The role of Hsp60, CD-8 and IFN-γ in immunopathobiogenesis of periapical granuloma in dental caries

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ABSTRACT

Background: The incidence of dental caries with periapical granulomas in Indonesia is quite high. However, the mechanism of the formation of periapical granulomas in dental caries caused by bacterial infection in immunopathobiogenesis cannot be explained completely. Thus, this explanation is necessary in order to be used as a basis for diagnostic, preventive and therapeutic measures.

Purpose: This research was aimed to determine the role of Hsp60, CD-8 and IFN-γ in immunopathobiogenesis of periapical granuloma in dental caries.

Methods: This research was an analytic observational study with cross sectional approach. Samples of this research were 36 teeth of patients with dental caries, consisting of 18 caries teeth with periapical granulomas and 18 caries teeth without periapical granulomas. The variables observed in this research were Hsp60, CD-8 and IFN-γ. Measurements were conducted by using immunohistochemical methods on periapical tissue.

Results: The mean of Hsp60, CD-8 and IFN-γ in granuloma group was significantly higher than those in non granuloma group (p<0.05). The positive role of IFN-γ on the incidence of granulomas appeared to be more prominent.

Conclusion: The study suggested that in immunopathobiogenesis of periapical granuloma in dental caries, Hsp60, CD-8 and IFN-γ played important roles, but the role of IFN-γ was found to be more prominent.

Key words: Dental caries, immunopathobiogenesis, periapical granuloma, Hsp60, CD-8, IFN-γ

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INTRODUCTION

Dental caries with chronic periapical can be considered as a multifactorial infectious disease caused by irritants in necrotic pulp moving into periapical tissues. Cases involving chronic periapical disease was necrotic pulp can be treated with root canal treatment. The continuous exposure of those irritants to the periapical tissues will stimulate host defense in the form of periapical granulomas. The healing process of periapical granulomas is actually associated with the body’s immune response. Thus, it is possible for recurrence despite the application of root canal treatment, or it may even develop into a radicular cyst, which is more difficult to treat. Therefore, if the process to being granuloma formation can be prevented, many difficulties in healing process of periapical granulomas can be overcome.

Unfortunately, the mechanism of immunopathobiogenesis of periapical granulomas caused by chronic periapical in dental caries still cannot be explained. Many researches only focus on the causes of dental pulp necrosis, such as tumor necrosis factor α (TNF-α) and interleukin-6 (IL-6) in periapical lesions. The mechanism of periapical granuloma is necessary to be studied since the number of cases of dental pulp necrosis is more commonly found in patients with dental caries.

Histological changes in the periapical tissues caused by bacterial invasion, moreover, will be indicated by the presence of granulation tissue containing with macrophages, lymphocytes, plasma cells, neutrophils, and fibrovascular elements in varying numbers. At the same time, there will also be damage to the periapical tissues and bone resorption. Periapical granuloma is actually composed of granulation tissue surrounded by a cell wall of fibrous connective tissue.

In histopathological examination of the chronic lesions, it can be known the existence of lymphocytes, plasma cells, neutrophils, histiocytes, eusinophils, and epithelial cell rests of Mallesse. Lymphocytes are the predominant cell type (50%), which number is closely related to the total number of the following cells, namely CDs -4 T cells and CD-8 T cells. Therefore, in chronic lesions the number of CD-8 T cells is increased. All of their structures are surrounded by a capsule of fibrous connective tissues consisting of CD-8 T lymphocytes. Therefore, it can be said that pathological periapical granuloma is frequently caused by dental caries.

Dental caries is considered as a chronic pathological process caused by microorganisms found in dental caries tissue as a potential immunogen leading to changes in the pulp tissue related to immune response. In other words, bacteria can be considered as an important factor in the development and growth of dental caries.

The presence of anaerobic bacteria considered as pathogenic can trigger macrophages to form periapical granulomas. In the process of periapical granuloma formation, various components have important roles, namely Hsp60 as chaperone playing a role in the fraction of proteins involved in APC. It is because Hsp60 synthesized in the exosome is used to assist the synthesis and maturation of proteins to become functional ones. Consequently, the processing of epitope can be run, and Hsp60 will be recognized by CTL/CD-8, which will then secrete IFN-γ.

