



## REVIEW ARTICLE

## A REVIEW: THERAPEUTIC, MEDICINAL AND FOOD USES OF ALOE VERA

Fatma A. Ahmed<sup>\*1,2</sup> , Taha A.I. El-Bassossy<sup>1</sup> , Ahmed A.M. Abdelgawad<sup>1,3</sup>

<sup>1</sup>Medicinal and Aromatic Plants Department, Desert Research Center, 11753, Cairo, Egypt.

<sup>2</sup>Regional Development Centers (RDC), Academy of Scientific Research and Technology (ASRT), Egypt.

<sup>3</sup>Department of Chemistry, College of Science, Jazan University, Jizan, Kingdom of Saudi Arabia

## Article Info:

## Abstract



## Article History:

Received: 3 October 2023

Reviewed: 9 November 2023

Accepted: 26 December 2023

Published: 15 January 2024

## Cite this article:

Ahmed FA, AI El-Bassossy T, Abdelgawad AAM. Therapeutic, medicinal and food uses of *Aloe vera*: A review. Universal Journal of Pharmaceutical Research 2023; 8(6):72-81.

<https://doi.org/10.22270/ujpr.v8i6.1045>

## \*Address for Correspondence:

Taha AI El-Bassossy, Medicinal and Aromatic Plants Department, Desert Research Center, 11753, Cairo, Egypt. Tel- +20-1000028656.  
 E-mail: [tahachemist2008@gmail.com](mailto:tahachemist2008@gmail.com)

The history for the use of *Aloe vera* for medicinal purposes starts from about 3000 years. Medicinally, this plant may be able to treat skin cancer as well as sunburns, burns, and small wounds. More than 104 compounds from various parts of this plant, including minerals, vitamins, amino acids, enzymes, sterols, anthraquinone, flavonoids, terpenoids, coumarins, polysaccharides, sugars, and polyphenols, have been isolated thus far, according to scientific reports on phytochemical analysis of this plant. The biological activities of these compounds are diverse and include anthelmintic, hepatoprotective, antidiabetic, diuretic, antibacterial, antiviral, antioxidant, antiseptic, anti-inflammatory, anticancer, and cosmetic effects for medical therapy. This article mainly emphasizes therapeutic, medicinal and food uses of *A. vera*.

**Keywords:** *Aloe vera*, food, medicinal, therapeutic, flavonoids.

## INTRODUCTION

*Aloe* is an individual from the 420-species of family Xanthorrhoeaceae, which has been utilized for medicinal purposes for around 3,000 years. *Aloe* is a genus of succulent herbs that grow to a height of 80–100 cm. They matured in 4 to 6 years and can live for just about 50 years in the right circumstances. Among the *Aloe* species, *A. vera* (L.) Burm. f. syn. *A. barbadensis* M. is the most physiologically active species<sup>1,2</sup>. According to the World Health Organization, medicinal plants will be the primary source of many different medications<sup>3</sup>. The plant was originally found in southern and eastern Africa, in the Sudan along the upper Nile. After that, it was brought to northern Africa, where it spread naturally to other parts of Africa and the Mediterranean. The plant is grown for commercial use in Venezuela, Aruba, Bonaire, Haiti, India, South Africa, and the United States of America<sup>4</sup>. While Southern California deserts are home to the highest-quality aloe plants. As long as its roots are not damaged, the plant can endure underneath frigid temperatures and temperatures as high as 104°F.

*Aloe vera* (L.) Burm f. is a perpetual succulent xerophyte that produces tissue that holds water in its

leaves in order to withstand arid environments with little to no precipitation. Additionally, this is a well-known medicinal plant that has been used extensively to cure different ailments in traditional Chinese medicine. There is a continuous search for new anti-pathogen compounds using plant-based extracts. A substantial percentage of recently created antibiotics available for purchase come from sources that are semi-synthetic or natural, and about 20% of all plants have undergone pharmacological or biological research<sup>5</sup>. The motivation behind this review is to introduce an exhaustive update on the utilization of *A. vera* in medication, food, and treatment.

## Taxonomy

The following is the taxonomic classification of *A. vera*<sup>6</sup>:

Kingdom: Plantae	Subkingdom: Viridiplantae
Division: Tracheophyta	Class: Magnoliopsida
Order: Asparagales	Family: Xanthorrhoeaceae
Genus: <i>Aloe</i>	Species: <i>vera</i>
Binomial name: <i>Aloe vera</i> L.	

## Synonyms

*Aloe vera* is also known as (synonyms): *A. perfoliata* L., *A. barbadensis* Miller, *A. chinensis* Bak., *A. indica* Royle, *A. elongata* Murray, *A. officinalis* Forsk., *A. rubescens* DC, and *A. vera* L. var. *littoralis*. König ex

Bak., *L. var. chinensis*, *A. vera* Lam, *A. vulgaris* Berger. *A. vera* (L.) Burm. f. is considered a synonym, whereas *A. barbadensis* Mill. is considered the correct species name in most formularies and reference books. Depending to the International Rules of Botanical Nomenclature (IRBN), *A. vera* (L.) Burm. f. is the correct name for this species<sup>7</sup>.

#### Botanical description

A sessile, three-sided stem, a shallow underground root growth, and beefy, toothed passes on assembled in a rosette 30 to 50 cm long and 10 cm wide at the base describe this succulent perennial herb, which has a pea-green tone. The radiant yellow, cylindrical blossoms are 25-35 cm long, with an axillary spike and stamens that oftentimes arise outside the sprouting tube. There are a lot of seeds in the fruits<sup>4</sup>.

#### Active ingredients

More than one hundred potential active ingredients from six different classes make up the active components of *A. vera*: phenylpropanoids, flavonoids, and coumarins, phenylpyrone and phenol derivatives, phytosterols, anthraquinone and their glycosidic derivatives, chromone and its glycosidic derivatives, and others<sup>8,9</sup>. The leaf is transverse section displays three cell layers: the outer, protective layer, and the colorless inner layer<sup>10</sup>.

#### The outer protective layers of the leaf

According to several studies, derivatives of hydroxy anthracene, anthraquinone, and the glycosides aloin A and B are available in the severe yellow plastic of pericyclic tubules in the external layer of the leaves in levels going from 15 to 40%<sup>11</sup>. Substances such as moisture, fiber, ash, lipids, protein, organic acids, minerals, free sugars, and polysaccharides were identified through chemical analysis of the basic ingredients of leaves of *A. vera*. Fructose, glucose and sucrose were the three basic free sugars. The primary organic acids were lactic, fumaric, acetic, malic, lactone, citric, and oxalic acids. *A. vera* has been used to isolate and characterize approximately 29 chromone, 32 anthraquinones, 13 flavonoids, 12 phenylpropanoid acids, 4 coumarins, 3 phenylpyrone, 1 naphthalene triglycosylated (alloferoside A), 1 methyltetraline (ferroxidine), and their derivatives<sup>9</sup>.

#### The middle layer of the leaf

It is vital to recognize this juice from *A. vera* gel, which is a comparable boring adhesive gel produced using parenchymatous leaf cells. The pericycle and

adjacent leaf parenchyma cells are the source of this juice, which flows spontaneously from the sliced leaf. This gel can be dried without the use of heat. Water (>98%), polysaccharides (cellulose, pectins, glucomannan, hemicellulose, acemannan), lipids, proteins, vitamins, enzymes, inorganic compounds, amino acids, and phytosterols (cycloartanol, lophenol, 24-methylenecycloartanol, 24-ethyl-lophenol, and 24-methyllophenol) make up the majority of the chemical constitution of *A. vera* gel<sup>12</sup>. Emodin of aloe and the anthraquinone glycosides Aloin A and B are among the principal active ingredients from *A. vera* juice, which are hydroxy-anthracenic products, which make up 15 to 40% of the overall constituents<sup>13</sup>. Furthermore, it has been discovered that the components Al, B, Ba, Fe, Ca, Na, P, Mg, Si, etc are available in *A. vera* gel<sup>14,15</sup>.

#### The inner layers of the leaf

In the center district of the leaf, which is a clear, delicate, soggy, and tricky tissue, huge, slight walled parenchyma cells hold water as tacky adhesive<sup>16,17</sup>. The inward layer of the leaf gel contains up to close to 100% water, with amino acids, glucomannan, lipids, and sterols<sup>18</sup>, vitamins (B1, B2, B6, and C)<sup>19</sup>, many mono- and polysaccharides, a number of inorganic substances, enzymes (lactate dehydrogenase, amylase, lipase, acid and alkaline phosphatase), and organic substances (barbaloin, aloin, and emodin)<sup>20</sup>. A lengthy chain of acetylated mannose serves as *A. vera* is primary functional component<sup>21,22</sup>.

