SECRETORY IMMUNOGLOBULIN A (SIGA) AND DENTAL CARIES OF CHILDREN WITH DIFFERENT DISEASES AND CONDITIONS INFLUENCING ORAL MEDIUM

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SUMMARY

SIGA is the main type immunoglobulin in the mixed saliva and is considered as the main secretion factor of the adaptive immunity in the mouth. The purpose of the study is an evaluation of the SIGA quantities and the connection of those antibodies with dental caries of children with different diseases and conditions influencing the oral medium. The study was performed with 116 children with diabetes, asthma, orthodontic problems (removable braces) and healthy controls. The following methods were used: (1) ELISA with “Salivary secretory IgA KIT of Salimetrics LLC USA; was used for quantitative determination of IgA-S in the saliva ;(2) Evaluation of the liquid oral medium (test “Saliva check-GC”) and dental status (DMF-T) of the children studied.

The results obtained came to show that 2/3 of the healthy children and the children with diabetes have low values of c(<100 µg/ml). In the group of healthy children there are no high values of SIGA; Middle values of SIGA (100 -300 µg/ml) are most frequently met in children with asthmatic disease; One half of the children with orthodontic apparatuses have high values of SIGA (>300 µg/ml); There is no dependence between secretory immunity and dental caries in children.

Key words: secretory IgA-antibodies, oral homeostasis, dental biofilm, diabetes, asthma, orthodontic treatment, dental caries, DMF.

INTRODUCTION

Secretory immunoglobulin A (SIGA) is the dominant immunoglobulin in external secretions and also in mixed saliva. It is often characterized as a component of the immune system “first-line of defense” against pathogenic microorganisms. Restricting microbes’adhesion, these antibodies react to dental biofilm formation and thus interfere into the defense of plaque-oral pathology (caries and periodontal diseases)(1, 2, 6, 8).

SIGA is an essential biomarker for the local defense of the mouth. Their secretion depends on the general health of the organism and its immunity. On the other hand, the rich oral antigen potential is also a stimulus for SIGA antibodies(3, 4, 5, 9).

The role of SIGA for the development of dental caries is determined by the ability of such antibodies to impede plaque microorganisms’ colonization on the enamel surface by selective connection. Thus they oppose all other mechanisms of microbe adhesion and interfere during formation of plaque biofilm(9). An interesting question is why residual microflora persists in oral cavity despite the presence of a high SIGA level in the saliva? It is considered that the stability of the flora is a result of its decreased activity to provoke immune reactions. It is assumed that the greater part of the oral microorganisms are not immunogenic for the organism since they live in symbiosis with the body following the long period of evolutionary adaptation(10,13).

It is the aim of the investigation to evaluate the quantities of SIGA in children with different diseases and conditions influencing oral medium as well as the connection of these antibodies with dental caries.

For the purpose we set the following tasks:
1. Study of the average values of SIGA in the saliva of children with different diseases and conditions influencing oral medium;
2. Study of the dependence between dental caries and SIGA in the children under investigation.

MATERIAL AND METHODS

The study was performed on 116 children: (1)30 diabetics children (av. age 12,2± 3,53 years old);(2) 25 children suffering asthma (av. age 8,84 ±3,02 years old);(3)4 healthy children (av. age 10,47± 2,75 years old); (27) undergoing orthodontic treatment (av. age 11,04 ± 1,26 years old).

The selected groups of children are suitable for this study because in the three of them we can expect a change in the oral immunity. In diabetic children this change is in relation to the main disease; in asthmatic children- allergic terrain and local influence of the inhalatory corticosteroids; the local irritating effect of the plastic in the orthodontic
appliances may modify the oral medium(12).

1. Clinical examination and evaluation of dental caries. All children were subjected to:
   • evaluation of the oral medium by tools of caries risk assessment(11);
   • registration of the dental caries – DMF(T+t);
   • saliva sample for SIgA study was taken.

2. Method of saliva collection. Saliva was collected in the morning, on empty stomach, after stimulation for 2 min the salivation by means of chewing standard neutral chewing gum (from the test “Saliva Chek” GC. Stimulated saliva was collected in a plastic container of 5 ml and then by a dropper a certain amount was taken and frozen in a refrigerator (-10°C).

