Abstract

The aim of the study was to ascertain the existence of intestinal metaplasia in gastric mucosa of patients with gastric carcinoma coupled with *H. pylori* positive chronic atrophic gastritis and possible connection of IM with the development of gastric carcinoma. The paper presents prospective study that included 50 patients with gastric carcinoma and 50 patients with chronic atrophic *H. pylori* positive gastritis. All the patients were subjected to gastroscopy as well as biopsy targeted at antrum, lesser curvature and corpus and at the area 1-2 cm removed from tumor lesion. Biopsy samples were sliced by microtome and stained. We analyzed presence, frequency and severity of inflammatory-regenerative, metaplastic and dysplastic changes in the mucosa and evaluated their prognostic value. We typed IM immunohistochemically.

This study confirmed responsibility of *H. pylori* for inflammatory events in gastric mucosa in patients with gastric carcinoma. According to our findings incomplete IM of types IIa and IIb as precancerous lesion is responsible for the development of gastric carcinoma and is associated with chronic atrophic gastritis grade I and II (92% of subjects, $p=0.0007$, $h=1$, $p=0.01$). Thus, the finding of incomplete intestinal metaplasia may be used as an indicator for early gastric carcinoma detection. Patients with patho-histologically verified incomplete intestinal metaplasia associated with active chronic atrophic gastritis of levels I and II represent risk group for the development of gastric carcinoma of intestinal type.

KEY WORDS: intestinal metaplasia (IM), epithelial dysplasia, *Helicobacter pylori* (*H. pylori*), chronic atrophic gastritis (CAG).
INTRODUCTION

_H. pylori_ induced chronic atrophic gastritis (CAG) is a precancerous condition accompanied by gradual loss in epithelia differentiation that eventually leads towards carcinoma. Intestinal metaplasia (IM) as precancerous lesion is either partial or complete transformation of gastric glands epithelial cells into the intestinal type. It may appear either in antrum or pylorus mucosa or diffusely. Various classifications are based on morphological, histochemical, enzymehistochemical, immunohistochemical and ultrastructural characteristics (1, 2).

Complete type of mature IM is characterized by histology, mucine histochemistry, enzymes and endocrine cells identical to those found in small intestine. Pathohistological features of this type are columnar absorptive cells, multiplied goblet cells that contain predominantly acid mucines, Paneth’s non-differentiated cells and M-cells. IM typing was performed. All the patients were examined for the presence of inflammatory-regenerative changes in gastric mucosa. Based on the defined histological criteria we graded inflammatory-regenerative changes as chronic superficial or chronic atrophic gastritis of the degree I, II and III. The degree of inflammation activity was determined based on leukocyte infiltration. It was defined either as active or dormant phase. When the changes were graded as inflammatory-regenerative or dysplastic the tissue was prepared for immunohistochemical analysis. The tissue was HE stained prior to the processing. IM was assigned to one of the three possible categories based on the detailed histochemical analysis of mucine type, PAS and HID/AB staining and morphological changes. The degree of IM dispersal was also analyzed. IM was considered focal when found in 1/3 of the biopsic sample, medium when found in 2/3 of the sample and severe when dispersed over the entire sample. The mucine type was examined within goblet cells that was either well developed, underdeveloped or no brush border. Paneth’s cells are either absent or extremely rare. Immature IM type is encountered in severe atrophic forms of gastritis due to rapid desquamation and regeneration as well as delayed cell differentiation. Hyperplastic glands laid with juvenile non-differentiated cells with hyper-chromic nuclei and increased mitotic index develop in the process. The non-differentiated cells may differentiate into various cell types in the presence of various promoters. IM is believed to develop due to the absence of antioxidative activities in the mucosa (3, 4, 5).

An issue is raised of whether _Helicobacter pylori_ ( _H. pylori_ ) presence is necessary for the development of IM. It is known that _H. pylori_ does not dwell in the area of advanced IM as it is not found in biopsic samples collected from antrum where the atrophy is pronounced. _H. pylori_ is found in the mucosa in B gastritis and IM. Positive correlation between _H. pylori_ infection and IM type II is noted in patients with gastric carcinoma.

