


Review

Chemical Constituents and Pharmacological Activities of Garlic (*Allium sativum* L.): A Review

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Abstract: Medicinal plants have been used from ancient times for human healthcare as in the form of traditional medicines, spices, and other food components. Garlic (*Allium sativum* L.) is an aromatic herbaceous plant that is consumed worldwide as food and traditional remedy for various diseases. It has been reported to possess several biological properties including anticarcinogenic, antioxidant, antidiabetic, renoprotective, anti-atherosclerotic, antibacterial, antifungal, and antihypertensive activities in traditional medicines. *A. sativum* is rich in several sulfur-containing phytoconstituents such as alliin, allicin, ajoenes, vinylthiins, and flavonoids such as quercetin. Extracts and isolated compounds of *A. sativum* have been evaluated for various biological activities including antibacterial, antiviral, antifungal, antiprotozoal, antioxidant, anti-inflammatory, and anticancer activities among others. This review examines the phytochemical composition, pharmacokinetics, and pharmacological activities of *A. sativum* extracts as well as its main active constituent, allicin.

Keywords: Garlic; *Allium sativum*; pharmacological activities; phytoconstituents; allicin; pharmacokinetics

1. Introduction

Medicinal plants have been a good source of new pharmacologically active molecules. For example, natural products could be a potential alternative for controlling the pathogen associated with diseases [1–5]. Recently, antibiotics and most drugs on the market have shown unwanted symptoms and the

emergence of resistant pathogenic microorganisms, toxic effects related to these drugs, and withdrawal issues restricting their use in many countries [6], therefore, much attention has been paid to the herbal extracts and pharmacologically active molecules extracted from different plant species that are used previously in the traditional medicine [7–11]. Many plant species have been reported to exert pharmacological properties due to their phytoconstituents such as glycosides, alkaloids, saponins, steroids, flavonoids, tannins, and terpenoids (e.g., monoterpenes, diterpenes, and sesquiterpenes). Nowadays, eighty percent of the world's populations depend on traditional medicines as an essential source of their primary health care [12,13]. Medicinal plant extracts and their constituents also possess various biological activities including virucidal, bactericidal, fungicidal, anti-inflammatory, analgesic, sedative, spasmolytic, and local anesthetic activities among others [14,15].

Garlic (*Allium sativum* L.; Family: Amaryllidaceae) is an aromatic herbaceous annual spice and one of the oldest authenticated and most important herbs that have been used from ancient times as traditional medicine [16,17]. It is considered the second broadly used *Allium* species with onion (*Allium cepa* L.), which is used as a remedy against several common diseases such as cold, influenza, snake bites, and hypertension [18]. *Allium* species and their active components are reported to reduce the risk of diabetes and cardiovascular diseases, protect against infections by activating the immune system and have antimicrobial, antifungal, anti-aging as well as anti-cancer properties which confirmed by epidemiological data from human clinical studies [19]. Garlic has been used for cooking purposes as a spice that can flavor foods during the cooking process. As well, it possesses therapeutic purposes including the treatment of lung disorders, whooping cough, stomach disorders, cold, earache, and assists in preventing cardiovascular disease [17]. While aged garlic extract (AGE), prepared from aged garlic is a folk herbal remedy that has been shown to enhance the immune system and thus inhibit cancer and heart disorders. Raw garlic and its transformed products have been reported to contain various sulfur compounds that have been included in several types of preparations [20]. Moreover, quercetin, the major flavonoid isolated from garlic, was found to interact with some medications such as vitamin E and C [21] and modify the in vitro as well as the in vivo transferases and cytochrome P450 isozymes activity, however, the in vivo studies revealed that garlic oil and its three allyl sulfide components increase the CYP3A1, 2B1, and 1A1 expression in the hepatic detoxification system [22].

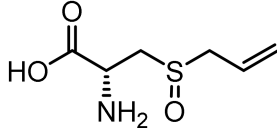
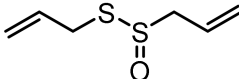
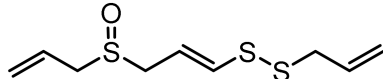
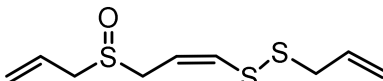
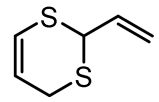
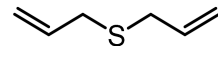
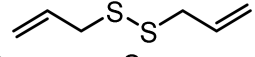
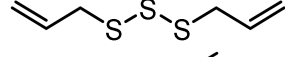
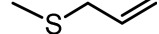
Alliin [S-(2-propenyl)-2-propene-1-sulfinothioate], the most biologically active sulfur-containing compound of garlic, is responsible for its smell and taste [23,24]. Alliin (S-allyl-L-cysteine sulfoxide) is the main precursor of alliin, which represents about 70% of total thiosulfinates existing in the crushed cloves [25]. Allyl mercaptan is the odorant molecule responsible for the garlic breath and results from the interaction of alliin or diallyl disulfide with cysteine in the presence of S-allyl-mercapto cysteine [25,26]. Alliin is a lipid-soluble sulfur compound, which can be easily damaged by cooking and has the ability to provoke intolerance, allergic reactions, and gastrointestinal disorders [27–29]. Various pharmacological activities have also been reported for the extracts and isolated compounds from garlic. The main aim of this article is to critically review the available scientific information about the traditional uses, chemical constituents, pharmacokinetics, and pharmacological activities of garlic.

2. Chemical Constituents of Garlic

Bulbs of *A. sativum* are reported to contain hundreds of phytochemicals including sulfur-containing compounds (Table 1) such as ajoenes (*E*-ajoene, *Z*-ajoene), thiosulfinates (alliin), vinyl dithiins (2-vinyl-(4H)-1,3-dithiin, 3-vinyl-(4H)-1,2-dithiin), sulfides (diallyl disulfide (DADS), diallyl trisulfide (DATS)) and others that accounted 82% of the overall garlic sulfur content [30]. Alliin, the main cysteine sulfoxide is transformed to alliin by allinase enzyme after cutting off the garlic and breaking down the parenchyma [31]. S-propyl-cysteine-sulfoxide (PCSO), alliin and S-methyl cysteine-sulfoxide (MCSO) are the main odoriferous molecules of freshly milled garlic homogenates [31]. PCSO can produce more than fifty metabolites depend on water content and temperature as well as allinase enzyme that can act on the mixture of MCSO, PCSO, and alliin to produce other molecules, such as allyl methane

thiosulfonates, methyl methanethiosulfonate, and further corresponding thiosulfonates (R-S-S-R'), by which R and R' are allyl, propyl, and methyls groups [31].

Table 1. List and structures of some of the sulfur-containing compounds isolated from *Allium sativum*.

Compounds	Molecular formula	Structure
Alliin	C ₆ H ₁₁ NO ₃ S	
Allicin	C ₆ H ₁₀ OS ₂	
E-Ajoene	C ₉ H ₁₄ OS ₃	
Z-Ajoene	C ₉ H ₁₄ OS ₃	
2-Vinyl-4H-1,3-dithiin	C ₆ H ₈ S ₂	
Diallyl sulfide (DAS)	C ₆ H ₁₀ S	
Diallyl disulfide (DADS)	C ₆ H ₁₀ S ₂	
Diallyl trisulfide (DATS)	C ₆ H ₁₀ S ₃	
Allyl methyl sulfide (AMS)	C ₄ H ₈ S	

S-alk(en)yl-L-cysteine sulfoxides are the secondary metabolites obtained from cysteine which accumulate in the plants of *Allium* genus [32]. Garlic formulations consist of several organosulfur compounds, N-acetylcysteine (NAC), S-allyl-cysteine (SAC) [33], and S-allyl-mercapto cysteine (SAMC), which are derived from alliin [34]. Notably, SAC has antioxidant, anti-inflammation, regulated redox, pro energetic, antiapoptotic, and signaling capacities [32,35], while SAMC shows an anticancer activity through preventing the cancer cells multiplication [36].

Allicin (allyl thiosulfonate), is a sulfenic acid thioester and its pharmacological effect is attributed to its antioxidant activity as well as its interaction with thiol-containing proteins [37]. In the allicin biosynthesis, cysteine is transformed to alliin that is hydrolyzed by the allinase enzyme [38]. This enzyme composed of pyridoxal phosphate (PLP) which splits alliin and produces ammonium, pyruvate, and allyl sulfenic acid that are highly reactive and unstable at room temperature, where two molecules were combined to form allicin [37,39].

3. Pharmacokinetics and Stability of Garlic Components

De Rooij et al. [40] reported the existence of N-acetyl-S-allyl cysteine (NASAC) in human urine after garlic ingestion that is created by converting SAC into the N-acetylated metabolite by N-acetyl transferase enzyme. Previous reports revealed the existence of diallyl disulfide (DADS), allyl methyl sulfide (AMS), diallyl sulfide (DAS), allyl methyl disulfide (AMDS), dimethyl sulfide, acetone and diallyl trisulfide (DATS) in tested participants breath after administration of 38 g of raw garlic. It was reported that DADS, DAS, DATS, and AMDS achieved the maximum concentrations within 2 to 3 h. Freeman and Kodera [41] examined allicin stability in simulated gastric fluid (SGF), blood, stimulated intestinal fluid (SIF) and various solvents such as; methanol, water, ethyl acetate at pH 1.2 and 7.5 and they observed that allicin affected the SGF and SIF at pH 1.2 and 7.5, respectively. These

results suggest that allicin degraded at room temperature and was more stable in methanol than in ethyl acetate. Furthermore, about 90% of the allicin stayed in the SIF (pH 7.5) and SGF (pH 1.2) after incubation at 37 °C for 5 h, while only a small amount could be detected after 5 min in the blood. About 62% and 80% of allicin remained one day after *Allium* administration without increasing the concentration of allicin decomposition products such as DADS [40]. The pharmacokinetic examination in rats using 35 S-labeled alliin, vinyl dithiols, and allicin, revealed that the alliin peak time (T_{max}) was lower than 10 min and was eliminated after 6 h from the blood, whereas the allicin peak time (T_{max}) was 30–60 min and the mean total fecal and urinary excretion was 85.5% after 72 h.

Allicin, a bright yellow oily liquid that possesses a distinctive garlic odor and it is very unstable, therefore it can be easily decomposed even at room temperature [42]. Previous studies reported that allicin can easily degrade under the influence of temperature to form ajoenes ((*E*)- and (*Z*)-4, 5, 9-trithiadodeca-1, 6, 11-triene-9-oxides) and vinyl dithiols which are more stable than allicin [43]. These degradation products are commonly isolated from oil, aqueous and chloroform garlic extracts and are present as (*E*) and (*Z*) isomers, where (*E*)-ajoene is usually found in double amounts [44].

4. Pharmacological Activities of Garlic and Its Related Compounds

4.1. Traditional Uses of Garlic

Garlic is one of the most important bulb vegetables that has a pungent flavor and widely used all over the world as a spice and flavoring agent. The organosulfur compounds like allicin and DADS are the main compounds responsible for its pungency effects and spicy aroma. Garlic is well-known to be used in food preparation, especially dried foods for storage and some types of soup and it can be utilized in both fresh and dehydrated states [45]. Traditionally, garlic and its related compounds have been stated to have several biological activities including anticarcinogenic, antioxidant [46], antidiabetic, renoprotective, anti-atherosclerotic, antibacterial, antifungal [47], and antihypertensive activities [17]. Moreover, garlic has been used in traditional medicine to treat indigestion, respiratory and urinary tract infections and cardiac disorders and it showed carminative, antipyretic, sedative, aphrodisiac, and diuretic effects [32].

