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## Ginger and its health benefits: A review

**Syeda Azmat, Beenish Gulzar and Tabasum Fatima**

### Abstract

For the last 2500 years, ginger has been known for its several scientific properties and valued in different parts of the globe. Ginger is a spice that has traditionally been treated as medicine in both Traditional Chinese Medicine and Ayurveda. Ginger has rich phytochemistry and several health promoting perspectives. In ginger family, *Zingiber officinalis* is one of most widely used species and it is found in several foods and beverages. Ginger has been used commonly to treat diarrhoea, stomach upset, indigestion and nausea. It also has anti-inflammatory and antioxidant properties. Ginger constituents include 17.77g carbohydrates, 1.7g sugars, 2g dietary fibre, 0.75g fat and 1.82g proteins per 100g. The chemistry of ginger is well documented with respect to its phytochemicals, oleoresins and volatile oils. This review presents the potential properties of ginger to treat numerous disorders including cancer due to its anti-inflammatory and anti-oxidant properties. It is also useful in controlling the process of aging. Ginger, one of the most commonly used spices and medicinal plants, has been demonstrated to improve diet-induced metabolic abnormalities. This scientific review favors ginger due to its rich phytochemistry; however, due to some ambiguities, it is recommended to conduct clinical trials of ginger with sound protocol design before claiming its efficacy.

**Keywords:** Ginger, health benefits

### Introduction

Ginger (*Zingiber officinale*) is a flowering plant whose rhizome, ginger root or ginger, is widely used as a spice and a folk medicine. It is a herbaceous perennial which grows annual pseudostems (false stems made of the rolled bases of leaves) about a meter tall bearing narrow leaf blades. The inflorescences bear pale yellow with purple flowers and arise directly from the rhizome on separate shoots. In 2017, global production of ginger was 3.3 million tonnes, led by India with 34% of the world total. Nigeria, China, and Indonesia also had substantial production.

### Ginger and Its Constituents

Ginger (*Zingiber officinale*), a member of the *Zingiberaceae* family, is a popular spice used globally especially in most of the Asian countries. Chemical analysis of ginger shows that it contains over 400 different compounds. The major constituents in ginger rhizomes are carbohydrates (50–70%), lipids (3–8%), terpenes, and phenolic compounds (Grzanna *et al* 2005) [31]. Terpene components of ginger include zingiberene,  $\beta$ -bisabolene,  $\alpha$ -farnesene,  $\beta$ -sesquiphellandrene, and  $\alpha$ -curcumene, while phenolic compounds include gingerol, paradols, and shogaol. These gingerols (23–25%) and shogaol (18–25%) are found in higher quantity than others. Besides these, amino acids, raw fiber, ash, protein, phytosterols, vitamins (e.g., nicotinic acid and vitamin A), and minerals are also present (Langner *et al* 1998, Shukla *et al* 2007) [21, 48]. The aromatic constituents include zingiberene and bisabolene, while the pungent constituents are known as gingerols and shogaols [58]. Other gingerol- or shogaol-related compounds (1–10%), which have been reported in ginger rhizome, include 6-paradol, 1-dehydrogingerdione, 6-ginger Dione and 10-gingerdione, 4-gingerdiol, 6-gingerdiol, 8-gingerdiol, and 10-gingerdiol, and diarylheptanoids (Govindrajan *et al*. 1982, Ali *et al* 2008). The characteristic odor and flavour of ginger are due to a mixture of volatile oils like shogaols and gingerols.

The rhizome of ginger plant has been used as a spice since several years across the globe. It was found that, ginger was one of widely used herbs in traditional Chinese, Ayurveda, Europe and America (Langner *et al.*, 1998; Avato *et al.*, 2000; Duke and Ayensu, 1985; Kapil *et al.*, 1990; Qureshi *et al.*, 1989; Blumenthal *et al.*, 1997; Kamtchoung *et al.*, 2000; Afzal *et al.*, 2011; Grzanna *et al.*, 2005) [31, 21, 48, 38, 1, 20, 63, 62]. The mode of administration of ginger is oral, intra muscular (IM) and topically (Barnes *et al.*, 2002; Yang and Chang, 1988;

Chrubasik *et al.*, 2005; Shukla and Singh, 2007) [11, 17, 68]. Historically, it has been used to treat nausea, vomiting, rheumatism, baldness, respiratory diseases and bleeding disorders (Young *et al.*, 2006; Suekawa *et al.*, 1984; Newall *et al.* 1996; Srivastava, 1984; Kim *et al.* 2005; Kelly *et al.*, 2009) [85, 71, 41, 40, 69].

