Case studies: Treatment of chronic wounds with cultured skin substitutes

Cultured skin substitutes prepared from allogenic human skin cells and collagen substrates may provide an alternative treatment for acceleration of healing in selected kinds of chronic wounds.

Abstract

This report of four patients (2 male, 2 female) presents preliminary results of treatment of chronic skin wounds with cultured skin substitutes prepared from allogenic human skin cells and collagen substrates. The patients ranged in age from 64 to 73 years (mean: 67.25). The etiologies of their wounds were saphenous veinectomy, venous stasis, peripheral vascular disease, and venous stasis, with wound sizes 10cm², 20cm², 36cm² and 184cm². After the wounds were debrided, the cultured grafts, overlaid with a porous, non-adherent dressing, were stapled to the wounds. The procedure included frequent irrigations with essential nutrients containing broad spectrum antimicrobials, followed by dressing with antimicrobial ointments, and periodic examination of the wounds. Complete healing was obtained in two patients, partial healing in one with gradual recurrence due to poor patient compliance, and no healing in one. The two healed wounds did not recur after more than one year of observation and achieved closure in shorter periods of time than their duration before treatment with the skin substitutes. These results suggest that grafts of cultured allogeneic skin cells and collagen may accelerate healing of qualified chronic wounds. Qualifications include, but are not limited to, sufficient perfusion of the affected site and adequate debridement to viable tissue.

Introduction

Chronic ulcerations of the skin are responsible for a significant portion of health care expenditures. It was estimated in the U.S. in 1979 that 128,000 patients were hospitalized primarily for stasis ulceration. The average length of stay was 11.4 days with an estimated cost of 300 million dollars.¹ A survey by Meehan² of 148
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Figure 1. Diagram of normal healing in skin. Wound repair proceeds by an orderly and timely process of hemostasis, inflammation, cell proliferation, development of fibrovascular tissue, epithelial closure, and resolution and remodeling of healed skin. Cosmetically-acceptable scar forms in healed wounds.

Figure 2. Diagram of abnormal healing in skin. Chronic wounds are established by failures in inflammation, proliferation, or epithelial closure with superimposition of microbial contamination. Keloid or hypertrophic scarring result from failure of the healing process to resolve. Continuation of a cycle of inflammation and proliferation produces an excessive scar.

hospitals reports a prevalence of pressure ulcers of 9.2 percent. A treatment that could safely and efficaciously reduce the healing time of chronic skin ulcers would be of great benefit to the patient and community.

Mechanisms of normal healing in skin wounds follow a timely and orderly sequence of hemostasis, inflammation, cell proliferation, development of fibro-vascular tissue, epithelial closure, resolution and remodeling of repaired tissue. (Figure 1) In abnormal healing of skin, vascular insufficiencies or autoimmune conditions may cause failure of inflammation and cell proliferation which is compounded by microbial contamination to establish chronic wounds. (Figure 2) Excessive healing that produces keloid and hypertrophic scarring is also abnormal, and results from failure of the inflammatory and proliferative phases of wound healing to resolve.

Successful transplantation of cultured epithelial autografts was first reported in 1981. Similarly, cultured epithelial allografts have been demonstrated to reduce pain and to accelerate healing of chronic cutaneous wounds.

Cultured epidermal keratinocytes may be combined with collagen-based substrates as a vehicle for transplantation and to promote repair of connective tissue. However, cultured keratinocytes used alone are associated with greater fragility and bullae formation than the combination of cells with a collagen implant.

This report presents preliminary results of treatment of chronic skin wounds with cultured skin substitutes. The skin substitutes consist of collagen-glycosaminoglycan (GAG) sponges populated with cultured epidermal keratinocytes and dermal fibroblasts from allogeneic donors. Complete healing was obtained in two of four patients, partial healing in one, and no healing in one. Patients with complete healing were able to resume normal activities, and healed wounds did not recur after more than one year of observation. These results
Figure 3. Histology of cultured skin substitute (CSS). CSS consists of a collagen-glycosaminoglycan sponge inoculated with human epidermal keratinocytes and dermal fibroblasts. CSS resembles split-thickness skin graft in total thickness and in separation of epithelial and connective tissues into distinct compartments. Scale bar = 0.1 mm.

suggest that cultured skin substitutes prepared from allogeneic human skin may provide an alternative treatment for acceleration of healing in selected kinds of chronic wounds.

Materials and methods

Cultured skin substitutes (CSS) were prepared from cultured skin cells and collagen-GAG substrates as previously described. Dermal substitutes were prepared in vitro from bovine collagen and chondroitin-6-sulfate by freeze-drying and sterilized gamma-irradiation. Separate cultures of human epidermal keratinocytes and dermal fibroblasts were prepared by isolation from human cadaveric skin obtained from the Ohio Valley Tissue and Skin Bank after confirmation of negative tests for transmissible pathogens.

