


Review

# Biochemical Properties and Cosmetic Uses of *Commiphora myrrha* and *Boswellia serrata*

Bssmah Ghazi Alraddadi<sup>1</sup> and Hyun-Jae Shin<sup>1,2,\*</sup> 

<sup>1</sup> Department of Beauty and Cosmetology, Graduate School of Industrial Technology Startup, Chosun University, 309 Pilmun-daero, Dong-gu, Gwangju 61452, Korea

<sup>2</sup> Department of Biochemical Engineering, Chosun University, 309 Pilmun-daero, Dong-gu, Gwangju 61452, Korea

\* Correspondence: shinhj@chosun.ac.kr

**Abstract:** Organic materials have been the subject of numerous recent cosmetics studies. Plant extracts and/or plant-derived resources in the Middle East have significant potential in many cosmetic fields because they exhibit useful properties that promote the repair of cellular functions and improve skin conditions. Multiple organic substances from these resources have long-lasting environmental effects, extensive storage capacities, and affect human physiological activity. Recent studies have revealed that *Commiphora myrrha* (myrrh) and *Boswellia serrata* (frankincense) extracts can prevent aging, acne, and spots while enhancing skin moisture and suppleness. Given that myrrh and frankincense exhibit anti-inflammatory and antioxidant characteristics that significantly improve human health, their utilization in various cosmetic products has received attention from global cosmetic corporations. This review details the bioactive compounds and cosmeceutical properties of myrrh and frankincense and their use in cosmetic formulations.

**Keywords:** *Commiphora myrrha*; *Boswellia serrata*; frankincense; cosmetics; antioxidant; antiaging



**Citation:** Alraddadi, B.G.; Shin, H.-J. Biochemical Properties and Cosmetic Uses of *Commiphora myrrha* and *Boswellia serrata*. *Cosmetics* **2022**, *9*, 119. <https://doi.org/10.3390/cosmetics9060119>

Academic Editor: Lucia Panzella

Received: 25 October 2022

Accepted: 10 November 2022

Published: 12 November 2022

**Publisher's Note:** MDPI stays neutral with regard to jurisdictional claims in published maps and institutional affiliations.



**Copyright:** © 2022 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (<https://creativecommons.org/licenses/by/4.0/>).

## 1. Introduction

Since antiquity, plants have been utilized in cosmetics and have been the focus of modern scientific research. Drawing increasingly complex images has become possible because of the improved understanding of the skin and plants over time. Plants are sophisticated organisms that respond to their surroundings by producing various metabolites. Phyto-molecules applied to the skin affects skin cells and influences the health and appearance of the skin. Physicochemical and ethnobotanical studies have identified various plants with the potential to improve contemporary cosmetic products [1]. Plants provide high-value active compounds for medicinal and aesthetic purposes. Targeted at improving skin problems and appearance, humans have utilized plants for thousands of years, which have been developed into modern cosmetics. The skin serves as the body's interface to the exterior world and actively defends against threats. Therefore, the ability of plants to produce chemicals that can protect and calm the skin is essential. Modern cosmetics can adjust skin elasticity in addition to hydrating the skin and reducing redness [2].

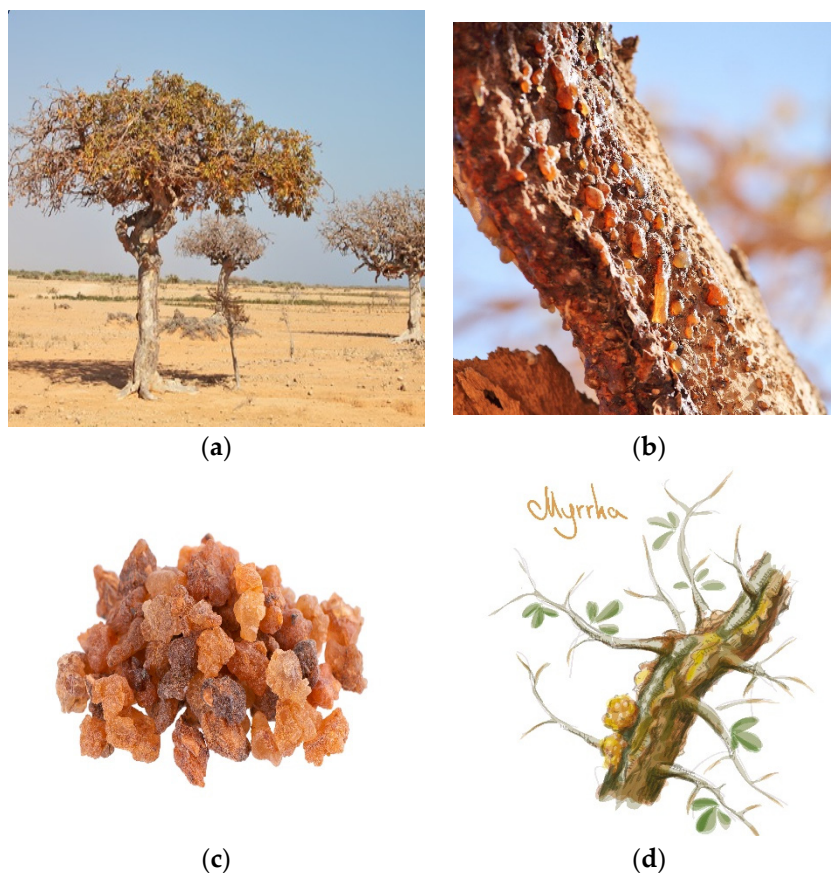
The family Burseraceae contains about 18 genera, an estimated 560–600 species [3,4]. Since antiquity, these species have been used for their aromatic properties and medicinal applications. Thus, their spruce trees and shrubs often play a prominent role in the ethnobotany of their surrounding environment. Although the Burseraceae species are spread worldwide, studies and information are limited to the genera in Asia and Africa, such as *Commiphora* (myrrh), *Canarium* (incense elemi), and *Boswellia* (frankincense). Indigenous tribes have employed the resins of this species for various uses in traditional medicine, including its application as an external agent (cosmetics), for treating wounds (dressings), preventing the deterioration of extremities and broken teeth, or as an antiseptic and disinfectant. Moreover, inhaling the smoke can induce anesthetic effects [5].

This review aims to introduce *C. Myrrha* and *B. Serrata* and determine their chemical composition, their activities as antioxidants, antimicrobials, and anti-inflammatories, as well as their influential role in promoting health in the human body. Specifically, and especially, their effective activity against acne and wrinkles, and many of the excellent effects and higher values of these plants in cosmetics.

## 2. *Commiphora myrrha*

The *Commiphora* species has high medical importance. It abounds mostly under mountains and in rocky settings in Saudi Arabia, notably east of Tihama [6,7]. *Commiphora* forms a particular element of Saudi flora [6], and its trees are spread on the sands of the Red Sea, west and southwest of the Kingdom of Saudi Arabia (KSA). They have a wide ecological range between 1 and 2100 m above sea level and are found in tropical and semi-tropical areas [6–9]. They are also found in Somalia [6–11]. Some species are located mainly in Africa (specifically in Ethiopia, Somalia, and Kenya) [10] and Asia (specifically India [6,8,12] and Pakistan [8]).

The family Burseraceae, including the *Commiphora* species, are trees or shrubs with vertical resinous channels protruding from the bark. The leaves are compounded, spirally organized, and densely packed at the twig tips. Small and solitary (or in groups), the flowers grow on the branches [3] (Figure 1). *Commiphora myrrha* is a reddish-brown oligomycin known as one of the oldest medicines used by humanity, as it dates back to 2800 BC, according to the medical texts of ancient Egypt [13–16].



**Figure 1.** (a) *Commiphora myrrha* tree; (b) resin appears from the tree trunk; (c) *C. Myrrha* resin; (d) illustration of *C. Myrrha* plant. All pictures and illustrations do not have any copyright issues.

The Arabic term “Murr”, which means “Bitter”, describes the flavor and balsamic aroma of myrrh [17]. Myrrh is an aromatic resin extracted from the *Commiphora* tree. Historically, myrrh was extremely expensive, as it cost more than gold. The most outstanding physician in the Middle Ages, the Muslim physician Al-Razi, used the myrrh plant to

treat kidney and bladder diseases and stomach bloating [12]. Its essential oils (EOs) have been used to manufacture perfumes and cosmetics and in aromatherapy as incense and for medicinal purposes for centuries [12,18,19]. Medicinally, it has been utilized as an emmenagogue, antispasmodic, antiparasitic, antitussive, and astringent. Leprosy, syphilis, and other infectious disorders, as well as cancer, have all been cured by myrrh. It is also a component of traditional medical practices in China, the Middle East, and Africa [17] and has been used since antiquity for several medicinal applications [20,21].

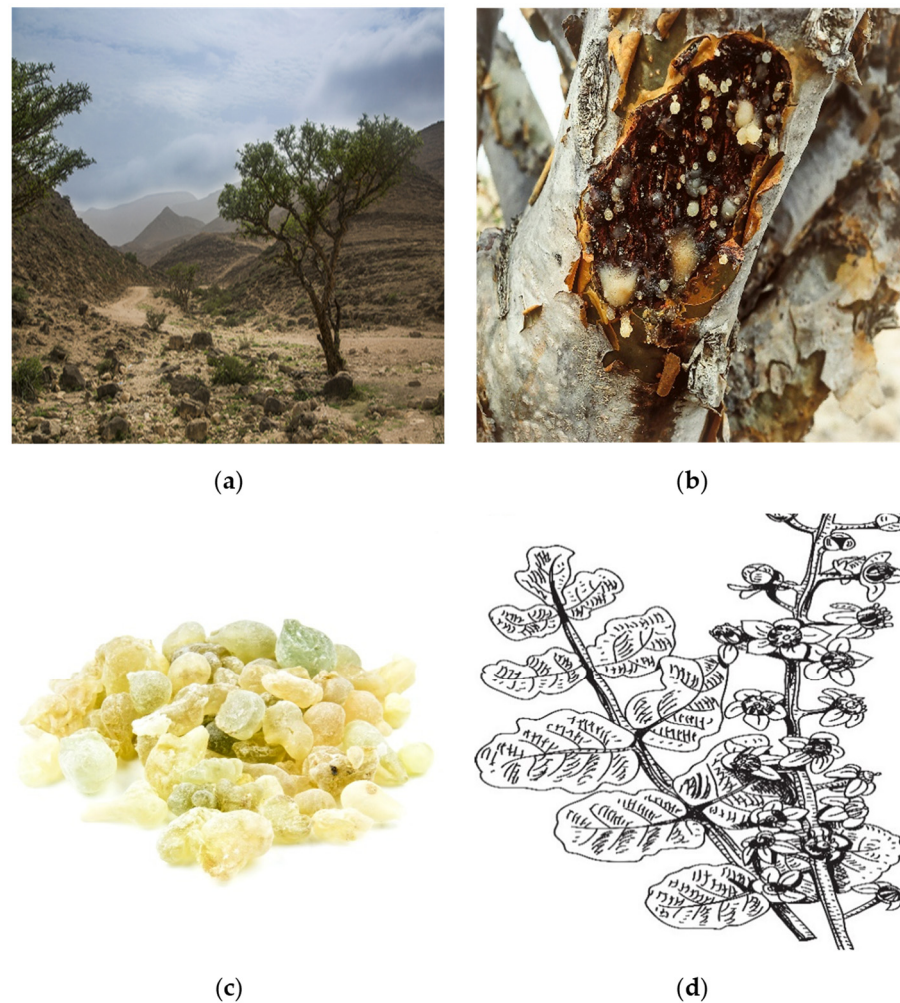
### 3. *Boswellia serrata*

*Boswellia serrata* resin, olibanum, or frankincense, is obtained from the Burseraceae family [22,23]. *Boswellia* is a relatively small genus with approximately 28 species [24]. Since antiquity, humans have employed medicinal plants, such as the Indian frankincense tree (*Boswellia serrata*), as medicines. *Boswellia serrata* is a medium-sized deciduous tree primarily found in India, Asia, and Africa. It has had a significant impact on curing various ailments [25–27] in Saudi Arabia [28], Southern Arabia, Oman [29–32], Pakistan [33], Nigeria, Yemen, and Somalia [31,34,35].

