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# The study of the beneficial effects of ginger on human health: A critical review

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#### Abstract

Ginger appears to have been used as a spice and a medicine from early times by the Chinese and the Indians. It has been studied for its antibacterian, antifungal, pain-relieving anti-ulcer, antitumor and other properties). The rhizome of Ginger has long been used in Ayurvedic and traditional Chinese medicine to treat a wide range of ailments including gastrointestinal disorders, mainly nausea and vomiting associated with motion sickness. It has been recommended by herbalists for use as a carminative, diaphoretic, expectorant and astringent. Like many medicinal herbs, much of the information has been handed down by word of mouth with little controlled scientific evidence to support the numerous claims. However, in the last few years, more organized scientific investigation have focused on the mechanism and targets of ginger and its various components. Ginger (Z. Officinal's Roscoc, Zingibera ceae) is not only widely used as a dietary condiment but it has also been extensively utilized as a traditional medicine. Ginger's anti inflammatory properties help relieve pain and reduce inflammation associated with arthritis rheumatism and muscle spasm. Ginger also has been found to increase gastric-juice secreation and the production of hypochloride. It mean that food is digested more quickly, creating an unfriendly environment for bacteria.

Keywords: Gastro intestinal, spice, medicine, ailments, inflammatory

## Introductions

Ginger root was first cultivated in Asia and has been used as a medicinal herb for at least 2000 years ago [1] Ginger is a natural dietary component which has antioxidant and anticarcinogenic properties [2] Fluid extracts of ginger have been used since the 1500s for the treatment of GI distress. In China, ginger root and stem are used as pesticides against aphids and fungal spores [3] The taste of Ginger being aromatic and pleasantry pungent, it finds wide employment as a spice in the preparation of dishes of a most diverce character varying from curries to ginger bread. By virtue of its action as a carminative and stimulant to the gastrointestinal tract, ginger plays a very useful part in pharmacy. Ginger Oleoresing is obtained by solvent extraction of dried ginger and is prepared both in certain industrialized western countries and in some of the spice producing countries, most notably in Australia. The ginger oil prepared from fresh ginger rhizomes was determined by gas chromatography (GC) and GC-MS techniques. The main sesquitepene hydrocarbons identified were  $\alpha$ -zingiberence (27.30%)  $\alpha$ -curcumene (8-9%),  $\beta$ -sesqui phellandrene (4.8%) nd birsabolene (3.2%) [4].

#### Botany

**Medicinal species:** *Zingiber officinale* common names: Ginger, African ginger, Black ginger, Cochin ginger, Ganjiang, Gegibre, Ingwer, Jamaican ginger.

**Botanical family:** Zingiberaceae Ginger is closely related to two other cooking spices, turmeric and cardamons.

**Plant description:** Ginger is a herbaceous perennial leafy shoots, which attain a height of about 1 to 3 ft. After the flowers have disappeared and the stems have withered, ginger is ripe for collection. The rhizomes is aromatic and the source of the dried powered spice.

# General compostion of the ginger rhizome

Ginger contains from 0.25 to 3 percent of a volatile oil of light yellow colours having a characteristic odour. Jamaican variety yields about 1 percent, African from 2 to 3 percent and the Indian about 3.5 percent [5].

Corresponding Author: Bijoy Kumar Dey Research Scholar, Seacom Skills University, Kendra Dangal, Bolepur, Birbhum, West Bengal, India A typical analysis of a market sample of green ginger gave the following values moisture, 80.9; protein, 2.3; fat, 0.9; carbohydrates, 12.3; fibre 2.4; and minerals, 1.2. (as percentages). The principal minerals and vitamins in mg/100g are ca, 20; p, 60; and fe, 2.6; the vitamins, thiamine, 0.06; riboflavin, 0.03; niacin, 0.6 'and ascorbic acid, 6.0. About 18.6% protein remains unextracted, the

extracted proteins contin 35.6% albumin, 16.9%; globulin, 11.0%; prolamine and 17.9% glutelin, on total proteins <sup>[6]</sup>. Commercial dried gingeres have been reported to provide olecoresins in yields of 3.5 - 10% and to contain 15-30% of volatile oil (Govindarajan 1982) Table 1.

