Effect of vacuum sealing drainage on healing time and inflammation-related indicators in patients with soft tissue wounds

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Abstract

The aim of this study was to investigate the therapeutic effects of vacuum sealing drainage (VSD) on wound repair time and inflammation-related indicators in patients with soft-tissue wounds in comparison with traditional treatment. From January 2018 to January 2020, 130 enrolled patients with soft-tissue wounds were randomly divided into two groups: VSD group (65 cases) and routine dressing change (RDC) group (65 cases). The inflammation-related indicators including erythrocyte sedimentation rate (ESR), C-reactive protein (CRP), white blood cell (WBC), and procalcitonin (PCT) of preoperative stage and postoperative day 3 (POD 3) and POD 7 were recorded. Wound healing was observed 3 and 7 days after treatment, and the clinical efficacy, changes in the wound (coverage rate and thickness of granulation tissue and bacterial clearance rate), wound-cleaning time, wound-healing time, and hospital stay time were recorded after treatment as well. No significant difference was observed in terms of the baseline between the two groups. On POD 3 and POD 7, CRP, WBC, and PCT levels in the VSD group were lower than those in the RDC group, while ESR levels were higher, with significant differences (\(P < .05\)). After treatment, the wound-cleaning time, wound-healing time, and hospital stay time of the VSD group were all lower than those of the RDC group, with significant differences (\(P < .05\)). VSD has a significant effect on the treatment of patients with soft-tissue wounds, which can effectively shorten the time of wound healing and reduce inflammation-related indicators. Compared with traditional RDC, VSD is more worthy of clinical application.

KEYWORDS

vacuum sealing drainage, soft-tissue wound, healing time, inflammation-related indicators, effect...
1 | INTRODUCTION

A soft tissue wound is a common and frequently occurring disease in clinical practice. Because of the complexity of the injuries inside the wound, it is often accompanied by severe injuries of the cloven hoof tissue, such as skin, fascia, tendon, and fat, which may damage internal organs in severe cases.\(^1\) When the above-mentioned natural protection barrier of the human body is damaged, the deep tissue is directly exposed to the external environment, which is prone to infection and secondary damage to the wound, making it difficult for the wound to heal and increasing the difficulty of clinical treatment.\(^2\) Without timely and effective treatment, local blood supply will be affected, leading to bacterial infection, tissue necrosis, and liquefaction, which is extremely difficult to cure. Currently, debridement and dressing change are often used in the clinic to heal soft-tissue wounds, and skin grafting is followed when certain conditions are achieved. However, because of the time and effort of debridement, the effect is not ideal, which seriously affects the treatment compliance of patients.

The vacuum sealing drainage (VSD) technique has been widely applied for effectively treating diverse wound surfaces, like tumour, chronic ulcer, and soft-tissue wounds.\(^3\)\(^-\)\(^5\) Shen’s team found that VSD could reduce the dressing change frequency, shorten the operation time and hospital stay, accelerate wound healing, and reduce postoperative infection and lower-limb deep venous thrombosis in patients with orthopaedic trauma.\(^6\) A meta-analysis also demonstrated that VSD is a more effective therapy and is associated with a greater decrease in wound size and shorter time to wound healing compared with traditional therapy of diabetic foot ulcers.\(^7\) Many studies on VSD have been carried out, and it is certain that the overall efficacy of this method is significantly better than that of traditional dressing change treatment.\(^8\) However, there are few studies on the healing time and inflammation-related indicators in patients with soft-tissue wounds treated with VSD. Here, this study aims to explore the specific efficacy and inflammation-related indicators of VSD in patients with soft-tissue wounds so as to provide reference for the treatment of clinical patients with soft-tissue wounds.

2 | MATERIALS AND METHODS

2.1 | Patients

All patients provided informed consent for undergoing treatment. A total of 130 patients with soft-tissue wounds, admitted in our hospital from January 2018 to January 2020, were categorised into a VSD group and a routine dressing change (RDC) group randomly and averagely according to the random number table method. This study conformed to the relevant requirements of the World Medical Association Declaration of Helsinki.

