Review Article

Anti-Aging Potential of Phytoextract Loaded-Pharmaceutical Creams for Human Skin Cell Longetivity

Saima Jadoon,1 Sabiha Karim,2 Muhammad Hassham Hassan Bin Asad,3 Muhammad Rouf Akram,4 Abida Kalsoom Khan,5 Arif Malik,6 Chunye Chen,7 and Ghulam Murtaza3

1Department of Natural Resources Engineering and Management, University of Kurdistan, Hewler 44003, Iraq
2University College of Pharmacy, University of the Punjab, Lahore 54000, Pakistan
3Department of Pharmacy, COMSATS Institute of Information Technology, Abbottabad 22060, Pakistan
4Department of Pharmacy, University of Sargodha, Sargodha 40100, Pakistan
5Department of Chemistry, COMSATS Institute of Information Technology, Abbottabad 22060, Pakistan
6Institute of Molecular Biology and Biotechnology, University of Lahore, Lahore 54000, Pakistan
7Key Laboratory of Biorheological Science and Technology, Chongqing University, Chongqing 400030, China

Correspondence should be addressed to Ghulam Murtaza; gmdogar356@gmail.com

Received 12 December 2014; Accepted 13 March 2015

Academic Editor: Gabriele Saretzki

Copyright © 2015 Saima Jadoon et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

The exposure to ultraviolet radiations (UVR) is the key source of skin sunburn; it may produce harmful entities, reactive oxygen species (ROS), leading to aging. The skin can be treated and protected from the injurious effects of ROS by using various pharmaceutical formulations, such as cream. Cream can be loaded with antioxidants to quench ROS leading to photo-protective effects. Moreover, modern medicines depend on ethnobotanicals for protection or treatment of human diseases. This review article summarizes various in vivo antioxidant studies on herbal creams loaded with phyto-extracts. These formulations may serve as cosmeceuticals to protect skin against injurious effects of UVR. The botanicals studied for dermatologic use in cream form include Acacia nilotica, Benincasa hispida, Calendula officinalis, Camellia sinensis, Camellia sinensis, Nelumbo nucifera, Capparis decidua, Castanea sativa, Coffea arabica, Crocus sativus, Emblica officinalis Gaertn, Foeniculum vulgare, Hippophae rhamnoides, Lithospermum erythrorhizon, Malus domestica, Matricaria chamomilla L., Moringa oleifera, Morus alba, Ocimum basilicum, Oryza sativa, Polygonum minus, Punica granatum, Silybum marianum, Tagetes erecta Linn., Terminalia chebula, Trigonella foenum-graecum, and Vitis vinifera. The observed anti-aging effects of cream formulations could be an outcome of a coordinating action of multiple constituents. Of numerous botanicals, the phenolic acids and flavonoids appear effective against UVR-induced damage; however the evidence-based studies for their anti-aging effects are still needed.

1. Etiologies and Types of Human Skin Aging

Skin aging is a dermatologic change that progresses as a person ages or is exposed to ultraviolet radiations (UVR) if no treatment is adopted. The extensive research activities are focused on this skin concern that involves the appearance of unpleasant, observable marks on skin surface due to proteolysis of cutaneous elastic fibers resulting in the reduced cell functions [1]. Skin aging can be divided into two types, that is, intrinsic aging or chronological aging (inevitable phenomenon) and extrinsic or premature or photoaging (evitable phenomenon) owing to the physiological and environmental factors, respectively [2–4]. Morphologically, photoaging is characterized by dry, rough, pigmented, and abraded skin especially of face and hands in individuals who live in sunny geographical regions and are chronically exposed to direct sunlight (Figure 1). Conversely, fine, smooth wrinkles on dry, pale skin impart the characteristics of intrinsic aging [1]. Diagnostically, intrinsic skin aging is identified by seborrheic keratosis which is not a biomarker of photoaging [5]. Pathologically, the photodamaged skin shows vascular damage that is absent in intrinsically aged skin. An increased
skin vascularization and angiogenesis are observed in the photoaged skin [6]. Microscopically, thicker epidermis is another feature of the photoaged skin [4]. It is noteworthy that the strength and resiliency of skin depend on proper and uniform arrangement of collagen (types I and III) fibrils and elastin in the dermis [7]; thus, collagen deficiency may result in skin aging due to the production of collagenase and thymine dimer in skin on exposure to UVR. Histologically, the extracellular matrix of intrinsically aged skin possesses diminished levels of elastin [2], while the elastin amassing in the photoaged skin is observed just below the dermal-epidermal junction [8]. Elastin is a fibrous protein that is reduced in thickness from deeper to superficial dermis. It provides natural elasticity and strength to human body. It also plays a role in tissue repair [9]. The basic and the major molecular unit involved in the construction of human skin is collagen that is produced from procollagen. Collagen is a protein that is present in the connective tissues of human body. The dermal fibroblasts generate the procollagen under the effect of transforming growth factor-β (TGF-β) and activator protein-1 (AP-1), where TGF-β and AP-1 govern the production and breakdown of collagen, respectively. Under the effect of UVR received from sun, the upregulation of matrix metalloproteinases (MMPs) enzymes secreted by keratinocytes, fibroblasts, and other cells promotes breakdown of collagen by AP-1 as well as decrease in collagen synthesis (Figure 1) [10, 11]. It results in breakdown of the connective tissues during photoaging [12–14]. During adulthood, there is about 1% decrease in collagen content per year, but this rate is higher in the aged people since old age people have higher levels of MMP [7].

2. Reactive Oxygen Species and Photoaging

The exposure to UVR is the main cause of oxidative stress in the skin and thus is an important risk factor for development of skin problems, for example, wrinkle formation, lesions, and cancer. On exposure to sunlight, skin molecules absorb UVR resulting in the generation of reactive oxygen species (ROS). There are two types of ROS: type 1 consists of a single, excited oxygen molecule (¹O₂) (Figure 3), while oxygen molecules with unpaired electron constitute second type of ROS. The examples of second type are presented in Table 1 that also describes the enzymes which are involved in the generation of these ROS [16]. Reactive oxygen entities exert a damaging effect on cellular fractions including cell walls, lipid membranes, mitochondria, nucleus, and DNA producing “oxidative stress,” that is, a difference between ROS and antioxidants, ROS being in excess leading to tissue injury and development of disease including aging, cancer, ischemia, liver injury, arthritis, and Parkinson's syndrome (Figure 2).

3. Benefits and Types of Antioxidants

The oxidative stress-mediated development of diseases is manageable by prolonged usage of the safe antioxidants [17]. The literature study reveals that numerous compounds have been investigated with the intention of exploring evidence against ROS-induced damage and noted their antiaging effect on skin. These compounds are efficient for overcoming sunlight-induced skin problems and making it fresh, healthy, and young through collagen synthesis [18]. Generally, the antioxidants behave as antiaging compounds in action because they are capable of scavenging ROS leaving healthy effect on skin. Since living systems have capability to maintain homeostasis of ROS in cell, the human skin is protected from UVR through complex antioxidant defense system comprising of two types of antioxidants, that is, endogenous and exogenous (consumed) antioxidants. The former category constitutes a network of protective antioxidants in skin; it includes melanin and some enzymes. Manganese-superoxide dismutase is a mitochondrial enzyme that destroys the superoxide ions produced by respiratory chain activity [19]. In
Table 1: Enzymes involved in the generation of ROS with an unpaired electron.

| Number | Name                     | Symbol | Enzymes                                                                 |
|--------|--------------------------|--------|-------------------------------------------------------------------------|
| 1      | Superoxide anion radicals| \( \cdot O_2 \) | NADPH oxidase and xanthine oxidase [22–24]                           |
| 2      | Hydroxyl radicals         | \( \cdot OH \)  | Superoxide dismutase (SOD) [25]                                       |
| 3      | Nitric oxide radicals     | \( \cdot NO \)  | Nitric oxide synthase (NOS) [26]                                       |
| 4      | Lipid peroxy radicals     | \( \cdot LOO \) | Superoxide dismutase [25]                                             |

Exposure to UV radiations

Extrinsic aging

Degradation of extrinsic antioxidants

Intrinsic aging

Inflammatory response

Exposure to UV radiations

Extracellular matrix degradation

Increased reactive oxygen species

Activation of AP-1

Suppression of TGFB receptor

Activation of MMPs

Suppression of procollagen synthesis

Degradation of collagen

Reduced level of collagen in dermal layer

Elastosis

Appearance of wrinkles

Photoaging

Figure 2: Mechanism of aging.

