

Original Research

View Article Online



Received 15 January 2025

Revised 17 February 2025

Accepted 21 March 2025

Available Online 05 June 2025

Edited by Sabya Sachi Das

KEYWORDS:

Anti-cancer properties

Breast cancer

Phytochemicals

Sustainable Development Goal

Therapeutics

Molecular pathways

Natr Resour Human Health 2025; 5(3):339-353
<https://doi.org/10.53365/nrhh/203182>

eISSN: 2583-1194

Copyright © 2025 Visagaa Publishing House

Key Insights on Therapeutic Potential of Various Phytochemicals against Breast Cancer

Dipika Pal¹, Pallavi Singh^{2,*}, Madhulika Esther Prasad³, Vijay Jagdish Upadhye⁴

¹Department of Biotechnology, Graphic Era Deemed to be University, India

²Department of Biotechnology, Graphic Era Deemed to be University, 566/6, Bell Road, Clement Town, 248002, Dehradun, India

³Department of Biochemistry and Biotechnology, Sardar Bhagwan Singh University, Dehradun-248161, Uttarakhand, India

⁴Centre of Research for Development, Department of Microbiology, Parul Institute of Applied Sciences (PIAS), Parul University, Vadodara-391760, Gujarat, India

ABSTRACT: Breast cancer poses a global health challenge and requires innovative approaches for prevention and treatment. Phytochemicals have gained attention for their potential anti-cancer properties by offering promising therapeutic interventions with their many advantageous effects and activity in promoting apoptotic pathways and other molecular pathways involved in breast cancer. Research indicates that phytochemicals have many beneficial effects, including anti-inflammatory, antioxidant, and antiproliferative effects. Phytochemicals have the potential to affect crucial signaling pathways of breast cancer. Promiscuous phytochemicals such as curcumin, resveratrol, apigenin, and epigallocatechin gallate (EGCG), among others, have demonstrated notable anti-cancer effects in preclinical studies. This review delves deep into the therapeutic action and mechanism of various phytochemicals against breast cancer. It highlights the promiscuous role of multiple phytochemicals, their sources, and their target molecules in breast cancer along with shedding light on some of the limitations of these phytochemicals. It discusses the key role of phytochemicals like Apigenin and Silibinin in combating the progression of Breast cancer and also emerging as crucial avenues to counter chemotherapeutic resistance. Effective treatments and cures should be developed to reduce the burden of non-communicable diseases and ensure human well-being and a good quality of life in order to accomplish sustainable development goals.

1. INTRODUCTION

In 130 countries worldwide, it is found that cancer is the second leading cause of premature death (Bray et al., 2021). In India, the mortality rate of cancer is 68% of the yearly occurrence, which means that 30% of the patients will only survive 5 years or longer after the diagnosis. (Ramani et al., 2023). Noncommunicable diseases (NCDs), which include cancer, diabetes and cardiovascular disease were first acknowledged as a serious health issue by the World Health Organisation (WHO) in 1999. In 2012, NCDs contributed to over 38 million annual deaths, or 68% of the total deaths globally. (Morgan et al., 2018). The United Nations' Sustainable Development Goal (SDG) 3 (Good health and well-being) promotes the well-being and ensures healthy lives of people of all ages, which is one of the top goals for the

entire). Furthermore, the World Health Assembly elevated cancer care on the global health agenda in 2017 by passing resolution 70.12, which allowed WHO to prioritize cancer prevention and control programs. (Murthy et al., 2024). The five most common dietary and behavioral hazards contribute to almost one-third of deaths caused by cancer one of which is low fruit and vegetable intake. Most of the prevalent cancers, that include breast cancer, have been cured when discovered early and have shown effective results (Pereira et al., 2021). One of the targets of SDG related to cancer is to achieve universal health improvement which will protect people financially, provide the opportunity for better healthcare services, and ensure that everyone has access to medications and vaccinations at a cheap, safe, and effective practices. (Frieden et al., 2020). Reports from the World Health Organization in 2020 depicted that 2.3 million women were detected with

* Corresponding author.

E-mail address: pallavisinh.22@gmail.com (Pallavi Singh)

This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

breast cancer internationally which resulted in 6,85,000 deaths. Data from Scientific investigation reflects that the average age of women who have been diagnosed with Breast cancer is 61 years. Breast cancer is extremely fatal with no effective therapeutics developed as of now. (Arnold et al., 2022). Breast cancer stands as a prevalent human malignancy and comes in second place after lung cancer (Sohel et al., 2023). Modern techniques of breast cancer treatment encompass a diverse range of methods, including radiotherapy, surgical procedures, chemotherapy, antiestrogen therapy, aromatase inhibitors, anti-angiogenesis, pharmaceuticals, and the utilization of anthracyclines (Cazzaniga & Bonanni, 2012). Although, these therapies have their negative effects and are very costly, and also cause negative health implications. To avoid this, natural products emerge as an effective alternative therapeutic, marked with fewer associated side effects (Cazzaniga & Bonanni, 2012). Plants naturally produce thousands of active natural compounds of low molecular weight, and have a structural diversity known as phytochemicals. They protect the plants from various kinds of bacterial, fungal, and viral infections, environmental stress, and UV exposure and produce an immune response against them. These can also regulate various kinds of molecular mechanisms that can prevent different kinds of diseases (Wink, 2015). Up to date, there are more than 4,500 phytochemicals that are on the record. They are categorized according to the protective roles they play. Common sources of phytochemicals are carrots, turmeric, green tea, tomatoes, grapes, cherries, garlic onion, whole wheat, cabbage, raspberries, beans and soy foods, etc. (Mazurakova et al., 2022).

2. CLASSIFICATION OF PHYTOCHEMICALS

Broadly, Phytochemicals are divided into primary and secondary metabolites based on their role in plant metabolism (Younas et al., 2018). The phytochemicals are further classified into sub-classes according to their structures (Hussein & El-Anssary, 2019). Classification of various phytochemicals has been given in Figure 1.

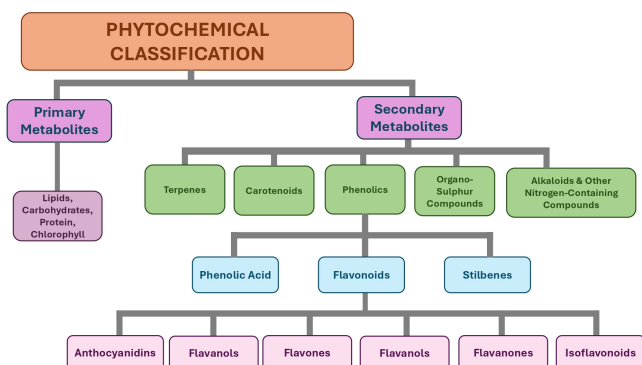


Figure 1. Classification of Phytochemicals.

3. DIFFERENT PHYTOCHEMICALS EFFECTIVE AGAINST BREAST CANCER

3.1. Curcumin

Curcumin is an orange-yellow pigmented crystallized compound that is found in the rhizome of turmeric (*Curcuma longa* L.) and is mainly utilized as a food coloring agent in Indian sub-continent (Michele Dei Cas). Curcumin is poorly soluble in water. When taken by the oral route, it is absorbed by the wall of gastrointestinal tract and its metabolism takes place in the liver. Curcumin holds the capability of modulating the various kinds of signaling pathways and molecular processes in breast cancer (Mishra et al., 2023). Studies have shown that curcumin can regulate various pathways including the PI3K/ATK, nuclear factor [NF]-kB and many more (S. Hu et al., 2018). Through inducing an arrest in the cell cycle at the G2/M phase, it can also promote the proliferation of cancerous cells (Mosieniak et al., 2012). Many studies have shown that curcumin efficiently causes TNBC cell lines to go through autophagy and apoptosis, which includes MDA-MB-231 (J. Zhu et al., 2024). When curcumin was tested on MCF7 cells the expression of pro-apoptotic protein and apoptosis regulator Bax was found to be high. (Y. Song et al., 2020)

3.2. Genistein

Native to sub-Indigenous continents, genistein is a natural isoflavone mainly found in soybeans. In 1899, Genistein was first extracted from *Genista tinctoria* (L.). It shows anti-oxidant and anti-cancerous properties (Tuli et al., 2019). Genistein can show the cell cycle arrest in G0/G1 and G2/M phase arrest through cyclin B (Jiang et al., 2018). It shows antiproliferative effects by first obstructing NF-kB pathways and then stimulating NF-kB. It can also modulate the eEGFR/Akt/NFkB pathway which contributes significantly to cell proliferation and ultimately causes cancer cells to die (Bhat et al., 2021).