2.2 | Inclusion and exclusion criteria

Inclusion criteria: All patients with (a) a wound unable to be sutured, heal by itself, or still present after debridement period I; (b) normal functions of important organs such as heart, liver, and kidney; and (c) normal spirit and good compliance were included.

Exclusion criteria: Patients with (a) malignant tumours, (b) poor blood sugar control, (c) ischaemic wounds accompanied by venous thrombosis, (d) sepsis or severe systemic soft-tissue injury, (e) severe secondary infection after trauma, and (f) serious complications and (g) women who were pregnant or were breastfeeding were excluded.

2.3 | Methods

All patients underwent debridement and repair of damaged skin, blood vessels, nerves, and tendons. RDC was applied once every 1 to 2 days: washing the wound with oxydol (3% hydrogen peroxide; State Food and Drug Administration [SFDA] approval number: H44023919), iodophor (Appliance Permit Number 2642634 of 2017 by Jiangsu Food and Drug Surveillance Authority), and normal saline (0.9% NaCl, 5 mL; SFDA approval number: S10870001) and smearing debridement adhesive (15 g, URGO HYDROGEL; Overseas Medical Devices Registration Record: 20193141636, United Kingdom). The wound was covered with normal saline gauze ((90-160) cm × (100-600) cm; Appliance Permit Number 2640347 of 2016 by Jiangsu Food and Drug Surveillance Authority) and bound with sterile dressing (Appliance Permit...
Number 2640032 of 2017 by Shanxi Food and Drug Surveillance Authority). RDC group patients were also treated with anti-inflammation, anti-infection, and nutrition support.

The VSD group was given RDCs, followed by closed negative-pressure drainage dressing according to the size of the wound, shape, depth, etc. If wounds were too large, more dressing and a vacuum therapeutic sponge adsorption pad (disposable closed negative-pressure drainage sponge; Appliance Permit Number 20140621 of 2020 by Henan Food and Drug Surveillance Authority) of a suitable size were applied to cover the wound surface to ensure complete coverage and no space, with a drainage tube (diameter: 4.67 mm [F14], type I; Appliance Permit Number 2140543 of 2016 by Jiangsu Food and Drug Surveillance Authority) connecting the pressure suction device. The wound surface was closed with a medical biological semipermeable membrane (medical transparent drug film; Appliance Permit Number 2641763 of 2014 by Hubei Food and Drug Surveillance Authority) tightly attached to the skin, with the outer edge of the membrane covering the normal skin outside of the wound edge by at least >2 cm. The negative pressure was set to be −150 to −300 mm Hg (1 mm Hg = 0.133 kPa) for 5 to 7 days, during which the patient’s condition should be continuously observed. Meanwhile, the level and colour of drainage fluid should be monitored, and the drainage bottle should be updated regularly.

2.4 | Observation indicators

The levels of inflammation-related indicators, including erythrocyte sedimentation rate (ESR), C-reactive protein (CRP), white blood cell (WBC), and procalcitonin (PCT), were recorded before treatment and 3 and 7 days after treatment. Wound healing was observed 3 and 7 days after treatment, and clinical efficacy, changes in the wound (coverage rate and thickness of granulation tissue and bacterial clearance rate), wound-cleaning time, wound-healing time, and hospital length of stay were recorded after treatment as well.

Inflammation-related indicators: 3 to 5 mL of elbow venous blood was collected for WBC detection using an HCC200Plus full-automatic analyser, and for ESR was detected using a PUC-2068 full-automatic dynamic ESR analyser. PCT and CRP were detected by TRACE immunoassay.

Wound healing: The area of the residual wound was measured using a right-angle measuring ruler.

Clinical efficacy: According to a literature study,9 we defined recovery as the healing of all wounds within 2 weeks with good epidermal coverage; marked effect as granulation and epithelial tissue of the wound growing well, and the area was reduced by more than 75%; effective as new granulation tissue was found on the wound surface, and the area was reduced by 40% to 75%; and ineffective as no significant improvement, and the wound area reduced by less than 40%.

Wound-cleaning time: No liquid effusion was observed for 3 days with clean wound and healthy granulation.