Patients were enrolled by informed consent into a protocol approved by the Institutional Review Boards of the University of Cincinnati and the Jewish Hospitals of Cincinnati. All wounds treated were on the distal lower extremity. Medical histories were obtained from subjects, and examinations were performed for evaluation of vascular competence. A summary of patient information is presented in Table 1. Wound areas ranged from 10 to 184 cm². Etiologies of wounds were peripheral vascular disease (ML), venous stasis (SM and JT), and dehiscence after saphenous veinectomy for coronary artery bypass surgery (JY). Patients were examined for perfusion of the affected leg by detection of dorsalis pedis pulse. Two patients (ML and SM) in whom no pulse was detected were further evaluated by Ankle/Brachial Pressure Ratios (A/BPR), and imaging of the affected leg by Impedance Plethysmography (IP) as described in Table 1. Duration of the wounds ranged from three to twelve months. Patient JY received one application of CSS, patients ML and JT received two applications, and patient SM received three applications.

Chronic wounds were debrided to viable subcutaneous tissue. CSS were overlaid with a porous, non-adherent dressing*. Cultured grafts were removed from dishes and transferred with the porous non-adherent dressing* as a support and stapled to the wound. Grafts were dressed with cotton gauze and transparent film** and CSS were irrigated with essential nutrients containing broad spectrum antimicrobials[5,11] four times daily for seven days. On day seven, all dressings and staples were removed, and grafts were exposed to air for 30 to 60 minutes to dry the graft surface.

Between days seven and fourteen, wounds were dressed with a non-adherent dressing containing equal parts of three ointments*. Grafted wounds were examined at two to three days and one week post surgery, and periodic intervals thereafter.

Results

Figure 3 shows the histologic appearance of cultured skin substitutes as used in this study. The porous side of the collagen-GAG sponge is populated with cultured human fibroblasts, and the non–porous side is covered with a stratified culture of epidermal keratinocytes. The total thickness of the cell-biopolymer implant is less than 0.5 mm, and epidermal and dermal components are organized similarly to a split-thickness skin graft.

Table 1 summarizes outcomes after treatment with CSS of three venous stasis and one diabetic leg ulcers. Patient ML healed approximately 85 percent of the wound area within five weeks of the first application of CSS. A second application of CSS to the remaining open area promoted complete closure of the wound by three months after grafting. Figures 4A through 4D show clinical photographs of patient ML. At one week (Figure 4B), a translucent epithelium covers the wound. Epithelial coverage of the wound is accomplished over the majority of the wound surface within three weeks after application. (Figure 4C) In addition to epithelial closure of the wound, the patient reported reduced pain that allowed resumption of normal ambulation. Cosmetic appearance of the healed wound was very
## Table 1. Patient Information and Outcomes

<table>
<thead>
<tr>
<th>Patient ID</th>
<th>Gender, Age</th>
<th>Wound Area (cm²)</th>
<th>Etiology</th>
<th>Associated Disease</th>
<th>Dorsalis Pedis Pulse</th>
<th>A/BPR</th>
<th>IP</th>
<th>Wound Duration</th>
<th>Treatments</th>
<th>Outcomes</th>
<th>Recurrence</th>
</tr>
</thead>
<tbody>
<tr>
<td>ML</td>
<td>Female, 73</td>
<td>36</td>
<td>Peripheral vascular disease</td>
<td>Diabetes</td>
<td>No*</td>
<td>Rt, 0.84; Lt, 0.60</td>
<td>Normal</td>
<td>4 months</td>
<td>Debridements CSS X 2</td>
<td>85% healed at 4 weeks after CSS #1 applied</td>
<td>Healed completely at 8 months after CSS #2 applied</td>
</tr>
<tr>
<td>SM</td>
<td>Male, 64</td>
<td>184</td>
<td>Venous stasis</td>
<td>Post phlebitic syndrome</td>
<td>No*</td>
<td>Rt, 0.79; Lt, 1.14</td>
<td>Bilateral venous thrombosis</td>
<td>12 months</td>
<td>Debridements CSS X 3, HBO₂</td>
<td>10% healed at 1 week</td>
<td>3 applications of CSS failed. Amputation after 3 months</td>
</tr>
<tr>
<td>JY</td>
<td>Male, 65</td>
<td>10</td>
<td>Saphenous veinectomy</td>
<td>Diabetes</td>
<td>Yes</td>
<td>N/A</td>
<td>N/A</td>
<td>3 months</td>
<td>Debridement CSS X 1</td>
<td>60% healed at 1 week</td>
<td>Healed completely at 4 weeks</td>
</tr>
<tr>
<td>JT</td>
<td>Female, 67</td>
<td>20</td>
<td>Venous stasis</td>
<td>Post phlebitic syndrome</td>
<td>Yes</td>
<td>N/A</td>
<td>N/A</td>
<td>12 months</td>
<td>Debridements CSS X 2, HBO₂</td>
<td>50% healed at 4 weeks after CSS #2 applied</td>
<td>Gradual recurrence due to poor patient compliance</td>
</tr>
</tbody>
</table>

* Ankle/Brachial Pressure Ratios (A/BPR) and Impedance Plethysmography (IP) were performed only on patients in whom no Dorsalis Pedis Pulse was detected. A/BPR values in boldface type indicate treated side.

