11β-Hydroxysteroid dehydrogenase blockade prevents age-induced skin structure and function defects

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Glucocorticoid (GC) excess adversely affects skin integrity, inducing thinning and impaired wound healing. Aged skin, particularly that which has been photo-protected, shares a similar phenotype. Previously, we demonstrated age-induced expression of the GC-activating enzyme 11β-hydroxysteroid dehydrogenase type 1 (11β-HSD1) in cultured human dermal fibroblasts (HDFs). Here, we determined 11β-HSD1 levels in human skin biopsies from young and older volunteers and examined the aged 11β-HSD1 KO mouse skin phenotype. 11β-HSD1 activity was elevated in aged human and mouse skin and in PE compared with donor-matched photo-protected human biopsies. Age-induced dermal atrophy with deranged collagen structural organization was prevented in 11β-HSD1 KO mice, which also exhibited increased collagen density. We found that treatment of HDFs with physiological concentrations of cortisol inhibited rate-limiting steps in collagen biosynthesis and processing. Furthermore, topical 11β-HSD1 inhibitor treatment accelerated healing of full-thickness mouse dorsal wounds, with improved healing also observed in aged 11β-HSD1 KO mice. These findings suggest that elevated 11β-HSD1 activity in aging skin leads to increased local GC generation, which may account for adverse changes occurring in the elderly, and 11β-HSD1 inhibitors may be useful in the treatment of age-associated impairments in dermal integrity and wound healing.

Introduction
Glucocorticoid (GC) excess, whether of endogenous (e.g., Cush- ing’s syndrome) or exogenous (topical or systemic therapy) origin, is associated with a diverse range of adverse side effects including adipose tissue redistribution, proteolysis, bone resorption, and hyperglycemia manifesting respectively as omental adiposity, muscle weakness, osteoporosis, and insulin resistance (1).

In skin, GC excess leads to marked atrophy with dermal and epidermal thinning according to both human (2–4) and rodent (5, 6) studies. GCs also induce a flattening of the normally undulating “rete ridges” at the dermal-epidermal junction (DEJ) (7). At a cellular level, dermal collagen content in human skin is decreased following topical and systemic GC treatment (3, 8, 9) and is similarly decreased in rats following subcutaneous dexamethasone injection (10). Subsequently, there is a profound increase in the transparency of skin with a tissue paper–like consistency, increased fragility, tearing and bruising (7), increased transepidermal water loss (TEWL) (11), poor wound healing (12, 13), and increased infection risk (14).

Many of these features are also shared with the phenotype of aging skin, including thinning and DEJ flattening (15), decreased dermal cellularity (16), reduced collagen content (17–20), and reduced dermal fibroblast proliferation and collagen secretion in cells from older donors (18). Consequently, aged skin also suffers from an impaired permeability barrier, with increased TEWL (21), altered mechanical properties (22), delayed wound healing (23, 24), and increased disease prevalence (25).

We hypothesize that with no appreciable changes in circulating cortisol concentrations with age, changes in tissue-specific prerreceptor regulation of local GC availability by 11β-hydroxysteroid dehydrogenase type 1 (11β-HSD1) may explain the phenotypic link between GC excess and aging skin. In intact cells, 11β-HSD1 functions exclusively as an NADPH-dependent oxoreductase, activating cortisol from cortisone. We recently characterized the expression of 11β-HSD1 in human and rodent skin, reporting increased 11β-HSD1 expression in primary human dermal fibroblasts (HDFs) from older donors (26). Increased expression was also detected in donor-matched photo-exposed (PE) versus photo-protected (PP) HDFs (26), suggesting that an increased capacity for local GC activation may represent a novel mediator of age-related changes in skin physiology and function common to both intrinsic and extrinsic aging.

Here, we investigated age-dependent changes in 11β-HSD1 human and murine skin tissue explants. Additionally, we examined alterations in dermal integrity, cellularity, and collagen content in murine skin from age-matched 11β-HSD1–null and WT littermates and report a striking reversal of age-induced dermal morphology in KO mice. We describe the GC-driven changes in collagen biosynthesis, modification, and processing gene expression that may underpin these in vivo observations. Finally, we
demonstrate the translational therapeutic potential of 11β-HSD1 blockade by reporting accelerated wound healing in young mice treated with a selective 11β-HSD1 inhibitor and in aged 11β-HSD1 KO mice relative to their WT littermates.