*Boswellia serrata* is a medium to large tree that grows up to 18 m tall and 2.4 m wide. The leaves are imparipinnate with opposing leaflets, alternating, packed at the terminals of the branches, and typically serrated. It is hermaphroditic; the flowers are tiny and white and grow in axillary racemes. Its thin, greenish-grey bark changes from yellow or red to deciduous petals that are 3–5 in number, imbricated, and eventually turn an ash color. The bark sheds in smooth, papery flakes of flaming red and secretes tiny drops of resin. After an injury or a natural break to the bark, exudate containing oleo gum resin is produced. The oleo gum resin is aromatic, clear, and golden yellow, which eventually transforms into crusts and brownish-yellow tears or drips [31,36–38] (Figure 2).

The therapeutic properties of *Boswellia serrata* have been established by numerous researchers and are well-acknowledged [38–42]. Its resin was recognized as one of the most significant medicinal herbs with exceptional efficacy in numerous therapeutic areas [43,44]. *B. Serrata* is a significant source of boswellic acid (BA) in the pharmaceutical sector. Paints and varnishes are made from high-quality turpentine from *B. Serrata*. However, the commercial applications of the *B. Serrata* oleo-primary resin are in the medication industry, western biomedicine, religious purposes, perfumes [45], and cosmetics [46]. The most significant medical applications for *B. Serrata* are the treatment of rheumatoid arthritis, osteoarthritis, Crohn's disease, and asthma [38,47–54]. It is also used to treat diseases like cough and bronchitis [37,55–57]. It is frequently prescribed in the Unani System of Medicine for skin conditions, corneal ulcers, osteoarthritis, dysentery, chronic inflammatory illnesses, wound healing, and diarrhea [36], used as an antiobesity [58] and anticonvulsant agent [59], and is beneficial in liver fibrosis [60] and several cancers [61–65]. *Boswellia* resin is used to prevent and treat colitis and ulcerative colitis. *B. Serrata* shows satisfactory antioxidant activity in the cerebrovascular system [66,67] and various inflammatory conditions [68–77] and exerts antipyretic, antiatherosclerotic, and analgesic activity [38,78–80] in inflammatory bowel disease, allergies, and arthritis, including osteoarthritis, and pain [81–85]. It has also been used as an antiseptic in mouthwashes [86] and for treating chronic inflammatory diseases [87–89]. Its acids have been evaluated for their ability to prevent the development of periodontal biofilm [90] or the prevention of infections of the skin and nails [91,92].

Additionally, it has biological effects, including antiulcer, antiulcerogenic, antibacterial, and psychopharmacological effects [35,93–98]. *Boswellia* extract was used in toxicological tests on animals, and the results indicated that it was safe to use in herbal remedies [99,100]. The principal active ingredient in BA is an aromatic compound with these therapeutic effects [42,101,102]. Moreover, the US Food and Drug Administration (USFDA) has acknowledged the safety of *Boswellia serrata* resin and authorized its use as an ingredient in feed additives [42].



**Figure 2.** (a) *Boswellia serrata* tree; (b) resin appears from the tree trunk; (c) *B. Serrata* resin; (d) Illustration of *B. Serrata* plant. All pictures and illustrations do not have any copyright issues.

Hindus, Babylonians, Persians, Romans, Chinese, Greeks, and residents of early American civilizations have all employed natural resins, chiefly for embalming and incense at ceremonial gatherings. They had a firm conviction that when these materials were burnt, the smoke and scent they release will appease their gods and comfort their souls. Burning these organic resins had become a significant aspect of their lifestyle; they burned them during sacrifice rites and daily rituals to protect their souls from the influence of evil spirits or to honor the deceased or living [103–105].

#### 4. Ingredients and Biological Activities of Myrrh and Frankincense

##### 4.1. Main Ingredients and Compounds in Myrrh and Frankincense

Phytochemical examination of different extracts of *C. Myrrha* and *B. Serrata* reveals major active chemical components in the EO, gums, and resins. The main features and compounds of *C. Myrrha* and *B. Serrata* extracts are summarized in Table 1.

**Table 1.** Main ingredients and compounds in *Commiphora myrrha* and *Boswellia serrata*.

| <i>Commiphora myrrha:</i>   | References   |
|---|--------------|
| Water-soluble gum, alcohol-soluble resins, polysaccharides, and proteins; steroids, sterols, and terpenes make up the volatile oil; and furanosesquiterpenes.   | [106]        |
| 3–8% essential oil, 30–60% water-soluble gum, and 25–40% alcohol-soluble resin.   | [16]         |
| Flavonoids, alkaloids, tannins, glycosides, steroids, saponins, and terpenoids. The methanolic extract is high in bioactive components, such as flavonoids and glycosides.  | [107]        |
| Myrrin, sesquiterpenes, aldehyde, eugenols, and protein.  | [7]          |
| $\alpha$ -, $\beta$ - and $\gamma$ -commiphoric acid; $\alpha$ - and $\beta$ -heerabomyrrhol; protocatechuic acid, and pyrocatechin   | [108]        |
| 2-Methoxy-5acetoxyfruranogermacr-1(10)-en-6-one and abietic acid.   | [109–111]    |
| Diterpenic acids, sesquiterpenes, 2-Methoxy-8,12-epoxygermacra-1(10) 7,11-triene-6-one, myrrhone, sandaracopimaric acid, abietic acid, dehydroabietic acid, and mansumbinone  | [112]        |
| <i>Boswellia serrata:</i>   |              |
| Arabinose, rhamnose, glucose, galactose, fructose, galacturonic acid, and $\beta$ sitosterol; phenol-o-cresol, m-cresol, p-cresol, thymol, and carvacrol; carboxylic acid- $\alpha$ -campholenic acid   | [113,114]    |
| 2,2,4-trimethylcyclopent-3-en-1-yl acetic acid, and campholytic acid.   |              |
| Portion (97.3%) in E- $\beta$ -ocimene and limonene, (2.7%) sesquiterpenes, in E-caryophyllene.   | [115]        |
| Resin acids (60–70%), water-soluble gum (about 20%), and monoterpene essential oil (3–10%)  | [116]        |
| Oil (45%), $\alpha$ -thujene (12%), $\alpha$ -pinene (8%), sabinene (2.2%), $\beta$ -pinene (0.7%), myrcene (3.8%), $\alpha$ -phellandrene (1%), pycmene (1%), limonene (1.9%), linalool (0.9%), perillene (0.5%), methylchavicol (11.6%), methyleugenol (2.1%), germacrene D (2.0%), kessane (0.9%), cembrene A (0.5%), and cembrenol (1.9%); a monoterpene 5,5-dimethyl-1-vinylbicyclo- hexane (2%), m-camphorene (0.7%); and p-camphorene (0.3%) | [117]        |
| Diterpenes, cembrenol (serratol), incensole, and incensole acetate  | [118,119]    |
| Lipophilic pentacyclic triterpene acids of the oleanane ( $\alpha$ -BAs), ursane-( $\beta$ -BAs) and lupane-type (lupeolic acids), and an ether-insoluble fraction polysaccharide (arabinose, galactose, xylose) soluble in water   | [38,120–123] |

## 4.2. Biological Function and Activity of the Plants

### 4.2.1. *Commiphora myrrha*

*Commiphora myrrha* exhibits antibacterial and antifungal properties and is high in chemicals that aid treatment [12]. Its ethanolic extracts exerted antibacterial action against microorganisms to some extent [107]. Myrthanol A, a monoterpene alcohol from myrrh EO, has been proven to be more effective than hydrocortisone at reducing inflammation [15,16,124,125]. Several studies have established the antimicrobial activities of myrrh, including fasciolocidal, antibacterial, antiviral, and antifungal activities [126–128]. These inherent antibacterial and antifungal properties are due to the sesquiterpenes and furanosesquiterpenoids it contains [129–131]. Myrrh preparations have been classified as weakly sensitizing in local lymph node assay (LLNA) studies [21,132]. Caryophyllene C15H24, in myrrh oil, has an antibacterial, antitumor, and anti-inflammatory activity [133].

