EFFICIENCY AND SAFETY OF SILICON-CONTAINING ENTEROSORBENT IN THE TREATMENT OF IRRITABLE BOWEL SYNDROME

Tkachenko E.I., Avalueva E.B., Skazyvaeva E.V., Ivanov S.V., Pushkina A.V., Lapinskii I.V.

1North-West State Medical University named after I.I.Mechnikov, Saint-Petersburg, Russia


1North-Western State Medical University named after I.I. Mechnikov, Kirochnaya str. 41, St.-Petersburg 191015, Russian Federation. Phone: 8 (812) 303-50-00, fax: 8 (812) 303-50-35. E-mail: rectorat@spbmapo.ru

SUMMARY:
Introduction: Irritable Bowel Syndrome (IBS) is a psychosocial functional gastrointestinal disorder significantly reducing the patient’s quality of life. The study objective was to assess the efficiency of using silicon-containing enterosorbent (Enterosgel) in IBS patients with diarrhea.

Materials and methods: 30 patients with diarrhea type of IBS (16 women and 14 men) were included in the study. All the patients were divided into two groups: the main group was treated by the enterosorbent "Enterosgel" (dosing 22.5 g 3 times a day, 1 hour after meals for 21 day), the reference group was treated with bismuth tripotassium dicitrate 1 tablet 3 times a day 30 minutes before meals and 1 tablet at night for 21 days.

To assess the treatment efficiency, patients’ complaints were evaluated using GSRS questionnaire, stool frequency and form were assessed as well as
myoelectric activity of the digestive tract (apparatus "Gastroenteromonitor-GEM-01").

Results: As a result of the treatment, patients of the main group have achieved a statistically significant reduction in the severity of complaints of abdominal pain, diarrhea syndrome, dyspepsia scales, and on the total GSRS scale. After therapy with the comparator, in patients of the reference group intensity of abdominal pain, reflux syndrome, diarrhea syndrome and dyspepsia significantly decreased. The most clinically significant changes observed in patients of the main group included normalized stool frequency (slowing to 7 times a week), normalization of stool form (up to type 4 on the Bristol scale) and a decrease of pain severity.

Conclusion: There were no significant differences between groups in assessing the digestive tract MEA, however, the aberrant GIT response of IBS patients with diarrhea to food intake was revealed: a decrease instead of an expected increase of the MEA was noted.

So, the study found that application of Enterosgel reduced the complaints of IBS patients with diarrhea and abdominal pain severity, thus contributing to the harmonization of quality of patient’s life. The treatment regimen, which can be recommended for use in patients with diarrhea-predominant IBS is: orally for 3 consecutive weeks at a dose of 22.5 g 3 times a day 60 minutes before meals (a dose of paste diluted in 200 ml of cool boiled water or washed down with 200 ml of cool boiled water).

Keywords: enterosorption, Enterosgel, irritable bowel syndrome, myoelectrical activity

Information about the authors:
Tkachenko Evgenii Ivanovich – MD, head of Department of medical microbiology, North-Western State Medical University named after I.I. Mechnikov, Saint Petersburg, Russia. Mobile phone: 8 (812) 543-95-38, e-mail: tkachenkoe@mail.ru.
Avalueva Elena Borisovna – MD, professor of Department of internal propaedeutic, North-Western State Medical University named after I.I. Mechnikov, Saint Petersburg, Russia. Mobile phone: 8 (921) 994-28-07 e-mail: avalueva@mail.ru.

Skazyvaeva Ekaterina Vasiljevna – PhD, associate professor of Department of internal propaedeutic, North-Western State Medical University named after I.I. Mechnikov, Saint Petersburg, Russia. Phone: 8 (812) 543-95-38, e-mail: skazyvaeva@yandex.ru.

Ivanov Sergey Vitalyevich – PhD, associate professor of Department of internal propaedeutic, North-Western State Medical University named after I.I. Mechnikov, Saint Petersburg, Russia. Phone: 8 (905) 209-60-01, 8 (812) 543-95-38, e-mail: ivanov.sv@mail.ru.

Lapinskii Igor Vadimovich – postgraduate student, North-Western State Medical University named after I.I. Mechnikov: Department of internal propaedeutic, Saint Petersburg, Russia. Mobile phone: 8 (911) 734-09-80, e-mail: lapinsky85@yandex.ru.