SUBJECTS AND METHODS

This paper presents prospective study, which includes three viewpoints at the possibility of gastric adenocarcinoma development: clinical, pathohistological and microbiological. The study included 50 patients with gastric carcinoma that are treated at the Gastroenterohepatology Clinic. All the patients were subjected to endoscopic examination where stomach was macroscopically divided into sections: corpus, antrum, antropyloric region and pylorus. The patients with cardiac and distal esophagus carcinoma were not included in the study. Following the macroscopic verification of the tumor we performed biopsy of the tumoral formation, as well as biopsy of the mucosa in the region 1-2 cm removed from the tumor. We also collected mucosa samples from corpus, antrum and pyloric region. Biopsy mucosa samples were conserved in 10% buffered neutral formalin, paraffin embedded and sliced with microtome into 5 µm thick sections. The sections were stained with Heamatoxylin and Eosin (HE staining), immunohistochemically for p53 antibodies. IM typing was performed. All the patients were examined for the presence of inflammatory-regenerative changes in gastric mucosa. Based on the defined histological criteria we graded inflammatory-regenerative changes as chronic superficial or chronic atrophic gastritis of the degree I, II and III. The degree of inflammation activity was determined based on leukocyte infiltration. It was defined either as active or dormant phase. When the changes were graded as inflammatory-regenerative or dysplastic the tissue was prepared for immunohistochemical analysis. The tissue was HE stained prior to the processing. IM was assigned to one of the three possible categories based on the detailed histochemical analysis of mucine type, PAS and HID/AB staining and morphological changes. The degree of IM dispersal was also analyzed. IM was considered focal when found in 1/3 of the biopsic sample, medium when found in 2/3 of the sample and severe when dispersed over the entire sample. The mucine type was examined within goblet and absorptive cells. Mucines may be either neutral or acid. HID/AB staining enables differentiation of acid mucines into sulfo- or sialomucins (sulfomucin – brown, sialomucin – blue). In PAS method acid mucines are stained blue while neutral mucines are stained red. We registered brush border on the surface of absorptive cells that was either well developed, underdeveloped or completely missing. Paneth’s cells were also identified and graded as numerous, reduced or missing. Villi development was also graded as: well developed, underdeveloped and undeveloped. In IM type I, mucosa architecture is mainly well preserved as well as Paneth’s cells abundance. Goblet cells mostly secrete sialomucines, some sulfomucines, while absorptive cells have no secretionary properties. Brush border at their surface is preserved. Mucosa surface is not villous although it may appear slightly villous in small number of cases. In IM type II mucosa architecture is significantly degraded, villi are well developed, glandular crypts are elongated and curved and coated in tall columnar cells with either underdeveloped or no brush border. Paneth’s cells are ei-
ther rare or missing. Goblet cells predominantly secrete sialomucines while poorly differentiated absorptive cells secrete either sulfomucines or neutral mucines. Absorptive cells are scarce. In IM type III mucosa architecture is completely degraded while the surface is exceptionally villous. Mucosa surface glands are elongated and curved to a higher degree compared to those in the IM type II. Gland branching and distortion is evident. Goblet cells are scarce and secrete mainly acid mucines, sulfo- and sialomucines, while poorly differentiated absorptive cells contain sulfomucines. Brush border is absent from absorptive cells. Paneth’s cells are missing. The control group included 50 patients suffering from chronic H. pylori positive gastritis. We endoscopically aimed biopsy at antrum, lesser curvature and corpus mucosa. We applied the same methodology as in the cases of gastric carcinoma in conserving, embedding and staining of the samples as well as in the analysis of inflammatory-regenerative and dysplastic changes in gastric mucosa. We used immunohistochemical methods in IM typing. HUT testing was used for verification of H. pylori presence in gastric mucosa in the areas of antrum and corpus.

RESULTS

The study included 100 subjects divided into two groups. The first group consisted of 50 patients with stomach carcinoma, 28 male and 22 female. Average age of patients was 58 for males and 62 for females. Anatomically, 66% of carcinoma formations were endoscopically identified in the distant part of the stomach. Intestinal metaplasia was differentiated according to regions.

