

INTERNATIONAL JOURNAL OF PHARMACEUTICAL SCIENCES

[ISSN: 0975-4725; CODEN(USA): IJPS00] Journal Homepage: https://www.ijpsjournal.com



Research Article

Anticancer Potential of Plant and Natural Product

Patil Sunita*, Kalal Laxmi

N.B.S. Institute of Pharmacy, Ausa, dist. Latur, India.

ARTICLE INFO

ABSTRACT

Published: 16 Mar. 2025 Keywords: Cancer cell, Natural products, Plants, Carcinoma, medicinal plant, Alkaloids, Anticancer agents. DOI: 10.5281/zenodo.15034793

One of the main causes of death is cancer, and its incidence is steadily rising throughout the world. Numerous illnesses, including cancer, have been treated with traditional medicine, its active compounds, and certain natural products. There are a number of medications on the market to treat different kinds of cancer, but none of them have been proven to be completely safe and effective. The toxicity of the current medications is the main issue with cancer chemotherapy. Nonetheless, plants and medicines derived from them have shown promise in the management and treatment of cancer. There are numerous natural compounds, and some of them have been found to have strong anticancer properties. Nowadays, the majority of cancer medication research involves...

INTRODUCTION

Plants have been used to treat illnesses for as long as civilization [1], and traditional medicines continue to play a significant role in the routine treatment of various illnesses [2]. Due to historical, cultural, and other factors, folk medicine has gained popularity recently, particularly in underdeveloped nations with little healthcare. Serious to negative access consequences could result, nevertheless, if medicinal herbs are not scientifically evaluated to support their usage [3]. Biologically active materials are thought to be mostly found in plants. Eighty percent of rural residents use medicinal their primary healthcare system, plants as

according to recent figures [4]. In both industrialized and developing nations, cancer is one of the most common illnesses. It is a class of diseases where body cells divide ab normally and uncontrollably, which can be fatal. Normal cells are typically invaded and destroye d by cancer cells. Therefore, we can regulate the development of cancer if we limit or stop this cell division.Despite spending billions of dollars on r esearch, we still do not fully understand what can cer is.Millions of people receive a cancer diagnos is each year, which results in their death. Approx imately 3500 million people wordwide lose their l ives to cancer.A number of chemopreventive me

*Corresponding Author: Patil Sunita

Address: N.B.S. Institute of Pharmacy, Ausa, dist. Latur, India.

Email : sunitaupatil02@gmail.com

Relevant conflicts of interest/financial disclosures: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

icines are used to treat cancer, but their toxicity li mits their applications.

What Causes Cancer?

Mutations in DNA, which tell cells how to divid e and proliferate, may be the first cause of cancer. Most DNA mutations can be fixed by normal cells, but those that cannot be fixed and cause t he cell to proliferate can turn into cancer [5]. The Traditional Medicine History of Medicinal plants have been used to treat various d iseases for thousands of years and in many parts of the world [6]. Humans used medicinal plants b efore 60,000 years ago, according to fossil recor ds [7]. Today, plants remain the main source of m edicine in rural areas of developing nations [8], a nd it is estimated that 80% of people in these cou ntries still use medicinal plants for their health care [9].

Ayurvedic Concept of Cancer

Both Charaka and Sushruta Samhita [10,11] referred to the cancer's counterpart as "Granthi" and "Arbuda" [12]. Depending on the dosage, "Granthi" and "Arbuda " can either be inflammatory or non-inflammatory [13]. Ayurveda defines health as the balanced coordination of the body, mind, and consciousness of the three doshas "Vatta, Pitta, and Kapha" that cause disease [14]. The loss of mutual coordination among the three main body humors causes a morbid condition, which is why tridoshicarbudas are typically malignant [15].

Role Of Plants as Medicinal and Anticancer Agent Since the beginning of human civilization, plants have been used as a source of food, shelter, and medicine. It is estimated that 80–85% of the world's population relies on traditional medicines for their primary health care needs, and it is assumed that a large portion of traditional therapy involves the use of plant extracts or their active principles [16,17,18]. While many recent studies have been conducted to advance the treatment and control of cancer progression, there is still much work to be done and room for improvement. The main drawbacks of synthetic drugs are the side effects they cause, but plant-derived products are much safer and have much lower toxicity than chemical synthetic drugs. When the cytotoxic podophyllotoxins were isolat ed and the vinca alkaloids (vinblastin and vincrist ine) were discovered and developed in the 1950s, the hunt for anticancer agents from plant sources began [19].

