Molecular Analysis of Aggressive Microdermabrasion in Photoaged Skin

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Objective: To investigate dermal remodeling effects of crystal-free microdermabrasion on photodamaged skin.

Design: Biochemical analyses of human skin biopsy specimens following microdermabrasion treatment in vivo.

Setting: Academic referral center.

Participants: Volunteer sample of 40 adults, aged 50 to 83 years, with clinically photodamaged forearms.

Intervention: Focal microdermabrasion treatment with diamond-studded handpieces of varying abrasiveness on photodamaged forearms and serial biopsies at baseline and various times after treatment.

Main Outcome Measures: Quantitative polymerase chain reaction, immunohistochemistry, and enzyme-linked immunosorbent assay were used to quantify changes in inflammatory, proliferative, and remodeling effectors of normal wound healing. Type I and type III procollagen served as the main outcome marker of dermal remodeling.

Results: Coarse-grit microdermabrasion induces a wound healing response characterized by rapid increase in induction of cytokeratin 16 and activation of the AP-1 transcription factor in the epidermis. Early inflammation was demonstrated by induction of inflammatory cytokines, antimicrobial peptides, and neutrophil infiltration in the dermis. AP-1 activation was followed by matrix metalloproteinase–mediated degradation of extracellular matrix. Consistent with this wound-healing response, we observed significant remodeling of the dermal component of the skin, highlighted by induction of type I and type III procollagen and by induction of collagen production enhancers heat shock protein 47 and prolyl 4-hydroxylase. Dermal remodeling was not achieved when microdermabrasion was performed using a medium-grit handpiece.

Conclusions: Microdermabrasion using a coarse diamond-studded handpiece induces a dermal remodeling cascade similar to that seen in incisional wound healing. Optimization of these molecular effects is likely the result of more aggressive treatment with a more abrasive handpiece.

Trial Registration: clinicaltrials.gov Identifier: NCT00111254

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MICRODERMABRASION IS A popular procedure for skin rejuvenation. It has been suggested that microdermabrasion can improve the appearance of wrinkles, atrophic acne scars, dyspigmentation, and other signs of aging skin. The proposed mechanism by which microdermabrasion exerts these effects is through remodeling the dermis with elaboration of new collagen and other matrix components.

Karimipour et al demonstrated that aluminum oxide microdermabrasion activates a dermal remodeling cascade involving cytokines, transcription factors, and matrix metalloproteinases (MMPs). However, stimulation of new collagen synthesis occurred in only a few subjects and was not statistically significant. A possible explanation for the lack of significant collagen production might relate to the minimal wounding of microdermabrasion. In fact, aggressive ablative resurfacing procedures characteristically result in significant collagen production.

Hence, the objective of this study was to investigate whether microdermabrasion could be improved through more aggressive (but still nonablative) perturbation of the epidermis. We hypothesized that increasing the wounding stimulus might enhance dermal remodeling, as observed with other aggressive procedures, and thereby elicit consistent induction of collagen production. To test this hypothesis, we used a diamond-studded handpiece with varying roughness as a wounding stimulus.
The microdermabrasion system used (Diamond-Tome; Altair Instruments, Camarillo, California) offers a unique way to vary the abrasive stimulus. The system differs from standard microdermabrasion systems that use aluminum oxide crystals as corundum. The system uses a handpiece that has diamond fragments embedded in the contact point of the wand with the skin. The wand’s roughness is controlled by the size of the diamond particles at the contact point.

We assessed the ability of medium-grit and coarse-grit handpieces to elicit molecular responses that occur during normal wound healing. Collagen production was quantified.

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**METHODS**

**HUMAN SUBJECT DESCRIPTION, TREATMENT, AND TISSUE PROCUREMENT**

This study was approved by the University of Michigan Medical School Institutional Review Board for Human Subjects Research. All subjects provided written informed consent. Forty subjects (26 males and 14 females), aged 50 to 83 years, received a single microdermabrasion treatment with the diamond-studded handpiece to photodamaged forearm skin using a medium-grit (100-µm particle size) or coarse-grit (125-µm particle size) wand. In each subject, three 2 × 2-cm areas of photodamaged forearm skin were treated with microdermabrasion. The microdermabrasion device was set at ~25 inches mercury, and the microdermabrasion wand was applied in horizontal, vertical, and oblique directions for a total of 3 passes. For this study, 3 passes represents 1 microdermabrasion treatment. Each pass was characterized as a back-and-forth motion lasting approximately 15 seconds to "paint" the test area in sequential rows; the handpiece was not lifted off the skin during each pass. This procedure is similar to how we perform microdermabrasion on a daily basis in our clinic. The treatment resulted in mild erythema in all subjects that would typically last less than 2 hours.