The polysaccharides in the parenchymatous tissue of the inward leaf of aloe plants have been linked to several of the therapeutic benefits of extracts from aloe leaves<sup>23,24</sup>, is thought that rather than being attributed to a single chemical component, these processes in biology ought to be attributed to the synergistic activity of the chemicals included within<sup>25</sup>.

#### *A. vera* root

Certain phenolic chemicals, particularly naphthaquinones and anthraquinones, have also been found in the case of *A. vera* root<sup>26-28</sup>.

#### Uses of ethnopharmacology

Historically, *A. vera* gel has been utilized topically to small burns, heal wounds, and skin irritations as well as orally to treat immune system deficiencies, coughing, constipation, ulcers, migraines, diabetes, and arthritis<sup>29</sup>. For millennia, *A. vera* has been treated medicinally in a several nations, including: Egypt, Greece, India, Japan, Mexico, and China<sup>30</sup>.

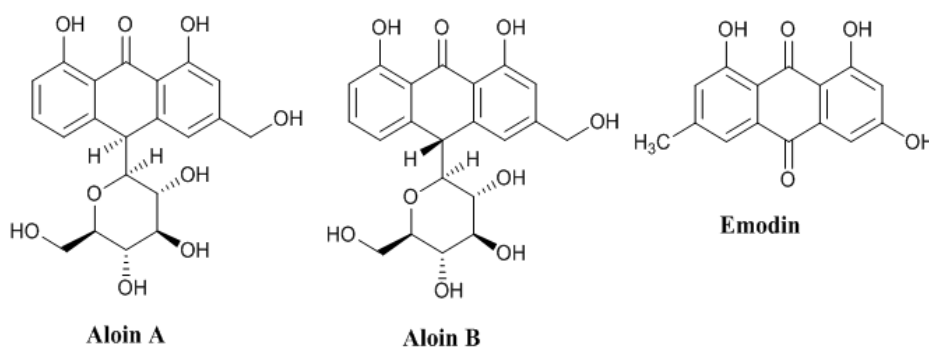


Figure 1: Potential active anthraquinone components of *A. vera*.

*A. vera* had been applied by the Egyptians to make papyrus-like scrolls and treat tuberculosis<sup>31</sup>. Nadkerni,<sup>32</sup> mentioned that a variety of *A. barbadensis* formulations, including lotions, confections, and juices, are effective treatments for a range of illnesses. Aloin, a combination of glucosides found in aloe, is the active ingredient in several medications. This plant is used to make Elio, an anthelmintic, purgative, and emmenagogue used to treat paediatric helminthiasis. Gel is helpful for pressure ulcers and ulcerative colitis<sup>33</sup>. The properties of *A. vera* include emmenagogue, deobstruent, stomachic, carminative, aperient, depurative, and diuretic effects. For almost 3,000 years, traditional medicine has made use of edible coating gels derived from the *A. vera* plant<sup>9</sup>.

## MEDICAL AND THERAPEUTIC USES

The present investigation aims to showcase the advantages and applications of leaf gel that have recently been identified. We shall provide a quick overview of *A. vera* known biological functions. According to certain claims, the polysaccharides present in the gel of *A. vera* have antibacterial, anti-inflammatory, radiation damage repair, antifungal, wound healing, immunostimulation, antiviral, antidiabetic, and antitumor activities, as well as antioxidant and hematopoiesis-stimulating effects. Furthermore, significant medicinal uses will be described, including the application of powdered dry *A. vera* gel as an excipient in measurements formulations for medications with prolonged release<sup>34,35</sup>.

### Anti-aging agent

The skin is capacity for moisture retention is attributed to muco-polysaccharides. Amino acids additionally mellow hard skin cells, and zinc goes about as an astringent to fix pores. Studies has demonstrated that using *A. vera* gel gloves to treat dry skin caused by industrial exposure can improve skin uprightness, decrease erythema, and lessen the emergence of acne-related wrinkles. *A. vera* gel hydrates and feels refreshing on the skin. It also contributes to skin regeneration and gerontology. Aloe is biogenic nature gives rise to this trait. In the cosmetics business, *A. vera* is used as a skin tonic<sup>1</sup>.

### Antibacterial activity

According to earlier research, using *A. vera* gel as an edible covering helps to minimize microbial spoiling and fruit deterioration. Nabigol and Asghari, discovered that *A. vera* gel repressed the development of mycelium (*Aspergillus niger* and *Penicillium digitatum*)<sup>36</sup>. It was discovered that *A. vera* gel at a dosage of 500 mL/L completely inhibited *P. digitatum* and 64% inhibited *A. niger*. In an alternative investigation<sup>37</sup>, *Shigella flexneri* and *Streptococcus progenies*, *S. aureus*, *P. aeruginosa*, *E. coli*, and *S. typhi* were reported to be inhibited in growth by *A. vera* leaf gel. There have likewise been reports of *A. vera* gel antibacterial properties against *Helicobacter pylori*<sup>38</sup>. Agarry et al.,<sup>39</sup> revealed that *Trichophyton mentagrophytes*' (20.0 mm) development was suppressed by the aloe gel. Conversely, *A. vera* extracts

did not demonstrate any antimicrobial activity against *Xanthomonas* species<sup>40</sup>.

Benitez et al.,<sup>41</sup> found that *A. vera* gel works better than chitosan and alginate to inhibit mesophilic bacteria, yeasts, and molds on kiwifruit slices. An *A. vera* gel covering was found to expand the timeframe of realistic usability of guava by approximately one week in a different study. This is because the edible covering inhibits the growth of microorganisms<sup>42</sup>. They were not knowing the precise method of action, but acemannan saponins, and anthraquinone derivatives found in *A. vera* are known to have antibacterial activity<sup>43</sup>. *A. vera* gel has inhibited two micro-organisms: *S. pyogenes* and *S. faecalis*<sup>44</sup>.

### Antidiabetic effects

An evaluation of the medical advantages of medicinal plants, including *A. vera*, focused on the function of active macromolecules with antidiabetic action. In mice with type-2 diabetes, the phytosterols and polysaccharides of *A. vera* increased insulin levels, which had anti-diabetic effects<sup>45-47</sup>. Clinical and experimental studies on *A. vera* sap have demonstrated a noteworthy hypoglycemic impact when consumed for 4-14 weeks<sup>48</sup>.

While some studies indicated that there may be no change in glucose levels, several preclinical (in animals) and clinical (in people) formulations have shown that *A. gelatinifera* formulations. *A. vera* in various structures (for example juice or as ingredients in baking, etc.) shows a reduction in the level of glucose in the blood. Different techniques for extracting and separating mucous *A. vera* gel from secretory anthraquinone may account for variations in the outcomes of these *in vivo* investigations. Furthermore, it can be challenging to connect the effect or lack of to the product in question because it isn't always evident which portion of an aloe leaf was assessed in a particular study. Through oral administration in rats of alcohol insoluble residue (gel of *A. vera*) extract led to a considerable reduction in hepatic transaminases, plasma and tissue cholesterol, triglycerides, free unsaturated fats, phospholipids, and fasting blood glucose, in addition to a noteworthy increase in plasma insulin levels. Following administration of the gel extract, mice treated with streptozotocin exhibited normal an increase in low-density lipoprotein cholesterol and high-density lipoprotein cholesterol levels in the plasma<sup>49</sup>.

### Antifungal activity

*A. vera* was tested for its antifungal properties on the mycelium development of *F. oxysporum*, *C. coccodes*, and *R. solani*. *F. oxysporum* was inhibited by the pulp at 104  $\mu\text{L}^{-1}$ , whereas *F. oxysporum*, *R. solani*, and *C. coccodes* showed lower colony growth rates in the liquid fraction at a concentration of 105  $\mu\text{L}^{-150}$ . Sitara et al.<sup>51</sup>, stated that, *A. niger*, *A. alternata*, *A. flavus*, *D. hawaiiensis*, and *P. digitatum* are the five plant pathogenic fungi that were the subject of a thorough investigation of the three separate doses of the antifungal ability of *A. vera* gel. It has been found that *D. hawaiiensis* and *A. alternata* growth is entirely inhibited by the maximum test dosage of *A. vera* gel (0.35%). As per an alternative investigation, the base

fungicidal groupings of *A. vera* against *P. gladioli*, *F. oxysporum*, *H. pruneti*, and *B. gladiolorum* varied based on the kind of fungus species and ranged from 80 to 100  $\mu\text{L}/\text{mL}$ <sup>52</sup>.