### Therapeutic Properties of Ginger

The review article was written with help from secondary data analysis. Information on searching databases, various journals, books, articles and key words were used during writing of therapeutic properties of ginger.

#### Cardiovascular effects

Gingerol and shogaol classes of compounds might have many therapeutic effects including anti-inflammatory, antioxidant, and hypocholesterolemic effects, as suggested by many studies. Ginger enhances blood circulation throughout the body by diluting circulating blood and by enhanced stimulation of the heart muscle. This improves cellular metabolism and helps to relief cramp and tension (Gong *et al.*, 1989; Pecoraro *et al.*, 1998; Frisch *et al.*, 1995; Yamahara *et al.*, 1989; Ernst and Pittler, 2000; Chaiyakunapruk *et al.*, 2006) [14, 30, 24, 58, 58]. *In vitro* research indicates that gingerols and the related shogaols exhibit cardio depressant activity at low doses and cardiostimulant properties at higher doses (Wang CC *et al.* 2003) [79]. Both (6)-shogaol and (6)-gingerol, and the gingerdiones, are reportedly potent enzymatic inhibitors of prostaglandin, thromboxane, and leukotriene biosynthesis (Rajesh Kumar Mishra *et al.* 2012).

#### Hypotensive effect

Many studies have proven the hypotensive effect of ginger when it was given at 0.3-3 mg/kg. It helps to reduce atrial blood pressure by blocking calcium channel or by acting on muscarinic receptor (Ernst and Pittler, 2004; Portoni *et al.*, 2003; Ozgoli and Goli, 2009; Vutyavanich *et al.*, 2001) [24, 59, 57, 77].

#### Antinociceptive Effects

(6)-shogaol has produced anti-nociception and inhibited the release of substance P in rats, seemingly via the same receptor to which capsaicin binds. However, it was observed to be 100 times less potent and to elicit half the maximal effect of capsaicin (MaJ *et al.* 2004).

#### Gastrointestinal Effects

There is evidence that ginger rhizome (root) increases stomach acid production. If so, it may interfere with antacids, sucralfate (Carafate), H<sub>2</sub> antagonists, or proton pump inhibitors. In contrast, other *in vitro* and animal studies have revealed gastro protective properties (Thomson M *et al.* 2002, Al Yahya *et al.* 1989) [5, 7] in addition, (6) shogaol, generally more potent than (6)- gingerol, has inhibited intestinal motility in intravenous preparations and facilitated gastrointestinal motility in oral preparations. Ginger extract has also been reported to inhibit the growth of *Helicobacter pylori* *in vitro* (Srivastava KC *et al.* 1984) [69]

#### Anti-hypercholesterolaemic effect

Ginger extracts interferes with cholesterol biosynthesis leading to decreasing cholesterol levels. Ginger extracts have antilipidemic effects, by reducing thermogenesis and

high lipids levels. It also helps to increase serum HDL-cholesterol (Ernst and Pittler, 2004; Portoni *et al.*, 2003; Ozgoli and Goli, 2009; Vutyavanich *et al.*, 2001; AlAwwadi, 2010; 2013) [24, 59, 57, 77]. Gastrointestinal effect of ginger Ginger is very useful in the treatment of several gastrointestinal diseases including peptic and duodenal ulcer. Ulcer is generally caused due to imbalance between defensive and offensive factors like acid, pepsin and *Helicobacter pylori*; and in this case, ginger is useful due to its anti-inflammatory properties. Ginger acts and protects gastric mucosa against several ulcerogenic agents. Ginger is also very useful in cases of ulcerogenesis due to its antioxidant activities (Lumb, 1994; Gull *et al.*, 2012; Dugasani *et al.*, 2010; Halvorsen *et al.*, 2002) [51, 32, 19, 34].