Furthermore, IFN-γ released by both Th-1 and CTL/CD-8 will induce macrophage activities. Thus, those macrophages will migrate around those containing intracellular bacteria, and then create granuloma. It is because the presence of pathogenic bacteria will trigger some histiocytes to develop into macrophages and APC leading to the formation of granulomas, while some others will develop into phagocytic causing no granulomas. The study was aimed to determine the role of Hsp60, CD-8 and IFN-γ in immunopaobiogenesis of periapical granuloma in dental caries.

MATERIALS AND METHODS

This research was an analytic observational research with cross sectional approach. The samples of this research were selected from the population based on inclusion criteria (purposive sampling). The samples were divided into two groups, consist of group with granulomas and group with no-granulomas. The subjects were 36 patients who were indicated for tooth extraction and to accept informed consent about the purposes and objectives of this research.

Those patients classified into group with granulomas must have a diagnosis of dental pulp necrosis with periapical granulomas. Diagnose was determined through clinical and radiographic examinations. Clinically the depth of cavity reached the pulp and radiographic figure showed that there was radiolucent area with distinct border with 0.5 mm-2 of diameter. In the other hand, those patients who were classified into non-granulomas group must have a diagnosis of dental pulp necrosis without periapical granulomas indicated clinically by the depth of pulp cavity and the radiographic figure showed no periapical radiolucency.

In the group with granulomas, periapical tissues around tooth extraction were taken by using a pair of tweezers. Meanwhile, in the group with non-granulomas, periapical tissues taken were tissues attached to the extracted tooth. Those periapical tissues were partially fixed in 10% buffered formalin, then histopathologically examined by using Hematoxylin Eusin (HE) staining, and observed by using a microscope with a magnification of 400 times.

Afterwards, the number of cells producing Hsp60, CD-8 and IFN-γ contained in those periapical tissues was calculated by using immunohistochemical method. In the immunohistochemical method, staining process was conducted by using monoclonal antibodies against Hsp60, CD-8 and IFN-γ. Then those were observed by using a light microscope magnification of 400 times, and also carried out by means of counters in five visual fields. Next, the results...
of the examination showed the number of cells producing Hsp60, CD-8 and IFN-γ that would give a positive reaction to monoclonal antibodies, divided by the total (number of cells) and multiplied by 100%.\textsuperscript{21}

Finally, multivariation statistical analysis was used to measure the differences of the average number of cells producing Hsp60, CD-8 and IFN-γ among the groups. Besides that, discriminant analysis was also conducted to obtain the mechanism of periapical granulomas and the patterns of variable contribution between in the group with granulomas and in the group with non-granulomas, so the dominant variable used to distinguish them could be determined.

**RESULTS**

The results of immunohistochemical examination using monoclonal antibody, anti-Hsp60, CD-8 lymphocytes and IFN-γ induced into granuloma and non-granuloma tissues can be seen in Figure 1. The calculation results of cells producing Hsp60, lymphocytes CD-8, and IFN-γ in the group with granulomas and in the group with non-granulomas were obtained by using immunohistochemical method. The mean and standard deviation of those three variables on each group can be seen in Table 1.

Based on the results, it can be said that the mean of Hsp60, CD-8 lymphocytes and IFN-γ in the group with
granulomas was larger than that in the group with non-granulomas (Table 1). The results of multivariate analysis using Wilks’ Lambda test shows p<.001. This means that there was a significant difference between the mean of Hsp60, CD-8 lymphocytes and IFN-γ in the group with granulomas and that in the group with non-granulomas.

Moreover, based on the results of a statistical test with stepwise method, it is known that IFN-γ was the only one dominant variable in distinguishing the group with granulomas and the group with non-granulomas. The minimum of F value of IFN-γ was about 0.1, while the F values of Hsp60 and CD-8 lymphocytes were 0.998 and 0.087. Since the F values of Hsp60 and CD-8 lymphocytes were higher than the maximum limit of 0.05 for inclusion in the analysis, Hsp60 and CD-8 lymphocytes did not become discriminative with the grouping of granulomas/non-granulomas. However, the discriminant analysis resulted in Fisher’s linear discriminant function as shown in Table 2.

Cross tabulation result from discriminant pattern can be seen on the Table 3. Based on Fisher’s linear discriminant function above, it was found that there were two groups of samples with granulomas incorrectly classified as the group with non-granulomas, while for the sample of non granulomas there was not any incorrect classification. The means of difference among those two groups was 94.4%, namely 88.9% for the group with granulomas and 100% for the group with non-granulomas. The results of cross-tabulations of the discriminant pattern can be seen in Table 3.