Additionally, prior research has demonstrated that combining *A. vera* gel with various homogenizers, including 0.15 g of glycerol starch, enhances the effectiveness of preventing fungal rot and weight reduction in cherry tomatoes<sup>53</sup>. Navarro *et al.*<sup>54</sup>, led a concentrate on nectarines using *A. vera* gel on alone or in conjunction with thymol, and saw that as the *A. vera* gel alone is more powerful at preventing the decay brought on by *P. digitatum*, *B. cinerea*, and *R. stolonifer*. In a prior study, *A. vera* gel coatings were examined for their ability to prevent deterioration and were found to dramatically reduce the amounts of yeast, mould, and mesophilic bacteria in a variety of vegetables and fruits, including tomatoes<sup>55,56</sup>, citrus fruits<sup>57</sup>, berries fruits<sup>58</sup>, blueberry<sup>59</sup>, strawberry<sup>60</sup>, and ready-to-eat pomegranate seeds<sup>61</sup>. A prepared *A. vera* gel slowed the growth of the *C. albicans* fungus<sup>62</sup>.

#### Anti-inflammatory Action

Bradykinase activity has been employed in research conducted *in vitro* and *in vivo* that demonstrate the gel of *A. vera* has beneficial anti-inflammatory properties<sup>63</sup>. Bradykinin is a fiery compound that causes torment, and the peptidase bradykinase that was segregated from aloe has been shown to separate it<sup>64,65</sup>. The active components of *A. vera* include sterols ( $\beta$ -sitosterol, campesterol, lupeol, and cholesterol) and mannose-6-phosphate<sup>65</sup> which have anti-inflammatory properties, assist in lowering pain and serving as a natural painkiller. Aloe has additional aspirin-like substance that gives it antibacterial and anti-inflammatory qualities. *A. vera* diminishes the union of prostaglandin E2 from arachidonic corrosive and hinders the cyclo-oxygenase pathway. Rats with carrageenin-induced paw oedema showed a substantial reduction in acute inflammation when treated with fresh *A. vera* gel, but not in chronic inflammation<sup>66</sup>.

The oedema generation was observed to be inhibited by *A. vera* aqueous and chloroform extracts in a manner comparable to those of popular anti-inflammatory medications (i.e. dexamethasone and indomethacin). Additionally, there was a strong correlation observed between the counter oedema properties of these two concentrates and their ability to diminish the number of neutrophils that move into the peritoneal depression<sup>67</sup>. An adjuvant-induced rat model of arthritic inflammation showed a 48% reduction in inflammation with adding 5.0% leaf homogenate from *A. vera* leave<sup>68</sup>. Potential exists for using *A. vera* to treat the inflammatory response of the stomach mucosa brought on by an *H. pylori* infection<sup>69</sup>.

#### Antioxidant effects

Several authors have reported that the unfractionated entire gel and the fractions of *A. vera* contain antioxidant properties. The cell reinforcement properties of *A. vera* gel might be because of the presence of phenolic cancer prevention agents, superoxide dismutase catalysts, and glutathione peroxidase movement. *A. vera* gel dose dependent antioxidant activity was shown by incubating

inflammatory colon mucosal biopsies and employing two cell-free *in vitro* techniques. This study evaluated the scavenging of peroxy and superoxide radicals using cell-free methodologies. *A. vera* gel likewise inhibited the production of prostaglandin E2 from inflammatory colorectal biopsies at a dosage of 1 in 50, while thromboxane B2 release remained unaltered<sup>70</sup>.

#### Antiseptic properties

Six germ-free specialists called lupeol, urea nitrogen, salicylic corrosive, cinnamonic corrosive, sulfur and phenols are what give *A. vera* its antiseptic properties. These substances inhibit the growth of bacteria, viruses, and fungus. Even though the majority of these applications are intriguing, more research is necessary to evaluate its efficacy across all illnesses<sup>71</sup>.

#### Anti-stress effect

*A. vera* juice is helpful for the legitimate working of the body frameworks<sup>72</sup>. It reduces the process of cells being damaged under stress and lessens the body physiological and biochemical alterations<sup>73</sup>. Synthetic responses that change a compounds oxidative state are alluded to as oxidative pressure. Certain cancer prevention agents are tracked down in the body normal administrative device, though dietary cell reinforcements come from food sources. *A. vera* is a prime example of a useful food that helps protect against oxidative stress<sup>74</sup>.

#### Antitumor activity

The primary purposes of *A. vera* gel are defence, hydration, insulation, and complement activation associated with polysaccharides. *In vitro* application of fresh gel induced both growth and adhesion of normal human cells, while a settled gel planning demonstrated cytotoxic to growth and typical human cells. The additional chemicals introduced to the gel during processing were blamed for this cytotoxicity<sup>75</sup>.

The anticancer and antiulcer properties of *A. vera* gel glycoproteins have additionally been displayed to support the development of solid human skin cells. Research indicates that benzopyrene cannot bind to primary rat hepatocytes due to the presence of the polysaccharide fraction, which inhibits the production of benzopyrene-DNA adducts that may cause cancer. There have also been reports of myristicphorbol acetate's tumor-promoting properties being inhibited and glutathione S-transferase being increased, which suggests that *A. vera* gel might be helpful in the chemoprevention of cancer<sup>76</sup>. Anthraquinone *A. vera* emodin has antineoplastic qualities since it can stop or slow the proliferation of malignant cancer cells. Nevertheless, the usefulness of *A. vera* gel on human health has been the subject of incredibly few and often contradicting statistically significant clinical trials<sup>77</sup>.

#### Antiviral Activity

Many researchers have also been interested in *A. vera* is antiviral properties because of its purported beneficial effects against strains of the HSV type 2 herpes simplex virus by Keivan *et al.*<sup>78</sup>, as well as against influenza A virus spreads by Li *et al.*<sup>79</sup>. It has been shown that a few parts of *A. vera* gel are strong antiviral specialists. Herpes simplex contamination was diminished in two developed target cell lines by acemannan<sup>80</sup>. Parts of *A. vera* gel called lectins straight

forwardly kept the cytomegalovirus from multiplying in cell culture, perhaps by blocking the blend of proteins<sup>81</sup>. Pure aloe emodin has been shown to be effective against all viruses, including the influenza, varicella, and pseudorabies viruses, as well as herpes simplex viruses of Type I and Type II. Electron microscopy study of the herpes simplex virus treated with anthraquinone revealed signs of partially torn envelopes. According to these results, anthraquinones that have been isolated from a variety of plants directly inhibit viruses that are enclosed. These actions may be due to indirect effect due to stimulation of the immune system. Numerous enveloped viruses, including varicella zoster, herpes simplex virus, and influenza, are similarly rendered inactive by anthraquinone aloin<sup>82,83</sup>.

#### **Application of cosmetic and skin protection**

*A. vera* is widely used as cosmetics and nutritional medicines<sup>84</sup>, as well as a protection against skin damage caused by radiation<sup>85</sup>. The skin releases the antioxidant protein metallothionein after applying *A. vera* gel. This protein absorbs hydroxyl radicals and prevents the skin from producing glutathione peroxidase and superoxide dismutase. Suppressive of the immune system cytokines, as interleukin-10 (IL-10), are produced when skin keratinocytes are inhibited. This terminates the inhibition of delayed type hypersensitivity caused by the UV<sup>86</sup>. *A. vera* gel has been applied topically to dermabraded skin, and various researchers have reported occurrences of burning skin sensations and contact dermatitis. It seems that these reactions were related to the anthraquinone pollutants present in this mixture<sup>87</sup>. Aloin and its gel are applied topically as a skin tonic for acne. Additionally, aloe sugars are utilised in moisturising formulas. Blended with specific essential oils, it creates a fantastic moisturiser that smoothes the skin, a sunscreen lotion that blocks UV rays, and a variety of other cosmetic items. *A. vera* concentrates might be helpful in the treatment of benign skin cysts, boils, and other minor skin disorders since it has been demonstrated that they suppress the development of organisms that cause fungus<sup>88</sup>.

#### **Effect on the secretion of gastric acid and ulcers**

Animal and human stomach ulcers have reportedly been treated or prevented with *A. vera* gel. Additionally, it was shown that ethanol-induced stomach ulcers in rats could not be stopped by aloe gel. Numerous possible mechanisms, such as those that lower inflammation, encourage healing, increase mucus production, and control stomach secretions, have been connected to *A. vera* anti-ulcer properties<sup>89</sup>. It was shown that either direct collaboration with the corrosive producing cells or conceivable connection with H<sub>2</sub>-receptors on the parietal cells was responsible for the concentration-dependent decrease of stomach acid discharges observed in the *A. vera* water-based extract made with ethanol. Gestural protective effect was only shown at the lowest tested dose. Some ideas suggest that at this low concentration, the *A. vera* extract has a cytoprotective effect, mitigating mucosal damage by a different mechanism than stomach acid neutralisation and inhibition. The mechanism of

cytoprotection has been the subject of numerous theories, some of which include increased mucus production, further developed mucosal blood stream, and expanded phospholipid content in the mucosal layer<sup>90</sup>.