In general, expression of antioxidant enzymes is found very high in the epidermal layer compared to that of stratum corneum and dermis. If there is imbalance between oxidants and endogenous antioxidants, exogenous antioxidants are helpful to restore the balance. The exogenous antioxidants comprise of compounds that cannot be synthesized by human body. Vitamins, ascorbate, carotenoids, and polyphenols constitute latter type of antioxidants which are also involved in the maintenance of oxidative homeostasis [20]. The endogenous antioxidants in dermal and epidermal layers of skin exposed to sunlight are depleted under the effect of elevated levels of UVR-generated ROS. Such depletion results in the diminished activity of these antioxidants leading to skin damage [21]. With age, endogenous antioxidants are steadily consumed increasing the risk of oxidative stress; then the use of exogenous antioxidants as prevention strategy is essential.

It is evident from the above discussion that skin cells are damaged by oxidative stress which might be decreased by action of the antioxidants.

4. Exogenous Antioxidants

The exogenous antioxidants include synthetic and natural compounds. The synthetic exogenous antioxidants include monoethanolamine, diethanolamine, sodium laureth sulfate, and triethanolamine, but these compounds have undesired effects including allergic and irritant contact dermatitis and contact dermatitis [19]. On the other hand, natural exogenous antioxidants are nontoxic in nature and produce no unwanted effect on skin.
5. Phytoantioxidants

The phyto-kingdom includes vegetables, fruits, whole grains, and beverages, for example, tea, chocolate, and wine. These products are rich in natural antioxidants. An important class of natural exogenous antioxidants is phytoantioxidants, that is, antioxidants found in plants [21]. Phytoantioxidants include terpenes or polyphenols (Figure 4). After synthesis in plants, these compounds are found to have important role in the metabolism and defense system of plants. Terpenes are known to have potential for managing the oxidative stress through their free radical scavenging potential. Moreover, polyphenols occur in all parts (roots to leaves) of the plants and protect them from environmental stress through their free radical scavenging property. There are various types (>8000 phenolic structures) of polyphenols on the basis of...
molecular weight and polarity [64]. The structural formula of polyphenols contains phenol group(s), that is, benzene ring possessing hydroxyl group. The antioxidant activity of various polyphenols depends on number and position of phenol groups [65].

6. Stratum Corneum as Target Site for Antioxidants

The normal human skin maintains homeostasis of water and other materials in body, principally due to the presence of stratum corneum [66]. The stratum corneum, a water barrier in function, consists primarily of lipids, that is, ceramides, cholesterol, free fatty acids, triglycerides, stearyl esters, and cholesterol sulfate. The cholesterol sulfate is responsible for intercellular adhesion, and its high concentration is known to inhibit desquamation. The synthesis of these lipids is affected by many factors, mainly related to enzymes, fatty acids, environment, cosmetics, and water contents. Other important constituents of stratum corneum are proteins (e.g., involucrin and loricrin), enzymes, and water (approximately 30%) [67]. Depending upon natural moisturizing factor of skin cells, some fraction of this water is tightly held in stratum corneum and is responsible for skin elasticity. The disturbance in level or nature of any of lipids, proteins, enzymes, and water might lead to skin problems including wrinkled and dry skin. Dry skin might be due to excessive transepidermal water loss that could be retained for maintaining the proper skin hydration by using skin moisturizer. It might exercise softening effect on skin. The skin moisturizer, however, should be inert, nonirritant, stable, and sterile [68]. On the other hand, skin wrinkles might be due to distorted elastic fibers, diminished collagen contents, and uneven types I and III collagen. There is decrease in type IV collagen protein at the wrinkle’s base; it could be due to activation of MMPs, the collagen-degrading enzymes. Alternatively, the activation of MMPs may lead to upregulation of collagenase, gelatinase, and stromelysin [69]. Thus, the skin wrinkles could be treated by using topical formulations loaded with the bioactive compounds having potential of inhibiting MMPs, thus increasing the collagen level. Moreover, skin colour depends on the kind and allocation of melanin in the skin, in addition to number and amount of melanocytes [67, 70]. Figure 5 shows the melanin synthesis involving tyrosinase and a series of oxidative reactions that could be inhibited by usage of skin-whitening agents. Thus, the stratum corneum is a primary target site for topical phytoantioxidants for skin protection against UVR-mediated oxidative stress [71]. The phytoantioxidants might have capability of stimulating the regeneration of stratum corneum to protect itself and the underlying epidermis and dermis from the injurious effects of UVR and promote growth of the skin [72, 73].

7. Skin Care Products

The pharmaceutical formulations used for skin care, termed as the cosmetics, could be herbal in nature. The herbal cosmetics might contain the isolated bioactive compounds or the crude phytoextracts [74]. Currently, there are extensive research activities in progress involving development and characterization of extract loaded formulations to concurrently achieve various goals such as anti-inflammatory and antiaging effect [75]. There are three types of bioactive compounds present in various phytoextracts; the compounds include polyphenols, flavonoids, and carotenoids. These compounds exert both the antioxidant and the UV protection effect [76].

8. Pharmaceutical Creams

Skin care products could be solid, semisolid, or liquid. The semisolid formulations include creams, ointments, and pastes. Cream is an emulsion of oil and water, prepared for skin applications [77]. Emulsions represent a class of disperse systems which comprise of two insoluble, thermodynamically stable phases, that is, continuous and dispersed phase [78]. The emulsion is water-in-oil if the dispersed phase is oil and vice versa. This type of emulsion is termed as simple emulsion. If simple emulsion is further dispersed in the dispersed phase medium, such type of system is termed as the multiple emulsion. Based on globule size of dispersed phase, emulsions can be grouped into various classes including macro- and microemulsion [79]. Emulsions constitute an exclusive class of cosmetics that produce a pleasant feeling to skin on application, acceptable for long-term use, improved spreadability of the ingredients, and remain stable during long storage period [80]. Owing to these characteristics, emulsions are extensively used as a vehicle in drug delivery, particularly across the skin. In particular, for dry skin, water-in-oil (W/O) emulsions are more broadly used for the treatment of dermatological concerns [81]. The addition of antioxidants as active ingredients endows these emulsions with features of cosmetics. For improved cosmetic features, the botanical extracts can be added to the cosmetic creams since the extracts comprise of a number of antioxidants that might produce synergistic effect [82].