### 4.2.2. *Boswellia serrata*

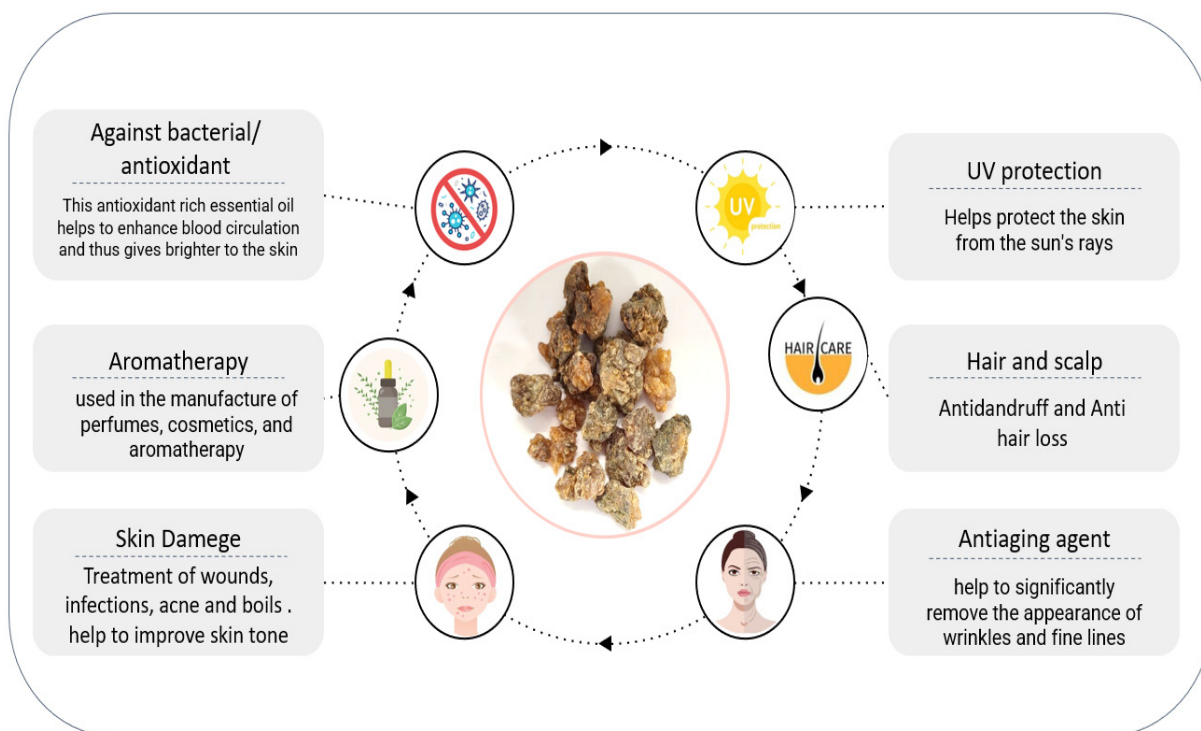
*B. Serrata* has been reported to possess anti-inflammatory and antiarthritic activities. The anti-inflammatory activity exerts 47% inhibitory action [30,134–136]. The analysis of the compounds of *B. Serrata* extracts revealed that its triterpenoids are effective against rheumatoid arthritis, chronic colitis, ulcerative colitis, skin allergies and ulcers, tumors, osteoarthritis, and inflammation [116,137]. BAs are a mixture of pentacyclic triterpenes acids, beta-BA, 3-acetyl beta BA, 11-ketobeta-BA, and 3-acetyl-11-keto-beta-BA, which have an essential role in its anti-inflammatory activity [63,81,138]. Various studies have reported that crude extracts of the *Boswellia* species contain an antimicrobial agent that inhibits microbial growth [139]. Silver nanoparticles prepared from an aqueous extract of *Boswellia* act as an antimicrobial agent on Gram-positive and Gram-negative bacteria [140]. Other studies have also disclosed that frankincense gum resin is active against *Staphylococcus aureus*, *Escherichia coli*, *Klebsiella* spp., *Pseudomonas aeruginosa*, *Proteus mirabilis*, and *Bacillus subtilis* [141]. In addition, concentrations of  $\alpha$ -pinene (38.41%) and myrcene (15.21%)

have been reported in *B. Serrata* EO, whereas *C. Myrrha* EO was distinguished by a high furanoeudesma-1,3-diene (17.65%) content, followed by curzerene (12.97%),  $\beta$ -element (12.70%), and germacrene- $\beta$  (12.15%), having both antimicrobial and antifungal activities [142]. The antimicrobial activity of *B. Serrata* extract has been demonstrated using disk diffusion or well plate and broth microdilution methods [49,90]. This activity has been investigated, and numerous therapeutic applications have revealed the pharmacological results, including many diseases, such as anti-inflammatory, antitumor, immunomodulatory, and inflammatory bowel diseases, in addition to its antioxidant activity [74,143]. Thus, it has been well established that *B. Serrata* possesses antibacterial and antifungal activity [44,144].

## 5. Cosmetic Applications of Myrrh and Frankincense

### 5.1. *Commiphora myrrha*

*Commiphora myrrha* is used to treat wounds, infections, acne, boils [21,145], hair, and the scalp [146,147]. It has been used to manufacture perfumes and cosmetics and used in aromatherapy [12]. The extract affected chapped and irritated skin types [148] and improved skin tone, firmness, and elasticity [149], and had a UV protection effect [150] (Figure 3).

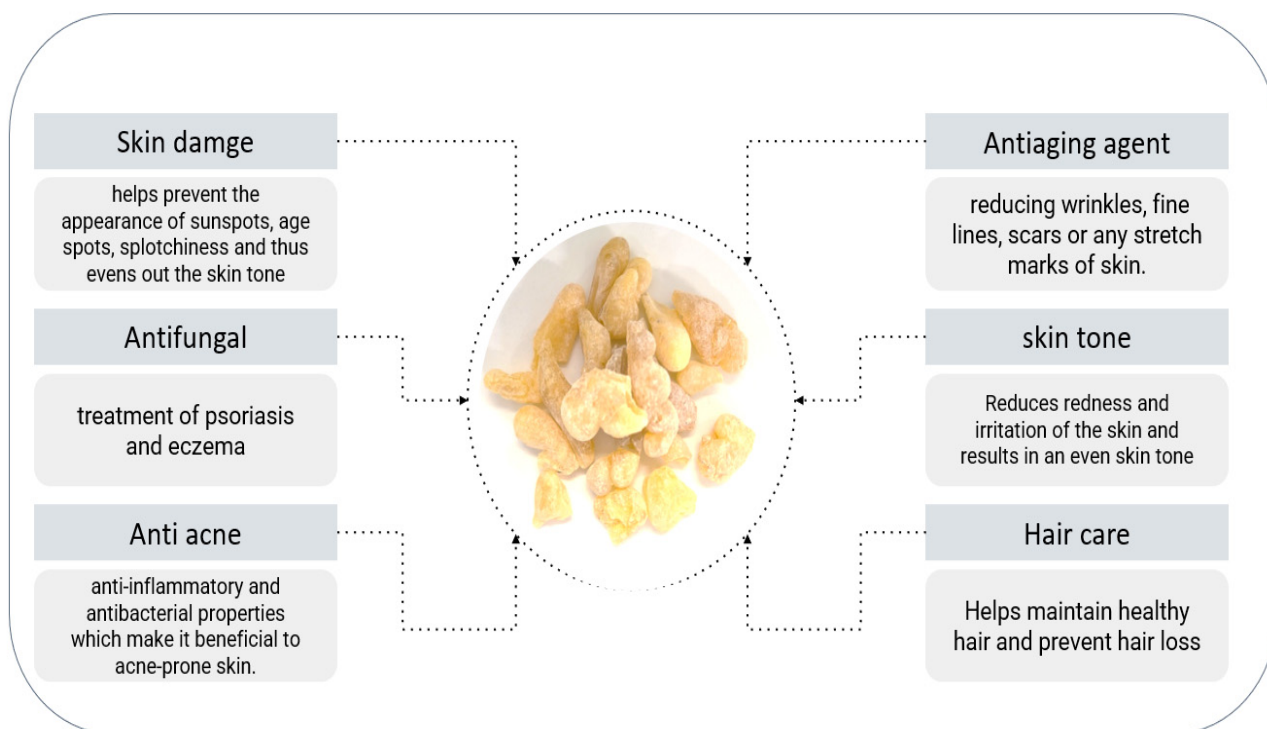


**Figure 3.** The illustrated summary of the cosmetic uses of *C. Myrrha*.

### 5.2. *Boswellia serrata*

Frankincense EO has anti-inflammatory and antibacterial properties for all skin types, such as acne and blemish-prone skin, aging skin, and dry skin, among others, making it particularly beneficial to acne-prone skin. It alleviates oily and acne-prone skin and provides lipids for aged skin. It is also a natural toner that improves skin tone and removes pores. It is a powerful astringent and is exceedingly effective for reducing wrinkles, fine lines, scars, or stretch marks on the skin. Frankincense EO promotes new cell generation, maintains skin elasticity, and soothes dry and chapped skin [151]. It lessens or prevents the appearance of sunspots, age spots, and splotches, thus evening out the skin tone [152]. It reduces skin redness and irritation, further promoting an even skin tone [153]. It has been used as a skin remedy for bruises and infected sores [154]. It is a potent antiwrinkle

and antiaging agent [155] and can be used for treating psoriasis, eczema [49,156], and skin diseases and as an antihair-loss remedy [38,50,51,53,54] (Figure 4).



**Figure 4.** The illustrated summary of the cosmetic uses of *B. Serrata*.

## 6. Value-Added Cosmetic Products from *C. Myrrha* and *B. Serrata*

Since antiquity, humans have mastered the art of exploring nature to obtain vital ingredients for meeting their fundamental requirements and care for their diseases and cosmetology. Many cultures utilize natural or synthetic cosmetics to improve their skin condition and facial attractiveness. “Cosmetics” originates from the Greek term “Kosmos”, meaning to adorn, and has referred to any substance intended to enhance or beautify. The botanical ingredient of a cosmetic or personal care product is derived from plants (herbs, roots, flowers, fruits, leaves, or seeds). Many plants are employed in the cosmetics industry in many Middle Eastern countries [157]. *C. Myrrha* and *B. Serrata* have several applications in cosmetics, ranging from antiaging and soothing to anti-inflammatory, encouraging faster skin renewal and increasing skin elasticity as a skin brightener. They have been used in antiaging skincare products; both plants have hit the Western market for skincare applications. The plant extracts and oils are also used for topical applications as they confer skincare benefits. Some of the cosmetic products containing *C. Myrrha* and *B. Serrata* extracts are summarized in Tables 2 and 3. In the following section, we will focus on the antioxidant, antiaging, and skin-whitening activities of *C. Myrrha* and *B. Serrata*.