4.2. Activities Related to Infectious Diseases

4.2.1. Antibacterial Activity

The antimicrobial activity of garlic is attributed to allicin activity that was reported toward a wide variety of microorganisms including antibiotic-resistant, Gram-positive and Gram-negative bacteria such as *Shigella*, *Escherichia coli* [48], *Staphylococcus aureus*, *Pseudomonas aeruginosa* [44], *Streptococcus mutans*, *S. faecalis*, *S. pyogenes*, *Salmonella enterica*, *Klebsiella aerogenes* [49], *Vibrio*, *Mycobacteria*, *Proteus vulgaris*, and *Enterococcus faecalis* [50]. Various garlic extracts (aqueous, chloroform, methanolic, and ethanolic extracts) were reported to inhibit the growth of several pathogenic bacteria with varying degrees of susceptibility. For instance, a study revealed that ethanolic garlic extract showed higher inhibitory effect against *E. coli* and *Sal. typhi* than the aqueous extract that showed little or no inhibition effect [51]. Meriga et al. [52] reported that aqueous garlic extract showed antibacterial activity toward Gram-negative (*Kl. pneumoniae* and *E. coli*) as well as Gram-positive (e.g., *Bacillus subtilis* and *S. aureus*) strains, whereas methanolic garlic extract showed antimicrobial activity against all tested strains except *S. aureus*. However, hexane, ethyl acetate, and chloroform extracts did not show any antibacterial effect. Moreover, garlic extracts prevented the growth of enterotoxigenic *E. coli* strains and other pathogenic intestinal bacteria, which are the main cause of diarrhea in humans and animals. Besides the antibacterial activity of garlic, it was reported to prevent the toxins produced by bacterial infection [53]. Allicin also showed effectiveness toward methicillin-resistant *S. aureus* (MRSA) [50]. Allicin's antimicrobial activity is due to its chemical interaction with enzymes containing thiol e.g., thioredoxin reductase, RNA polymerase, and alcohol dehydrogenase [54] by oxidizing protein cysteine

or glutathione residues under physiological conditions. Allicin is a dose-related biocide that can influence essential metabolism of cysteine proteinase, and thus, kill all eukaryotic cells due to the presence of thiol groups in all living cells.

4.2.2. Antifungal Activity

Garlic extracts showed a broad spectrum fungicidal effect against a wide range of fungi including *Candida*, *Torulopsis*, *Trichophyton*, *Cryptococcus*, *Aspergillus*, *Trichosporon*, and *Rhodotorula* species. Recently, garlic extract was found to inhibit the *Meyerozyma guilliermondii* and *Rhodotorula mucilaginosa* germination and growth [55]. Another study reported the antifungal activity of various *A. sativum* extracts namely aqueous, ethanolic, methanolic, and petroleum ether against human pathogenic fungi such are *Trichophyton verrucosum*, *T. mentagrophytes*, *T. rubrum*, *Botrytis cinerea*, *Candida* species, *Epidermophyton floccosum*, *Aspergillus niger*, *A. flavus*, *Rhizopus stolonifera*, *Microsporum gypseum*, *M. audouinii*, *Alternaria alternate*, *Neofabraea alba*, and *Penicillium expansum* [56]. The garlic extract acted by affecting the fungal cell wall and causing irreversible ultrastructural changes in the fungal cells, which lead to loss of structural integrity and affected the germination ability. These changes in the cytoplasmic content lead to nucleus and cell organelles damage that ultimately leads to cell death. Moreover, allicin and garlic oil showed potent antifungal effects against *Candida albicans*, *Ascosphaera apisin*, and *A. niger* [44] and they acted by penetrating the cellular membrane as well as organelles membranes like the mitochondria and leading to organelles destruction and cell death [57]. DADS and DATS separated from garlic essential oil showed antifungal activity against a number of fungi (*C. albicans*, *C. tropicalis*, and *Blastoschizomyces capitatus*). In addition to that, saponins extracted from *A. sativum* exhibited antifungal activity against *Botrytis cinerea* and *Trichoderma harzianum* [58].

4.2.3. Anti-Protozoal Activity

Various studies reported the anti-protozoal activity of garlic extracts and its phytochemicals against several protozoan parasites. For instance, an in vitro study revealed that the aqueous, ethanolic, and dichloromethane *A. sativum* extracts exhibited anthelmintic activity against *Haemonchus contortus* and the ethanolic extract was the most effective one, while aqueous garlic extract showed potent activity against *Trichuris muris* and *Angiostrongylus cantonensis* [58]. Garlic was also examined in vivo and in vitro against *Taenia taeniaeformis*, *Hymenolepis microstoma*, *H. diminuta*, *Echinostoma caproni*, and *Fasciola hepatica* [59]. Abdel-Hafeez et al. [60] showed that garlic extract inhibited the growth of *Blastocystis* spp. in vivo and this activity attributed to that garlic extracts contains several phytochemicals e.g., thiosulfinates are one of the bioactive compounds that possess antibacterial activity that is related to thiol enzymes inhibition which presents in several microorganisms. Allicin also acts by preventing the parasite's RNA as well as DNA and protein synthesis. Moreover, allicin and DATS, phytochemicals isolated from garlic extract, showed antiparasitic activity against *Entamoeba histolytica*, *Plasmodium falciparum*, *Babesia*, *Theileria*, *Trypanosoma brucei*, and *Giardia lamblia* [58]. Ajoene also exhibited antiparasitic activity by inhibiting the human glutathione reductase and *T. cruzi* trypanothione reductase [61]. Hazaa et al. [62] reported the activity of garlic oil toward broad-spectrum microorganisms such are *Cochlospermum planchonii*, *Plasmodium*, *Giardia*, *Leishmania*, and *Trypanosoma*.

4.2.4. Antiviral Activity

The antiviral activity of garlic extracts has been evaluated against influenza B, human rhinovirus type 2, human cytomegalovirus (HCMV), Parainfluenza virus type 3, herpes simplex type 1 and 2, vaccinia virus, and vesicular stomatitis virus [63]. Interestingly, in vivo experiment exhibited the antiviral activity of garlic extract and they reported that garlic showed protective activity against influenza viruses by improving the production of neutralizing antibodies when given to mice and this activity was based on the presence of several phytochemicals namely, ajoene, allicin, allyl methyl thiosulfinate, and methyl allyl thiosulfinate [64]. Allicin acts by preventing several thiol enzymes, while ajoene's antiviral activity was due to the prevention of adhesive interaction and fusion of leukocytes.

Moreover, DATS was effective against the HCMV replication and viral immediate-early gene expression and it acts by enhancing natural killer-cell (NK-cell) activity that destroys virus-infected cells [58].

4.3. Antioxidant and Anti-inflammatory Activities

4.3.1. Antioxidant Activity

Asdaq and Inamdar [33] reported that the frequent garlic intake promotes internal antioxidant activities and reduces oxidative adverse effects either by increasing the endogenous antioxidant synthesis or reducing the production of oxidizers such as oxygen-free radical species (ORS). Gentamycin is an antibiotic that has been used to treat several types of bacterial infections and was reported to promote hepatic damage through raising aspartate transaminase and alanine aminotransferase enzymes in addition to lowering the plasma albumin level. It is demonstrated that garlic protects against gentamycin- as well as acetaminophen-induced hepatotoxicity by improving antioxidant status, and regulating oxidative stress [50]. As ROS seems to be at the core of many ailments, it is justified to assume that the antioxidant effect of garlic might be through modulation of ROS, increasing glutathione and cellular antioxidant enzymes [53]. Moreover, garlic extract was found to increase the activities of some antioxidant enzymes (e.g., superoxide dismutase (SOD)) and decrease glutathione peroxidase (GSH-Px) in hepatic tissues of rats. Notably, several reports indicated that AGE rich in flavonoid, phenol, and different sulfur compounds e.g., SAC shows high radical scavenging activity [65]. Additionally, AGE acted by stimulating the expression of different antioxidant enzymes, namely glutamate-cysteine ligase modifier (GCLM) and heme oxygenase-1 (HO-1) subunit by the nuclear factor erythroid-2 related factor 2 (Nrf2)-antioxidant response element (ARE) pathway that is responsible for human endothelial cells protection against oxidative stress [66]. Alliin, the major compound isolated from AGE, showing wide-spectrum antioxidant activities by controlling ROS generation and preventing mitogen-activated protein kinase (MAPK). Moreover, it was reported to prevent ROS production by inhibiting NADPH oxidase 1, and thus, inhibiting the osteoclast fusion caused by receptor activator of nuclear factor-kappa B ligand (RANKL) [67]. Allicin, DADS, and DATS are the main antioxidative compounds that showed an antioxidant effect in lower doses at the physiological level [54]. Saponins extracted from garlic were reported to scavenge intracellular ROS and protect mouse-derived C2C12 myoblasts towards growth inhibition and H₂O₂-induced DNA damage [68]. Interestingly, Abdel-Daim et al. [69] reported that DAS exhibited potent antioxidant and cytoprotective activities and these activities may be due to suppressing the enzymatic activity of cytochrome P450-2E1 and thereby reducing the generation of reactive oxygen and nitrogen species or by inducing the mRNA expression of Nrf2 and heme-oxygenase 1 enzyme.

4.3.2. Anti-Inflammatory Activity

Garlic extracts and its related phytochemicals have been reported to possess anti-inflammatory activity. A study reported that the garlic extracts remarkably impaired the liver inflammation and damage caused by *Eimeria papillate* infections [70]. Hobauer et al. [71], as well as Gu et al. [72], observed that the anti-inflammatory activity of garlic is caused by inhibiting the emigration of neutrophilic granulocytes into epithelia. Aged black garlic (ABG) exhibited potent antioxidant activities and these activities may be responsible for its anti-inflammatory activity. The ABG chloroform extract acts by reducing NF- κ B activation in human umbilical vein endothelial cells caused by tumor necrosis factor- α (TNF- α). Moreover, ABG methanolic extract was reported to prevent the cyclooxygenase-2 (COX-2) and prostaglandin E₂ (PGE₂) production by NF- κ B inactivation [73]. You et al. [74] investigated the anti-inflammatory effect of ABG and they reported that this activity may be attributed to the direct suppression of toll-like receptor 4 (TLR4) signaling cascade activation in macrophages, reducing nuclear NF- κ B level and improving the NF- κ B and I κ B cytosolic levels in LPS-activated RAW264.7 cells. Additionally, they revealed that ABG extract may act by another mechanism of action by inhibiting the iNOS and COX-2 expression, and thus, prevented the NO, interleukin-6 (IL-6) and

TNF- α formation of in LPS-activated RAW264.7 cells and TPA-mediated dermatitis in mice. Allicin demonstrated a defensive mechanism against pathogens by its ability to enhance the activity of immune cells and influence signaling pathways associated with these immune cells. Moreover, allicin works on T-cell lymphocytes by inhibiting the SDF1 α chemokine which is associated with the weakness of the dynamic structure of the actin cytoskeleton [75], in addition to this, it leads to inhibit the Transendothelial migration of neutrophils. Notably, Abdel-Daim et al. [76] reported that the anti-inflammatory activity of DAS induced by diminishing the expression of the inflammatory cytokines (e.g., NF- κ B, IL-1 β , and TNF- α), and the ROS generation by suppressing CYP-2E1 hepatic enzyme. Another report indicated that thiacremonone (a sulfur compound isolated from garlic) prevents neuroinflammation and amyloidogenesis by blocking the NF- κ B activity, and therefore can be used to treat neurodegenerative disorders (e.g., Alzheimer's disease) related to inflammation [77].