9. Preparation and Characterization of Phytoextract Loaded Creams

The creams are topically used to protect and treat the skin problems including hyperpigmentation and wrinkles. Beside these advantages, the creams may produce skin problems such as infection, photosensitivity, erythema, contact dermatitis, cancer, and/or change in skin colour. During the development of such antiaging creams, the researchers should be more focused on the elucidated sources, structures, and interactive modes of the composite active constituents with the skin to achieve maximum formulation efficacy and skin safety [83].

The preparation of herbal creams may involve the modified methodology using isolated phytochemicals or the extracts along with appropriate composition of the mandatory constituents essentially employed for creams with desirable features [84]. Due to technical sophistication of cream development, it is accentuated that the phytochemicals...
Postinflammatory, hyperpigmentation, pregnancy, drugs, photosensitizing agents, UV light, or systemic diseases (e.g. Addison’s disease, liver disease, and pituitary tumors)

Figure 5: Melanin synthesis involving tyrosinase and a series of oxidative reactions.

maintain their bioactivity during extreme processing. To ensure effective and stable formulations, exhaustive analytical strategies are adopted. Various physicochemical characterization parameters include stability, pH, and viscosity testing [85]. Table 3 shows various types of equipment used for in vivo characterization of botanical creams.

10. Phytoextract Loaded Creams

Due to the presence of numerous bioactive ingredients in phytoextracts, extract loaded creams are considered more efficacious with lesser side effects against aging in comparison to creams loaded with specific individual antioxidant. Owing to tremendous antioxidant potential, phytoextracts are extensively used in numerous cream formulations. Up to now, Acacia nilotica, Benincasa hispida, Calendula officinalis, Camellia sinensis, Nelumbo nucifera, Capparis decidua, Castanea sativa, Coffea arabica, Crocus sativus, Emblica officinalis Gaertn, Foeniculum vulgare, Hippophae rhamnoides, Lithospermum erythrorhizon, Malus domestica, Matricaria chamomilla L., Moringa oleifera, Morus alba, Ocimum basilicum, Oryza sativa, Polygonum minus, Punica granatum, Silybum marianum, Tagetes erecta Linn., Terminalia chebula, Trigonella foenum-graecum, and Vitis vinifera have successfully been used in developing the stable cream formulations with excellent antioxidant effect, possibly due to presence of multiple antioxidant phytochemicals. In this review article, the documented phytoextract loaded creams with their characterization on human skin have been discussed with special emphasis on their bioactive constituents (Table 2).

10.1. Acacia nilotica. Acacia nilotica (Mimosaceae) contains tannins, gallic acid, phlobatannin, pyrocatechol, (+)-catechin, protocatechuic acid, (−)-epigallocatechin-7-gallate,
Table 2: Phytoextracts loaded to creams with antioxidant features studied in human.

| Botanical name               | Family                  | Part used | Nature of extracts | Antioxidants                         | Nature of cream | Oil phase          | Emulsifier   | References         |
|------------------------------|-------------------------|-----------|--------------------|--------------------------------------|-----------------|--------------------|--------------|--------------------|
| Acacia nilotica              | Mimosaceae              | Bark      | Ethanol            | Phlobatannin, pyrocatechol, (+)-catechin, proocatechuic acid, (-)-epigallocatechin-7-gallate, and (-)-epigallocatechin-5, 7-digallate | Simple W/O cream†† | Paraffin oil       | ABIL EM 90    | [27]               |
| Benincasa hispida            | Cucurbitaceae           | Fruit     | Petroleum ether    | Caffeic acid                         | Simple W/O cream†† | Cetyl alcohol      | Polysorbate   | [28]               |
| Calendula officinalis        | Compositae              | Flowers   | Ethanol            | Isohamnetin, quercetin, myricetin, and kaempferol | Simple W/O cream†† | Paraffin oil       | ABIL EM 90    | [29, 30]           |
| Camellia sinensis            | Theaceae                | Leaves    | Ethanol            | Epigallocatechin gallate             | Simple W/O cream†† | Paraffin oil       | ABIL EM 90    | [31, 32]           |
| Camellia sinensis (green tea) and Nelumbo nucifera (lotus) |  | Leaves | Ethanol for Camellia sinensis; methanol for Nelumbo nucifera | Epigallocatechin gallate in green tea; hyperin, isoquercetin, and astragalin in lotus | W/O/W nanomultiple-emulsions | Paraffin oil | ABIL EM 90, polyoxyethylene (20) cetyl ether, Cetomacrogol 2000 | [33–35] |
| Capparis decidua             | Capparidaceae           | Full plant | Methanol          | Isoginkgetin and ginkgetin          | Simple W/O cream†† | Paraffin oil       | ABIL EM 90    | [36]               |
| Castanea sativa              | Fagaceae                | Leaves    | Ethanol            | Catechin and myricetin derivatives   | Simple W/O cream†† | Surfactant-free formulation | ABIL EM 90 | [37]               |
| Coffea arabica               | Rubiaceae               | Berry     | Ethanol            | Chlorogenic acid, condensed proanthocyanidins, quinic acid, and ferulic acid | Simple cream†† | Information not available | Information not available | [38]               |
| Crocus sativus               | Iridaceae               | Flowers   | Ethanol            | Zeaxanthin, lycopene, carotenes, crocetin, and picrocrocin | Simple W/O cream†† | Paraffin oil       | ABIL EM 90    | [39]               |
| Emblica officinalis Gaertn   | Euphorbiaceae           | Fruit     | Hydroalcoholic     | Emblicanin A, emblicanin B, puniglucosin, and pedunculagin | Simple W/O cream†† | Paraffin oil       | ABIL EM 90    | [40]               |
| Foeniculum vulgare           | Apiaceae                | Seeds     | Ethanol            | Gallic acid, caffeic acid, ellagic acid, quercetin, and kaempferol | Simple W/O cream†† | Paraffin oil       | ABIL EM 90    | [41, 42]           |
| Hippophae rhamnoides         | Elaeagnaceae            | Fruit     | Hydroalcoholic     | Isohamnetin, quercetin, myricetin, and kaempferol | Simple W/O cream†† | Paraffin oil       | ABIL EM 90    | [43]               |
| Botanical name               | Family               | Part used | Nature of extracts | Antioxidants | Nature of cream       | Oil phase                      | Emulsifier                     | References |
|-----------------------------|----------------------|-----------|--------------------|--------------|-----------------------|--------------------------------|--------------------------------|------------|
| *Lithospermum erythrorhizon* | Boraginaceae         | Root      | Ethanol            | Naphthoquinone (shikonin, acetylshikonin, deoxyshikonin, b-acetoxyisovaleryshikonin, isobutylshikonin, b,b-dimethyl acrylshikonin, 2-methyl-n-butyrylshikonin, and isovaleryshikonin) | Simple O/W cream†*†         | Cyclomethicone, caprylic/capric triglyceride, phytosphingosine (0.005%), and cholesterol | Sodium lauroyl lactylate         | [44]      |
| *Malus domestica*           | Rosaceae             | Fruit     | Methanol: formic acid: double distilled water (70:2:28) | Hesperetin   | Simple W/O cream††    | Paraffin oil                   | ABIL EM 90                      | [45, 46]   |
| *Matricaria chamomilla L.*  | Asteraceae           | Whole plant | Hydroalcoholic    | α-Bisabolol and apigenin | Simple W/O cream††      | Cetyl alcohol                  | Sodium lauroyl lactylate         | [47]      |
| *Moringa oleifera*          | Moringaceae          | Leaves    | Hydroalcoholic    | Epigallocatechin gallate, myricetin, quercetin, rutin, morin, taxifolin, chrysins, baicaletin, fisetin, biochanin A, genistein, kaempferol, emodin anthraquinone, caffeic acid phenethyl ester, and octyl and dodecyl gallates | Simple O/W cream††          | Paraffin oil                   | ABIL EM 90                      | [48–50]    |
| *Morus alba*                | Moraceae             | Fruit     | Hydroalcoholic    | Rutin, quercetin, isoquercitrin, and quercetin | Simple O/W cream††       | Paraffin oil                   | ABIL EM 90                      | [51, 52]   |
| *Ocimum basilicum*          | Lamiaceae            | Seeds     | Ethanol            | Quercetin, isoquercetin, kaempferol, caffeic acid, rosmarinic acid, rutin, catechin, ferulic acid, rutinsoide, and apigenin | Simple W/O cream††         | Paraffin oil                   | ABIL EM 90                      | [53]      |
| *Oryza sativa*              | Poaceae              | Grains    | Ethanol            | Gallic acid, pyrogallol, apigenin, and rutin | Niosomes loaded cream    | Information not available      | Information not available      | [54]      |
| *Polygonum minus*           | Polygonaceae         | Leaves    | Aqueous            | Caffeic acid and quercetin | Simple W/O cream††      | Isoparaffin                    | Laureth-7                       | [55]      |
| *Punica granatum*           | Punicaceae           | Seeds     | Ethanol            | Ellagic acid | Nanotransfersomes loaded cream | Cetyl alcohol | Span 60 and tween 80 | [56]      |
| *Silybum marianum*          | Asteraceae           | Seeds     | Ethanol            | Silymarin (silybin, silydianin, and silychristin) | W/O emulsion cream      | Paraffin oil                   | ABIL EM 90                      | [57, 58]   |
| Botanical name       | Family       | Part used | Nature of extracts | Antioxidants<sup>†</sup> | Nature of cream | Oil phase                          | Emulsifier                     | References |
|---------------------|--------------|-----------|--------------------|--------------------------|----------------|-----------------------------------|--------------------------------|------------|
| *Tagetes erecta* Linn. | Asteraceae   | Flowers   | Ethyl acetate      | Lutein                   | Nanostructured lipid carrier loaded cream | Glyceryl monostearate, stearic acid, octylidodecanol, and mineral oil | Tween, span, or triethanolamine stearate | [59]       |
| *Terminalia chebula* | Combretaceae | Seeds     | Methanol           | Gallic acid, ellagic acid, tannic acid, ethyl gallate, chebulic acid, chebulagic acid, corilagin, and ascorbic acid | Simple W/O cream<sup>††</sup> | Paraffin oil                      | ABIL EM 90                     | [60]       |
| *Trigonella foenum-graecum* | Fabaceae   | Seeds     | Methanol           | Kaempferol derivatives such as 3-O--D-glucosyl(1-2) - -D-galactoside | Simple W/O cream<sup>††</sup> | Paraffin oil                      | ABIL EM 90                     | [61, 62]   |
| *Vitis vinifera*    | Vitaceae     | Shoot     | Ethanol            | Resveratrol, delphinidin,peonidin, petunidin, malvidin, and (+)-catechin | Simple W/O cream<sup>††</sup> | Cetyl alcohol                     | Span 60                         | [63]       |