#### **Hepatoprotective activities**

Aqueous extract from *A. vera* aerial dried parts reversed some biochemical parameters and dramatically decreased the liver damage caused by carbon tetrachloride in mice. Histopathological examinations verified the healing effectiveness of *A. vera* water extract against liver damage caused by carbon tetrachloride, as demonstrated by the relapse of centrilobular rot, full scale vascular greasy modifications, and scattered lymphomononuclear cell penetration in the hepatic parenchyma. Additionally, a rise in bile solids and bile flow indicates that the extract therapy appears to boost the secretory function from liver cells. The antioxidant activity of the hepatoprotective function was also linked to the preservation of the liver metabolising enzymes<sup>91</sup>.

#### **Immunomodulatory Effects**

The immunomodulatory properties of the sugars in *A. vera* gel, namely acemannan, have been shown in a number of studies. These studies suggest that activation of macrophage cells, characterized by cell surface markers and cytokine release (TNF- $\alpha$ , IL-1, IL-6, interleukin-1 or IL-1, and interferon- $\gamma$  or INF- $\gamma$ ), is what causes these actions<sup>92-94</sup>. Numerous low-molecular-weight human activated neutrophils can also be prevented from releasing reactive oxygen free radicals by substances<sup>95</sup>. Certain immune-modulatory effects have been demonstrated to be associated with aloe gel's glycoproteins, specifically lectins. Alprogen prevents calcium from entering mast cells, which prevents mast cell production of leukotriene and histamine through the action of antigen-antibody complexes<sup>96</sup>.

It was shown that applying aloe gel after UV exposure can avoid the reduction of both local and systemic immunity as well as postpone certain types of hypersensitivity reactions to alloantigens and *Candida albicans*. The polysaccharides in the gel have an immune-protective impact, although the mechanism is different from that of antioxidants, anti-inflammatories, and DNA-repair enzymes. *A. vera* has demonstrated anti-inflammatory characteristics compounds, but the polysaccharides did not successfully lessen UV-induced edema and inflammation or speed up the removal and repair of UV-induced cyclo-butyl pyrimidine dimers. Furthermore, the effectiveness of antioxidants requires their presence in the skin prior to UV radiation, whereas aloe polysaccharides continue to work even 24 hours after UV exposure. Therefore, the immunological defense mechanism acts downstream of DNA damage and repair, potentially through modifying signal transduction pathways that are triggered by DNA damage. Therefore, the component of activity of sugars was made sense of through their impact on antigen-introducing cells and the cytokine series<sup>97</sup>.

### Laxative effects

Anthraquinone, a potent laxative included in latex, is known to increment digestive water content, actuate bodily fluid discharge, and work with gastrointestinal peristalsis<sup>98</sup>. The 1, 8-dihydroxy-anthracene glycosides, or aloin A and B, were originally known as barbaloin and are mainly responsible for the aloe<sup>63</sup>. The dynamic metabolites (aloe-emodin 9-anthrone being the prevalent dynamic metabolite) are produced in the colon by intestinal bacteria hydrolyzing aloins A and B, which are not caught up in the upper digestive tract, following oral medication<sup>66</sup>, similar to senna, it both stimulates and irritates the digestive tract. *A. vera* plastic is commonly known for its purgative properties. Seldom does aloe work as a laxative until six hours after oral use, and perhaps not at all. It can take up to 24 hours to complete this.

### Moisturizing and skin hydration effects

The moisturizing properties of *A. vera* gel demonstrated that the only formulations that raised the stratum corneum's water content after just one application were those with more noteworthy focuses (0.25% w/w and 0.5% w/w). All of the formulations, including those with concentrations of 0.1% w/w, 0.25% w/w, and 0.5% w/w of gel *A. vera* powder, showed the same results after two weeks of application twice daily. Comparing the *A. vera* gel formulations to the vehicle employed in the formulations, however, revealed no difference in the trans epidermal water loss. It was suggested that the items containing *A. vera* gel further developed skin hydration conceivably through a humectant system<sup>99</sup>.

### Wound healing effects

The powerful course of wound mending happens in three phases. The main stage was portrayed by aggravation, hyperemia, and leukocyte invasion. The subsequent step requires the expulsion of dead tissue. The third phase of multiplication is described by the development of stringy tissue and epithelial recovery<sup>100</sup>. As indicated by a later assessment, body of evidence favours using *A. vera* to treat burns that are between the first and second degree<sup>101</sup>.

Acemannan<sup>21,22</sup>, glucomannan, mannose-6-phosphate, glycoprotein<sup>102</sup> gibberellins and plant growth hormone are the main functional constituents of *A. vera*, which accelerate wound healing, interact with fibroblast growth factor receptors, and reduce radiation-induced skin reactions<sup>103</sup>, promote the fibrogenic cytokines to be released<sup>98</sup>, and encourage prolonged granulation tissue stimulation. Aloe extract treatment affects collagen cross-linking and composition (more type III), which improves breaking strength and wound contraction<sup>18</sup>. In the granulation tissue of the healing wound, it also promotes the synthesis of dermatan sulphate and hyaluronic acid<sup>104</sup>.

Aloe gel has been utilized to treat radiation consumes and radiation ulcers; two radiation consume patients have shown full recuperation subsequent to utilizing this therapy. When contrasted with consumes treated with oil jam cloth (18.2 days), sores treated with new aloe gel recuperated quicker (11.8 days) than those treated with cream<sup>105,106</sup>. A widely available product is powdered *A. vera* concentrate. To balance the dynamic

cutaneous ischemia brought about by consumes, frostbite, electrical shocks, and intra-blood vessel drug use, aloe gel is utilized topically. These gel capabilities as an inhibitor of thromboxane A<sub>2</sub>, a middle person of improved tissue harm, as per an *in vivo* assessment of these wounds<sup>107</sup>.

### Adverse reactions

An overdose might result in colicky stomach spasms and discomfort, as well as the creation of thin, watery faeces, stomach spasms and pain can happen even after just one dose. Chronic overuse of laxatives containing anthraquinone stimulants can result in hepatitis<sup>108</sup> as well as abnormalities related to electrolytes, malabsorption, loss of weight, albuminuria, metabolic acidosis, and haemorrhage<sup>109</sup>. Recurring usage of stimulant laxatives may aggravate older individual's weakness and orthostatic hypotension. Following an increase in dosage, renal tubular injury may result in secondary aldosteronism. There have also been reports of osteomalacia of the vertebral column, increased discharge of calcium in the stools, and protein-losing gastroenteropathy with hypoalbuminemia and diarrhoea. People who use anthraquinone laxatives for prolonged periods of time have been seen to have pseudomelanotic colouring of the intestinal mucosa (*Pseudomelanosis coli*). When the medication is stopped, the pigmentation is normally reversible within 4 to 12 months and is clinically harmless<sup>110</sup>.

### Uses in the food industries

It has been employed in the food manufacturing industry to make useful food sources, as a fixing in other food items, and to make gel-based wellbeing beverages and refreshments<sup>35</sup>. Edible coatings leave a thin film on the fruit's surface that keeps moisture and gases from the atmosphere at bay<sup>111,112</sup>. Aloe gels serve in food preservation by reducing fresh produce's respiration and transpiration and delaying the deterioration of food after harvest<sup>113</sup>. Many researchers have been done on the utilization of *A. vera* gel after harvest to date<sup>9</sup>.

### Uses in pharmaceutical industries

*A. vera* has been utilized in the pharmaceutical industry to produce tablets and cases as well as skin prescriptions including salves and gel plans<sup>114</sup>. Recently, significant medicinal effects of *A. vera* gel and entire leaf remove were viewed as ready to build the bioavailability of co-regulated nutrients in human subjects<sup>115</sup>. Because of the ways in which *A. vera* gel enhances absorption, it can be used to efficiently provide medications that are not well absorbed orally. Moreover, direct compressible matrix type tablets were effectively made using the dehydrated powder that was extracted from *A. vera* gel. Due to their ability to discharge a model compound slowly over a drawn out timeframe, these grid style tablets have shown potential for use as an excipient in changed discharge dose structures<sup>116</sup>.