Wound-healing time: Successful skin grafting or flap transfer surgery was performed.

Granulation tissue coverage: Before treatment and 7 days after treatment, Adobe Photoshop CS10.0 software was used to calculate granulation tissue coverage. Granulation tissue coverage = granulation tissue coverage area/wound area ×100%.

Bacterial clearance rate: Wound secretions were taken for bacteriological culture before treatment and 7 days after treatment. Bacterial clearance rate = (the number of bacteria before debridement − the number of bacteria after 7 days of treatment)/the number of bacteria before debridement ×100%.

Granulation tissue thickness: Granulation tissue thickness was measured using a micrometre before and 7 days after treatment. Granulation tissue thickness = granulation tissue thickness of 7 days after treatment − granulation tissue thickness before debridement.

2.5 | Statistical analysis

IBM Microsoft SPSS 21.0 software was used for data statistical analysis. The measurement data were expressed as and a t test was used for comparison. Enumeration data were expressed as ratio (%) using a chi-square test. was considered statistically significant.

3 | RESULTS

3.1 | Baseline characteristics

The baseline characteristics of the two groups were comparable. The age range of the VSD group was 22 to 53 years, and the mean age was 44.29 ± 6.32 years. Patients in the RDC group ranged in age from 21 to 52 years, with an average age of 43.97 ± 6.47 years (Table 1).

3.2 | Clinical efficacy

After treatment, the clinical efficacy of the VSD group was obvious compared with that of the RDC group
3.3 | Wound changes

After treatment, the granulation tissue coverage rate, bacterial clearance rate, and granulation tissue growth thickness of the VSD group were significantly higher than those of the RDC group ($P < .05$), as shown in Table 3.

3.4 | Wound healing

The wound-healing areas of the VSD group and RDC group were $11.76 \pm 2.08 \text{ cm}^2$ and $8.32 \pm 1.65 \text{ cm}^2$ on postoperative day (POD) 3 and $15.38 \pm 1.09 \text{ cm}^2$ and $12.21 \pm 1.23 \text{ cm}^2$ on POD 7, respectively, with significant differences ($P < .05$).

3.5 | Inflammation-related indicators

Before treatment, the levels of ESR, CRP, WBC, and PCT in the two groups were not significantly different ($P > .05$). Three and 7 days after treatment, CRP, WBC, and PCT levels in the VSD group were lower than those in the RDC group, while ESR levels were higher than those in the RDC group, with significant differences ($P < .05$), as shown in Table 4.

3.6 | Time of wound cleaning, wound healing, and hospital stay

After treatment, the wound-cleaning time, wound-healing time, and hospital length of stay of the VSD
group were all lower than those of the RDC group, with significant differences ($P < .05$), as shown in Table 5.

### DISCUSSION

Soft-tissue wound healing follows the pattern of the four phases of haemostasis, inflammation, proliferation, and maturation/matrix remodelling.¹⁰ There are many interfering factors, which will impede the progress of wound healing.¹¹ For the treatment of soft-tissue injury, the wound is usually cleaned thoroughly and then covered with gauze directly. The main theoretical basis is that atmospheric oxygen is involved in wound healing in a dry environment, and the permeability of gauze can provide enough oxygen for the wound to ensure cell growth.¹² However, clinical studies have confirmed that this method will lead to dewatering of wounds, resulting in the loss of active substances and the formation of scab. In addition, it is easier to adhere dressings to fresh tissues, and there will be pain when changed, which may even lead to re-injury and prolonged healing time.¹³ At the same time, an open wound is vulnerable to bacterial invasion, which has certain limitations.

The VSD technique can promote the growth of granulation tissues, eliminate necrotic tissues and effusion, and decrease the incidence of bacterial infection around wounds by increasing local blood perfusion. The application of VSD could reduce oedema and inflammation in wounds, which accelerates the healing of wounds. Because of the above advantages, the technique has been prevalent in the treatment of chronic wounds like postoperative incision infection, soft-tissue cleft, open fracture or infection, ulcers, acne, purulent infection of body surface or joint cavity, and skin graft.¹⁴⁻¹⁷ Although VSD has certain clinical advantages in the treatment of soft tissue of traumatic wounds, most of the studies were retrospective analyses, and the observation indicators were not comprehensive. Therefore, this prospective cohort study was designed to investigate the effects of VSD on the wound repair time and inflammation-related indicators in patients with soft-tissue wounds.