Gives the composition of ginger, spent singer and by products in commercials ginger samples.

Table 1: Composition of ginger, spent ginger, and by-products (commercial samples).

					Ash				Protein				
	Moisture	NVEE	VEE	Fibre	Total	Sand	Lime	Crude starch	Nx6.25)	NVEE	VEE	Alcohol extract	Cold water extract
Jamaican													
Natural	11.20	3.91	1.79	3.72	4.17	0.22	0.28	57.59	7.85	7.30	3.23	4.95	15.54
Limed	10.56	3.12	1.27	2.37	8.31	0.02	2.72	57.31	9.34			1	1
Cochin													
Rough	10.43	3.70	2.09	3.62	3.86	0.10	0.48	59.08	8.15	6.68	7.03	6.32	14.30
Scraped, limed	9.97	2.95	1.49	2.60	5.38	0.08	1.29	62.42	7.50				
Callcut										6.42	4.62	7.64	13.08
African	9.97	5.35	2.73	4.66	4.00	0.11	0.25	56.74	7.92	8.49	7.17	6.36	12.62
Japanese	10.30	3.94	0.96	2.73	6.19	0.71	1.666	0.55	5.40	7.01	7.39	8.37	14.40
Washes													
Scraggy	4.99	9.55	6.05	13.18	8.05	0.89	0.61	31.38	7.00				
Cuttings	3.19	2.76	7.06	8.69	9.20	1.81	1.06	40.23	8.69				
Residue (ginger after manufacture)	10.61	3.86	1.61	5.17	2.12	0.18		59.86	6.94				
Residue (extract)	8.02	0.54	0.13		5.05	1.50							

**Note:** NVEE-non-volatile other extract; VEE-volatile other extract.

# Medicinal and pharmacological properties Anticancer properties

Ginger, a natural dietary component, has been known to have antioxidant and anti carcinogenic properties. A study conducted by Abdullah, provided evidence that ginger acts as a potent growth inhibitory compound in human colon adenocarcinoma cells and the study supports the possibility of chemopreventive potential of ginger in colon cancer cells. The cytotoxic effect could be as a result of the active component. Azoxy methane induced intestinal carcinogenesis in rats was significantly suppressed by dietary administration of gingerol [7].

It is evidence that the anit tumour effects on colon cancer cells were exercerted by ginger by suppressing their growth, arresting the G0/G1-Phase, reducing DNA synthesis and inducing apoptosis [8].

#### **Anti-inflammatory effect**

Oral dried ginger or ginger extract (solvent 80% ethanol) or (6) shogaol reduced carrageenan induced paw swelling paw edema induced by compound 48/80 or serotonin was also significantly inhibited by intraperitoneal administration of a hydroalcoholic extract [9-10].

Ginger oil given orally for 26 days caused a significant suppression of paw and joint swellings in rates treated with Mycobacterium tuberculosis to induce severe arthritis in the knee and paw [11].

# Platelet aggregation

Ten studies were included, comprising eight clinical trails and two observational studies. Of the eight clinical trials, four reported that ginger reduced platelet aggregation, while the remaining four reported no effect. The two observational studies also reported mixed findings [12].

**Antioxidant effect:** Ginger and some specific constituents have demonstrated antioxidant effects in cell culture system.

Gingers extract inhibited hydroxyl radicals by 79.6% at 37 °C and 74.8% at 80 °C which showed a higher antioxidant activity than quercetin [13].

In rats ginger extract also ameliorated acetic acid – induced ulcerative colitis, likely due to antioxidant actions [14].