**Table 2.** Cosmetic products containing *C. Myrrha*.

| Product Name  | <i>C. Myrrha</i> /Extract Included | Function  |
|---|------------------------------------|---|
| Complexion Polish—Natural Face Exfoliant, Sydney, Australia   | Resin Extract                      | This technique helps to purify the skin and remove toxins. It is highly effective on mature skin because it helps in cell renewal |
| Knight Co.—Repairing Face Oil, Los Angeles, USA   | Resin Extract                      | Antiaging.  |
| Hey honey—911 PRO GEL, Sarasota, Florida, USA   | Resin Extract                      | Soothe and correct acne, inflammation, and scarring.  |
| Beb organic—Soothing Serum, Simi Valley, California, USA  | Resin Extract                      | Treat spots from acne and damaged skin.   |
| SheaMoisture—Raw shea butter face & body bar soap, Amityville, New York, USA                        | Frankincense and Myrrh Extract     | Glowing skin.   |
| Hey honey—Don't miss a spot, Sarasota, Florida, USA   | Resin Extract                      | Natural antiseptic, astringent, and disinfectant.   |
| Biologique Recherche—Gommage P50 Corps, Paris, France   | Resin Extract                      | Replenishes the epidermis, antioxidant.   |
| Biologique Recherche—Masque Gommage Mains, Paris, France  | Resin Extract                      | Treatment of pigmentation.  |
| Biology Recherche—Lotion P50, Paris, France   | Resin Extract                      | Brightening.  |
| Biologique Recherche—Creme Grand Millesime, Paris, France   | Resin Extract                      | Antioxidant.  |
| Biologique Recherche—Serum Renovateur 14, Paris, France   | Resin Extract                      | Antioxidant.  |
| SheaMoisture—Raw Shea Butter Hydrating Moisturizer W/Frankincense & Myrrh Amityville, New York, USA | Resin Extract                      | Anti-inflammatory, antibacterial and antiaging properties.  |
| Biologique Recherche—Le Grand Serum, Paris, France  | Resin Extract                      | Antiaging.  |
| Bliss—Pore Patrol Oil Free Hydrator With Willow Bark, New York, USA                                 | Resin Extract                      | Antimicrobial and antibacterial.  |
| Biologique Recherche—Masque Creme Biofixine, Paris, France  | Resin Extract                      | Antiwrinkle and muscle-relaxing.  |
| Sunday 2 Sunday—Root Refresh Micellar Rinse, Airtech Pkwy, Plainfield, USA                          | Resin Extract                      | Cleans the hair and scalp.  |
| Colleen Rothschild Beauty—Restorative Hand Cream, Texas, USA  | Resin Extract                      | Soothes the skin.   |
| Kiss the Moon—Glow Night-Time Face Oil, Richmond, UK  | Resin Extract                      | Helps glow, nourish, and restore dehydrated skin.   |
| Country Comfort—Herbal Savvy Goldenseal Myrrh, Nuevo, California, USA                               | Myrrh gum powder                   | Disinfecting ointment for psoriasis, eczema, burns, infections, cuts, wounds, scrapes, and dry skin.                              |
| Jean D'Arcel—Anti Wrinkle Lift Stick, Melbourne, Australia  | Resin Extract                      | Antimicrobial/antibacterial.  |
| Anne Semonin—Super Active Serum, Paris, France  | Resin Extract                      | Reduces the depth of wrinkles and skin plumping.  |
| Kuwait shop—Myrrh And Turmeric Lightening And Clarifying Cream, Kuwait, Kuwait                      | Resin Extract                      | Contributes to the unification and natural lightening of the skin tone.   |
| Lush—Beauty Sleep, Vancouver, Canada  | Resin Extract                      | Antimicrobial and antibacterial.  |



**Table 3.** Cosmetic products containing *B. Serrata*.

| Product Name   | <i>B. Serrata</i> /Extract Included | Function   |
|--|-------------------------------------|--|
| Formula Dra Norma Bustos—Anti Stress Dérmico, Buenos Aires, Argentina                                | <i>B. Serrata</i> Extract           | Soothes skin.  |
| Bioearth—Siero Idratante Lenitivo, Fornovo di Taro, Italy  | <i>B. Serrata</i> Extract           | Soothes skin.  |
| Smooth-E—Acne Treatment Hydrogel, Bangkok, Thailand  | <i>B. Serrata</i> Extract           | Relieve inflammation of acne, treat acne, and gently soothes the skin.                 |
| Cantabria Labs Biretix—Gel Ultra Purifying, Madrid, Spain  | <i>B. Serrata</i> Extract           | Hydrating and soothing activity.   |
| Dermedics—YOUTH EXPERT™ Instant Relief Eye Serum, Selangor, Malaysia                                 | <i>B. Serrata</i> Extract           | Anti-irritant.   |
| Dermedics—MESO CALM Instant Soothing Elixir, Selangor, Malaysia                                      | <i>B. Serrata</i> Extract           | Soothes irritations.   |
| Dermedics—YOUTH EXPERT™ Physiological Micellar Water, Selangor, Malaysia                             | <i>B. Serrata</i> Extract           | Anti-irritant.   |
| Ossola Skincare—Prickly Pear & Marula Tinted Emulsion, Carmel-By-The-Sea, California, USA            | <i>B. Serrata</i> Extract           | Soothes skin.  |
| Aveda—outer peace™ acne relief pads, New York, USA   | <i>B. Serrata</i> Extract           | Help wipe out blackheads and prevent new breakouts.                                    |
| I image—CLEAR CELL clarifying salicylic tonic, Lantana, Florida, USA                                 | <i>B. Serrata</i> Extract           | Refresh and purify clogged pores and help soothe irritated skin.                       |
| Aveda—men pure-formance™ conditioner, New York, USA  | <i>B. Serrata</i> Extract           | Refreshes scalp.   |
| Dermadoctor—Ain'T Misbehavin' Intensive 10% Sulfur Acne Mask & Emergency Spot Treatment, Kansas, USA | <i>B. Serrata</i> Extract           | Helps reduce the appearance of spots and blackheads. It also helps the skin to recover |
| Dermalex—Rosacea Treatment, London, UK   | <i>B. Serrata</i> Extract           | Relieve different skin conditions.   |
| NPURE—Day Cream Centella, Jakarta, Indonesia   | <i>B. Serrata</i> Extract           | Antiacne.  |
| Flexpower—Soothe Lotion, Santa Monica, California, USA   | <i>B. Serrata</i> Extract           | Soothes, anti-inflammatory.  |
| Found—Marshmallow Calming Face Serum, Los Angeles, USA   | <i>B. Serrata</i> Extract           | Skin conditioning.   |
| Ausceuticals—Collagen Serum, Perth, Australia  | <i>B. Serrata</i> Extract           | Skin conditioning.   |
| Aveda—Outer Peace™ Foaming Cleanser, New York, USA   | <i>B. Serrata</i> Extract           | Cleans deep into pores without irritation or over-drying skin.                         |
| Biodermal—Couperose Crème, Rotterdam, Netherlands  | <i>B. Serrata</i> Extract           | Soothes skin.  |
| ITreatSkin—Neem Cream, Manchester, UK  | <i>B. Serrata</i> Extract           | Soothes and reduces skin Inflammation.   |
| Kate Ryan—Collagen Booster Intense Repair Serum, Dayton, USA   | <i>B. Serrata</i> Extract           | Reduces lines and wrinkles.  |
| LANCER—Soothe and Hydrate Serum, Beverly hills, USA  | <i>B. Serrata</i> Extract           | Improves skin redness while balancing skin tone.                                       |
| Mitomo—Hyaluronic Acid + Lithospermum Facial Essence Mask, Tokyo, Japan                              | <i>B. Serrata</i> Extract           | Brightening, moisturizing, and refining.   |
| Skin Actives—Collagen Serum, Chandler, USA   | <i>B. Serrata</i> Extract           | Helps improve skin tone and texture.   |
| Kate Somerville—Liquid Exfolikate, Moreno Valley, California, USA                                    | <i>B. Serrata</i> Extract           | Soothes skin.  |
| Yadah—Anti-T Mist, Seoul, Korea  | <i>B. Serrata</i> Extract           | Revitalize and smoothens skin.   |
| New Vitality Lumatone—Anti-Aging Eye Cream, New York, USA  | <i>B. Serrata</i> Extract           | Antiaging.   |

## 7. Summary and Future Aspect

This review summarizes the chemical structure and biological activities of *C. Myrrha* and *B. Serrata* and their uses as cosmetic ingredients. The results of previous studies re-

vealed their nutritional, pharmaceutical, and cosmetic benefits, as they were used over time. These plants are a natural source of medicines and cosmetics. They have anti-inflammatory, antibacterial, antioxidant, and anti-irritant properties. These plants are also antiaging, antiacne, and skin brightening and soothing. Additionally, they reduce the effects of spots and acne. It is also effective for hair and scalp health and antihair-loss.

Further developing value-added cosmetics may prevent many common skin disorders, such as eczema, chapped skin, wrinkles, dark spots, and hair loss. This review highlights the practical uses of *C. Myrrha* and *B. Serrata* as new and safe value-added products and cosmetic ingredients. Thus, using these plants for cosmetic value helps the growth of the cosmetic industry and the optimum exploitation of these natural resources.

From an economic perspective, finding natural sources with cosmetic properties and using them in the development of cosmetics helps to achieve the best use of these plants and is likely to reduce environmental losses, such as burning plants and dumping environmental waste, and helps expand the sources of income for countries exporting these plants.

**Author Contributions:** B.G.A. was involved in writing the original draft, and H.-J.S. provided supervision and writing revision. All authors have read and agreed to the published version of the manuscript.

**Funding:** This research was funded by the Foreign Ministry of Higher Education, Technical and Vocational Training Corporation, Saudi Arabia, grant number “1442/115281 on 8/2021”.

**Institutional Review Board Statement:** Not applicable.

**Informed Consent Statement:** Not applicable.

**Data Availability Statement:** Not applicable.

**Acknowledgments:** The authors thank the Department of Beauty and Cosmetology, Graduate School of Industrial Technology Startup, Chosun University, 309 Pilmun-daero, Dong-gu, Gwangju 61452, Korea.