<sup>†</sup> Including but not limited to.

<sup>††</sup> Simple cream is the cream that is not loaded with some novel carrier system such as nanotransfersomes.
and (−)-epigallocatechin-5,7-digallate. All parts of this plant, from root to flowers, possess various medicinal activities such as antidiabetic, antiasthmatic, and anticancer [86]. Ali et al. prepared water-in-oil cream of bark extract of Acacia nilotica and applied it to photoaged skin [27]. They found improved mechanical features of skin, that is, reduced levels of roughness, scaliness, smoothness, and wrinkles of photoaged skin. The investigators attributed this antiaging activity to the phenolic compounds present in extract; the phenolics have capability to quench ROS.

10.2. Benincasa hispida. The major constituents of Benincasa hispida (Cucurbitaceae) are triterpenoids, flavonoids, glycosides, saccharides, carotenes, vitamins, β-sitosterin, and uronic acid [87]. The fruit of this plant is effective for different diseases including cardiac disease, diabetes, inflammation, and cancer [88]. The water-in-oil cream of fruit extract of Benincasa hispida showed antioxidant activity revealing its potential to retard symptoms of aging [28].

10.3. Calendula officinalis. Calendula officinalis belongs to family Compositae. This plant is rich in active compounds including terpenoids, carotenoids, flavonoids, and volatile oils [89]. The water-in-oil cream of flower extract of Calendula officinalis exhibited aptitude of stimulating skin tightness and improved skin elasticity leading to delayed aging process. Moreover, this preparation enhances the skin hydration level, as evident from reduced TEWL values, which is crucial for normal cutaneous metabolism; it prevents early aging [29]. In addition, the reduction in skin melanin contents and the decrease in skin sebum level were also observed after application of this formulation [30].

10.4. Camellia sinensis (Green Tea) and Nelumbo nucifera (Lotus). Potent antioxidants have been isolated from both green tea and lotus. Green tea is rich in a polyphenol, catechins, especially EGCG which is a strong antioxidant [90]. The water-in-oil cream loaded with ethanolic leaf extract of Camellia sinensis significantly reduced the sebum production as compared to base formulation (formulation without extract) [31]. In another study, Mahmood et al. reported reduced TEWL values and thus improved skin hydration level using this formulation [32]. Furthermore, the W/O/W nano-multiple emulsion containing alcoholic extracts of Camellia sinensis (Theaceae) leaves and Nelumbo nucifera (Nelumbonaceae) plant, alone or in combination, was formulated and tested on human skin. Both extracts were found to have diminished sebum secretions for mono (green tea or lotus) and combined treatments (green tea plus lotus) after applying same formulations to volunteers’ skin. Green tea plus lotus together produced statistically better antisebum effect [33]. It indicates that active ingredients in lotus add a synergistic effect to the activity of green tea that can be attributed to 5α-reductase inhibition activity of both extracts [91]. Zinc compounds, polyphenolics, flavonoids, tannic acid, and linoleic acid are excessively found in lotus [92]. Linoleic acid, a polyunsaturated fatty acid, and EGCG have an advantage of inhibiting sebum production due to their 5α-reductase inhibition activity [93]. In another study, the researchers noted the significant effectiveness of nano-multiple emulsion against skin wrinkles, roughness, and scaliness leading to skin revitalization effect. Green tea and lotus combined in multiple emulsions exerted a superior synergistic antiaging effect [34]. In addition, Mahmood and Akhtar also reported that this formulation had an advantage of reducing melanin and improving hydration contents of skin without causing erythema [35].

10.5. Capparis decidua. Capparis decidua belongs to family Capparidaceae [94]. This plant is rich in active compounds including isothiocyanate glucoside, glucocapparin, stachydrine, n-triacontane, β-carotene, and β-sitosterol [95]. The water-in-oil emulsion cream of plant extract of Capparis decidua exhibited reduction in skin sebum level after application on skin [36].

10.6. Castanea sativa. Castanea sativa (Fagaceae) contains catechin, myricetin 3-O-glucoside, quercetin 3-O-rutinoside, quercetin 3-O-glucoside, kaempferol 3-O-rutinoside, and kaempferol 3-O-glucoside. The surfactant-free cream of ethanolic extract of Castanea sativa leaves has been found to produce moisturizing effect on human skin by controlling transepithelial water loss [37].

