The Local Immunity of the Oral Cavity in Women with Recurrent Aphthous Stomatitis Associated with Urogenital Infection

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Abstract

This paper studied the dynamics of local immunity factors of the oral cavity in women with recurrent aphthous stomatitis against the background of urogenital infection. The research proves the maximum efficacy of hydroxyzine hydrochloride, aminodihydrophthalasindione sodium and Eplan used in multiple treatments of recurrent aphthous stomatitis in patients with urogenital infection as evidenced by a marked reduction in the coefficient of local immunity factors balance, which is observed as early as on day 7 of the therapy provided. Moreover, the findings show the growth of immunological parameters of oral fluid (lysozyme and SIgA), the normalization of the coefficient of local immunity factors balance in the treatment of recurrent aphthous stomatitis in women with urogenital infection, the improvement being observed 3 months after the treatment suggesting a favorable local immunity of the oral cavity.

Keywords

Hydroxyzine Hydrochloride, Aminodihydrophthalasindione Sodium, Eplan, Chronic Recurrent Aphthous Stomatitis, Coefficient of Local Immunity Factors Balance, Secretory IgA, Lysozyme, Immunoglobulin A, G

1. Introduction

Recurrent aphthous stomatitis (RAS) is frequent in dental practice [1] [2] [3] [4]
The disease is long lasting and characterized by frequent recurrences, resistance to the treatment provided [1] [4] [6]. Some scientists distinguish a variety of etiological factors, among them there are immune, neurogenic, infection-allergic and other theories of RAS origin [1] [2] [5]. Reports in literature give a variety of different techniques and medications for RAS treatment. However, RAS cure is still an urgent problem [1] [2] [3] [4] [5] [7].

In this regard, our study aimed at optimizing the methods used to treat chronic recurrent aphthous stomatitis (CRAS). To accomplish the purpose, we set a problem of proving the use of some medications in RAS therapy based on the study of the changes; a number of immunological parameters of oral fluid undergo during the treatment.

2. Materials and Methods

The present study was carried out at the Department of Therapeutic Dentistry, Nizhny Novgorod State Medical Academy for 12 years. The study included data from a survey of 1000 women aged 25 - 35 years with chronic recurrent aphthous stomatitis, 500 practically healthy women were included in the control group.

Among them, groups of patients with chronic recurrent aphthous stomatitis (CRAS) were identified who applied to the dental clinic NizhGMA, MLPU “Women’s Consultation No. 5” in Nizhny Novgorod, the NOCC, the Research Institute of Preventive Medicine NizhGMA, OC AIDS with complaints of pain in the SDR and from the urogenital tract reproductive problems, as well as for preventive examination. We performed a clinical examination of 300 women with RAS associated with urogenital infection, aged from 25 to 35 years (mean age 28.2 ± 1.4) followed by a treatment. Exclusion criteria were severe somatic diseases (diabetes, HIV, cancer, blood diseases, etc.), refusal of examination and treatment, age to 25 and after 35 years. Eligibility criteria are women from 25 to 35 years old with gynecological diseases and CRAS.

CRAS patients were randomized in three groups, 100 women in each group, depending on the treatment techniques used: group I (main group) with drug therapy including general treatment combined with therapeutic agents: hydroxyzine hydrochloride, aminodihydrophthalasindione sodium, and local treatment: Eplan applied on problem oral mucosa areas; group II with drug therapy consisting in general treatment combined with therapeutic agents: hydroxyzine hydrochloride, aminodihydrophthalasindione sodium, and local treatment—problem areas of oral mucosa were covered by solcoseryl (dental adhesive paste); and group III with drug therapy including general treatment with antihistamines and multivitamin preparations administered, and local treatment: solcoseryl (dental adhesive paste) applied on problem oral mucosa areas.

Patients of all 3 groups (100 in each) with CRAS for anamnestic data, complaints for primary treatment, age, baseline values of dental and gynecological status were comparable. Patients of the examined groups were examined hor-
monal and immune status, biochemical and general blood test, determination of
blood groups and Rh factor. A comparative evaluation of various diagnostic
techniques was carried out, in particular, the reaction of direct immunofluores-
cence and polymerase chain reaction in real time, enzyme immunoassay, bacte-
riological culture. All patients received informed consent to participate in the
study and receive Galavit, Atarax and Eplan in the treatment of CRAS, as well as
the permission of the Ethics Committee.

The study included several items:
1) Dental examination of women, diagnosis of CRAS, the study of the occur-
rence of dental diseases in women with gynecological pathology evaluation of
dental status in women with gynecological pathology.
2) General clinical methods of examination.
3) Study of the features of the etiology of urogenital infection in prolific and
infertile women with CRAS, combination of dental pathology with infections of
the urogenital tract.
4) Conducting a comparative evaluation of the immune status in prolific and
infertile patients of CRAS.
5) Study of hormonal status in patients with CRAS in combination with uro-
genital infection and infertility.
6) Detection of the nuances of fluctuations in changes in the oral fluid and
blood during dental pathology against the background of gynecological diseases.

