

Preoperative Shock Wave Therapy Reduces Ischemic Necrosis in an Epigastric Skin Flap Model

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Abstract: Extracorporeal shock wave therapy (ESWT) has recently been demonstrated to improve skin flap survival. In all these studies ESWT was applied immediately after the surgical intervention. Thus, the purpose of this study was to determine the preoperative effect of ESWT as a noninvasive technique to precondition flap tissue in a rat epigastric skin flap model.

ESWT and control groups each contained 10 animals. ESWT was applied 7 days before the surgical intervention, whereas the control group received no treatment. Follow-up evaluation was performed on postoperative day 5. The mean area of flap necrosis, expressed as a percentage of the total flap area, was calculated. A significant reduction of the average flap necrosis area was observed in the ESWT group ($27.2\% \pm 9.6\%$) compared with the control group ($46.1\% \pm 7.9\%$) ($P < 0.05$).

In summary, this study indicates that preoperative ESWT may enhance skin flap survival in a rodent model.

Key Words: shock wave therapy, ischemic reperfusion injury, preconditioning, improving flap survival rate

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Failure of composite flaps, due to ischemia injury, remains a serious clinical problem. During the last decade several techniques have been established to improve the survival of flaps by an increase of the resistance to further ischemia. All these techniques seem to have in common that they expose and precondition the tissue to a sublethal degree of environmental stress, which may be applied by a variety of physical and pharmacological stimuli.¹

Recent studies reported an induction of angiogenesis together with release of angiogenic factors mediated through extracorporeal shock wave (ESW).^{2–5} Animal studies in a rat model showed that ESW enhances the distal area of skin flaps, while ESW application was admitted immediately after the operation.^{2,3,6} The present study was thus designed to examine whether ESW therapy also is effective in the preoperative treatment to ameliorate flap survival.

MATERIALS AND METHODS

Male adult Wistar rats with a weight between 200 and 300 g were used. All animals were maintained in accordance to the guidelines of the German Animal Welfare Act. The experimental protocol was approved by a review committee of the state of

Baden-Württemberg, Germany. Seven days before the surgical intervention all animals were anesthetized with an intraperitoneal injection of ketamine hydrochloride (100 mg/kg; Ketanest 100 mg/mL; Fort Dodge Laboratories, Fort Dodge, Iowa) and xylazine (5 mg/kg; Rampun; Bayer Corp, Stilwell, Kansas) and randomly divided into 2 groups (ESW group and control group) of 10 animals each.

Epigastric Skin Flap Surgical Procedure

The operative procedure was identical for all groups. All operative procedures were performed under aseptic conditions. Animals were anesthetized with an intraperitoneal injection of ketamine hydrochloride (100 mg/kg; Ketanest 100 mg/mL; Fort Dodge Laboratories) and xylazine (5 mg/kg; Rampun; Bayer Corp). With the animals in a state of deep anesthesia, an extended epigastric adipocutaneous flap (6 × 10 cm) based on the left superficial epigastric vessels was raised. The right superficial epigastric vessels were ligated and the flap was sutured back to its native configuration and placed onto a silicon sheet to prevent neovascularization. To avoid autocannibalism, protective collars made of plastic film were applied to all the animals.⁷

ESW Group

Rats were placed in a supine position and an epigastric flap measuring 10 × 6 cm was outlined with a permanent marker on the abdominal skin. Ultrasound transmission gel was used as a contact medium between the ESW apparatus and skin. Shock wave was applied one-time in a dose of 500 impulses at 0.11 mJ/mm² and 8 Hz, using an electrohydraulic shock wave device with focusing applicator (dermaPACE, Sanuwave Inc., Marietta, GA) to the right upper corner of the flap. This area represents the random portion of the flap that predictably undergoes necrosis.^{3,6,7}

The reason for the dosage and the timing that we applied ESW of 500 impulses at 0.11 mJ/mm² followed previous experiences.^{2,3,8–11}

Control Group

In this group the flap was raised and sutured back without preoperative ESW treatment.

Evaluation

Follow-up evaluation was performed on postoperative day 5. Therefore all animals were killed and standardized digital pictures of the flap were taken and transferred to the computer (Fig. 1A, B). The mean area of flap necrosis, defined by surgical borders (expressed as a percentage of the total flap area) was calculated for each animal, using a self-programmed planimetry software. The results were expressed as percentage relative to surface area of the total flap.

All values were calculated as mean ± SD. Statistical analysis was performed with F and Student *t* test. A *P* value of less than 0.05 was considered statistically significant.