3.3. Quercetin

Quercetin, a polyphenolic compound predominantly found in nuts, teas as well as herbs. It demonstrates various positive impacts that include, antioxidant, anti-diabetic, anti-inflammatory, and anti-proliferative effects. Quercetin is a lipophilic compound that can cross the cellular membrane and initiate several intracellular signaling pathways. Numerous studies have concluded that Quercetin has the anti-cancer properties (Shafabakhsh & Asemi, 2019). Research has demonstrated that it can limit the expansion of breast cancer cells in humans (Maugeri et al., 2023). Quercetin demonstrated dose-dependent anticancer effects in LoVo and MCF-7 cells. The addition of Quercetin-5 to 8-disulphonate can significantly increase the water solubility and anticancer properties of Quercetin. In cancer cells, quercetin induces apoptosis and cell cycle arrest at the S phase (H. Zhang et al., 2012).

3.4. Resveratrol

Resveratrol, a polyphenol phytoalexin is found in various foods including grapes, pistachios, and in small amounts in bilberries and blueberries. It demonstrates a wide range of activities that include anti-oxidative, anti-inflammatory, anti-aging, and cardiovascular protective properties (Meng et al., 2021). It is a compound that has the potential to improve the effects of breast cancer and open up the way for the development of new therapies (Ren et al., 2021). Resveratrol can modulate and target the STAT3 pathway in breast cancer cells. Additionally, it can also influence STAT3's upstream regulators, that can affect the growth, advancement, and spread of breast cancer cells (Kohandel et al., 2021). The genes that show differential methylation were involved in key cellular pathways including Wnt, MAPK, and JAK/STAT. (Lubecka et al., 2016). With the treatment of Resveratrol, oncogenes including AKT1, STAT3, and NOTCH2 show hypermethylation, which aligns with the pathways identified in comprehensive genomic analysis (Kurzava Kendall et al., 2024).

3.5. Kaempferol

Kaempferol is a flavonoid that has multiple beneficiary health effects. It is present in plants including tea trees, saffron, aloe vera, onion leaves, brown mustard, pumpkin, and many others (Periferakis et al., 2022). Kaempferol and its derivatives have shown many beneficiary effects including anticancer, antioxidant, antidiabetic, neuroprotective, and antitumor characteristics (Imran et al., 2019). Kaempferol inhibits the development of MCF-7 breast cancer cells, potentially by stimulating apoptosis and by downregulating the expression of Bcl2 (Yi et al., 2016). Kaempferol inhibits the cell cycle in cancerous breast cells at the G2/M stage by downregulating CDK1 (X. Wang et al., 2019). It was observed that kaempferol reduces IQGAP3 levels in breast cancer cells, causing increased cell death and suppressed cell proliferation (G. Hu et al., 2019).

3.6. Apigenin

Apigenin, (4',5,7,-trihydroxyflavone) is a common flavonoid that is present in various kinds of plants, fruits, and herbs. It shows many beneficiary effects including anti-apoptotic, antioxidant, and anti-cancerous properties (Lee et al., 2023). It has proved its effectiveness in breast cancerous cells. Firstly, in estrogen-responsive breast cancer cells, include those that have become resistant to anti estrogen medications like fulvestrant or tamoxifen. Increased concentrations of apigenin suppress the proliferation of cancerous tissue or cells and demonstrate synergistic effects when administered together with fulvestrant and tamoxifen, indicating that it can handle breast cancer that is resistant to estrogen. Apigenin exhibits significant inhibitory activity in the breast cancer cells which have the overexpressing of HER2/neu by inducing apoptosis on a time and depending on the dosage system. The compound's capacity to modulate PI3K activity, Akt function, and HER2/neu autophosphorylation emphasizes its future potential as a treatment for HER2/neu-positive breast cancer.

Additionally, apigenin inhibits proliferation and stimulates apoptosis by inducing death in breast cancer cells through the caspase cascade, activating caspase-9 subsequently followed by caspase-3 (Rahmani et al., 2022). Apigenin offers the potential of utilizing transcriptome reprogramming which deals with the aggressive and resistant treatment of triple-negative breast cancer (TNBC). Apigenin modulates the alternative splicing (AS) patterns. Abnormal alternative splicing (AS) events foster the development of tumors and resistance to cell death constitutes the characteristic features of TNBC. By broadly reprogramming TNBC-related AS to favor isoforms associated with cell death while sparing normal breast cells, apigenin counteracts this through molecular interactions with core spliceosome subunits and RNA-binding proteins (RBPs) (Sudhakaran et al., 2023). It suppresses the two essential enzymes, aromatase and fatty acid synthase that contribute to the development of breast cancer. Furthermore, through the induction of progesterin, apigenin can prevent angiogenesis by the suppression of VEGF expression. Apigenin inhibits chemokines and inflammatory cytokines, thereby preventing metastasis, a major cause of death from breast cancer. Also, by inhibiting the YAP/TAZ pathway, apigenin reduces the proliferation, migration, and stemness characteristics of TNBC cells (Adel et al., 2022). Figure 2 denotes effect of Apigenin on pathways and progression of Breast Cancer.

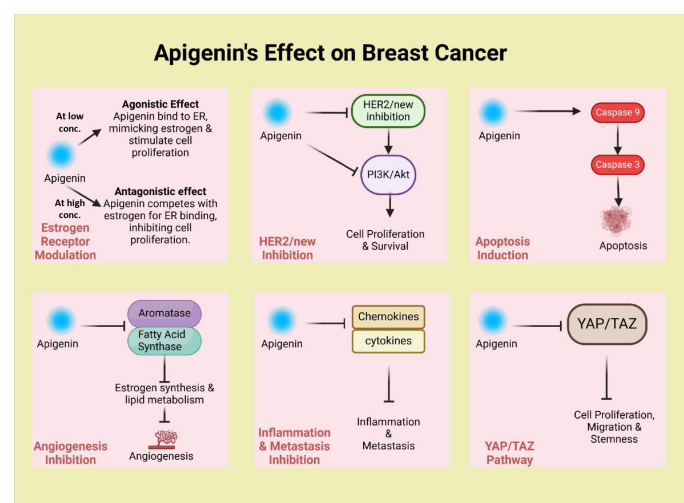


Figure 2. Effect of Apigenin on Breast Cancer

3.7. Silibinin

Silibinin is a common flavonoid present in milk thistle. It shows many positive impacts in regulating diabetes, and liver damage and shows anti-tumor activities. Although its advantages are widely recognized, its precise anti-tumor mechanism is yet to be determined. By application of network pharmacology, many researchers looked into the mechanisms and targets of silibinin's cancer-treating effects. This was supported by the experimentation on mice and adenoid cystic carcinoma cells. Li et al.'s 2024 study provides insights into the potential application of silibinin in the treatment of cancer

by examining the tumor-treating mechanism of the drug for the first time through the use of network pharmacology and experimental validation (P. Li et al., 2024). Silibinin causes cell cycle arrest by targeting the significant signaling pathways involving MAPK and AKT pathways while causing much less damage to normal cells. It has been shown to interact with tumour-suppressive microRNAs such as miR-133a, currently however, it is unknown how precisely these interactions are contributing to the suppression of breast cancer. Given its impact on miR-133a and associated downstream targets, like EGFR, silibinin may still prove to be a useful therapeutic component but to be able to fully understand its effect on miR-133a in the therapy of breast cancer, more research will need to be done (Hossein-Tehrani et al., 2024). Silibinin is a potential therapeutic agent for the treatment of breast cancer as it helps regulate miRNA, causes apoptosis, and decreases the chances of cell invasion and proliferation (Ray et al., 2024). Figure 3 denotes effect of Silibinin on Breast Cancer.