IM in antral region was identified in 32 subjects (64%). 31.13 % of the subjects were suffering from superficial chronic gastritis in active phase. 65.63 % of the subjects had intestinal metaplasia associated with chronic atrophic gastritis grade I. 56.25 % of those were in active phase. 32 % had intestinal metaplasia associated with chronic atrophic gastritis grade II in active phase. IM was identified in 24 patients with epithelial dysplasia. IM was associated with slight epithelial dysplasia in 18 cases and with mild epithelial dysplasia in 6 cases. Correlation is significant and high, p=0.0356, d=22. Rank correlation (Wilcoxon test) between the level of dysplasia and IM in antral region was significant and high (significance level p=0.05). IM in the region around carcinoma was confirmed in 30 subjects (60%). In 80 % of cases IM was associated with chronic atrophic gastritis grade I. All of those were in active phase. In 20% cases IM was associated with chronic atrophic gastritis grade II in active phase. With respect to epithelial dysplasia, it was slight in 13 IM patients and mild in 7 cases. In 10 patients epithelial dysplasia was not found. Rank correlation (Wilcoxon test) between IM and the degree of epithelial dysplasia was not significant (p=0.05). IM in the region of corpus was found in 7 patients (14%). In 3 patients IM was associated with superficial and in 4 patients with CAG grade I and grade II. In 14.29 % patients gastritis was in active phase. In the corpus region correlation between IM and gastritis grade was significantly high at p=0.01 level of significance. Rank correlation (Wilcoxon test) between IM and the degree of epithelial dysplasia was high and significant (p=0.05). Control group included 29 men and

<table>
<thead>
<tr>
<th>Region</th>
<th>Superficial</th>
<th>CAG gr. I</th>
<th>CAG gr. II</th>
<th>CAG gr. III</th>
</tr>
</thead>
<tbody>
<tr>
<td>Intestinal Metaplasia</td>
<td>1 (3.13%)</td>
<td>21 (65.63%)</td>
<td>10 (32%)</td>
<td>0</td>
</tr>
<tr>
<td>Active Phase</td>
<td>1 (3.13%)</td>
<td>18 (56.25%)</td>
<td>10 (32%)</td>
<td>0</td>
</tr>
<tr>
<td>Dormant Phase</td>
<td>0</td>
<td>3 (9.32)</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>

**Table 1.** Distribution of intestinal metaplasia according to chronic gastritis grade and the level of activity in the group of subjects – antral region

<table>
<thead>
<tr>
<th>Region</th>
<th>Superficial</th>
<th>CAG gr. I</th>
<th>CAG gr. II</th>
<th>CAG gr. III</th>
</tr>
</thead>
<tbody>
<tr>
<td>Intestinal Metaplasia</td>
<td>0</td>
<td>0</td>
<td>24 (80%)</td>
<td>6 (20%)</td>
</tr>
<tr>
<td>Active Phase</td>
<td>0</td>
<td>24 (80%)</td>
<td>6 (20%)</td>
<td>0</td>
</tr>
<tr>
<td>Dormant Phase</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>

**Table 2.** IM distribution according to chronic gastritis grade and the level of activity in the group of subjects – region around carcinoma

<table>
<thead>
<tr>
<th>Region</th>
<th>Superficial</th>
<th>CAG gr. I</th>
<th>CAG gr. II</th>
<th>CAG gr. III</th>
</tr>
</thead>
<tbody>
<tr>
<td>Intestinal Metaplasia</td>
<td>3 (52.86%)</td>
<td>3 (42.96%)</td>
<td>1 (14.29%)</td>
<td>0</td>
</tr>
<tr>
<td>Active Phase</td>
<td>2 (28.57%)</td>
<td>2 (28.57%)</td>
<td>1 (14.29%)</td>
<td>0</td>
</tr>
<tr>
<td>Dormant Phase</td>
<td>1 (14.29%)</td>
<td>1 (14.29%)</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>

**Table 3.** IM distribution according to chronic gastritis grade and the level of activity – corpus region
21 women, 60 and 61 years average age respectively. In the control group 19 subjects had IM in the region of antrum (38%). In 68.42 % subjects IM was associated with CAG grade I in active phase. In 19 subjects it was associated with epithelial dysplasia. Correlation between IM and epithelial dysplasia was significantly high at $p<0.05$ significance threshold. Intestinal metaplasia in the region of lesser curvature was found in 8 subjects (16%). Of those, 62 % had CAG grade I. Active phase was ascertained in 37.5 % subjects. Rank correlation was significant between the degree of epithelial dysplasia and IM presence at $p<0.05$ significance threshold. IM in the corpus region was found in one patient along with CAG grade I in active phase. Correlation between IM and epithelial dysplasia was not significant.