RESULTS

No animal was eliminated from the study because of infection, seroma, hematoma formation, or autocannibalization. There

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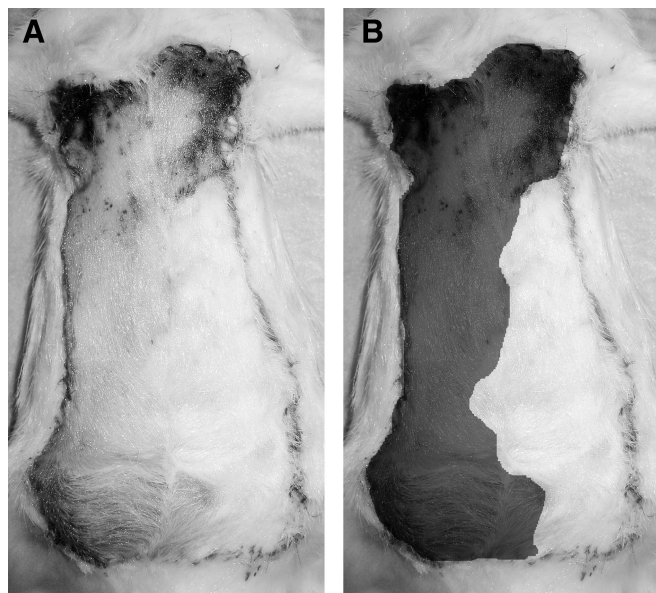


FIGURE 1. A, B, Characteristic sample of the control group at day 5 after surgery. The necrotic zone of the flap was measured by using a self programmed planimetry software.

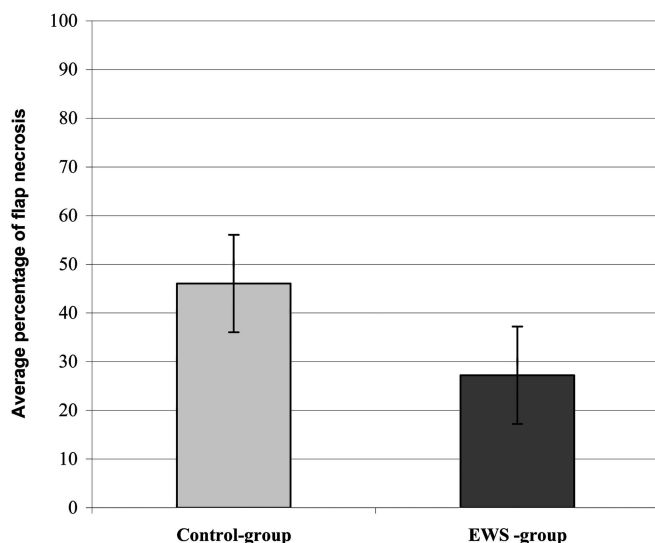


FIGURE 2. Diagram showing the percentage of flap necrosis in the EWS group compared with the control group.

was a significant increase of the average surviving area of the ESW-group compared with the control group ($P < 0.05$). The ESW group showed an average flap necrosis of $27.2\% \pm 9.6\%$, whereas the control group had an average area of flap necrosis of $46.1\% \pm 7.9\%$ (Fig. 2).

DISCUSSION

Skin flap necrosis in the distal flap areas is still a major challenge in reconstructive surgery and several techniques have been described to improve blood supply and tissue perfusion in compromised tissues.

Previous animal studies suggest that ESW treatment has a positive effect in rescuing ischemic zone of flap by an increase of tissue perfusion. ESW treatment was first introduced into medicine for the treatment of urolithiasis, but has been widely adopted for a number of musculoskeletal disorders including nonunion of long bone fractures, calcifying tendonitis, or plantar fasciitis.^{12–15} Low energy shock ESW therapy effectively induced angiogenesis and improved chronic ischemia in animals and humans.^{16–18} Further studies showed a significant rise of growth factors such as VEGF as well as endothelial nitric oxide synthase and proliferating cell nuclear antigen after ESW treatment.² With regard to plastic surgery perspectives, ESW therapy has recently been demonstrated to improve skin flap survival.^{2,3,6} In all these studies ESW was applied immediately after the surgical intervention. Thus, the purpose of this study was to determine the preoperative effect of ESW as a noninvasive technique to precondition flap tissue. According to the findings from earlier reports, our current study confirms an enhanced flap survival in the animal model. Although the exact mechanism of ESW therapy remains unknown, we hypothesize that mechanical forces of the ESWT stimulate cell proliferation and vascular remodeling in living skin. As cell growth and vascular supply are critical to wound healing and tissue expansion, devices applying controlled mechanical loads to tissues may be a powerful therapy to increase growth factors induced in-growth of neovascular formation. This is an early study in the preoperative treatment with ESW for flap tissue. Because of limitation of our experimental design further studies are planned to determine the mechanisms which are involved in the effectiveness of ESWT. Although several approaches have been developed to reduce ischemic skin flap necrosis, the potential of ESW treatment looks very promising avoiding complications or adverse effects.^{4,16,17,19}

As a result, the study indicates that preoperative ESW treatment may enhance skin flap survival in a rodent model, proving its potential to become a viable, noninvasive, and cost-effective method to improve the blood supply in ischemic tissue.

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