#### Antiemetic effect of ginger

Ginger shows strong antiemetic property by enhancing intestinal motility and inhibiting serotonin receptors. It stimulates peripheral anti-cholinergic and ant-histaminic receptors and antagonises 5- hydroxytryptamine receptors in the GIT (Lumb, 1994; Gull *et al.*, 2012; Dugasani *et al.*, 2010; Halvorsen *et al.*, 2002) [51, 34, 19, 32]. Ginger anti-nausea effect due to chemotherapy Chemotherapy is known to cause severe nausea and vomiting. It has been proved that ginger is effective in preventing nausea and vomiting caused by chemotherapy. Gingerols the key ingredients responsible for the activity have shown pharmacological effect. It is also used to treat nausea after surgery and same has been proved in several randomised clinical trials. This effect is seen due to its action on the 5-HT<sub>3</sub> receptor (Ajith *et al.*, 2007; Krim *et al.*, 2013; Waggas, 2009; Sabina *et al.*, 2011; Ahmed *et al.*, 2008) [3, 43, 78, 66, 4]. Morning sickness FDA classifies ginger as safe for the treatment of morning sickness and it is widely used during early pregnancy. It reduces symptoms of morning sickness if same is taken in the recommended amount. The German Commission and Europe does not consider it as safe due to lack of published data (El-Sharaky *et al.*, 2009; Nasri *et al.*, 2013; Ajith *et al.*, 2008; El-Abhar *et al.*, 2008; Kyung *et al.*, 2006) [23, 55, 4, 22, 46]. Hematologic (platelets) effects of ginger Scientific evidence is still pending; however it was found that ginger is having anti-thrombotic and strong antiinflammatory effect due to increased fibrinolytic activity when same has been taken at about 5 g. It was found that Gingerols and Paradol have good anti-platelet and COX-I inhibitor properties (Mehdizadeh *et al.*, 2012; Jagetia *et al.*, 2004; Jagetia *et al.*, 2003) [13, 35, 54, 6]. The effect of the ginger is different if it is consumed dry or fresh. Regulation of blood glucose and lipid levels Ginger is very effective in lowering blood glucose level when same has been taken in dried form. It also decreases cholesterol and triglyceride level. Long term usage helps to increase high-density lipoprotein cholesterol concentrations (Duke and Ayensu, 1985; Afzal *et al.*, 2011; Kim *et al.*, 2007; Li *et al.*, 2012) [20].

#### Rheumatologic effect of ginger

Ginger exerts its anti-inflammatory effects by the mechanisms which explain the role of inhibition of preinflammatory factor like prostaglandin and leukotriene biosynthesis which can decline pain associated with rheumatoid and osteoarthritis. It is having proven history of treatment of rheumatic conditions (Duke and Ayensu, 1985; Avato *et al.*, 2000; Afzal *et al.*, 2011; Ha *et al.*, 2012) [20, 13, 15]. Headache Ginger is used for the treatment of headache

and having Al-Awwadi 113 good effect on reducing symptoms of pain. This effect is due to reduction in prostaglandin synthesis. It also has been reported that ginger suppresses leukotriene biosynthesis by inhibiting 5-lipoxygenase (Ernst and Pittler, 2004; Nasri *et al.*, 2013; Tjendraputra *et al.*, 2001) [24, 25, 55, 72]. Anti-Inflammatory effect Ginger is showing anti-inflammatory effect by suppression of PG synthesis and also interference in cytokine signalling (Duke and Ayensu, 1985; Uz *et al.*, 2009; Mahmoud *et al.*, 2012) [20, 73, 52].

#### Antigen toxic Activity

Norethandrolone and oxandrolone were investigated for their genotoxic effect on human lymphocyte chromosomes using chromosomal aberrations and sister chromatid exchanges as parameters and subsequently Genistein and [6]-gingerol were used as antigenotoxic agents to ameliorate the genotoxicity induced by the steroids. Norethandrolone and oxandrolone were studied at 5, 10, 20, 30 and 40 µM, respectively and were found to be significantly genotoxic at 30 and 40 µM. Genistein and [6] gingerol proved to be equally effective in reducing genotoxic damage at appropriate doses (Beg T *et al* 2008) [12].

#### Antimicrobial Effect

Due to phenolic compounds, ginger has shown excellent antimicrobial properties and effective in controlling virus, bacteria, fungal disease. In many countries, ginger is used to preserve food (Ernst and Pittler, 2004; Liao *et al.*, 2012; Chen *et al.*, 2009) [25, 50].

#### Antibacterial Effect

Ginger has shown good antimicrobial effect against both Gram positive and negative bacteria; however, severally, this effect is reduced due to heating (Jagetia *et al.*, 2004; Ha *et al.*, 2012; Tjendraputra *et al.*, 2001; Kubra *et al.*, 2013) [4, 16, 32, 6].

#### Antiparasitic Action

Ginger acts as anti-parasitic; study shows the *in vivo* potential of methanolic extract of *Zingiber officinale* in the treatment of trypanosomiasis (Halvorsen *et al.*, 2002; Jagetia *et al.*, 2003; Kubra *et al.*, 2013; Duarte, 2016; Kumar *et al.*, 2015; Choi *et al.*, 2013; Saraswat, 2010; Pushpanathan, 2008) [12, 14, 73, 37, 35, 18]. 114 J Pharmacognosy Phytother.