The results of this study showed that, after treatment, the clinical efficacy of the VSD group was significant ($P < .05$), indicating that VSD can effectively improve the clinical symptoms of patients with soft-tissue wounds and is conducive to the growth of fresh granulation tissue, with significant effect. Hou et al¹⁸ confirmed that the

### TABLE 4  The comparison of inflammation-related indicators in two groups

| Time                  | Vacuum sealing drainage group | Routine dressing change group | $t$  | $P$ value |
|-----------------------|-------------------------------|-------------------------------|------|-----------|
| Erythrocyte sedimentation rate (mm/h) | Preoperative 5.76 ± 1.04       | 5.72 ± 0.98                   | 0.226| .822      |
|                       | Postoperative day (POD) 3 48.07 ± 4.32 | 39.93 ± 3.29                  | 12.086| <.001     |
|                       | POD 7                         | 50.87 ± 1.02                   | 6.131| <.001     |
| C-reactive protein (mg/L) | Preoperative 16.83 ± 1.98    | 16.84 ± 1.53                   | −0.032| .975     |
|                       | POD 3                         | 7.87 ± 0.67                    | −43.297| <.001     |
|                       | POD 7                         | 6.07 ± 0.21                    | −51.995| <.001     |
| White blood cell ($\times10^9$/L) | Preoperative 9.08 ± 0.38     | 9.07 ± 0.32                    | 0.162| .872      |
|                       | POD 3                         | 7.65 ± 0.23                    | −24.851| <.001     |
|                       | POD 7                         | 6.83 ± 0.18                    | −15.863| <.001     |
| Procalcitonin (ng/L)  | Preoperative 2.39 ± 0.23      | 2.38 ± 0.24                    | 0.243| .808      |
|                       | POD 3                         | 0.23 ± 0.07                    | −51.594| <.001     |
|                       | POD 7                         | 0.08 ± 0.02                    | −86.533| <.001     |

### TABLE 5  The comparison of wound cleaning time, wound healing time, and hospital length of stay (x ± s, days)

|                  | Vacuum sealing drainage group | Routine dressing change group | $t$  | $P$ value |
|------------------|-------------------------------|-------------------------------|------|-----------|
| Cleaning time    | 12.94 ± 0.32                  | 14.32 ± 0.18                  | −30.303| <.001     |
| Healing time     | 16.98 ± 0.65                  | 20.76 ± 0.43                  | −39.103| <.001     |
| Hospital stay    | 16.54 ± 0.19                  | 19.87 ± 0.32                  | −72.140| <.001     |
frequent use of negative-pressure drainage for soft-tissue wounds could promote wound healing. Patients treated with VSD 1 to 3 times were more in line with the therapeutic indications of surgical skin grafting. Agel et al\(^{19}\) confirmed that VSD can promote the blood circulation of the wound, clean up foreign bodies completely, facilitate the formation of granulation tissue, create conditions for tissue plasticity, and ultimately promote wound healing. This study showed that, after treatment, the granulation tissue coverage, bacterial clearance rate, and thickness of the granulation tissue growth were higher in the VSD group \((P < .05)\), which illustrated that VSD can effectively eliminate bacteria from patients with soft-tissue wounds, provide the conditions for the growth of granulation tissue, and then promote granulation tissue growth. In this study, wound healing was further analysed, and the results showed that wound healing in the VSD group was better than that in the RDC group on PODs 3 and 7 \((P < .05)\), indicating that the implementation of VSD is beneficial in healing patients with soft-tissue wounds. This might be because VSD can achieve zero aggregation of exudates and necrotic tissues on the wound, and sufficient drainage of foreign bodies is conducive to the rapid and good growth of granulation tissues on the wound, thus facilitating wound healing. In addition, the reduced frequency of dressing changes in the VSD method can decrease the contact of the wound with the outside environment and the workload of medical staff, thus minimising the interference to wound healing.