### DISCUSSION

Progression of chronic active gastritis is accompanied by changes in phenotype of gastric mucosa epithelial cells. Under the influence of the modified environment epithelial metaplasia occurs due to gastric mucosa capacity for regenerative and metaplastic changes. *H. pylori* is believed to promote hyper-proliferation of epithelial cells by its presence and action upon mucosa (6,7,8). Proliferative properties of germinative cells ensure regeneration of surface layer cells in normal mucosa. These cells, referred to as base or stem cells, are located in lower segments of epithelial glands, which explains absence of mitotic activity in surface layer of mucosa (9). Known properties of these cells include auto-regulation, which accounts for adaptive reaction to tissue damage. These cells divide slowly; they are extremely sensitive to radiation and are programmed to self-destruct when their DNA pattern is damaged. They have the ability to preserve genome by selectively maintaining original, paternal DNA copy (10,11,12). Other cell types differentiate from those cells since they are multipotent and located in neck parts of glands in the lower crypts segments. Compensation of the lost cells in crypts requires time for regeneration. Cells migrate from the neck part upwards towards the surface. Glandular atrophy occurs in the scope of prolonged chronic atrophic gastritis as inflammation spreads and affects deeper mucosa layers. In atrophic gastritis, gastric mucosa is continuously desquamated (peeled off) and regenerated (renewed). This process is accompanied by compromised differentiation, atypical regeneration, which may lead to carcinoma. The measure of gastritis activity is presence of polymorphonuclear infiltrate, which correlates with *H. pylori* finding. It is widely believed that *H. pylori* is an important factor in gastritis progression. Along with the progression of atrophy, gastric mucosa suffers loss of gastric antral and corpus glands. In the process, either partial or complete transformation of gastric glands epithelial elements into intestinal type occurs. In the analysis of the mucosa we registered absorptive enterocytes that alternate with sialomucines secreting goblet cells. Such type of epithelial metaplasia is referred to as complete or intestinal metaplasia type I (13,14). Further progression is marked with glands being coated with columnar cells, scarce presence of Paneth’s cells and sialomucines presence in goblet cells. Absorptive cells are less differentiated and secrete sulfomucine and neutral mucine. This type is denoted IM type II. In IM type III mucosa surface is extremely villous, goblet cells are scarce and secrete acid, sulfo- and sialomucines, while less differentiated absorptive cells secrete sulfomucine. Thus, changes in gastric cells phenotype accompany inflammation and atrophy of gastric mucosa. The normal gastric epithelium is replaced by cells with morphological properties resembling those of the intestinal type. IM progression generates dysplasia which may progress into gastric carcinoma (15). Gastric mucosa atrophy introduces marked increase in cell replication along with the expression of fetal antigens, which influence loss of cell differentiation. Our research targeted verification of the presence, severeness and frequency of intestinal metaplasia occurrence in various regions of gastric mucosa. We found intestinal metaplasia in antral region in 32 out of 50 patients (64%) with gastric carcinoma. In 21 subjects it was associated with chronic atrophic gastritis grade I, which was in active phase in 18 subjects. In 10 subjects we found chronic atrophic gastritis grade II in active phase. We found 30 cases of IM in the region around carcinoma (60%). 24 patients had CAG grade I while 6 had CAG grade II, all of those in active phase. In corpus region we found 7 IM (14%). In 3 cases IM was superficial, 3 had CAG grade I and one had CAG grade II. Four of those were in active phase. In the control group, for compar-