#### Antineoplastic

Ginger is a powerful antineoplastic agent. In several studies, extracts of ginger suppress cell proliferation and act against resistance of cancerous cells (Barnes *et al.*, 2002; Newall *et al.* 1996; Ernst and Pittler, 2000; Nasri *et al.*, 2013; Kumar *et al.*, 2015; Saraswat, 2010) [24, 55]. Antioxidant Ginger is having powerful antioxidant activity due to its oil which has protective effect on DNA damage. They have demonstrated this effect in many cell culture (Chaiyakunapruk *et al.*, 2006; Ramkisson *et al.*, 2012; Kabuto *et al.*, 2005; Mahmoud *et al.*, 2012; AlAwwadi, 2010; 2013) [14, 37, 42]. Ginger is a scavenger of free radicals Ginger oil has scavenging effects due to volatile oils and same has been proved in many studies (Duke and Ayensu, 1985; Avato *et al.*, 2000; Kamtchoung *et al.*, 2000; Kumar *et al.*, 2015; Pushpanathan, 2008) [20, 32, 16, 64, 34]. Lipid peroxidation Ginger has preventive effect on lipid

peroxidation and it inhibits or breaks its chain (Duke and Ayensu, 1985; Afzal *et al.*, 2011; Verma *et al.*, 1993) [20].

#### Anti-ulcerogenic effect of ginger

This has both many benefits and drawbacks. Prostaglandin has been shown to have housekeeping and gastro-protective function by maintaining gastric mucosal integrity (Duke and Ayensu, 1985; Qureshi *et al.*, 1989; El-Sharaky *et al.*, 2009; Ajith *et al.*, 2008; Duarte, 2016) [23, 20, 4, 18]. Modulation of biological activities by ginger Ginger modulates genetic pathway, acts on tumour suppression of genes and modulates biological Activities (Duke and Ayensu, 1985; Jagetia *et al.*, 2004; Ha *et al.*, 2012; Duarte, 2016) [20, 6, 18].

#### Conclusion

Although the medicinal properties of ginger have been known for thousands of years, a significant number of *in vitro*, *in vivo*, and epidemiological studies further provide substantial evidence that ginger and its active compounds are effective against wide variety of human diseases. This marvellous spice and medicinal plant, ginger, is constrained severely by the absence of seed set, and the breeder is left with the alternative of clonal selection or induced mutations with all its uncertainty and limitations. Biotechnology opened up many potential avenues such as tissue culture, somaclonal variation, *in vitro* mutagenesis and selection, molecular fingerprinting, recombinant DNA technology, and genetic modification through transgenic for creating disease-resistant lines. Concerted efforts are needed to solve the serious problems besetting this "great medicine" and "universal cure" as described in the Indian systems of medicine, which is a great spice unparalleled in the range of applications and uses. Moreover, most of the known activities of ginger components are based only on *in vitro* and *in vivo* studies, except for a few clinical studies in human subjects. Therefore, more extensive and well-controlled human studies are required to demonstrate its efficacy as a safe and cost-effective alternative.

#### References

1. Afzal M, al-hadidi D, Menon M, Pesek J, Dhama MS. Ginger: An Ethnomedical, Chemical and Pharmacological Review. *Drug Interact.* 2011; 18:159-190.
2. Ahmed RS, Suke SG, Seth V, Chakraborti A, Tripathi AK, Banerjee BD. Protective effects of dietary ginger (*Zingiber officinale* Rosc.) on lindane-induced oxidative stress in rats. *Phytother. Res.* 2008; 22(7):902-906.
3. Ajith TA, Aswathy MS, Hema U. Protective effect of *Zingiber officinale* roscoe against anticancer drug doxorubicin-induced acute nephrotoxicity. *Food Chem. Toxicol.* 2008; 46(9):3178-81.
4. Ajith TA, Nivitha V, Usha S. *Zingiber officinale* Roscoe alone and in combination with alpha-tocopherol protect the kidney against cisplatin-induced acute renal failure. *Food Chem. Toxicol.* 2007.
5. Al Yahya, Rafatullah MA, Mossa S, Ageel JS, Parmar AMNS, Tariq M. Gastroprotective activity of ginger *Zingiber officinale* rosc., in albino rats. *Am J Chin Med.* 1989; 17(1-2):51-56. 29.
6. Al-Awwadi 115 Huang Q, Iwamoto M, Aoki S, Tanaka N, Tajima K, Yamahara J. Anti-5-hydroxytryptamine effect of galanolactone, diterpenoid isolated from