Results

11β-HSD1 expression increases in aging skin. 11β-HSD1 activity in human skin obtained from older versus younger donors increased by 42% and 26% in PP and PE biopsies, respectively (Figure 1A). Interestingly, activity in donor-matched PE versus PP samples from young (n = 20) and aged (n = 20) donors. (B) 11β-HSD1 activity (pmol/mg/h) was also greater in older (91–99 weeks; n = 5) versus younger (11–20 weeks; n = 10) mouse skin. *P < 0.05; **P < 0.01.

Figure 1
11β-HSD1 activity increases in aging skin ex vivo. (A) 11β-HSD1 activity (percentage of conversion of 100 nM cortisone to cortisol) was greater in aged (>60 years) versus young (20–30 years) human skin in both PP (n = 20) and PE (n = 20) biopsies. Activity was also greater in donor-matched PE versus PP samples from young (n = 20) and aged (n = 20) donors. (B) 11β-HSD1 activity (pmol/mg/h) was also greater in older (91–99 weeks; n = 5) versus younger (11–20 weeks; n = 10) mouse skin. *P < 0.05; **P < 0.01.

Furthermore, immunohistochemical studies identified increased 11β-HSD1 protein expression in dermal fibroblasts from PE relative to donor-matched PP skin sections, with increased expression also observed in the epidermal compartment (Figure 2, A and B). Analysis of 11β-HSD1 staining in PE sections
confirmed a 2.5-fold and 3.5-fold increase in dermal and epidermal expression, respectively (Figure 2C), and a 4-fold increase in endogenous epidermal melanin staining (internal control). We also examined the expression of hexose-6-phosphate dehydrogenase, the enzyme supplying NADPH cofactor for 11β-HSD1, and the GC receptor, but found them to be unaffected by donor age and site in human skin or by age and 11β-HSD1 KO in mice (data not shown).

Reversal of dermal atrophy in aged 11β-HSD1–null mice. Aged WT mouse skin displayed a loose collagen network with noticeable atrophy and a sponge-like appearance characterized by large, vacant interfibril spaces (Figure 3A). In some areas of tissue sections from aged mice, the collagen meshwork appeared shredded, with little or no structural integrity or organization. 11β-HSD1 activity in KO mouse skin was reduced to background levels, with negligible mRNA expression (Figure 3B). Strikingly, aged KO mice displayed a histological skin profile more comparable to that of the young WT littermates compared with the aged WT littermates, with improved dermal integrity and more orderly collagen organization (Figure 3C).

Collagen density as assessed with Masson trichrome staining was also improved in 11β-HSD1 KO (Figure 4B) compared with WT mice (Figure 4A). Quantification of collagen staining intensity revealed a 2-fold increase in the aged KO mice (Figure 4C).

GC regulation of collagen biosynthesis and processing gene expression. Our in vivo studies identified dermal collagen integrity and structural organization as important GC targets in aging skin. The regulation of collagen homeostasis (e.g., posttranslational modification, transport, and extracellular maturation) by physiological concentrations of endogenous GC had not been previously investigated. Cortisol treatment (100 nM) in HDFs (n = 5) downregulated mRNA expression of genes at various stages in the synthesis and processing of mature collagen (Table 1). These included prolyl (LEPREL1, LEPREL2, and P4HA2, with an 80%, 33%, and 36% decrease, respectively) and lysyl (PLOD2; 53% decrease) hydroxylases, which hydroxylate the proline and lysine residues on nascent collagen propeptides that facilitate helical trimeric procollagen assembly. We also noted a trend toward reduced expression of LEPREL1, P4HB, and PLOD1 hydroxylase mRNA.