**Conflicts of Interest:** The authors declare no conflict of interest.

## References

1. Faccio, G. Plant complexity and cosmetic innovation. *IScience* **2020**, *23*, 101358. [[CrossRef](#)] [[PubMed](#)]
2. Ahshawat, M.S.; Saraf, S.; Saraf, S. Preparation and characterization of herbal creams for improvement of skin viscoelastic properties. *Int. J. Cosmet. Sci.* **2008**, *30*, 183–193. [[CrossRef](#)] [[PubMed](#)]
3. Gachathi, F.N. Recent advances on classification and status of the main gum-resin producing species in the family Burseraceae. In Proceedings of the Regional Conference for Africa on Conservation, Management and utilisation of Plant gums, Resins and essential oils, Nairobi, Kenya, 6–10 October 1997.
4. Nielsen, I.; Hedberg, I.; Edwards, S. Flora Ethiopia vol. 3, Pittosporaceae to Araliaceae. *Nord. J. Bot.* **1989**, *12*, 402. [[CrossRef](#)]
5. Rüdiger, A.L.; Siani, A.C.; Junior, V.V. The chemistry and pharmacology of the South America genus Protium Burm. f. (Burseraceae). *Pharmacogn. Rev.* **2007**, *1*, 93–104.
6. Shalabi, L.F.; Otaif, F.S. Commiphora Jacq (Burseraceae) in Saudi Arabia, Botanical, Phytochemical and Ethnobotanical Notes. *Ecologies* **2022**, *3*, 38–57. [[CrossRef](#)]
7. Omer, A.S.; Mai, A.; ElAmin, E.M.; Promy, V.P.; Dagestani, M.; AlEisa, N.; AlMahasna, A. Effect of various level of dietary Commiphora myrrha on rats. *Wulfenia J.* **2014**, *21*, 1–12.
8. Kumar, S.P.; Anjana, S.; Shukla, N.P. Phenol Free and FTA card based Genomic DNA Extraction from Stems and Leaves of Commiphora wightii and C. myrrha. *J. Life Sci.* **2018**, *11*, 25–30.
9. Haffor, A.S.A. Effect of myrrh (*Commiphora molmol*) on leukocyte levels before and during healing from gastric ulcer or skin injury. *J. Immunotoxicol.* **2010**, *7*, 68–75. [[CrossRef](#)]
10. Leminih, M.; Teketay, D. Frankincense and myrrh resources of Ethiopia: I distribution, production, opportunities for dryland development and research needs. *SINET Ethiop. J. Sci.* **2003**, *26*, 63–72. [[CrossRef](#)]
11. Helal, E.G.; Mahmoud, A.; El-Badawy, E.E.; Kahwash, A.A. Effect of Commiphora myrrha extract on some physiological parameters and histological changes in diabetic albino rats. *Egypt. J. Hosp. Med.* **2005**, *20*, 148–162. [[CrossRef](#)]
12. Gadir, S.A.; Ahmed, I.M. Commiphora myrrha and commiphora Africana essential oils. *J. Chem. Pharm. Res.* **2014**, *6*, 151–156.
13. Hassanzadeh\_Taheri, M.; Hosseini, M.; Dorrani pour, D.; Afshar, M.; Moodi, H.; Salimi, M. The Oleo-Gum-Resin of Commiphora myrrha ameliorates male reproductive dysfunctions in streptozotocin-induced hyperglycemic rats. *Pharm. Sci.* **2019**, *25*, 294–302. [[CrossRef](#)]
14. Su, S.; Hua, Y.; Wang, Y.; Gu, W.; Zhou, W.; Duan, J.A.; Tang, Y. Evaluation of the anti-inflammatory and analgesic properties of individual and combined extracts from *Commiphora myrrha*, and *Boswellia carterii*. *J. Ethnopharmacol.* **2012**, *139*, 649–656. [[CrossRef](#)]

15. Shalaby, M.A.; Hammouda, A.A.E. Analgesic, anti-inflammatory and anti-hyperlipidemic activities of *Commiphora molmol* extract (Myrrh). *J. Intercult. Ethnopharmacol.* **2014**, *3*, 56. [[CrossRef](#)] [[PubMed](#)]
16. Kavousi, A.; Nikkhah, E.; Tayarani-Najaran, Z.; Javadi, B. Wound healing effects and related mechanisms of action of methanol extracts of *Boswellia sacra* and *Commiphora myrrha* oleo-gum-resins on adult human dermal fibroblasts (HDFa). *SSRN* **2021**, *1*, 1–17. [[CrossRef](#)]
17. Alhussaini, M.S.; Saadabi, A.M.; Alghonaim, M.I.; Ibrahim, K.E. An evaluation of the Antimicrobial activity of *Commiphora myrrha* Nees (Engl.) oleo-gum resins from Saudi Arabia. *J. Med. Sci.* **2015**, *15*, 198. [[CrossRef](#)]
18. Lemenith, M.; Teketay, D. Frankincense and myrrh resources of Ethiopia: II. Medicinal and industrial uses. *SINET Ethiop. J. Sci.* **2003**, *26*, 161–172. [[CrossRef](#)]
19. Tipton, D.A.; Lyle, B.; Babich, H.; Dabbous, M.K. In vitro cytotoxic and anti-inflammatory effects of myrrh oil on human gingival fibroblasts and epithelial cells. *Toxicol. Vitro.* **2003**, *17*, 301–310. [[CrossRef](#)]
20. Madia, V.N.; De Angelis, M.; De Vita, D.; Messore, A.; De Leo, A.; Ialongo, D.; Costi, R. Investigation of *Commiphora myrrha* (Nees) Engl. oil and its main components for antiviral activity. *Pharmaceuticals* **2021**, *14*, 243. [[CrossRef](#)]
21. Zhou, Q.; Liu, Y.; Tang, Y.; Shokoohinia, Y.; Chittiboyina, A.G.; Wang, M.; Avonto, C. Identification of potential skin sensitizers in myrrh. *Cosmetics* **2019**, *6*, 47. [[CrossRef](#)]
22. Thulin, M.; Warfa, A.M. The frankincense trees (*Boswellia* spp., Burseraceae) of northern Somalia and southern Arabia. *Kew Bull.* **1987**, *42*, 487–500. [[CrossRef](#)]
23. Hosain, N.A.; Ghosh, R.; Bryant, D.L.; Arivett, B.A.; Farone, A.L.; Kline, P.C. Isolation, structure elucidation, and immunostimulatory activity of polysaccharide fractions from *Boswellia carterii* frankincense resin. *Int. J. Biol. Macromol.* **2019**, *133*, 76–85. [[CrossRef](#)]
24. Tolera, M.; Sass-Klaassen, U.; Eshete, A.; Bongers, F.; Sterck, F.J. Frankincense tree recruitment failed over the past half century. *For. Ecol. Manag.* **2013**, *304*, 65–72. [[CrossRef](#)]
25. Chaurasia, A.R.T.I.; Gharia, A.N.I.L. Antifungal activity of medicinal plant *Boswellia Serrata*. *J. Ultra Chem.* **2017**, *13*, 88–90. [[CrossRef](#)]
26. Hamm, S.; Bleton, J.; Connan, J.; Tchaplal, A. A chemical investigation by headspace SPME and GC–MS of volatile and semi-volatile terpenes in various olibanum samples. *Phytochemistry* **2005**, *66*, 1499–1514. [[CrossRef](#)]
27. Raja, A.F.; Ali, F.; Khan, I.A.; Shawl, A.S.; Arora, D.S. Acetyl-11-keto- $\beta$ -boswellic acid (AKBA); targeting oral cavity pathogens. *BMC Res. Notes* **2011**, *4*, 406. [[CrossRef](#)]
28. Kohoude, M.J.; Gbaguidi, F.; Agbani, P.; Ayedoun, M.A.; Cazaux, S.; Bouajila, J. Chemical composition and biological activities of extracts and essential oil of *Boswellia dalzielii* leaves. *Pharm. Biol.* **2017**, *55*, 33–42. [[CrossRef](#)]
29. Shah, B.A.; Qazi, G.N.; Taneja, S.C. Boswellic acids: A group of medicinally important compounds. *Nat. Prod. Rep.* **2009**, *26*, 72–89. [[CrossRef](#)]
30. Suchita, W.; Raman, D.M.R.; Kaur, C.D. A Review on phytochemistry and pharmacological activities of *Boswellia serrata*: A natural remedy. *Int. J. Pharmacogn.* **2021**, *8*, 454–461.
31. Maupetit, P. New constituents in olibanum resinoid and essential oil. *Perfum. Flavorist* **1984**, *9*, 19–37.
32. Leung, A.Y.; Foster, S. *Encyclopedia of Common Natural Ingredients Used in Food, Drugs, and Cosmetics*; John Wiley & Sons, Inc.: New York, NY, USA, 1996.
33. Ghorpade, R.P.; Chopra, A.; Nikam, T.D. In vitro zygotic embryo germination and propagation of an endangered *Boswellia serrata* Roxb., a source of boswellic acid. *Physiol. Mol. Biol. Plants* **2010**, *16*, 159–165. [[CrossRef](#)] [[PubMed](#)]
34. Khan, I.A.; Abourashed, E.A. *Leung's encyclopedia of common natural ingredients: Used in food, drugs and cosmetics*, 3rd ed.; John Wiley & Sons: Hoboken, NJ, USA, 2010.
35. Singh, S.; Khajuria, A.; Taneja, S.C.; Khajuria, R.K.; Singh, J.; Johri, R.K.; Qazi, G.N. The gastric ulcer protective effect of boswellic acids, a leukotriene inhibitor from *Boswellia serrata*, in rats. *Phytomedicine* **2008**, *15*, 408–415. [[CrossRef](#)]
36. Alam, M.; Khan, H.; Samiullah, L.; Siddique, K.M. A review on phytochemical and pharmacological studies of Kundur (*Boswellia serrata* Roxb ex Colebr.)-A Unani drug. *J. Appl. Pharm. Sci.* **2012**, *2*, 148–156.
37. Ramani, P.V.; Godhani, P.J. To determine antimicrobial and phytochemical properties of *Boswellia serrata* leaf extracts. *J. Pharmacogn. Phytochem.* **2020**, *9*, 45–47.
38. Siddiqui, M.Z. *Boswellia serrata*, a potential anti-inflammatory agent: An overview. *Indian J. Pharm. Sci.* **2011**, *73*, 255.
39. Schrott, E.; Laufer, S.; Lammerhofer, M.; Ammon, H.P.T. Extract from gum resin of *Boswellia serrata* decreases [IA. sub. 2]-antibody in a patient with “Late onset Autoimmune Diabetes of the Adult” (LADA). *Phytomedicine Int. J. Phytother. Phytopharm.* **2014**, *21*, 786–787.
40. Umar, S.; Umar, K.; Sarwar, A.H.M.G.; Khan, A.; Ahmad, N.; Ahmad, S.; Khan, H.A. *Boswellia serrata* extract attenuates inflammatory mediators and oxidative stress in collagen induced arthritis. *Phytomedicine* **2014**, *21*, 847–856. [[CrossRef](#)]
41. Zhao, W.; Entschladen, F.; Liu, H.; Niggemann, B.; Fang, Q.; Zaenker, K.S.; Han, R. Boswellic acid acetate induces differentiation and apoptosis in highly metastatic melanoma and fibrosarcoma cells. *Cancer Detect. Prev.* **2003**, *27*, 67–75. [[CrossRef](#)]
42. Montaser, M.M.; El-Sharnouby, M.E.; El-Noubi, G.; El-Shaer, H.M.; Khalil, A.A.; Hassanin, M.; El-Araby, D.A. *Boswellia serrata* resin extract in diets of Nile tilapia, *Oreochromis niloticus*: Effects on the growth, health, immune response, and disease resistance to *Staphylococcus aureus*. *Animals* **2021**, *11*, 446. [[CrossRef](#)]
43. Hewedy, W.A. Effect of *Boswellia serrata* on rat trachea contractility in vitro. *Nat. Prod. J.* **2020**, *10*, 33–43. [[CrossRef](#)]

