoscopic screening for mild iron deficiency anemia. Gastroscopy revealed diffuse hyperemia, edema and nodularity of the fundic and corpus mucosa. Due to non-specific endoscopic findings and iron-deficiency anemia our preliminary diagnosis was diffuse type of gastric carcinoma or gastric lymphoma. Biopsy specimens of gastric mucosa showed inflammatory infiltrate rich in Mott cells, consisting entirely of cytoplasmic RB. Absence of nuclear atypia and mitosis of the plasma cells, polyclonal pattern of the Mott cells and negative staining for cytokeratins favored diagnosis of RBG. The patient was treated with proton-pump inhibitor for 8 wk. Long-term clinical and endoscopic surveillance was scheduled. Albeit, there was no improvement in endoscopic features of the gastric mucosa in three consecutive gastroscopies, histopathological findings demonstrated that the chronic inflammatory infiltrate in the fundic mucosa is less pronounced, rich in plasma cells, with almost absent RB and Mott cells.

**CONCLUSION**

The prognosis of this entity is uncertain, that is why these patients are subjects of
introduction

The first case of Helicobacter pylori (H. pylori)-related Russell body gastritis (RBG) was announced in 1998 by Tazawa et al.[1]. They described that H. pylori-positive gastritis is characterized by localized accumulation of plasma cells containing Russell bodies (RB) in gastric mucosa[1]. RB represent nondegradable, condensed immunoglobulin disposed in endoplasmic dilated rough reticulum cistern of plasma cells[1].

Most of the cases reported in the English literature so far are about H. pylori-positive RBG[1][8]. The main theory regarding the pathogenesis of RBG includes chronic infection with H. pylori leading to abnormal secretion of immunoglobulin or their derivates by plasma cells and subsequent formation of intracellular RB[8].

The few H. pylori-negative RBG cases that have been published were associated with HIV infection, alcohol and drug abuse, concomitant carcinoma and plasma cell neoplasms.

The clinical and endoscopic manifestation of RBG are variable and non-specific[1][3]. This rare type of chronic gastritis should be distinguished from other malignancies of the stomach such as carcinoma, lymphoma and plasmacytoma.

We present a case from our practice with H. pylori-negative RBG, who underwent endoscopic and histological surveillance three times over a period of one year.

We also made a review of twenty-one cases of H. pylori-negative RBG published in the literature up to now with their specific and unique clinical, endoscopic and histopathological features.

Case Presentation

Chief complaints

We present a case of a 51-year-old male who underwent endoscopic screening for mild iron deficiency anemia. The patient had no upper and lower gastrointestinal (GI) complaints.

History of present illness

The iron-deficiency anemia was diagnosed 2 mo before the admission to the hospital.

History of past illness

He was without other co-morbidities and past history for illness.
**Physical examination**

From the physical examination, he had pale skin and visible mucous membranes.

**Laboratory examinations**

The laboratory work-up showed hemoglobin = 107 g/L, serum iron was 10.2 µmol/L (11.6-31.3 µmol/L), ferritin 18.43 ng/mL (30-400 ng/mL), total iron binding capacity 83.2 µmol/L (45-72 µmol/L). Inflammatory serological markers were within the normal limits–CRP was 0.30 ml/L (0-5 mg/L). Other biochemistry test results as well as carcinoembryonic antigen and carbohydrate antigen 19-9, were within the normal limits.

**Imaging examinations**

The colonoscopy was unremarkable. However, upper GI endoscopy revealed diffuse hyperemia, edema and nodularity of the gastric mucosa in the fundus and body, with a clear demarcation line between the body and the antrum. (Figure 1) The duodenal mucosa was intact. Due to non-specific endoscopic findings and iron-deficiency anemia our preliminary diagnoses were diffuse gastric carcinoma or gastric lymphoma. Therefore, multiple biopsies were taken from the stomach. We did not obtain duodenal, ileal and colonic biopsies, as there were no endoscopic abnormalities of the mucosa. We performed whole-body computed tomography with contrast enhancement. It showed neither pulmonary and abdominal space-occupying lesions, nor bone lytic lesions and enlarged lymph nodes.

