

VOL.35 | ISSUE-1 | CHAPTER-10

PUBLISHER: INTERNATIONAL JOURNAL FOUNDATION (IJF)



# Genital tract candidiasis in patients with Helicobacter Pylori (HP) acid-related disease after providing eradicative therapy.

\* Viacheslav Kravtsov<sup>1,2</sup>, Taame Maria<sup>3,2</sup>, Grukhin Yuriy<sup>4</sup>, Surovtceva Tatiana<sup>4</sup>

<sup>1</sup>Nikiforov Russian Center of Emergency and Radiation Medicine, EMERCOM, Saint-Petersburg, Russian Federation <sup>2</sup>S.M. Kirov Military Medical Academy, Saint-Petersburg, Russian Federation <sup>3</sup> University of Eastern Finland (UEF), Institute of Public Health and Clinical Nutrition, Finland <sup>4</sup> Saint-Petersburg State public health Institution "City Clinical Hospital" № 20, Saint-Petersburg, Russian Federation

#### **ABSTRACT**

Women who diagnosed with Helicobacter pylori -acid related diseases and assigned into two lines of eradicative therapy were tested positive for vaginal candidiasis in comparison to healthy women who didn't go through eradicative therapy against HP.

Thus, the second line of eradicative therapy of HP had an influence on the normal micloflora by decreasing the immune system of genital female tract. Medical professionals should pay attention of the importance in maintaining the immune system of female patients during eradication therapy of HP acid related diseases as well as following the hygiene methods.

Keywords: Helicobacter pylori (HP), eradicative therapy, genital female tract, vaginal candidiasis

## **ORIGINAL RESEARCH ARTICLE**

ISSN: 2456-1045 (Online) (ICV-MDS/Impact Value): 72.30 (GIF) Impact Factor: 5.188

**Publishing Copyright** @ International Journal Foundation **Journal Code:** ARJMD/MDS/V-35.0/I-1/C-10/MAR-2019

**Category:** MEDICAL SCIENCE

Volume: 35.0/Chapter- X/Issue -1 (MARCH-2019) Journal Website: www.journalresearchijf.com

Paper Received: 26.03.2019 Paper Accepted: 06.04.2019 Date of Publication: 15-04-2019

Page: 51-53

# Name of the Corresponding author:

#### Viacheslav Kravtsov\*

Nikiforov Russian Center of Emergency and Radiation Medicine, EMERCOM, Saint-Petersburg, Russian Federation

# **CITATION OF THE ARTICLE**



Kravtsov V; Maria T; Yuriy G; Tatiana S; (2019) Genital tract candidiasis in patients with Helicobacter Pylori (HP) acid-related disease after providing eradicative therapy.; Advance Research Journal of Multidisciplinary Discoveries; 35(10) pp. 51-53

# I. INTRODUCTION

Studies show that Candida colonization may lead to vulvovaginitis after antibiotics treatment in women. However, the pathogenesis on how antibiotics may lead to vulvovaginitis is not understood clearly and larger studies should be conducted in this field. [1][2]

The implication of Helicobacter pylori (HP) in progression of stomach and duodenum diseases is proved and generally accepted by numerous research studies [3],[4],[5]. There is two-line eradicative antihelicobacter therapy applied in order to treat HP-associated acid-related disease. The standard provided treatment is constantly regulated by The Maastricht Treaty. In the event of a failure to eradicate HP after providing the first course, the second course of antihelicobacter therapy is administered. It includes Bismuth medications together with PPI, amoxicillin and tetracycline. The administration lasts 10-14 days. [6]

Clinical practice demonstrates that after providing anti-helicobacter therapy by the standard scheme, the result is not only HP eradication, but oftentimes the provocation of candidiasis in the gastrointestinal tract (GIT) associated with rising gastric acid pH after taking PPI [7]. With the contraction of candidiasis in the gastrointestinal tract (GIT) we can expect the possibility of provocation of candidiasis in the female genital tract. It is reasonable to expect the risk of appearance of candida in the female genital tract after providing the second course of anti-helicobacter therapy.