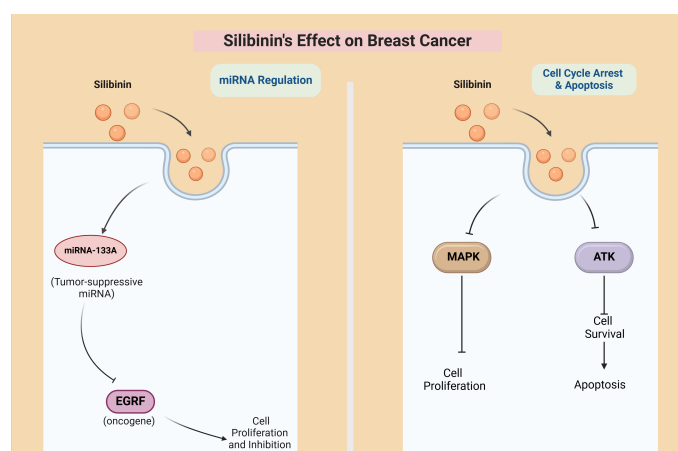


Figure 3. Effect of Silibinin on Breast Cancer.

3.8. EGCG

Epigallocatechin-3-gallate (EGCG), is a polyphenolic compound that is present in green tea. It has been said to show great levels of anti-cancerous activities and also has the ability to regulate various signaling pathways. It has also been observed that when EGCG is combined with other natural and synthetic drugs, it will boost the efficiency of traditional medicine and also reduce their side effects (M. Zhang et al., 2007). Additionally, EGCG has demonstrated a wide range of impacts on various essential signaling pathways which may include phosphatidylinositol-3 kinase (PI3K) as well as MAP kinase. Furthermore, EGCG also decreases the escalated levels of ERK1/2 and Jak/STAT3 pathways which are strongly related to malignant processes (Ediriweera et al., 2017). EGCG alters signaling pathways that affect cell growth and differentiation and inhibits the cell cycle progression. In addition, EGCG suppresses VEGF transcription, triggers apoptosis, and adversely affects several stages of metastasis (Romano & Martel, 2021).

3.9. Lycopene

Lycopene a natural red pigment compound of the carotenoid family that is present in watermelon, papaya, pink grapefruit, and guava (Carvalho et al., 2021). The behavior of MCF-7 cells was studied in relation to lycopene at various concentrations and treatment times. Researchers discovered that lycopene dramatically reduced cell growth using MTT assays and that this effect was connected with both dosage and exposure duration. Notably, throughout 72 hours, a 16 mM concentration resulted in the most noticeable decrease in cell proliferation. Subsequently, flow cytometry was used to detect the apoptosis and the results revealed a significant variation in the rate of cell death between the lycopene-treated group and the control group. The rise in apoptotic response is induced by the increased lycopene doses which indicate a clear interrelation between the dosage and exposure duration. Semi-quantitative mRNA level assessment, which was also examined in this study, revealed that cells treated with lycopene had greater levels of p53 and Bax, both of which were associated with duration and dose in breast cancerous cells (Peng et al., 2017).

3.10. Thymoquinone

Thymoquinone is a compound isolated mainly from black cumin seeds. Thymoquinone shows many beneficiary effects including antioxidant, anti-cancerous, pain-relieving, as well as fever-reducing properties (Ali et al., 2021). The 35 derivatives of thymoquinone showed acceptable physicochemical and absorption properties without hepatotoxicity, which qualified them as oral medications. Based on molecular docking against 3PP0 proteins, the derivatives of thymoquinone demonstrated greater activity as compared to thymoquinone. TQ 15 was the one with the greatest anticancer activity among them, as indicated by its highly substantial negative reactive species value. Lipophilic and steric factors were found to be essential in determining the activity of the 35 thymoquinone derivatives by QSPR analysis (Wulandari et al., 2021). The upregulation of PTPRR inhibits the p38-MAPK and MAPK pathways while downregulating Bcl-2, and upregulating the TP53, thus enhancing apoptosis. TQ also affects PI3K/AKT, Wnt, MAPK, and p53 which are important pathways that are necessary for survival and the growth of cancer-causing cells. It works in combination with paclitaxel and tamoxifen to increase their effectiveness by inducing apoptosis and modifying tumor suppressor genes. As TQ inhibits TGF- β and mesenchymal indicators while restoring epithelial markers, it has anti-metastatic characteristics that prevent the spread of cancer (Younas et al., 2018).

3.11. 3'-diindolylmethane

Brussels sprouts, broccoli, cauliflower, and cabbage contain heterocyclic and biologically active compounds like indole-3-carbinol (I3C) and 3,3'-diindolylmethane (DIM) (Reyes-Hernández et al., 2023). Through a variety of methods, 3,3'-Diindolylmethane (DIM), which is produced from indole-3-carbinol, exhibits hope in the treatment of breast cancer.

DIM significantly reduces the growth of cancerous cells which was proved by investigations utilizing cell lines of breast cancer (MCF-7, BT-474, T47D, BT-20), by promoting p27kip protein expression before the commencement of apoptosis. DIM also modifies signaling pathways. It also reduces Akt activity and increases p27kip transcript levels in BT-20 cells that had Her-2 overexpression and activated Akt. These results point to the probability of DIM as a chemopreventive drug for breast cancer, regardless of the estrogen receptor expression or Her-2 (Z. Wang et al., 2008).

3.12. α Mangostin

The fruit regarded as the "queen of fruits," *Garcinia mangostana* Linn, contains α -mangostin, a xanthone derivative that is isolated from its skin. In various cancers, including those of the breast, colon, lung, pancreatic, skin, and blood, it has demonstrated multiple anticancer capabilities (Meylina et al., 2021). Mangosteen peel is rich in natural polyphenols such as xanthenes as well as vitamins, minerals, phosphorus, and iron. The class of oxygenated heterocycles is made up of the secondary metabolites of xanthenes. [1,3,6,7-tetrahydroxy-2,8-di (3-methyl-2-butenyl)] one α mangostin metabolite found in xanthenes. C₂₄H₂₂O₆ is the chemical formula for α mangostin (Herdiana et al., 2021). The research by Simon examined how α -mangostin breast cancer cells, and it revealed the dose-related reduction in cells viability, especially in spheroid models. According to flow cytometry analysis, α -mangostin causes MCF-7 cells to exhibit apoptosis indicated by positive annexin-V staining, caspase activation, as well as nuclei condensation. Myc-BCL-XL clones showed resistance to α -mangostin-induced apoptosis, while stable MCF-7 clones overexpressing HA-MOAP-1 showed increased susceptibility. In MCF-7 cells, BCL-XL opposes the activation of MOAP-1-mediated apoptosis caused by α -mangostin (Simon et al., 2022). To improve efficacy and lessen the endometrial side effects, a study looked at ER+ breast cancer patients being treated with α -mangostin (AM) in addition to tamoxifen. It prevented 4-OH-TMX-induced CCND1 overexpression in endometrial cells and downregulated oncogenic genes (KCNH1, CCND1, BIRC5) in breast cancer cells. These results demonstrate AM's potential to supplement ER+ breast cancer therapy in a way that would improve treatment outcomes while minimizing side effects (Vargas-Castro et al., 2023). The molecular mechanisms of alpha mangostin are shown in the given Figure 4.

3.13. Daidzein

As a multifunctional herbal therapy, Daidzein is a phytoestrogen categorized as a nonsteroidal oestrogen and a component of the isoflavone class, cures an array of ailments which include cancer, inflammation, and osteoporosis. Its various therapeutic properties comprise anti-inflammatory, antioxidant, and antithrombotic properties. By its modulation of oestrogen control, Daidzein exhibits preventive effects against breast cancer. Despite its effects on oestrogen receptors, it also

safeguards nerves and the skin, reduces inflammation, combat cancer, and reduces oxidative damage. It is naturally found in foods based on soy, including tempeh, wheat, and maize-based fortified supplemental flours, textured soy protein, miso, and tofu (Magee et al., 2016). Research conducted by Kumar, V et al., shows how Daidzein increases the level of pro-apoptotic protein Bax by increasing the apoptosis through the intrinsic pathway, which results in changes in mitochondrial membrane permeability and the release of cytochrome c. This activates caspase 3/7 and caspase 9 which leads to apoptosis (Kumar & Chauhan, 2021).