The Use of Plants in Cancer Therapy

cancer patients is constantly increasig Approxima tely 2to3 percent of deaths worldwide are caus ed by various forms of cancer [20], and the avai lable treatment options include surgery, chemothe rapy, and radiation [21]. The rising costs of traditi onal treatments, such as chemotherapy and radiati on, and the lack of effective medications to treat s olid tumors have led to a greater reliance on tradit ional medicine, which is based on the use of medi cinal plants [22]. These plants have an almost li mitless capacity to produce substances that draw r esearchers looking for new and innovative chemo therapeutics [23]. Despite the fact that some plant are used in cancer therapy, plant products derived anticancer agents only make up onefourt h of the available treatmentoptions. Nine plantder ived compoundsVinblastin, Vincristin, navelbine, Etoposide, Teniposide, Taxol, Taxotere, topoteca n, and irinote can have been approved for use in cancer therapy in the United States since 1961 [24]. Plant-Derived Anticancer Agents in Clinical Use.







1.Vinca Alkaloids

Vinca alkaloids are a significant family of antica ncer medications. Vinblastine (VLB) and vincristine (VCR), two significant nat urally occurring chemicals derived from the Mad agascar periwinkle, Catharanthus roseus G. Don. (Apocynaceae), were the first agents to be used in therapeutic settings. These medications were fou nd while looking for oral hypoglycemics.Researc hers found that plant extracts triggered bone marr ow depression in rats and dramatically decreas ed white blood cell numbers, albeit they we re unable to substantiate this activity.Mice wit h transplantable lymphocytic leukemia also liv e longer thanks to plant extract. Vincristine and v inblastine, two active alkaloids, were isolated afte r additional extraction and fractionation. Although the plant was native to Madagascar, samples use d to identify vincristine and 1479inblastine were gathered in Jamaica and the Philippines. The semi synthetic analogues of vinca alkaloids vindesine (VDS) and vinorelbine (VRLB) have been dev eloped recently. These are mostly used to treat a ra nge of malignancies, either by themselves or in c onjunction with other chemotherapy medications. VLB is used to treat Kaposi's sarcoma, lymphom

as, leukemias, breast cancer, testicular cancer, a nd lung cancer.Additionally, VCR demonstrated effectiveness against leukemia, specifically juven ile acute lymphocytic leukemia [25,26].When co mpared to other vinca alkaloids, vinflunine, a bifl uorinated derivative of vinorelbine, has better anti cancer activity.Both vinflunine and vinorelbine sh ow decreased toxicity in animal models, and this new vinca alkaloid is presently undergoing phase II clinical studies [27, 28].





2.Allium Sativum

In India, Allium sativum, commonly known as ga rlic or ginger, is used to treat a wide range of illne sses.Allicin is the main ingredient in raw garlic, a



nd allicin rearranges to produce ajoene. Its cytotox in action has been evaluated using human primar y fibroblasts, a tumorigenic lymphoid cell line obt ained from a burkit lymphoma, and a permanent nontumorigenic cell line produced from baby la mster kidney cells. The range of the cytotoxic acti vity was 250 ug/m l29.In a number of animal mo dels, some organo sulfur compounds found in gar lic, such as sallylesteine, have been shown to tra ck the development of chemically produced and t ransplantable tumors [30].Garlic may have antica ncer effects, according to the National Cancer Ins titute, which is part of the NIH.Garlic (250 mg/kg , p.o., three times a week) significantly inhibited t he development of tongue carcinogenesis caused by nitroquinoline1oxide in male wistar rats, as evi denced by the absence of carcinomas during the i nitiation phase and their decreased occurrence dur ing the postinitiation phase [31].Garlic's sulphydr yl components have the power to prevent the dev elopment of chemicals that cause cancer.Increase d garlic consumption has been linked in a number of population studies to a lower risk of pancreati c, esophageal, colon, stomach, and breast cancers [32].



Fig.3 [53]

3. Silybum Marianum

The milk thistle plant, Silybum marianum, yielde d the flavonoid compound silymarin.Silymarin w as investigated for its ability to prevent UV induced skin cancer in mice.Silymarin treatment demonstrated a strong anticancer effect, lowering the likelihood of tumor incidence to 60%, tumor multiplicity to 78%, and tumor volume per mouse to 90% [33].Cell cycle arrest at the G1/Sphase, i nduction of cyclindependent kinase inhibitors, do wnregulation of antiapoptotic gene 1480product, i nhibition of cellsurvival kinases, and inhibition of inflammatory transcription factors are some of th e ways that silymarin suppresses the growth of tu mor cells.Additionally, it was discovered that sily marin inhibited gene products linked to tumor cel l proliferation, invasion, angiogenesis, and metast asis [34]. One of the main components of Silybum marianum fruits, silibinin (flavanolignan), is also helpful in treating breast cancer in humans [35].