The transport and secretion of newly formed procollagen helices are assisted by the collagen-specific chaperone HSP47 (SERPINH1), which also exhibited reduced expression follow-
ing GC treatment. Following secretion into the extracellular environment, the nonhelical procollagen "ends" are cleaved by membrane-bound collagen peptidases (ADAMTS2 and BMP1), which did not appear to be GC regulated. The resulting tropocollagen peptides are cross-linked by lysyl oxidases (LOX and LOXL1-4), which use lysine and hydroxylysine residues to form covalent bonds in the resultant collagen fibril. Interestingly, mRNA expression for all 5 lysyl oxidases was reduced by GC treatment, with decreases ranging from 31% to 53%. The stability of collagen fibrils is further supported by proteoglycans such as decorin (DCN), which displayed a 2.7-fold induction of mRNA expression following GC treatment. Collagen-degrading MMPs and their inhibitors were also GC regulated, with reduced expression of MMP1 (71%), MMP12 (44%), and TIMP3 (64%).

Table 1
GC-regulated target genes in cultured fibroblasts impair collagen biosynthesis and processing

<table>
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<tr>
<th>Gene symbol</th>
<th>Gene title</th>
<th>Function</th>
<th>Fold change</th>
<th>P value</th>
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<td><strong>Pre-propeptide expression</strong></td>
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<td>COL1A1</td>
<td>Collagen type 1, alpha 1</td>
<td>Primary matrix structural component</td>
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<td>0.078</td>
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<td>COL3A1</td>
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<td>Secondary matrix structural component</td>
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<td>PEPD</td>
<td>Peptidase D</td>
<td>Proline recycling</td>
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<td><strong>Procollagen assembly</strong></td>
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<td>LEPRE1</td>
<td>Leucine proline-enriched proteoglycan (leprecan) 1</td>
<td>Intrafibril stability</td>
<td>0.84</td>
<td>0.059</td>
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<td>↓ LEPREL1</td>
<td>Leprecan-like 1</td>
<td>Intrafibril stability</td>
<td>0.2</td>
<td>B</td>
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<td>↓ LEPREL2</td>
<td>Leprecan-like 2</td>
<td>Intrafibril stability</td>
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<td>Prolyl 4-hydroxylase, alpha polypeptide 1</td>
<td>Intrafibril stability</td>
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<td>P4HB</td>
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<td>PLOD1</td>
<td>Procollagen-lysine, 2-oxoglutarate 5-dioxygenase 1</td>
<td>Intrafibril stability</td>
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<td>0.081</td>
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<td>Procollagen-lysine, 2-oxoglutarate 5-dioxygenase 2</td>
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<td>A</td>
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<td>PLOD3</td>
<td>Procollagen-lysine, 2-oxoglutarate 5-dioxygenase 3</td>
<td>Intrafibril stability</td>
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<td><strong>Procollagen transport</strong></td>
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<td>↓ SERPINH1</td>
<td>Serpin peptidase inhibitor, clade H (heat shock protein 47), member 1, (collagen binding protein 1)</td>
<td>Collagen chaperone</td>
<td>0.81</td>
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<td><strong>Tropocollagen formation</strong></td>
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<td>ADAMTS2</td>
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<td>N-propeptide cleavage</td>
<td>0.96</td>
<td>0.58</td>
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<td>BMP1</td>
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<td>C-propeptide cleavage</td>
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<td>↓ LOX</td>
<td>Lysyl oxidase</td>
<td>Interfibril cross-linking</td>
<td>0.69</td>
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<td>↓ LOXL1</td>
<td>Lysyl oxidase-like 1</td>
<td>Interfibril cross-linking</td>
<td>0.47</td>
<td>B</td>
</tr>
<tr>
<td>↓ LOXL2</td>
<td>Lysyl oxidase-like 2</td>
<td>Interfibril cross-linking</td>
<td>0.58</td>
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<td>Lysyl oxidase-like 3</td>
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<td>Interfibril cross-linking</td>
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<td>↑ DCN</td>
<td>Decorin</td>
<td>Interfibril cross-linking</td>
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<td><strong>Collagen fibril remodeling</strong></td>
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<td>↓ MMP1</td>
<td>Matrix metalloprotease 1 (interstitial collagenase)</td>
<td>Cleaves collagen I, II, III VII and X</td>
<td>0.29</td>
<td>C</td>
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<td>Matrix metalloprotease 2 (gelatinase A, 72 kDa gelatinase, 72 kDa type IV collagenase)</td>
<td>Cleaves collagen IV</td>
<td>0.87</td>
<td>0.2</td>
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<td>MMP3</td>
<td>Matrix metalloprotease 3 (stromelysin 1, progelatinase)</td>
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<td>MMP8</td>
<td>Matrix metalloprotease 8 (neutrophil collagenase)</td>
<td>Cleaves fibronectin, laminin, gelatin I, III, IV, and V, collagen III, IV, X, and IX and cartilage proteoglycans. Activates procollagenase.</td>
<td>0.76</td>
<td>0.14</td>
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<td>MMP10</td>
<td>Matrix metalloprotease 10 (stromelysin 2)</td>
<td>Cleaves fibronectin, gelatin of I, III, IV, and V and weakly collagens III, IV, and V</td>
<td>2.3</td>
<td>0.066</td>
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<td>↓ MMP12</td>
<td>Matrix metalloprotease 12 (macrophage elastase)</td>
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<td>↓ MMP13</td>
<td>Matrix metalloprotease 13 (collagenase 3)</td>
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<tr>
<td>TIMP1</td>
<td>TIMP metalloprotease inhibitor 1</td>
<td>Inhibits MMP1-3, 7-13 and 16</td>
<td>0.93</td>
<td>0.29</td>
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<td>TIMP2</td>
<td>TIMP metalloprotease inhibitor 2</td>
<td>Inhibits MMP1-3, 7-10, 13-16 and 19</td>
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<td>0.22</td>
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<td>↓ TIMP3</td>
<td>TIMP metalloprotease inhibitor 3</td>
<td>Inhibits MMP1-3, 7 and 9</td>
<td>0.36</td>
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<td>↓ TIMP4</td>
<td>TIMP metalloprotease inhibitor 4</td>
<td>Inhibits MMP1-3, 7, 9 and 13-16</td>
<td>4.61</td>
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Treatment of primary HDFs with 100 nM cortisol downregulated mRNA expression of key mediators of collagen fibril assembly, transport, extracellular cross-linking and remodeling (n = 5). *P < 0.05; **P < 0.01; ***P < 0.001.
and a 4.6-fold induction of TIMP4. We found that MMP8 and MMP10 also displayed a trend toward GC-mediated induction and reduction, respectively.