Furthermore, the positive role of IFN-γ on the incidence of granulomas appeared more clearly as seen in Figure 2, where the mean and standard deviations of both variables

distinguishing the group with granulomas and the group with non-granulomas granuloma were calculated based on Fisher’s function.

**DISCUSSION**

Based on Table 1, it is known that the average number of cells producing Hsp60 in the group suffering from granulomas (5.68) was significantly different that in the group suffering from non-granulomas (0.95). Similarly, a research conducted by Suzuki et al. also showed that there was an increase in the local expression of Hsp60 in periapical inflammatory lesions. This increasing of Hsp60 expressions in patients with periapical granulomas is caused by immune response to the inflammatory later leading to the expression of MHC class I. The producing of the expression of MHC class I then will trigger a series of processes to produce several cytokines, including IFN-γ, TNF-α and TNF-β. On the other hand, the presence of MHC class I expression on the cell surface will also be recognized by CD-8 lymphocytes, thereby increasing the expression of CD-8. However, in the cases of periapical granulomas, the increasing of CD-8 expression does not occur since the release of IFN-γ activates monocytes (macrophages) which will surround intracellular bacteria. It happened because there is stimulation of IFN-γ, especially MAF, which then will lead to the formation of granuloma.

### Table 1. The mean and standard deviation of immunity components, namely Hsp60, CD-8 lymphocytes and IFN-γ

<table>
<thead>
<tr>
<th>Immunity Components (%/1)</th>
<th>Granuloma (N = 18)</th>
<th>Non-granuloma (N = 18)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean</td>
<td>Standard Deviation</td>
</tr>
<tr>
<td>Hsp60</td>
<td>5.68</td>
<td>7.47</td>
</tr>
<tr>
<td>CD-8</td>
<td>3.71</td>
<td>4.56</td>
</tr>
<tr>
<td>IFN-γ</td>
<td>5.61</td>
<td>2.73</td>
</tr>
</tbody>
</table>

Wilks’ Lambda p = 0.000

### Table 2. Fisher’s linear discriminant function with variable IFN-γ

<table>
<thead>
<tr>
<th>Immunity Components</th>
<th>Group</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Granulomas</td>
</tr>
<tr>
<td>IFN – γ</td>
<td>146.604</td>
</tr>
<tr>
<td>(constant)</td>
<td>-4.808</td>
</tr>
</tbody>
</table>

**Figure 2.** Discrimination pattern of IFN-γ.
The average number of cells producing CD-8 in the group with granulomas (3.71) was significantly different that in the group with non-granulomas (0.19). The difference is caused by high immunity possessed by those patients with granulomas. As a result, bacteria entering to the body will be phagocytized by monocytes, so they will get phagolysosome and be neutralized by enzyme system (lysozyme). Bacteria that are resistant to lysozyme will be destroyed by monocytes with reactive oxygen species (ROS) system, especially super oxidant radicals, so the bacteria will get oxidative stress. To protect proteins contained in the body of bacteria from damage, the bacteria then will release stress proteins (Hsp).

When the attacks of intracellular bacteria occur continuously, the host cell will release excessive ROS. As a result, anti-ROS produced by the host will become unbalanced. In other words, ROS is greater than anti-ROS (scavenger enzyme), so the host cells will get oxidative stress. The host cells that distress then will lead to the damage of proteins in the host cells. Thus, to prevent the damage, the host cell releases stress protein, Hsp60. Hsp60 is synthesized in the exosome and serves as chaperones, which will help folding and can be used to assist the synthesis and maturation of proteins to become functional ones. Consequently, the processing of epitope can be run, and Hsp60 will be recognized by CTL / CD-8, which then will secrete IFN-γ.1-8

CD-8 lymphocyte can be considered as a specific phenotype of mononuclear cells used to differentiate asymptomatic lesions and symptomatic lesions in patients with periapical lesions. According to Colic et al.,12 the percentage of CD-8 cells in symptomatic lesions is higher than that in asymptomatic lesions. In other words, the research proves the existence of CD-8 lymphocytes in asymptomatic lesions although it does not distinguish further the percentage of CD-8 cells in granuloma tissue and that in non-granuloma tissue in patients with asymptomatic periapical lesions.