- ginger. Chem. Pharm. Bull. Jagetia G, Baliga M, Venkatesh P. 2004; 39:397-399.
7. Al-Yahya MA, Rafatullah S, Mossa JS, Ageel AM, Parmar NS, Tariq M. Gastroprotective activity of ginger *Zingiber officinale* Rosc, in albino rats. Am. J Chin. Med. Al-Awwadi NAJ 2013. 1989; 17:51-56.
  8. Anonymous. Monographs on the medicinal uses of plants. Exeter: European Scientific Cooperative on Phytotherapy. Avato P, Tursil E, Vitali C, Miccolis V, Cadido V (2000). Allyl Sulphide Constituents of Garlic Volatile Oil as Antimicrobial Agents. Phytomedicine. 1997; 7:239-243.
  9. Anti diabetics effect of Achillea Santolina aqueous leaves extract, Al-Awwadi NAJ (2010). Effects of Achillea Santolina extracts and fractions on human platelet aggregation *in vitro* and on rat arteriovenous shunt thrombosis *in vivo*, Thi-Qar Med. J (TQMJ) 2010, 4(7):151-156.
  10. BH Ali, G Blunden MO Tanira, A Nemmar Some phytochemical, pharmacological and toxicological properties of ginger (*Zingiber officinale* Roscoe): a review of recent research,” Food and Chemical Toxicology, 2008; 46(2):409-420.
  11. Barnes KK, Kolpin DW, Meyer MT, Thurman EM, Furlong ET, Zaugg SD, *et al.* Water-quality data for pharmaceuticals, hormones, and other organic wastewater contaminants in U.S. streams, U.S. Geological Survey Open-File Report. 1999-2000. 02-94.
  12. Beg T, Siddiqe YS, Ara G, Gupta J, Afzal M, Antigenotoxic Effect of Genistein and gingerol on genotoxicity induced by norethandrolone and oxandrolone in cultured human lymphocytes. Int J Pharmaco. 2008; 4:177-183.
  13. Blumenthal M, Busse W, German CE. Monographs: Therapeutic Monographs on Medicinal Plants for Human Use. Austin, TX: American Botanical Council. 1997.
  14. Chaiyakunapruk N, Kitikannakorn N, Nathisuwan S, Leeprakobboon K, Leelasattagool C, The efficacy of ginger for the prevention of postoperative nausea and vomiting: a meta-analysis. Am. J Obstet. Gynecol. 2006; 194(1):95-99.
  15. Chen BH, Wu PY, Chen KM, Fu TF, Wang HM, Chen CY. Antiallergic potential on RBL-2H3 cells of some phenolic constituents of *Zingiber officinale* (Ginger) J Nat. Prod. 2009; 72:950-953.
  16. Choi YY, Kim MH, Hong J, Kim SH, Yang WM. Dried Ginger (*Zingiber officinalis*) Inhibits Inflammation in a Lipopolysaccharide Induced Mouse Model. Evidence-Based Complement. Altern. Med. 2013, 914563.
  17. Chrubasik S, Pittler MH, Roufogalis BD. Zingiberis rhizoma: a comprehensive review on the ginger effect and efficacy profiles. Phytomedicine. 2005; 12(9):684-701.
  18. Duarte, MC. Antileishmanial activity and mechanism of action from a purified fraction of *Zingiber officinalis* Roscoe against *Leishmania amazonensis*. Exp. Parasitol. 2016; 166:21-28.
  19. Dugasani S, Pichika MR, Nadarajah VD, Balijepalli MK, Tandra S, Korlakunta JN, Comparative antioxidant and anti-inflammatory effects of [6]-gingerol, [8]-gingerol, [10]-gingerol and [6]-shogaol. J Ethnopharmacol. 2010; 127:515-520.
  20. Duke JA, Ayensu ES. Medicinal Plants of China. Medicinal Plants of the World. Algonac, MI: Reference Publications, Inc. 1985; 1: 362.
  21. E Langner, S Greifenberg, J Gruenwald, Ginger: history and use,” Advances in Therapy, 1998; 15(1)25-44.
  22. El-Abhar HS, Hammad LN, Gawad HS. Modulating effect of ginger extract on rats with ulcerative colitis. J Ethnopharmacol. 2008; 118(3):367-172.
  23. El-Sharaky AS, Newairy AA, Kamel MA. Protective effect of ginger extract against bromobenzene-induced hepatotoxicity in male rats. Food Chem. Toxicol. 2009; 47(7):1584-1590.
  24. Ernst E, Pittler MH. Efficacy of ginger for nausea and vomiting: A systematic review of randomized clinical trials. Br. J Anaesth. 2000; 84(3):367-371.
  25. Ernst E, Pittler MH. Randomized controlled trial of ginger to treat nausea and vomiting in pregnancy. Obstet. Gynecol. 2004; 103(4):639-645.
  26. Flavonoids with gastroprotective activity. Molecules, 14: 979-1012.
  27. Kim JK, Kim Y, Na KM, Surh YJ, Kim TY. [6]-gingerol prevents UVB-induced ROS production and COX-2 expression *in vitro* and *in vivo*. Free Radic Res. 2007; 41(5):603-614.
  28. Frisch C, Hasenohrl RU, Mattern CM, Hacker R, Huston JP. Blockade of lithium chloride-induced conditioned place aversion as a test for antiemetic agents: comparison of metoclopramide with combined extracts of *Zingiber officinale* and Ginkgo biloba. Pharmacol. Biochem. Behav. 1995; 52:321-327.
  29. Ginger (*Zingiber officinale* Rosc.), a dietary supplement, protects mice against radiation-induced lethality: Mechanism of action. Cancer Biother Radiopharm. 19(4):422-435.
  30. Gong QM, Wang SL, Gan C. A clinical study on the treatment of acute upper digestive tract hemorrhage with wen-she decoction. Chung Hsi I Chieh Ho Tsa Chih. 1989; 9:272-273, 260.
  31. Grzanna R, Lindmark L, Frondoza CG. Ginger – An herbal Medical Product with Broad Anti- Inflammatory Action. J Med. Food. 2005; 8:125-132.
  32. Gull I, Saeed M, Shaukat H, Aslam SM, Samra ZQ, Athar AM. Inhibitory effect of Allium sativum and *Zingiber officinale* extracts on clinically important drug resistant pathogenic bacteria. Ann. Clin. Microbiol. Antimicrob. 2012; 11:8.
  33. Ha SK, Moon E, Ju MS, Kim DH, Ryu JH, Oh MS, Kim SY. 6- Shogaol, a ginger product, modulates neuroinflammation: a new approach to neuroprotection. Neuropharmacology. 2012; 63(2):211-23.
  34. Halvorsen BL, Holte K, Myhrstad MC, Barikmo I, Hvattum E, Remberg SF, *et al.* A systematic screening of total antioxidants in dietary plants. J Nutr. 2002; 132(3):461-471.  
[https://en.wikipedia.org/wiki/Ginger#cite\\_ref-fao\\_19-0](https://en.wikipedia.org/wiki/Ginger#cite_ref-fao_19-0)
  35. Jagetia GC, Baliga MS, Venkatesh P, Ulloor JN. Influence of ginger rhizome (*Zingiber officinale* Rosc.) on survival, glutathione and lipid peroxidation in mice after whole-body exposure to gamma radiation. Radiat Res. 2003; 160(5):584-592.
  36. Jung HW, Yoon CH, Park KM, Han HS, Park YK, Hexane fraction of *Zingiberis rhizoma* Crudus extract inhibits the production of nitric oxide and proinflammatory cytokines in LPS-stimulated BV2