3.14. Secoisolariciresinol

Secoisolariciresinol diglucoside, derived from flaxseed, exhibits promising pharmaceutical effects like antioxidative and anti-inflammatory effects. Research highlights its ability to treat various cancers, that include lung and breast cancers, by inducing apoptosis in cancer cells. This process involves the reactive oxygen species (ROS) production, that causes the potential of the mitochondrial membrane to break and DNA damage in cells with acute lymphoblastic leukemia. Moreover, it inhibits the cell cycle proteins to promote the growth of breast cancer cells by upregulating p21WAF1/CIP1 and downregulating G1-phase cyclins and CDKs (Chen et al., 2022).

3.15. Hesperetin

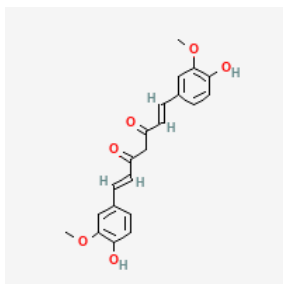
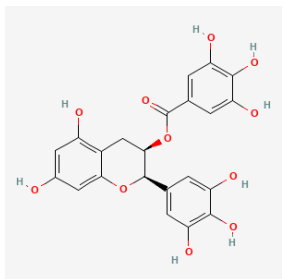
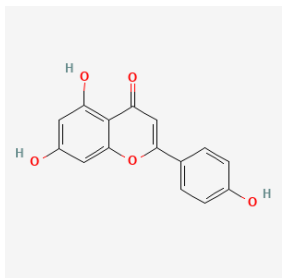
Citrus fruits, including grapefruits, lemons, and oranges, are excellent sources of two flavonoid compounds hesperetin and its aglycone, hesperidin. Pharmacological actions ranging from cardioprotective to hepatoprotective, neuroprotective, anti-tumor, anti-diabetic, antimicrobial, antioxidant have been documented (Yap et al., 2021). Hesperetin works similarly against all of these subtypes and is more effective against HER2-positive cells because it inhibits the activity of HER2 tyrosine kinase. In models where tumor growth is dependent on estrogen, hesperetin also inhibits aromatase, lowering estrogen levels. In breast cancer, both flavonoids show promise in reducing metastasis and preserving tissue architecture (Yap et al., 2021).

4. SOURCE AND MODE OF ACTION OF DIFFERENT PHYTOCHEMICALS

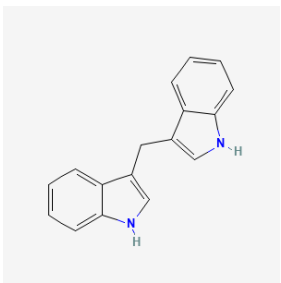
Different phytochemicals have different sources and modes of action, which are listed in the table below. The 2D structures in the table have all been retrieved from PubChem. (National Center for Biotechnology Information (NCBI), n.d.)

Table 1

Source of Different Phytochemicals and Various cancer pathways targeted by these Phytochemicals

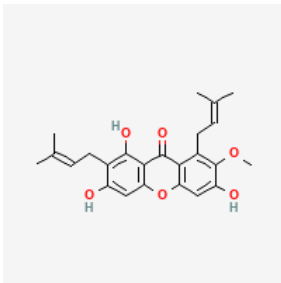
S.No.	Phytochemical	Type	2D-Structure	Sources	Breast Cancer Pathways affected by Phytochemicals	Ref.
1.	Curcumin	Polyphenolic	 <p>A structure of a chemical formula Description automatically generated</p>	<i>Curcuma longa</i> (Turmeric)	P53, Ras, AKT, Wnt- β catenin, mTOR, JAK/STAT, NF- κ B, apoptosis-related signaling pathway, ERK pathway, Bcl-2, caspase, EMT pathway	Banerjee and Mandal (2022); S. Hu et al. (2018); Lambring et al. (2023); Morgan et al. (2018); X. Song et al. (2019)
2.	EGCG	Catechin	 <p>A structure of a chemical formula Description automatically generated</p>	<i>Camellia sinensis</i> (Green Tea)	Wnt, MAP kinase, TGF- β , MTOR, P53, PI3K, EGF receptor signaling pathway, actin cytoskeleton pathway regulation, JAK/STAT, insulin signaling pathway, axon guidance pathway, cadherin pathway, PDGF signaling pathway, notch, NF- κ B, EGFR, Ras	Morgan et al. (2018); Z. Wang et al. (2008)
3.	Apigenin	Flavone	 <p>A structure of a chemical formula Description automatically generated</p>	<i>Petroselinum crispum</i> (Parsley), <i>Apium graveolens</i> (celery), <i>Lactuca sativa</i> (lettuce)	Estrogen signaling pathway, HER2/neu signaling pathway, Caspase cascade (caspase-9, caspase-3), PI3K/Akt signaling pathway, Transcriptome reprogramming, Alternative splicing regulation, Apoptosis induction (intrinsic, extrinsic pathways), p53 activation, Receptor tyrosine kinase inhibition, Aromatase inhibition, Fatty acid synthase suppression, Angiogenesis inhibition (VEGF suppression), Inflammation modulation (cytokines, chemokines), Metastasis suppression, YAP/TAZ pathway blockade.	Adel et al. (2022); Morgan et al. (2018); Sudhakaran et al. (2023); Ubaid et al. (2023)

4. 3,3'-diindolylmethane Indoles



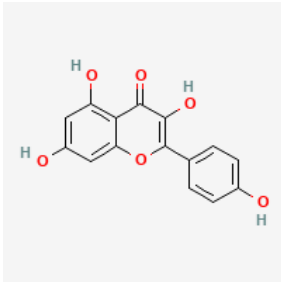
A molecule of a chemical structure Description automatically generated with medium confidence

Cruciferous vegetables In MCF10A breast cancer cells inhibit AKT activation, induce p27kip protein and apoptosis. [Morgan et al. \(2018\); Vargas-Castro et al. \(2023\)](#)
5. Alpha-mangostin Xanthonoid



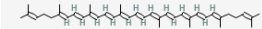
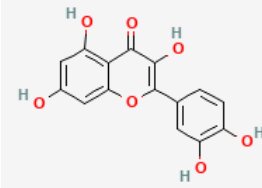
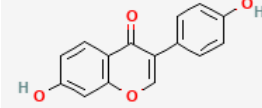
A structure of a chemical formula Description automatically generated

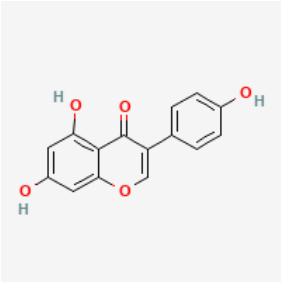
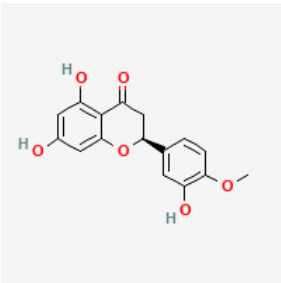
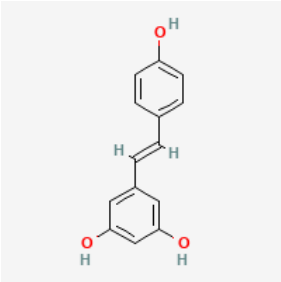
Garcinia mangostana (Mangosteen fruit) Enhance the anti-proliferative effect of tamoxifen, mitochondrial apoptotic pathway. [Chen et al. \(2022\); Morgan et al. \(2018\); Roy et al. \(2019\)](#)
6. Kaempferol Flavonoid