44. Salman, K.A.; Jawad, S.M.; Abbas, S.H. Evaluation of Antibacterial Activity of Boswellia Serrata Extract Against some of the Oral Pathogenic Bacteria. *Indian J. Forensic Med. Toxicol.* **2021**, *15*, 3371.
45. Mothana, R.A.; Hasson, S.S.; Schultze, W.; Mowitz, A.; Lindequist, U. Phytochemical composition and in vitro antimicrobial and antioxidant activities of essential oils of three endemic Soqotraen Boswellia species. *Food Chem.* **2011**, *126*, 1149–1154. [[CrossRef](#)]
46. Gupta, M.; Verma, S.K.; Singh, S.; Trivedi, L.; Rout, P.K.; Vasudev, P.G.; Misra, L. Anti-proliferative and antibacterial activity of oleo-gum-resin of Boswellia serrata extract and its isolate 3-hydroxy-11-keto- $\beta$ -boswellic acid. *J. Herb. Med.* **2022**, *32*, 100546. [[CrossRef](#)]
47. Brendler, T.; Brinckmann, J.A.; Schippmann, U. Sustainable supply, a foundation for natural product development: The case of Indian frankincense (*Boswellia serrata* Roxb. ex Colebr.). *J. Ethnopharmacol.* **2018**, *225*, 279–286. [[CrossRef](#)]
48. Majeed, M.; Prakash, L. *Boswellin CG: Chemistry, Toxicology & Clinical*; Sabinsa Corporation: Piscataway, NJ, USA, 2007.
49. Mishram, S.; Bishnoi, R.S.; Maurya, R.; Jain, D. Boswellia serrata ROXB—A bioactive herbs with various pharmacological activities. *Asian J. Pharm. Clin. Res.* **2020**, *13*, 33–39. [[CrossRef](#)]
50. Miller, A.L. Effects of Boswellia serrata on asthma. *Altern. Med. Rev.* **2001**, *6*, 20–47.
51. Mathe, C.; Culioli, G.; Archier, P.; Vieillescazes, C. Characterization of archaeological frankincense by gas chromatography–mass spectrometry. *J. Chromatogr. A* **2004**, *1023*, 277–285. [[CrossRef](#)]
52. Ahmed, S.; Alam, A.; Shahabuddin, M.; Khan, I.; Ali, H. Versatile pharmacological action and compound formulation of Kundur in Unani medicine: A review. *Int. J. Pharmacogn.* **2014**, *1*, 627–631.
53. Sharma, S.; Thawani, V.; Hingorani, L.; Shrivastava, M.; Bhate, V.R.; Khiyani, R. Pharmacokinetic study of 11-keto  $\beta$ -Boswellic acid. *Phytomedicine* **2004**, *11*, 255–260. [[CrossRef](#)] [[PubMed](#)]
54. Hostanska, K.; Daum, G.; Saller, R. Cytostatic and apoptosis-inducing activity of boswellic acids toward malignant cell lines in vitro. *Anticancer Res.* **2002**, *22*, 2853–2862.
55. Upaganlawar, A.; Ghule, B. Pharmacological activities of *Boswellia serrata* Roxb.—Mini review. *Ethnobot. Leaflet.* **2009**, *6*, 10.
56. Behnamrasuli, M.; Hoseinzadeh, H.; Ghafarimoghadam, G. Empowering effect of Frankincense extract on memory. *Tarbiat Moalem Univ. J. Sci.* **2001**, *1*, 1–13.
57. Ahangarpour, A.; Heidari, H.; Fatemeh, R.A.A.; Pakmehr, M.; Shahbazian, H.; Ahmadi, I.; Mehrangiz, B.H. Effect of Boswellia serrata supplementation on blood lipid, hepatic enzymes and fructosamine levels in type2 diabetic patients. *J. Diabetes Metab. Disord.* **2014**, *13*, 29. [[CrossRef](#)] [[PubMed](#)]
58. Zutshi, U.; Rao, P.G.; Kaur, S.; Atal, G.S.C. Mechanism of cholesterol lowering effect of Salai guggal ex-*Boswellia serrata*. *Indian J. Pharm.* **1980**, *12*, 59.
59. Wildfeuer, A.; Neu, I.S.; Safayhi, H.; Metzger, G.; Wehrmann, M.; Vogel, U.; Ammon, H.P. Effects of boswellic acids extracted from a herbal medicine on the biosynthesis of leukotrienes and the course of experimental autoimmune encephalomyelitis. *Arzneim.-Forsch.* **1998**, *48*, 668–674.
60. Ahmed, H.H.; El-Alfy, N.Z.; Mahmoud, M.F.; Yahya, S.M. Boswellia serrata oleo-gum resin: A natural remedy for retrogradation of liver fibrosis in rats. *Der Pharm. Lett.* **2015**, *7*, 134–144.
61. Liu, Z.; Liu, X.; Sang, L.; Liu, H.; Xu, Q.; Liu, Z. Boswellic acid attenuates asthma phenotypes by downregulation of GATA3 via pSTAT6 inhibition in a murine model of asthma. *Int. J. Clin. Exp. Pathol.* **2015**, *8*, 236.
62. Housen, M.E.; Ragab, A.; Mesbah, A.; El-Samanoudy, A.Z.; Othman, G.; Moustafa, A.F.; Badria, F.A. Natural anti-inflammatory products and leukotriene inhibitors as complementary therapy for bronchial asthma. *Clin. Biochem.* **2010**, *43*, 887–890. [[CrossRef](#)]
63. Roy, N.K.; Parama, D.; Banik, K.; Bordoloi, D.; Devi, A.K.; Thakur, K.K.; Kunnumakkara, A.B. An update on pharmacological potential of boswellic acids against chronic diseases. *Int. J. Mol. Sci.* **2019**, *20*, 4101. [[CrossRef](#)]
64. Suther, C.; Daddi, L.; Bokoliya, S.; Panier, H.; Liu, Z.; Lin, Q.; Zhou, Y. Dietary Boswellia serrata Acid Alters the Gut Microbiome and Blood Metabolites in Experimental Models. *Nutrients* **2022**, *14*, 814. [[CrossRef](#)]
65. Ranjbarnejad, T.; Saidijam, M.; Moradkhani, S.; Najafi, R. Methanolic extract of Boswellia serrata exhibits anti-cancer activities by targeting microsomal prostaglandin E synthase-1 in human colon cancer cells. *Prostaglandins Other Lipid Mediat.* **2017**, *131*, 1–8. [[CrossRef](#)] [[PubMed](#)]
66. Assimopoulou, A.N.; Zlatanov, S.N.; Papageorgiou, V.P. Antioxidant activity of natural resins and bioactive triterpenes in oil substrates. *Food Chem.* **2005**, *92*, 721–727. [[CrossRef](#)]
67. Yassin, N.A.Z.; El-Shenawy, S.M.A.; Mahdy, K.A.; Gouda, N.A.M.; Marrie, A.E.-F.H.; Farrag, A.R.H.; Ibrahim, B.M.M. Effect of Boswellia serrata on Alzheimer’s disease induced in rats. *J. Arab Soc. Med. Res.* **2013**, *8*, 1–11.
68. Gupta, I.; Gupta, V.; Parihar, A.; Gupta, S.; Lüdtkke, R.; Safayhi, H.; Ammon, H.P. Effects of Boswellia serrata gum resin in patients with bronchial asthma: Results of a double-blind, placebo-controlled, 6-week clinical study. *Eur. J. Med. Res.* **1998**, *3*, 511–514. [[PubMed](#)]
69. Ammon, H.P.T. Boswellic acids in chronic inflammatory diseases. *Planta Med.* **2006**, *72*, 1100–1116. [[CrossRef](#)]
70. Abdel-Tawab, M.; Werz, O.; Schubert-Zsilavec, M. Boswellia serrata. *Clin. Pharmacokinet.* **2011**, *50*, 349–369. [[CrossRef](#)]
71. Pop, R.M.; Popolo, A.; Trifa, A.P.; Stanciu, L.A. Phytochemicals in cardiovascular and respiratory diseases: Evidence in oxidative stress and inflammation. *Oxidative Med. Cell. Longev.* **2018**, *2018*, 1603872. [[CrossRef](#)]
72. Etzel, R. Special extract of Boswellia serrata (H 15) in the treatment of rheumatoid arthritis. *Phytomedicine* **1996**, *3*, 91–94. [[Cross-Ref](#)]