3. ELISA method for study of SIgA in the saliva(14). For the quantitative determination of the SIgA in the saliva we employed the ELISA method with „Salivary secretory IgA KIT” of Salimetrics LLC - USA. The method is indirect, a constant quantity of goat anti-human SIgA conugated with horseradish peroxidase(HRP) being used. The antibody-conjugate is added in test-tubes with specific dilututions of standards or samples from the saliva studied. From each sample sprinkling is made in separate small holes with human SIgA grouped on the same plaque. The quantities of free antibodies are in reverse proportion to the quantity of SIgA. An enzyme reaction is evoked, which causes blue colouring that changes into yellow after the end of the reaction. By means of standard registration as per ELISA the optical thickness is measured with a length of the wave 450nm and 650 nm, the average value being taken into account. After certain calculations (described by the firm producer of the test) a standard curve is built from the values of the six controls and the standards. By means of the equation composed on the basis of the curve the concentrations of SIgA are calculated in µg/ml.

Fig. 1. Standard curve obtained from the study

The statistical processing was done with the help of the computer program SPSS 16.

RESULTS

1. Comparative study of SIgA in the saliva of children studied. The average values of SIgA in the saliva of the healthy children (third group) are 121.34 ±15.027 µg/ml. For children with diabetes (first group) the average quantities of SIgA in the saliva are 133.95±160.504 µg/ml, close to those of the control group. Due to the initial phase of diabetes development, the system endocrine disease seems not to influence the local secretory oral immunity. Children with asthma exhibit statistically verifiable higher quantities of IgA-S (196.456±145.308 µg/ml), compared to the healthy children (P<0,05). The average values of SIgA in children with orthodontic apparatuses are even higher (208.874±125.953 µg/ml). The results can be explained by the stimulating action of the plastic apparatus, alien to the oral environment or by the conditions created in the mouth that lead to more difficult maintenance of the oral hygiene, thus enforcing the bacterial antigenic stimulus.

We distributed the examined children in the following groups:

(1) low SIgA(up to 100 µg/ml);
(2) medium SIgA(100 - 300 µg/ml);
(3) high SIgA (> 300 µg/ml) values of SIgA.

The distribution of the children examined into the groups thus formed is shown in the following table.

Fig. 2. Distribution of the children examined according to the quantity of SIgA in the saliva

The results obtained show that in the group of healthy children are absent any high values of SIgA (> 300 µg/ml). Two thirds(64,7%) of these children are with low and 1/3(35,3%) with medium values of SIgA (up to 300 µg/ml). In the two group with proven above the average values of SIgA (children with asthma and with orthodontic apparatuses) a very interesting distribution of data is observed – half of the children with asthma(56%) belong to the category with medium values, accordingly, half of the children with orthodontic apparatuses(48,1%) belong
to the category with high values. The distribution of the low values in the children with diabetes is similar to that of healthy children, but unlike the healthy children in 1/5 of the children with diabetes (20%) there also very high values (> 300 µg/ml).

**2. Study of the dependence between dental caries and SIgA.** In order to investigate the dependence of SIgA and the dental caries in clinical conditions, we followed DMF (T+t) by grouping the children studies in two ways:

- In the mentioned above three groups according to SIgA:
  1. low SIgA (up to 100 µg/ml);
  2. medium SIgA (100 - 300 µg/ml);
  3. high SIgA (> 300 µg/ml) values of SIgA.
- In two groups according to their cariosity:
  - DMF(T+t) up to 6 teeth;
  - DMF (T+t) above 6 teeth.

**2.1. DMF(T+t) in children grouped according to the quantities of SIgA.** To eliminate the influence of the age factor, in addition to the curiosity, we compared the average age of the children in the groups formed. The results are shown in the next table.