A structure of a chemical formula Description automatically generated

(*Camellia sinensis*) Tea, (*Phaseolus nanus* L.) beans, *Malus spp.* (apples), and *Aloe barbadensis* (aloe vera) ERK1/2 pathway, suppresses proliferation, increases the level of gH2AX, cleave caspase 3, caspase 9 and p-ATM, activates p53, downregulates PI3K-AKT. [Lee et al. \(2023\); Morgan et al. \(2018\); Sengupta et al. \(2022\); Vasconcelos et al. \(2020\)](#)

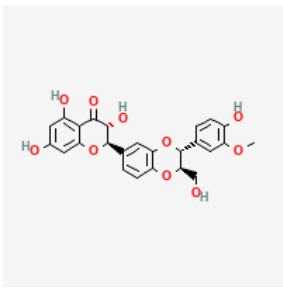
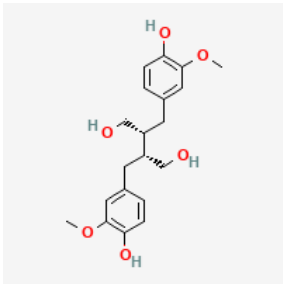
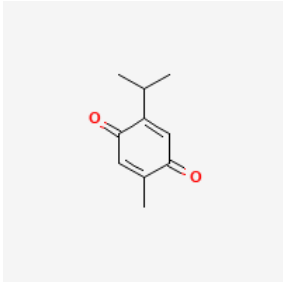
- | | | | | | | |
|----|-----------|--------------|--|--|---|--|
| 7. | Lycopene | Tetraterpene |  | <i>Solanum lycopersicum</i> (Tomatoes), <i>Cucumis melo</i> (melons). | NF-KB pathway was inhibited in HMC3 cells treated with nanoLEG. | Jang et al. (2022);
Morgan et al. (2018) |
| 8. | Quercetin | Flavonoid |  | Citrus fruits, <i>Allium cepa</i> L. (onions), <i>Brassica oleracea</i> (broccoli) | Inhibit PI3K-ATK pathway which that regulate NF-kB activity, PI3K, inhibit AKT pathway, mTOR, STAT3 | Bhat et al. (2021);
Morgan et al. (2018) |
| 9. | Daidzein | Isoflavone |  | <i>Glycine</i> (Soybeans) soy products | Estrogen receptor modulation, inflammation modulation, antioxidant pathways. | Magee et al. (2016);
Morgan et al. (2018) |
- A structure of a chemical formula Description automatically generated
- A structure of a chemical formula Description automatically generated
- A structure of a chemical formula Description automatically generated

- | | | | | | | |
|-----|-------------|---------------------|---|--|--|--|
| 10. | Genistein | Isoflavonoid |  | Soy-based edibles | Wnt, suppresses tyrosine kinase, modulate hedgehog-Gle1 signaling, seizes AKT and MEK signaling. | Morgan et al. (2018) ; Shoaib et al. (2023) ; H. Zhang et al. (2012) |
| 11. | Hesperetin | Flavanone glycoside |  | Citrus fruits | Apoptosis induction via ROS-mediated mitochondrial dysfunction and DNA damage, cell cycle regulation through G1-phase cyclin and CDK downregulation, and p21WAF1/CIP1 upregulation | Behroozaghdam et al. (2021) ; Morgan et al. (2018) |
| 12. | Resveratrol | Stilbenes |  | <i>Malus spp.</i> (Apples),
<i>Prunus subg</i> (plums). | JAK/STAT pathway, STAT3 pathway, Wnt pathway, MAPK pathway. | Kurzava Kendall et al. (2024) ; Lubecka et al. (2016) ; Morgan et al. (2018) ; Periferakis et al. (2022) |

A structure of a chemical formula Description automatically generated

A structure of a chemical formula Description automatically generated

A structure of a chemical formula Description automatically generated

13.	Silibinin	Flavonoid		<i>Silybum marianum</i> (Milk thistle)	MAPK signaling pathway, apoptosis induction, miRNA regulation, AKT signaling pathway, EGFR signaling pathway, chemo-resistance reversal, metastasis suppression, cell cycle regulation.	Hossein-Tehrani et al. (2024) ; Morgan et al. (2018) ; Ray et al. (2024) ; M. Zhang et al. (2007)
14.	Secoisolariciresinol	Lignan		<i>Brassica oleracea</i> (broccoli)	Apoptosis induction, potential breakdown of mitochondrial membrane, DNA damage.	Morgan et al. (2018) ; Yap et al. (2021)
15.	Thymoquinone	Antioxidant		<i>Nigella sativa</i> (Black seeds)	TQ stimulate breast cancer cells to paclitaxel and reverses resistance of tamoxifen.	Morgan et al. (2018) ; Younas et al. (2018)

A structure of a chemical formula Description automatically generated

A structure of a chemical formula Description automatically generated

A black and red molecule Description automatically generated

Table 2

A comparative analysis of enlisted Phytochemical's Limitations

S.NO	PHYTOCHEMICAL	LIMITATIONS	REFERENCE
1.	EGCG	poor bioavailability and absorption, and unpredictable distribution in human tissues.	X. Song et al. (2019)
2.	Resveratrol	Rapid absorption and metabolism, poor bioavailability, and low aqueous solubility, low potency.	Behroozaghdam et al. (2021)
3.	Curcumin	poor bioavailability and absorption, may be toxic at higher doses	Agarwal et al. (2023); Silva et al. (2022)
4.	Kaempferol	Poor bioavailability, low water solubility, rapid metabolism	Da et al. (2019)
5.	Genistein	Rapid metabolism, poor bioavailability, and low aqueous solubility	Emerich and Thanos (2006); Ferrado et al. (2023)
6.	Quercetin	Poor solubility and bioavailability, rapid metabolism	H. Zhang et al. (2012)
7.	Lycopene	Poor absorption, Lack of vitamin A activity,	van Breemen (2019)
8.	α mangostin	high frequency of adverse effects, restricted efficacy, multidrug resistance, and extreme toxicity to developing healthy cells	Herdiana et al. (2021)
9.	Thymoquinone	Poor bioavailability, low water solubility, rapid metabolism	Chehl et al. (2009); Kundu et al. (2014)
10.	Apigenin	Poor bioavailability, dose-dependent effect	Patel and Patel (2015); Shukla and Gupta (2010)
11.	Hesperetin	poor bioavailability and rapid metabolism, lack sufficient selectivity for cancer cells over normal cells	Russo et al. (2020); Y. Song et al. (2020)
12.	Silibinin	Poor bioavailability and pharmacokinetic properties limit its efficacy, poor solubility, rapid metabolism	Kaur et al. (2020); Ting and Jiang (2020)
13.	Secoisolariciresinol	variable bioavailability, metabolism differences among individuals	Bergamo and Goglia (2021)
14.	Daidzein	poor bioavailability and undergoes extensive metabolism in the body	Magee et al. (2016)
15.	3,3'-diindolylmethane	Complex metabolism and variability	Fan et al. (2021)

5. CONCLUSION

Concerning data on the mortality rate from cancer especially breast cancer highlights the urgency of searching for novel and efficient therapeutic approaches. This review reveals the pathways by which phytochemicals affect breast cancer biology and includes them in therapeutic approaches that may enhance patient outcomes and contribute to the goals outlined in the Sustainable Development Goal 3 of the United Nation's goals. The natural compounds reviewed here, which include curcumin, apigenin, genistein, quercetin, resveratrol, kaempferol, hesperitin, and others offer promising potential therapeutics for breast cancer. The anti-cancer capabilities of every compound have been attributed to their distinctive mechanisms of action. Curcumin and EGCG, for instance, modulate several signaling pathways, that include NF- κ B and MAPK, which have significance for the survival and growth of cancerous cells. Through the production of oxidative stress and disruption of the cell cycle, resveratrol and quercetin inhibit the growth of cancerous cells, correspondingly. Through the production of oxidative stress and disruption of the cell

cycle, resveratrol and quercetin inhibit the growth of cancerous cells, correspondingly. By the stimulation of apoptosis and controlling the cell cycle progression, apigenin and DIM shows a particular potential in targeting hormone receptor-positive and triple-negative breast cancer cells. In compounds like α -mangostin and silibinin, apoptosis can be induced through mitochondrial pathways which helps to overcome the drug resistance. These organic substances help in promoting universal health care and well-being for cancer treatment by showing therapeutic and preventive effects.

