TOXICITY ANALYSIS OF CRAB SHELL CHITOSAN ARGINYLGLYCYLASPARTIC ACID SCAFFOLD MEMBRANE AND ITS EFFECT ON HUMAN DENTAL PULP CELL VIABILITY

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ABSTRACT

Objective: Crab shell chitosan is a biomaterial used for scaffolding. In Indonesia, Badan Tenaga Nuklir Nasional has made a crab shell chitosan arginylglycylaspartic acid (RGD) scaffold membrane. The purpose of adding RGD was to enhance cell attachment to the scaffold. The objective of this research is to analyze the toxicity of crab shell chitosan RGD scaffold membrane on human dental pulp cells and its effect on their viability.

Methods: Human dental pulp cells were cultured for 5 days in Minimum Essential Medium Alpha (α-MEM) complete containing amphotericin B, penicillin, streptomycin, and fetal bovine serum. Then, the treatment group was exposed to crab shell chitosan RGD scaffold membrane and crab shell chitosan scaffold membrane incubated for 24 h. The toxicity of the crab shell chitosan RGD scaffold membrane was analyzed with a 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide assay.

Result: The result of this research is that crab shell chitosan RGD scaffold membrane did not decrease the percentage of viability of human dental pulp cells.

Conclusion: It is concluded that crab shell chitosan RGD scaffold membrane does not have toxic effects on human dental pulp cells.

Keywords: Crab shells, Chitosan, Scaffold, Arginylglycylaspartic acid, Toxicity, Human dental pulp cells.

INTRODUCTION

Bone defects can happen due to trauma, periodontal disease, tumors, cysts, and congenital abnormalities, leading to conditions that can change a patient’s quality of life [1]. There are a variety of treatments available to cure these conditions.

One such treatment is the use of a bone graft. Bone grafts can be obtained from patients themselves (autograft) or from other animal species (xenograft). Bone grafts have been used for many years and are considered a standard treatment for bone defects; however, autografts show some disadvantages such as insufficient supply and donor site pain [1]. These disadvantages lead to the development of alternative technologies such as tissue engineering. Tissue engineering is a technique that is used for repairing or generating new tissue and consists of three main components: Growth factor, stem cells, and scaffold (Amir et al.). Growth factor is a transcription factor that originates from gene activation during the tissue engineering process [2]. Stem cells are unspecialized cells that have a high proliferation rate and can generate complex tissues and organs [3]. Stem cells can originate from dental pulp [4]. A scaffold is a temporary, extracellular matrix that facilitates cell attachment, cell proliferation, and cell differentiation [5].

One material that can be used as a scaffold is chitosan, the biopolymer derivative of chitin that can be obtained from the exoskeleton of crustaceans such as crab and shrimp. It is biocompatible, biodegradable, and has osteoinductive abilities [6]. However, chitosan lacks bioactive signals that are important for cell attachment, proliferation, and differentiation. For this reason, a chitosan scaffold is often modified with some bioactive molecules such as arginine-glycine-aspartic acid (RGD).

RGD is a cell adhesion peptide that can be found in extracellular matrix proteins such as laminin and collagen. The objective of adding RGD to a chitosan scaffold is to improve cell attachment to the scaffold. Nowadays, Badan Tenaga Nuklir Nasional (BATAN) develops crab shell chitosan RGD scaffold in the form of a membrane combined with RGD. The purpose of forming this scaffold into a membrane is to obtain a scaffold that has high stability when used to repair bone defects, especially large bone defects. However, before this scaffold can be used in tissue engineering, studies must ensure that this scaffold is biocompatible, meaning that this scaffold is non-toxic. The objective of this study is to analyze the toxicity of crab shell chitosan RGD scaffold membrane on human dental pulp cell viability.