PCT and CRP were important acute reactive proteins, the synthesis and levels of which will increase when the body has an acute reaction. \(^{20}\) PCT is the propeptide substance of serum calcitonin and a sugar protein composed of 116 amino acids. PCT is induced by inflammatory factors such as bacterial endotoxin and interleukin in the first 2 to 4 hours of systemic inflammation, and its level is closely related to the severity of disease. \(^{21}\) CRP is mainly synthesised by hepatocytes and has the function of activating complement, strengthening phagocytic macrophages, and promoting granulocytes. It is extremely low in a normal body, and its concentration in blood rises rapidly when the body has an infection or acute trauma. \(^{22}\) The ESR is a kind of non-specific inflammation-related indicator, indicating the sedimentation rate of red blood cells under certain conditions, and its level is relatively stable in normal people. When inflammation, tissue damage, or infection occurs in the body, red blood cells will overlap with each other, reducing the resistance by plasma and promoting the increase of ESR levels in the body. \(^{23}\) WBC is a clinically commonly used marker for the detection of inflammatory response. \(^{24}\) It can resist pathogenic microorganisms in the body directly and increases when the body is invaded by pathogenic microorganisms. However, because of the difference in age, gender, and other factors, its detection specificity is low. \(^{25}\) In recent years, ESR, CRP, WBC, and PCT levels have often been used in clinical studies on postoperative infection, \(^{26}\) but there are few studies on the effect of inflammation-related indicators in patients with soft-tissue wounds treated by VSD. Therefore, this study showed that on PODs 3 and 7, the CRP, WBC, and PCT levels in the VSD group were lower than those in the RDC group \((P < .05)\), indicating that VSD could effectively reduce the inflammation-related indicators in patients with soft-tissue wounds. However, the ESR level was higher than those in the VSD group because there might be many factors affecting red blood cell overlap, such as immunoglobulins or fibrinogen. \(^{27}\) The efficacy of VSD is obvious, which might be because the negative-pressure closure drainage could isolate the external environment using biological translucent membrane closure, forming a barrier to prevent cross-infection, reducing the level of serum inflammation-related indicators and the use of antibiotics.

Duan and his team\(^{4}\) illustrated the outstanding effect of VSD with instillation in removing the debris of necrotic tissue on the wound bed, in the continual and complete drainage of wound exudates, and in prompting wound healing. Moreover, a VSD-assisted irrigation technique used in the treatment of a severe multiple-space infection in the oral and maxillofacial cervical regions showed favourable clinical effects with short treatment duration, lesser pain experience, and high clinical and therapeutic efficacy. \(^{28}\) Our results are consistent with the results of previous studies, indicating that the use of VSD can effectively reduce the time of wound cleaning, wound healing, and hospital stay for patients with soft-tissue wounds. However, clinically, it is worth noting that because of the complexity of the soft-tissue wound, such as the irregular shape and different depths of the wound, the application of VSD is prone to a situation where the material is not fully applied to the wound and the wound is difficult to seal. Furthermore, it was reported that the combination of an experimental vascular protective shield with VSD could reduce influences on systolic and diastolic capacities of the myocardium and avoid multiple compressions of exposed vessels, thus contributing to early vascularisation of wounds and wound repair. \(^{29}\)

In conclusion, VSD can improve the clinical symptoms of patients with soft tissue wounds effectively by improving bacterial clearance rate, granulation tissue coverage, and granulation tissue thickness, reducing
inflammation-related indicators, and shortening the time of wound cleaning, wound healing, and hospital stay. It should be acknowledged that there were still some limitations in this study, like the small sample size, being conducted in a single centre, and short research time. In the follow-up study, we will improve these limitations and obtain more evidence to support the conclusion of this experiment. Nevertheless, the VSD treatment method is worthy of promotion and implementation in the clinic.

DATA AVAILABILITY STATEMENT
The datasets generated and analyzed during the current study are available from the corresponding author on reasonable request.

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