4.4. Anticancer Activity

Raw garlic extract was found to be the most effective and highly specific anticancer drug when compared with 33 raw vegetable extracts against different cancer cells without affecting the non-cancerous cells [78]. Shang et al. [68] reported that the anticancer mechanisms of garlic extracts were attributed to the inhibition of cell growth and proliferation, regulation of carcinogen metabolism, stimulation of apoptosis, prevention of angiogenesis, invasion, and migration and thus reducing the anticancer agent's negative effects. Interestingly, in 1960, tumor cells were reported to be killed when incubated in an allicin solution [63]. Allicin isolated from garlic was reported to suppress colorectal cancer metastasis through enhancing the immune function and preventing the formation of tumor vessels as well as survivin gene expression to enhance the cancer cell's apoptosis. It also can enhance the treatment of pancreatic cancer thereby invert gene silencing and restrain cancer cell proliferation [79]. Furthermore, Zhang et al. [80] revealed that allicin can prevent gastrointestinal cancer cells MGC 803 proliferation and induce apoptosis, which can be accomplished through enhancing p38 expression and cleaved caspase 3. Allicin-derived polysulfanes have been reported to target microtubules, which lead to interruption of the cell-cycle and finally to apoptosis. Several studies reported the activity of allicin in preventing cell proliferation [81] by targeting tubulin that shapes the mitotic spindle and thus inhibits cell division [82]. Iciek et al. [83] have reported the anti-tumor properties of organo-sulfur compounds (OSC) including allicin, DADS, alliin, DAS, allyl mercaptan (AM), and S-allyl cysteine (SAC), isolated from garlic. In addition, garlic powders inhibited the DNA damage caused by N-nitrosodimethylamine in the liver when administered to rats by 35% and this effect was due to the high concentration of alliin up to 60% in the samples [84]. Notably, Fleischauer and Arab [85] reported that continuous garlic intake could decrease different kinds of cancer propagation such as lung, colon, stomach, breast, and prostate. Piscitelli et al. [86] reported that garlic reduced the plasma concentrations of saquinavir by about 50% in healthy participants, after 3-week of a garlic supplement uptake, in addition to this, many researchers evaluated the antitumor and cytotoxic actions of garlic and its related constituents in vitro and in vivo. Moreover, Z-ajoene has shown anti-proliferative activities against different types of cancers and it inhibits the growth of human breast cancer cells and glioblastoma multiforme cancer stem cells (GBM CSC) [68]. It was found to stimulate apoptosis in human leukemic cells by promoting the peroxide production, caspase-3-like and caspase-8 activities [87].

4.5. Anti-Alzheimer's Disease Activity

Alzheimer's disease (AD) is the main cause of dementia in the elderly with neurodegenerative and cerebrovascular disorders [88]. Acetylcholinesterase (AChE) is the main enzyme that converts the acetylcholine (ACh) in the nervous system to acetate and choline [89]. ACh depletion in the central nervous system has been involved in the pathophysiology noticed in AD [90], therefore, donepezil (AChE inhibitor) was effective in the management/prevention of AD. Surprisingly, oil from garlic bulbs suppressed AChE activity of cerebral cortex synaptosome and exhibits antioxidant properties, thus, inhibiting AChE activity in vitro [91] as well as their ability to scavenge diphenyl-1-picrylhydrazyl

(DPPH) free radical that are used to evaluate the compound's ability to act as hydrogen donors or free radical scavengers and to assess the antioxidant activity of food [92] and reduce Fe^{3+} to Fe^{2+} could be suggested as the possible mechanism of action for their neuroprotective potential [86].

Noteworthy, the inclusion of garlic in cholesterol-fed rats' diet remarkably reduced the total glycosaminoglycans (GAGs) concentration in heart and aorta. This may be due to the enhanced GAGs degrading enzyme activity such as hyaluronidase, β -N-acetylhexosaminidase arylsulfatase and β -glucuronidase [20]. Sulfated GAGs are involved in lipid aggregation in the lesion development due to their ability to bind to plasma lipoproteins, mainly LDL. Moreover, sulfated GAGs stimulated the neurotoxic activities of various amyloidogenic peptides such as A in AD [20]. Borek [93] evaluated the neuroprotective effect of AGE using an animal model and they showed that AGE protected the brain from neurodegenerative diseases by preventing brain injury following ischemia, saving neurons toward apoptosis, and inhibiting oxidative death caused by β -amyloid [94]. Moreover, Mbyirukira and Gwebu [95] reported that AGE or SAC inhibits the brain's frontal lobe degeneration, promotes memory and learning retention, and prolongs the lifespan.

Based on the amyloid hypothesis, aggregated β -amyloid ($\text{A}\beta$) accumulation in the brain is believed to be the pathological factors that drive the onset of AD. It has been suggested that the formation of the neurofibrillary tangles contain τ -protein and synaptic degradation caused by the imbalance consequences between $\text{A}\beta$ clearance and $\text{A}\beta$ production. Haider et al. [96] reported that the prolonged garlic uptake is related to promoting the memory function by affecting the levels of the neurotransmitter, serotonin. The in vivo consumption of *A. sativum* extracts have shown that it improves memory by eliminating free radicals that cause oxidative damage and inhibit AChE enzyme [97]. It was noted that allicin inhibits AChE and butyrylcholinesterase (BuChE) enzymes (enzymes that break down neurotransmitter choline) which successively increased ACh concentration in the brain. Thus, delayed cognitive decline and dementia [98].

Garlic is also investigated to have immunomodulatory, anti-inflammatory, and antioxidant effects and this focused on the question of whether the known effect of processed garlic and its related compounds mainly allicin in inhibiting AChE and BuChE enzymes [29]. Combination therapy of allicin with cholinesterase inhibitors (ChEIs) including; rivastigmine, galantamine, and donepezil are now the most commonly used for the treatment of AD [99] as they have the ability to correct the cholinergic deficiency seen with AD. Antioxidants such as tocopherol, selegiline, and ascorbic acid (vitamin C) were examined as a possible preventive therapy for AD, and they show delayed functional deterioration in AD patients [100]. Anti-inflammatory drugs such as NSAIDs have been used as a potential treatment in AD because of their capacity to bind to and stimulate the nuclear receptor peroxisome proliferator-activated receptor (PPAR)- γ as well as their direct effects on the amyloid formation [101].

It should be noted that AChE inhibitors could be part of any combination therapy against AD [100]. For instance, Millard et al. [102] reported that AChE incubated with allicin produced rapid inactivation that was concentration and time-dependent. Many results showed concentration-dependent inhibition of bovine AChE by allicin complementing the previous finding. However, different cholinesterase inhibitors such as donepezil, rivastigmine, and tacrine are used to treat AD, and their side effects are becoming increasingly remarkable [103,104]. Therefore, the search for new derivatives extracted from the natural product with a dual function and lower side effects could be useful for patients with AD.

Allicin is a small lipophilic molecule that can suppress BuChE and AChE, and therefore, enhances ACh concentration, which is decreased remarkably in AD patient's brains [103]. Recently, allicin has been shown to have a protective effect on ischemic or traumatic neuronal damage controlled by apoptosis and oxidative stress pathways [105].

4.6. Activities Related to Metabolic Diseases

4.6.1. Effect on Dyslipidemia

Dyslipidemia is known to be the main cause of myocardial infarction and cardiovascular diseases and it is defined by high levels of triglyceride (TG), LDL, total cholesterol (TC), and low HDL level [106]. Interestingly, various evidence encourages the significant and crucial role of garlic preparations and its phytochemicals in treating hypercholesterolemia by preventing the cholesterol biosynthesis in the liver as well as inhibiting low-density lipoproteins (LDL and HDL) oxidation. Moreover, garlic reduces the cholesterol level either by stimulating the acidic and neutral steroids excretion or by reducing the cholesterogenic and lipogenic effects of fatty acid synthase, 3-hydroxy-3-methyl-glutaryl-CoA reductase, malic, and glucose-6 phosphate dehydrogenase in hepatocytes [107]. Garlic was found to have an important effect on dyslipidemia by significantly decreased serum TC, TG, and LDL levels and moderately elevated HDL cholesterol [108]. Various experimental and clinical trials were performed in animals and humans using various garlic preparations and they exhibited disputable results. They claimed that these variable results were attributed to the differences in garlic preparation composition, amount of active sulfur compounds exist in each preparation and the mechanism by which they act. For instance, Iweala et al. [108] reported that ethanolic garlic extract uptake to albino rabbits resulting in decreased their cholesterol level and body weight. Campbell et al. [109] reported that AGE significantly prevented the development of thickened, lipid-filled lesions in the preformed neointima generated from balloon-catheter harm of the right carotid artery in rabbits fed with cholesterol. In clinical trials in patients, Sobenin et al. [110,111] revealed that garlic administration at a dose of 300 and 60 mg/day for 12 months and 12 weeks, respectively decreased TC, TG, and LDL while increased HDL. Moreover, Ashraf et al. [112] garlic tablets administration at a dose of 600 mg/day for 12 weeks in diabetic patients with dyslipidemia results in high HDL and low LDL and TC levels.

4.6.2. Effect on Diabetes Mellitus

Ethanolic garlic extracts exhibited an antidiabetic effect against streptozotocin- and alloxan-induced diabetic mice and rabbits by activating the insulin secretion from parietal cells of the pancreas [113]. Another clinical study examined the antidiabetic effect of garlic pills administration at 900 mg/day in patients with type II diabetes and hyperlipidemia and they reported that garlic pills decrease the cholesterol, serum lipids, and fasting blood sugar [114]. Moreover, allyl propyl disulfide, allicin, cysteine sulfoxide, and S-allyl cysteine sulfoxide decreased the blood glucose level by preventing the insulin activation caused by liver, enhancing the secretion of insulin from pancreatic beta cells, isolation of insulin from the bonded forms, and increasing the cell sensitivity to insulin [114]. Zhai et al. [115] reported that the activity of alliin in reducing diabetes mellitus in rats was similar to that demonstrated by glibenclamide and insulin. Garlic oil also was reported to decrease the serum amylase, serum aspartate and alanine transferases, and serum alkaline and acidic phosphatase in diabetic rats.

4.6.3. Effect on Obesity

Obesity is the most common health problems that may lead to many ailments like hypertension, dyslipidemia, cardiovascular disorders, and metabolic syndrome. Garlic extracts have been reported for their activity in reducing body weight, adipose tissue mass and improved plasma lipid profiles in mice with high-fat diet-induced obesity and these effects mediated by the downregulation of multiple genes expression that is included in adipogenesis along with upregulation of the mitochondrial inner membrane proteins expression [116]. Moreover, Lee et al. [116] revealed that the antiobesity effect of garlic extracts attributed to stimulation of AMP-activated protein kinase (AMPK) as well as increased thermogenesis and decreased multiple genes expression that is included in adipogenesis. Ajoene isolated from garlic extracts was found to stimulate apoptosis, decrease the fat accumulation in 3T3-L1 adipocytes and dramatically decrease the body weight gain in mice without affecting the amount of food intake [117]. 1,2-vinyldithiin also has been reported to prevent the human preadipocytes

differentiation and decrease lipid accumulation by decreasing the C/EBP α , PPAR γ 2, and LPL expression and the PPAR γ effect in human adipocytes [118].

4.6.4. Antihypertensive Activity

Varshney and Budoff [119] reported the essential function of garlic in the control of cardiovascular risk factors as it is known to significantly decrease systolic as well as diastolic blood pressure. Garlic formulations have been broadly used to inhibit and relieve cardiovascular disorders such as hypertension, arrhythmia, thrombosis, hyperlipidemia, and atherosclerosis [19,120]. Several experimental and human studies reported the antihypertensive effect of garlic extracts and its derived bioactive molecules. For example, Sobenin et al. [121] showed the plasma fibrinolytic activity of garlic extracts and they found that it increased fibrinolytic activity in both healthy and acute myocardial infarction participants. Moreover, in vivo experiment exhibited the antihypertensive effect of aqueous garlic extract in '2 kidney 1-clip' model of hypertension in rat by reducing thromboxane B2 and prostaglandin E2 level and thereby reduced hypertension in tested rats [122]. Garlic administration at a dose of 100 mg/kg for 5 days resulted in complete prevention of acute hypoxic pulmonary vasoconstriction caused by endothelin-1 in isolated rat pulmonary arteries and they found that garlic acts by reducing endothelin 1 and angiotensin II production [120]. The mechanism of antihypertensive effect of garlic extracts is that garlic contains many active sulfur molecules that have been shown to stimulate endothelium-constricting and -relaxing factors leading to lower blood pressure. Garlic has also been shown to stimulate the production of both nitric oxide (NO) and hydrogen sulphide (H₂S) that finally leads to vasodilation. Therefore, garlic is used as a medicinal plant for controlling blood pressure worldwide [123]. Furthermore, garlic exhibited a significant role in inhibiting thrombosis as well as platelet adhesion or aggregation in humans. The AGE was reported to prevent both ADP-activated platelets binding to immobilized fibrinogen and platelet aggregation by inhibiting GPIIb/IIIa receptor and increasing cAMP [124]. Furthermore, garlic has been reported to reduce the risk of plasma viscosity, unstable angina, and peripheral arterial occlusive disorders and increase the elasticity of the blood vessels and perfusion of capillaries [87]. The gamma-glutamylcysteine isolated from garlic was reported to decrease the blood pressure by inhibiting the angiotensin-converting enzyme (ACE). Dubey et al. [122] revealed that allicin shows remarkable activity in reversing systolic blood pressure caused by dexamethasone and enhances body weight and food intake in hypertension caused by dexamethasone in rats.

Few pharmacological effects of garlic and its related bioactive compounds are shown in Table 2. Some of the mechanisms of action related to these activities are shown in Figure 1.

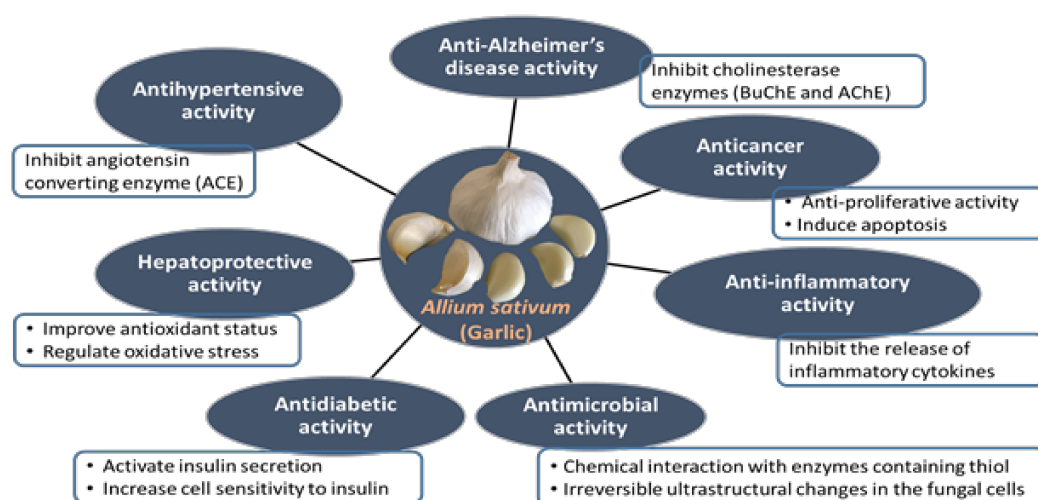


Figure 1. Schematic representation of different pharmacological activities of garlic (*Allium sativum*) and their mechanisms.

Table 2. The pharmacological activity of garlic (*Allium sativum*) and its related compounds.

Activities	Bioactive Compound	Mechanism of Action	References
Antibacterial	Allicin	Chemical interaction with enzymes containing thiol	[54]
Antifungal	DADS	Irreversible ultrastructural changes in the fungal cells, loss of structural integrity and affected the germination ability	[44]
	DATS		
Antiviral	Allicin	Chemical interaction with enzymes containing thiol	[58]
	DATS	Enhancing Natural killer-cell (NK-cell) activity that destroys virus-infected cells	
Antiprotozoal	Allicin	Preventing the parasite's RNA, DNA and protein synthesis.	[58]
	DATS		
Antioxidant	Ajoene	Inhibiting the human glutathione reductase and <i>T. cruzi</i> trypanothione reductase	[61]
	Allicin, DADS, and DATS	Modulation of ROS, increasing glutathione and cellular antioxidant enzymes	[54]
	Alliin	Controlling ROS generation and preventing mitogen-activated protein kinase (MAPK)	[67]
	DAS	Suppressing the enzymatic activity of cytochrome P450-2E1, reducing the generation of reactive oxygen and nitrogen species	[69]
Anti-inflammatory	Allicin	Enhancing the immune cell activity \uparrow , inhibiting the SDF1 α chemokine and Transendothelial migration of neutrophils	[60]
	DAS	Diminishing the expression of the inflammatory cytokines (e.g., NF- κ B, IL-1 β , and TNF- α), and ROS generation by suppressing CYP-2E1 hepatic enzyme	[76]
	Thiocremonone	Blocking the NF- κ B activity	[77]
Anti-cancer	Allicin, alliin, DADS, DAS	Enhancing p38 expression and cleaved caspase 3.	[80]
	Z-Ajoene	Stimulating apoptosis in human leukemic cells, promoting the peroxide production, caspase-3-like, and caspase-8 activities	[87]
Immunomodulatory	Allicin	Suppressing BuChE and AChE	[105]
Anti-obesity	Ajoene	Decreasing the fat accumulation in 3T3-L1 adipocytes and dramatically decreases the body weight gain	[117]
	1,2-Vinyldithiin	Decreasing the C/EBP α , PPAR γ 2, and LPL expression and the PPAR γ effect in human adipocytes	[118]
Antidiabetic	Allyl propyl disulfide, allicin, cysteine sulfoxide, and S-allyl cysteine sulfoxide, alliin	Decreasing the insulin secretion from pancreatic cells, increasing liver metabolism, and thus enhancing the short-acting insulin production	[114,115]
Hypolipidemic, hypocholesterolaemic	Different garlic preparations	Decreasing serum TC, TG, and LDL levels and moderately elevating HDL cholesterol	[107]
Anti-Atherosclerotic, antithrombotic	Different garlic preparations	Preventing ADP-activated platelets binding to immobilized fibrinogen and platelet aggregation, inhibiting GPIIb/IIIa receptor and increasing cAMP	[120]
Antihypertensive	Gamma-glutamylcysteine	Inhibiting the angiotensin-converting enzyme	[87]

4.7. Recommended Dose and Toxic Side Effects of Garlic

4.7.1. Recommended Dose

The generally recommended doses of the daily garlic uptake for the elderly are 4 g of raw garlic or 7.2 g of AGE or one dried garlic powder tablet twice to thrice per day [125]. Rana et al. [82] revealed that oral or intraperitoneal administration 50 mg/kg of garlic to rats did not show any effect on liver and lung tissue, while intake garlic at 250, 500, and 1000 mg/kg per day led to acute deformities in the rat's liver and lung tissue, suggesting the dose-related toxicity. While garlic intake at a dose of 500 and 1000 mg/kg/day remarkably decreased the auto-antioxidants without changing the lipid peroxidation level, whereas the daily intake of 1000 mg/kg resulted in morphological deformities in the liver under light microscopy and ultrastructural levels. Moreover, histological examination revealed nonspecific focal injury to the hepatocytes. In addition to this, Mikaili et al. [51] reported that garlic bulb extracts ingestion to male and female rats at 300 and 600 mg for 21 days, led to delayed growth and affects the biological and histological parameters. In particular, Asdaq and Inamdar. [126] indicated that the combination therapy of 250 mg/kg of garlic with hydrochlorothiazide shows synergistic antihypertensive and cardioprotective activities against toxicity caused by fructose and isoproterenol. While the combination therapy of 250 mg/kg of garlic with propranolol revealed a remarkable elevation in the antioxidant enzymes activities throughout ischemic injury [33].

4.7.2. Adverse Effects and Toxicity

Although the US Food and Drug Administration (FDA) considers garlic safe for humans, it can induce gastric agitation especially if ingested in high doses by sensitive people. To assess the safety of garlic, randomized controlled trials were performed, side effects such as insomnia, vomiting, heartburn, dizziness, diarrhea, tachycardia, nausea, bloating, flushing, headache, mild orthostatic hypotension, sweating, offensive body odor, and flatulence were observed [82]. Ingestion of raw garlic in high doses on an empty stomach can induce changes in the intestinal flora, flatulence and gastrointestinal upset [127]. Moreover, blisters dermatitis and burns were observed from raw garlic local applications [127]. Garlic does not seem to affect the drug metabolism, although recent reports on healthy participants show inconsistent results regarding the garlic effect in the pharmacokinetics of protease inhibitors, as well as anticoagulants due to its antithrombotic properties [46]. Many surgeons recommended stopping garlic administration in high doses up to 7 to 10 days prior to operation due to its effect to prolong the bleeding time that was observed in one patient with epidural spontaneous hematoma [46].

Previous in vivo experiments revealed that prolonged feeding of raw garlic in high doses led to weight loss and anemia due to red blood cells (RBCs) lysis, while administration of 5 mL/kg of raw garlic juice resulted in stomach injury that led finally to death [20]. Additionally, the chronic administration of 50 mg garlic powder per day led to anti-androgenic effects by inhibiting spermatogenesis in rats, leading to decrease sialic acid concentration in the seminal vesicles, testes, and epididymis with reduced Leydig cell function [82]. Oxidative hemolysis is the main toxicological mechanism of *Allium*-derived sulfur compounds and it is distinguished by methemoglobinemia development and Heinz body formation in the RBCs [128]. Initially, several clinical symptoms were observed including depression, vomiting, loss of appetite, abdominal pain, diarrhea, as well as anemia associated with pale mucous membrane, jaundice, rapid heart and respiratory rates, weakness, and hemoglobinuria [128]. *Allium* poisoning symptoms may appear after one day or several days of its ingestion based on the amounts taken [129].

Previous reports have reported the cardiovascular effects of garlic including potentially irreversible antiplatelet activity, anticoagulant, fibrinolytic activity, a remarkable decrease in platelet accumulation and mixed activity on fibrinolytic effectiveness [130]. Chen et al. [131] revealed that dehydrated raw garlic powder when administered orally resulted in acute injury to the gastric mucosa, whilst Yuncu et al. [132] reported that AGE, the sulfur-free compound, protects the intestinal mucosa of

experimental animals. Clinical studies reported that low doses of garlic are safe, whereas therapeutic doses might cause mild gastrointestinal disorders, while high doses have been reported to cause liver damage [82,123,133].

Allicin is a membrane-permeable compound that can enter cells easily and interact with cellular thiols such as glutathione or cysteine residues in proteins [37,89,90] as well as enzymes containing reactive cysteine and this may be the potential interpretation of allicin's toxicity [63]. Interestingly, Rana et al. [82] revealed that garlic powder or allicin at a concentration of 200 mg/mL can cause significant cell damages in the isolated rat liver.