- microglial cells via the NF kappaB pathway. Food Chem. Toxicol. 2009; 47:1190-197.
37. Kabuto H, Nishizawa M, Tada M, Higashio C, Shishibori T, Kohno M. Zingerone [4-(4-hydroxy-3-methoxyphenyl)-2-butanone] sprevents 6-hydroxydopamine-induced dopamine depression in mouse striatum and increases superoxide scavenging activity in serum. Neurochem. Res. 2005; 30:325-232.
  38. Kamtchouing P, Mbongue FGY, Dimo T, Jasta HB. Evaluation of androgenic activity of *Zingiber officinale* Andpentadiplandra Brazzearea In Male. Asian J Androl. 2000; 4:299-301.
  39. Kapil U, Sood Ak, Gaur DR. Maternal Beliefs Regarding Diet during Common Childhood Illnesses. Indian Pediatr. 1990; 27:595-599.
  40. Kelly S, Guilherme E, Meri E, Anderson L, Alba R, Clélia A, José M, Leônia M, 2009.
  41. Kim S, Kundu J, Shin Y, Park J, Cho M, Kim T. Gingerol inhibits COX-2 expression by blocking the activation of p38 MAP kinase and NF- B in phorbol ester stimulated mouse skin. Oncogene. 2005; 24:2558-2567.
  42. Koo KL, Ammit AJ, Tran VH, Duke CC, Roufogalis BD. Gingerols and related analogues inhibit arachidonic acid-induced human platelet serotonin release and aggregation. Thromb Res. 2001; 103(5):387-397.
  43. Krim M, Messaadia A, Maida I, Ouachari O, Saka S. Protective effect of ginger against toxicity induced by chromate in rats. Ann. Biol. Clin. (Paris). 2013; 71(2):165-173.
  44. Kubra IR, Murthy PS, Rao LJ. *In vitro* antifungal activity of dehydrozingerone and its fungitoxic properties. J Food Sci. 2013; 78(1):64- 69.
  45. Kumar A, Goyal R, Kumar S, Jain S, Jain N, Kumar P. Estrogenic and Anti-Alzheimer's studies of *Zingiber officinalis* as well as Amomum subulatum Roxb: the success story of dry techniques. Med. Chem. Res. 2015; 24(3):1089-1097.
  46. Kyung KS, Gon JH, Geun KY, Sup JJ, Suk WJ, Ho KJ. 6- Shogaol, a natural product, reduces cell death and restores motor function in rat spinal cord injury. Eur. J Neurosci. 2006; 24(4):1042-1052.
  47. Langner E, Greifenberg S, Gruenwald, J Ginger History, and Use. Adv. Ther. 15:25-44
  48. Lantz RC, Chen GJ, Sarihan M, Sóllyom AM, Jolad SD, Timmermann BN. The effect of extracts from ginger rhizome on inflammatory mediator production. Phytomedicine. 2007; 14:123-128.
  49. Li F, Nitteranon V, Tang X, Liang J, Zhang G, Parkin KL, *et al.* *In vitro* antioxidant and anti-inflammatory activities of 1-dehydro-[6]-gingerdione, 6-shogaol, 6-dehydroshogaol and hexahydrocurcumin. Food Chem. 2012; 135(2):332-337.
  50. Liao YR, Leu YL, Chan YY, Kuo PC, Wu TS. Anti-platelet aggregation and vasorelaxing effects of the constituents of the rhizomes of *Zingiber officinale*. Molecules. 2012; 17(8):8928-8937.
  51. Lumb AB. Effect of dried ginger on human platelet function. Thromb Haemost. 1994; 71:110-111.
  52. Mahmoud MF, Diaai AA, Ahmed F. Evaluation of the efficacy of ginger, Arabic gum, and Boswellia in acute and chronic renal failure. 116 J Pharmacognosy Phytother. Ren Fail. 2012; 34:73-82.