Pushkina Anna Vasilievna – postgraduate student, North-Western State Medical University named after I.I. Mechnikov: Department of internal propaedeutic, Saint Petersburg, Russia. Mobile phone: 8 (981) 701-76-23, e-mail: pushkina.anna.md@mail.ru.

Medvedeva Olga Ivanovna – PhD, associate professor of Department of internal propaedeutic, North-Western State Medical University named after I.I. Mechnikov, Saint Petersburg, Russia. Phone: 8 (812) 543-95-38, e-mail: olga.medvedeva@spbmapo.ru.

**INTRODUCTION**

Irritable bowel syndrome refers to psychosocial functional gastrointestinal disorders manifested by recurrent abdominal pain and impaired bowel evacuation frequency in the absence of its organic changes [1]. Deterioration of QOL of IBS patients influences the disability duration and, as a consequence, deteriorates
economic and social losses; it should be noted that the highest incidence is among young working people (at the age from 24 to 41) [2].

Nowadays, considering the absence of the unified diagnostic test (marker), it is a common practice to verify IBS diagnosis according to Rome criteria (version III), where the disease is defined as recurrent abdominal pain or discomfort for at least three days per month for three consecutive months that should be noted for at least six months. IBS is associated with the following two or more characteristics:

1) Improvement after defecation

2) Change in stool frequency during the disease

3) Change in stool form with the disease onset [17]. Considering Bristol stool form scale and Rome III criteria, it has been proposed to divide IBS patients into the following groups: IBS with constipation, IBS with diarrhea, mixed type of IBS. It is important to note that in a patient different IBS subtypes can vary throughout the illness.

Application of Roman criteria allows to adhere to a specific algorithm for IBS verification, which is caused by insufficient study of its pathophysiology. Most researchers tend to believe that in IBS pathophysiology violation of colonic motility, visceral hypersensitivity (threshold of nociception or densification of pain impulses with the normal sensitivity threshold) and changes in the functioning of links along the brain-gut axis are crucial. In the recent years, it has been found that IBS pathogenesis is more complicated. Such factors as inflammation [4] (including development and maintenance of mild inflammation in the intestinal wall, for example in patients with IBS postinfection [5]), immunological [6] and dietary factors [7], colon microbiota disorders [8] as well as changes in enteroendocrine system and its cells [9] play an important role in the disease progression [10].

To date, IBS treatment schedules are not standardized, and there is no single treatment algorithm of various IBS forms. It is believed that this is due to the problem of search and objective assessment of the drug efficiency and is associated with lack of knowledge, IBS pathophysiology complexity and a variety of clinical manifestations. On administration of treatment for IBS patients, a detailed study of the history of the disease and differentiation of complaint groups (abdominal, general, psychological, etc.) is of great importance, which will largely determine the tactics of therapy. Normalization of lifestyle and diet regime, psychotherapy are important components in the treatment of this disease. For IBS treatment, various groups of drugs are used, including antispasmodics, laxatives, anti-diarrheal drugs, pro-and prebiotics, antidepressants.
A search of new effective ways of complex IBS therapy, in particular the use of enterosorbsents, seems promising. Sorbents are substances possessing high adsorption capacity, which are not degraded in the GIT and capable of binding exogenous and endogenous toxic substances (microorganisms and their toxins, poisons, excessive metabolites and other harmful substances) by adsorption and absorption, ion exchange or complexation. The story of enterosorbents goes back to antiquity: healers of the Ancient Egypt, India and Greece already used charcoal, clay, pounded tuffs and dead-burnt horn for the treatment of poisoning, dysentery, jaundice and other diseases. With this purpose, birch or bone charcoal was used by the healers of ancient Russia [11].

The term "enterosorption" was proposed by Nikolaev V.G. et al. in 1983 [12, 13] to describe a new method of sorption therapy consisting of daily oral administration of large (20-50 g) doses of highly active synthetic coal spherical granulation obtained by pyrolytic processing of various polymer resins. It is also important that the authors have identified classical property of enterosorbents – non-invasion of the gastrointestinal mucosa, i.e. absence of systemic pharmacokinetics [13].

Among the considered drug class, siliceous sorbents are especially noteworthy, Enterosgel being the most widespread among them and known for its selective properties.

**Study objectives**

The purpose of this study is to determine efficiency and safety of application of Enterosgel, indicated for treatment of patients with diarrhea-predominant irritable bowel syndrome.