<table>
<thead>
<tr>
<th>Group</th>
<th>n</th>
<th>Age - years</th>
<th>X ± SD</th>
<th>∆</th>
</tr>
</thead>
<tbody>
<tr>
<td>1st group</td>
<td>52</td>
<td>10,84</td>
<td>3,07</td>
<td>0,421</td>
</tr>
<tr>
<td>2nd group</td>
<td>39</td>
<td>10,56</td>
<td>3,34</td>
<td>0,535</td>
</tr>
<tr>
<td>3rd group</td>
<td>25</td>
<td>10,65</td>
<td>2,49</td>
<td>0,499</td>
</tr>
<tr>
<td>T</td>
<td>P</td>
<td>n</td>
<td>P &gt; 0,05</td>
<td></td>
</tr>
</tbody>
</table>

It should be noted that though the average age of the children does not differ in the three groups (P > 0,05).

Dental caries (DMF) is approximately equal for children with very low and very high values of SIgA. For the group with children with average values, the DMF-index is lower, and the differences are not statistically verifiable (P > 0,05).

**Fig. 3. DMF(T+t) of children divided to the quantities of SIgA**

These results show a lack of dependence between the secretory immunity and dental caries. This conclusion is confirmed also when comparing SIgA of the children grouped according to DMF(T+t).

**2.2. Comparing SIgA of the children grouped according to DMF(T+t).** The results are shown in Table 3.

<table>
<thead>
<tr>
<th>Cariosity DMF(T+t)</th>
<th>Number of children (n)</th>
<th>SIgA-S µg/ml</th>
<th>X</th>
<th>± SD</th>
<th>∆</th>
</tr>
</thead>
<tbody>
<tr>
<td>up to 6</td>
<td>82</td>
<td>150,02</td>
<td>132,33</td>
<td>14,61</td>
<td></td>
</tr>
<tr>
<td>above 6</td>
<td>33</td>
<td>160,94</td>
<td>143,36</td>
<td>24,95</td>
<td></td>
</tr>
<tr>
<td>T</td>
<td>P</td>
<td>T=0,391</td>
<td>P=0,697</td>
<td>(P &gt; 0,05)</td>
<td></td>
</tr>
</tbody>
</table>

There are no proven differences between SIgA of children with lower and those with higher cariosity.

**2.3. Correlation dependence between DMF(T+t) and SIgA.** We searched for the presence of correlation dependence between DMF(T+t) and SIgA. The coefficient of Pearson showed very low values which confirm the lack of dependence between caries and SIgA.

<table>
<thead>
<tr>
<th>Pearson correlation</th>
<th>SIgA</th>
</tr>
</thead>
<tbody>
<tr>
<td>DMF(T+t)</td>
<td>0,032</td>
</tr>
</tbody>
</table>
DISCUSSION

For the first time in our country a study has been performed on the local secretory immunity as a part of the complex evaluation of the oral risk environment in children with different diseases. The differences in the average values show that the secretory immunity is stimulated more than the local immunogenic factors and the immunopathological processes (asthma) and is not influenced by systemic diseases like diabetes. Plastic orthodontic apparatuses as local antigenic irritators provide greatest stimulus on oral secretory immunity.

No dependence was found between the secretory immunity and dental caries in children. The role of SIgA in the development of dental caries is determined by the ability of these antibodies to selectively connect with plaque microorganisms and thus impede their colonization on the enamel surface. In this way they counteract to all other mechanisms of microbe adhesion and impede the formation of plaque biofilm. Secretory IgA is connected mostly with S.mutans and antigens of its enzymes and metabolic products.

In real oral environment however this at first glance considerable preventive action of SIgA with respect to dental caries is not so efficient due to the constant washing off action of saliva and the impossibility to maintain sufficient concentration of SIgA on the enamel surface.

CONCLUSIONS:

1. 2/3 of the healthy children and the children with diabetes have low values of c(<100 µg/ml). In the group of healthy children there are no high values of SIgA;
2. Middle values of SIgA (100 -300 µg/ml) are most frequently met in children with asthmatic disease;
3. One half of the children with orthodontic apparatuses have high values of SIgA (>300 µg/ml);
4. There is no dependence between secretory immunity and dental caries in children.

REFERENCES: