

Topical Single-dose Vascular Endothelial Growth Factor has No Effect on Soft Tissue Healing

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Abstract

Background: Vascular endothelial growth factor (VEGF) production in dermal wounds has been evaluated for evidence that it plays a probable role in wound healing. Events such as increased vascular permeability and concentration of inflammatory cells on the site of injury, produced by VEGF, were linked to tissue repair. **Aim:** The present study aimed to evaluate the effects of single-dose topical administration of VEGF on wound healing. **Materials and Methods:** A total of 30 male Wistar albino rats weighing 250-280 g were used in this study. In addition, 2-cm-long skin incisions were created over bilaterally exposed skin of the tibia region in each rat. VEGF plasmid 2 µg was administered locally into the right side wound bed of each animal. No other procedure besides skin closure was administered on the left side. To determine histologic assessments, skin samples were obtained from six anesthetized rats at each interval (4, 8, 12, 16 and 30 days) through excisional biopsy. The tissues were fixed in 10% neutral-buffered formalin for 1 week and then embedded in paraffin wax. Transverse sections of the embedded tissue 5-7 µm thick were stained with hematoxylin and eosin (H and E). **Results:** There was no significant difference regarding necrosis, epithelialization, inflammation, fibroblast activity, ulcerative formation, or hemorrhage between experimental and control groups. No statistically significant difference was found between the groups regarding granulation tissue formation and epidermal thickness. **Conclusion:** The administration method and dosage of VEGF is a major factor in terms of its effectiveness. The results of the present study did not evaluate the effectiveness of single-dose 2 µg topical administration of VEGF; however, various doses of VEGF plasmid should be tested in future studies in order to provide beneficial effects from topical administration of VEGF.

Keywords: Angiogenesis, Vascular endothelial growth factor, Vascular permeability, Wound healing

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Introduction

Angiogenesis, the formation of new blood vessels, is critical to physiological processes such as wound healing, tissue growth, intrauterine development, and reproductive functions. Wound healing presents the formation of granulation tissue, which may be characterized with fibrovascular tissue, including fibroblasts, collagen, and blood vessels. Angiogenesis is

regulated by the expression of various vascular growth factors and modulators, which are defined as engines driving wound repair. Among them, vascular endothelial growth factor (VEGF) is a potent, multifunctional cytokine that exerts some vital and independent actions on vascular endothelium. Being more potent than histamine, it serves as an inducer of vascular permeability. Increased vascular permeability occurs during the early phases of wound repair. Events such as increased vascular permeability and concentration of inflammatory cells on the site of injury, produced by VEGF, actively provide tissue repair. Increasing the migration and proliferation of pre-existing endothelial cells also assists in this process.^[1-3]

The VEGF gene encodes a protein that is often found as a disulfide-linked homodimer. The VEGF protein is a glycosylated mitogen that specifically acts on

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endothelial cells and has various effects, including mediating increased vascular permeability; inducing angiogenesis, vasculogenesis, and endothelial cell growth; promoting cell migration; and inhibiting apoptosis. The identification of increased vascular permeability accompanied with increased VEGF production in dermal wounds provided evidence for a probable role of VEGF in wound healing. VEGF seems to be a direct angiogenic factor that stimulates endothelial cell migration. Furthermore, various animal studies have been published investigating the upregulation of VEGF production in wound healing.^[4]

The present study was carried out to evaluate the effects of single-dose topical administration of VEGF plasmid on the healing of induced rat skin wounds.

Materials and Methods

The study was conducted at the Department of Genetics, Institute of Experimental Medicine, Istanbul University. The study protocol was approved by the Istanbul University Animal Care Ethical Committee. All procedures were conducted in accordance with the Istanbul University ethical guidelines for the treatment and welfare of experimental animals. In total, 30 male Wistar albino rats weighing 250-280 g were used in this study. The animals were housed at 21°C and were given tap water and standard rat food.

Plasmid construction

The VEGF-A expression plasmid was constructed by ligating full-length human VEGF-A (Gene ID: 7422) cDNA into pcDNA4 (Invitrogen, Carlsbad, USA). Briefly, total RNA was isolated from cultured cells using a total RNA purification kit according to the manufacturer's recommendations (QiagenCo. Hilden, Germany). After reverse transcription, the VEGF-A cDNA PCR was amplified with the forward primer 5' AATTCGCCG ACATGACGGACAGACAG ACAGACACCGCC 3' containing EcoRI sites and the reverse primer 5' TCTAGATCAC CGCCTCGGCTTGCA CATCTGC 3' with an modified XbaI site at their 5' and 3' ends. The purified PCR fragments and pcDNA4 were double-digested with EcoRI and XbaI restriction enzymes. The digested fragments recovered from 1% low-melting agarose gel by cutting, and DNA was eluted from the sliced agarose gel using an extraction kit (Qiagen Co. Hilden, Germany). In addition, 50 ng of EcoRI and XbaI digested, dephosphorylated vector DNA and 1 µl of PCR product were incubated overnight with DNA ligase buffer containing T4 DNA ligase at 16°C. Further, 3 µl of ligation reaction and 100 µl of DH5 alpha competent cells were heat-shock transformed at 42°C and spread on agar plates containing 100 µg/ml ampicillin

(Invitrogen, Carlsbad, USA). Plates were incubated overnight at 37°C. PCR-tested and selected positive clones were confirmed with sequencing (Refgen Co., Turkey). Plasmid was purified using an Endo-free Maxi plasmid kit (Qiagen Co. Hilden, Germany). Then, 2 µg of the plasmid constructs, bearing a human VEGF-A DNA, were administered to the rat. Empty pcDNA4 plasmid (2 µg) was used as a control.^[5,6]

Surgical procedure

The animals were anesthetized via intramuscular injection of ketamine hydrochloride (50-100 mg per kg of body weight). Bilateral tibial regions of the rats were shaved with electric clippers and cleansed with 10% povidone iodine and 70% alcohol swabs before manipulation. Next, 2-cm-long skin incisions were created over bilaterally exposed skin of the tibia region in each rat.

VEGF plasmid (2 µg) was administered locally into the right side wound bed of each animal (experimental group). No extra procedure besides skin closure was administered on the left side (control group). Incision lines were primarily sutured with 3.0 silk (Dogsan Medical Supplies Industry, Trabzon, Turkey) immediately after surgery in all rats.

To determine histologic assessments, skin samples were obtained from six anesthetized rats at each interval (4, 8, 12, 16 days and 1 month) through excisional biopsy, including incision lines, cleared of surrounding mesentery and fat and washed with saline. The anesthetized rats were then sacrificed using intraperitoneal 135 mg/kg sodium pentothal.

Histological analysis

The tissues were fixed in 10% neutral-buffered formalin for 1 week and then embedded in paraffin wax. Transverse sections of the embedded tissue, 5-7 µm thick, were stained with hematoxylin and eosin (H and E). Histological assessment was carried out by an experienced pathologist.

The sections were examined under light microscopy regarding necrosis, epithelialization, inflammation, fibroblast activity, ulceration and hemorrhage. The mean area percentage of granulation tissue was detected in three randomly chosen fields for each section under microscope (X200 magnification) and scored as the following: (+) for 0-10 %; (++) for 10-30%, and (+++) for 30-100 %. Epidermal thickness was calculated in millimeters.

Statistical study

The differences in epidermal thickness between the treatment and control groups at time intervals of the

4, 8, 12, 16 and 30 days, based on all z-scores, were compared using the Mann-Whitney U-test. At each time interval, the granulation scores of the treatment group were compared with the granulation scores of the control group using Pearson's chi-square test across treatment groups or Fisher's exact test, if 20% of the expected counts were < 5. The treatment comparison was performed at an α level of 0.05. All data were analyzed using SPSS for Windows version 21 (IBM SPSS Statistics, New York, USA).

Results

There was no mortality among the animals during the study period. The rats were macroscopically observed on each day of the study period. No complications, such as infection and exposure around the wound margin, which result in secondary healing, were observed. During the early period of wound healing, variables were evaluated and compared at intervals (4, 8, 12, 16 days and 1 month). There was no significant difference with respect to necrosis, epithelialization, inflammation, fibroblast activity, ulcerative formation, or hemorrhage between experimental and control groups. No necrosis was found in any of the rats. The mean area percentage of granulation tissue was more than 30% in the 12th- and 16th-day groups (experimental and control), whereas the mean area percentage of granulation tissue was less than 30% in the remaining groups; however, there was no statistically significant difference between the groups with respect to granulation tissue formation ($P > 0.05$) [Table 1]. Epidermal thickness scores were given in Table 2. No statistically significant difference was found in epidermal thickness between the groups ($P > 0.05$) [Table 2].

4th postoperative day

In the experimental group, a few ulcerations and desiccate formations were observed on the surface of the epidermis. A number of vascular sections filled with red blood cells and surrounded by chronic inflammatory cells infiltration were observed. Fresh bloody focal points were present in one sample [Figure 1].

In the control group, ulcerative formation in epidermis was present in all rats. There was desiccate formation on the epidermal surface in two out of six rats. Vascular sections and intense inflammatory cell infiltration was observed in the edematous connective tissue under the desiccated areas [Figure 1].

8th postoperative day

Both groups showed similar wound-healing outcomes. Continuing ulcerative formation was observed in one rat in the experimental group and three rats in the control

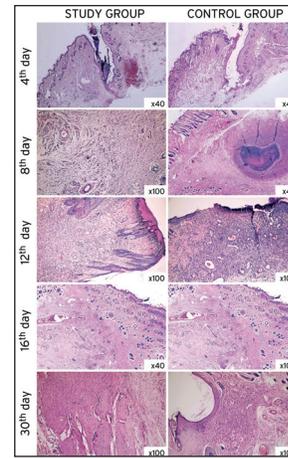


Figure 1: Histological sections of the study and control groups according to the time intervals

Table 1: Granulation scores of the study and control groups at each time intervals

| | | SCORES | | | P |
|--------|---------|--------|----|-----|----------------------------|
| | | + | ++ | +++ | |
| Day 4 | Control | 0 | 0 | 6 | No statistics are computed |
| | Study | 0 | 0 | 6 | |
| Day 8 | Control | 0 | 5 | 1 | 0, 50 |
| | Study | 0 | 4 | 2 | |
| Day 12 | Control | 0 | 1 | 5 | 0, 30 |
| | Study | 0 | 0 | 6 | |
| Day 16 | Control | 0 | 1 | 5 | 0, 30 |
| | Study | 0 | 0 | 6 | |
| Day 30 | Control | 0 | 6 | 0 | No statistics are computed |
| | Study | 0 | 6 | 0 | |

Table 2: Epidermal thickness of the study and control groups at each time intervals (mm)

| | Group | Mean | SD | Z value | P |
|--------|---------|--------|--------|---------|------|
| Day 4 | Control | 0.0270 | ,00179 | -0.971 | 0.33 |
| | Study | 0.0300 | ,01789 | | |
| Day 8 | Control | 0.0267 | ,01633 | -0.968 | 0.33 |
| | Study | 0.0360 | ,00179 | | |
| Day 12 | Control | 0.0333 | ,01211 | -0.686 | 0.49 |
| | Study | 0.0393 | ,00103 | | |
| Day 16 | Control | 0.0400 | ,01095 | 0.000 | 1.0 |
| | Study | 0.0400 | ,01789 | | |
| Day 30 | Control | 0.0400 | ,01789 | -0.490 | 0.62 |
| | Study | 0.0450 | ,01225 | | |

group. Abscess formation in dermis was observed in one rat in the experimental group; inflammatory cell infiltration of the connective tissue was present in the remaining rats in the experimental group. Intense abscess formation and inflammatory cell infiltration of connective tissue was observed in all samples in the control group [Figure 1].

12th postoperative day

In the experimental group, the initial phase of epidermis regeneration was seen in all samples, and continuing abscess formation in connective tissue was seen in one individual. Dermal infiltration of inflammatory cells was reduced compared to 8th-day outcomes, and the initial phase of fibrotic activity was observed [Figure 1].

In the control group, there was no exposed area on the epidermal surface, except for one individual with continuing ulcerative formation on the surface. Fibrosis was observed in a few areas in the connective tissue [Figure 1].

16th postoperative day

Significant reduction of inflammatory infiltration in connective tissue was observed in the experimental group. Epidermal healing was also observed in all rats in the control group. Fibrotic focal spots became marked in all samples in both groups [Figure 1].

30th postoperative day

Complete regeneration of epidermis and intense activity of dermis were both observed. Fibrosis penetration into deep tissues was observed in the experimental group [Figure 1].

In the control group, epidermal regeneration was almost completed in all samples, and fibrosis of superficial areas in connective tissue was clearly seen [Figure 1].

Discussion

VEGF has been proven to be an effective factor in wound healing immediately after injury. Acute inflammation, re-epithelialization, formation of granulation tissue, and tissue remodeling are phases of wound healing that occur on an ongoing basis. VGEF has a role in wound healing by means of an angiogenic process. Angiogenic activity is maximally observed between days 3 and 7; VEGF is up-regulated to promote the early stages of angiogenesis.^[1,7] In the present study, the findings of early angiogenesis could be detected when 4th-day results and 8th-day results were examined, while the features of normal wound repair such as fibrovascular tissue, collagen and blood vessels were observed in all groups throughout the 30-day experiment. However, the ineffectiveness of the administration of VEGF on the following days seems unpromising.

Granulation tissue is eventually remodeled, as vessels are resorbed and fibroblasts disappear. As the wound is granulated, the process of angiogenesis stops, and blood vessels decline as endothelial cells undergo apoptosis.^[1,7] In our study, granulation tissue was observed most

between days 12 and 16; area percentage of granulation tissue was detected to be less than 30% in the remaining groups. Contrary to our findings, Corral *et al.*, found that single-dose treatment at 30 µg per wound improved granulation tissue formation.^[8] This conflict may be due to the fact that receptor regulation may be more important than growth factor regulation during wound healing.^[9] Wound surface is lined with macrophages that secrete VEGF. Collagen deposition is one of the responses of the endothelial cells to this. Collagen deposition is expected to be excessive in the first 2 weeks and to regress by the end of the second week;^[10] however, there was no difference between the groups.

In the present study, even though some inflammation is needed for wound angiogenesis,^[10] the utmost care was taken to avoid any possibility of trauma and inflammation, which may cause cytokine release and thus affect wound healing. No deposition of unhealthy granulation tissue and delayed closure of the wound were observed. The epidermis regained its thickness in treated and non-treated groups during the phases of the experimental process. Nevertheless, no significant differences were found between any of the groups with regard to epidermal thickness, meaning that a single dose of VEGF plasmid injection, known as a mitogen selective for endothelial cells, was not adequate to increase the capability of inhibiting apoptosis. Corral *et al.*, also claimed that VEGF had no effect on new epithelium formation in either ischemic or nonischemic wounds, whereas Galiano reported that topical VEGF application accelerated cutaneous repair.^[8,11] However, a short dosing interval of every-other-day administration for five doses in total was reported to be used in Galiano's study.

VEGF induces interstitial collagenase expression in human endothelial cells.^[12] When it is considered that the collagen deposition revealed no difference between the groups, it may be stated that single-dose topical administration of VEGF could neither induce endothelial cells grown on the surface of the collagen matrix nor stimulate the proliferative response.

The authors of this paper are aware that the effect of VEGF is highly dependent on its administration method and dosage.^[8,11] While the results of the present study did not evaluate the effectiveness of single-dose 2 µg topical administration of VEGF plasmid, various doses of VEGF should be tested in future studies in order to provide beneficial effects from topical administration of VEGF.

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