  53. Mallikarjuna K, Sahitya Chetan P, Sathyavelu Reddy K, Rajendra W. Ethanol toxicity: Rehabilitation of hepatic antioxidant defense system with dietary ginger. Fitoterapia. 2008; 79:174-178.
  54. Mehdizadeh M, Dabaghian F, Nejhad A, Fallah-Huseini H, Choopani S, Shekarriz N, *Zingiber Officinale* Alters, methylenedioxymethamphetamine-Induced Neurotoxicity in Rat Brain. Cell J Fall 2012; 14(3):177-184.
  55. Nasri H, Nematbakhsh M, Ghobadi S, Ansari R, Shahinfard N, Rafieian Kopaei M. Preventive and curative effects of ginger extract against histopathologic changes of gentamicin-induced tubular toxicity in rats. Int. J Prev. Med. 2013; 4(3):316-321.
  56. Newall CA, Anderson LA, Phillipson JD. Herbal Medicines: A Guide for Healthcare Professionals. London: Pharmaceutical Press, 1996.
  57. Ozgoli G, Goli M. Effects of ginger capsules on pregnancy, nausea, and vomiting. J Altern Complement Med. 2009; 15(3):243-246.
  58. Pecoraro A, Patel J, Guthrie T, Ndubisi B. Efficacy of ginger as an adjunctive anti-emetic in acute chemotherapy-induced nausea and vomiting. ASHP Midyear Clinical Meeting. 1998; 33:429.
  59. Portoni G, Chng LA, Karimi-Tabesh L, Koren G, Tan MP, Einarson A. Prospective comparative study of the safety and effectiveness of ginger for the treatment of nausea and vomiting in pregnancy. Am. J Obstet. Gynecol. 2003; 189(5):1374-1377.
  60. Pushpanathan T. The essential oil of *Zingiber officinalis* Linn (*Zingiberaceae*) as a mosquito larvicidal and repellent agent against the filarial vector *Culex quinquefasciatus* Say (Diptera: Culicidae). Parasitol. Res. 2008; 102(6):1289-1291.
  61. Qian DS, Liu ZS). Pharmacologic studies of antimotion sickness actions of ginger. Chung Kuo Chung Hsi I Chieh Ho Tsa Chih 1992; 12:95- 98.
  62. Qureshi S, Shah AH, Tariq M, Ageel AM. Studies on Herbal Aphrodisiacs Used In Arab System of Medicine. Am. J Chin. Med. 1989; 17:57-63.
  63. R Grzanna, L Lindmark, CG Frondoza, Ginger an herbal medicinal product with broad anti-inflammatory actions, Journal of Medicinal Food. 2005 8(2):125–132.
  64. Ramkissoon JS, Mahomoodally MF, Ahmed N, Subratty AH. Relationship between total phenolic content, antioxidant potential, and antiglycation abilities of common culinary herbs and spices. J Med. Food. 2012; 15(12):1116-1123.
  65. SV Nair, Ziaullah, HP Rupasinghe, Fatty acid esters of phloridzin induce apoptosis of human liver cancer cells through altered gene expression,” PLoS ONE, 2012; 19:9, Article ID e107149, 2014.
  66. Sabina EP, Pragasam SJ, Kumar S, Rasool M. 6-gingerol, an active ingredient of ginger, protects acetaminophen-induced hepatotoxicity in mice. Zhong Xi Yi Jie He Xue Bao. 2011; 9(11):1264-1269.
  67. Saraswat M. Antiglycating potential of *Zingiber officinalis* and delay of diabetic cataract in rats. Mol. Vision. 2010; 16(165-66):1525-1537.
  68. Shukla Y, Singh M. Cancer Preventive Proberties of Ginger: A Brief Review. Food Chem. Toxicol. 2007; 45(5):683-690.
  69. Srivastava KC. Aqueous extracts of onion, garlic and ginger inhibit platelet aggregation and alter arachidonic