Punch biopsy specimens (4 mm) were obtained from treated and untreated (control) skin at different intervals ranging from 4 hours to 14 days after treatment. Each subject’s baseline (no treatment) biopsy specimen served as his or her control. Immediately after biopsy, skin specimens were embedded in optimal cutting temperature (OCT) embedding medium (Tissue-Tek OCT compound; Miles, Naperville, Illinois), frozen in liquid nitrogen, and stored at −80°C until processing.

RNA extraction from skin biopsy specimens, reverse transcription, and messenger RNA (mRNA) quantification by real-time reverse transcription–polymerase chain reaction were performed as previously described. Custom primers and probes were used for type I procollagen (COL1A1) (GenBank Z74615), type III procollagen (COL3A1) (GenBank X15332), and the housekeeping gene 36B4 (GenBank AB007187). All other primer-probe sets were validated gene expression assays (TaqMan; Applied Biosystems, Foster City, California). Results are presented as fold change in treated vs untreated skin samples (normalized to transcript levels of 36B4) or as fold change vs 36B4 (2^ΔΔCt), where Ct indicates cycle threshold and is the end point of the real-time polymerase chain reaction.

**IMMUNOHISTOCHEMISTRY**

Frozen tissue embedded in OCT medium was cut into 7-µm-thick sections. Immunohistochemical staining was performed using the following primary antibodies: cytokeratin 16 (Novocastra; Leica Microsystems, Bannockburn, Illinois), cjun (Abtransduction Laboratories, Lexington, Kentucky), JunB (Santa Cruz Biotechnology, Santa Cruz, California), MMP type I (MMP1), prolyl 4-hydroxylase, type I procollagen (Chemicon, Temecula, California), neutrophil elastase (DAKO, Carpinteria, California), heat shock protein 47 (HSP47; BioGenex, San Ramone, California), and fibroblast activation protein (Bender MedSystems, Burlingame, California). Tissue-bound primary antibody was visualized with a secondary antibody–peroxidase complex, and the amount of staining was quantified using commercially available software (Image-Pro; Media Cybernetics, Inc, Bethesda, Maryland). To assess the specificity of staining, substitution of isotype γ-globulin for the primary antibody was used. There was no staining visualized with any isotype γ-globulin.

**PROTEIN EXTRACTION AND TYPE I PROCOLLAGEN ENZYME-LINKED IMMUNOSORBENT ASSAY**

Frozen sections (50 µm thick) were collected from OCT-embedded skin samples onto polyethylene naphthalene foil-coated slides (Leica Microsystems). For each sample, the dermal area was measured using software and associated with the laser capture microdissection microscope (Leica ASLMD), and skin sections were isolated from surrounding OCT by microdissection. Samples were collected into ice-cold protein extraction buffer (50mM Tris hydrochloride [pH 7.4], 0.15M sodium chloride, 1% Triton X-100, and protease inhibitors [Complete Mini, Roche Diagnostics, Indianapolis, Indiana]). Extraction products were centrifuged for 5 minutes at 10 000 g at 4°C, and supernatants were assayed for type I procollagen using an enzyme-linked immunosorbent assay kit (Panvera, Madison, Wisconsin). Type I procollagen protein concentrations were normalized to sample volumes.

**STATISTICAL ANALYSIS**

Changes in biomarkers over time compared with baseline levels were statistically evaluated using repeated-measures analysis of variance. Dunnett multiple comparisons procedure was used to test the significance of specific comparisons. The type 1 error rate was set at .05 for a 2-tailed hypothesis. Descriptive statistics included means, ranges, and standard errors. These data were analyzed using commercially available statistical software (SAS; SAS Institute, Inc, Cary, North Carolina).

**RESULTS**

We demonstrate that aggressive nonablative microdermabrasion is an effective procedure to stimulate collagen production in human skin in vivo. The beneficial molecular responses, with minimal downtime, suggest that aggressive microdermabrasion may be a useful procedure to stimulate remodeling and to improve the appearance of aged human skin.