Histologically, in the biopsy of fundic mucosa, we observed inflammatory infiltrate rich in plasma cells with numerous eosinophilic hyaline bodies (RB) and so-called mature plasma cells (Mott cells), consisting entirely of cytoplasmic RB (Figure 2A). Several hyperplastic lymphoid follicles were also observed. The plasma cells expressed CD79a (Figure 3A), CD138 and they showed polyclonal pattern, both expressed kappa (Figure 3B) and lambda IgM light immunoglobulin chains (Figure 3C). A large number of eosinophils were seen and the neutrophilic leucocytes were rare.

There was no evidence of nuclear atypia and mitosis of the plasma cells, which ruled out lymphoma. Giemsa staining for *H. pylori* infection was negative, as well as immunohistochemical detection. Negative staining for cytokeratin AE1/AE3 excluded the signet-ring cell carcinoma. This microscopic finding corresponds to the so-called RBG.

We ruled out HIV, HCV and HBV infections, as well as autoimmune diseases, such as autoimmune thyroiditis and rheumatoid arthritis. However we did not check for M protein in the serum, we did not perform Bence-Jones protein urine test, TB-spot, EBER in situ hybridization or trephine biopsy of the bone marrow.

**FINAL DIAGNOSIS**

Histopathological findings confirmed *H. pylori*-negative RBG.

**TREATMENT**

The patient was treated with proton-pump inhibitor (PPI) (Esomeprazole)-20 mg bid for 8 wk and intravenous iron medication.

**OUTCOME AND FOLLOW-UP**

Long-term clinical and endoscopic surveillance was scheduled. Three months later, he came for follow-up. His blood tests showed slight increase of his hemoglobin level (117 g/L). He underwent second gastroscopy with endoscopic findings identical to the previous one. Diathermic snare was used which allowed obtaining of larger and deeper tissue specimen of gastric mucosa. Histology report revealed dense accumulation of plasma cells in lamina propria, with decreased distribution of RB (Figure 2B). Intestinal metaplasia was observed in the areas with plasma cell infiltration but without dysplasia. Histopathological findings from third gastroscopy, performed twelve months after the initial diagnosis, demonstrated that the chronic inflammatory infiltrate in the fundic mucosa is less pronounced, rich in plasma cells,
Figure 1  Endoscopic appearance of *Helicobacter pylori*-negative Russell body gastritis.

Figure 2  Russell body gastritis of the fundus mucosa in the standard histological stain Hematoxylin-eosin, × 400. A: The initial biopsy specimen shows abundant plasma cell inflammatory infiltrate, rich in Russell body and mature plasma cell; B: The follow-up biopsy revealed no change in plasma cell inflammatory infiltrate, but with decreased distribution Russell body and mature plasma cells; C: Third biopsy, twelve months after the initial diagnosis, demonstrated that chronic inflammatory infiltrate in the fundus mucosa is less pronounced, rich in plasma cells, with almost absent Russell body and mature plasma cells.

Figure 3  Immunohistochemical stains in Russell body gastritis (initial biopsy) × 200. A: The inflammatory infiltrate in the gastric fundus mucosa is CD79a positive, which is in support of its homogeneous plasmocytic nature; B: Kappa; C: Lambda. The plasma cells are polyclonal both kappa (B) and lambda (C) light chains are positive.

with almost absent RB and Mott cells (Figure 2C).

**DISCUSSION**

RBG is a rare form of chronic gastritis which mostly affects the antrum and has a male predominance[8,9]. The diagnosis is histologic, and it is characterized by accumulation of plasma cells containing RB and Mott cells in gastric mucosa. According to the
literature, mucosal infiltration with RB and Mott cells may be associated with infectious or autoimmune processes\cite{8,15,20}. The diagnostic and differential-diagnostic histological algorithm includes immunohistochemical detection of plasma cell nature of the inflammatory infiltrate (CD138 and CD79a positivity), as well as its polyclonality (both kappa and lambda light immunoglobulin M chains expression). These immunohistochemical profiles, as well as the absence of nuclear atypia, mitotic activity, lymphoepithelial lesions and monoclonal infiltrates are most in keeping with a benign process and not with a lymphoproliferative disease (lymphoma or myeloma).