10.7. Coffea arabica. Coffea arabica belongs to family Rubiaceae. This coffee plant is rich in active compounds including chlorogenic acid, condensed proanthocyanidins, quinic acid, and ferulic acid [96]. All of these polyphenolic compounds possess antioxidant property [97]. McDaniel prepared cream of berry extract of Coffea arabica and found reduced levels of MMP-1 and IL-1β [38]. Moreover, upregulated gene expression for four collagen structural proteins and downregulated gene expression for three MMPs were also observed concluding reparative effects of Coffea arabica extract upon photoaged skin.

Table 3: Equipment used for in vivo characterization of botanical creams.

| Number | Equipment   | Purpose of use                      |
|--------|-------------|-------------------------------------|
| 1      | Mexamer     | Erythema analysis                   |
| 2      | Tewameter   | Transepideral water loss (TEWL) evaluation |
| 3      | Corneometer | Detection of skin hydration         |
| 4      | Evaporimeter| Barrier function test               |
| 5      | Sebumeter   | Assessment of skin surface sebum/lipid contents |
| 6      | Visiometer  | Wrinkle test                        |
| 7      | Cutometer   | Measurement of skin mechanical properties/elasticity |
| 8      | Chromamer   | Skin colour test                    |
10.8. *Crocus sativus*. Water-in-oil emulsion based cream loaded with ethanolic extract of *Crocus sativus* flowers has been found to produce moisturizing effect on human skin by controlling transepithelial water loss [39]. *Crocus sativus* (Iridaceae) contains zeaxanthin, lycopene, carotenes, crocetin, and picrocrocin.

10.9. *Emblica officinalis Gaertn*. Some potent antioxidants including gallotannins, for example, emblicanin A, emblicanin B, punigluconin, and pedunculagin, are present in *Emblica officinalis* Gaertn (Euphorbiaceae), generally known as Amla [98]. This plant exists in China, India, and Indonesia. All parts of this plant, from root to fruit, are medicinally effective in different health problems; for example, diarrhea and jaundice are effectively treated by using its fruit [99]. The hydroalcoholic fruit extract of *Emblica officinalis* has been formulated as water-in-oil cream that is found effective for reducing transdermal water loss, as observed by using Tewameter [40]. Since transdermal water loss plays crucial role in aging process this formulation can be employed as an antiaging product. This property of cream could be due to presence of strong antioxidants in *Emblica officinalis*.

10.10. *Foeniculum vulgare*. On application of water-in-oil emulsion based cream loaded with ethanolic extract of *Foeniculum vulgare* (Apiaceae) seeds for eight months, reduced transepithelial water loss was observed which resulted in improved moisture contents on human skin [41]. Moreover, there was an improvement in skin mechanical properties, that is, reduced levels of roughness, scaliness, smoothness, and wrinkling of photoaged skin. In another study, Rasul et al. applied same formulation on hyperpigmented human skin and observed the decreased melanin level, sebum production, and erythema of the treated skin [42]. The decrease in skin melanin level leads to skin-whitening effect. Beside flavonoids, they attributed this effect to linoleic acid present in the used extract. This unsaturated fatty acid, a main constituent of biological cell membranes, has an advantage of accelerating process of tyrosinase degradation resulting in reduced melanin synthesis due to low tyrosinase levels [100]. Furthermore, this cream may be useful for skin acne due to its diminishing effect on skin sebum level. Oleic acid, linolenic acid, and linoleic acid, present in *Foeniculum vulgare* extract, could be responsible for this effect [101]. These unsaturated fatty acids have inhibitory effect on sebum production owing to selective inhibition of 5-reductase which is involved in production of sebum [102]. Lastly, there was decrease in erythema on treated skin; this effect elaborates the anti-inflammatory action of this cream.

10.11. *Hippophae rhamnoides*. By using Cutometer, Khan et al. reported the improvement in facial skin mechanical parameters, for example, skin elasticity, indicating antiaging effect after using water-in-oil based hydroalcoholic cream loaded with fruit extract of *Hippophae rhamnoides* [43]. Another study reported the antisemum secretion effect of same formulation [103]. Moreover, the extracts of *Hippophae rhamnoides* and *Cassia fistula* were also found effective in the reduction of skin sebum content (antiacne effects) in human with grade I and grade II acne vulgaris [104]. Khan et al. reported that this formulation improves barrier function of human skin as tested by Tewameter and Corneometer [105]. The possible antiaging effect was pointed out as a feature of antioxidants such as carotene, particularly β-carotene, vitamin C, and vitamin E present in extract. Vitamin C occurs in a concentration of 28–2500 mg/100 g of *Hippophae rhamnoides* extract [106] and plays a role in the stimulation of dermal fibroblasts to synthesize collagen which is responsible for holding water contents in skin [107]. The *Hippophae rhamnoides* extract affects skin mechanical properties through increased expression of cell surface integrins that promote collagen contraction. Moreover, Khan et al. reported reduced TEWL and thus increased skin hydration level for same formulation [108]. Furthermore, same researchers also described reduction in skin melanin level and erythema by using this cream [109].

10.12. *Lithospermum erythrorhizon*. Oil-in-water emulsion based cream loaded with ethanolic extract of *Lithospermum erythrorhizon* root has been found to produce moisturizing effect on human skin by controlling transepithelial water loss. *Lithospermum erythrorhizon* (Boraginaceae) contains shikonin, acetylshikonin, deoxyshikonin, b-acetoxyisovalerylshikonin, isobutylshikonin, b,b-dimethyl acrylshikonin, 2-methyl-n-butyrylshikonin, and isovalerylshikonin [44].

10.13. *Malus domestica*. The reduced transepithelial water loss has been observed leading to improved moisture contents in human skin after application of water-in-oil emulsion based cream loaded with hydroalcoholic extract of *Malus domestica* (Rosaceae) seeds for eight months [45]. Moreover, there was an improvement in skin mechanical properties, that is, reduced levels of roughness, scaliness, smoothness, and wrinkling of the photoaged skin. In another study, Khan et al. applied same formulation on the hyperpigmented human skin and observed decreased melanin level, sebum production, and erythema of the treated skin. These antiaging effects on skin could be attributed to flavonoid, quercetin, and hesperetin, present in *Malus domestica* extract [46].

10.14. *Matricaria chamomilla L*. After skin treatment with water-in-oil emulsion based cream loaded with hydroalcoholic extract of *Matricaria chamomilla* plant for eight weeks, reduced transepithelial water loss was observed. Since an increase in transepithelial water loss shows disruption of the stratum corneum and loss of intercellular lipids, accordingly this formulation possibly repairs the stratum corneum and improves the moisture contents in human skin [47]. Moreover, there was an improvement in skin mechanical properties, that is, diminished levels of roughness, scaliness, smoothness, and wrinkling of photoaged skin [110, 111]. The investigators attributed this antiaging activity to the phenolic compounds present in extract. *Matricaria chamomilla*
belongs to Asteraceae family and contains some bioactive ingredients including terpenes, polysaccharides, and flavonoids such as α-bisabolol and apigenin.