CONFLICTS OF INTEREST

The authors declare that there is no conflict of interest.

AUTHORSHIP CONTRIBUTIONS

DP conducted initial research, data collection, manuscript drafting, and drawing the figures.

MEP contributed to data analysis and data interpretation.

VJU contributed to manuscript editing.

PS conceptualised the study, supervised the manuscript writing process, figures, and contributed to the critical review of the manuscript.

All authors read and approved the final manuscript.

ORCID

Dipika Pal 0009-0000-1203-0897

Pallavi Singh 0000-0003-4537-4607

Madhulika Esther Prasad 0000-0002-8862-3557

Vijay Jagdish Upadhye 0000-0002-8821-1720

REFERENCES

- Adel, M., Zahmatkeshan, M., Akbarzadeh, A., Rabiee, N., Ahmadi, S., Keyhanvar, P., Rezaayat, S.M., Seifalian, A.M., 2022. Chemotherapeutic effects of apigenin in breast cancer: Preclinical evidence and molecular mechanisms; enhanced bioavailability by nanoparticles. *Biotechnology reports (Amsterdam, Netherlands)*. 34, e00730. <https://doi.org/10.1016/j.btre.2022.e00730>
- Agarwal, V., Gaur, P., Mohanty, S., 2023. Clinical effects of curcumin in enhancing cancer therapy: A systematic review. *BMC Cancer*. <https://doi.org/10.1186/s12885-023-11032-6>
- Ali, M.Y., Akter, Z., Mei, Z., Zheng, M., Tania, M., Khan, M.A., 2021. Thymoquinone in autoimmune diseases: Therapeutic potential and molecular mechanisms. *Biomedicine & pharmacotherapy = Biomedecine & pharmacotherapie*. 134, 111157. <https://doi.org/10.1016/j.biopha.2020.111157>
- Arnold, M., Morgan, E., Rungay, H., Mafra, A., Singh, D., Laversanne, M., Vignat, J., Gralow, J.R., Cardoso, F., Siesling, S., Soerjomataram, I., 2022. Current and future burden of breast cancer: Global statistics for 2020 and 2040. *Breast (Edinburgh, Scotland)*. 66, 15-23. <https://doi.org/10.1016/j.breast.2022.08.010>
- Banerjee, S., Mandal, A.K.A., 2022. Role of epigallocatechin-3-gallate in the regulation of known and novel microRNAs in breast carcinoma cells. *Frontiers in genetics*. 13, 995046. <https://doi.org/10.3389/fgene.2022.995046>
- Behroozaghdam, M., Dehghani, M., Hashemi, M., Zabolian, A., Kamali, D., Javanshir, S., Hasani Sadi, F., 2021. Resveratrol in breast cancer treatment: From cellular effects to molecular mechanisms of action. *Cellular and Molecular Life Sciences*. 78, 761-778. <https://doi.org/10.1007/s00018-020-03520-1>
- Bergamo, P., Goglia, P.R., 2021. Secoisolariciresinol diglucoside (sdg) from flaxseed as a potential preventive and therapeutic agent in breast cancer: Insights from recent studies. *Cancer Treatment and Research Communications*. 28, 100443. <https://doi.org/10.1016/j.ctarc.2021.100443>
- Bhat, S.S., Prasad, S.K., Shivamallu, C., Prasad, K.S., Syed, A., Reddy, P., Cull, C.A., Amachawadi, R.G., 2021. Genistein: A potent anti-breast cancer agent. *Current issues in molecular biology*. 43(3), 1502-1517. <https://doi.org/10.3390/cimb43030106>
- Bimonte, S., Cascella, M., Barbieri, A., Arra, C., Cuomo, A., 2020. Current shreds of evidence on the anticancer role of egcg in triple negative breast cancer: an update of the current state of knowledge. *Infectious agents and cancer*. 15, 2. <https://doi.org/10.1186/s13027-020-0270-5>
- Bray, F., Laversanne, M., Cao, B., Varghese, C., Mikkelsen, B., Weiderpass, E., Soerjomataram, I., 2021. Comparing cancer and cardiovascular disease trends in 20 middle- or high-income countries 2000-19: A pointer to national trajectories towards achieving sustainable development goal target 3.4. *Cancer treatment reviews*. 100, 102290. <https://doi.org/10.1016/j.ctrv.2021.102290>
- Carvalho, G.C., Sábio, R.M., Chorilli, M., 2021. An overview of properties and analytical methods for lycopene in organic nanocarriers. *Critical reviews in analytical chemistry*. 51(7), 674-686. <https://doi.org/10.1080/10408347.2020.1763774>
- Cazzaniga, M., Bonanni, B., 2012. Breast cancer chemoprevention: old and new approaches. *Journal of biomedicine & biotechnology*. 2012, 985620. <https://doi.org/10.1155/2012/985620>
- Chehl, N., Chipitsyna, G., Gong, Q., Yeo, C.J., 2009. Molecular targets of dietary agents for prevention and therapy of cancer. *Biochemical Pharmacology*. 78(10), 1179-1184. <https://doi.org/10.1016/j.bcp.2009.06.039>
- Chen, T., Wang, Z., Zhong, J., Zhang, L., Zhang, H., Zhang, D., Xu, X., Zhong, X., Wang, J., Li, H., 2022. Secoisolariciresinol diglucoside induces pyroptosis by activating caspase-1 to cleave gsdmd in colorectal cancer cells. *Drug development research*. 83(5), 1152-1166. <https://doi.org/10.1002/ddr.21939>
- Da, J., Xu, M., Wang, Y., Li, W., Lu, M., Wang, Z., 2019. Kaempferol promotes apoptosis while inhibiting cell proliferation via androgen-dependent pathway and suppressing vasculogenic mimicry and invasion in prostate cancer. *Analytical Cellular Pathology*. 2019, 1-10. <https://doi.org/10.1155/2019/1>
- Dei Cas, M., Ghidoni, R., 2019. Dietary curcumin: Correlation between bioavailability and health potential. *Nutrients*. 11(9), 2147. <https://doi.org/10.3390/nu11092147>
- Ediriweera, M.K., Tennekoon, K.H., Samarakoon, S.R., Thabrew, I., de Silva, E.D., 2017. Protective effects of six selected dietary compounds against leptin-induced proliferation of oestrogen receptor positive (mcf-7) breast cancer cells. *Medicines (Basel, Switzerland)*. 4(3), 56. <https://doi.org/10.3390/medicines4030056>
- Emerich, D.F., Thanos, C.G., 2006. The pinpoint promise of nanoparticle-based drug delivery and molecular diagnosis. *Biomolecular Engineering*. 23(4), 171-184. <https://doi.org/10.1016/j.bioeng.2006.05.026>
- Fan, J., Meng, Q., Guo, D., Wang, W., Yu, M., Zhang, X., Fang, J., 2021. 3,3'-diindolylmethane and cancer therapy: Preclinical and clinical progress, molecular mechanisms and future perspectives. *Expert Opinion on Investigational Drugs*. 30(6), 573-586. <https://doi.org/10.1080/13543784.2021.1918102>
- Ferrado, J.B., Perez, A.A., Menegon, M., 2023. Pegylation of genistein-loaded bovine serum albumin nanoparticles and its effect on in vitro cell viability and genotoxicity properties. *Colloids and Surfaces B: Biointerfaces*. 222, 113082. <https://doi.org/10.1016/j.colsurfb.2022.113082>
- Frieden, T.R., Cobb, L.K., Leidig, R.C., Mehta, S., Kass, D., 2020. Reducing premature mortality from cardiovascular and other non-communicable diseases by one third: Achieving sustainable development goal indicator 3.4.1. *Global heart*. 15(1), 50. <https://doi.org/10.5334/gh.531>
- Herdiana, Y., Wathoni, N., Shamsuddin, S., Muchtaridi, M., 2021. α -mangostin nanoparticles cytotoxicity and cell death modalities in breast cancer cell lines. *Molecules (Basel, Switzerland)*. 26(17), 5119. <https://doi.org/10.3390/molecules26175119>
- Hossein-Tehrani, M., Abbasalipourkabir, R., Ziamajidi, N., 2024. The role of mir-133a in silibinin-mediated inhibition of the pi3k/akt/mtor pathway in mcf-7 breast carcinoma cells. *Molecular biology research communications*. 13(2), 79-83. <https://doi.org/10.22099/MBRC.2024.48818.1903>
- Hu, G., Liu, H., Wang, M., Peng, W., 2019. Iq motif containing gtpase-activating protein 3 (iqgap3) inhibits kaempferol-induced apoptosis in breast cancer cells by extracellular signal-regulated kinases 1/2