73. Roy, S.; Khanna, S.; Shah, H.; Rink, C.; Phillips, C.; Preuss, H.; Sen, C.K. Human genome screen to identify the genetic basis of the anti-inflammatory effects of Boswellia in microvascular endothelial cells. *DNA Cell Biol.* **2005**, *24*, 244–255. [[CrossRef](#)]
74. Alluri, V.K.; Kundimi, S.; Sengupta, K.; Golakoti, T.; Kilari, E.K. An anti-inflammatory composition of Boswellia serrata resin extracts alleviates pain and protects cartilage in monoiodoacetate-induced osteoarthritis in rats. *Evid.-Based Complement. Altern. Med.* **2020**, *2020*, 7381625. [[CrossRef](#)]
75. Engels, G. Frankincense, herbal gram. *J. Am. Bot. Coun.* **2010**, *88*, 1–4.
76. Kuo, R.Y.; Qian, K.; Morris-Natschke, S.L.; Lee, K.H. Plant-derived triterpenoids and analogues as antitumor and anti-HIV agents. *Nat. Prod. Rep.* **2009**, *26*, 1321–1344. [[CrossRef](#)]
77. Asante-Kwatia, E.; Yeboah Mensah, A.; Frimpong Baidoo, M. Analgesic and anti-inflammatory effect of Ghanaian medicinal plants. In *Medicinal Plants-Use in Prevention and Treatment of Diseases*, 2nd ed.; Bassam, A.R.H., Ed.; IntechOpen: London, UK, 2020; Volume 3, p. 192.
78. Pirotta, M. Arthritis disease: The use of complementary therapies. *Aust. Fam. Physician* **2020**, *39*, 638–640.
79. Singh, G.B.; Atal, C.K. Pharmacology of an extract of salai guggal ex-Boswellia serrata, a new non-steroidal anti-inflammatory agent. *Agents Actions* **1986**, *18*, 407–412. [[CrossRef](#)] [[PubMed](#)]
80. Kumar, R.; Singh, S.; Saksena, A.K.; Pal, R.; Jaiswal, R.; Kumar, R. Effect of Boswellia serrata extract on acute inflammatory parameters and tumor necrosis factor- $\alpha$  in complete Freund's adjuvant-induced animal model of rheumatoid arthritis. *Int. J. Appl. Basic Med. Res.* **2019**, *9*, 100. [[CrossRef](#)]
81. Iram, F.; Khan, S.A.; Husain, A. Phytochemistry and potential therapeutic actions of Boswellic acids: A mini review. *Asian Pac. J. Trop. Biomed.* **2017**, *7*, 513–523. [[CrossRef](#)]
82. Vishal, A.A.; Mishra, A.; Raychaudhuri, S.P. A double blind, randomized, placebo controlled clinical study evaluates the early efficacy of Aflapin<sup>®</sup> in subjects with osteoarthritis of knee. *Int. J. Med. Sci.* **2011**, *8*, 615. [[CrossRef](#)]
83. Sengupta, K.; Alluri, K.V.; Satish, A.R.; Mishra, S.; Golakoti, T.; Sarma, K.V.; Raychaudhuri, S.P. A double blind, randomized, placebo-controlled study of the efficacy and safety of 5-Loxin<sup>®</sup> for treatment of osteoarthritis of the knee. *Arthritis Res. Ther.* **2008**, *10*, R85. [[CrossRef](#)]
84. Sengupta, K.; Krishnaraju, A.V.; Vishal, A.A.; Mishra, A.; Trimurtulu, G.; Sarma, K.V.; Raychaudhuri, S.P. Comparative efficacy and tolerability of 5-Loxin<sup>®</sup> and Aflapin<sup>®</sup> against osteoarthritis of the knee: A double blind, randomized, placebo controlled clinical study. *Int. J. Med. Sci.* **2010**, *7*, 366. [[CrossRef](#)]
85. Kulkarni, R.R.; Patki, P.S.; Jog, V.P.; Gandage, S.G.; Patwardhan, B. Treatment of osteoarthritis with a herbomineral formulation: A double-blind, placebo-controlled, cross-over study. *J. Ethnopharmacol.* **1991**, *33*, 91–95. [[CrossRef](#)]
86. Ali, N.A.A.; Wurster, M.; Arnold, N.; Teichert, A.; Schmidt, J.; Lindequist, U.; Wessjohann, L. Chemical composition and biological activities of essential oils from the oleogum resins of three endemic Socotran Boswellia species. *Rec. Nat. Prod.* **2008**, *2*, 6.
87. Ammon, H.P.T. Boswellic acids and their role in chronic inflammatory diseases. *Anti-Inflamm. Nutraceuticals Chronic Dis.* **2016**, *928*, 291–327.
88. Ernst, E. Frankincense: Systematic review. *BMJ* **2008**, *337*, a2813. [[CrossRef](#)]
89. Majeed, M.; Majeed, S.; Narayanan, N.K.; Nagabhushanam, K. A pilot, randomized, double-blind, placebo-controlled trial to assess the safety and efficacy of a novel Boswellia serrata extract in the management of osteoarthritis of the knee. *Phytother. Res.* **2019**, *33*, 1457–1468. [[CrossRef](#)]
90. Vahabi, S.; Hakemi-Vala, M.; Gholami, S. In vitro antibacterial effect of hydroalcoholic extract of Lawsonia inermis, Malva sylvestris, and Boswellia serrata on aggregatibacter actinomycetemcomitans. *Adv. Biomed. Res.* **2019**, *8*, 22. [[CrossRef](#)]
91. Sadhasivam, S.; Palanivel, S.; Ghosh, S. Synergistic antimicrobial activity of Boswellia serrata Roxb. ex Colebr.(Bursaceae) essential oil with various azoles against pathogens associated with skin, scalp and nail infections. *Lett. Appl. Microbiol.* **2016**, *63*, 495–501. [[CrossRef](#)]
92. Jaroš, P.; Vrublevskaia, M.; Lokočová, K.; Michailidu, J.; Kolouchová, I.; Demnerová, K. Boswellia serrata Extract as an Antibiofilm Agent against Candida spp. *Microorganisms* **2022**, *10*, 171. [[CrossRef](#)] [[PubMed](#)]
93. Menon, M.K.; Kar, A. Analgesic and psychopharmacological effects of the gum resin of Boswellia serrata. *Planta Med.* **1971**, *19*, 333–341. [[CrossRef](#)] [[PubMed](#)]
94. Gupta, M.; Rout, P.K.; Misra, L.N.; Gupta, P.; Singh, N.; Darokar, M.P.; Bhakuni, R.S. Chemical composition and bioactivity of Boswellia serrata Roxb. essential oil in relation to geographical variation. *Plant Biosyst.-Int. J. Deal. All Asp. Plant Biol.* **2017**, *151*, 623–629.
95. Khan, M.A.; Singh, M.; Khan, M.S.; Najmi, A.K.; Ahmad, S. Caspase mediated synergistic effect of Boswellia serrata extract in combination with doxorubicin against human hepatocellular carcinoma. *Biomed Res. Int.* **2014**, *2014*, 294143. [[CrossRef](#)]
96. Ayub, M.A.; Hanif, M.A.; Sarfraz, R.A.; Shahid, M. Biological activity of Boswellia serrata Roxb. oleo gum resin essential oil: Effects of extraction by supercritical carbon dioxide and traditional methods. *Int. J. Food Prop.* **2018**, *21*, 808–820. [[CrossRef](#)]
97. Van Vuuren, S.F. Antimicrobial activity of South African medicinal plants. *J. Ethnopharmacol.* **2008**, *119*, 462–472. [[CrossRef](#)] [[PubMed](#)]
98. Gaurea, S.H.; Bapat, U.C. Study of antibacterial activity of resins of Boswellia serrata roxb ex colebr., Commiphora mukul (hooks ex-stocks) engl., Gardenia resinifera roth. And Shorea robusta gaertn. *Int. J. Pharm. Pharm. Sci.* **2016**, *8*, 29–31.

99. Lalithakumari, K.; Krishnaraju, A.V.; Sengupta, K.; Subbaraju, G.V.; Chatterjee, A. Safety and toxicological evaluation of a novel, standardized 3-O-acetyl-11-keto- $\beta$ -boswellic acid (AKBA)-enriched *Boswellia serrata* extract (5-Loxin<sup>®</sup>). *Toxicol. Mech. Methods* **2006**, *16*, 199–226. [[CrossRef](#)]
100. Katragunta, K.; Siva, B.; Kondepudi, N.; Vadaparthy, P.R.; Rao, N.R.; Tiwari, A.K.; Babu, K.S. Estimation of boswellic acids in herbal formulations containing *Boswellia serrata* extract and comprehensive characterization of secondary metabolites using UPLC-Q-Tof-MSe. *J. Pharm. Anal.* **2019**, *9*, 414–422. [[CrossRef](#)] [[PubMed](#)]
101. Camarda, L.; Dayton, T.; Di Stefano, V.; Pitonzo, R.; Schillaci, D. Chemical composition and antimicrobial activity of some oleogum resin essential oils from *Boswellia* spp. (Burseraceae). *Ann. Di Chim. J. Anal. Environ. Cult. Herit. Chem.* **2007**, *97*, 837–844.
102. Mannino, G.; Occhipinti, A.; Maffei, M.E. Quantitative determination of 3-O-acetyl-11-keto- $\beta$ -boswellic acid (AKBA) and other boswellic acids in *Boswellia sacra* Flueck (syn. *B. carteri* Birdw) and *Boswellia serrata* Roxb. *Molecules* **2016**, *21*, 1329. [[CrossRef](#)] [[PubMed](#)]
103. Murthy, T.K.; Shiva, M.P. Salai guggul from *Boswellia serrata* Roxb.—Its exploitation and utilization. *Indian For.* **1977**, *103*, 466–474.
104. Verghese, J. Olibanum in focus. *Perfum. Flavorist* **1988**, *13*, 1–12.
105. Holmes, P. Frankincense oil. *Int. J. Arom.* **1999**, *9*, 56–61.
106. Evans, W.C. A taxonomic approach to the study of medicinal plants and animal derived drugs. Trease and Evans Pharmacognosy. *WB Saunders* **2002**, *15*, 33–35.
107. Chandrasekharnath, N.; Mahlakshmi, Y.V.; Jayalakshmi, L.; Venkanna, B.; Uma, A. Screening and isolation of bioactive factors from *Commiphora myrrha* and evaluation of their antimicrobial activity. *IJERA* **2013**, *3*, 1291–1294.
108. Suliman, R.S.; Alghamdi, S.S.; Ali, R.; Aljatli, D.; Aljammaz, N.A.; Huwaizi, S.; Rahman, I. The Role of Myrrh Metabolites in Cancer, Inflammation, and Wound Healing: Prospects for a Multi-Targeted Drug Therapy. *Pharmaceuticals* **2022**, *15*, 944. [[CrossRef](#)] [[PubMed](#)]
109. Su, S.; Duan, J.; Chen, T.; Huang, X.; Shang, E.; Yu, L.; Tang, Y. Frankincense and myrrh suppress inflammation via regulation of the metabolic profiling and the MAPK signaling pathway. *Sci. Rep.* **2015**, *5*, 13668. [[CrossRef](#)] [[PubMed](#)]
110. Su, S.; Wang, T.; Duan, J.A.; Zhou, W.; Hua, Y.Q.; Tang, Y.P.; Qian, D.W. Anti-inflammatory and analgesic activity of different extracts of *Commiphora myrrha*. *J. Ethnopharmacol.* **2011**, *134*, 251–258. [[CrossRef](#)]
111. Fatani, A.J.; Alrojaye, F.S.; Parmar, M.Y.; Abuhashish, H.M.; Ahmed, M.M.; Al-Rejaie, S.S. Myrrh attenuates oxidative and inflammatory processes in acetic acid-induced ulcerative colitis. *Exp. Ther. Med.* **2016**, *12*, 730–738. [[CrossRef](#)]
112. Abukhader, R.; Al tawaha, A. Amazing Benefits of Myrrh. *Int. J. Pharm. Res.* **2021**, *13*, 303–308.
113. Rastogi Ram, P.; Mehrotra, B.N. Compendium of Indian medicinal plants. *New Delhi Lucknow NISCAIR* **1993**, *4*, 1.
114. Rastogi, R.P.; Mehrotra, B.N. *Compendium of Indian Medicinal Plants*; Central Drug Research Institute: Lucknow, India, 1990; Volume 4.
115. Al-Harrasi, A.; Al-Saidi, S. Phytochemical analysis of the essential oil from botanically certified oleogum resin of *Boswellia sacra* (Omani Luban). *Molecules* **2008**, *13*, 2181–2189. [[CrossRef](#)]
116. Rashan, L.; Hakkim, F.L.; Idrees, M.; Essa, M.; Velusamy, T.; Al-Baloshi, M.; Hasson, S.S. *Boswellia* gum resin and essential oils: Potential health benefits—An evidence based review. *Int. J. Nutr. Pharmacol. Neurol. Dis.* **2019**, *9*, 53–71. [[CrossRef](#)]
117. Al-Yasiry, A.R.M.; Kiczorowska, B. Frankincense-therapeutic properties. *Adv. Hyg. Exp. Med./Postep. Hig. I Med. Dosw.* **2016**, *70*, 380–391. [[CrossRef](#)] [[PubMed](#)]
118. Niebler, J.; Buettner, A. Identification of odorants in frankincense (*Boswellia sacra* Flueck.) by aroma extract dilution analysis and two-dimensional gas chromatography–mass spectrometry/olfactometry. *Phytochemistry* **2015**, *109*, 66–75. [[CrossRef](#)] [[PubMed](#)]
119. Al-Harrasi, A.; Csuk, R.; Khan, A.; Hussain, J. Distribution of the anti-inflammatory and anti-depressant compounds: Incensole and incensole acetate in genus *Boswellia*. *Phytochemistry* **2019**, *161*, 28–40. [[CrossRef](#)] [[PubMed](#)]
120. Herrmann, A.; Lechtenberg, M.; Hensel, A. Comparative isolation and structural investigations of polysaccharides from *Boswellia serrata* ROXB. and *Boswellia carteri* BIRDW. *Planta Med.* **2007**, *73*, 09. [[CrossRef](#)]
121. Sharma, R.A.; Varma, K.C. Studies on gum obtained from *Boswellia serrata* Roxb. *Indian Drugs* **1980**, *17*, 225–227.
122. Bhuchar, V.M.; Agrawal, A.; Sharma, S.K. Constituents of the gum obtained from *Boswellia serrata* exudate. *Indian J. Technol.* **1982**, *20*, 38–40.
123. Gangwal, M.L.; Vardhan, D.K. Carbohydrate contents of *Boswellia serrata*. *Asian J. Chem.* **1995**, *7*, 677.
124. Hana, D.B.; Kadhim, H.M.; Jasim, G.A.; Latif, Q.N. Antibacterial activity of *Commiphora molmol* extracts on some bacterial species in Iraq. *Sch. Acad. J. Pharm.* **2016**, *5*, 406–412.
125. Maver, T.; Maver, U.; Stana Kleinschek, K.; Smrke, D.M.; Kreft, S. A review of herbal medicines in wound healing. *Int. J. Dermatol.* **2015**, *54*, 740–751. [[CrossRef](#)]
126. Massoud, A.; El Sisi, S.A.W.S.A.N.; Salama, O. Preliminary study of therapeutic efficacy of a new fasciolocidal drug derived from *Commiphora molmol* (myrrh). *Am. J. Trop. Med. Hyg.* **2001**, *65*, 96–99. [[CrossRef](#)]
127. Buckley, S.A.; Evershed, R.P. Organic chemistry of embalming agents in Pharaonic and Graeco-Roman mummies. *Nature* **2001**, *413*, 837–841. [[CrossRef](#)] [[PubMed](#)]
128. Cenci, E.; Messina, F.; Rossi, E.; Epifano, F.; Marcotullio, M.C. Antiviral furanosesquiterpenes from *Commiphora erythraea*. *Nat. Prod. Commun.* **2012**, *7*, 143–144. [[CrossRef](#)] [[PubMed](#)]