**COARSE-GRIT MICRODERMABRASION INDUCES EARLY EPIDERMAL INJURY RESPONSE**

Induction of cytokeratin 16 by interfollicular keratinocytes is a well-characterized response to epidermal injury. A single microdermabrasion treatment with the
Coarse-grit microdermabrasion induces cytokeratin 16 (CK16) messenger RNA (mRNA) and protein expression in human skin in vivo. Photodamaged forearm skin was treated with coarse-grit microdermabrasion. Samples were obtained at the indicated times from untreated and treated skin sites. Connections were established between cytokeratin 16 gene expression and IL-8 gene expression. Positive control: Interleukin 1β messenger RNA (mRNA) levels were quantified by real-time reverse transcription–polymerase chain reaction (n=20). Interleukin 1β protein expression was maximal 64-fold 6 hours after a single treatment (Figure 3A and B). Interleukin 1β is a potent neutrophil chemoattractant that is involved in wound healing. Interleukin 8 is one such cytokine and is a potent neutrophil chemoattractant. A single microdermabrasion treatment using the coarse-grit handpiece resulted in statistically significant increases in IL-8 gene expression (GenBank NM000584.2). Coarse-grit microdermabrasion treatment using the coarse-grit handpiece resulted in statistically significant increases in IL-8 gene expression (GenBank NM000584.2). Interleukin 8 expression was maximal at 6 hours after a single treatment (P<.05) (Figure 4). One of the early inflammatory cellular events following epidermal injury is neutrophil infiltration into the wound site. Neutrophil elastase was used as a marker for infiltrating neutrophils. Neutrophil elastase protein expression was not detectable at baseline but was readily detectable 6 and 24 hours after a single treatment (Figure 5). Neutrophil elastase expression was induced throughout the reticular dermis. Increased neutrophil infiltration into the skin is consistent with observed elevations in IL-8 level.
COARSE-GRIT MICRODERMABRASION INDUCES ANTIMICROBIAL PEPTIDES

Antimicrobial peptides constitute a large diverse group of small-molecular-weight proteins that function in host defense against infection by directly killing microorganisms and by modulating innate and adaptive immunity.16 Recent studies17,18 have demonstrated altered expression of antimicrobial peptides in inflammatory skin diseases and following skin injury. A single treatment with coarse-grit microdermabrasion stimulated gene expression of antimicrobial peptides human defensin α1 (DEFA1) (GenBank NM004084.2), human β-defensin 2 (HBD2) (GenBank NM018661.3), and human β-defensin 3 (HBD3) (GenBank NM004942.2), and human defensin -defensin 2 (HBD2) (GenBank NM002422.3) increased 33-fold 6 hours after treatment (Figure 6A). Defensin α1 mRNA was near the limit of detection in untreated skin and was increased approximately 33-fold 6 hours after treatment (P < .05) (Figure 6A). Defensin α1 mRNA levels remained elevated 19-fold for at least 24 hours (P < .05). HBD2 and HBD3 mRNA expression was also rapidly induced following a single coarse-grit microdermabrasion treatment. HBD2 (Figure 6B) and HBD3 (Figure 6C) mRNA levels were elevated 15-fold 6 hours after treatment (P < .05); gene expression of both antimicrobial peptides remained elevated 24 hours after treatment (76- and 15-fold vs untreated skin, respectively; P < .05).

COARSE-GRIT MICRODERMABRASION INDUCES DERMAL REMODELING MMPs AND FIBROBLAST ACTIVATION PROTEIN

Matrix metalloproteinases break down structural proteins that comprise the dermal extracellular matrix (ECM) and are critical for dermal remodeling during wound healing.19-21 We examined 3 MMPs that are known to be inducible in human skin. Interstitial collagenase (MMP1) initiates collagen degradation by generating 2 smaller fragments, which are then further degraded by stromelysin 1 (MMP3) and gelatinase B (MMP9). A single coarse-grit microdermabrasion treatment resulted in 333-fold induction of MMP1 mRNA (GenBank NM002421.2) at 6 hours and 99-fold induction at 24 hours (P < .05) before trending toward baseline values (Figure 7A). MMP1 protein induction was localized in the papillary dermis (Figure 7A). Similarly, MMP3 gene expression (GenBank NM002422.3) increased 345-fold 6 hours and 39-fold 24 hours after a single treatment (P < .05) (Figure 7B). MMP9
gene expression (GenBank NM004994.2) followed a similar course, with 27-fold induction 6 hours after treatment ($P < .05$), dropping close to baseline within 24 hours after treatment (Figure 7C).