10.15. *Moringa oleifera*. *Moringa oleifera* (Moringaceae) contains carotene, vitamin C, vitamin B, vitamin A, carotenoids, myricetin, quercetin, kaempferol, gallic acid, syringic acid, and rutin. All parts of this plant, from root to leaves, possess various medicinal activities such as antibacterial, anticancer, and antioxidant [112, 113]. Ali et al. prepared water-in-oil cream of hydroalcoholic extract of *Moringa oleifera* leaves and applied it to photoaged skin [48, 49]. The investigators found reduced undesirable skin sebum contents of skin and diminished skin transepidermal water loss leading to increased skin hydration, particularly for dry skin using Sebumeter and Corneometer, respectively. In addition, same formulation was also found effective against skin wrinkles, roughness, and scaliness leading to skin revitalization effect [50]. They tagged the antiaging characteristic of *Moringa oleifera* to the phenolic compounds present in extract, since the phenolics have capability to scavenge ROS.

10.16. *Morus alba*. *Morus alba* belongs to family Moraceae. This plant is rich in active compounds including anthocyanins, gallic acid, flavonoids and tannins, citric acid, vitamin C, and palmitic acid [114, 115]. Akhtar et al. prepared oil-in-water cream of hydroalcoholic extract of *Morus alba* fruit followed by application to the photoaged skin of the human volunteers [53]. After 8 weeks, the studied skin areas were tested using Mexameter and Corneometer. The results indicated the reduction in melanin contents of skin, without producing erythema, attributing this activity to the presence of anthocyanin and flavonoids. These phenolics have tyrosinase inhibition activity, one of the modes of antiaging activity [51, 52, 116]. In another study, Akhtar et al. reported decrease in erythema and melanin contents in the treated skin using same formulation [52].

10.17. *Ocimum basilicum*. Rasul and Akhtar reported the improvement in facial skin mechanical (viscoelasticity) and biochemical parameters (superoxide dismutase, catalase, total protein, and ascorbic acid level) when skin was treated with water-in-oil emulsion based cream loaded with ethanolic extract of *Ocimum basilicum* (Lamiaceae) seeds [53]. Moreover, the reduction in malondialdehyde level was also noted. The possible antiaging effect was pointed out as a feature of antioxidants such as anthocyanins, ellagic acid, and hydrolysable tannins present in extract.

10.18. *Oryza sativa*. Some strong antioxidants including ferulic acid, gamma-oryzanol, and phytic acid are present in *Oryza sativa* (Poaceae). The grains extract of *Oryza sativa* was loaded to niosomes followed by the preparation of water-in-oil cream using these niosomes. This cream was found effective for reducing transepidermal water loss, as observed by using Corneometer and Evaporimeter. Since transepidermal water loss plays crucial role in aging process by enhancing skin hydration, this formulation can be employed as an antiaging product. This feature of cream could be due to strong antioxidants present in *Oryza sativa* extract [54].

10.19. *Polygonum minus*. Haris et al. reported the improvement in facial skin elasticity as well as reduced wrinkles after treating with water-in-oil emulsion based cream loaded with aqueous extract of *Polygonum minus* (Polygonaceae) seeds [55]. The possible antiaging effect could be due to the antioxidants such as caffeic acid and quercetin [118].

10.20. *Punica granatum*. Kaur and Saraf reported the improvement in facial skin mechanical (viscoelasticity) and biochemical parameters (catalase and ascorbic acid concentration) [56]. Moreover, the reduction in malondialdehyde level was also noted. These results indicate the antiaging effect of nanotransfersomes loaded cream. The investigators used nanotransfersomes loaded with ethanolic extract of *Punica granatum* (Punicaceae) seeds to prepare a novel cream beside development of conventional cream. The antiaging effect of various formulations was in this decreasing order: nanotransfersomal cream > conventional cream > blank nanotransfersomal cream > base cream. The possible antiaging effect was pointed out as a feature of antioxidants such as anthocyanins, ellagic acid, and hydrolysable tannins present in extract.

10.21. *Silibum marianum*. On application of water-in-oil emulsion based cream loaded with ethanolic extract of *Silibum marianum* (Asteraceae) seeds for eight months, reduced transepithelial water loss was observed which resulted in improved moisture contents in human skin [57]. Moreover, there was an improvement in skin mechanical properties, that is, reduced levels of roughness, scaliness, smoothness, and wrinkles of photoaged skin. In another study, Rasul et al. applied same formulation on hyperpigmented human skin and observed the decreased melanin level, sebum production, and erythema of the treated skin [58].

10.22. *Tagetes erecta* Linn. *Tagetes erecta* Linn. (Asteraceae) contains quercetagetin, syringic acid, lutin, quercetin, and gallates. This plant possesses different medicinal activities including antiaging, anticancer, and anti-inflammatory [119–121]. Leelapornpisid et al. prepared ethyl acetate extract of *Tagetes erecta* flowers and loaded it to nanostructured lipid carriers [59]. These carriers were then formulated as cream and applied to photoaged skin. Using Visiometer, the researchers observed reduction in skin wrinkles of photoaged skin without producing skin irritation after eight weeks of cream usage. The investigators attributed this antiaging activity to the antioxidants present in extract.

10.23. *Terminalia chebula*. After skin treatment with water-in-oil emulsion based cream loaded with hydroalcoholic extract of *Terminalia chebula* plant for eight weeks, reduced transepithelial water loss was observed. Moreover, there was decrease in skin melanin contents also [60]. These features could be due to the phenolic compounds present in extract.
Terminalia chebula belongs to Combretaceae family and contains some bioactive ingredients including gallic acid, ellagic acid, tannic acid, ethyl gallate, chebulic acid, chebularic acid, corilagin, and ascorbic acid [122].

10.24. Trigonella foenum-graecum. Trigonella foenum-graecum belongs to family Fabaceae. This medicinal plant is rich in active compounds including polyphenols, galactomannans and flavonoid, protodioscin, trigonoside, diosgenin, and yamogenin. Galactomannan has an advantage of improving skin hydration [123]. Waqas et al. and Akhtar et al. prepared water-in-oil cream of methanol extract of Trigonella foenum-graecum seeds [61, 62]. After applying this cream to human skin for predetermined time, the former authors observed an improvement in facial skin mechanical parameters without producing erythema, whereas the latter authors reported reduction in skin melanin contents and maintenance of skin hydration.

10.25. Vitis vinifera. Various bioactive compounds such as sarmentine are present in Vitis vinifera (Vitaceae). The hydroalcoholic shoot extract of Vitis vinifera was formulated as water-in-oil cream that was found effective for improving clinical signs of photoaged skin. This property of cream could be due to the presence of strong antioxidants in Vitis vinifera extract [63].

11. Conclusion

Due to constant exposure of human skin to the UV radiations present in sunlight, several pathobiological alterations in cells occur. The photoprotection is the main approach for managing the photoaging, but cosmeceuticals could also be used as an alternative therapy. The selection of therapeutic approaches depends on the nature of these injurious molecular changes. Large number of botanical extract loaded creams have been prepared and assessed (Table 1) for their antiaging potential. The observed antiaging effects of cream formulations could be an outcome of a coordinating action of multiple constituents. Of numerous botanicals, the phenolic acids and flavonoids appear effective against UV radiation-induced damage; however, the evidence-based studies for their antiaging effects are still needed. Since environment affects the skin-cream interaction, the cautious assessment of their clinical efficacy should be conducted in future.

Conflict of Interests

The authors declare that there is no conflict of interests regarding the publication of this paper.

References

[1] B. A. Gilchrest, “Skin aging and photoaging: an overview,” Journal of the American Academy of Dermatology, vol. 21, no. 3, pp. 610–613, 1989.

[2] I. M. Braverman and E. Fonferko, “Studies in cutaneous aging. I. The elastic fiber network,” Journal of Investigative Dermatology, vol. 78, no. 5, pp. 434–443, 1982.