HBO₂ – Hyperbaric Oxygen

CSS – Cultured Skin Substitute

acceptable by eight weeks after grafting. (Figure 4D)

SM received three applications of CSS, and all failed. Each set of CSS had initial adherence and epithelial survival during irrigations with nutrients and antimicrobials. After irrigations were discontinued on day seven, CSS degenerated rapidly with no healing of the wound by one month. After failure of all conventional and experimental alternatives for closure of this large wound, SM elected below-the-knee amputation.

JY received one application of CSS to a non-healing, dehiscent wound subsequent removal of the saphenous vein for cardiac bypass. Complete healing was accomplished within four weeks, with no recurrence after one year.

JT received two applications and had approximately 50 percent healing within four weeks after the second appli-
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Figure 4. Clinical photographs showing closure of a chronic diabetic ulcer with CSS. A) Chronic wound persisting for four months before treatment. B) One week after grafting with CSS shows development of a thin, translucent epithelium over the wound surface. C) By three weeks after grafting, stable epithelium has restored the protective barrier function of epidermis over > 80 percent of the wound surface. D) After eight weeks, the wound is fully healed, and the patient can ambulate freely without pain.

cation. Gradual recurrence of the treated wound was associated with poor compliance with wound care protocols and continued use of tobacco and alcohol.

Discussion

Data presented here suggest that grafts of cultured allogeneic skin cells and biopolymers may accelerate healing of qualified chronic wounds. Qualifications of wounds include, but are not limited to, sufficient perfusion of the affected site and adequate debridement to viable tissue. Additional qualifications of patient behavior are equally important and include compliance with wound care protocols, adequate nutrition, and management of associated disease. Two patients treated in this pilot study achieved wound closure in a shorter period of time than the duration of their wounds before treatment with CSS. This result suggests that CSS facilitated wound closure and may accelerate healing. However, determination of accelerated healing requires comparative investigation in a larger population of patients.

Closure of chronic wounds of large surface area within relatively short periods of time after grafting, e.g., one month, strongly suggests that transplanted allogeneic cells engraft directly. However, long term closure of chronic wounds is believed to result from replacement of allogeneic cells by autologous keratinocytes. Within three to four months after grafting, no allogeneic keratinocytes have been found, but wounds remain

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healed. This consistent finding in grafting of allogeneic keratinocytes suggests gradual replacement of allogeneic cells by circumferential ingrowth of autologous cells from the wound perimeter. This mechanism is consistent with the biological process termed "creeping substitution" in which parenchymal cells from the host repopulate the graft site, but allogeneic cells are not destroyed by T-cell cytotoxic action of the recipient. Reduced immunogenicity of cultured skin substitutes containing allogeneic keratinocytes is believed to result from depletion of donor leukocytes during culture of keratinocytes for transplantation.

Development of cell therapies will depend on culture and banking of allogeneic skin cells. Although use of cultured cells for transplantation is not regulated presently by FDA, new guidelines for use of allogeneic skin tissue also apply to cells isolated from these tissues. As with other experimental therapies, safety and efficacy of prospective therapies will be required. For cells cultured from allogeneic tissues, requirements for safety will include negative tests for viral and microbial pathogens, and no detectable disease after autopsy. Demonstration of efficacy requires comparative assessments of experimental therapies with prevailing standards of care to determine equivalence or superiority. It may be expected that continued preclinical and clinical studies with allogeneic cells will lead to improvements in care of chronic wounds. Establishment of standards and guidelines for allogeneic cells may be expected to serve as a foundation for transplantation of genetically modified cells for targeted therapy of wounds or metabolic diseases. With these goals ahead, the findings of this study provide an important demonstration that advanced therapies with allogeneic cells will provide new alternatives to healing of chronic skin wounds.

References


* N-Terface™; Winfield Laboratories, Richardson, TX
** OpSite™, Smith & Nephew United, Largo, FL
† Adaptic™, Johnson & Johnson Medical, Inc., Arlington, TX
‡ Neosporin™, E. Fougera & Co., Melville, NY
§ Bactroban™, Smith Kline–Beechum, Philadelphia, PA
∥ Nystatin™, Schein Pharmaceuticals, Inc., Brewster, NY