- acid metabolism. *Biomed. Biochim. Acta.* 1984; 43(8-9):335-346.
70. Stewart JJ, Wood MJ, Wood CD, Mims ME. Effects of ginger on motion sickness susceptibility and gastric function. *Pharmacology.* 1991; 42:111-120.
71. Suekawa M, Ishige A, Yuasa K, Sudo K, Aburada M, Hosoya E. Pharmacological studies on ginger. I. Pharmacological actions of pungent constituents, (6)-gingerol and (6)-shogaol. *J Pharmacobiodyn.* 1984; 7(11):836-848.
72. Tjendraputra E, Tran VH, Liu-Brennan D, Roufogalis BD, Duke CC. Effect of ginger constituents and synthetic analogues on cyclooxygenase-2 enzyme in intact cells. *Bioorganic Chem.* 2001; 29:156-63.
73. Uz E, Karatas OF, Mete E, Bayrak R, Bayrak O, Atmaca AF, *et al.* The effect of dietary ginger (*Zingiber officinalis* Rosc.) on renal ischemia/reperfusion injury in rat kidneys. *Ren Fail.* 2009; 31(4):251-260.
74. VE Tyler, *The Therapeutic Use of Phytomedicinals, Pharmaceutical Products Press, New York, NY, USA, 1994.*
75. VS Govindarajan, Ginger-chemistry, technology, and quality evaluation. Part 1," *Critical reviews in food science and nutrition.* 1982; 17(1)1-96.
76. Verma SK, Singh J, Khamesra R, Bordia A. Effect of ginger on platelet aggregation in man. *Indian J Med. Res.* 1993; 98:240-242.
77. Vutyavanich T, Kraissarin T, Ruangsri RA. Ginger for nausea and vomiting in pregnancy: randomized, double-masked, placebo controlled trial. *Obstet. Gynecol.* 2001; 97:577-82.
78. Waggas AM. Neuroprotective evaluation of extract of ginger (*Zingiber officinale*) root in monosodium glutamate-induced toxicity in different brain areas male albino rats. *Pak. J Biol. Sci.* 2009; 12(3):201-212.
79. Wang CC, Chen LG, Lee LT, Yang LL. Effects of 6-gingerol, an antioxidant from ginger, on inducing apoptosis in human leukemic HL-60 cells. *In Vivo (Athens Greece).* 2003; 17(6): 641-645.
80. Y Shukla, M Singh, Cancer preventive properties of ginger: a brief review, *Food and Chemical Toxicology.* 2007; 45(5):683-690.
81. Yamahara J, Huang QR, Li YH, Xu L, Fujimura H. Gastrointestinal motility enhancing effect of ginger and its active constituents. *Chem. Pharm. Bull. (Tokyo)* 1990; 38:430-431.
82. Yamahara J, Miki K, Chisaka T. Cholagogic effect of ginger and its active constituents. *J Ethnopharmacol.* 1985; 13:217-225.
83. Yamahara J, Rong HQ, Iwamoto M, Kobayashi G, Matsuda H, Fujimura H. Active components of ginger exhibiting anti-serotonergic action. *Phytother. Res.* 1989; 3:70-71.
84. Yang R, Chang CS. Plants used for pest control in China: a literature review. *Econ. Bot.* 1988; 42(3):376.
85. Young HY, Liao JC, Chang YS, Luo YL, Lu MC, Peng WH. Synergistic effect of ginger and nifedipine on human platelet aggregation: A study in hypertensive patients and normal volunteers. *Am. J Chin. Med.* 2006; 34(4):545-551.