Fig.4 [54] 4. Podophyllotoxin Derivative

Podophyllum peltatum Linnaeus and P. emodii W allich are two species of the Podophyllaceae fami ly that have a long history of therapeutic use, incl uding the treatment of warts and skin cancers.Nat ive American people have utilized podophyllum p eltatum to treat "cancer."The discovery in the 194 Os that a topical application of an alcohol extract of the dried roots, known as podophyllin, treats v enereal warts sparked interest.Although podophyl lotoxins were the main cytotoxic medicinal ingre dients and were isolated for the first time in 1880, it was not until the 1950s that spectroscopic tech niques advanced enough to reveal its true compos ition.During this time, other closely related podop



hyllotoxins, such as lignans, were also isolated an d put into clinical trials; however, their ineffectiv eness and intolerable toxicity led to their withdra wal.In the 1960s and 1970s, Sandoz Laboratories in Switzerland conducted extensive research that resulted in the development of etoposide and teni poside as clinical agents that are currently being u sed to treat lymphomas as well as bronchial and t esticular cancers. As of July 2004, the NCI had re corded 2069 anticancer clinical trials in progress, of which more than 150 were drug combinations t hat included etoposide against various cancers [3 6,37].





5.Taxanes

The creation of a class of molecules known as tax anes represents a more recent development in the creation of chemotherapeutic agents derived from plants.The bark of Taxus brevifolia Nutt. (Taxace ae) is where paclitaxel, commonly known as taxol , was initially isolated.They entered the market in the 1990s after their structure was initially discov ered in 1971.While the leaves of T. baccata are us ed in Ayurveda, with one use in treating "cancer," several Native American tribes have documented using various parts of T. brevifolia and other Tax us species (e.g., T. canadensis Marshall, T. baccat a L.) to treat certain noncancerous conditions.A si gnificant, renewable natural source of this signifi cant class of medications is the readily semisynthetic conversion of the relatively abundant ba ccatins to paclitaxel and active paclitaxel analogs, like docetaxel (Taxotere). Paclitaxel is biosynthe sised and found in the leaves of several Taxus spe cies.In addition to being effective against Kaposi sarcoma, paclitaxel is used to treat a wide range o f malignancies, such as breast, ovarian, and nonsmallcell lung cancer.Additionally, it has garnere d interest because of its potential for treating rheu matoid arthritis, multiple sclerosis, and psoriasis [38].It was discovered that docetaxel, a semisynth etic derivative of paclitaxel, was more efficient. Patients who are resistant to paclitaxel may be tre ated with docetaxel.For patients with ovarian, bre ast, or metastatic cancer, docetaxel and paclitaxel are utilized as first- and secondline treatments, res pectively [39].





6.Curcumin

Curcumin (diferuloylmethane), a polyphenolic co mpound that was isolated from the Indian plant sp ecies Curcuma longa (commonly called turmeric) , is now being used as a potential anticancer comp ound. While turmeric is primarily promoted as an antiinflammatory herbal remedy, some scientists think that the antioxidant curcumin in turmeric m ay prevent or slow the growth of many cancers, in cluding tumors of the esophagus, stomach, and in



testine, breast cancer, and skin cancer in experime ntal animals. Curcumin is involved in modulating the cell cycle pathway and induced apoptosis of various cancer cells, though the precise mechanis m of action has not yet been thoroughly studied. Phase I/II trials are currently being conducted to d etermine how curcumin affects colorectal cancer, multiple myeloma, and pancreatic cancer [40,41]. Known for its antiinflammatory, antiviral, antibac terial, antifungal, and anticancer properties, curcu min is used in Chinese medicine and as the yello w coloring ingredient in curry, a traditional India n dish. It may also help treat diabetes, allergies, ar thritis, and Alzheimer's disease.discussed curcumi n and proposed that agents that target multiple ge nes, like curcumin, are necessary for cancer preve ntion and treatment because the majority of cance rs are caused by dysregulation of up to 500 differ ent genes [42].