5. Combination Therapy with Other Drugs

Recently, Mohammadi et al. [134] revealed the potent activity of garlic and ezetimibe combined treatment in reducing plasma LDL-C and TC, and thus, inhibiting the absorption of intestinal cholesterol and reducing the cardiovascular disorders risk factors. Asdaq and Inamdar [33] reported the combined effect of garlic homogenate and propranol in attenuating the isopropanol-mediated cardiac β_1 -receptors excessive stimulation, myocardial hypoperfusion, electrolyte imbalance, glycogen depletion, free radical injury, thermogenesis, lipid peroxidation, lipid accumulation, and electrocardiographic disturbances. They indicated that garlic homogenate is a good combination therapy as it reduces the dose and toxic side effects of propranol, which may assist in decreasing repeated higher doses of propranol. Mikaili et al. [51] reported the combined effect of allicin with polymyxin B against various yeasts and filamentous fungi and this combination therapy was found to increase the permeability of plasma membrane in *Saccharo cerevisiae*. Moreover, the combination treatments of garlic with captopril showed a higher synergistic effect regarding ACE inhibition [135]. Notably, the combination treatment of AGE with methotrexate showed improved activity against the significant increase in liver function enzymes, proinflammatory cytokines and antioxidants [136]. Recent researches reported that the fresh garlic extracts and antibiotics combination therapy resulted in high antibacterial activity. For instance, Ismail et al. [137] revealed that aqueous garlic extract-ampicillin combined treatment exhibited a potent synergetic effect towards *Kl. pneumoniae*, *Sal. typhi*, *E. coli*, and *P. aeruginosa*. Moreover, Vathsala and Murthy [138] revealed the potent immunomodulatory and anti-plasmodial effect of garlic-artemether combination treatment. They reported that this combined therapy may have a potential role in reducing organ injury and protecting against *Plasmodium* species by affecting NO production, suggesting novel treatment options against malaria [138,139].

6. Conclusions

This review focused on the chemical constituents and pharmacological activities of *A. sativum*. Sulfur-containing compounds such as alliin, allicin, ajoenes, vinyldithiins, and sulfides, are the main constituents isolated from *A. sativum* extracts. Extracts and isolated compounds from *A. sativum* reported to possess several biological properties including anticarcinogenic, antioxidant, antidiabetic, renoprotective, anti-atherosclerotic, antibacterial, antifungal, antiprotozoal, and antihypertensive activities. Garlic is also well-known to have immunomodulatory and anti-inflammatory activities. Allicin, the active substance of the garlic, can induce gastric agitation especially if administered in high doses. In addition to that, *A. sativum* has been reported to affect the pharmacokinetics of antiretroviral drugs, as well as anticoagulants. Thus, proper consideration should be taken when using garlic as a medicine for the treatment of different diseases.

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Abbreviations

AGE	aged garlic extract
PCSO	S-propyl cysteine-sulfoxide
MCSO	S-methyl cysteine-sulfoxide
NAC	N-Acetylcysteine
SAC	S-Allyl-cysteine
SAMC	S-ally-mercapto cysteine
PLP	pyridoxal phosphate
DAS	Diallyl sulfide
DADS	Diallyl disulfide
DATS	Diallyl trisulfide
AMS	Allyl methyl sulfide
AMDS	allyl methyl disulfide
SGF	simulated gastric fluid
SIF	stimulated intestinal fluid
HCMV	Human <i>Cytomegalovirus</i>
NK-cell	Natural killer-cell
ORS	oxygen-free radical species
SOD	superoxide dismutase
GSH-Px	glutathione peroxidase
GCLM	glutamate-cysteine ligase modifier
HO-1	heme oxygenase-1 subunit
Nrf2	nuclear factor erythrobia-2 related factor 2
ARE	antioxidant response element
MAPK	mitogen-activated protein kinase
RANKL	receptor activator of nuclear factor-kappa B ligand
ABG	Aged black garlic
TNF- α	tumor necrosis factor- α
COX-2	cyclooxygenase-2
PGE ₂	prostaglandin E ₂
TLR4	toll-like receptor 4
IL-6	interleukin-6
GBM CSC	Glioblastoma multiforme cancer stem cells
AChE	Acetylcholinesterase
Ach	acetylcholine
AD	Alzheimer's disease
BuChE	butyrylcholinesterase
TG	triglyceride
TC	total cholesterol
UCP	mitochondrial inner membrane proteins
AMPK	AMP-activated protein kinase
NO	nitric oxide; H ₂ S: hydrogen sulphide
FDA	Food and Drug Administration
RBCs	red blood cells
ACE	Angiotensin-converting enzyme
IUPAC	International Union of Pure and Applied Chemistry

References

1. Ríos, J.L.; Recio, M.C. Medicinal plants and antimicrobial activity. *J. Ethnopharmacol.* **2005**, *100*, 80–84. [[CrossRef](#)] [[PubMed](#)]

2. Beshbishy, A.M.; Batiha, G.E.S.; Adeyemi, O.S.; Yokoyama, N.; Igarashi, I. Inhibitory effects of methanolic *Olea europaea* and acetic *Acacia laeta* on the growth of *Babesia* and *Theileria*. *Asian Pac. J. Trop. Med.* **2019**, *12*, 425–434.
3. Batiha, G.E.S.; Beshbishy, A.A.; Tayebwa, D.S.; Shaheen, M.H.; Yokoyama, N.; Igarashi, I. Inhibitory effects of *Syzygium aromaticum* and *Camellia sinensis* methanolic extracts on the growth of *Babesia* and *Theileria* parasites. *Ticks Tick Borne Dis.* **2019**, *10*, 949–958. [[CrossRef](#)] [[PubMed](#)]
4. Batiha, G.E.S.; Beshbishy, A.A.; Adeyemi, O.S.; Nadwa, E.; Rashwan, E.; Yokoyama, N.; Igarashi, I. Safety and efficacy of hydroxyurea and eflornithine against most blood parasites *Babesia* and *Theileria*. *PLoS ONE* **2020**, *15*, e0228996.
5. Batiha, G.-S.; Beshbishy, A.M.; Alkazmi, L.M.; Adeyemi, O.S.; Nadwa, E.H.; Rashwan, E.K.; El-Mleeh, A.; Igarashi, I. Gas chromatography-mass spectrometry analysis, phytochemical screening and antiprotozoal effects of the methanolic *Viola tricolor* and acetic *Laurus nobilis* extracts. *BMC Complement. Altern. Med.* **2020**, in press. [[CrossRef](#)]
6. Batiha, G.E.S.; Beshbishy, A.M.; Tayebwa, D.S.; Adeyemi, O.S.; Shaheen, H.; Yokoyama, N.; Igarashi, I. Evaluation of the inhibitory effect of ivermectin on the growth of *Babesia* and *Theileria* parasites in vitro and in vivo. *Trop. Med. Health* **2019**, *47*, 42. [[CrossRef](#)]
7. Essawi, T.; Srouf, M. Screening of some Palestinian medicinal plants for antibacterial activity. *J. Ethnopharmacol.* **2000**, *70*, 343–349. [[CrossRef](#)]
8. Batiha, G.E.S.; Beshbishy, A.M.; Tayebwa, D.S.; Adeyemi, O.S.; Shaheen, H.; Yokoyama, N.; Igarashi, I. The effects of *trans*-chalcone and chalcone 4 hydrate on the growth of *Babesia* and *Theileria*. *PLoS Negl. Trop. Dis.* **2019**, *13*, e0007030. [[CrossRef](#)]
9. Beshbishy, A.M.; Batiha, G.E.; Yokoyama, N.; Igarashi, I. Ellagic acid microspheres restrict the growth of *Babesia* and *Theileria* in vitro and *Babesia microti* in vivo. *Parasites Vectors* **2019**, *12*, 269. [[CrossRef](#)]
10. Batiha, G.E.S.; Beshbishy, A.A.; Tayebwa, D.S.; Adeyemi, O.S.; Yokoyama, N.; Igarashi, I. Anti-piroplasmic potential of the methanolic *Peganum harmala* seeds and ethanolic *Artemisia absinthium* leaf extracts. *J. Protozool. Res.* **2019**, *29*, 8–27.
11. Batiha, G.-S.; Beshbishy, A.M.; Adeyemi, O.S.; Nadwa, E.H.; Rashwan, E.M.; Alkazmi, L.M.; Elkelish, A.A.; Igarashi, I. Phytochemical screening and antiprotozoal effects of the methanolic *Berberis vulgaris* and acetic *Rhus coriaria* extracts. *Molecules* **2020**, *25*, 550. [[CrossRef](#)] [[PubMed](#)]
12. Dorman, H.J.; Deans, S.G. Antimicrobial agents from plants: Antibacterial activity of plant volatile oils. *J. Appl. Microbiol.* **2000**, *88*, 308–316. [[CrossRef](#)] [[PubMed](#)]
13. Batiha, G.-S.; Alkazmi, L.M.; Nadwa, E.H.; Rashwan, E.K.; Beshbishy, A.M. Physostigmine: A plant alkaloid isolated from *Physostigma venenosum*: A review on pharmacokinetics, pharmacological and toxicological activities. *J. Drug Deliv. Ther.* **2020**, *10*, 187–190. [[CrossRef](#)]
14. Bakkali, F.; Averbeck, S.; Averbeck, D.; Idaomar, M. Biological effects of essential oils—A review. *Food Chem. Toxicol.* **2008**, *46*, 446–475. [[CrossRef](#)] [[PubMed](#)]
15. Batiha, G.-S.; Alkazmi, L.M.; Wasef, L.G.; Beshbishy, A.M.; Nadwa, E.H.; Rashwan, E.K. *Syzygium aromaticum* L. (Myrtaceae): Traditional uses, bioactive chemical constituents, pharmacological and toxicological activities. *Biomolecules* **2020**, *10*, 202. [[CrossRef](#)] [[PubMed](#)]
16. Ayaz, E.; Alposy, H.C. Garlic (*Allium sativum*) and traditional medicine. *Turkiye Parazitoloji Derg.* **2007**, *31*, 145–149.
17. Badal, D.S.; Dwivedi, A.K.; Kumar, V.; Singh, S.; Prakash, A.; Verma, S.; Kumar, J. Effect of organic manures and inorganic fertilizers on growth, yield and its attributing traits in garlic (*Allium sativum* L.). *J. Pharmacogn. Phytochem.* **2019**, *8*, 587–590.
18. Barnes, J.; Anderson, L.A.; Phillipson, J.D. *Herbal Medicines*, 2nd ed.; Pharmaceutical Press: London, UK, 2002; Volume 14.
19. Rahman, K. Historical perspective on garlic and cardiovascular disease. *J. Nutr.* **2001**, *131*, 977S–979S. [[CrossRef](#)]
20. Mathew, B.; Biju, R. Neuroprotective effects of garlic a review. *Libyan J. Med.* **2008**, *3*, 23–33.
21. Al-Jaber, N.A.; Awaad, A.S.; Moses, J.E. Review some antioxidant plants growing in Arab world. *J. Saudi Chem. Soc.* **2011**, *15*, 293–307. [[CrossRef](#)]
22. Wanwimolruk, S.; Prachayasittikul, V. Cytochrome P450 enzyme mediated herbal drug interactions (Part 1). *EXCLI J.* **2014**, *13*, 347–391. [[PubMed](#)]

