Transcriptional changes in organoculture of full-thickness human skin following topical application of all-trans retinoic acid

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Synopsis

OBJECTIVE: Retinoids are used as therapeutic agents for numerous skin diseases, for example, psoriasis, acne and keratinization disorders. The same substances have also been recognized in the treatment for hyperpigmentation disorders such as melasma.

Other studies on photo-damaged skin have shown that retinoids reduce wrinkles, surface roughness, mottled pigmentation, and visual skin appearance as a whole. We tested the hypothesis that an organoculture of full-thickness human skin could be used as a preclinical model to investigate the retinoid transcriptional profile in human skin in vitro.

METHODS: Full-thickness skin explants were exposed to topically applied all-trans retinoic acid (RA) for 24 h. The gene expression profile was analysed using oligonucleotide microarrays, and data were validated with real-time (RT) PCR.

RESULTS: We showed that the expression of 93 genes was significantly altered more than twofold. Several of the altered genes, for example, KRT4, CYP26 and LCN2, have previously been shown to be affected by RA in keratinocyte monolayers, reconstructed epidermis and skin biopsies from patients treated topically or orally with RA. In addition, genes, such as SCEL, NRIP1, DGAT2, RDH12 EfnB2, MAPK14, SAMD9 and CEACAM6 not previously reported to be affected by RA in human skin, were identified for the first time in this study.

CONCLUSION: The results in the present study show that full-thickness human explants represent a valuable pre-clinical model for studying the effects of retinoids in skin.

Résumé

OBJECTIF: Les rétinoïdes sont utilisés comme agents thérapeutiques pour de nombreuses maladies de la peau, p.ex. le psoriasis, l’acné et les troubles de la kératinisation. Les mêmes substances ont également été reconnues dans le traitement des troubles de l’hyperpigmentation tels que le melasma.

D’autres études sur la peau photo-endommagée ont montré que les rétinoïdes réduisaient les rides, la rugosité de la surface, la pigmentation tachetée, et l’aspect visuel de la peau dans son ensemble. Nous avons testé l’hypothèse selon laquelle une organoculture de épaisseur intégrale de la peau humaine pourrait être utilisée comme modèle préclinique pour étudier le profil transcriptionnel des rétinoïdes dans la peau humaine in vitro.

MÉTHODES: Des explants de peau intégrale ont été exposés à l’application topique de all-trans acide rétinoïque (RA) pendant 24 heures. Le profil d’expression des gènes a été analysé en utilisant des puces DNA array et les données ont été validées par (RT) -PCR.

RÉSULTATS: Nous avons montré que l’expression de 93 gènes a été modifiée de façon significative par plus d’un facteur 2. Plusieurs des gènes modifiés par exemple KRT4, CYP26 et LCN2 ont été montrés précédemment d’être affectés par RA dans les monocultures de kératinocytes, dans l’épiderme reconstruit et dans des biopsies cutanées de patients traités par voie topique ou par voie orale avec RA. En outre, des gènes tels que SCEL, NRIP1, DGAT2, RDH12 EfnB2, MAPK14, SAMD9 et CEACAM6 non signalés précédemment, sont touchés par RA dans la peau humaine, et ont été identifiés pour la première fois dans cette étude.

CONCLUSION: Les résultats de la présente étude montrent que les explants humains de pleine épaisseur représentent un modèle préclinique précieux pour l’étude des effets des rétinoïdes dans la peau.

Introduction

One of the most studied classes of skin-targeted compounds is retinoids. All-trans retinoic acid (RA) also referred to as tretinoin, and its synthetic or natural derivatives (retinoids) affect epidermal growth and differentiation [1]. Synthetic retinoids are today used widely in the treatment of psoriasis and other disorders of keratinization [2]. Also, acne has been widely treated with various forms of natural and synthetic retinoids for more than 40 years [3].

Later, retinoids has been used in the treatment for hyperpigmentation disorders such as post-inflammatory hyperpigmentation and melasma [4].