Such a hypothesis seems to be reasonable, as in our clinical practice (the practice of gynecology and gastroenterology) we had to deal with vaginal candidiasis which had appeared after the second course of antibiotics for HP eradication. For this reason, for patients with Helicobacter pylori (HP) acid-related disease who underwent treatment in the Saint-Petersburg State public health Institution "City Clinical Hospital" №.20 in Saint Petersburg, we organized visits to the gynecologist, where standard cytological smears were obtained for candida tests.

In the present report, the research results included the incidence of vaginal candidiasis in patients with Helicobacter pylori (HP)acid-related disease after the second line of anti-helicobacter therapy.

# II. MATERIALS AND METHODS

The subjects of research were 152 women aged between 32 and 60 with Helicobacter pylori (HP) acid-related disease, who underwent the standard course of eradicative therapy over the course of ten days. The standard course of eradicative therapy includes taking PPI (pariet) and antibiotics (clarithromycin and amoxicillin)during 10 days.

The second treatment course was assigned to 31 patients, where 28 of them suffered chronic gastritis (C.29.6 in ICD-10) and the other 3 patients had peptic ulcer disease (C.27 in ICD-10) and didn't assign to HP eradication after providing the first course of anti-helicobacter therapy. The second treatment course was administered according to the schedule: rabeprazole (40 mg per day), amoxicillin (1000 mg per day), tetracycline (300 mg per day), bismuthate tripotassium dicitrate (240 mg per day) lasting 14 days.

After completion of the first line of treatment, each woman had a volitional gynecological examination. All patients were interviewed and none of them complained about genital discomfort at the time of the interview.

The patients visited the gynecologist between seven to fourteen days after providing the second course of anti-helicobacter therapy in order to have a brush biopsy via a cytological smear.

The referral group consisted of 120 apparently healthy women aged between 27 and 52 who had the preventive examinations at the gynecologist's.

The clinical series didn't include patients with diabetes and women who were prescribed intravaginal hygienical and medical procedures with the use of antiseptic agents.

The cytological smears were stained with hematoxylin-eosin and azure-2-eosin.

The microscopic examinations were held under a microscope with 400 and 1000 times magnifications.

We followed the methodological recommendations to detect Candida albicans [8].

### III. RESULTS AND DISCUSSION

The results were found after seven to fourteen days from providing the second course of anti-helicobacter therapy and found out seven of the 31 women complaining of pruritus, and some complained about heat and pain in the genital region. Upon gynecological examination of the vaginal tyroid fluor, vaginal mucous membrane hypermia and puffiness in the vaginal mucous coating were detected. Elements of Candida fungus, both in the form of budding yeast and in the form of pseudomycelium were found in the smears from the surface of the cervix uteri and lateralis vaginal vault of all the patients. In some cases, as many as 20-30 elements of Candida fungus were detected in the visual field (400x-magnification).

The clinical implication of the result is the diagnosis of vaginal candidiasis (C.37.3 in ICD-10) in more than one fifth of the patients with Helicobacter pylori (HP) acid-related disease after the antibiotic therapy.

As for clinical classification it is most likely that the detected cases of Candida vulvovaginitis date back to acute Candida vulvovaginitis [9].

None of the apparently healthy women who were presented in the referral group and had the preventative examinations at the gynecologist's tested positive for Candida, and none had neither objective nor subjective signs of Candida vulvovaginitis.

Thus, the results of our research point to the fact that providing the second course of anti-helicobacter therapy in patients with Helicobacter pylori (HP) acid-related disease triggers Candida vulvovaginitis in 22 percent of patients.