- (erk1/2) signaling activation. *Medical Science Monitor : International Medical Journal of Experimental and Clinical Research*. 25, 7666-7674. <https://doi.org/10.12659/MSM.915642>
- Hu, S., Xu, Y., Meng, L., Huang, L., Sun, H., 2018. Curcumin inhibits proliferation and promotes apoptosis of breast cancer cells. *Experimental and therapeutic medicine*. 16(2), 1266-1272. <https://doi.org/10.3892/etm.2018.6345>
- Hussein, R.A., El-Anssary, A.A., 2019. Plants Secondary Metabolites: The Key Drivers of the Pharmacological Actions of Medicinal Plants.. *IntechOpen*, p. 11-30. <https://doi.org/10.5772/intechopen.76139>
- Imran, M., Salehi, B., Sharifi-Rad, J., Aslam Gondal, T., Saeed, F., Imran, A., Shahbaz, M., Tsouh Fokou, P.V., Umair Arshad, M., Khan, H., Guerreiro, S.G., Martins, N., Estevinho, L.M., 2019. Kaempferol: A key emphasis to its anticancer potential. *Molecules (Basel, Switzerland)*. 24(12), 2277. <https://doi.org/10.3390/molecules24122277>
- Jang, W.Y., Kim, M.Y., Cho, J.Y., 2022. Antioxidant, anti-inflammatory, anti-menopausal, and anti-cancer effects of lignans and their metabolites. *International journal of molecular sciences*. 23(24), 15482. <https://doi.org/10.3390/ijms232415482>
- Jiang, H., Fan, J., Cheng, L., Hu, P., Liu, R., 2018. The anticancer activity of genistein is increased in estrogen receptor beta 1-positive breast cancer cells. *OncoTargets and therapy*. 11, 8153-8163. <https://doi.org/10.2147/OTT.S182239>
- Kaur, R., Kaur, M., Singh, J., 2020. Limitations of silibinin in breast cancer: Insights from recent studies. *Frontiers in Pharmacology*. 11, 586448. <https://doi.org/10.3389/fphar.2020.586448>
- Kohandel, Z., Farkhondeh, T., Aschner, M., Pourbagher-Shahri, A.M., Samarghandian, S., 2021. Stat3 pathway as a molecular target for resveratrol in breast cancer treatment. *Cancer cell international*. 21(1), 468. <https://doi.org/10.1186/s12935-021-02179-1>
- Kumar, V., Chauhan, S.S., 2021. Daidzein induces intrinsic pathway of apoptosis along with er α/β ratio alteration and ros production. *Asian Pacific journal of cancer prevention : APJCP*. 22(2), 603-610. <https://doi.org/10.31557/APJCP.2021.22.2.603>
- Kundu, J., Chun, K.S., Aruoma, O.I., 2014. Potential for therapeutic use of hydrogen sulfide in oxidative stress-induced neurodegenerative diseases. *International Journal of Medical Sciences*. 11(5), 553-559. <https://doi.org/10.7150/ijms.8431>
- Kurzava Kendall, L., Ma, Y., Yang, T., Lubecka, K., Stefanska, B., 2024. Epigenetic effects of resveratrol on oncogenic signaling in breast cancer. *Nutrients*. 16(5), 699. <https://doi.org/10.3390/nu16050699>
- Lambring, C.B., Chen, L., Nelson, C., Stevens, A., Bratcher, W., Basha, R., 2023. Oxidative stress and cancer: Harnessing the therapeutic potential of curcumin and analogues against cancer. *European journal of biology*. 82(2), 317-325. <https://doi.org/10.26650/eurbjbiol.2023.1348427>
- Lee, I.G., Lee, J., Hong, S.H., Seo, Y.J., 2023. Apigenin's therapeutic potential against viral infection. *Frontiers in bioscience (Landmark edition)*. 28(10), 237. <https://doi.org/10.31083/j.fbl2810237>
- Li, C., Xu, Y., Zhang, J., Zhang, Y., He, W., Ju, J., Wu, Y., Wang, Y., 2023. The effect of resveratrol, curcumin and quercetin combination on immuno-suppression of tumor microenvironment for breast tumor-bearing mice. *Scientific reports*. 13(1), 13278. <https://doi.org/10.1038/s41598-023-39279-z>
- Li, P., Wang, D., Yang, X., Liu, C., Li, X., Zhang, X., Liu, K., Zhang, Y., Zhang, M., Wang, C., Wang, R., 2024. Anti-tumor activity and mechanism of silibinin based on network pharmacology and experimental verification. *Molecules (Basel, Switzerland)*. 29(8), 1901. <https://doi.org/10.3390/molecules29081901>
- Lubecka, K., Kurzava, L., Flower, K., Buvala, H., Zhang, H., Teegarden, D., Camarillo, I., Suderman, M., Kuang, S., Andrisani, O., Flanagan, J.M., Stefanska, B., 2016. Stilbenoids remodel the dna methylation patterns in breast cancer cells and inhibit oncogenic notch signaling through epigenetic regulation of maml2 transcriptional activity. *Carcinogenesis*. 37(7), 656-668. <https://doi.org/10.1093/carcin/bgw048>
- Magee, P.J., Allsopp, P., Samaletdin, A., Rowland, I.R., 2016. Daidzein, r-(+)-equol and s-(-)-equol inhibit the invasion of mda-mb-231 breast cancer cells potentially via the down-regulation of matrix metalloproteinase-2. *European Journal of Nutrition*. 55(2), 713-722. <https://doi.org/10.1007/s00394-015-0903-y>
- Maugeri, A., Calderaro, A., Patanè, G.T., Navarra, M., Barreca, D., Cirmi, S., Felice, M.R., 2023. Targets involved in the anti-cancer activity of quercetin in breast, colorectal and liver neoplasms. *International journal of molecular sciences*. 24(3), 2952. <https://doi.org/10.3390/ijms24032952>
- Mazurakova, A., Koklesova, L., Samec, M., Kudela, E., Kajo, K., Skuciova, V., Csizmar, S.H., Mestanova, V., Pec, M., Adamkov, M., Al-Ishaq, R.K., Smejkal, K., Giordano, F.A., Büsselberg, D., Biringer, K., Golubnitschaja, O., Kubatka, P., 2022. Anti-breast cancer effects of phytochemicals: primary, secondary, and tertiary care. *The EPMA journal*. 13(2), 315-334. <https://doi.org/10.1007/s13167-022-00277-2>
- Meng, T., Xiao, D., Muhammed, A., Deng, J., Chen, L., He, J., 2021. Anti-inflammatory action and mechanisms of resveratrol. *Molecules (Basel, Switzerland)*. 26(1), 229. <https://doi.org/10.3390/molecules26010229>
- Meylina, L., Muchtaridi, M., Joni, I.M., Mohammed, A.F.A., Wathoni, N., 2021. Nanoformulations of β -mangostin for cancer drug delivery system. *Pharmaceutics*. 13(12), 1993. <https://doi.org/10.3390/pharmaceutics13121993>
- Mishra, A.P., Swetanshu, Singh, P., Yadav, S., Nigam, M., Seidel, V., Rodrigues, C.F., 2023. Role of the dietary phytochemical curcumin in targeting cancer cell signalling pathways. *Plants (Basel, Switzerland)*. 12(9), 1782. <https://doi.org/10.3390/plants12091782>
- Morgan, G.W., Foster, K., Healy, B., Opie, C., Huynh, V., 2018. Improving health and cancer services in low-resource countries to attain the sustainable development goals target 3.4 for noncommunicable diseases. *Journal of global oncology*. 4, 1-11. <https://doi.org/10.1200/JGO.18.00185>
- Mosieniak, G., Adamowicz, M., Alster, O., Jaskowiak, H., Szczepankiewicz, A.A., Wilczynski, G.M., Ciechomska, I.A., Sikora, E., 2012. Curcumin induces permanent growth arrest of human colon cancer cells: link between senescence and autophagy. *Mechanisms of ageing and development*. 133(6), 444-455. <https://doi.org/10.1016/j.mad.2012.05.004>
- Murthy, S.S., Trapani, D., Cao, B., Bray, F., Murthy, S., Kingham, T.P., Are, C., Ilbawi, A.M., 2024. Premature mortality trends in 183 countries by cancer type, sex, who region, and world bank income level in 2000-19: a retrospective, cross-sectional, population-based study. *The Lancet. Oncology*. 25(8), 969-978. [https://doi.org/10.1016/S1470-2045\(24\)00274-2](https://doi.org/10.1016/S1470-2045(24)00274-2)
- National Center for Biotechnology Information (NCBI), n.d.. PubChem. <https://pubchem.ncbi.nlm.nih.gov/> Accessed: [insert date if available].
- Patel, D., Patel, K., 2015. Apigenin and cancer chemoprevention: Progress, potential and promise (review). *International Journal of Oncology*. 46(2), 743-755. <https://doi.org/10.3892/ijo.2014.2757>
- Peng, S.J., Li, J., Zhou, Y., Tuo, M., Qin, X.X., Yu, Q., Cheng, H., Li, Y.M., 2017. In vitro effects and mechanisms of lycopene in mcf-7 human breast cancer cells. *Genetics and molecular research : GMR*. 16(2). <https://doi.org/10.4238/gmr16029434>
- Pereira, M., Camanho, A., Marques, R., Figueira, J., 2021. The