Separate from the effect of retinoids in skin disease, Kligman and Willis were the first to introduce retinoids for use as antiphotaging agents [5]. After its application, the authors noticed improvement in skin depigmentation and rejuvenation [6]. Today, retinoids are extensively used for this indication [7, 8].

Further studies have shown that RA’s clinical effects include improvement in wrinkles, surface roughness, mottled pigmentation and skin appearance as a whole when used on photo-damaged skin [9–11]. However, topical use of RA has also shown to develop erythema, scaling and burning/pruritus [12]. Therefore, there is a constant thrive within the cosmetic industry to develop retinoid-like anti-ageing ingredients with significantly reduced side effects.
At a cellular level, retinoids are known to modulate the proliferation and differentiation of epidermal keratinocytes [13, 14, 15] by binding to nuclear retinoic acid receptors (RARs) and 9-cis-retinoic acid receptors (RXRs), which results in upregulation or downregulation of the transcription of target genes [16]. RA is the biologically active form of vitamin A (retinol). Endogenous production of RA by epidermal keratinocytes involves the uptake of preformed vitamin A from the surroundings followed by a series of metabolic activation steps involving retinol dehydrogenases (RDH1, RDH4, RDH10, RDH12 and DHR59) and retinal dehydrogenases (RALDH1, RALDH2 and RALDH3) [17–19]. DHR59 specifically is encoding an enzyme that mediates conversion of retinol into RA.

Whilst endogenously produced or exogenously applied, RA is transported through the cytoplasm by specific intracellular retinoid-binding proteins (CRABP2) [8].

Once synthesized, the cellular levels of RA are controlled by several cytochrome P450-dependent enzymes (CYPs) – CYP26 A1, B1 and C1 – which metabolize RA into 4-hydroxy-RA, 4-oxo-RA and 18-OH-RA [20, 21]. CYP26B1 is known to be higher expressed than both CYP26A1 and CYP26C1 in human keratinocytes [19, 22].

Over the last quarter century, 532 genes have been validated to be regulated by RA [23]. In some cases, this control is direct, driven by a liganded RAR:RXR heterodimer bound to a DNA response element; in others, it is indirect, reflecting the actions of intermediated transcription factors, non-classical associations of receptors with other proteins or even more distant mechanisms.

Cell cultures have been studied extensively to characterize and prophesy the effect of retinoids on epidermal differentiation and growth [13, 15, 24–27]. In addition, a selection of genes using RT-PCR has been studied in full-thickness skin treated with retinoids [28, 29].

Today, recent gene array techniques allow the characterization of the mRNA expression status of a large number of genes in cells or tissues after retinoid treatments with more than 170 studies conducted.

Some studies using gene arrays have also been carried out on reconstructed epidermis and human epidermis in vivo [30]. However, to our knowledge, gene array analysis following the treatment of RA on organoculture of full-thickness skin has not been investigated earlier.

The aim of the present study was to determine whether it is possible to use an organoculture of full-thickness skin as a preclinical model to evaluate modulation of gene expression profile following topical application of a commercially available retinoid cream.

**Materials and methods**

**Ex vivo study – skin explant model**

Skin from surgical waste material from breast reductions was collected and immediately put in DMEM at 8°C. All subcutaneous fat was removed with a scalpel to ensure full diffusion of nutrients in the tissue. 8-mm punch biopsies were taken from the skin and referred to as full-thickness skin explants. Thereafter, the explants were put on Millipore Millicell Culture Plate Inserts (12 mm Ø) (Millipore Corporation, Stockholm, Sweden). Inserts containing skin explants were put in 6-well plates (1 insert/well), and 1 mL of supplemented keratinocyte medium (M154) (Life Technologies, Stockholm, Sweden) was added to each well to allow survival of the explants.

Three donors were used for this study. They were all healthy Caucasian females of 23, 41 and 60 years of age. The collection of skin samples was approved by the local Ethics Committee of Stockholm. All participants gave written informed consent for the skin sample collections, and the studies followed the guidelines of the World Medical Association’s Declaration of Helsinki (2000) concerning biomedical research involving human subjects.

Exhibiting an enhanced functional barrier function compared with reconstructed skin [31], the use of viable skin explants allows topical application of finished product to study cosmetic effects on a molecular level. In this study, 5 mg cm⁻² of cream containing RA (tretinoin) (0.05%) (Aberela, Jannsen-Cilag) was applied topically on each explant, using positive displacement pipette. 5 mg cm⁻² is a typical finite dose applied on skin penetration test and considered as a non-occlusive dose [32, 33].

The study was vehicle controlled with a cream containing all ingredients except for RA, referred to as placebo. The placebo cream was applied in the same conditions as the test cream.

The explants were incubated with the RA cream for 24 h at 37°C in 5% CO₂ humidified air. In the end of the incubation period, two Ø 3 mm biopsies were taken from each explants following RNA extraction.

**Skin explant viability**

Alamar blue® (Invitrogen, Stockholm, Sweden) was used as reagent for measuring skin viability. Ø 3 mm biopsies were used for corresponding Alamar blue tests according to manufacturer’s protocol. Briefly, Ø 3 mm biopsy was taken from each organoculture and put in 200 µL of Alamar blue dye and media at 1:10 ratio for 15 h at 37°C. Absorbance was measured at wavelengths of 570 nm and 600 nm to calculate Alamar Blue reduction, which is a measure of reduction reagents of cellular metabolism.

**RNA isolation**

Total RNAs were extracted from skin samples using QIAzol Lysis Reagent (Qiagen). Isolation begins with immersion of skin explants into QIAzol Lysis Reagent (Qiagen) followed by mechanical homogenization of the explants with a rotator–stator homogenizer (TissueRuptor, Qiagen) on ice for 20 s. RNA isolation was performed using the RNasey mini kit (Qiagen), and DNase I digestion was performed following the manufacturer’s instruction.

**Microarray expression analysis**

RNA concentration was measured with ND-1000 spectrophotometer (NanoDrop Technologies, Wilmington, DE), and RNA quality was evaluated using the Agilent 2100 Bioanalyzer system (Agilent Technologies Inc, Palo Alto, CA).

The 250 nanograms of total RNA from each sample were used to prepare biotinylated fragmented cRNA according to the GeneChip® 3' IVT Express Kit Manual (Affymetrix Inc., Santa Clara, CA). Affymetrix GeneChip® expression arrays (Human Genome U133 Plus 2.0) were hybridized for 16 h in a 45°C incubator, rotated at 60 rpm. The arrays were washed and stained using the Fluidics Station 450 and finally scanned using the GeneChip® Scanner 3000 7G (Affymetrix).

**Microarray data analysis**

Subsequent analysis of the gene expression data was carried out in the freely available statistical computing language R (http://www.
Regulation of individual probe sets in gene ontology categories using gene-set enrichment analysis

The enrichment analysis [39] was performed to determine the probability that gene ontology groups are specifically overexpressed in RA-treated skin as compared to vehicle-treated skin. For this analysis, we used the public DAVID Bioinformatics database: http://david.abcc.ncifcrf.gov/ [40]. The enrichment score of each gene was determined by calculating the relative expression in the RA-treated skin compared with placebo-treated skin.

Genes that were significantly upregulated or downregulated more than twofold were included in this analysis to evaluate enriched functional gene ontology groups.

To determine the overrepresented categories, a GO category-specific list for each category of interest was compiled containing those probe sets identified as differentially expressed. A GO enrichment score of >1 was set as a limit to be discussed as a finding.

As a further step, genes that were more than 10-fold differentially expressed were analysed manually to investigate which genes were affected most predominantly by RA.

Real-time reverse-transcriptase polymerase chain reaction (RT-PCR)

To validate the mRNA microarray results, three significantly upregulated genes were selected and their expression was analysed with real-time quantitative polymerase chain reaction (RT-qPCR) (Bio-Rad, Stockholm, Sweden). Skin explants from facial or breast skin from nine patients with a median age of 37 years were used for this conformational study. It is noteworthy that these donors differ from the donors used for the micro-array study. Total RNA from these skin explants treated with RA, as described above, were prepared with RNeasy mini kit (Qiagen) and treated with DNase I (Qiagen) to remove contaminating genomic DNA. Reverse transcription was performed with iScript cDNA synthesis kit (Bio-Rad). SsoFast EvaGreen supermix (Bio-Rad) was used in the RT-qPCR together with specific primers (from Qiagen) for the genes of interest, that is, DHRS9, CYP26B1 and HBEGF. Hot-start two-step PCR, consisting of 95°C for 3 min, followed by 40 cycles of 95°C for 10 s, 57°C for 30 s was performed using CFX96 Real-Time PCR Detection System (Bio-Rad). Relative quantification of gene expression in the different explants was calculated using the delta-delta-Ct method using the TATA box-binding protein (TBP) as a reference gene.

Results

Alamar blue assay

Explants from the three donors tested for the gene array were also tested for viability using Alamar blue assay. The viability of explants was investigated after topical application of RA for 24 h. The results showed that RA was not affecting the viability of the explants during a 24-h application (Fig. 1).

Identification of RA-responsive genes by 24 h stimulation ex vivo (Step 1)

We compared the transcriptional profiles of RA-treated and placebo-treated skin explants (n = 3) using Affymetrix oligonucleotide microarrays. The microarrays simultaneously measure the levels of 22,000 mRNAs. Among them were 93 significantly regulated by RA, within the limits of our selection criteria (less than twofold differentially expressed, P < 0.05). Of these 93 genes, 60 genes were upregulated and 33 downregulated.

To understand the biological meaning behind such a large list of genes, functional annotation clustering analysis was performed using DAVID Bioinformatics resources. The clusters with the smallest P-value (P < 0.05 adjusted using the method of Benjamini and Hochberg [38]) which were retrieved after 24-h stimulation with RA are presented in Fig. 2. When we summarized the functional categories of the genes upregulated by RA, we found that more than 30% of them fall into one of the following ontological categories: clusters (lipid, hormone and retinoid metabolism) and in development (organ, tissue and epithelial development) (Table I).

Cluster 1: Metabolic process (including RA metabolic genes) (enrichment score 3.6)

As shown by the functional annotation clustering analysis, metabolic process was the top cluster identified in our study (enrichment score 3.6) highlighting RA metabolism pathways with totally 11 genes included (Table I). Two main enzymes were identified, involved in the metabolism of RA metabolism, that is, cytochrome P450 26 A1 (CYP26A1), which was highly upregulated and to a lesser extend, and CYP26B1.

Also, the SDR family member dehydrogenase reductase 9 (DHR9), acyl-CoA synthetase long-chain family member 3 (ACSL3), protein convertase subtilisin/kexin type 5 (PCSK5), and nuclear receptor subfamily 3 (NR3C1) was found to be upregulated in our study.

Genes that shown to be downregulated in this cluster were the inflammatory gene arachidonate 12-lipoxygenase (ALOX12), diacylglycerol O-acyltransferase (DGAT2), mitogen-activated protein kinase 14 (MAPK14), serine carboxypeptidase 1 (SCPEP1) and retinol dehydrogenase 12 (RDH12).
Figure 2  Heat map of hierarchical clustering of 93 genes in RA-treated explants compared with placebo. Hierarchical clustering was performed using the GENESIS software [67], with default settings (e.g. Euclidian distance, average linkage) of genes differentially expressed more than less than twofold, that is, 93 genes. Green indicates reduced expression, black indicates the unaltered expression, and red indicates increased expression in RA-treated as compared to placebo-treated explants (n = 3). The colour scale bar is shown at the top of each figure. Cluster analyses of altered genes involved in metabolism genes and development genes. Hierarchal clustering analysis was performed in both the gene (row) and experiment (column) dimension. Yellow arrows indicate genes enriched in gene annotation clustering (David Bioinformatics) which resulted in cluster 1 representing genes involved in metabolism (Enrichment score 3.6). 5 genes of 10 genes in the metabolism cluster were defining genes involved in metabolism of RA. Blue arrows indicate genes enriched in cluster 2 which were defined as involved in development (Enrichment score 2.0) by David Bioinformatics.
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Table 1 Summary of the corresponding genes to the clusters within Fig. 2