It is most likely that Candida vulvovaginitis comes after anti-helicobacter therapy because PPI changes gastric acid pH and this triggers dysbacteriosis including candidiasis, as it facilitates the growth of Candida. It is possible that Candida enters in the mucous coating of the genital tract from the GIT as a result of the effect of the antibiotics. In other words, the medications of the antihelicobacter therapy from one hand make a possibility for Candida's growth in GIT (PPI), and from the other hand have an effect on tissue immunity of mucous coat of genital tract by changing its normal microflora with the antibiotics (klacid, amoxicilan).

# IV. CONCLUSION

Providing the second course of anti-helicobacter therapy in the patients with Helicobacter pylori (HP) acid-related disease heightens the risk of the emergence of Candida vulvovaginitis (for 22% of cases).

It is appears that the standards of providing the HP eradication should consider the sex of the patients and include preventive services in order to reduce the risk of

the emergence of Candida vulvovaginitis. Healthcare providers should be aware about possibility of high risk increase of vulvoginatis candidiasis in the female general tract after second line eradication of HP therapy in female genital tract.

# V. FUNDING

None

## VI. REFERENCES

- [1] Marie V. Pirotta, Suzanne M. Garland. Genital Candida Species detected in Samples from Women in Melbourne, Australia, before and after Treatment with Antibiotics. J Clin Microbiol. 2006; 44(9): 3213–3217. doi: 10.1128/JCM.00218-06
- [2] Xu J, Schwartz K, Bartoces M, Monsur J, Severson RK, Sobel JD. Effect of antibiotics on vulvovaginal candidiasis: a MetroNet study. J Am Board Fam Med. 2008; 21(4):261-8. doi: 10.3122/jabfm.2008.04.070169.
- [3] Hajimahmoodi M, Shams-Ardakani M, Saniee P, Siavoshi F, Mehrabani M, Hosseinzadeh H, Foroumadi P, Safavi M, Khanavi M, Akbarzadeh T, Shafiee A, Foroumadi A. In vitro antibacterial activity of some Iranian medicinal plant extracts against Helicobacter pylori. Nat Prod Res. 2011;25:1059–1066. doi: 10.1080/14786419.2010.501763.
- [4] **Buzás GM.** Metabolic consequences of Helicobacter pylori infection and eradication. World J Gastroenterol. 2014;20:5226–5234. doi: 10.3748/wjg.v20.i18.5226.
- [5] Alakkari A, Zullo A, O'Connor HJ. Helicobacter pylori and nonmalignant diseases. Helicobacter. 2011;16 Suppl 1:33–37. doi: 10.1111/j.1523-5378.2011.00878.x.
- [6] Malfertheiner P, Mégraud F, O'Morain C, Hungin AP, Jones R, Axon A, Graham DY, Tytgat G. Current concepts in the management of Helicobacter pylori infection--the Maastricht 2-2000 Consensus Report. European Helicobacter Pylori Study Group (EHPSG). Aliment Pharmacol Ther. 2002;16(2):167-80. https://doi.org/10.1046/j.1365-2036.2002.01169.x
- [7] **Ingrid Kohlstadt.** Advancing Medicine with Food and Nutrients, 2d ed. Boca Raton, Florida: CRC Press; 2012.

- [8] Pappas PG, Kauffman CA, Andes D, Benjamin DK, Jr., Calandra TF, Edwards JE, Jr., Filler SG, Fisher JF, Kullberg BJ, Ostrosky-Zeichner L, Reboli AC, Rex JH, Walsh TJ, Sober JD. Clinical Practice Guidelines for the Management of Candidiasis: 2009 Update by the Infectious Diseases Society of America. Clinical Infectious Diseases. Oxford University Press. 2009; 48(5): 503-535.
- [9] Gonçalves B, Ferreira C, Alves CT, Henriques M, Azeredo J, Silva S. Vulvovaginal candidiasis: Epidemiology, microbiology and risk factors. Crit Rev Microbiol. 2016; 42(6):905-27. doi: 10.3109/1040841X.2015.1091805.

\*\*\*\*\*

OF