**Materials and methods**

The study was conducted in the gastroenterology clinic of Northwest State Medical University named after I.I. Mechnikov. 30 patients with diarrhea-prevalent IBS (D-IBS) confirmed according to Rome III criteria were included in the study. The average age of the patients was 38.0 ± 10.2 years. Gender distribution of patients: men - 14, women - 16. The patients were divided into 2 groups: the main group (n = 15): treatment with Enterosgel one dose of pasta 22.5 g (active substance polymethylsiloxane polyhydrate) three times a day after meal during three weeks; control group: treatment - bismuth tripotassium dicitrate. The drug was
administered at a dose one tablet three times a day 30 minutes before meals and one tablet in the evening for 21 days.

Manifestation dynamics of gastrointestinal complaints was assessed using the standard questionnaire and GSRS questionnaire. GSRS questionnaire (Gastrointestinal Symptom Rating Scale) has been developed by department of studies on life quality (QOL), ASTRA Hassle (I.Wiklund, 1998) and is used for QOL assessment of patients with gastrointestinal diseases. The questionnaire consists of 15 points organized in five scales: abdominal pain (questions 1, 4), reflux syndrome (questions 2, 3, 5), diarrheal syndrome (questions 11, 12, 14), dyspeptic syndrome (questions 6, 7, 8, 9), constipation syndrome (questions 10, 13, 15), total measurement scale (questions 1-15). The scale values range from 1 to 7, higher values correspond to more expressed symptoms and lower QOL.

In all the patients, stool form and consistency was evaluated as well (Bristol scale) before and after treatment.

In all the patients, study of the GIT motor-evacuation function was performed with epektogastroenterorography method (EGG) using the apparatus «Gastroenteromonitor GEM-01». Peripheral computer EGG is an evaluation method of electric activity of various GIT segments. Considering the data obtained in EGG, the following values were evaluated: total level of GIT electrical activity; level of electrical activity for each one of the GIT segments studied. A standard study of GIT electrical activity consists of two stages: the first is a study on an empty stomach during 40 minutes, the second stage is a study after the standard breakfast during 40 minutes. After the treatment course the study was repeated, and the dynamics of basal values of GIT electrical activity was assessed. To evaluate the GIT motor activity, a special coefficient was calculated after the food load, equal to ratio of total postprandial MEA to total fasting GIT MEA (postprandial coefficient – PPC).

Statistical data processing was performed using the programme SPSS 17.0 (SPSS Inc., USA) with the help of Wilcoxon test (for paired observations) and Mann-Whitney test (for non-paired observations).
The dynamics of the GIT complaints in patients of the main and control group on the scales of GSRS questionnaire are presented in Table 1.

No differences between the groups in relation to the dynamics of GSRS questionnaire scale values were received (p> 0.05).

In patients of the main group after treatment with the study medication, a statistically significant reduction was achieved in the severity of complaints on the scales of abdominal pain, diarrhea syndrome, dyspeptic syndrome, and on the scale of the total score. Patients in the control group after therapy with the reference drug the intensity of complaints significantly decreased for all the scales: abdominal pain, reflux syndrome, diarrhea syndrome, constipation and on the total score scale.

Table 1

<table>
<thead>
<tr>
<th>Scale</th>
<th>Time of observation (beginning / end of treatment)</th>
<th>Number of patients, abs. (% of )</th>
<th>Number of patients, abs. (% of )</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Main group, n=15</td>
<td>Main group, n=15</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Me (Q₁; Q₃)</td>
<td>Difference of significance within a group, p</td>
</tr>
<tr>
<td>Abdominal pain</td>
<td>beginning</td>
<td>5 (5; 7,5)</td>
<td>0,003</td>
</tr>
<tr>
<td></td>
<td>end</td>
<td>4 (2; 5)</td>
<td></td>
</tr>
<tr>
<td>Reflux syndrome</td>
<td>beginning</td>
<td>6 (4,5; 6)</td>
<td>0,343</td>
</tr>
<tr>
<td></td>
<td>end</td>
<td>5 (3; 6)</td>
<td></td>
</tr>
</tbody>
</table>
The most clinically significant changes observed in patients of the main group were normalization of stool frequency (decrease up to 7 times a week), normalization of stool form (up to stool type 4 on the Bristol scale) and a decrease in bodily pain. Unlike reference drug (bismuthate tripotassium dicitrate), in the treatment of patients with Enterosgel® beneficial effect was detected not only on subjective clinical IBS manifestations, but also in normalization of stool frequency and form.