| Gene name | Gene symbol | Gene ID | Fold change |
|-----------|-------------|---------|-------------|
| Cluster 1a: Metabolism | | | |
| Cytochrome P450, family 26, subfamily A, polypeptide 1 | CYP26A1 | 206424_at | 22.61 |
| Dehydrogenase/reductase (SDR family) member 9 | Dhrs9 | 224009_x_at | 11.96 |
| Cytochrome P450, family 26, subfamily B, polypeptide 1 | CYP26B1 | 219825_at | 5.26 |
| Acyl-CoA synthetase long-chain family member 3 | ACSL3 | 201652_s_at | 3.66 |
| Proprotein convertase subtilisin/kexin type 5 | PCSK5 | 213652_at | 2.69 |
| Nuclear receptor subfamily 3, group C, member 1 | NR3C1 | 201866_s_at | 2.36 |
| Mitogen-activated protein kinase 14 | MAPK14 | 211561_s_at | 2.13 |
| Retinol dehydrogenase 12 (all-trans/9-cis/11-cis) | RDH12 | 242998_at | 2.42 |
| Diacylglycerol O-acyltransferase homologue 2 (mouse) | DGAT2 | 226064_s_at | 4.58 |
| Arachidonate 12-lipoxigenase | ALOX12 | 207206_s_at | 4.73 |
| Serine carboxypeptidase 1 | SCPEP1 | 218217_at | 6.41 |
| Cluster 1 b: Retinoic acid metabolism | | | |
| Chromosome 10 open-reading frame 54 (c10orf54) was upregulated 11-fold. | C10orf54 | 224009_x_at | 11.96 |
| Cytochrome P450, family 26, subfamily A, polypeptide 1 | CYP26A1 | 206424_at | 22.61 |
| Dehydrogenase/reductase (SDR family) member 9 | Dhrs9 | 224009_x_at | 11.96 |
| Cytochrome P450, family 26, subfamily B, polypeptide 1 | CYP26B1 | 219825_at | 5.26 |
| Retinol dehydrogenase 12 (all-trans/9-cis/11-cis) | RDH12 | 242998_at | 2.42 |
| Serine carboxypeptidase 1 | SCPEP1 | 218217_at | 6.41 |
| Cluster 2: Development | | | |
| Keratin 4 | KRT4 | 213240_s_at | 26.97 |
| Cytochrome P450, family 26, subfamily A, polypeptide 1 | CYP26A1 | 206424_at | 22.61 |
| Dehydrogenase/reductase (SDR family) member 9 | Dhrs9 | 224009_x_at | 11.96 |
| Epithelial membrane protein 1 | EMP1 | 213895_at | 7.31 |
| F-box protein 32 | FBXO32 | 241762_at | 6.39 |
| Paired-like homeodomain 1 | PITX1 | 209587_at | 4.54 |
| Scilin | SCIL | 1554921_a_at | 4.06 |
| Nuclear-receptor-interacting protein 1 | NRIP1 | 202600_s_at | 3.70 |
| Heparin-binding EGF-like growth factor | HBEFG | 38037_at | 10.8 |
| Ephrin-B2 | EFNB2 | 202669_s_at | 3.18 |
| Proprotein convertase subtilisin/kexin type 5 | PCSK5 | 213652_at | 2.69 |
| Nuclear receptor subfamily 3, group C, member 1 | NR3C1 | 201865_s_at | 2.11 |
| (glucocorticoid receptor) | | | |
| Sprouty homologue 1, antagonist of FGF signalling (Drosophila) | SPRY1 | 212558_at | 2.12 |
| Mitogen-activated protein kinase 14 | MAPK14 | 211561_s_at | 2.13 |
| Dystroglycan 1 | DAG1 | 205417_s_at | 2.28 |
| (dystrophin-associated glycoprotein 1) | | | |
| Crystallin, alpha B | CRYAB | 209283_at | 2.66 |
| Prolactin receptor | PRLR | 227629_at | 3.28 |
| Nebulin | NEB | 205054_at | 4.53 |
| Chromosome 1 open-reading frame 68 | C1orf68 | 217087_at | 5.92 |

Table includes gene name, gene symbol, gene ID and fold change compared with placebo. Explain more the redundancy.