Signet-ring stomach adenocarcinoma, where the cells resemble RB, but show nuclear atypia, mitotic activity and cytokeratin expression must also be excluded\cite{4,12,13}.

Once the histologic diagnosis and differential-diagnosis have been made, the pathologist must prove or rule out the association with \textit{H. pylori} infection. This is done with Gimsa staining or immunohistochemical detection.

Literature data for RBG histologic follow up are scarce.

We have the opportunity to follow up in three consecutive biopsies the histological evolution of gastric mucosa in RBG. Our results showed a tendency from decrease up to complete extinction of RB and Mott cells in this chronic gastritis over time and under the influence of treatment. Our histologic follow up results indicate that RBG is probably an inconstant and dynamic morphological finding developing within a rich of plasma cells chronic gastritis. Like other authors we observe decreased number of RB and Mott cells in the time\cite{8}. In contrast to this study, our results show that the reason for decreased number of RB in the stomach is not \textit{H. pylori} eradication. Probably, the factor causing RB formation is not only \textit{H. pylori} infection, other factors may also play role such as local degenerative or vascular-circulatory phenomena.

In our study, a total of 15 cases of RBG with polyclonal plasma cells, containing RB (Mott cells) have been described\cite{8}. Our case also showed polyclonal proliferation of plasma cells with RB with uneventful clinical follow up. The decreased number of Mott cells in the stomach after \textit{H. pylori} eradication shows that \textit{H. pylori} is one of the factors causing RB formation. In practical terms, the latter can be used as an additional diagnostic sign in contrast to the increased follow up distribution of RB in the context of multiple myeloma-associated RBG\cite{8}.

Diffuse infiltration of plasma cells with RB in the gastric mucosa requires differential diagnosis with several diseases. Cytokeratin negativity and CD79a positivity exclude signet-ring cell carcinoma of the stomach. Kappa and lambda polyclonal immunoreactive pattern exclude lymphoplasmacytic lymphoma\cite{8,15,20,21}, plasmacytoma and monoclonal gammopathy of undetermined significance\cite{8}. The lesion can be differentiated from MALT lymphoma by absence of nuclear atypia and lymphoepithelial lesions\cite{8,15,20}.

There are many unclear points about the etiology of RBG. According to Hasegawa\cite{21}, immunoglobulin accumulation could be in a result of an over or altered production as well as aberrant secretion and impaired excretion\cite{21}.

Most of the cases published in the literature have demonstrated a connection between \textit{H. pylori} infection and antigenic stimulation of plasma cells\cite{4,12}. On the other hand, \textit{H. pylori}-negative RBG is rarely reported. To the best of our knowledge, only twenty-one cases are published in the literature so far. There are insufficient data about the etiology and progress of this entity. A possible relationship between \textit{H. pylori}-negative RBG and the immune status has been proposed, with a number of cases reported in patients with HIV\cite{10,11,12,13}, alcohol\cite{10,12,13,20} and drug abuse\cite{12,13,16} and post-transplant patients\cite{18}.

Apart from chronic infections and immunosuppressive treatment, cancer could also be a trigger of immune dysregulation and RB and has been reported in patients with signet-ring cell carcinoma\cite{18}. For this reason, it is of great importance to be able to discriminate between cancer-induced mucosal changes and RBG. During upper endoscopy this entity should be kept in mind because of the vast majority of differential diagnosis such as plasma cell neoplasms, signet-ring cell carcinoma and MALT lymphoma. It is also of great significance to obtain biopsies according to Sydney system.

Our case is about 51-year-old man with iron-deficiency anemia and \textit{H. pylori}-negative RBG. We performed many diagnostic tests to rule out chronic infections, autoimmune diseases associated with B cell proliferation and different malignancies of gastric mucosa. We came to the conclusion that RBG in our patient is a benign process with uncertain prognosis and long-term clinical and endoscopic follow-up was scheduled. For a period of one year we observed histologic regression, most probably in a result of his 8-wk-treatment with PPI. Hence, PPI therapy could be considered as a feasible option of treatment for this rare type of gastritis.