Age-induced changes in collagen biosynthesis gene expression are rescued in 11β-HSD1 KO mice. In mouse skin, collagen (Col1a1) expression was unaffected by age or 11β-HSD1 genotype (Figure 5D), but we found a 64% age-induced decrease in mouse skin prolyl hydroxylase (Leprel1) expression, which was fully restored in aged 11β-HSD1 KO mice to levels comparable to those of young animals (Figure 5A). Similarly, a greater than 90% decrease in lysyl hydroxylase (Plod1) expression in aged WT skin was also partially rescued in the KO littermates (Figure 5B). Timp4 expression was also decreased by approximately 80% in the aged WT mice, but not in the KO mice (Figure 5C). By contrast, the age-induced downregulation of Mmp2 and Mmp3 expression was unaltered in the KO animals (Figure 5, E and F).

Local blockade of endogenous GC activation accelerates wound healing in vivo. Body weight at each time point was indistinguishable between the vehicle- and inhibitor-treated groups (data not shown). Gross wound morphology revealed accelerated closure in 11β-HSD1 inhibitor–treated mice compared with vehicle-treated controls at each time point (Figure 6A). Indeed, accelerated re-epithelialization (ranging from 35% to greater than 100%) was observed in inhibitor-treated animals at each time point measured. For example, the area of neoepidermis at 14 days after wounding was 23% in the vehicle-treated mice, but increased to 50% following 11β-HSD1 blockade (P < 0.001; n = 12) (Figure 6B). Moreover, while vehicle-treated mice required 18 days to reach 40% re-epithelialization, 11β-HSD1 blockade reduced this time by over a week, attaining a similar level of recovery by day 9.

Importantly, we also observed faster wound closure in the aged 11β-HSD1 KO mice compared with their WT littermates (Figure 7A). 11β-HSD1 KO wound areas were reduced by approximately 50% at 4 days (P < 0.05, n = 6), while WT wound areas were still comparable to those on day 0, with a similar trend observed on day 2 (Figure 7B).