23. Slusarenko, A.J.; Patel, A.; Portz, D. Control of plant diseases by natural products: Allicin from garlic as a case study. *Eur. J. Plant Pathol.* **2008**, *121*, 313. [[CrossRef](#)]
24. Rahman, M.S. Allicin and other functional active components in garlic: Health benefits and bioavailability. *Int. J. Food Prop.* **2007**, *10*, 245–268. [[CrossRef](#)]
25. Kaye, A.D.; De Witt, B.J.; Anwar, M.; Smith, D.E.; Feng, C.J.; Kadowitz, P.J.; Nossaman, B.D. Analysis of responses of garlic derivatives in the pulmonary vascular bed of the rat. *J. Appl. Physiol.* **2000**, *89*, 353–358. [[CrossRef](#)] [[PubMed](#)]
26. Lawson, L.D.; Gardner, C.D. Composition, stability, and bioavailability of garlic products used in a clinical trial. *J. Agric. Food Chem.* **2005**, *53*, 6254–6261. [[CrossRef](#)] [[PubMed](#)]
27. Rishton, G.M. Natural products as a robust source of new drugs and drug leads: Past successes and present day issues. *Am. J. Cardiol.* **2008**, *101*, S43–S49. [[CrossRef](#)]
28. Batiha, G.E.S.; Beshbishy, A.A.; Tayebwa, D.S.; Shaheen, M.H.; Yokoyama, N.; Igarashi, I. Inhibitory effects of *Uncaria tomentosa* bark, *Myrtus communis* roots, *Origanum vulgare* leaves and *Cuminum cyminum* seeds extracts against the growth of *Babesia* and *Theileria* in vitro. *Jpn. J. Vet. Parasitol.* **2018**, *17*, 1–13.
29. Sobolewska, D.; Podolak, I.; Makowska-Was, J. *Allium ursinum*: Botanical, phytochemical and pharmacological overview. *Phytochem. Rev.* **2015**, *14*, 81–97. [[CrossRef](#)]
30. Al-Snafi, A. Pharmacological effects of *Allium* species grown in Iraq. An overview. *Int. J. Pharm. Health Care Res.* **2013**, *1*, 132–147.
31. Zeng, Y.; Li, Y.; Yang, J.; Pu, X.; Du, J.; Yang, X.; Yang, T.; Yang, S. Therapeutic role of functional components in Alliums for preventive chronic disease in human being. *Evid. Based Complement. Altern. Med.* **2017**, *2017*, 9402849. [[CrossRef](#)]
32. Souza, G.A.; Ebaid, G.X.; Seiva, F.R.; Rocha, K.H.; Galhardi, C.M.; Mani, F.; Novelli, E.L. N-acetylcysteine an *Allium* plant compound improves high-sucrose diet-induced obesity and related effects. *Evid. Based Complement. Altern. Med.* **2011**, *2011*, 643269. [[CrossRef](#)] [[PubMed](#)]
33. Asdaq, S.M.B.; Inamdar, M.N. Pharmacodynamic and pharmacokinetic interactions of propranolol with garlic (*Allium sativum*) in rats. *Evid. Based Complement. Altern. Med.* **2011**, *2011*, 824042. [[CrossRef](#)] [[PubMed](#)]
34. Tran, G.B.; Dam, S.M.; Le, N.T.T. Amelioration of single clove black garlic aqueous extract on dyslipidemia and hepatitis in chronic carbon tetrachloride intoxicated Swiss Albino mice. *Int. J. Hepatol.* **2018**, *2018*, 9383950. [[CrossRef](#)] [[PubMed](#)]
35. Liu, Y.; Yan, J.; Han, X.; Hu, W. Garlic-derived compound S-allylmercaptocysteine (SAMC) is active against anaplastic thyroid cancer cell line 8305C (HPACC). *Technol. Health Care* **2015**, *23*, S89–S93. [[CrossRef](#)]
36. Cao, X.; Cao, L.; Ding, L.; Bian, J.S. A new hope for a devastating disease: Hydrogen sulfide in Parkinson's disease. *Mol. Neurobiol.* **2017**, *55*, 3789–3799. [[CrossRef](#)]
37. Miron, T.; Rabinkov, A.; Mirelman, D.; Wilchek, M.; Weiner, L. The mode of action of allicin: Its ready permeability through phospholipid membranes may contribute to its biological activity. *Biochim. Biophys. Acta* **2000**, *1463*, 20–30. [[CrossRef](#)]
38. Borlinghaus, J.; Albrecht, F.; Gruhlke, M.C.; Nwachukwu, I.D.; Slusarenko, A.J. Allicin: Chemistry and biological properties. *Molecules* **2014**, *19*, 12591–12618. [[CrossRef](#)]
39. Shimon, L.J.; Rabinkov, A.; Shin, I.; Miron, T.; Mirelman, D.; Wilchek, M.; Frolow, F. Two structures of alliinase from *Allium sativum* L.: Apo form and ternary complex with aminoacrylate reaction intermediate covalently bound to the PLP cofactor. *J. Mol. Biol.* **2007**, *366*, 611–625. [[CrossRef](#)]
40. De Rooij, B.M.; Boogaard, P.J.; Rijkse, D.A.; Commandeur, J.N.; Vermeulen, N.P. Urinary excretion of N-acetyl-S-allyl-L-cysteine upon garlic consumption by human volunteers. *Arch. Toxicol.* **1996**, *70*, 635–639. [[CrossRef](#)]
41. Freeman, F.; Kodera, Y. Garlic chemistry: Stability of S-(2-propenyl)-2-propene-1-sulfinothioate (Allicin) in blood, solvents, and simulated physiological fluids. *J. Agric. Food Chem.* **1995**, *43*, 2332–2338. [[CrossRef](#)]
42. Ilić, D.P.; Stojanović, S.; Najman, S.; Nikolić, V.D.; Stanojević, L.P.; Tačić, A.; Nikolić, L.B. Biological evaluation of synthesized allicin and its transformation products obtained by microwaves in methanol: Antioxidant activity and effect on cell growth. *Biotechnol. Biotechnol. Equip.* **2015**, *29*, 189–194. [[CrossRef](#)] [[PubMed](#)]
43. Thomson, M.; Ali, M. Garlic [*Allium sativum*]: A review of its potential use as an anti-cancer agent. *Curr. Cancer Drug Targets* **2003**, *3*, 67–81. [[CrossRef](#)] [[PubMed](#)]

44. Kuda, T.; Iwai, A.; Yano, T. Effect of red pepper *Capsicum annuum* var. conoides and garlic *Allium sativum* on plasma lipid levels and cecal microflora in mice fed beef tallow. *Food Chem. Toxicol.* **2004**, *42*, 1695–1700. [[CrossRef](#)] [[PubMed](#)]
45. Tesfaye, A.; Mengesha, W. Traditional uses, phytochemistry and pharmacological properties of garlic (*Allium Sativum*) and its biological active compounds. *Int. J. Sci. Res. Eng. Technol.* **2015**, *1*, 142–148.
46. Rahman, K.; Lowe, G.M. Garlic and cardiovascular disease: A critical review. *J. Nutr.* **2006**, *136*, 736s–740s. [[CrossRef](#)]
47. Davis, S.R. An overview of the antifungal properties of allicin and its breakdown products—the possibility of a safe and effective antifungal prophylactic. *Mycoses* **2005**, *48*, 95–100. [[CrossRef](#)]
48. Ross, Z.M.; O’Gara, E.A.; Hill, D.J.; Sleightholme, H.V.; Maslin, D.J. Antimicrobial properties of garlic oil against human enteric bacteria: Evaluation of methodologies and comparisons with garlic oil sulfides and garlic powder. *Appl. Environ. Microbiol.* **2001**, *67*, 475–480. [[CrossRef](#)]
49. Cutler, R.; Wilson, P. Antibacterial activity of a new, stable, aqueous extract of allicin against methicillin-resistant *Staphylococcus aureus*. *Br. J. Biomed. Sci.* **2004**, *61*, 71–74. [[CrossRef](#)]
50. Wallock-Richards, D.; Doherty, C.J.; Doherty, L.; Clarke, D.J.; Place, M.; Govan, J.R.; Campopiano, D.J. Garlic revisited: Antimicrobial activity of allicin-containing garlic extracts against Burkholderia cepacia complex. *PLoS ONE* **2014**, *9*, e112726. [[CrossRef](#)]
51. Mikaili, P.; Maadirad, S.; Moloudizargari, M.; Aghajanshakeri, S.; Sarahroodi, S. Therapeutic uses and pharmacological properties of garlic, shallot, and their biologically active compounds. *Iran. J. Basic Med. Sci.* **2013**, *16*, 1031–1048.
52. Meriga, B.; Mopuri, R.; MuraliKrishna, T. Insecticidal, antimicrobial and antioxidant activities of bulb extracts of *Allium sativum*. *Asian Pac. J. Trop. Med.* **2012**, *5*, 391–395. [[CrossRef](#)]
53. Shokrzadeh, M.; Ebadi, A.G. Antibacterial effect of garlic (*Allium sativum* L.) on *Staphylococcus aureus*. *Pak. J. Biol. Sci.* **2006**, *9*, 1577–1579.
54. Gruhlke, M.C.; Nwachukwu, I.; Arbach, M.; Anwar, A.; Noll, U.; Slusarenko, A.J. Allicin from garlic, effective in controlling several plant diseases, is a reactive sulfur species (RSS) that pushes cells into apoptosis. In Proceedings of the Modern fungicides and antifungal compounds VI. 16th International Reinhardsbrunn Symposium, Friedrichroda, Germany, 25–29 April 2010.
55. Pârvu, M.; Moț, C.A.; Pârvu, A.E.; Mircea, C.; Stoeber, L.; Roșca-Casian, O.; Țigu, A.B. *Allium sativum* extract chemical composition, antioxidant activity and antifungal effect against *Meyerozyma guilliermondii* and *Rhodotorula mucilaginosa* causing onychomycosis. *Molecules* **2019**, *24*, 3958. [[CrossRef](#)]
56. Fufa, B. Anti-bacterial and anti-fungal properties of garlic extract (*Allium sativum*): A review. *Microbiol. Res. J. Int.* **2019**, *28*, 1–5. [[CrossRef](#)]
57. Pai, S.T.; Platt, M.W. Antifungal effects of *Allium sativum* (garlic) extract against the *Aspergillus* species involved in otomycosis. *Lett. Appl. Microbiol.* **1995**, *20*, 14–18. [[CrossRef](#)] [[PubMed](#)]
58. Zhen, H.; Fang, F.; Ye, D.Y.; Shu, S.N.; Zhou, Y.F.; Dong, Y.S.; Nie, X.C.; Li, G. Experimental study on the action of allitridin against human cytomegalovirus in vitro: Inhibitory effects on immediate-early genes. *Antiviral Res.* **2006**, *72*, 68–74. [[CrossRef](#)]
59. Abdel-Ghaffar, F.; Semmler, M.; Al-Rasheid, K.A.; Strassen, B.; Fischer, K.; Aksu, G.; Klimpel, S.; Mehlhorn, H. The effects of different plant extracts on intestinal cestodes and on trematodes. *Parasitol. Res.* **2011**, *108*, 979–984. [[CrossRef](#)]
60. Abdel-Hafeez, E.H.; Ahmad, A.K.; Kamal, A.M.; Abdellatif, M.Z.; Abdelgelil, N.H. In vivo antiprotozoan effects of garlic (*Allium sativum*) and ginger (*Zingiber officinale*) extracts on experimentally infected mice with *Blastocystis* spp. *Parasitol. Res.* **2015**, *114*, 3439–3444. [[CrossRef](#)]
61. Gallwitz, H.; Bonse, S.; Martinez-Cruz, A.; Schlichting, I.; Schumacher, K.; Krauth-Siegel, R.L. Ajoene is an inhibitor and subversive substrate of human glutathione reductase and *Trypanosoma cruzi* trypanothione reductase: Crystallographic, kinetic, and spectroscopic studies. *J. Med. Chem.* **1999**, *42*, 364–372. [[CrossRef](#)]
62. Hazaa, I.K.K.; Al-Taai, N.A.; Khalil, N.K.; Zakri, A.M.M. Efficacy of garlic and onion oils on murin experimental *Cryptosporidium parvum* infection. *Al-Anbar J. Vet. Sci.* **2016**, *9*, 69–74.
63. Gruhlke, M.C.; Nicco, C.; Batteux, F.; Slusarenko, A.J. The effects of allicin, a reactive sulfur species from garlic, on a selection of mammalian cell lines. *Antioxidants* **2016**, *6*, 1. [[CrossRef](#)] [[PubMed](#)]