[3] W. Montagna, S. Kirchner, and K. Carlisle, “Histology of sun-damaged human skin,” Journal of the American Academy of Dermatology, vol. 21, no. 5, pp. 907–918, 1989.

[4] R. Warren, V. Gartstein, A. M. Kligerman, W. Montagna, R. A. Allendorf, and G. M. Ridder, “Age, sunlight, and facial skin: a histologic and quantitative study,” Journal of the American Academy of Dermatology, vol. 25, no. 5 I, pp. 751–760, 1991.

[5] D. R. Bielenberg, C. D. Bucana, R. Sanchez, C. K. Donawho, M. L. Kripke, and I. J. Farid, “Molecular regulation of UVB-induced cutaneous angiogenesis,” Journal of Investigative Dermatology, vol. 111, no. 5, pp. 864–872, 1998.

[6] K. Yano, H. Oura, and M. Detmar, “Targeted overexpression of the angiogenesis inhibitor thrombospondin-1 in the epidermis of transgenic mice prevents ultraviolet-B-induced angiogenesis and cutaneous photo-damage,” Journal of Investigative Dermatology, vol. 118, no. 5, pp. 800–805, 2002.

[7] J. Y. Seo and J. H. Chung, “Thermal aging: a new concept of skin aging,” Journal of Dermatological Science Supplement, vol. 2, pp. S13–S22, 2006.

[8] S. L. Mera, C. R. Lovell, R. Russell Jones, and J. D. Davies, “Elastin fibres in normal and sun-damaged skin: an immunohistochemical study,” British Journal of Dermatology, vol. 117, no. 1, pp. 21–27, 1987.

[9] M. Udompatakul, P. Sripigor, and P. Palungwachira, “An oral nutraceutical containing antioxidants, minerals and glycosaminoglycans improves skin roughness and fine wrinkles,” International Journal of Cosmetic Science, vol. 31, no. 6, pp. 427–435, 2009.

[10] G. J. Fisher, Z. Q. Wang, S. C. Datta, J. Varani, S. Kang, and J. J. Voorhees, “Pathophysiology of premature skin aging induced by ultraviolet light,” The New England Journal of Medicine, vol. 337, no. 20, pp. 1419–1428, 1997.

[11] J. Varani, R. L. Warner, M. Gharaee-Kermani et al., “Vitamin A antagonizes decreased cell growth and elevated collagen-degrading matrix metalloproteinases and stimulates collagen accumulation in naturally aged human skin,” Journal of Investigative Dermatology, vol. 114, no. 3, pp. 480–486, 2000.

[12] G. J. Fisher, S. C. Datta, H. S. Talwar et al., “Molecular basis of sun-induced premature skin ageing and retinoid antagonism,” Nature, vol. 379, no. 6563, pp. 335–339, 1996.

[13] J. H. Chung, J. Y. Seo, H. R. Choi et al., “Modulation of skin collagen metabolism in aged and photoaged human skin in vivo,” Journal of Investigative Dermatology, vol. 117, no. 5, pp. 1218–1224, 2001.

[14] H. C. Jin, Y. S. Jin, K. L. Mi et al., “Ultraviolet modulation of human macrophage metalloelastase in human skin in vivo,” Journal of Investigative Dermatology, vol. 119, no. 2, pp. 507–512, 2002.

[15] V. Cheynier, “Polyphenols in foods are more complex than often thought,” The American Journal of Clinical Nutrition, vol. 81, no. 1, pp. 223S–229S, 2005.

[16] H. Masaki, “Role of antioxidants in the skin: anti-aging effects,” Journal of Dermatological Science, vol. 58, no. 2, pp. 85–90, 2010.

[17] R. M. Lavker, “Cutaneous aging: chronologic versus photoaging,” in Photodamage, B. Gilchrest, Ed., vol. 1, pp. 123–135, Blackwell, 1995.

[18] Y. H. Kim, K. S. Kim, C. S. Han et al., “Inhibitory effects of natural plants of Jeju Island on elastase and MMP-1 expression,” International Journal of Cosmetic Science, vol. 29, pp. 487–488, 2007.
[19] C. S. Sander, H. Chang, S. Salzmann et al., “Photoaging is associated with protein oxidation in human skin in vivo,” *Journal of Investigative Dermatology*, vol. 118, no. 4, pp. 618–625, 2002.

[20] Y. Shindo, E. Witt, and L. Packer, “Antioxidant defense mechanisms in murine epidermis and dermis and their responses to ultraviolet light,” *Journal of Investigative Dermatology*, vol. 100, no. 3, pp. 260–265, 1993.

[21] F. McArdle, L. E. Rhodes, R. Parslew, C. I. A. Jack, P. S. Friedmann, and M. J. Jackson, “UVR-induced oxidative stress in human skin: effects of oral vitamin C supplementation,” *Free Radical Biology and Medicine*, vol. 33, no. 10, pp. 1355–1362, 2002.

[22] B. M. Babior, J. D. Lambeth, and W. Nauseef, “The neutrophil NADPH oxidase,” *Archives of Biochemistry and Biophysics*, vol. 397, no. 2, pp. 342–344, 2002.

[23] D. N. Granger, “Role of xanthine oxidase and granulocytes in ischemia-reperfusion injury,” *American Journal of Physiology*, vol. 255, no. 6, part 2, pp. H1269–H1275, 1988.

[24] A. G. Fantel, R. E. Person, R. W. Tumbic, T.-D. Nguyen, and B. Mackler, “Studies of mitochondria in oxidative embryotoxicity,” *Teratology*, vol. 52, no. 4, pp. 190–195, 1995.

[25] M. Comporti, “Lipid peroxidation and biogenic aldehydes: from the identification of 4-hydroxynonenal to further achievements in biopathology,” *Free Radical Research*, vol. 28, no. 6, pp. 623–635, 1998.

[26] C. F. Nathan and J. B. Hibbs Jr., “Role of nitric oxide synthesis in macrophage antimicrobial activity,” *Current Opinion in Immunology*, vol. 3, no. 1, pp. 65–70, 1991.

[27] A. Ali, N. Akhtar, and H. M. S. Khan, “Enhancement of human cheek skin texture by *Acacia nilotica* bark extract cream,” *Tropical Journal of Pharmaceutical Research*, vol. 12, no. 3, pp. 323–327, 2013.

[28] V. Sabale, H. Kunjwani, and P. Sabale, “Formulation and *in vitro* evaluation of the topical antiaging preparation of the fruit of *Benincasa hispida*,” *Journal of Ayurveda and Integrative Medicine*, vol. 2, no. 3, pp. 124–128, 2011.

[29] N. Akhtar, S. U. Zaman, B. A. Khan, M. N. Amir, and M. A. Ebrahimzadeh, “Calendula extract: effects on mechanical parameters of human skin,” *Acta Poloniae Pharmaceutica—Drug Research*, vol. 68, no. 5, pp. 693–701, 2011.

[30] N. Akhtar, S. U. Zaman, B. A. Khan et al., “Evaluation of various functional skin parameters using a topical cream of *Calendula officinalis* extract,” *African Journal of Pharmacy and Pharmacology*, vol. 5, no. 2, pp. 199–206, 2011.

[31] T. Mahmood, N. Akhtar, B. A. Khan, H. M. S. Khan, and T. Saeed, “Outcomes of 3% green tea emulsion on skin sebum production in male volunteers,” *Bosnian Journal of Basic Medical Sciences*, vol. 10, no. 3, pp. 260–264, 2010.