Discussion

Our previous studies reported an increase in 11β-HSD1 expression in HDFs obtained from older compared with younger donors and in PE cells relative to donor-matched PP cells in culture (26). Here, we provide new evidence that these in vitro observations translate into increased 11β-HSD1 expression and activity in aging skin both in humans and mice. Our data support a role for 11β-HSD1 in both intrinsic (e.g., PP skin) and “accelerated” extrinsic aging (e.g., PE skin); with increased 11β-HSD1 expression and activity in PE skin further exacerbating the age-induced increase in PP skin.

The increased 11β-HSD oxoreductase activity measured in our human cohort reflects the “net” effect of both 11β-HSD isozymes resulting from increased 11β-HSD1 expression and activity in aging skin both in humans and mice. Our data support a role for 11β-HSD1 in both intrinsic (e.g., PP skin) and “accelerated” extrinsic aging (e.g., PE skin); with increased 11β-HSD1 expression and activity in PE skin further exacerbating the age-induced increase in PP skin.

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isozyme are exclusively responsible. Immunohistochemical studies in human skin demonstrated 11β-HSD1 staining predominantly in PP epidermis, with increased staining in photo-aged keratinocytes in addition to induced expression in PE dermal fibroblasts. Although our subsequent studies focused primarily on the dermal consequences of age-induced increased 11β-HSD1 activity, our observation suggests that both cell types contribute to this increase. Further studies should elucidate whether impaired epidermal function in aged skin is also a consequence of increased GC activation.

To the best of our knowledge, our group was the first to report an association between 11β-HSD1 and aging using primary human osteoblasts (27). Subsequently, a growing body of evidence has suggested that the age-related increase in 11β-HSD1 occurs in a variety of cells and tissues, including lymphocytes (28), bone (29), brain (30), and adipose (31), implicating a role for 11β-HSD1 in systemic aging. In these tissues, increased local GC activation is in part responsible for the phenotypic consequences of aging skin, with aged 11β-HSD1 KO mice displaying a reversal of increased GC activation.

Reduced expression of prolyl hydroxylases by GC in skin was previously limited to studies in rats (40, 41). Recently, Vranka et al. characterized the phenotype of the prolyl 3-hydroxylase 1–null (LEPRE1-null) mouse which, in addition to bone and tendon abnormalities, displayed a less densely packed dermis, with fewer collagen fibrils in the skin of newborn KO mice and skin thinning in adults, with collagen fibril clumping and interfibril spaces (42). These findings are endorsed by mutations in LEPRE1 in humans, which cause a recessive metabolic bone disorder resembling lethal/severe osteogenesis imperfecta (43), with patients also exhibiting overmodification of type I procollagen chains in dermal fibroblast cultures from affected individuals (44).

Mutations in genes encoding lysyl hydroxylases are known to cause Ehlers-Danlos syndrome type VI, characterized by neonatal kyphoscoliosis, generalized joint laxity, skin fragility, and severe muscle hypotonia at birth (45), as well as Bruck syndrome (46), affecting bone collagen cross-linking (47), with overexpression associated with skin fibrosis (46). Our results show a decrease in the expression of all 5 known lysyl oxidase isoforms following GC treatment, in agreement with previous rodent studies evaluating the effect of local GC treatment (41). Decreased lysyl oxidase activity has been identified as a cause of Ehlers-Danlos syndrome variant type IX and the clinically distinct Menkes syndrome, both manifesting with skin laxity and hyperextensibility (48).

Figure 6
Accelerated wound healing in 11β-HSD1 inhibitor–treated mice. (A) Representative images showing improved wound closure in mice treated every 2–3 days with 1% RO151. (B) Accelerated re-epithelialization was recorded at each time point measured (n = 5–12). *P < 0.05; **P < 0.01; ***P < 0.001.

- HSD1 inhibitors (33, 34).
We also report GC-induced differential regulation of several enzymes involved in collagen turnover and remodeling, including downregulation of MMP1, the elastin-degrading MMP12 as previously reported (49, 50), and a similar trend for MMP10. Conversely, TIMP4, a potent inhibitor of the type IV collagenase MMP2 (51), was upregulated, whereas an opposing effect was observed for TIMP3. Although still poorly understood, altered expression of these highly complex mediators of collagen remodeling through increases in local GC availability are likely to play an important role in aging skin phenotype manifestation. Collectively, the identification of these critical regulators of collagen biosynthesis as novel GC targets in HDF raises the possibility that altered local GC availability could contribute to several aspects of altered collagen structural organization in aging skin.