Fibroblast activation protein (FAP) is a membrane-bound serine protease that can degrade denatured collagen fragments.\textsuperscript{22,23} It is involved in matrix remodeling and cell motility. A single microdermabrasion treatment with the coarse-grit handpiece dramatically induced FAP protein expression 6 hours (3.9-fold) and 24 hours (4.6-fold) after treatment (Figure 8).

**COARSE-GRIT MICRODERMABRASION INDUCES COLLAGEN BIOSYNTHETIC PATHWAYS**

Microdermabrasion with the coarse-grit wand induced collagen biosynthetic pathways. HSP47 is a molecular chaperone protein necessary for intracellular transport and processing of procollagen within dermal fibroblasts.\textsuperscript{24} Microdermabrasion with the coarse-grit wand resulted in significant increases in HSP47 protein expression. HSP47 protein staining was increased 7.5-fold ($P < .05$) throughout the dermis (Figure 9A) 14 days after microdermabrasion treatment with the coarse-grit wand. Prolyl 4-hydroxylase catalyzes hydroxylation of proline residues within procollagen. Hydroxylation of proline is necessary to stabilize the triple helix structure of procollagen.\textsuperscript{25} Prolyl 4-hydroxylase protein expression was near the limit of detection at baseline but was readily detectable throughout the reticular and papillary dermis 14 days after treatment (Figure 9B).

Consistent with elevated expression of HSP47 and prolyl 4-hydroxylase, coarse-grit microdermabrasion induced type I and type III procollagen expression. Type I and type III procollagen transcripts were induced 3.2-fold and 2.6-fold, respectively, 14 days after a single microdermabrasion treatment ($P < .05$) (Figure 10A and B). Type I procollagen protein expression was induced 3.7-fold 14 days after treatment as measured by enzyme-linked immunosorbent assay ($P < .01$) (Figure 10C). Type I procollagen production was induced throughout the papillary and deeper dermis 14 days after treatment (Figure 10D).

**MEDIUM-GRIT MICRODERMABRASION FAILS TO STIMULATE REPAIR RESPONSES**

Microdermabrasion with the medium-grit handpiece did not result in statistically significant changes in any of the evaluated molecular components of the wound response or dermal remodeling cascade. In untreated and medium-grit microdermabrasion–treated forearm skin, the transcript level or protein expression was quantified at the various times after treatment for the following: MMP1, MMP3, MMP9, cytokeratin 16, type I and type III procollagen, and cytokines IL-1$\beta$ and tumor necrosis factor. Medium-grit microdermabrasion did not result in protein production or significantly alter mRNA levels in any of these genes ($n = 20$) (data not shown).
As the population ages, skin rejuvenation has become an area of significant interest. Patients prefer cosmetic procedures that require minimal disruption of their normal lifestyle.26 Microdermabrasion is a popular method of superficial skin resurfacing that is used to treat various cosmetic ailments, including wrinkles, atrophic scars, and dyspigmentation.1-3 It is associated with minimal morbidity, making it an ideal procedure for patients who want treatment that does not require prolonged healing.27 Kari- minpour et al5 demonstrated that aluminum oxide microdermabrasion stimulates a dermal remodeling cascade involving AP-1, MMPs, and cytokines; however, the procedure failed to consistently induce collagen production. Type I collagen is the major structural protein in the dermis, accounting for approximately 90% of dermal mass.25 Fragmentation of collagen fibrils with concomitant impairment of structural integrity is a characteristic feature of photoaged and aged skin.30 In addition, stimulation of collagen production seems to be a prerequisite for effective treatments that objectively improve the wrinkled appearance of skin.7,30 Therefore, the failure of aluminum oxide microdermabrasion to reliably induce new collagen production suggests a minimal clinical effect. We investigated whether microdermabrasion can be improved to provide reliable remodeling with increased collagen production, while retaining the characteristic “minimal downtime.” We hypothesized that increasing the wounding stimulus might enhance activation of the dermal remodeling cascade and result in increased collagen generation.