7.Camellia Sinensis (Green Tea)

Green tea and occasionally black tea include poly phenols that aid in the death of malignant cells an d the prevention of their spread.In green tea, epig allocatechin3gallate (EGCG) is the most prevalen t polyphenol.Patients with chronic lymphocytic le ukemia (CLL), a kind of blood cancer, had fewer leukemia cells because to green tea's epigalloacat echin gallate (EGCG). According to certain epide miological research, EGCG can prevent human c olon and oral cancer cells from invading and migr ating.Reduced MMP2, MMP9, and uPA synthesis may be partially responsible for the effects of EG CG [43].Additionally, it was found that EGCG in hibited the growth of cancer cell lines such as hep atocellular carcinoma by inducing cell cycle arres t [44], inhibited the growth of cancer cells in ovar ian carcinoma cell lines HEY and OVCA, and inh ibited the growth of cancer cells in human colon a nd rectal cancer cell lines HT29 and HCA7 [45,4 6,47,48]. In addition, studies found that women w ho drank powdered green tea had a lower risk of bladder cancer, and that men who drank the most

green tea had a 37 percent lower risk of pancreati c cancer. A large Chinese clinical study found tha t the risk of prostate cancer decreased as the frequ ency and quantity of green tea consumption incre ased, but that green tea could decrease the likelih ood of breast cancer recurrence but not prevent or improve breast cancer [50].

CONCLUSION

This review article offers details on natural medic ines and herbs that may be able to treat or slow th e spread of cancer.

A number of clean, efficient agents for the treatm ent of various cancers have been found in natural materials and their derivatives. It is well known th at medicinal plants have strong anticancer propert ies. When discussing anticancer treatments, they are frequently mentioned as a major source of syn thetic or herbal ingredients. The use of natural me dicines derived from medicinal plants is crucial in the fight against cancer. In animal models of sarc oma, skin cancer, and leukemia, they have demon strated anticancer activity.

REFRENCES

- Fabricant, D. & Farnsworth, N. (2001). The value of plants used in traditional medicine for drug discovery. Environmental Health Perspectives, 109 (1), 69-75.
- Alviano, D. & Alviano, A. (2009). Plant extracts: search for new alternative to treat microbial diseases. Current Pharmaceutical Biotechnology, 10, 106-121.
- Souza, G., Hass, A., Poser, G., Schapoval, E., & Elisabetsky, E. (2004).
 Ethnopharmacological studies of antimicrobial remedies in the south of Brazil. Journal of Ethnopharmacology, 90 (1), 135-143.
- Sakarkar, D. & Deshmukh, V. (2011). Ethnopharmacological review of traditional medicinal plants for anticancer activity. International Journal of Pharma Tech Research, 3, 298 308.

- Krishnamurthi K. Screening of natural products for anticancer and antidiabetic properties. Health Administrator. XX (1&2): 69, (2000).
- 6. Palombo, E. (2009). Traditional medicinal plant extracts and natural products with activity against oral bacteria: potential aplication in the prevention and treatment of oral diseases. Evidence-based Complementary and Alternative doi:10.1093/ecam/nep067. Medicine,
- Fabricant, D. & Farnsworth, N. (2001). The value of plants used in traditional medicine for drug discovery. Environmental Health Perspectives, 109 (1), 69-75.
- Chitme, H., Chandra, R., & Kaushik, S.(2003). Studies on anti-diarrheal activity of Calotropis Gigantea R. Br. in eperimental animals. Journal of Pharmacy and Pharmaceutical sciences, 7, 70-75.
- 9. Kim, H. (2005). Do not put too much value on conventional medicines. Journal of Ethnopharmacology, 100, 37-39.
- 10. Charaka. Charaka Samhita. Chaukhamba Publications, Varanasi, India; 700 BC: 215.
- 11. Susruta. Susruta Samhita. Chaukhamba Publications, Varanasi, India ; 700 BC: 520.
- Misra B. Bhawa Prakash Nighantu. Chaukhamba Publications, Varanasi, India; 1600 AD: 29
- 13. Kapoor L. D. Handbook of ayurvedic medicinal plants. CRC Press, Florida; 1990.
- 14. Balachandran P, Govindarajan R., Cancer- an ayurvedic perspective. Pharmacology Research, 51. 19. 2005.
- 15. Singh R. H., An assessment of the ayurvedic concept of cancer and a new paradigm of anticancer treatment in Ayurveda. Journal of Alternative & Complementary Medicine, 8. 609. 2002.
- 16. Ignacimuthu S, Ayyanar M, Sivaraman S.K., Ethnobotanical investigations among tribes in

Madurai district of Tamil Nadu (India). Journal of Ethnobiology and Ethnomedicine, 2. 1. 2006.