In our article we present all published cases in English literature with \textit{H. pylori}-
negative RBG so far.

A summary of 22 reported *H. pylori*-negative RBG is described in Table 1. Aggregated data from all the reported cases show that patients are within the age range of 20 to 82 years with predominant age group between 70-80 years (40.9%). Male to female ratio was 2.14:1. Most of the patients had GI symptoms such as abdominal pain, dyspepsia and nausea. Endoscopic findings include vast spectrum of nonspecific features such as erythema, edema, erosions and ulcerations or well-formed nodular lesions. In eight of the patients *H. pylori*-negative RBG was localized in the gastric antrum (36.36%), in three of them it was in body of the stomach (13.63%), in the cardia (13.63%), in more than one region of the stomach (13.63%) and without specific localization in three of the patients (13.63%). According to our data, RBs distribution is the rarest in the fundus (9.09%) (Figure 4). Of all the twenty-two cases, two patients had anemia as concomitant disease, three reported alcohol abuse and three with HIV infection (Figure 5). Other associated conditions include multiple myeloma, chronic kidney failure, drug abuse, post kidney transplant, diabetes mellitus and colonic adenoma. Majority of the cases (ten) with *H. pylori*-negative RBG showed no evidence of concurrent medical conditions. In this literature review it is well visible that two of the cases of *H. pylori*-negative RBG are associated with lymphoproliferative disease (multiple myeloma) [6,24-28].

CONCLUSION

We would like to summarize that there are few cases of *H. pylori*-negative RBG described in the literature. This condition should be kept in mind during endoscopic surveillance and differentiated from other benign and malignant entities. To the best of our knowledge, our report is the first that present a long-term follow up in a patient *H. pylori*-negative RBG, treated with PPI. We came to conclusion that PPI therapy leads to significant reduction of RB and plasma cells in the gastric mucosa.

So far little is known about its etiology and pathogenesis, thus larger studies must be conducted. The outcome of chronic stimulation of the Mott cells is unknown, therefore it is of paramount importance to actively follow up these patients.
### Table 1: A summary of 22 reported *Helicobacter pylori*-negative Russell body gastritis