Validation of microarray data with qRT-PCR

To confirm independently the microarray results obtained, we performed quantitative RT-PCR analysis of three representative genes, that is, CYP26B1, HRS9 and HBEFG (Fig. 3). These three genes were chosen based on following criteria; they were significantly upregulated by RA and included in the gene ontology groups that were detected via functional annotation clustering using DAVID Bioinformatics resources. These genes are also known to be regulated by RA and expressed in the two top 2 clusters of genes in our expant study: vitamin A metabolism (CYP26B1, DHR9) and development (HBEFG). Results showed upregulation of CYP26B1 (1.52-fold), HBEFG (3.22-fold) and DHR9 (3.35-fold) by topical application of RA.

Step 2: Analysis of genes affected more than 10-fold by RA

Upon analysing the gene array manually for genes affected more than 10-fold, RA was shown to affect 8 genes more than 10-fold (Table II).

| Gene name | Gene symbol | Gene ID | Fold change |
|-----------|-------------|---------|-------------|
| Keratin 4 | KRT4 | 213240_s_at | 26.97 |
| Cytochrome P450, family 26, subfamily A, polypeptide 1 | CYP26A1 | 206424_at | 22.61 |
| (c10orf54), CYP26A1, DHRS9, NR3C1 and MAPK14 were identified in both cluster 1 and 2.

Discussion

Several previous reports describe the genetic targets of retinoids using gene array techniques [41]. However, the genetic profile of RA has not earlier been studied in explants of full-thickness skin.

Cluster 2: Development genes (enrichment score 2.0)

The second cluster was identified as the development genes including tissue development and epithelial development (Table I and Fig. 2). In total, 19 genes were identified in this cluster. Keratin 4 (KRT4) was found to be highly upregulated. Within the same cluster, heparin-binding EGF-like growth factor (HBEFG), CYP26A1, DHRS9, epithelial membrane protein 1 (EMP1), F-box protein 32 (FBXO32), paired-like homeodomain 1 (PITX1), scilin (SCIL), nuclear-receptor-interacting protein 1 (NRIP1), ephrin-B2 (EFNB2), proprotein convertase subtilisin/kexin type 5 (PCSK5) and NR3C1 were identified as genes induced by RA.

Downregulated genes within the development cluster were as follows: sprouty homologue 1 (SPRY1), MAPK14, dystroglycan (DAG1), crystallin, alpha B (CRYAB), prolactin receptor (PRLR), nebulin (NEB), and chromosome 1 open-reading frame 68 (C1orf68), CYP26A1, DHR9, NR3C1 and MAPK14 were identified in both cluster 1 and 2.
In this study, we developed an ex vivo skin explant organoculture model to study the complete gene expression following RA treatment, adopting cDNA micro-array analysis.

Concomitantly, we described an approach for recovering RNA of full-thickness skin taken from the explant model skin. This procedure permits a direct quantitative and qualitative assessment of topical treatment with dermatological active ingredients, such as RA.

We found that 93 genes were significantly more than two-fold regulated by RA in three organocultures of human skin. The marked alterations of the expressions of 93 genes were found to be involved in mainly two clusters, that is, lipid and retinoid metabolism and in development, that is, organ, tissue and epithelial development. Comparing to previous studies in monocultures of keratinocytes, Lee et al. showed that these cells responded to RA very fast with 315 genes affected already at 1 h. Please note that these authors used a threshold of 1.62-fold change to show effect of RA [41].