#### **Gastrointestinal effects**

The antiemetic effect of acetone and hydroalcoholic ginger extracts and their antiemetic constituents (shogaols, gingerols) experimented on four animals, one study shows reveral of the inhibitory effect of cisplatin on gastric emptying in rats by ginger acetone or ethanol extracts. 15-16 Another anticular compound in <sup>[6]</sup> ginger ulfonic acid with weaker pungency but more potent antiulcer activity than <sup>[6]</sup> gingerol and <sup>[6]</sup> Shogaol <sup>[17]</sup>.

## **Antimicrobial activity**

Ginger has strong antibacterial in addition to some antifungal properties. It has been reported *in vitro* studies to suppress the growth of a variety of common infections bacteria including *staphylococcus aureus and listena monocy to genes* [18]. It is found in animal studies ginger extracts exhibited the capacity to protect mice against infections [19].

# Osteoarthritic pain

Several animal studies show evidence that ginger and its active ingredients have the capacity to decrease symptoms of inflammation associated conditions such as arthritis [20, 21]. But further studies are necessary to prove the efficacy of ginger preparation in the treatment of osteoarthritic pain.

## Cardiovascular effect

Ginger reduced the blood pressure and decreased cardiac workload and thromboxane thus lowering the clotting ability of the blood <sup>[22]</sup>. One study reported that potential of different extracts (ethanolic, hexane and aqueous) of ginger

and the essential oil in 5-HT<sup>3</sup> receptor antagonistic effects <sup>[6]</sup> Gingerol showed maximum potential <sup>[23]</sup>.

## Nausea and vomiting during pregnancy

A flew studies demonstrated that 05. or 1g of ginger powder or an extract (solvent not stated) may be effective in treating nausea and/or emesis during pregnancy. Preliminary studies suggest that ginger may be effective for mind to moderate nausea and vomiting of pregnancy, when used at a recommended dose of 1-g dried ginger per day [24-25]. The quality and integrity of ginger preparations manufactured for use by women during pregnancy to be carefully determined.

# Regulation of blood glucose and lipid levels

Ginger is used to decrease cholesterol and triglyceride level. The ginger constituent zingiberont also produced lower blood glucose levels, body weight, in ovariectomized rats. (26) But one human study, in which ginger powder was administered in 4-g daily doses for 3 months to patients with cononary artery disease did not show any change in either blood glucose or blood lipid leves [27].

### **Numerious properties in treating Cinv**

Ginger demonstrates numerous properties that may be beneficial in treating chemotherapy induced nausea, retching and vomiting (CINV) including reversing the inhibitory effect of cisplatin on gastric emptying in rats [28-29]. There are multiple results the effect of Ginger or CINV. One study showing no effect [30], another with mixed results [31], and two other with positive out comes [32-33].

### **Toxicology**

There are no reports of severe toxicity in human from the ingestion of ginger root. A comprehensive review of human trials concluded that ginger at doses upto 2 g/d resulted in minimum toxicity for human. (34) Although progress in determining the active components and metabolites of ginger and understanding their pharmacokinetics has been made, more work is clearly needed.

## Discussion

Ginger is one of the most commonly consumed dietarcondiments in the world (Surh *et al.* 1999). Although ginger is generally considered to be safe (Kaul and Joshi 2001), the lack of a complete understanding of its mechanisms of action suggests caution in its the therapeutic use (Wilkin son 2000a). The medicinal chemical, and pharmacological properties of ginger have been extensively reviewed. (Surh, Lee, and Lee 1998: Ernst and Pittle) 2000; Afzal *et al.* 2001; and Roufogalis 2005; Eliopoulos 2007; White 2007; Nicoll and He nlin 2009). Ginger has been used for thousand of years for the treatment of hundreds of ailments from colds to cancer. The potential of ginger in the culinary, non culinary and medicinal field in based on the chemistry of volatile oil and non-volatile pungent principles.

The main compounds are zingiberene (29.5%) and sesquiphellandrene (18.4%). Ginger has proven anti-inflammatory, antioxidant and antiulcer principles [35].