- Elujoba A. A., Odeleye O. M., Ogunyemi C. M., Traditional medicine development for medical and dental primary health care delivery system in Africa. African Journal of Traditional, Complementary and Alternative Medicines, 2. 46. 2005.
- Tomlinson T. R., Akerele O., Medicinal plants: their role in health and biodiversity. University of Pennsylvania Press, Philadelphia, 1998.
- 19. Gordon M. C., David J., Plants as a source of anti-cancer agents. Journal of Ethnopharmacology, 100. 72. 2005.
- 20. Madhuri, S. and Pandey G. (2009).Some anticancer medicinal plants of foreign origin.Current science,96(6),779-782.
- 21. Tannock F.(1998).Conventional Cancer therapy:Promise broken or promisedelayed?. Lancet,352,9-16.
- 22. Wood-Sheldon,J,.,Balick,M.and Laird,S.(91997).Medicinal plants:Can utilization and conservation Coexist?.The New York Botanical Garden.USA.
- 23. Reed, J. and Pellecchia, M. (2005). Apoptosisbased therapies for hematologic malignancies, blood, 106, 408-418.
- Lee, K. (1999). Anticancer drug design based on plant-derived natural products. Journal of Biomedical Science, 6, 236-350.
- 25. Gueritte F., Fahy J., The vinca alkaloids. In Anticancer Agents from Natural Products, edited by Cragg GM, Kingston DGI, Newman DJ. Brunner-Routledge Psychology Press, Taylor & Francis Group, Boca Raton, Chapter 7. 23. 2005.
- 26. Cragg,G.M., Newman, D.J.,(2005). Plants as a source of anticancer agenta. J Ethnopharmacol.,100:72-79.



- Okouneva, T.,Hill, B.T., WILSON, L., Jordan, M.A., (2003). The effects of vinflunine,vinorelbine, and vinblastine on centromere dynamics. Mol cancer ther.,2:427-436.
- 28. Simoens, C., Lardon, F., Pauwels, B., De Pooter, C.M.J., Lambrechts, H.A.J., Pattyn,G.G.O., Breillout, F., Vermorken, J.B., (2008). Comparative study of the radiosensitising and cell cycle effect of vinflunine and vinorelbine, in vitro. BMC cancer.,8:65.
- 29. Scharfenberg K., Wagner R., Wagner K.G., The cytotoxic effect of ajoene, a natural product from garlic, investigated with different cell lines. Cancer Letters, 53(2-3). 103. 1990.
- Thomson M., Ali M., Garlic (Allium sativum): a review of its potential use as an anti-cancer agent. Current Cancer Drug Targets, 3(1). 67. 2003.
- Banasenthil S., Ramachandran C.R., Nagini S., Prevention of 4 nitroquinoline-1-oxide induced rat tongue carcinogenesis by garlic. Fitoterapia, 72. 524. 2001.
- Ranjani R, Ayya RM. Anticancer Properties of Allium sativum A Review, Asian Journal of Biochemical and Pharmaceutical Research. 2012; 3 (2): 190-196.
- 33. Agarwal R., Agarwal C., Ichikawa H., Singh R.P., Agarwal B.B., Anticancer potential of silymarin: from bench to bed side. Anticancer Research, 26(6B). 4457. 2006.
- 34. Kim S., Choi J. H., Lim H. I., Lee S. K., Kim W.W., Kim J. S., et al., Silibinin prevents TPA-induced MMP-9 expression and VEGF secretion by inactivation of the Raf/MEK/ERK pathway in MCF-7 human breast cancer cells, Phytomedicine, 16(6-7). 573. 2009.
- 35. Li Y. L., Gan G. P., Zhang H. Z., Wu H. Z., Li C. L., Huang Y. P., Liu Y. W., Liu J. W., A flavonoid glycoside isolated from Smilax china L. rhizome in vitro anticancer effects on

human cancer cell lines. Journal of Ethnopharmacology, 113(1). 115. 2007.