### Other uses

*A. vera* extracts can also be used to dilute semen for artificially fertilizing sheep and as a preservative for fresh food<sup>43</sup> and applied in small farms to conserve water. The gel of *A. vera* contains compounds called soapy saponins that have antibacterial and cleaning

qualities. Strong anti-microbial activity of the saponins is demonstrated against bacteria, viruses, fungi, and yeasts<sup>117</sup>.

#### Usage risks

Patients with incendiary gut conditions, like diverticulitis, Crohn's illness, ulcerative colitis, peevish inside disorder, or a ruptured appendix, shouldn't accept aloe; Aloe should also not be taken by children under the age of ten. After weighing the benefits and risks, *A. vera* should only be used under medical supervision during pregnancy or nursing. People encountering cramps, colic, hemorrhoids, nephritis, or other unexplained gastrointestinal side effects like uneasiness, sickness, or retching ought to try not to utilize aloe<sup>118</sup>.

#### CONCLUSIONS

*A. vera* is beneficial to humans in daily life because it relieves a range of skin conditions, including minor cuts, bug stings, wounds, poison ivy, and dermatitis. It also moisturizes and prevents ageing of the skin, supports the health of the digestive system, blood and lymphatic circulation, and supports the function of the liver, kidney, and gall bladder. *A. vera* is known as the "wonder plant" for a variety of uses, counting as an antibacterial, a calming, a therapy for diabetes and disease, and a restorative. Increased research effort on the plant is necessary to improve its utility for human use. Without a doubt, *A. vera* is a gift from nature to humans that may be used for therapeutic, cosmetic, and burn purposes. It is up to us to learn more about this plant and to give thanks to the natural world for its endless supply.

#### ACKNOWLEDGEMENTS

Academy of Scientific Research and Technology is acknowledged by the authors for providing adequate funding to carry out this research under the grant named "Maximize the Utilization of Succulent Plants for Community Development in Matrouh Governorate".

#### AUTHORS' CONTRIBUTIONS

**Ahmed FA:** writing original draft, conceptualization., **AI El-Bassossy T:** supervision, editing. **Abdelgawad AAM:** writing, review, and editing. The final manuscript was read and approved by all authors.

#### DATA AVAILABILITY

The data supporting the findings of this study are not currently available in a public repository but can be made available upon request to the corresponding author.

#### CONFLICT OF INTEREST

None to declare.

#### REFERENCES

- West DP, Zhu YF. Evaluation of *Aloe vera* gel gloves in the treatment of dry skin associated with occupational exposure. *Am J Infect Control* 2003;31(1):40-42. <https://doi.org/10.1067/mic.2003.12>
- Yagi A, Kabash A, Mizuno K, Moustafa SM, Khalifa TI, Tsuji H. Radical scavenging glycoprotein inhibiting Cyclooxygenase-2 and Thromboxane A2 Synthase from *Aloe vera* Gel. *Planta Med* 2003; 69(3):269-271. <https://doi.org/10.1055/s-2003-38481>
- Lanka S. A review on *Aloe vera*- The wonder medicinal plant. *J Drug Deliv Ther* 2018; 8(5-s):94-99.4. <http://dx.doi.org/10.22270/jddt.v8i5-s.1962>
- Yeh GY, Eisenberg DM, Kaptchuk TJ, Phillips RS. Systematic review of herbs and dietary supplements for glycemic control in diabetes. *Diabetes Care* 2003; 26(4):1277-1294. <https://doi.org/10.2337/diacare.26.4.1277>
- Mothana RA, Linclequist U. Antimicrobial activity of some medicinal plants of the Island Soqatra. *J Ethnopharmacol* 2005;96(1-2):177-181 <https://doi.org/10.1016/j.jep.2004.09.006>
- Hossain M, Mamunor Rashid A, Towfique N, Sen M. A review on ethnopharmacological potential of *Aloe vera* L. *J Intercult Ethnopharmacol* 2013;2(2):113. <http://dx.doi.org/10.5455/jice.20130612035300>
- African Pharmacopoeia. Vol. 1, 1<sup>st</sup> ed. Organization of African Unity, Scientific, Technical & Research Commission, Lagos, 1985.8. Choi SW, Son BW, Son YS, Park YI, Lee SK, Chung MH. The wound-healing effect of a glycoprotein fraction isolated from *Aloe vera*. *Br J Dermatol* 2001;145(4):535-545. <https://doi.org/10.1046/j.1365-2133.2001.04410.x>
- Choi SW, Son BW, Son YS, Park YI, Lee SK, Chung MH. The wound-healing effect of a glycoprotein fraction isolated from *Aloe vera*. *British J Dermatol* 2001; 145(4): 535-545. <https://doi.org/10.1046/j.1365-2133.2001.04410.x>
- Kahramanoğlu İ, Chen C, Chen J, Wan C. Chemical constituents, antimicrobial activity, and food preservative characteristics of *Aloe vera* gel. *Agronomy* 2019; 9(12):831.10. <https://doi.org/10.3390/agronomy9120831>
- Brown JP. A Review of the genetic effects of naturally occurring flavonoids, anthraquinones and related compounds. *Mutation Res* 1980; 75(3): 243-277. [https://doi.org/10.1016/0165-1110\(80\)90029-9](https://doi.org/10.1016/0165-1110(80)90029-9)
- Saccu D, Bogoni P, Procida G. Aloe Exudate: Characterization by reversed phase HPLC and headspace GC-MS. *J Agric Food Chem* 2001; 49(10):4526-4530.
- Tanaka M, Misawa E, Ito Y, *et al.* Identification of five phytosterols from *Aloe vera* gel as anti-diabetic compounds. *Biol Pharm Bull* 2006;29(7):1418-1422.
- Bozzi A, Perrin C, Austin S, Arce Vera F. Quality and authenticity of commercial *Aloe vera* gel powders. *Food Chem* 2007; 103(1):22-30.14. <https://doi.org/10.1016/j.foodchem.2006.05.061>
- Yamaguchi T, Takamura H, Matoba T, and Terao J. HPLC Method for evaluation of the free radical scavenging activity of foods by using 1,1-Diphenyl-2- Picrylhydrazyl. *Biosci Biotech Biochem* 1998; 62(6): 1201-1204. <https://doi.org/10.1271/bbb.62.1201>
- Femenia A, Sanchez ES, Simal S, Rossello C. Compositional features of polysaccharides from *Aloe vera* (*Aloe barbadensis* Miller) plant tissues. *Carbohydrate Polymers* 1999; 39(2):109-117.
- Newton LE. Aloes in habitat. In: Reynolds T, ed. *Aloes: the genus Aloe*. CRC Press: Boca Raton; 2004:3-36.
- López A, de Tangil M, Vega-Orellana O, Ramírez A, Rico M. Phenolic constituents, antioxidant and preliminary antimycoplasmic activities of leaf skin and flowers of *Aloe vera* (L.) Burm. f. (syn. *A. barbadensis* Mill.) from the Canary Islands (Spain). <https://doi.org/10.3390/molecules18054942>