Subsequently, we analyzed the expression of GC-regulated genes in our KO mouse model and showed a reversal of age-related decreases in Leprel1, Plod1 and Timp4 expression. Procollagen type 1 expression (Col1a1) was largely unaffected by age or genotype, suggesting that altered posttranslational processing of collagen remodeling through increases in local GC availability are likely to play an important role in aging skin phenotype manifestation. Collectively, the identification of these critical regulators of collagen biosynthesis as novel GC targets in HDF raises the possibility that altered local GC availability could contribute to several aspects of altered collagen structural organization in aging skin.

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Although the central theme of the current research focused on chronological aging, our finding demonstrating elevated 11β-HSD1 activity and expression in PE skin presents 11β-HSD1 as a candidate indicator of photo-aging and extends the scope of pharmacological intervention and/or treatment to photo-damaged skin. Finally, our studies may have implications in the context of slow-healing wounds occurring in diabetics or sufferers of chronic stress, where limiting local GC generation could also improve healing.

Methods

Materials were purchased from Sigma-Aldrich unless otherwise stated.

Tissue preparation and cell culture

Human tissue. A group of 40 healthy Caucasians consisting of 20 young (year ± SD, 25.7 ± 3.0; 9 female and 11 male) and 20 older (72.2 ± 8.2; 11 female and 9 male) donors underwent 3-mm biopsies of PE (lower outer) and PP (inner upper) arm skin. The skin was sterilized and covered with Supadrape (Westfield Medical Ltd.) containing a central aperture prior to local anesthetic injection with 2 ml of 2% lidocaine HCI (National Veterinary Services). Following dispersion, skin was tested for numbness before obtaining 2 biopsies per site using a Stiefel 3-mm Biopsy Punch (Medisave). Cotton gauze was used to contain bleeding before Inadine patch and Mepore dressing application (Oncall Medical Supplies). Tissue was stored in complete media (RPMI-1640, 10% FCS, 2 mM L-glutamine, 1% Pen-Strep, 1% sodium pyruvate, and 1% nonessential amino acids) and either analyzed immediately or snap-frozen and stored at –80°C.

These findings suggest that blockade of the age-related increase in 11β-HSD1 activity may promote improved structural and functional properties in aging skin. Selective 11β-HSD1 inhibitors have already been deployed by many pharmaceutical companies, largely on the premise that they may reverse or prevent obesity-related diabetes mellitus (53–57). Our collaborative research using a highly selective and effective 11β-HSD1 inhibitor, RO151, demonstrates accelerated re-epithelialization and wound healing and presents the exciting possibility that topical 11β-HSD1 inhibitors might be used to combat age-related impairments in wound healing. Our observation that aged 11β-HSD1 KO mice exhibit accelerated wound healing relative to WT littermates further supports this possibility.
One biopsy per site per human donor (4–10 mg) was analyzed for 11β-HSD1 (oxoreductase) activity. Mouse skin explants (~6 mg) were analyzed in duplicate. Intact tissue activity was measured in 10-ml glass tubes containing 0.5 ml of complete media, 100 pmol of cortisone or 11-dehydrocorticosterone, and kinetic amounts of [3H] cortisone to cortisol (24 hours). In these samples, activity was not normalized for tissue weight — which was increased in PE samples (Supplemental Figure 1A; supplemental material available online with this article; doi:10.1172/JCI64162DS1), owing to a deeper dermal layer composed primarily of inert structural proteins, e.g., collagen — as this did not correlate with enzyme activity in either group (Supplemental Figure 1B). Furthermore, punch biopsies standardized the same surface area between samples. For mouse samples, enzyme activity was normalized for tissue weight (pmol corticosterone/mg tissue/h) as surface area varied.