METHODS

In this research, crab shell chitosan RGD scaffold produced by BATAN, Indonesia, was used. The scaffold had the following specifications: In membrane form (2.5 cm × 1 cm); in white color; degree of deacetylation was 94.5%; viscosity 34.40 cps; and 4 mg RGD/50 ml chitosan 2%. The scaffold was produced using a freeze-drying method (Fig. 1). The toxicity of crab shell chitosan RGD scaffold membrane on the viability of human dental pulp cells was analyzed with a 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) test. Two kinds of chitosan materials were used: Crab shell chitosan RGD scaffold membrane and crab shell chitosan scaffold membrane without RGD. Crab shell chitosan RGD scaffold membranes with 1 mg and 2 mg concentration, were described in human dental pulp cell cultures and incubated for 24 h. Human dental pulp cell viability was expressed as a percent of the control. Test difference between groups was then analyzed using the one-way analysis of variance test (ANOVA test).
Cell culture
First premolar or impacted third molars were removed from patients at the Dental Hospital, Faculty of Dentistry, Universitas Indonesia. The human teeth were soaked in Minimum Essential Medium Alpha (α-MEM) complete. Every 50 ml of α-MEM complete contains L-glutamine, ribonucleosides and deoxyribonucleosides with 10% fetal bovine serum, penicillin 10.000 UI/ml, streptomycin 100 mg, and amphotericin B 25 μg/ml. Furthermore, the teeth were washed with phosphate buffer saline containing antibiotics. Dental pulp tissue was obtained using an extirpation needle and collected on a petri dish containing α-MEM without serum. Human dental pulp cells (10⁶) were cultured in a 25 cm² flask with α-MEM complete for up to 5 days. The medium was replaced every 2 days.

MTT
Toxicity tests were analyzed using MTT. The MTT assay is used as an indirect measure of cell toxicity based on the number of living cells left after being exposed to a test material. Human dental pulp cells, with a density of 2 ×10⁴, were placed on a 96-well plate and cultured for 5 days. With the aim of advancing cell growth, the medium was replaced every 2 days. Next, the treatment group was presented with the crab shell chitosan RGD scaffold membrane with and without RGD (1 mg and 2 mg). A 96-well plate was incubated in a CO₂ incubator (37°C, 5% CO₂) for 24 h. After incubation, an MTT test was conducted by adding 1.5 ml solution of MTT (5 mg/ml) to all test groups.

Further, incubation in a CO₂ incubator (37°C, 5% CO₂) was conducted for 3 h. Next, 100 µl of acidified isopropanol was added to each well, followed by incubation at room temperature for 1 h on an orbital shaker to dissolve the formazan crystals. Samples absorbance values (OD) were read using an enzyme-linked immunosorbent assay reader with a wavelength of 490 nm. Data were corrected with a culture medium without cells (blank value). The experiment was repeated twice on each test group that was at least Duplo. Cell viability was calculated using the following formula:

\[ \text{Cell viability} = \frac{\text{mean absorbance value treatment group}}{\text{average value of absorbance of control group}} \times 100\% \]

Statistical Analysis
The data were presented as mean values and standard deviations. First, the data were tested using a Shapiro-Wilk normality test. Significant differences in the data were analyzed using a one-way ANOVA test (p<0.05).

RESULT
Effect of rotary cell culture system on human dental pulp cells
Human dental pulp cells used in this study were cultured for 5 days (Fig. 2).

Based on one-way ANOVA statistical analyses, there were significant differences in cell viability between the control group and all chitosan-treated groups.

The viability of human dental pulp cells markedly increased after exposure to crab shell chitosan RGD scaffold membrane compared to the control group.

Based on one-way ANOVA statistical analyses, there was a significant difference in cell viability between the crab shell chitosan RGD scaffold membrane 1 mg treated group and the crab shell chitosan scaffold membrane 1 mg treated group (p<0.05) and between the crab shell...
chitosan RGD scaffold membrane 2 mg treated group and the crab shell chitosan scaffold membrane 2 mg treated group (p < 0.05).