Lipid and retinoid metabolism

As shown by the functional annotation clustering analysis, metabolic process was the top cluster identified in our study, highlighting lipid and RA metabolism pathways. Two main enzymes were identified with regard to retinoic acid metabolism, that is, CYP26A1, which was highly upregulated (22.6-fold) and to a lesser extend, and CYP26B1 (5.3). The importance of these enzymes with regard to retinoid metabolism in the human skin has been confirmed by other authors [22, 42]. For example, Lee et al. showed induction of CYP26B1 (12.8-fold) by RA in epidermal keratinocytes after 4-h stimulation [41]. In contrast to our study, these authors have showed that CYP26B1 is the most abundant RA metabolizing CYP26 enzyme in unstimulated keratinocytes at the mRNA level [22, 42, 43]. On the other hand, they suggested that CYP26A1 may play a role during situations of excess of RA in epidermis which is the situation in our study.

The expression of DHRS9 was also increased in our study (11.96-fold) and is in agreement with Lee et al.’s study on
epidermal keratinocytes where DHRS9 was induced 3.5-fold after 24 h [41].

Other genes affected involved in metabolism were ACSS3 which was upregulated in our study which is concomitant with reports on the effect of RA on rat hepatoma cells [44]. However, to our knowledge, such an effect of RA has not been reported in human skin to date.

The lipoygenase, ALOX12, involved in fatty acid oxidation, was reduced after RA treatment. These results are compatible with Lee et al.'s study in human keratinocytes where ALOX12B was reduced after RA treatment [41].

Moreover, we found DGAT2 to be reduced (4.58-fold). Same accounts for the expression of RDH12, which was also reduced (2.42-fold) in response to exogenous RA in our study. To our knowledge, RDH12 has not been described in human skin previously.

In addition, MAP kinase 14, which is essential for embryonic development and regulates different cellular functions such as proliferation, differentiation, cell death, adhesion, migration, as well as the response to stress, and many metabolic pathways were found to be downregulated in our study. A similar enzyme, MAPK13, was downregulated in human keratinocytes by RA treatment. The inhibitory effect of RA on MAPK is documented by other authors and has been hypothesized to have a role in the inhibitory action of RA on UV-induced skin damage [45].

Regulation of development genes by RA

Retinoid signalling is known to be important in the development of the eye [46], lungs [47], heart [48] and skin [49, 50]. Adult skin continues to grow throughout the life of humans, and therefore, it is not surprising that genes involved in epidermal development are stimulated after RA application.

In our study, development genes were identified as the second cluster of importance by gene ontology analysis. In this cluster, KRT4 was found to be highly upregulated which is also well known in human keratinocytes [41]. It has even been suggested as a future marker for retinoid bioactivity in epidermis in vivo [41]. Within the same cluster, HBEGF was increased which also is consistent with literature data on the proliferative effect of RA in human keratinocytes [41] and in reconstituted human epidermis [30].

HBEGF was confirmed to be regulated by RA by qPCR in skin explants (Fig. 3). The discrepancy in fold change (10.8 in the microarray study and 3.22 by qPCR) could be due to the difference in donors as well as skin site (facial and breast). In fact, Cyp26a1, Cyp26b1, Rbp1, Crabp1, Hbegf and Krt4 are all known marker for retinoid activity (Gericke et al i.e. ref 65).

Interestingly, EMP1 was found to be highly upregulated (7.31-fold), which is consistent with reports in F9 teratocarcinoma stem cells [51]. EMP1 has been prominently found in the gastrointestinal tract, skin, lung and brain but not in liver. The proteins of this family are likely to serve similar functions possibly related to cell proliferation and differentiation in a variety of cell types [52].

Another gene in the development cluster, scellin (SCEL), a precursor of the cornified envelope [53], was highly upregulated. The specific function of this gene in human skin is not known to date, but deletion of this gene in mice did not affect normal barrier function [54].