[32] T. Mahmood, N. Akhtar, B. A. Khan, M. Ahmad, H. M. S. Khan, and S. U. Zaman, “Application of stable green tea extract cream on human cheeks,” *International Journal of Academic Research*, vol. 2, no. 2, pp. 121–126, 2010.

[33] T. Mahmood and N. Akhtar, “Combined topical application of lotus and green tea improves facial skin surface parameters,” *Rejuvenation Research*, vol. 16, no. 2, pp. 91–97, 2013.

[34] T. Mahmood, N. Akhtar, and C. Moldovan, “A comparison of the effects of topical green tea and lotus on facial sebum control in healthy humans,” *Hippokratia*, vol. 17, no. 1, pp. 64–67, 2013.

[35] T. Mahmood and N. Akhtar, “Short term study of human skin irritation by single application closed patch test: assessment of four multiple emulsion formulations loaded with botanical extracts,” *Cutaneous and Ocular Toxicology*, vol. 32, no. 1, pp. 35–40, 2013.

[36] S. U. Zaman, N. Akhtar, B. A. Khan et al., “Development of a sebum control cream from a local desert plant *Capparis decidua*,” *Journal of Medicinal Plants Research*, vol. 6, no. 5, pp. 744–748, 2012.

[37] I. F. Almeida, J. Maleckova, R. Saffi et al., “Characterization of an antioxidant surfactant-free topical formulation containing *Castanea sativa* leaf extract,” *Drug Development and Industrial Pharmacy*, vol. 41, no. 1, pp. 148–155, 2013.

[38] D. H. McDaniel, “Clinical safety and efficacy in photoaged skin with CoffeeBerry extract, a natural antioxidant,” *Cosmetic Dermatology*, vol. 22, no. 12, pp. 610–616, 2009.

[39] N. Akhtar, H. M. Khan, S. Ashraf, I. S. Mohammad, N. Saqib, and K. Bashir, “Moisturizing effect of stable cream containing *Crocus sativus* extracts,” *Pakistan Journal of Pharmaceutical Sciences*, vol. 27, no. 6, pp. 1881–1884, 2014.

[40] N. Akhtar, A. Zaman, A. Ali et al., “Effects of *Emblica officinalis* extract cream on human skin trans-epidermal water loss measured with non invasive probe,” *Journal of Pharmacy and Alternative Medicine*, vol. 1, no. 1, pp. 32–37, 2012.

[41] A. Rasul, N. Akhtar, B. A. Khan, T. Mahmood, S. Uz Zaman, and H. M. Shoab Khan, “Formulation development of a cream containing fennel extract: in vivo evaluation for anti-aging effects,” *Pharmazie*, vol. 67, no. 1, pp. 54–58, 2012.

[42] A. Rasul, N. Akhtar, M. Iqbal et al., “Sebumetric and mexametric evaluation of a fennel based cream,” *ScienceAsia*, vol. 38, no. 3, pp. 262–267, 2012.

[43] A. Ali Khan, N. Akhtar, and V. A. Braga, “Anti-aging effects of *Hippophae rhamnoides* emulsion on human skin,” *Tropical Journal of Pharmaceutical Research*, vol. 11, no. 6, pp. 955–962, 2012.

[44] M.-J. Chang, H.-C. Huang, H.-C. Chang, and T.-M. Chang, “Cosmetic formulations containing *Lithospermum erythrorhizon* root extract show moisturizing effects on human skin,” *Archives of Dermatological Research*, vol. 300, no. 6, pp. 317–323, 2008.

[45] H. M. S. Khan, N. Akhtar, F. Rasool, B. A. Khan, T. Mahmood, and M. S. Khan, “In vivo evaluation of stable cream containing flavonoids on hydration and TEWL of human skin,” *World Academy of Science, Engineering and Technology*, vol. 47, pp. 896–899, 2010.

[46] H. M. Shoab Khan, N. A. F. Rasool, A. Madni, T. Saeed, B. A. Khan, and T. Mahmood, “Investigation of a new sebum control cream containing apple juice extract,” *Asian Journal of Chemistry*, vol. 23, no. 2, pp. 810–812, 2011.

[47] A. T. Nóregra, T. A. L. Wagemaker, and P. M. B. G. M. Campos, “Antioxidant activity of *Matricaria chamomilla* L. extract and clinical efficacy of cosmetic formulations containing this extract and its isolated compounds,” *Biomedical and Biopharmaceutical Research*, vol. 10, no. 2, pp. 249–261, 2013.

[48] A. Ali, N. Akhtar, M. S. Khan, T. Khan, A. Ullah, and M. I. Shah, “Effect of *Moringa oleifera* on undesirable skin sebum secretions of sebaceous glands observed during winter season in humans,” *Biomedical Research*, vol. 24, no. 1, pp. 127–130, 2013.

[49] A. Ali, N. Akhtar, M. S. Khan et al., “Moisturizing effect of cream containing *Moringa oleifera* (Sohajana) leaf extract by biophysical techniques: in vivo evaluation,” *Journal of Medicinal Plants Research*, vol. 7, no. 8, pp. 386–391, 2013.
[116] Z.-R. Yu, C.-C. Hung, Y.-M. Weng, C.-L. Su, and B.-J. Wang, "Physiochemical, antioxidant and whitening properties of extract from root cortices of mulberry as affected by membrane process," *LWT—Food Science and Technology*, vol. 40, no. 5, pp. 900–907, 2007.

[117] S. K. Marwat, M. S. Khan, S. Ghulam, N. Anwar, G. Mustafa, and K. Usman, “Phytochemical constituents and pharmacological activities of sweet *Basil-Ocimum basilicum* L. (Lamiaceae),” *Asian Journal of Chemistry*, vol. 23, no. 9, pp. 3773–3782, 2011.

[118] M. Maizura, A. Aminah, and W. M. Wan Aida, “Total phenolic content and antioxidant activity of kesum (*Polygonum minus*), ginger (*Zingiber officinale*) and turmeric (*Curcuma longa*) extract,” *International Food Research Journal*, vol. 18, pp. 529–534, 2011.

[119] R. M. Pérez Gutiérrez, H. H. Luna, and S. H. Garrido, “Antioxidant activity of *Tagetes erecta* essential oil,” *Journal of the Chilean Chemical Society*, vol. 51, no. 2, pp. 883–886, 2006.

[120] W. Li, Y. Gao, J. Zhao, and W. Qi, “Phenolic, flavonoid, and lutein ester content and antioxidant activity of 11 cultivars of Chinese marigold,” *Journal of Agricultural and Food Chemistry*, vol. 55, no. 21, pp. 8478–8484, 2007.

[121] A. Phrutivorapongkul, K. Kiattisin, P. Jantrawut, S. Chansakaow, S. Vejabhikul, and P. Leelapornpisid, “Appraisal of biological activities and identification of phenolic compound of African marigold (*Tagetes erecta*) flower extract,” *Pakistan Journal of Pharmaceutical Sciences*, vol. 26, no. 6, pp. 1071–1076, 2013.

[122] A. K. Mishra and P. Chattopadhyay, "Herbal cosmeceuticals for photoprotection from ultraviolet B radiation: a review," *Tropical Journal of Pharmaceutical Research*, vol. 10, no. 3, pp. 351–360, 2011.

[123] T. Aburjai and F. M. Natsheh, “Plants used in cosmetics,” *Phytotherapy Research*, vol. 17, no. 9, pp. 987–1000, 2003.