In larger bone defects, especially one-wall periodontitis defects, the use of bone grafts is ineffective due to disadvantages such as immunology effects that can happen after transplantation [1]. Three decades ago, tissue engineering was developed as an alternative technique to treat larger bone defect [7]. Tissue engineering consists of three major components: Scaffold, stem cells, and growth factor. A material that can be used as a scaffold in bone tissue engineering is chitosan. It is biocompatible, biodegradable, and non-toxic. However, to increase chitosan’s efficacy, chitosan needs to be combined with a bioactive molecule such as RGD. A previous study reported that a chitosan scaffold combined with RGD could enhance the attachment and proliferation of osteoblasts in vitro.

In this research, crab shell chitosan RGD scaffold membrane and crab shell chitosan scaffold membrane produced by BATAN, Indonesia, and human dental pulp cells were used. Toxicity analyses were conducted using the MTT assay method. Based on this research, it was determined that crab shell chitosan RGD scaffold membranes did not decrease the cell viability of human dental pulp cells. This result was supported by other research using chitosan scaffolds with and without RGD. A preliminary study also showed that the proliferation of human dental pulp did not decrease after being exposed to a crab shell chitosan RGD scaffold membrane.

Fernandes et al. discovered that chitosan scaffolds were not toxic to rat osteoblasts [8]. Tsai et al. determined that the addition of RGD to chitosan scaffolds stimulated osteoblast cells to proliferate [9]. Prabhaharan and Sivashankari found that chitosan scaffolds combined with RGD were not toxic to chondrocyte and fibroblast cells [10]. A study by Chen (2016) showed that the viability of osteoblasts increased after being exposed to chitosan scaffolds combined with RGD [11].

To identify a dose-effect relationship, we used crab shell chitosan scaffold membranes with or without RGD in two different concentrations; 1 mg and 2 mg. As seen in Fig. 4, there is no significant difference in cell viability between the crab shell chitosan RGD scaffold membrane 1 mg treated group and the crab shell chitosan RGD scaffold membrane 2 mg treated group and between the crab shell chitosan scaffold membrane 1 mg treated group and the crab shell chitosan scaffold membrane 2 mg treated group.

Fig. 4 also shows that the viability of human dental pulp cells (%) treated with the crab shell chitosan scaffold membrane 1 mg treated group was significantly higher than the dental pulp cells treated with the crab shell chitosan RGD scaffold membrane 1 mg treated group. Like the 1 mg concentration conditions, the viability of human dental pulp cells (%) treated with the crab shell chitosan scaffold membrane 2 mg treated group was also higher than the dental pulp cells treated with the crab shell chitosan RGD scaffold membrane 1 mg treated group. Based on a statistical analysis using a one-way ANOVA, the difference was significant.

These results are in agreement with the preliminary study that used crab shell chitosan scaffold membrane and crab shell chitosan RGD scaffold membrane 1 mg and 2 mg, indicating that the level of cell proliferation of the human dental pulp cells was higher in the crab shell chitosan scaffold 1 mg group. Based on the data above, the viability percentage of human dental pulp cells treated with crab shell chitosan scaffold membrane without RGD was higher than the viability percentage of human dental pulp cells treated with crab shell chitosan RGD scaffold membrane. This result cannot be explained because only two samples were used in this study, which does not allow for the exploration of a dose and effect relationship.

In clinical applications, membrane scaffolds coat all damaged bone surfaces to transfer nutrients for the cells to achieve localized tissue repair. The addition of RGD to crab shell chitosan scaffold membranes is used to enhance cell attachment to the scaffold along with obtaining a good adhesion to the network. Cell adhesion is the first stage before cells can proliferate. This study shows that the addition of RGD to the crab shell chitosan RGD scaffold membrane did not have toxic effects on human dental pulp cells. In future studies, relations between effects and doses need to be determined to obtain optimal cell numbers and chitosan scaffold membrane levels. When scaffolds and implanted cells are transplanted to areas of bone damage, based on results on these types of studies, cell proliferation and cell differentiation can be ideal.

**CONCLUSION**

This study showed that the addition of RGD to the crab shell chitosan membrane did not have toxic effects on human dental pulp cells. Crab shell chitosan RGD scaffold membrane and crab shell chitosan scaffold membrane showed different effects on human dental pulp cells.

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