- 36. Lee K.H., Xiao Z., Podophyllotoxins and analogs. In Anticancer Agents from Natural Products, edited by Cragg GM, Kingston DGI, Newman DJ. Brunner-Routledge Psychology Press, Taylor & Francis Group, Boca Raton, Chapter 5. 71. 2005.
- 37. Shoeb, M., MacManus,S.M.,Jaspars, M.,TRevidadu, J., Nahar, L., Thoo-Lin, P.K., Sarker, S.D., (2006). Montamine, a unique dimeric indole alkaloid, from the seeds of Centaurea montana (Asteraceae), and its in vitro cytotoxin activity against the CaCo2 colon cancer cells.Tetrahedron.,62:11172-77
- 38. Kingston D. G. I., Taxol and its analogs. In Anticancer Agents from Natural Products, edited by Cragg GM, Kingston DGI, Newman DJ. Brunner-Routledge Psychology Press, Taylor & Francis Group, Boca Raton, Chapter 6. 89. 2005.
- Hait,W.N., Rubin, E., Alli, E.,Goodin, S., (2007). Tubulin targeting agents. Update on cancer therapeutics.,2:1-18.
- Sa, G., Das, T., Banerjee, S., Chakraborty, J., (2010). Curcumin:from exotic spice to modern anticancer drug. AI Ameen J Med Sci.,3:21-37.
- 41. Goel, A., Kunnumakkara, A.B., Aggarwal, B.B.,(2008). Curcumin as "Curecumin": from kitchen to clinic. Biochem. Pharmacol.,75:787-809.
- 42. Kunnumakkara AB, Anand P, Aggarwal BB. Curcumin inhibits proliferation, invasion, angiogenesis and metastasis of different cancers through interaction with multiple cell signalling proteins. Cancer Lett. 2008; 269: 199-225.
- 43. Ho Y. C., Yang S.F., Peng C. Y., Chou M. Y., Chang Y. C., Epigallocatechin-3-gallate inhibits the invasion of human oral cancer cells and decreases the productions of matrix

etalloproteinases and urokinase-plasminogen activator. Journal of Oral Pathol Medicine, 36. 588. 2007.

- 44. Nishikawa T., Nakajima T., Moriguchi M., Jo M., Sekoguchi S., Ishii M. et al., A green tea polyphenol, epigalocatechin-3-gallate, induces apoptosis of human hepatocellular carcinoma, possibly through inhibition of Bcl-2 family proteins. Journal of Hepatology, 44. 1074. 2006.
- 45. Spinella F., Rosano L., Decandia S., Di C.V., Albini A., Elia G., Natali P.G., Bagnato A., Antitumor effect of green tea polyphenol epigallocatechin-3-gallate in ovarian carcinoma cells: evidence for the endothelin-1 as a potential target. Exp Biol Med, 231. 1123. 2006 a.
- 46. Spinella F., Rosano L., Di C.V., Decandia S., Albini A., Nicotra M.R., Natali P.G., Bagnato A., Green tea polyphenol epigallocatechin-3gallate inhibits the endothelin axis and downstream signaling pathways in ovarian carcinoma. Mol Cancer Therapy, 5. 1483. 2006 b.
- 47. Hwang J. T., Ha J., Park I. J., Lee S. K., Baik H. W., Kim Y. M., Park O. J., Apoptotic effect of EGCG in HT-29 colon cancer cells via AMPK signal pathway. Cancer Letters, 247. 115. 2007.
- 48. Peng G., Dixon D.A., Muga S. J., Smith T.J., Wargovich M. J., Green tea polyphenol (-)epigallocatechin-3-gallate inhibits cyclooxygenase-2 expression in colon carcinogenesis. Molecular Carcino, 45. 309. 2006.
- 49. Hu Z., Yang Y., Ho P.C., Chan S. Y., Heng P. W., Chan E., Duan W., Koh H.L., Zhou S., Herb-drug interactions: a literature review. Drugs, 65. 1239. 2005.
- 50. Ishii T, Mori T, Tanaka T, Mizuno D, Yamaji R, Kumazawa S, Nakayama T, Akagawa M. Covalent modification of proteins by green tea

polyphenol (-)-epigallocatechin-3-gallate through autoxidation. Free Radic Biol Med. 2008; 45: 1384-1394.

- 51. https://images.app.goo.gl/or2z1kbMLNqiQDi q9
- 52. https://images.app.goo.gl/B6sUETGKWqEpy myV7
- 53. https://images.app.goo.gl/2XpR9v47vvgHfhe W9.

HOW TO CITE: Patil Sunita, Kalal Laxmi, Anticancer Potential of Plant and Natural Product, Int. J. of Pharm. Sci., 2025, Vol 3, Issue 3, 1477-1485. https://doi.org/10.5281/zenodo.15034793