18. Reynolds T, Dweck AC. *Aloe vera* leaf gel: A review update. *J Ethnopharmacol* 1999; 68(1-3): 3-37. [https://doi.org/10.1016/s0378-8741\(99\)00085-9](https://doi.org/10.1016/s0378-8741(99)00085-9)
19. Vogler BK, Ernst E. *Aloe vera*: A systematic review of its clinical effectiveness. *The British J General Pract* 1999; 49(447): 823-828.
20. Hayes SM. Lichen Planus: Report of successful treatment with *Aloe vera*. *General Dentistry* 1999; 47(3): 268-272.
21. Djeraba A, Quere P. *In vivo* macrophage activation in chickens with acemannan, a complex carbohydrate extracted from *Aloe vera*. *Int J Immunopharmacol* 2000; 22(5): 365-372. [https://doi.org/10.1016/s0192-0561\(99\)00091-0](https://doi.org/10.1016/s0192-0561(99)00091-0)
22. Lee JK, Lee, MK, et al. Acemannan purified from *Aloe vera* induces phenotypic and functional maturation of immature dendritic cells. *Int Immunopharmacol* 2001; 1(7): 1275-1284. [https://doi.org/10.1016/S1567-5769\(01\)00052-2](https://doi.org/10.1016/S1567-5769(01)00052-2)
23. Habeeb F, Shakir E, Bradbury F, et al. Screening methods used to determine the anti-microbial properties of *Aloe vera* inner gel. *Methods* 2007; 42(4): 315-320. <https://doi.org/10.1016/j.ymeth.2007.03.004>
24. Ni Y, Turner D, Yates KM, Tizard I. Isolation and characterization of structural components of *Aloe vera* L. leaf pulp. *Int Immunopharmacol* 2004; 4(14): 1745-1755. <https://doi.org/10.1016/j.intimp.2004.07.006>
25. Dagne E, Bisrat D, Viljoen A, Van WykBE. Chemistry of aloe species. *Current Org Chem* 2000; 4(10): 1055-1078. <http://dx.doi.org/10.2174/1385272003375932>
26. Lobo R, Prabhu KS, Shirwaikar A, et al. A HPTLC densitometric method for the determination of aloeverose in *Aloe vera* gel. *Fitoterapia* 2010; 81(4): 231-233. <https://doi.org/10.1016/j.fitote.2009.09.001>
27. Induli M, Cheloti M, Wasuna A, et al. Naphthoquinones from the roots of *Aloe secundiflora*. *Phytochem Lett* 2012; 5(3): 506-509. <https://doi.org/10.1016/j.phytol.2012.04.014>
28. Quispe C, Villalobos M, Bórquez J, Simirgiotis M. Chemical composition and antioxidant activity of *Aloe vera* from the Pica Oasis (Tarapacá, Chile) by UHPLC-Q/Orbitrap/MS/MS. *J Chem* 2018; 1-12. <https://doi.org/10.1155/2018/6123850>
29. Eshun K, He Q. *Aloe vera*: A valuable ingredient for the food, pharmaceutical and cosmetic industries- A review. *Crit Rev Food Sci Nutrition* 2004; 44(2): 91-96. <http://dx.doi.org/10.1080/10408690490424694>
30. Marshall JM. *Aloe vera* gel: What is the evidence?" *The Pharm J* 1990; 24: 360-362.
31. Baker OT. *The Amazing Ancient to Modern Useful Plant Aloe vera: Amazing Plant of the Magic Valley*, R. Prevost, Lemon Grove, 1975.
32. Nadkerni KM. *Indian Materia Medica*, 3<sup>rd</sup> Edition, Bombay Popular Prakashan Private Limited, Mumbai, 1976.
33. Langmead L, Feakins RM, and Goldthorpe S. Randomized, double blind, placebo-controlled trial of oral *Aloe vera* gel for active ulcerative colitis. *Alimentary Pharmacol Therap* 2004; 19(7): 739-747. <http://dx.doi.org/10.1111/j.1365-2036.2004.01902.x>
34. Boudreau MD, Beland FA. An evaluation of the biological and toxicological properties of *Aloe barbadensis* (Miller), *Aloe vera*. *J Environmen Sci Health, Part C* 2006; 24(1): 103-154. <https://doi.org/10.1080/10590500600614303>
35. Hamman J. Composition and applications of *Aloe vera* leaf gel. *Molecules* 2008; 13(8): 1599-1616. <https://doi.org/10.3390/molecules13081599>
36. Nabigol A, Asghari A. Antifungal activity of *Aloe vera* gel on quality of minimally processed pomegranate arils. *Int J Agronomy Plant Produc* 2013; 4(4): 833-838.
37. Ferro VA, Bradbury F, Cameron P, et al. *In vitro* susceptibilities of *Shigella flexneri* and *Streptococcus pyogenes* to inner gel of *Aloe barbadensis* Miller. *Antimicro Agents Chemother* 2003; 47: 1137-1139. <http://dx.doi.org/10.1128/AAC.47.3.1137-1139.2003>
38. Cellini L, Di Bartolomeo S, Di Campli E, et al. *In vitro* activity of *Aloe vera* inner gel against *Helicobacter pylori* strains. *Letters App Microbiol* 2014; 59(1): 43-48. <https://doi.org/10.1111/lam.12241>
39. Agarry OO, Olaleye MT, Bello-Michael CO. Comparative antimicrobial activities of *Aloe vera* gel and leaf. *African J Biotech* 2005; 4(12): 1413-1414.
40. Satish S, Raveesha KA, Janardhana GR. Antibacterial activity of plant extracts on phytopathogenic *Xanthomonas campestris* Pathovars. *Lett Appl Microbiol* 1999; (28)2: 145-147. <http://dx.doi.org/10.1046/j.1365-2672.1999.00479.x>
41. Benítez S, Achaerandio I, Pujolà M, Sepulcre F. *Aloe vera* as an alternative to traditional edible coatings used in fresh-cut fruits: A case of study with kiwifruit slices. *LWT - Food Sci Tech* 2015;61(1): 184-193. <https://doi.org/10.1016/j.lwt.2014.11.036>
42. Krishnan SA, Ullas A, Sagarika N, Oommen TE, Sunaila K. Development of *Aloe vera* based edible coating. *Int J Pure App Biosci* 2017; 5: 796-801.
43. Serrano M, Miguel J, Guillen F, Castillo S, Martinez-Romero D, Valero D. Use of *Aloe vera* gel coating preserves the functional properties of table grapes. *J Agri Food Chem* 2006; 54(11): 3882-3886. <http://dx.doi.org/10.1021/jf060168p>
44. Robson MC, Hegggers JP, Hagstrom WJ. Myth, magic, witchcraft or fact? *Aloe vera* revisited. *J Burn Care Res* 1982; 3(3): 157- 163. <http://dx.doi.org/10.1097/00004630-198205000-00005>
45. Yagi A, Sato Y, Miwa Y, et al. Ribosomal DNA sequence analysis of different geographically distributed *Aloe vera* Plants: Comparison with clonally regenerated plants. *Saudi Pharm J* 2006; 14( 3-4): 208-211.
46. Noor A, Gunasekaran S, Manickam AS, Vijayalakshmi MA. Antidiabetic activity of *Aloe vera* and histology of organs in streptozotocin-induced diabetic rats. *Curr Sci* 2008; 94(8):1070- 1076.
47. Arokiyaraj S, RadhaR S, Martin S, Perinbam K. Phytochemical analysis and anti-diabetic activity of *Cadaba fruticososa*. *R. Br Indian J Sci Tech* 2008; 1(6): 1-4.
48. Ghannam N, Kingston M, Al-Meshaal IA, Tariq M, Parman NS, Woodhouse N. The antidiabetic activity of Aloes: Preliminary clinical and experimental observations. *Hormone Res* 1986; 24(4): 286-294. <http://dx.doi.org/10.1159/000180569>
49. Rajasekaran S, Ravi K, Sivagnanam K, Subramanian S. Beneficial effects of *Aloe vera* leaf gel extract on lipid profile status in rats with streptozotocin diabetes. *Clin Exp Pharmacol Physiol* 2006; 33: 232-237.
50. de Rodríguez DJ, Hernández-Castillo D, Rodríguez García R, Angulo-Sanchez JL. Antifungal activity *in vitro* of *Aloe vera* pulp and liquid fraction against plant pathogenic fungi. *Indust Crops Products* 2005; 21(1): 81-87. <http://dx.doi.org/10.1016/j.indcrop.2004.01.002>
51. Sitara U, Hassan N, Naseem J. Antifungal activity of *Aloe vera* gel against plant pathogenic fungi. *Pak J Bot* 2011; 43: 2231-2233.
52. Rosca-Casian O, Parvu M, Vlase L, Tamas M. Antifungal activity of *Aloe vera* leaves. *Fitoterapia* 2007; 78(3): 219-222. <https://doi.org/10.1016/j.fitote.2006.11.008>
53. Ortega-Toro R, Collazo-Bigliardi S, Roselló J, Santamarina P, Chiralt A. Antifungal starch-based edible films containing *Aloe vera*. *Food Hydrocoll* 2017; 72: 1-10. <https://doi.org/10.1016/j.foodhyd.2017.05.023>
54. Navarro D, Díaz-Mula HM, Guillén F, et al. Reduction of nectarine decay caused by *Rhizopus stolonifer*, *Botrytis cinerea* and *Penicillium digitatum* with *Aloe vera* gel alone or with the addition of thymol. *Int J Food Microbiol* 2011; 151(2): 241-246. <https://doi.org/10.1016/j.ijfoodmicro.2011.09.009>
55. García MA, Ventosa M, Díaz R, Falco S, Casariego A. Effects of *Aloe vera* coating on post harvest quality of tomato. *Fruits* 2014; 69(2): 117-126. <https://doi.org/10.1051/fruits/2014001>
56. Chrysargyris A, Nikou A, Tzortzakis N. Effectiveness of *Aloe vera* gel coating for maintaining tomato fruit quality. *New Zealand J Crop Horti Sci* 2016; 44(3): 203-217. <https://doi.org/10.1080/01140671.2016.1181661>