# Conclusion

The review article is based on current and past research done on the powerful therapeutic effect of ginger for the various indications. Ginger is considered to be a safe herbal medicine, Herbal medicine is still the mainstay of about 75-80% of the world population, mainly in developing countries because of better culture acceptability, better compatibility with the human body and lesser side effects. Although the medicinal properties of ginger have been known for thousand of years. Therefore more extensive and well controlled human studies are sought before approving its use as a supplement for treatment of the diseases in order to give ginger a deserving place.

**Conflict of Interests:** The author declares that there is no conflict of interest.

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#### Reference

- Ody P. The complete medicinal herbal Dorling Kindersley; New York 1993.
- Manju V, Nalini N. Chemopre Ventive efficacy of ginger, a naturally occurring anti carcinogen during the initiation, Post. Initiatin stages of 1, 2-dimethylhydra zine induced colon cancer Clinica Chimica Aata 2005;358(1-2):60-67.
- 3. Yang R2, et al. Econ Bot 1988;42(3):376.
- Antonious GF, Kochhar TS. Zingiberence and curcumene in wild tomato. Journal of Environmental Science and Health Part B, Presticides, Food contaminants and Agricultural wastes 2003;38(4):489-500
- Chopra's Indigenous Drugs of India Academic Publishers, Kolkata. Third Reprint 2006. ISBN: 81-85086-80-X.
- 6. Govindarajan VS. Ginger-Chemistry, technology and quality evaluation Part I, CRC Critical Reviews 1982;17(1):1-96.
- 7. Abdulla S, Abidin SA2, Murad NA, Makpol S, Ngah WZW, Yusof YAM. Ginger extract (*Zingiber officinale*) triggers apoptosis and G0/G1 cells arrest in HCT 116 and HY 29 colon cancer cell lines. A fr J Bio Chem Res 2010;4:134-142.
- 8. Yoshimi N, Wang A, Morishi tay, Tanaka T, Sugie S, Kawaik, *et al.* Modifying effects of fungal and herb metabolites on azoxymethane induced intestinal carcinogenesis in rats. J Cancer Res 1992;83:1273-1278
- 9. Jana U, Chattopadhyay RN, Shaw BP. preliminary studies on anti-inflammatory activity of *Zingiber officinale* Ross; vitex negundo Linn, and Tinospora cordifolia (Willid) miers in albino rats. Indian J Pharmacol 1999;31:232-233.
- 10. Mascolo N, Jain SC, Capasso F. Ethnopharmacologic investigation of ginger. J. Ethno Pharmacol 1989;27;129-140.
- 11. Sharma JN, Srivastava KC, Gan EK. suppressive effects of eugenol and ginger oil on arthritic rats pharmacology 1994;49:314-318.
- 12. Mark W, MC Kavanagh D, MC Carthy AL, Bird R, Ried K, *et al.* The Effect of Ginger (*Zingiber officinale*) on Platelet Aggregation: A Systematic Literature