Wound healing involves several overlapping phases.15 Microdermabrasion with aluminum oxide crystals or a diamond-tipped wand triggers molecular responses that are observed during wound healing and should be considered a form of superficial wound. In previous work, we suggested that minimal barrier disruption combined with physical movement of the skin by vacuum is likely responsible for generating a wound healing response.5,31

![Figure 9](image) Figure 9. Coarse-grit microdermabrasion induces expression of HSP47 and prolyl 4-hydroxylase proteins involved in collagen synthesis in human skin in vivo. Photodamaged forearm skin was treated with coarse-grit microdermabrasion, and immunohistochemistry was performed for HSP47 (n=20) (A) and prolyl 4-hydroxylase (n=10) (B). Insets represent original magnification x120.

![Figure 10](image) Figure 10. Coarse-grit microdermabrasion induces type I and type III procollagen expression in human skin in vivo. Photodamaged forearm skin was treated with coarse-grit microdermabrasion. Samples were obtained at the indicated times from untreated and treated skin sites. A, Type I procollagen messenger RNA (mRNA) levels were quantified by real-time reverse transcription–polymerase chain reaction. *P<.005 vs untreated control. B, Type III procollagen mRNA levels were quantified by real-time reverse transcription–polymerase chain reaction. *P<.01 vs untreated control. C, Type I procollagen protein levels were determined by enzyme-linked immunosorbent assay. *P<.01. D, Type I procollagen protein expression was localized by immunohistochemistry. Images are representative of 20 subjects. Insets represent original magnification x120.
In our study, medium-grit microdermabrasion did not induce cytokeratin 16 expression nor did it induce other components of the repair pathways. Subjects with the largest increases in cytokeratin 16 expression also demonstrated a trend toward some of the largest increases in type I collagen expression.

Cytokines may serve as initiators of inflammation because they elaborate proinflammatory cytokines under many different conditions. Nickoloff and Naidu demonstrated that injury of the epidermis by tape stripping resulted in cytokeratin 16 gene expression and the induction of several epidermal matrix metalloproteinases (MMPs) into the dermis, with resultant dermal remodeling. COLOase indicates collagenase; GELase, gelatinase B; IL, interleukin; Strom, stromelysin 1; and PMN, polymorphonuclear leukocyte.

Figure 11. Schematic of the wound-healing cascade induced by microdermabrasion using a coarse-grit diamond-studded handpiece. Stratum corneum is preserved, and there is generation of cytokeratin 16 (CK16), antimicrobial peptide (AMP), cytokines, and presumed leakage of epidermal matrix metalloproteinases (MMPs) into the dermis, with resultant dermal remodeling. COLOase indicates collagenase; GELase, gelatinase B; IL, interleukin; Strom, stromelysin 1; and PMN, polymorphonuclear leukocyte.

Handpiece.
Stimulation of the wound-healing response may be responsible for the beneficial effects noted after microdermabrasion. Orringer et al. demonstrated that the use of carbon dioxide (10600 nm) or Nd:YAG (1320 nm) lasers can also stimulate a remodeling response with collagen production. The molecular responses seen following coarse-grit microdermabrasion are quantitatively less than those seen with carbon dioxide resurfacing but are quantitatively far superior to the results of nonablative resurfacing lasers like the 1320-nm Nd:YAG and aluminum oxide microdermabrasion. Ablative carbon dioxide laser resurfacing clearly yields significant clinical photorejuvenation, while nonablative resurfacing lasers demonstrate less, if not questionable, efficacy.

To the extent that molecular changes can predict clinical outcome, aggressive coarse-grit microdermabrasion should elicit significant skin rejuvenation. Further studies will determine whether microdermabrasion, if performed aggressively, has the capacity to become a worthwhile resurfacing procedure that results in noticeable cosmetic improvement while minimizing patient morbidity and downtime.

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Author Contributions: Drs Karimipour, Rittie, Hammerberg, and Fisher had full access to all the data in the study and take responsibility for the integrity of the data and the accuracy of the data analysis. Study concept and design: Karimipour, Voorhees, Orringer, Sachs, and Fisher. Acquisition of data: Karimipour, Rittie, Hammerberg, Min, and Fisher. Analysis and interpretation of the data: Karimipour, Voorhees, Hamilton, and Fisher. Drafting of the manuscript: Karimipour, Hammerberg, and Fisher. Critical revision of the manuscript for important intellectual content: Rittie, Hammerberg, Min, Voorhees, Orringer, and Sachs. Statistical analysis: Hamilton. Obtained funding: Karimipour. Study Supervision: Karimipour, Voorhees, Orringer, Hamilton, and Fisher.

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