Aminodihydropthalasindione sodium was administered in the form of sub-
lingual tablets according to the scheme: daily for 10 days, 4 tablets per 24 h, and
during the following 10 days every other day at the same dose. Thus, the therapy
regimen was 30 days. Hydroxyzine hydrochloride was given t.i.d.: by 12.5 mg in
the morning and afternoon, and 25 mg in the evening within 4 weeks. The oral
cavity was treated by antiseptics followed by Eplan applications for 20 - 30 mi-
utes 3 - 4 times a day until the epithelialization of affected areas. The patients
were followed up for 12 months after the therapy provided. This scheme is ge-
erally accepted and its effectiveness is proved in use, as is shown in the annota-
tion to the drug and publications.

During the laboratory stage we determined the level of secretory immunoglo-
bulin A (SIgA) by radial immunodiffusion (RID) in gel—G. Mancini, A. Carbo-
nara (1965) using guidelines by Е.В. Tchernokhvostova, S.I. Golderman (1975);
serum immunoglobulins (G. A) in oral cavity using radial immunodiffusion
(RID) in gel (G. Mancini, A. Carbonara, 1965); lysozyme in oral cavity by pho-
tonephelometry (V. G. Dorofeichuk, 1968), as well as a coefficient of local im-
munity factors balance (C_b) developed by V.G. Dorofeichuk and Tolkacheva et
al. (1987) [8].

C_b was used for an integral assessment of local oral immunity. C_b formula was
devised considering the functional relationships of lysozyme with IgG and IgA.

\[
C_b = \frac{IgG \cdot 0.4\%}{IgA \cdot 0.6 \cdot la}
\]
where:

- \( IgG, IgA \) is concentration of immunoglobulins;
- \( la \) is lysozyme activity in secretion;
- 40% is conditional norm of lysozyme activity;
- 0.6 is \( IgG/IgA \) relationship, which was true in most healthy people.

For the present study we used oral fluid samples, 3 - 5 ml, taken in the morning, fasting, with no stimulation, and kept in a fridge at \(-20^\circ C\) till the laboratory stage.

The findings were statistically processed using the methods for findings reliability assessment, analysis of variance, and autocorrelation according to standard techniques. The obtained data were processed and analyzed using Microsoft Office (Excel) application programs, and a software statistical package “Stadia” and “Statistica 7.0”.

3. Results

Table 1 shows the findings of immunological parameters of oral fluid in the treatment of 300 patients with RAS and urogenital infection, and 50 apparently healthy subjects (control).

The represented data show a slight IgA decrease in the examined women compared to controls, on day 7 after the treatment there was a significant decrease of IgA level \((p \leq 0.05)\), with the following increase by month 3.

IgG amount was lower in the examined women compared to those of the control group, and decreased sharply by month 3 \((p \leq 0.05)\).

Lysozyme level was significantly reduced in RAS patients compared to the controls \((p \leq 0.05)\). Significant lysozyme decrease was found by day 7 after therapy \((p \leq 0.05)\), while on day 90 its level was significantly higher than the initial one.

In patients with RAS and urogenital infection \( C_b \) was significantly higher compared to the controls \((p \leq 0.05)\) that argued for moderate local immunity of the oral cavity. By day 7 after therapy \( C_b \) slightly decreased, still remaining significantly higher compared to that in controls \((p \leq 0.05)\). \( C_b \) sharply decreased by month 3 after therapy \((p \leq 0.05)\) reaching the values similar to those in the control group \((p \leq 0.05)\), i.e., there was favorable local immunity of the oral cavity.

SIgA level in RAS patients was significantly higher compared to controls, on day 7 and day 90 after therapy SIgA was found to have a significant increase \((p \leq 0.05)\), it approaching the values in the control group.

Thus, a significant increase of immunological parameters of the oral fluid \((IgG, SIgA, C_b)\) was found in the treatment of recurrent aphthous stomatitis in women with urogenital infection, the increase being the most significant on month 3 after therapy.

The analysis of \( C_b \) changes in the oral fluid depending on a treatment method showed the following.
Table 1. Immunological parameters of oral fluid in recurrent aphthous stomatitis and urogenital infection (N = 350).

<table>
<thead>
<tr>
<th>Oral fluid parameters</th>
<th>Time periods before and after therapy (days)</th>
<th>Control group</th>
<th>Before therapy</th>
<th>7</th>
<th>90</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>M ± m</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>IgA, g/l</td>
<td>0.06 ± 0.002</td>
<td>0.058 ± 0.003</td>
<td>0.041 ± 0.002*</td>
<td>0.053 ± 0.002</td>
<td></td>
</tr>
<tr>
<td>IgG, g/l</td>
<td>0.08 ± 0.003</td>
<td>0.065 ± 0.008</td>
<td>0.06 ± 0.01</td>
<td>0.033 ± 0.004*</td>
<td></td>
</tr>
<tr>
<td>Lysozyme, %</td>
<td>45.2 ± 2.3</td>
<td>32.6 ± 1.4*</td>
<td>25.3 ± 1.2*</td>
<td>37.0 ± 2.2*</td>
<td></td>
</tr>
<tr>
<td>Cb</td>
<td>1.54 ± 0.17</td>
<td>4.595 ± 0.543*</td>
<td>3.258 ± 0.423*</td>
<td>1.534 ± 0.184</td>
<td></td>
</tr>
<tr>
<td>SIgA, g/l</td>
<td>0.33 ± 0.09</td>
<td>0.089 ± 0.012*</td>
<td>0.125 ± 0.002</td>
<td>0.242 ± 0.013</td>
<td></td>
</tr>
</tbody>
</table>

*—significant differences in relation to controls (p < 0.05).