- Review 2015. https://doi.org/ID.1371/ journal. Pone 0141119
- 13. Stoilova I, Krastanv A, Stoyanova A, Deney P, Gargova S. Antioxidant activity of a ginger extract (*Zingiber officinale*) Food Chemistry 2007;102:764-770.
- 14. El-Abhar H, Hammad L, Gawad H. Modulating effect of ginger extract on rats with ulcerative colitis. J Ethropharma col. 2008;118:367-372.
- Yamahara J, Rong HQ, Naitoh Y, Kitani T, Fuirimura H. Gastrointestinal motility enhancing effect of ginger and its active constituents. Cham. Pharm. Bull (Tokyo) 1990;38:430-431.
- 16. Sharma SS, Gupta YK. Rever sal of Cisplatin induced delay in gastric emptying in rats by ginger (*Zingiber officinale*). J Ethnopharma col 1998;62:49-55.
- 17. Yoshikawa M, Yamaguchi S, Kunimi K, Matsuda H, Okuno Y, Yamahara J, *et al.* Stomachic principles in ginger. III. An antiulcer principle, 6-gingesul fonic acid and three monoacyl digalactosylglyceerols, ginger sulfonic acid, and three monoacyldigalactosylglycerols, ginger glycolipids A, B and C from Zingiberis rhizome originating in Taiwan, Chem. Pharm. Bull (Tokyo) 1994;42;1226-1230.
- 18. Norajit K, Laonakunjit N, Kerdchoenchuen O. Antibacterial effect of five Zingiberacecac essential oils. Molecules 2007;12:2047-2060.
- 19. Jageta G, Baliga M, Venkatesh P, Ulloor J. Influence of ginger rhizome (*Zingiber officinale* Rose) on survival, glutathione and lipid peroxidation in mice after whole body irradiation, Radiat Res 2003;160:584-592.
- Sharma J, Srivastav K, Gan E. suppressive effects of eugenol and ginger oil on arthritic rats. Pharmacology 1994;49:314-318.
- 21. Mustafa T, Srivastavak, Jensen K. Drug development report: Pharmacology of ginger, *Zingiber officinale*. J Drug Devel 1993;6:25-89.
- 22. Meena MR. M.Sc. Thesis: Indian Agricultural Research Institute, New Delhi 1992.
- Abdel Aziz H, Nahrstedt A, Prtereit F, Windeck T, Plouch M, Verspohl EJ. receptor blocking activity of arylakanes isolated from the rhizome of *Zingiber* officinale. Planta Medica 2005;71(7):609-916.
- 24. Sripramote M, Lekhyananda N. A randomized comparison of ginger and vitamin B<sub>6</sub> in the treatment of nausea and vomiting of pregnancy. J Med. Assoc. Thai 2003;86:846-853.
- 25. Chubrasik S, Pittler M, Roufogalis B. Zingiberris: a comprehensive review on the ginger effect and efficacy profiles, phytomedicine 2005;12:684-701.
- Han L, Mortomoto C, Zheng Y, Li W, Asami E, Okuda H, et al. Effects of Zingiber one on fat storage in ovarieetomized rats. Yakugaku Zasshi 2008:128:1195-1201.
- 27. Ginger. An overview of Health benefits; Keith Singletary, PhD., Nutrition Today 2010;45:4.
- 28. Gupta YK, Sharma M. Reversal of P. Yrogallol induced delay in gastric emptying in rats by ginger (*Zingiber officinale*) Methods Find Exp clin Pharma col. 2001;23(9):501-503. Doi: 10.1358/mf. 2001. 23.9.662137 (PubMed)
- 29. Sharma SS, Gupta YK. Reversal of cisplatin-induced delay in gastric emptying in rats by ginger (*Zingiber*

- *officinale*) J Ethnopharmacol 1998;62(1):49-55. doi: 10.1016/S0378 8741(98) 00053-1 (PubMed)
- 30. Manusirivithayas, Sripramole M, Tangjitgamols, *et al.* Antiemetic effect of ginger in gynecologic oncology patients receiving cisplatin, Int. Gyne. Col cancer 2004;14(6):1063-1069, doi:1111/J.1048-891X.2004;14603.X. (PubMed)
- 31. Pecoraro A, Patel J, Guthrie Ndubis B. Efficacy of ginger as an adjunctive anti-emetic in acute Chemotherapy-induced nausea and vomiting, ASHP Mid year clinical Meeting 1998, 429E.
- 32. Pace J. Oral ingestion of encapsulated ginger and reported self care action for the relief of chemotherapy-associated N & E. Dissertation Abstracts International 1987;47:3297-B.
- 33. Sontakk S, Thawani V, Naik MS. Ginger as an antiemetic in nausea and vomiting induced by chemotherapy: a randomized, cross over, double-blind study. Indian J Pharmacol 2003:35:32-36.
- 34. Chubrasik S, Pittler M. Addendum to a recent systematic review on ginger. Forsch Komplem en tarmed klass Naturheilkd 2005;12:168.
- 35. @CAB, Internation Chemistry of Spices (eds. V.A. Partha Sarathy, B. Champakam and T.J. Zachariah 2008.