The average number of cells producing IFN-γ in the group with granulomas was 5.61 and that in the group with non-granulomas was 0.42 (Table 1). It means that the increasing of immunity indicates the increasing of the ability of cells to eliminate bacteria. Besides that, it also means that intracellular signals will be increased, so cytokines playing a role in delivering extracellular signal, namely IL-12, will be improved. The cytokines also play a role in inducing T lymphocytes to secrete IFN-γ. The secretion of IFN-γ then will stimulate the activation of macrophages playing a role to prevent the diffusion of bacteria, thus, the cells containing intracellular bacteria will be surrounded by macrophages, and then granuloma will be formed.8,11 Similarly, a research conducted by Breloer et al.13 also showed that the in vitro release of IFN-γ can dramatically be increased as the increasing of Hsp60 moving to T cells and macrophages, but either the production of IL-2 or the proliferation of T cells will not be increased. It means that the induction of IFN-γ is dependent entirely on the ability of macrophages to produce IL-12.

In patients suffering from dental caries with periapical granuloma, immunity components, namely Hsp60, CD-8 lymphocytes and IFN-γ, have an important contribution as indicated with the increasing of cells producing CD-8, IFN-γ and Hsp60.

Immunopathobiogenesis of periapical granulomas can show a relationship between those three components to the pattern of cells producing high CD-8, and IFN-γ, and Hsp60. It means that those three immunity components have positive contribution in the immunopathobiogenesis of periapical granuloma. For instance, CD-8 lymphocyte expression will directly trigger the apoptosis of CD-8 lymphocytes, and then serve as a proapoptotic to inhibit CD-4 (Th2 cells). These barriers will affect the balance resulting in the increasing of Th1.14 On the other hand, Hsp60 will also stimulate Th1 through IL-12, so cytokines (IFN-γ) will be secreted. Cells producing IFN-γ will trigger the proliferation of CD-8, so CD-8 lymphocytes will be increased and IFN-γ produced will also be high.

Hsp60 is a mediator of immune stimulation with different mechanisms, and can be affected by LPS. Hsp60 and LPS, according to Osterloh et al.,15 can increases cytokine (IFN-γ) produced by Th1 cells. The presence of Hsp60 will induce the production of IFN-α, so that Hsp60 functions as a classic signal that induces tissue damage to the innate immune system. If there is a bacterial infection, then Hsp60 binds to LPS and facilitates the detection of microbes by recognizing Pathogen-Associated Molecular Pattern (PAMP) early and increasing TLR signaling. The result of a research conducted by Suzuki et al.,10 showed that the epithelial residue of Malassez had a negative reaction to Hsp60 indicating no protective function of Hsp60 against inflammation. This finding explains another function of Hsp60 in immunopathobiogenesis of periapical granulomas since Hsp60 is previously more often associated with epithelial proliferation and migration process into periapical lesions.

Similarly, the results of an in vitro research conducted by Breloer et al.13 showed that Hsp60 led to a pro-inflammatory response of cells in the adaptive immune system. It is also known that Hsp60 triggers specific IFN-γ secretion with a high number of T cells in Peritoneal Exudates Cells (PEC) as the cells displaying the antigen (APC). In short, these results indicate that Hsp60 as endogenous molecules can cause inflammatory response. These also indicate that the activation of cells of the innate and adaptive immune system by Hsp60 is strongly influenced by the type of APC.

The formation of granuloma can be considered as a chronic inflammatory process due to the failure of the acute inflammatory process, causing activation of persistent antigens and accumulation of macrophages continuously. It is because IFN-γ released by T cells causes the transformation of macrophages into epitheloid cells and data cells.11 Similarly, although the ability of macrophages to neutralize the bacteria that lead to a very high majority of bacteria will be destroyed by lysozyme system, then...
causing no inflammatory reaction excessive, there is still humoral immunologic reaction. In this immunological reaction, macrophages will secrete IL-12, which will induce lymphocytes (Th1), so IFN-γ will be produced. Then the expression of IFN-γ will trigger proliferation of CD-8 lymphocytes. Consequently, it will secrete IFN-γ and activate macrophages. Therefore, in patients suffering from dental caries with periapical granulomas, there will be no destruction of macrophages containing intracellular bacteria, but the activity of macrophages to surround those macrophages containing intracellular bacteria will be increased.8

The study suggested that in the immunopathobiogenesis of periapical granuloma in dental caries, Hsp60, CD-8 and IFN-γ played important rules, but the role of IFN-γ was found to be more prominent.

REFERENCES

3. Radics T. The role of inflammatory and immunological processes in the development of chronic apical periodontitis. Universitas of Debrecen Medical and Health Science Center Faculty of Dentistry. 2004.