**Immunohistochemistry**

11β-HSD1 protein detection in frozen human skin sections (fixed in 3:1 acetone/ethanol) was analyzed by immunohistochemistry using a primary mAb (Abcam) and an alkaline phosphatase-conjugated goat anti-mouse secondary Ab (Dako). Fast Red TR/Naphthol AS-MX Tablets were used to visualize the staining according to the manufacturer’s guidelines. Briefly, slides were incubated for 15 minutes in 56:1 methanol/hydrogen peroxide to inactivate endogenous peroxidases, rinsed with deionized water, and blocked for 1 hour in TBSAT (10 ml 10x TBS [121.2 g Tris/160.4 g sodium chloride, pH 7.6, in 2 liters of deionized water], stored in aliquots at ~20°C), 3 g of BSA, 50 μl of Triton X100, and 10% goat serum. Slides were drained and incubated for 2 hours in fresh TBSAT with the primary Ab diluted 1:100 before washing under agitation in three 10-minute washes of TBST (~800 ml 1x TBS plus 0.05% Tween-20). Slides were drained and incubated for 30 minutes in fresh TBSAT with the secondary Ab diluted 1:200 before repeating washes. Following visualization, slides were air-dried before applying aqueous mounting medium and coverslips. Staining was observed in similar structures to those previously described using a polyclonal Ab (26) and was negligible when substituting equimolar concentrations of isotype control (mouse IgM; AbCam) for the primary Ab (Supplemental Figure 2).

Sections were photographed using a Nuance multispectral imaging system (CRi, Advanced Molecular Vision). Nuance image analysis software was used to annotate the DEJ and stratum corneum-granulosum interface (CRi, Advanced Molecular Vision). Staining intensity was measured in the epidermis and upper dermis.

**Histology**

Tissue was stored in 10% neutral buffered formalin (4 g sodium monophosphate, 6.5 g sodium diphosphate in 900 ml of distilled water, and 100 ml of 37% formaldehyde) before paraffin embedding and sectioning. Paired sections from each age and genotype were cut per slide. Following dewaxing and rehydration (26), the slides were stained with H&E, mounted, coverslipped, and analyzed in a blinded fashion by 2 independent researchers. Sections were stained for collagen using a Masson trichrome kit according to the manufacturer’s guidelines. Collagen stain intensity was determined using the Nuance system described above.

**RNA extraction and reverse transcription**

Total RNA was extracted from primary cultures of HDFs using the Tri-Reagent method (Sigma-Aldrich), with generation of cDNA by reverse transcription as previously described (26). Tissue RNA was extracted using an RNasy Fibrous Tissue kit (QIAGEN) using a PowerGen 125 homogenizer (Fisher Scientific) according to the manufacturer’s protocol.

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**Figure 8**

GC treatment impairs multiple elements of collagen biosynthesis and processing. GC treatment of primary HDFs decreased collagen transcription (1) and altered collagen metabolism (7) as previously shown. We found additional GC targets, including reduced expression of collagen posttranslational hydroxylase enzymes (3), collagen chaperone (4), and all lysyl oxidase enzyme isoforms required for collagen inter- and intrafibril cross-link formation (6).

**Mouse tissue.** Studies were conducted as previously described (26). Mouse skin from male young (11–20 weeks of age, n = 10) and aged (91–99 weeks of age, n = 5) WT mice and from littersmates with global deletion of 11β-HSD1 (generated in-house; n = 15) (58) was obtained by dissection of shaved skin from the lower dorsal region. 11β-HSD1 KO mice were backcrossed (greater than 9 generations) into C57BL6/J mice, with appropriate strain controls used.

**HDF culture and treatment.** HDFs were cultured from lower outer arm biopsies from Caucasian donors of variable age by incubation for 8 days at 37°C in a 5% CO2 atmosphere in T25 flasks containing 2 ml of complete media.

Upon attaining 80% confluence, cells were passaged at a ratio of 1:3 as required. For GC treatment, cells were seeded into 6-well plates at 70% confluence and incubated overnight. Cells were treated with 100 nM cortisol required. For GC treatment, cells were seeded into 6-well plates at 70% confluence and incubated overnight. Cells were treated with 100 nM cortisol required. For GC treatment, cells were seeded into 6-well plates at 70% confluence and incubated overnight. Cells were treated with 100 nM cortisol required. For GC treatment, cells were seeded into 6-well plates at 70% confluence and incubated overnight. Cells were treated with 100 nM cortisol required.
Real-time PCR gene expression analysis