NRIP1 was also highlighted in the gene ontology analysis for development of genes, and interestingly, this biomarker has not been found in skin to date. However, it has been described in context of regulation of mesenchymal stem cell differentiation by retinoids as NRIP1 is induced by RA in stem cells [55]. The upregulation of EfnB2 seen by retinoic acid in our study could implicate the regulation angiogenesis and blood vessel development as this has been reported in the literature [56]. However, the role of EfnB2 has not been investigated to date in human skin.

It is noteworthy that NR3C1 also known as glucocorticoid receptor is upregulated in human skin by topical application of RA in our study. To our knowledge, this mechanism has not been reported in human skin. However, a study in cultured chick embryonic skin showed that the addition of 2 μM retinol resulted in a threefold increase in specific dexamethasone binding to the glucocorticoid receptor [57]. Whether the same mechanism applies to our model remains to be elucidated.

Prolactin receptor (PRLR) was profoundly downregulated which is in line with a study in patients with acne where isotretinoin caused suppression of several pituitary hormones such as prolactin in patients' sera [58].

Upon analysing the gene array manually for genes affected more than 10-fold by RA without RA was found to upregulate 8 genes more than 10-fold. One of these was FMO, which was upregulated 50.17-fold after 24-h stimulation with RA. FMO has been found to be present in human skin [59]. It is known to metabolize xenobiotics, but to our knowledge, not known to metabolize retinoids.

SAMD9 was upregulated 40.02-fold by topical application of RA for 24 h. The function of SAMD9 in human skin is to date unknown, but it has been suggested to have a role in regulation of cell death in malignant glioma [60].

Furthermore, CEACAM6/cd66c was induced 34.64-fold. CEACAM6 is found on epithelial and other cells. It has to our knowledge not been described to be expressed in human skin. Other genes from the same family, for example CEACAM1, have shown to be important for the wound healing process in human skin [61].

KRT4 was also highly upregulated (26.97-fold) and was linked to the cluster of development genes by analysis using DAVID Bioinformatics. Same accounts for cytochrome P450, family 26, subfamily A (CYP26A), which was already identified in the metabolism cluster. C10orf54 was upregulated 14.42-fold. This gene encodes for platelet receptor G24. The function of this receptor in human skin is to our knowledge unknown. As already described, DHRS9 was highly upregulated (11.96-fold). LCN2 was upregulated 10.73-fold. The induction of LCN by RA has also been confirmed by other authors. The confirming studies were investigating the effect of oral application of isotretinoin in patients with acne [62, 63] but also of RA in monocytes of keratinocytes [41].

In conclusion, we have in the current study shown that a full-thickness organoculture of human skin is an efficient non-invasive model to study the effects of topically applied compounds at the gene expression level.

We have in this study confirmed several genes affected by RA, which has been previously found by other authors in monocytes of keratinocytes [41], reconstructed epidermis [30] and in skin biopsies after oral application of isotretinoin [64], for example, CYP26A1/B1, DHRS9, ALOX12, KRT4, HBEGF and LCN2. In fact, KRT4, HBEGF and CYP26A1/B1 are considered to be retinoid target genes [65] (Gericke et al., 2013) where KRT 4 is a marker for keratinocyte differentiation [66] (Wanner et al. 1996), CYP26 for retinoid metabolism and HBEGF for increased keratinocyte proliferation and epidermal thickness [8].
We have also disclosed several genes that have never been identified in human skin before, for example SCEL, EfnB2, SAMD9 and CEACAM6. Consequently, the roles of these genes in skin have yet to be discovered.

Overall, we believe that this organoculture model possessing a physiological complexity such as a fully functioning barrier function compared with cell cultures and reconstructed skin models could serve as a model to evaluate the effect of topical application of cosmetic ingredients. The model has been shown in this study to be suitable for assessing short-term stimulation with RA. Therefore, this model can be used as a first screen to identify more effective therapeutic strategies by testing functional connections between drugs and genes using databases such as connectivity map (Cmap, Broad Institute). However, further validation will be needed including a larger sample group to confirm this hypothesis.

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