<table>
<thead>
<tr>
<th>Scale</th>
<th>Time of observation (beginning / end of treatment)</th>
<th>Number of patients, abs. (% of )</th>
<th>Number of patients, abs. (% of )</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Me $(Q_{1}; Q_{3})$</td>
<td>Me $(Q_{1}; Q_{3})$</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Difference of significance within a group, p</td>
<td>Difference of significance within a group, p</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Diarrhea syndrome</td>
<td>beginning</td>
<td>12 $(9,5; 15)$</td>
<td>0,003</td>
</tr>
<tr>
<td></td>
<td>end</td>
<td>7 $(3,5; 10)$</td>
<td></td>
</tr>
<tr>
<td>Dyspepsy</td>
<td>beginning</td>
<td>15 $(10,5; 17)$</td>
<td>0,003</td>
</tr>
<tr>
<td></td>
<td>end</td>
<td>8 $(6; 10.5)$</td>
<td></td>
</tr>
<tr>
<td>Constipation syndrome</td>
<td>beginning</td>
<td>5 $(4; 6)$</td>
<td>0,591</td>
</tr>
<tr>
<td></td>
<td>end</td>
<td>5 $(3; 7)$</td>
<td></td>
</tr>
<tr>
<td>Total score</td>
<td>beginning</td>
<td>46 $(35; 50.5)$</td>
<td>0,004</td>
</tr>
<tr>
<td></td>
<td>end</td>
<td>29 $(23; 40)$</td>
<td></td>
</tr>
</tbody>
</table>
In the patients of the main group value of the total fasting GIT myoelectric activity before treatment was $Me = 21.4 \text{ mV (Q1 = 16.0; Q3 = 42.5)}$, postprandial - $Me = 32.4 \text{ mV (Q1 = 16.7; Q3 = 59.6)}$. During the treatment with Enterosgel® total fasting value of GIT myoelectric activity was $Me = 23.9 \text{ mV (Q1 = 11.2; Q3 = 46.9)}$, postprandial - $Me = 21.3 \text{ mV (Q1 = 14.4; Q3 = 40.9)}$. Values before and after treatment were not statistically different ($p> 0.05$).

In patients of the control group value of the total GIT myoelectric activity before treatment was fasting $Me = 21.2 \text{ mV (Q1 = 8.5; Q3 = 51.5)}$, postprandial - $Me = 18.2 \text{ mV (Q1 = 6.4; Q3 = 48.8)}$. after a course of the comparator drug total fasting value of GIT myoelectric activity was $Me = 39.4 \text{ mV (Q1 = 8.3; Q3 = 53.4)}$, postprandial - $Me = 11.8 \text{ mV (Q1 = 10.1; Q3 = 38.3)}$ The values before and after treatment were also not statistically different ($p> 0.05$).

PPC in patients of the main and control group before and after treatment is shown in Figure 7.

![Figure 7](image-url)
observation group, vertical axis - the value of PPC, units. The dotted line is the postprandial factor equal to 1.0.

Figure captions: <Beginning of treatment – End of treatment

AUC value, U – Main group – Control group>

in 12 patients (40%) a decrease in PPC was observed (<1.0), reaching in 7 patients (23%) values <0.6. After treatment in 16 patients (53%) a reduction in PPC was observed (<1.0), while PPC increase (> 1.5) was observed in 8 patients (27%), of which in 5 patients (17%) PPC was > 2.0. It is possible that after treatment, on the background of clinical improvement, reduction of diarrhea and normalization of evacuation function a trend towards significant PPC decrease and postprandial power decrease is expected.

Conclusion

The study of efficacy and safety of application of Enterosgel® administered for treatment of patients with diarrhea-predominant irritable bowel syndrome had demonstrated safety and efficiency of the study medication.

Evaluation according to GSRS showed significant reduction on the scales of abdominal pain, diarrhea syndrome, dyspeptic syndrome as well as on the total measure scale. According to EGG, while studying MEA, a tendency towards normalization of the GIT motor-evacuation function is evident.

Summary

Enterosgel® has a positive effect on the clinical performance, reducing abdominal pain severity and normalizing stool frequency in D-IBS patients, thus contributing to the harmonization of quality of life.

Treatment regimens that can be recommended for use in patients with D-IBS: 22.5 g of the paste with the active substance polymethylsiloxane polyhydrate 3 times a day 1 hour after meal orally for 3 consecutive weeks (paste dose should be dissolved in 200 ml of cool boiled water or washed down with at least 200 ml of cool boiled water).

References