HDF studies. The BioMark system (Fluidigm) was used to analyze the expression of 33 genes involved in collagen biosynthesis and processing. Specific target preamplification was conducted in 5-µl volumes consisting of a 2.5-µl TaqMan PreAmp Master Mix (2x; Applied Biosystems), a 1.25-µl pooled TaqMan gene expression assay (1 µl for each assay and 4 µl 1x TE buffer, 0.2x final assay concentration), and 1.25 µl (25–50 ng) of cDNA (or 1x TE for the negative control). Following brief vortexing and centrifugation, samples were heated to 95°C for 10 minutes followed by 14 cycles of 95°C for 15 seconds and 60°C for 4 minutes. Samples were then diluted 1.5 with 20 µl of 1x TE buffer. Following preamplification, 1 µl of each sample was also pooled and serially diluted in 1x TE buffer (1:10, 1:100, 1:1,000, and 1:10,000) for standard curve generation.

Samples (in triplicate) and standards (in duplicate) were prepared for loading onto the chip in 8-µl volumes in a 96-well plate: 4 µl of TaqMan Universal PCR Master Mix (2x); 0.4 µl of GE Sample Loading Reagent (20x, Fluidigm), and 3.6 µl of preamplified cDNA per standard. Plates were vortexed, centrifuged, and stored at 5°C until required. Gene expression assays were prepared in 8-µl volumes: 4 µl of TaqMan Gene Expression Assay (20x) and 4 µl of Assay Loading Reagent (2x; Fluidigm), giving final concentrations of 9 µM and 2.5 µM for the primers and probe per assay, respectively. Samples were vortexed, centrifuged and stored at 5°C. Chips were primed using the IFC controller according to the manufacturer’s protocol (Fluidigm), loaded with 5 µl of Sample Mix, 5 µl of Assay Mix, and activated using the IFC controller. Chips were processed on a BioMark quantitative PCR reader (Fluidigm). Following standard curve verification, ΔCt values were determined for each sample by normalizing to the geometric mean of 4 housekeeper genes (RPL13, TBP, B2M, and PPIA). Mouse tissue: Lepre1, Plod1, Timp4, Col1a1, Mmp2, and Mmp3 mRNA levels were measured (in duplicate) by conventional real-time PCR using an ABI 7500 system (PerkinElmer). PCR was performed in 10-µl reactions in 96-well plates. Reactions contained 5 µl of 2x TaqMan Universal PCR Mastermix, 0.5 µl of assay (900 nM primer and 250 nM probe final concentrations), 3.5 µl of distilled water, and 1 µl of cDNA (25–50 ng). Reactions were standardized to the housekeeper 18S (all Applied Biosystems). Reactions were as follows: 50°C for 2 minutes, 95°C for 10 minutes, 40 cycles of 95°C for 15 seconds, and 60°C for 1 minute.

Wound-healing studies

11β-HSD1 inhibitor studies. Inhibitor studies were conducted in 8-week-old male Swiss mice (n = 5–12) using RO151, known to selectively inhibit 11β-HSD1 activity in vivo (59), with an IC50 of 40 nM (± 4 nM SE; n = 17) against the murine homolog (determined by cellular assays using 3T3-L1 adipocytes; data not shown). One percent RO151 was formulated in 9.1% β11-HSD1 activity in vivo (59), with an IC50 of 40 nM (± 4 nM SE; β11) = 17) using RO151, known to selectively inhibit β11-HSD1 inhibitor studies. Inhibitor studies were conducted in 8-week-old male Swiss mice (n = 5–12) using RO151, known to selectively inhibit β11-HSD1 activity in vivo (59), with an IC50 of 40 nM (± 4 nM SE; n = 17) against the murine homolog (determined by cellular assays using 3T3-L1 adipocytes; data not shown). One percent RO151 was formulated in 9.1% β11-HSD1 activity in vivo (59), with an IC50 of 40 nM (± 4 nM SE; β11) = 17). Data represent the means ± SEM. The null hypothesis was rejected at a significance level of P < 0.05, with data values as follows: *P < 0.05; **P < 0.01; and ***P < 0.001.

Study approval

Animal studies were conducted under Home Office license and following approval of the Joint Ethics and Research Governance Committee of the University of Birmingham (Birmingham, United Kingdom) in accordance with the UK Animals (Scientific Procedures) Act, 1986. Human studies were conducted following ethical approval (Black Country Research Ethics Committee, Manchester, United Kingdom) and according to the Declaration of Helsinki Principles. All participants gave their written informed consent before study enrolment.

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