Group II demonstrated Cb downtrend on day 7. By month 3 and month 6 Cb in this group was significantly lower than the initial value (p ≤ 0.05), on month 12 it remained stable, its value being significantly lower than that before the treatment (p ≤ 0.05). Group I was found to have Cb decrease on day 7. On month 3 the parameter’s value went down significantly compared to its level before the therapy (p ≤ 0.05); by month 6 there was a slight growth of Cb level, while by month 12 it decreased again remaining lower than that before the treatment (p ≤ 0.05).

In group III Cb level slightly decreased by day 7 and by day 90. On day 180 we observed its growth (p ≤ 0.05), however, 12 months after the treatment Cb level significantly decreased (p ≤ 0.05).

During all the periods of observation Cb values were nearly the same in groups I and II, Cb decrease being accentuated in group I.

Thus, RAS therapy resulted in Cb decrease in all 3 groups, the maximum effect was found in group I as early as on day 7 after the treatment initiation.

4. Discussion

There are a lot of reviews on RAS [4] [6] [9] [10] [11] [12]. A number of various treatment strategies has been evaluated using drug therapy, topical agents and mouth washes [13] [14] [15] [16] [17]. In studies laser therapy is one of the suggested treatments to reduce patients’ discomfort [10] [15] [18]. In another study the application of ozone on RAS lesions was used for treatment [19].

In the literature available to us and to the best of our knowledge, this study is the first of such type, which demonstrates the local immunity of the oral cavity in women with RAS associated with urogenital infection.

In the present examination a significant increase in the immunological indices of the oral fluid (IgG, SIgA, Xb.) was observed in the treatment of patients with aphthous stomatitis and urogenital infection with the scheme proposed by us, using atarax, eplan and galavit, most pronounced at 3 months of treatment. These changes may be associated with the effect of the drug’s use.
So, the mechanism of Galavit action is associated with its ability to regulate the functional and metabolic activity of innate and adaptive immunity (monocytes, macrophages, neutrophils, natural killers, and others). Galavit normalizes the phagocytic activity of monocytes/macrophages, the neutrophils bactericidal activity and the cytotoxic activity of NK cells. At the same time, restoring the reduced activity of immune cells, the drug increases the body’s resistance to infectious diseases of bacterial, viral and fungal etiology, contributes to a more rapid elimination of the pathogen from the body, and reduces the frequency, severity and duration of infections. In addition, Galavit* normalizes antibody production, increases the functional antibodies activity (affinity), indirectly regulates the production of endogenous interferons (IFN-α, IFN-γ) by producing cells. In inflammatory diseases, the drug reversibly (for 6 - 8 hours) inhibits the excessive synthesis of hyperactive macrophages of tumor necrosis factor-α, interleukin-1, interleukin-6 and other proinflammatory cytokines, the level of which determines the degree of inflammatory reactions, their cyclicity, as well as the severity of intoxication of the organism. Galavit* reduces the production of reactive oxygen species by hyperactive macrophages, thereby reducing the level of oxidative stress and protecting tissues and organs from the damaging effects of radicals. Normalization of excessively increased functional activity of phagocytic cells leads to the restoration of their antigen presenting and regulating function, reducing the level of auto-aggression.

Atarax (a derivative of diphenylmethane), has a moderate anxiolytic activity; also has a sedative, antiemetic, antihistamine and m-anticholinergic action. It blocks central m-cholinergic and histamine H1 receptors and inhibits the certain subcortical zones activity.

Eplan has a bactericidal, anti-inflammatory, wound-healing, regenerating and analgesic effect, which is provided by the active ingredients of the drug.

Thus, the combined use of these drugs has an impact on different pathogenesis of recurrent aphthous stomatitis (RAS), leading, inter alia, to an increase in the immunological resistance.

5. Conclusion

The study carried out proves maximum efficiency of hydroxyzine hydrochloride, aminodihydrophthalasindione sodium, and Eplan (group I) used in the combination therapy of RAS in women with urogenital infection as evidenced by significant Cb decrease in group I as early as on day 7 of therapy. Moreover, the findings suggest the increase of immunological parameters of the oral fluid (lysozyme and SIgA), the normalization of Cb values during the treatment of recurrent aphthous stomatitis in women with urogenital infection, the normalization being the most significant on month 3 after therapy, indicating a favourable local immunity of the oral cavity.

Conflicts of Interest

The authors declare no conflicts of interest regarding the publication of this paper.
References