57. Jhalegar Md, Sharma J, Singh D. Antifungal efficacy of botanicals against major postharvest pathogens of Kinnow mandarin and their use to maintain postharvest quality. *Fruits* 2014; 69(3): 223–237.  
<https://doi.org/10.1051/fruits/2014012>
58. Hassanpour H. Effect of *Aloe vera* gel coating on antioxidant capacity, antioxidant enzyme activities and decay in raspberry fruit. *LWT - Food Sci Tech* 2015; 60(1): 495–501.  
<https://doi.org/10.1016/j.lwt.2014.07.049>
59. Vieira MJ, Flores-López ML, de Rodríguez DJ, Sousa MC, Vicente AA, Martins JT. Effect of chitosan–*Aloe vera* coating on postharvest quality of blueberry (*Vaccinium corymbosum*) fruit. *Postharvest Biol Tech* 2016; 116: 88–97.  
<https://doi.org/10.1016/j.postharvbio.2016.01.011>
60. Nasrin TAA, Rahman MA, Hossain MA, Islam MN, Arfin MS. Postharvest quality response of strawberries with *Aloe vera* coating during refrigerated storage. *The J Horticult Sci Biotech* 2017; 92(6): 598–605.  
<https://doi.org/10.1080/14620316.2017.1324326>
61. Martínez-Romero D, Castillo S, Guillén F, Díaz-Mula HM, Zapata PJ, Valero D, Serrano M. *Aloe vera* gel coating maintains quality and safety of ready-to-eat pomegranate arils. *Postharvest Biol Tech* 2013; 86: 107–112.  
<https://doi.org/10.1016/j.postharvbio.2013.06.022>
62. Heggors JP, Pineless GR, Robson MC. Dermaide *Aloe/Aloe vera* Gel: Comparison of the antimicrobial effects. *The American J Med Tech* 1979; 41: 293–294.
63. Tyler VE. *Herbs of Choice*. Pharmaceutical Products Press, New York, 1994.
64. Ito S, Teradaira R, Beppu H, Obata M, Nagatsu T, Fujita K. Properties and pharmacological activity of carboxypeptidase in *Aloe arborescens* Mill. var. *Natalensis* Berger. *Phytother Res* 1993; 7(7): S26–S29.  
<http://dx.doi.org/10.1002/ptr.2650070710>
65. Haller JS. A drug for all seasons, medical and pharmacological history of *Aloe*. *Bulletin of the New York Academy of Medicine*. 1990; 66: 647–659.
66. Che QM, Akao T, Hattori M, Kobashi K, Namba T. Isolation of human intestinal bacteria capable of transforming barbaloin to *Aloe-Emodin* Anthrone. *Planta Medica* 1991;57(1): 15–19. <http://dx.doi.org/10.1055/s-2006-960007>
67. Vázquez B, Avila G, Segura D, Escalante B. Antiinflammatory activity of extracts from *Aloe vera* gel. *J Ethnopharmacol* 1996; 55(1): 69–75.  
[https://doi.org/10.1016/s0378-8741\(96\)01476-6](https://doi.org/10.1016/s0378-8741(96)01476-6)
68. Davis RH, Parker WL, Samson RT, Murdoch DP. Isolation of a stimulatory system in an *Aloe* extract. *J American Ped Medical Assoc* 1991; 81(9): 473–478.  
<https://doi.org/10.7547/87507315-81-9-473>
69. Prabhjone R, Thong-Ngam D, Wisedopas N, Chatsuwat T, Patumraj S. Anti-inflammatory effects of *Aloe vera* on leukocyte-endothelium interaction in the gastric microcirculation of *Helicobacter pylori*-infected rats. *Clin Hemorheol Microcirc* 2006; 35: 359–366.
70. Langmead L, Makins RJ, Rampton DS. Anti-inflammatory effects of *Aloe vera* gel in human colorectal mucosa *in vitro*. *Alimentary Pharmacol Amp Therap* 2004; 19(5): 521–527.  
<https://doi.org/10.1111/j.1365-2036.2004.01874.x>
71. Zawahry ME, Hegazy MR, Helal M. Use of *Aloe* in treating leg ulcers and dermatoses. *Int J of Dermatol* 1973; 12(1): 68–73.  
<http://dx.doi.org/10.1111/j.1365-4362.1973.tb00215.x>
72. Saroj PL, Dhandar DG, Singh RS. *Indian Aloe* Central Institute for Arid Horticulture, Bikaner, 2004.
73. Foster S. *Aloe vera*: The succulent with skin soothing cell protecting properties, herbs for health magazine, 1999.  
<http://www.healthy.net/library/articles/hfh/Aloe.htm>
74. El-Shemy HA, Aboul-Soud MA, Nassr-Allah AA, Aboul-Enein KM, Kabash A, Yagi A. Antitumor properties and modulation of antioxidant enzymes' activity by *Aloe vera* leaf active principles isolated via supercritical carbon dioxide extraction. *Current Med Chem* 2010; 17(2): 129–138.  
<http://dx.doi.org/10.2174/092986710790112620>
75. Davis RH, Di Donato JJ, Hartman GM, Hass RC. Anti-inflammatory and wound healing activity of a growth substance in *Aloe vera*. *J American Podiat Medical Association* 1994; 84(2): 77–81.  
<https://doi.org/10.7547/87507315-84-2-77>
76. Steenkamp V, Stewart MJ. Medicinal applications and toxicological activities of *Aloe* products. *Pharm Biol* 2007; 45(5): 411–420.  
<https://doi.org/10.1080/13880200701215307>
77. Fenig E, Nordenberg J, Beery E, Sulkes J, Wasserman L. Combined effect of *aloe-emodin* and chemotherapeutic agents on the proliferation of an adherent variant cell line of Merkel cell carcinoma. *Oncol Reports* 2004; 11(1): 213–217.  
<https://doi.org/10.3892/or.11.1.213>
78. Keivan Z, isup Moloud AZ, Kohzad S, Zahra R. Antiviral activity of *Aloe vera* against herpes simplex virus type 2: An *in vitro* study. *African J Biotech* 2007; 6(15): 1770–1773.  
<https://doi.org/10.5897/ajb2007.000-2276>
79. SWLi, Yang TC, Lai CC, et al. Antiviral activity of *Aloe-emodin* against influenza A virus via galectin-3 up-regulation. *European J Pharmacol* 2014; 738: 125–132.  
<https://doi.org/10.1016/j.ejphar.2014.05.028>
80. Kemp MC, Kahlon JB, Chinnah AD, Carpenter RH, McAnalley BH, McDaniel HR, Shannon WM. *In vitro* evaluation of the antiviral effects of acemannan on the replication and pathogenesis of HIV-1 and other enveloped viruses: Modification of the processing of glycoprotein glycoprotein precursors. *Antiviral Res* 1990;13(1): 83–96.  
[http://dx.doi.org/10.1016/0166-3542\(90\)90156-2](http://dx.doi.org/10.1016/0166-3542(90)90156-2)
81. Saoo K, Miki H, Ohmori M, Winters WD. Antiviral activity of *Aloe* extracts against cytomegalovirus. *Phytotherapy Res* 1990; 10(4): 348–350.  
[https://doi.org/10.1002/\(SICI\)1099-1573\(199606\)10:4%3C348::AID-PTR836%3E3.0.CO;2-2](https://doi.org/10.1002/(SICI)1099-1573(199606)10:4%3C348::AID-PTR836%3E3.0.CO;2-2)
82. Sydiskis RJ, Owen DG, Lohr JL, Rosler KH, Blomster RN. Inactivation of enveloped viruses by anthraquinones extracted from plants. *Antimicrobial Agents Chemoth* 1991; 35(12): 2463–2466.  
<http://dx.doi.org/10.1128/AAC.35.12.2463>
83. Alves DS, Pérez-Fons L, Estepa A, Micol V. Membrane-related effects underlying the biological activity of the anthraquinones *emodin* and *barbaloin*. *Biochem Pharmacol* 2004; 68(3): 549–561.  
<https://doi.org/10.1016/j.bcp.2004.04.012>
84. Cragg GM, Newman DJ. Natural product drug discovery in the next millennium. *Pharm Biol* 2001; 39(sup1): 8–17.  
<https://doi.org/10.1076/phbi.39.s1.8.0009>
85. Roberts DB, Travis EL. Acemannan- containing wound dressing gels reduce radiation-induced skin reactions in C3H mice. *Int J Radiation Oncol Biol Physiol* 1995; 32(4): 1047–1052. [http://dx.doi.org/10.1016/0360-3016\(94\)00467-Y](http://dx.doi.org/10.1016/0360-3016(94)00467-Y)
86. Bosley C, Smith J, Baratti P. A Phase III trial comparing an Anionic Phospholipid based (APP) cream and *Aloe vera*-based gel in the prevention and treatment of radiation dermatitis. *Int J Radiation Oncol Biol Physics* 2003; 57 (2): 34–38. [http://dx.doi.org/10.1016/S0360-3016\(03\)01404-4](http://dx.doi.org/10.1016/S0360-3016(03)01404-4)
87. Siegers CP. Anthranoid laxative abuse—A risk for colorectal cancer. *Gut* 1993; 34(8): 1099–1101.  
<http://dx.doi.org/10.1136/gut.34.8.1099>
88. SumbulS, Ahmed SW, AzharI. Anti fungal activity of *Allium*, *Aloe*, and *Solanum* species. *Pharm Biol* 2004;42(7), 491–498.  
<http://dx.doi.org/10.3109/13880200490891845>
89. Suvitayavat W, Sumrongkit C, Thirawarapan SS, Bunyapraphatsara N. Effects of *Aloe* preparation on the histamine-induced gastric secretion in rats. *J Ethnopharmacol* 2004; 90(2–3): 239–247.  
<https://doi.org/10.1016/j.jep.2003.09.044>
90. Yusuf S, Agunu A, Diana M. The effect of *Aloe vera* A. berger (Liliaceae) on gastric acid secretion and acute gastric mucosal injury in rats. *J Ethnopharmacol* 2004; 93: 33–37.  
[https://doi.org/10.1016/s0378-8741\(04\)00119-9](https://doi.org/10.1016/s0378-8741(04)00119-9)
91. Chandan BK, Saxena AK, Shukla S, Sharma N, Gupta DK, Suri KA, Suri J, Bhadauria M, Singh B. Hepatoprotective potential of *Aloe barbadensis* Mill. against carbon tetrachloride induced hepatotoxicity. *J Ethnopharmacol* 2007; 111(3): 560–566.  
<https://doi.org/10.1016/j.jep.2007.01.008>