son, IM in antral region was found in 19 out of 50 examined patients (38%). Of those 13 (26%) had CAG grade I, 5 (10%) had CAG grade II and one had superficial gastritis. All of those were in active phase. 8 (16%) cases of IM in lesser curvature region were identified. Five of those were associated with CAG grade I, one with CAG grade II and two with superficial gastritis. We found one case of IM in corpus region associated with CAG grade I in active phase. Comparing with the group of patients with gastric carcinoma, no significant difference in distribution of intestinal metaplasia in antral region was found in our research. There is significant difference in frequency (p=0.0007, h=1 for significance threshold p=0.01). Significant difference exists in frequency of IM in corpus region. IM is more frequent in patients with gastric carcinoma. The difference is significant with p=0.00278 at significance threshold p=0.01. We used immunohistochemical staining to type IM. 17 (34%) subjects had complete IM type I. 20 (40%) of them had incomplete IM type IIa. Four (8%) of patients had incomplete metaplasia type IIb. According to the above, IM was found in 41 out of 50 patients with gastric carcinoma. In 27 (54%) patients it was associated with CAG grade I, of those, 48% were in active phase. In 11 (22%) patients we found CAG grade II in active phase. In 17 (34%) complete IM was focal. Incomplete IM type IIa was of focal appearance in 4 (8%) subjects. In 14% of subjects it spread over 2/3 of the mucosa sample while 18% subjects had diffuse form. Incomplete IM type IIb took up 2/3 of the sample in 2 (4%) patients. In 2 (4%) patients it was diffuse. Goblet cells more frequently secreted acid sialomucines in incomplete IM type IIa and IIb. Absorptive cells secreted neutral or acid sulfomucines, with their brush border less developed and Paneth’s cells either absent or scarce. In the control group, complete IM type I was found in 13 (26%) subjects while incomplete IM type IIa was found in 7 (14%) patients. They were also associated with CAG grade I and II in active phase. IM was predominantly located in antral region – in 16 of 20 (80%) control group subjects. Four subjects had IM in corpus region. IM was focal in 4 (8%) subjects, spread over 2/3 of sample in 15 (30%) and diffuse in one subject. In 18 (36%) subjects goblet cells secreted acid sialomucine. They were found to secrete acid sulfomucine in 2 (4%) subjects. Absorptive cells secreted acid sialomucine in 2 (4%) subjects, acid sulfomucine in 5 (10%) subjects and neutral mucines in 10 (20%) subjects. Brush cover was well developed in 5 (10%), underdeveloped in 13 (26%) and absent in 2 (4%) subjects. Paneth’s cells were absent in 9 (18%) and scarce in 11 (22%) subjects. We were not able to find significant differences in Wilcoxon rank test between groups per regions by gastritis type (p=0.4192, h=0). There is significant difference between groups in IM occurrence in antral region with p=0.0007, h=1 at the significance threshold p=0.01. No significant difference in gastritis presence in corpus region was found. Analyzing earlier published studies we found that Reis (13) studied IM and concluded that classic sequential paths from IM I to IM III lead through IM II. In our research we found incomplete IM type IIb (type III according to certain authors) in only 4 (8%) patients with gastric carcinoma. No IM type III cases were found in the control group. Safele-Ribiero stated IM type I as the most frequent in their studies. It is suggested (16,17) suggest that metaplastic cell lines ensue as a response of gastric mucosa to chronic damage and precede epithelial dysplasia and gastric adenocarcinoma. In our subjects, we found IM type I in 34% patients with gastric carcinoma and in 26% patients with H. pylori positive gastritis. 40% patients had incomplete IM or type II while 8% had IM type III. In the control group though, 14% subjects had incomplete IM type II. In our research IM was identified in 41 (82%) patients with gastric carcinoma. Incomplete IM type IIa was more frequent in patients with gastric carcinoma. According to the severeness of the changes incomplete IM types IIa and IIb were either diffuse or spread over 2/3 of the biopsic specimen width. In comparison with the control group no significant difference in IM frequency in antral region was found. In the region around carcinoma IM was found in 39 (60%) of the subjects and it was associated with chronic atrophic gastritis grade I and II in active phase.

CONCLUSION

Based on the results of our research we may conclude that IM as precancerous lesion is responsible for the development of gastric carcinoma. When associated with chronic atrophic gastritis grade I and II it can serve as an indicator of the events that lead to intestinal gastric carcinoma development. Special attention is required in patients with pathohistologically verified incomplete IM type IIa associated with slight or mild epithelial dysplasia due to its significantly frequent occurrence in our patients with gastric carcinoma.
References