64. Sawai, T.; Itoh, Y.; Ozaki, H.; Isoda, N.; Okamoto, K.; Kashima, Y.; Kawaoka, Y.; Takeuchi, Y.; Kida, H.; Ogasawara, K. Induction of cytotoxic T-lymphocyte and antibody responses against highly pathogenic avian influenza virus infection in mice by inoculation of a pathogenic H5N1 influenza virus particles inactivated with formalin. *Immunology* **2008**, *124*, 155–165. [[CrossRef](#)] [[PubMed](#)]
65. Jang, H.J.; Lee, H.J.; Yoon, D.K.; Ji, D.S.; Kim, J.H.; Lee, C.H. Antioxidant and antimicrobial activities of fresh garlic and aged garlic by-products extracted with different solvents. *Food Sci. Biotechnol.* **2017**, *27*, 219–225. [[CrossRef](#)]
66. Liu, J.; Guo, W.; Yang, M.L.; Liu, L.X.; Huang, S.X.; Tao, L.; Zhang, F.; Liu, Y.S. Investigation of the dynamic changes in the chemical constituents of Chinese “laba” garlic during traditional processing. *RSC Adv.* **2018**, *8*, 41872–41883. [[CrossRef](#)]
67. Chen, Y.; Sun, J.; Dou, C.; Li, N.; Kang, F.; Wang, Y.; Cao, Z.; Yang, X.; Dong, S. Alliin attenuated RANKL-induced osteoclastogenesis by scavenging reactive oxygen species through inhibiting Nox1. *Int. J. Mol. Sci.* **2016**, *17*, 1516. [[CrossRef](#)] [[PubMed](#)]
68. Shang, A.; Cao, S.Y.; Xu, X.Y.; Gan, R.Y.; Tang, G.Y.; Corke, H.; Mavumengwana, V.; Li, H.B. Bioactive compounds and biological functions of garlic (*Allium sativum* L.). *Foods* **2019**, *8*, 246. [[CrossRef](#)] [[PubMed](#)]
69. Abdel-Daim, M.M.; Shaheen, H.M.; Abushouk, A.I.; Toraih, E.A.; Fawzy, M.S.; Alansari, W.S.; Aleya, L.; Bungau, S. Thymoquinone and diallyl sulfide protect against fipronil-induced oxidative injury in rats. *Environ. Sci. Pollut. Res. Int.* **2018**, *25*, 23909–23916. [[CrossRef](#)]
70. Ahmad, T.A.; El-Sayed, B.A.; El-Sayed, L.H. Development of immunization trials against *Eimeria* spp. *Trials Vaccinol.* **2016**, *5*, 38–47. [[CrossRef](#)]
71. Hobauer, R.; Frass, M.; Gmeiner, B.; Kaye, A.D.; Frost, E.A. Garlic extract (*Allium sativum*) reduces migration of neutrophils through endothelial cell monolayers. *Middle East J. Anaesthesiol.* **2000**, *15*, 649–658.
72. Gu, X.; Wu, H.; Fu, P. Allicin attenuates inflammation and suppresses HLA-B27 protein expression in ankylosing spondylitis mice. *BioMed Res. Int.* **2013**, *2013*, 171573. [[CrossRef](#)]
73. Jeong, Y.Y.; Ryu, J.H.; Shin, J.H.; Kang, M.J.; Kang, J.R.; Han, J.; Kang, D. Comparison of anti-Oxidant and anti-Inflammatory effects between fresh and aged black garlic extracts. *Molecules* **2016**, *21*, 430. [[CrossRef](#)] [[PubMed](#)]
74. You, B.R.; Yoo, J.M.; Baek, S.Y.; Kim, M.R. Anti-inflammatory effect of aged black garlic on 12-O-tetradecanoylphorbol-13-acetate-induced dermatitis in mice. *Nutr. Res. Pract.* **2019**, *13*, 189–195. [[CrossRef](#)] [[PubMed](#)]
75. Sela, U.R.; Ganor, S.; Hecht, I.; Brill, A.; Miron, T.; Rabinkov, A.; Wilchek, M.; Mirelman, D.; Lider, O.; Hershkovich, R. Allicin inhibits SDF-1 α -induced T cell interactions with fibronectin and endothelial cells by down-regulating cytoskeleton rearrangement, Pyk-2 phosphorylation and VLA-4 expression. *Immunology* **2004**, *111*, 391–399. [[CrossRef](#)] [[PubMed](#)]
76. Abdel-Daim, M.M.; Abushouk, A.I.; Bungău, S.G.; Bin-Jumah, M.; El-Kott, A.F.; Shati, A.A.; Aleya, L.; Alkahtani, S. Protective effects of thymoquinone and diallyl sulphide against malathion-induced toxicity in rats. *Environ. Sci. Pollut. Res.* **2020**, 1–8. [[CrossRef](#)]
77. Jin, P.; Kim, J.A.; Choi, D.Y.; Lee, Y.J.; Jung, H.S.; Hong, J.T. Anti-inflammatory and anti-amyloidogenic effects of a small molecule, 2,4-bis(p-hydroxyphenyl)-2-butenal in Tg2576 Alzheimer’s disease mice model. *J. Neuroinflamm.* **2013**, *10*, 767. [[CrossRef](#)]
78. Li, Z.; Le, W.; Cui, Z. A novel therapeutic anticancer property of raw garlic extract via injection but not ingestion. *Cell Death Discov.* **2018**, *4*, 108. [[CrossRef](#)]
79. Chhabria, S.V.; Akbarsha, M.A.; Li, A.P.; Kharkar, P.S.; Desai, K.B. In situ allicin generation using targeted alliinase delivery for inhibition of MIA PaCa-2 cells via epigenetic changes, oxidative stress and cyclin-dependent kinase inhibitor (CDKI) expression. *Apoptosis* **2015**, *20*, 1388–1409. [[CrossRef](#)]
80. Zhang, X.; Zhu, Y.; Duan, W.; Feng, C.; He, X. Allicin induces apoptosis of the MGC-803 human gastric carcinoma cell line through the p38 mitogen-activated protein kinase/caspase-3 signaling pathway. *Mol. Med. Rep.* **2015**, *11*, 2755–2760. [[CrossRef](#)]
81. Prager-Khoutorsky, M.; Goncharov, I.; Rabinkov, A.; Mirelman, D.; Geiger, B.; Bershadsky, A.D. Allicin inhibits cell polarization, migration and division via its direct effect on microtubules. *Cell Motil. Cytoskelet.* **2007**, *64*, 321–337. [[CrossRef](#)]
82. Rana, S.V.; Pal, R.; Vaiphei, K.; Sharma, S.K.; Ola, R.P. Garlic in health and disease. *Nutr. Res. Rev.* **2011**, *24*, 60–71. [[CrossRef](#)]

83. Iciek, M.; Kwiecień, I.; Chwatko, G.; Sokołowska-Jeżewicz, M.; Kowalczyk-Pachel, D.; Rokita, H. The effects of garlic-derived sulfur compounds on cell proliferation, caspase 3 activity, thiol levels and anaerobic sulfur metabolism in human hepatoblastoma HepG2 cells. *Cell Biochem. Funct.* **2012**, *30*, 198–204. [[CrossRef](#)] [[PubMed](#)]
84. Singh, V.; Belloir, C.; Siess, M.H.; Le Bon, A.M. Inhibition of carcinogen-induced DNA damage in rat liver and colon by garlic powders with varying alliin content. *Nutr. Cancer* **2006**, *55*, 178–184. [[CrossRef](#)] [[PubMed](#)]
85. Fleischauer, A.T.; Arab, L. Garlic and cancer: A critical review of the epidemiologic literature. *J. Nutr.* **2001**, *131*, 1032s–1040s. [[CrossRef](#)] [[PubMed](#)]
86. Piscitelli, S.C.; Burstein, A.H.; Welden, N.; Gallicano, K.D.; Falloon, J. The effect of garlic supplements on the pharmacokinetics of saquinavir. *Clin. Infect. Dis.* **2002**, *34*, 234–238. [[CrossRef](#)]
87. Bayan, L.; Koulivand, P.H.; Gorji, A. Garlic: A review of potential therapeutic effects. *Avicenna J. Phytomed.* **2014**, *4*, 1–14.
88. VinayKumar, D.K. *Robbins Basic Pathology* [Edited by] Vinay Kumar, Ramzi S. Cotran, Stanley L. Robbins; with Illustrations by James A. Perkins; Saunders: Philadelphia, PA, USA, 2003.
89. Dall'Acqua, S.; Maggi, F.; Minesso, P.; Salvagno, M.; Papa, F.; Vittori, S.; Innocenti, G. Identification of non-alkaloid acetylcholinesterase inhibitors from *Ferulago campestris* (Besser) Grecescu (Apiaceae). *Fitoterapia* **2010**, *81*, 1208–1212. [[CrossRef](#)]
90. Lu, S.H.; Wu, J.W.; Liu, H.L.; Zhao, J.H.; Liu, K.T.; Chuang, C.K.; Lin, H.Y.; Tsai, W.B.; Ho, Y. The discovery of potential acetylcholinesterase inhibitors: A combination of pharmacophore modeling, virtual screening, and molecular docking studies. *J. Biomed. Sci.* **2011**, *18*, 8. [[CrossRef](#)]
91. Akinyemi, A.J.; Lekan Faboya, A.P.; Awonegan, I.O.; Anadozie, S.; Oluwasola, T.A. Antioxidant and anti-Acetylcholinesterase activities of essential oils from garlic (*Allium sativum*) Bulbs. *Int. J. Plant Res.* **2018**, *31*. [[CrossRef](#)]
92. Singh, P.; Shukla, R.; Prakash, B.; Kumar, A.; Singh, S.; Mishra, P.K.; Dubey, N.K. Chemical profile, antifungal, antiaflatoxic and antioxidant activity of *Citrus maxima* Burm. and *Citrus sinensis* (L.) Osbeck essential oils and their cyclic monoterpene, DL-limonene. *Food Chem. Toxicol.* **2010**, *48*, 1734–1740. [[CrossRef](#)]
93. Borek, C. Garlic reduces dementia and heart-disease risk. *J. Nutr.* **2006**, *136*, 810S–812S. [[CrossRef](#)]
94. Jackson, R.; McNeil, B.; Taylor, C.; Holl, G.; Ruff, D.; Gwebu, E. Effect of aged garlic extract on human recombinant caspase-3 activity. *J. Ala. Acad. Sci.* **2003**, *74*, 121–122.
95. Mbyirukira, G.; Gwebu, E.T. Aged garlic extract protects serum-deprived PC12 cells from apoptosis. *J. Ala. Acad. Sci.* **2003**, *74*, 127–128.
96. Haider, S.; Naz, N.; Khaliq, S.; Perveen, T.; Haleem, D.J. Repeated administration of fresh garlic increases memory retention in rats. *J. Med. Food* **2008**, *11*, 675–679. [[CrossRef](#)] [[PubMed](#)]
97. Mukherjee, D.; Banerjee, S. Learning and memory promoting effects of crude garlic extract. *Indian J. Exp. Biol.* **2013**, *51*, 1094–1100.
98. Kumar, S. Dual inhibition of acetylcholinesterase and butyrylcholinesterase enzymes by allicin. *Indian J. Pharmacol.* **2015**, *47*, 444–446. [[CrossRef](#)]
99. Hogan, D.B. Progress update: Pharmacological treatment of Alzheimer's disease. *Neuropsychiatr. Dis. Treat.* **2007**, *3*, 569–578.
100. Schmitt, B.; Bernhardt, T.; Moeller, H.J.; Heuser, I.; Frölich, L. Combination therapy in Alzheimer's disease. *CNS Drugs* **2004**, *18*, 827–844. [[CrossRef](#)]
101. Weggen, S.; Eriksen, J.L.; Das, P.; Sagi, S.A.; Wang, R.; Pietrzik, C.U.; Findlay, K.A.; Smith, T.E.; Murphy, M.P.; Bulter, T.; et al. A subset of NSAIDs lower amyloidogenic A β 42 independently of cyclooxygenase activity. *Nature* **2001**, *414*, 212. [[CrossRef](#)]
102. Millard, C.B.; Shnyrov, V.L.; Newstead, S.; Shin, I.; Roth, E.; Silman, I.; Weiner, L. Stabilization of a metastable state of *Torpedo californica* acetylcholinesterase by chemical chaperones. *Protein Sci.* **2003**, *12*, 2337–2347. [[CrossRef](#)]
103. Inglis, F. The tolerability and safety of cholinesterase inhibitors in the treatment of dementia. *Int. J. Clin. Pract. Suppl.* **2002**, *127*, 45–63.
104. Jann, M.W.; Shirley, K.L.; Small, G.W. Clinical pharmacokinetics and pharmacodynamics of cholinesterase inhibitors. *Clin. Pharmacokinet.* **2002**, *41*, 719–739. [[CrossRef](#)] [[PubMed](#)]