<table>
<thead>
<tr>
<th>Cases</th>
<th>Ref.</th>
<th>Age/sex</th>
<th>History</th>
<th>Endoscopic findings</th>
<th>Symptoms</th>
<th>Histology</th>
<th>Immunology</th>
<th>Follow up</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Erbersdobler <em>et al</em>[16]</td>
<td>80/Female</td>
<td>Alcohol and analgesic abuse, Candida esophagitis</td>
<td>Circumscribed, irregular mucosal swelling at the back side of the fundus (lesion up to 3 cm)</td>
<td>Epigastric pain and nausea</td>
<td>Confirmed candida and showed plasma cells with RBs</td>
<td>Poly NR</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>Drut <em>et al</em>[10]</td>
<td>34/Male</td>
<td>HIV+, Drug addict, Alcohol abuse</td>
<td>2-cm-raised area located at the major curvature of the stomach, presenting a central 1 cm rounded macule</td>
<td>Epigastric pain, acute diarrhoea, blood-stained stools</td>
<td>Moderate-to-severe gastritis with RBs</td>
<td>Poly NR</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>Habib <em>et al</em>[22]</td>
<td>75/Male</td>
<td>Alcohol use, Renal failure Dyslipidaemia, Rhabdomyolysis</td>
<td>Oesophagitis and nodular chronic active gastritis in the antrum</td>
<td>Reflux complaints, intermittent coffee-ground emesis</td>
<td>Regenerative changes and a dense chronic inflammatory infiltrate composed of numerous RBs</td>
<td>Poly NR</td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>Del Gobbo <em>et al</em>[7]</td>
<td>78/Female</td>
<td>NR</td>
<td>Hyperaemia in the antral and GEJ mucosa</td>
<td>Epigastric pain</td>
<td>Moderate chronic inflammation in the mucosa of the cardia showed RBs</td>
<td>Poly NR</td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>Coyne <em>et al</em>[13]</td>
<td>49/Male</td>
<td>Drug addict, HCV and Diabetes mellites</td>
<td>Severe erosive gastritis with oedematous mucosal folds</td>
<td>Nausea, epigastric pain, weight loss</td>
<td>RBG</td>
<td>Mono (κ chain, IgM)</td>
<td>NR</td>
</tr>
<tr>
<td>6</td>
<td>Bhalia <em>et al</em>[12]</td>
<td>82/Male</td>
<td>HIV+</td>
<td>Gastritis</td>
<td>Dyspepsia, loose stools, loss of appetite and weight</td>
<td>RBs present in gastric mucosa</td>
<td>Poly NR</td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>Klair <em>et al</em>[24]</td>
<td>76/Female</td>
<td>Anemia, Multiple myeloma</td>
<td>Multiple small polyps in the fundus were seen on retroflexion, along with cobblestoned erythematous and irregular mucosa</td>
<td>Bone pains and adynamia</td>
<td>Oxyntic mucosa with chronic, inactive gastritis, with plasma cells</td>
<td>Poly NR</td>
<td></td>
</tr>
<tr>
<td>8</td>
<td>Zhang <em>et al</em>[17]</td>
<td>78/Male</td>
<td>NR</td>
<td>Gastritis with uneven mucosa in the antrum, corpus and incisura angularis</td>
<td>Heartburn</td>
<td>RBG with moderate chronic inflammation</td>
<td>Mono (κ chain)</td>
<td>Clinical follow-up evaluations were uneventful</td>
</tr>
<tr>
<td>9</td>
<td>Zhang <em>et al</em>[17]</td>
<td>28/Male</td>
<td>NR</td>
<td>Erythema in antrum</td>
<td>Epigastric pain</td>
<td>RBG with mild chronic inflammation</td>
<td>Mono (κ chain)</td>
<td>NR</td>
</tr>
<tr>
<td>10</td>
<td>Zhang <em>et al</em>[17]</td>
<td>24/Female</td>
<td>NR</td>
<td>Erythema in antrum</td>
<td>Abdominal discomfort</td>
<td>RBG with mild chronic inflammation</td>
<td>Mono (κ chain)</td>
<td>NR</td>
</tr>
<tr>
<td>11</td>
<td>Zhang <em>et al</em>[17]</td>
<td>66/Male</td>
<td>NR</td>
<td>Ulceration stage A2 in Forrest classification in incisura angularis</td>
<td>Haematochezia</td>
<td>RBG with moderate glandular atrophy and mild chronic inflammation</td>
<td>NR</td>
<td>NR</td>
</tr>
<tr>
<td>12</td>
<td>Muthukumarana <em>et al</em>[23]</td>
<td>44/M</td>
<td>Status post pancreatic and Kidney transplant Diabetes mellites</td>
<td>Diffuse mild erythematous gastric mucosa, non-crateted duodenal ulcer</td>
<td>Watery diarrhoea with abdominal pain, nausea and vomiting</td>
<td>Stomach, duodenum, terminal ileum, colon mucosa with RBs</td>