92. Chow JTN, Williamson DA, Yates KM, Goux WJ. Chemical characterization of the immunomodulating polysaccharide of *Aloe vera* L. Carbohydrate Res 2005;340(6): 1131–1142. <https://doi.org/10.1016/j.carres.2005.02.016>
93. Im SA, Oh ST, Song S, et al. Identification of optimal molecular size of modified Aloe polysaccharides with maximum immunomodulatory activity. Int Immunopharmacol 2005; 5(2): 271–279. <https://doi.org/10.1016/j.intimp.2004.09.031>
94. Pugh N, Ross SA, ElSohly MA, Pasco DS. Characterization of Aloeride, a new high- molecular weight polysaccharide from *Aloe vera* with potent immunostimulatory activity. J Agricul Food Chem 2001; 49(2): 1030-1034. <http://dx.doi.org/10.1021/jf001036d>
95. Hart LA, Nibbering PH, van den Barselaar MT, van Dijk HAJ, Labadie RP. Effects of low molecular constituents from *Aloe vera* gel on oxidative metabolism and cytotoxic and bactericidal activities of human neutrophils. Int J Immunopharmacol 1990; (12)4: 427- 434. [http://dx.doi.org/10.1016/0192-0561\(90\)90026-J](http://dx.doi.org/10.1016/0192-0561(90)90026-J)
96. Hansel R, Keller K, Rimpler H, Schneider G. Hagers Handbuch der Pharmazeutischen Praxis. Monograph: Valeriana, 5<sup>th</sup> Edition, Springer, Berlin, 1994. <http://dx.doi.org/10.1007/978-3-642-57881-6>
97. Strickland FM. Immune regulation by polysaccharides: Implications for skin cancer. J Photochem Photobiol B: Biol 2001; 63(1–3): 132–140. [https://doi.org/10.1016/s1011-1344\(01\)00210-x](https://doi.org/10.1016/s1011-1344(01)00210-x)
98. Ishii Y, Tanizawa H, Takino Y. Studies of Aloe. V. mechanism of cathartic effect. Biol Pharm Bulletin, 1994; 17(5): 651-653. <http://dx.doi.org/10.1248/bpb.17.651>
99. DalBelo SE, Rigo Gaspar L, Berardo Gonçalves PM. Moisturizing effect of cosmetic formulations containing *Aloe vera* extract in different concentrations assessed by skin bioengineering techniques. Skin Res Tech 2006; 12(4): 241–246. <https://doi.org/10.1111/j.0909-752x.2006.00155.x>
100. Reddy UmaCH, Reddy SK Reddy J. *Aloe vera*- A wound healer. Asian J Oral Health Allied Sci 2011; 1: 91-92.
101. Maenthaisong NR, Chaiyakunapruk, Niruntraporn S. The efficacy of *Aloe vera* for burn wound healing: A systematic review. Burns 2007; 33(6): 713-718. <http://dx.doi.org/10.1016/j.burns.2006.10.384>
102. Hegggers JP. Beneficial effect of Aloe on wound healing in an excisional wound healing model. J Alt Comp Med 1996; 2(2): 271-277. <http://dx.doi.org/10.1089/acm.1996.2.271>
103. de Witte P. Metabolism and Pharmacokinetics of Anthranoids. Pharmacol 1993; 47(1): 86-97. <http://dx.doi.org/10.1159/000139847>
104. Chithra P, Sajithal GB, Chandrakasan G. Influence of *Aloe vera* on Glycosaminoglycans in the matrix of healing dermal wounds in rats. J Ethanopharmacol 1998; 59(3): 179-186. [http://dx.doi.org/10.1016/S0378-8741\(97\)00112-8](http://dx.doi.org/10.1016/S0378-8741(97)00112-8)
105. Syed TA, Afzal M, Ashfaq AS. Management of genital herpes in men with 0.5% *Aloe vera* extracts in a hydrophilic cream: A placebo- controlled double-blind study. J Dermatol Treat 1997; 8(2): 99-102. <http://dx.doi.org/10.3109/095466397091602>
106. Jia Y, Zhao G, Jia J. Preliminary evaluation: The effects of *Aloe ferox* Miller and *Aloe arborescens* Miller on wound healing. J Ethnopharmacol 2008; 120(2): 181–189. <https://doi.org/10.1016/j.jep.2008.08.008>
107. Antherton P. *Aloe vera*: Magic or medicine?. Nursing Standard 1998; 12(41): 49-54.
108. Beuers U, Spengler U, Pape GR. Hepatitis after chronic abuse of senna. Lancet 1991; 337 (8737): 472. [http://dx.doi.org/10.1016/0140-6736\(91\)91012-J](http://dx.doi.org/10.1016/0140-6736(91)91012-J)
109. Muller-Lissner SA. Adverse effects of laxatives: Facts and fiction. Pharmacol 1993; 47(1): 138-145. <http://dx.doi.org/10.1159/000139853>
110. Heizer WD, Warshaw AL, Waldmann TA, et al. Protein-losing gastroenteropathy and malabsorption associated with factitious diarrhea. Annals Int Med 1968; 68(4): 839-852. <http://dx.doi.org/10.7326/0003-4819-68-4-839>
111. McHugh TH, Senesi E. Apple Wraps: A novel method to improve the quality and extend the shelf life of fresh-cut apples. J Food Sci 2000;65(3): 480–485. <https://doi.org/10.1111/j.1365-2621.2000.tb16032.x>
112. Michailides TJ, Manganaris GA. Harvesting and handling effects on postharvest decay. Stewart Postharvest Rev 2009; 5(2): 1–7. <https://doi.org/10.2212/spr.2009.2.3>
113. Kahramanoğlu I. Introductory chapter: Postharvest physiology and technology of horticultural crops. In Postharvest Handling; Kahramanoğlu, I., Ed.; In Tech Open: London, UK, 2017; 1–5. <https://doi.org/10.5772/intechopen.69466>
114. HeQ, Changhong L, Kojo E, Tian Z. Quality and safety assurance in the processing of *Aloe vera* gel juice. Food Cont 2005; 16(2): 95–104. <https://doi.org/10.1016/j.foodcont.2003.12.001>
115. Vinson JA, Al Kharrat H, Andreoli L. Effect of *Aloe vera* preparations on the human bioavailability of vitamins C and E. Phytomed 2005; 12(10): 760–765. <https://doi.org/10.1016/j.phymed.2003.12.013>
116. Jani GK, Shah DP, Jain VC, Patel MJ, Vithalan DA. Evaluating mucilage from *Aloe Barbadosis* Miller as a pharmaceutical excipient for sustained-release matrix tablets. Pharm Technol 2007; 31: 90-98.
117. Peter A. *Aloe vera* Myth or Medicine? Positive Health Publications, 2002.
118. Bisset NG. *Sennae folium*. Max Wichtl's Herbal Drugs & Phytopharmaceuticals. CRC Press, Boca Raton 1994.