129. Mohamed, A.G.; Abbas, H.M.; Kassem, J.M.; Gafour, W.A.; Attalah, A.G. Impact of myrrh essential oil as a highly effective antimicrobial agent in processed cheese spreads. *Int. J. Dairy Sci.* **2016**, *11*, 41–51. [[CrossRef](#)]
130. Marcotullio, M.C.; Rosati, O.; Lanari, D. Phytochemistry of *Commiphora erythraea*: A Review. *Nat. Prod. Commun.* **2018**, *13*, 1209–1212. [[CrossRef](#)]
131. Ajiteru, O.; Lee, O.J.; Kim, J.H.; Lee, Y.J.; Lee, J.S.; Lee, H.; Park, C.H. Fabrication and characterization of a myrrh hydrocolloid dressing for dermal wound healing. *Colloid Interface Sci. Commun.* **2022**, *48*, 100617. [[CrossRef](#)]
132. Lv, L.; Yan, G.Y.; Zhao, Y.L.; He, X.J.; Jiang, X.; Zhuo, Y.Q.; Cen, X.B. Investigation of the dermal sensitizing potential of traditional medical extracts in local lymph node assays. *Exp. Biol. Med.* **2009**, *234*, 306–313. [[CrossRef](#)] [[PubMed](#)]
133. Kamil, N.B.; Al-Ghaban, N.M. Evaluation the effects of local exogenous application of a mixture of Myrrh and Sage oil on incisional wound healing on the skin of the face (Histological, Histochemical and Histomorphpmetrical study in rabbits). *J. Pharm. Sci. Res.* **2018**, *10*, 1030–1035.
134. Atal, C.K.; Singh, G.B.; Batra, S.; Sharma, S.; Gupta, O.P. Salai Guggal ex-Boswellia serrata a promising anti-hyperlipidemic and anti-arthritic agent. *Indian J. Exp. Biol.* **1980**, *12*, 59.
135. Atal, C.K.; Gupta, O.P.; Singh, G.B. Salai-guggal a promising anti-arthritic and anti-hyperlipidemic agent. *Br. J. Pharmacol.* **1981**, *74*, 203–204.
136. Ammon, H.P.T.; Safayhi, H.; Mack, T.; Sabieraj, J. Mechanism of antiinflammatory actions of curcumine and boswellic acids. *J. Ethnopharmacol.* **1993**, *38*, 105–112. [[CrossRef](#)]
137. Hussain, H.; Al-Harrasi, A.; Csuk, R.; Shamraiz, U.; Green, I.R.; Ahmed, I.; Ali, Z. Therapeutic potential of boswellic acids: A patent review (1990–2015). *Expert Opin. Ther. Pat.* **2017**, *27*, 81–90. [[CrossRef](#)] [[PubMed](#)]
138. Hajhashemi, V.; Safaei, S. Effect of a selection of skin penetration enhancers on topical anti-inflammatory effect of Boswellic acids in carrageenan-induced paw edema in rats. *Adv. Biomed. Res.* **2021**, *10*, 18. [[CrossRef](#)] [[PubMed](#)]
139. Haghaei, H.; Soltani, S.; Hosseini, S.A.; Rashidi, M.R.; Karima, S. Boswellic acids as promising leads in drug development against Alzheimer's disease. *Pharm. Sci.* **2020**, *27*, 14–31. [[CrossRef](#)]
140. Kora, A.J.; Sashidhar, R.B.; Arunachalam, J. Aqueous extract of gum olibanum (*Boswellia serrata*): A reductant and stabilizer for the biosynthesis of antibacterial silver nanoparticles. *Process Biochem.* **2021**, *47*, 1516–1520. [[CrossRef](#)]
141. Ismail, S.M.; Aluru, S.; Sambasivarao, K.R.S.; Matcha, B. Antimicrobial activity of frankincense of *Boswellia serrata*. *Int. J. Curr. Microbiol. App. Sci.* **2014**, *3*, 1095–1101.
142. De Rapper, S.; Van Vuuren, S.F.; Kamatou, G.P.P.; Viljoen, A.M.; Dagne, E. The additive and synergistic antimicrobial effects of select frankincense and myrrh oils—A combination from the pharaonic pharmacopoeia. *Let. Appl. Microbiol.* **2012**, *54*, 352–358. [[CrossRef](#)]
143. Alluri, V.K.; Dodda, S.; Kilari, E.K.; Golakoti, T.; Sengupta, K. Toxicological assessment of a standardized *Boswellia serrata* gum resin extract. *Int. J. Toxicol.* **2019**, *38*, 423–435. [[CrossRef](#)]
144. Di Stefano, V.; Schillaci, D.; Cusimano, M.G.; Rishan, M.; Rashan, L. In vitro antimicrobial activity of frankincense oils from *Boswellia sacra* grown in different locations of the Dhofar region (Oman). *Antibiotics* **2020**, *9*, 195. [[CrossRef](#)]
145. Grbić, M.L.; Unković, N.; Dimkić, I.; Janačković, P.; Gavrilović, M.; Stanojević, O.; Vukojević, J. Frankincense and myrrh essential oils and burn incense fume against micro-inhabitants of sacral ambients. Wisdom of the ancients? *J. Ethnopharmacol.* **2018**, *219*, 1–14. [[CrossRef](#)]
146. El Ashry, E.S.H.; Rashed, N.; Salama, O.M.; Saleh, A. Components, therapeutic value and uses of myrrh. *Die Pharm.-Int. J. Pharm. Sci.* **2003**, *58*, 163–168.
147. Shameem, I. Phytochemical & therapeutic potentials of Murr makki (*Commiphora myrrha*): A review. *Indian J. Appl. Res.* **2018**, *8*, 102–104.
148. Ashry, K.M.; El-Sayed, Y.S.; Khamiss, R.M.; El-Ashmawy, I.M. Oxidative stress and immunotoxic effects of lead and their amelioration with myrrh (*Commiphora molmol*) emulsion. *Food Chem. Toxicol.* **2010**, *48*, 236–241. [[CrossRef](#)] [[PubMed](#)]
149. Happy, A.A.; Jahan, F.; Momen, M.A. Essential Oils: Magical Ingredients for Skin Care. *J. Plant Sci.* **2021**, *9*, 54–64.
150. Chakravarty, N.; Kellogg, C.; Alvarez, J.; Equils, O.; Morgan, M. UV Protection by Natural Products: C. myrrha Oil Versus Sunscreen. *J. Drugs Dermatol.* **2018**, *17*, 905–907.
151. Hamidpour, R.; Hamidpour, S.; Hamidpour, M.; Shahleri, M. Frankincense (乳香 *Ru Xiāng*; *Boswellia* species): From the selection of traditional applications to the novel phytotherapy for the prevention and treatment of serious diseases. *J. Tradit. Complement. Med.* **2013**, *3*, 221–226. [[CrossRef](#)]
152. Mikhaeil, B.R.; Maatooq, G.T.; Badria, F.A.; Amer, M.M. Chemistry and immunomodulatory activity of frankincense oil. *Z. Nat. C* **2003**, *58*, 230–238. [[CrossRef](#)]
153. Eyre, H.; Hills, M.J.; Watkins, S.D. Compositions containing *Boswellia* extracts. *Quest Int. BV* **2003**, *6*, 516.
154. Michie, C.A.; Cooper, E. Frankincense and myrrh as remedies in children. *J. R. Soc. Med.* **1991**, *84*, 602. [[CrossRef](#)]
155. Kaur, A.; Singh, T.G.; Dhiman, S.; Arora, S.; Babbar, R. Novel herbs used in cosmetics for skin and hair care: A. review. *Plant Arch.* **2020**, *20*, 3784–3793.
156. Togni, S.; Maramaldi, G.; Di Piero, F.; Biondi, M. A cosmeceutical formulation based on boswellic acids for the treatment of erythematous eczema and psoriasis. *Clin. Cosmet. Invest. Dermatol.* **2014**, *7*, 321.
157. Sawicka, B.; Noaema, A. Cosmetics active ingredients used in the Middle East. In Proceedings of the II National Scientific Conference “Modern Technologies and Treatments in Cosmetology”, Tomsk, Russia, 29–30 September 2015.