<td>Poly NR</td>
<td></td>
</tr>
<tr>
<td>13</td>
<td>Saraggi <em>et al</em>[20]</td>
<td>66/Male</td>
<td>NR</td>
<td>Los Angeles class A esophagitis. Multiple biopsy has been taken</td>
<td>Heartburn</td>
<td>Mild lymphoplasmacytic inflammation in the mucosa of the cardia</td>
<td>Poly NR</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Authors</td>
<td>Age</td>
<td>Sex</td>
<td>Diagnosis</td>
<td>Symptoms</td>
<td>Findings</td>
<td>Follow-up</td>
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</tr>
<tr>
<td>14</td>
<td>Antunes et al. [20]</td>
<td>79</td>
<td>Female</td>
<td>NR</td>
<td>Hematemesis; RBG</td>
<td>NR; NR</td>
<td></td>
<td></td>
</tr>
<tr>
<td>15</td>
<td>Imai et al. [27]</td>
<td>64</td>
<td>Male</td>
<td>Chronic renal failure on dialysis</td>
<td>Flare, swollen mucous membrane and multiple verrucous erosion in gastric antrum</td>
<td>Poor appetite and blood eosinophilia; Infiltration of plasma cell containing RBs and eosinophils</td>
<td>NR</td>
<td></td>
</tr>
<tr>
<td>16</td>
<td>Trna et al. [28]</td>
<td>77</td>
<td>Male</td>
<td>NR</td>
<td>Several areas of different and mildly prominent mucosa in the GEJ and cardia</td>
<td>Non-cardiac chest pain and mild dysphagia; Nondysplastic intestinal metaplasia with mild chronic inflammatory infiltrate with RBs and plasma cells</td>
<td>NR; Follow-up endoscopy with biopsies - without any difference</td>
<td></td>
</tr>
<tr>
<td>17</td>
<td>Altindag et al. [6]</td>
<td>81</td>
<td>Female</td>
<td>Multiple myeloma (diagnosed from bone marrow 3 years after endoscopy)</td>
<td>Gastritis in the antrum; Dyspepsia</td>
<td>Mild inflammation of gastric mucosa with RB; Poly</td>
<td>NR; Histology report revealed increased distribution in RBs in follow-up endoscopy</td>
<td></td>
</tr>
<tr>
<td>18</td>
<td>Altindag et al. [6]</td>
<td>79</td>
<td>Female</td>
<td>NR</td>
<td>Gastritis in the antrum and gastric tubular adenoma with LGD</td>
<td>GI bleed; Mild glandular atrophy, moderate intestinal metaplasia, severe inflammation of gastric mucosa with RB</td>
<td>Poly; NR</td>
<td></td>
</tr>
<tr>
<td>19</td>
<td>Altindag et al. [6]</td>
<td>72</td>
<td>Male</td>
<td>NR</td>
<td>Gastritis in the antrum</td>
<td>Dyspepsia; Mild inflammation of gastric mucosa with RB and mild glandular atrophy</td>
<td>Poly; NR</td>
<td></td>
</tr>
<tr>
<td>20</td>
<td>Altindag et al. [6]</td>
<td>64</td>
<td>Male</td>
<td>Colonic tubular adenoma, HGD</td>
<td>Gastritis in the antrum</td>
<td>Epigastric pain, suspicion of gastric tumor; Moderate glandular atrophy, moderate intestinal metaplasia and moderate inflammation of gastric mucosa with RB</td>
<td>Poly; NR</td>
<td></td>
</tr>
<tr>
<td>21</td>
<td>Qiao et al. [11]</td>
<td>28</td>
<td>Male</td>
<td>HIV+, pancytopenia, splenomegaly, hepatomegaly</td>
<td>Erosions, erythematous mucosa, and vascular congestion in the gastric body and antrum</td>
<td>Abdominal pain, fatigue, rectal bleeding; Chronic inactive gastritis with RB infiltration in the mucosa</td>
<td>Poly; NR</td>
<td></td>
</tr>
<tr>
<td>22</td>
<td>Present study</td>
<td>52</td>
<td>Male</td>
<td>Anemia</td>
<td>Diffuse hyperemia and edema of the gastric mucosa in the fundus and body</td>
<td>Iron-deficiency anemia; Abundant plasma cell inflammatory infiltrate, rich in RB and Mott cell</td>
<td>Poly; Without endoscopic improvement, histology report showed decreased RB in second follow-up and almost absent RB in third follow-up</td>
<td></td>
</tr>
</tbody>
</table>

Gl: Gastrointestinal; NR: Not reported; RB: Russell body; IgA: Immunoglobulin A; GEJ: Gastro-oesophageal junction; RBG: Russell body gastritis.
Figure 4 Distribution of Russell bodies in the stomach: in all cases from the literature. RB: Russell body.

Figure 5 Associated conditions in patients with Helicobacter pylori-negative Russell body gastritis according to the available literature.

REFERENCES


Surveilling Russell body H. pylori-negative gastritis