105. Liu, S.G.; Ren, P.Y.; Wang, G.Y.; Yao, S.X.; He, X.J. Allicin protects spinal cord neurons from Glutamate-induced oxidative stress through regulating the heat shock protein 70/inducible nitric oxide synthase pathway. *Food Funct.* **2015**, *6*, 321–330. [[CrossRef](#)] [[PubMed](#)]
106. Hosseini, A.; Hosseinzadeh, H. A review on the effects of *Allium sativum* (Garlic) in metabolic syndrome. *J. Endocrinol. Investig.* **2015**, *38*, 1147–1157. [[CrossRef](#)] [[PubMed](#)]
107. Qidwai, W.; Ashfaq, T. Role of garlic usage in cardiovascular disease prevention: An evidence-based approach. *Evid. Based Complement. Altern. Med.* **2013**, *2013*, 125649. [[CrossRef](#)]
108. Iweala, E.E.; Akubugwo, E.I.; Okeke, C.U. Effect of ethanolic extracts of *Allium sativum* Linn. Liliaceae on serum cholesterol and blood sugar levels of albino rabbits. *Plant Prod. Res. J.* **2005**, *9*, 14–18.
109. Campbell, J.H.; Efendy, J.L.; Smith, N.J.; Campbell, G.R. Molecular basis by which garlic suppresses atherosclerosis. *J. Nutr.* **2001**, *131*, 1006S–1009S. [[CrossRef](#)]
110. Sobenin, I.A.; Nedosugova, L.V.; Filatova, L.V.; Balabolkin, M.I.; Gorchakova, T.V.; Orekhov, A.N. Metabolic effects of time-released garlic powder tablets in type 2 diabetes mellitus: The results of double-blinded placebo-controlled study. *Acta Diabetol.* **2008**, *45*, 1–6. [[CrossRef](#)]
111. Sobenin, I.A.; Pryanishnikov, V.V.; Kunnova, L.M.; Rabinovich, Y.A.; Martirosyan, D.M.; Orekhov, A.N. The effects of time-released garlic powder tablets on multifunctional cardiovascular risk in patients with coronary artery disease. *Lipids Health Dis.* **2010**, *9*, 119. [[CrossRef](#)]
112. Ashraf, R.; Aamir, K.; Shaikh, A.R.; Ahmed, T. Effects of garlic on dyslipidemia in patients with type 2 diabetes mellitus. *J. Ayub. Med. Coll. Abbottabad.* **2005**, *17*, 60–64.
113. Patel, D.K.; Prasad, S.K.; Kumar, R.; Hemalatha, S. An overview on antidiabetic medicinal plants having insulin mimetic property. *Asian Pac. J. Trop. Biomed.* **2012**, *2*, 320–330. [[CrossRef](#)]
114. Faroughi, F.; Mohammad-Alizadeh Charandabi, S.; Javadzadeh, Y.; Mirghafourvand, M. Effects of garlic pill on blood glucose level in borderline gestational diabetes mellitus: A triple blind, randomized clinical trial. *Iran. Red. Crescent Med. J.* **2018**, *20*, e60675. [[CrossRef](#)]
115. Zhai, B.; Zhang, C.; Sheng, Y.; Zhao, C.; He, X.; Xu, W.; Huang, K.; Luo, Y. Hypoglycemic and hypolipidemic effect of S-allyl-cysteine sulfoxide (alliin) in DIO mice. *Sci. Rep.* **2018**, *8*, 3527. [[CrossRef](#)] [[PubMed](#)]
116. Lee, M.S.; Kim, I.H.; Kim, C.T.; Kim, Y. Reduction of body weight by dietary garlic is associated with an increase in uncoupling protein mRNA expression and activation of AMP-activated protein kinase in diet-induced obese mice. *J. Nutr.* **2011**, *141*, 1947–1953. [[CrossRef](#)] [[PubMed](#)]
117. Han, C.Y.; Ki, S.H.; Kim, Y.W.; Noh, K.; Lee da, Y.; Kang, B.; Ryu, J.H.; Jeon, R.; Kim, E.H.; Hwang, S.J.; et al. Ajoene, a stable garlic by-product, inhibits high fat diet-induced hepatic steatosis and oxidative injury through LKB1-dependent AMPK activation. *Antioxid. Redox Signal.* **2011**, *14*, 187–202. [[CrossRef](#)]
118. Keophiphath, M.; Priem, F.; Jacquemond-Collet, I.; Clément, K.; Lacasa, D. 1,2-Vinyldithiin from garlic inhibits differentiation and inflammation of human preadipocytes. *J. Nutr.* **2009**, *139*, 2055–2060. [[CrossRef](#)]
119. Varshney, R.; Budoff, M.J. Garlic and Heart Disease. *J. Nutr.* **2016**, *146*, 416S–421S. [[CrossRef](#)]
120. Drobiova, H.; Thomson, M.; Al-Qattan, K.; Peltonen-Shalaby, R.; Al-Amin, Z.; Ali, M. Garlic increases antioxidant levels in diabetic and hypertensive rats determined by a modified peroxidase method. *Evid. Based Complement. Altern. Med.* **2011**, *2011*, 703049. [[CrossRef](#)]
121. Sobenin, I.; Andrianova, I.; Ionova, V.; Karagodin, V.; Orekhov, A. Anti-aggregatory and fibrinolytic effects of time-released garlic powder tablets. *Med. Health Sci. J.* **2012**, *10*, 47–51. [[CrossRef](#)]
122. Dubey, H.; Singh, A.; Patole, A.M.; Tenpe, C.R. Antihypertensive effect of allicin in dexamethasone-induced hypertensive rats. *Integr. Med. Res.* **2017**, *6*, 60–65. [[CrossRef](#)]
123. Ried, K.; Fakler, P. Potential of garlic (*Allium sativum*) in lowering high blood pressure: Mechanisms of action and clinical relevance. *Integr. Blood Press. Control* **2014**, *7*, 71–82. [[CrossRef](#)]
124. Rahman, K.; Lowe, G.M.; Smith, S. Aged garlic extract inhibits human platelet aggregation by altering intracellular signaling and platelet shape change. *J. Nutr.* **2016**, *146*, 410S–415S. [[CrossRef](#)] [[PubMed](#)]
125. Tattelman, E. Health effects of garlic. *Am. Fam. Physician* **2005**, *72*, 103–106. [[PubMed](#)]
126. Asdaq, S.M.B.; Inamdar, M.N. The potential benefits of a garlic and hydrochlorothiazide combination as antihypertensive and cardioprotective in rats. *J. Nat. Med.* **2011**, *65*, 81–88. [[CrossRef](#)]
127. Piasek, A.; Bartoszek, A.; Namiesnik, J. Phytochemicals that counteract the cardiotoxic side effects of cancer chemotherapy. *Postepy Hig. Med. Dosw.* **2009**, *63*, 142–158.
128. Salgado, B.; Monteiro, L.; Rocha, N. *Allium* species poisoning in dogs and cats. *J. Venom. Anim. Toxins Incl. Trop. Dis.* **2011**, *17*, 4–11. [[CrossRef](#)]

129. Lee, L.S.; Andrade, A.S.; Flexner, C. Interactions between natural health products and antiretroviral drugs: Pharmacokinetic and pharmacodynamic effects. *Clin. Infect. Dis.* **2006**, *43*, 1052–1059. [[CrossRef](#)]
130. Borrelli, F.; Capasso, R.; Izzo, A.A. Garlic (*Allium sativum* L.): Adverse effects and drug interactions in humans. *Mol. Nutr. Food Res.* **2007**, *51*, 1386–1397. [[CrossRef](#)]
131. Chen, K.; Xie, K.; Liu, Z.; Nakasone, Y.; Sakao, K.; Hossain, A.; Hou, D.X. Preventive effects and mechanisms of garlic on dyslipidemia and gut microbiome dysbiosis. *Nutrients* **2019**, *11*, 1225. [[CrossRef](#)]
132. Yuncu, M.; Eralp, A.; Celik, A. Effect of aged garlic extract against methotrexate-induced damage to the small intestine in rats. *Phytother. Res.* **2006**, *20*, 504–510. [[CrossRef](#)]
133. Almogren, A.; Shakoor, Z.; Adam, M.H. Garlic and onion sensitization among Saudi patients screened for food allergy: A hospital-based study. *Afr. Health Sci.* **2013**, *13*, 689–693. [[CrossRef](#)]
134. Mohammadi, A.; Vafaei, S.A.; Moradi, M.N.; Ahmadi, M.; Pourjafar, M.; Oshaghi, E.A. Combination of ezetimibe and garlic reduces serum lipids and intestinal niemann-pick C1-like 1 expression more effectively in hypercholesterolemic mice. *Avicenna J. Med. Biochem.* **2015**, *3*, e23205. [[CrossRef](#)]
135. Liu, M.Z.; Zhang, Y.L.; Zeng, M.Z.; He, F.Z.; Luo, Z.Y.; Luo, J.Q.; Wen, J.G.; Chen, X.P.; Zhou, H.H.; Zhang, W. Pharmacogenomics and herb-drug interactions: Merge of future and tradition. *Evid. Based Complement. Altern. Med.* **2015**, *2015*, 321091. [[CrossRef](#)] [[PubMed](#)]
136. Badr, G.M.; Arafa, N.S. Synergetic effect of aged garlic extract and methotrexate on rheumatoid arthritis induced by collagen in male albino rats. *Indian J. Exp. Biol.* **2020**, *58*, 33–38.
137. Ismail, R.M.; Saleh, A.H.A.; Ali, K.S. GC-MS analysis and antibacterial activity of garlic extract with antibiotic. *J. Med. Plants Stud.* **2020**, *8*, 26–30.
138. Vathsala, P.G.; Murthy, P.K. Immunomodulatory and antiparasitic effects of garlic–arteether combination via nitric oxide pathway in *Plasmodium berghei*-infected mice. *J. Parasit. Dis.* **2019**, 49–61. [[CrossRef](#)] [[PubMed](#)]
139. Beshbishy, A.M.; Batiha, G.E.-S.; Alkazmi, L.; Nadwa, E.; Rashwan, E.; Abdeen, A.; Yokoyama, N.; Igarashi, I. Therapeutic effects of atranorin towards the proliferation of *Babesia* and *Theileria* parasites. *Pathogen* **2020**, *9*, 127. [[CrossRef](#)] [[PubMed](#)]



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