- convergence of the world health organization member states regarding the united nations' sustainable development goal 'good health and well-being'. Omega-international Journal of Management Science. 104, 102495. <http://dx.doi.org/10.1016/j.omega.2021.102495>
- Periferakis, A., Periferakis, K., Badarau, I.A., Petran, E.M., Popa, D.C., Caruntu, A., Costache, R.S., Scheau, C., Caruntu, C., Costache, D.O., 2022. Kaempferol: Antimicrobial properties, sources, clinical, and traditional applications. International journal of molecular sciences. 23(23), 15054. <https://doi.org/10.3390/ijms232315054>
- Rahmani, A.H., Alsahli, S.A., Almatroudi, A., Almogbel, M.A., Khan, A.A., Anwar, S., Almatroodi, S.A., 2022. The potential role of apigenin in cancer prevention and treatment. Molecules (Basel, Switzerland). 27(18), 6051. <https://doi.org/10.3390/molecules27186051>
- Ramani, V.K., Jayanna, K., Naik, R., 2023. A commentary on cancer prevention and control in india: Priorities for realizing sdgs. Health science reports. 6(2), e1126. <https://doi.org/10.1002/hsr2.1126>
- Ray, P.P., Islam, M.A., Islam, M.S., Han, A., Geng, P., Aziz, M.A., Mamun, A.A., 2024. A comprehensive evaluation of the therapeutic potential of silibinin: a ray of hope in cancer treatment. Frontiers in pharmacology. 15, 1349745. <https://doi.org/10.3389/fphar.2024.1349745>
- Ren, B., Kwah, M.X., Liu, C., Ma, Z., Shanmugam, M.K., Ding, L., Xiang, X., Ho, P.C., Wang, L., Ong, P.S., Goh, B.C., 2021. Resveratrol for cancer therapy: Challenges and future perspectives. Cancer letters. 515, 63-72. <https://doi.org/10.1016/j.canlet.2021.05.001>
- Reyes-Hernández, O.D., Figueroa-González, G., Quintas-Granados, L.I., Gutiérrez-Ruiz, S.C., Hernández-Parra, H., Romero-Montero, A., Del Prado-Audelo, M.L., Bernal-Chavez, S.A., Cortés, H., Peña-Corona, S.I., Kiyekbayeva, L., Ateşşahin, D.A., Goloshvili, T., Leyva-Gómez, G., Sharifi-Rad, J., 2023. 3,3'-diindolylmethane and indole-3-carbinol: potential therapeutic molecules for cancer chemoprevention and treatment via regulating cellular signaling pathways. Cancer cell international. 23(1), 180. <https://doi.org/10.1186/s12935-023-03031-4>
- Romano, A., Martel, F., 2021. The role of egcg in breast cancer prevention and therapy. Mini reviews in medicinal chemistry. 21(7), 883-898. <https://doi.org/10.2174/1389557520999201211194445>
- Roy, R., Hahm, E.R., White, A.G., Anderson, C.J., Singh, S.V., 2019. Akt-dependent sugar addiction by benzyl isothiocyanate in breast cancer cells. Molecular carcinogenesis. 58(6), 996-1007. <https://doi.org/10.1002/mc.22988>
- Russo, M., Milioto, A., Spagnuolo, C., 2020. Dietary flavonoids in cancer therapy and prevention: Substrates and inhibitors of cytochrome p450 enzymes involved in drug metabolism. Current Drug Metabolism. 21(10), 749-762. <https://doi.org/10.2174/138920022166621082611513>
- Sengupta, B., Biswas, P., Roy, D., Lovett, J., Simington, L., Fry, D.R., Travis, K., 2022. Anticancer properties of kaempferol on cellular signaling pathways. Current topics in medicinal chemistry. 22(30), 2474-2482. <https://doi.org/10.2174/1568026622666220907112822>
- Shafabakhsh, R., Asemi, Z., 2019. Quercetin: a natural compound for ovarian cancer treatment. Journal of ovarian research. 12(1), 55. <https://doi.org/10.1186/s13048-019-0530-4>
- Shoaib, S., Khan, F.B., Alsharif, M.A., Malik, M.S., Ahmed, S.A., Jamous, Y.F., Uddin, S., Tan, C.S., Ardianto, C., Tufail, S., Ming, L.C., Yusuf, N., Islam, N., 2023. Reviewing the prospective pharmacological potential of isothiocyanates in fight against female-specific cancers. Cancers. 15(8), 2390. <https://doi.org/10.3390/cancers15082390>
- Shukla, S., Gupta, S., 2010. Apigenin: A promising molecule for cancer prevention. Pharmaceutical Research. 27(6), 962-978. <https://doi.org/10.1007/s11095-010-0089-7>
- Silva, M.A.M., Oliveira, M.C., Santos, R., 2022. Antitumor properties of curcumin in breast cancer based on preclinical studies: A systematic review. Cancers. 14(9), 2165. <https://doi.org/10.3390/cancers14092165>
- Simon, S.E., Lim, H.S., Jayakumar, F.A., Tan, E.W., Tan, K.O., 2022. Alpha-mangostin activates moap-1 tumor suppressor and mitochondrial signaling in mcf-7 human breast cancer cells. Evidence-based complementary and alternative medicine : eCAM. 2022, 7548191. <https://doi.org/10.1155/2022/7548191>
- Sohel, M., Aktar, S., Biswas, P., Amin, M.A., Hossain, M.A., Ahmed, N., Mim, M.I.H., Islam, F., Mamun, A.A., 2023. Exploring the anti-cancer potential of dietary phytochemicals for the patients with breast cancer: A comprehensive review. Cancer medicine. 12(13), 14556-14583. <https://doi.org/10.1002/cam4.5984>
- Song, X., Zhang, M., Dai, E., Luo, Y., 2019. Molecular targets of curcumin in breast cancer (review). Molecular medicine reports. 19(1), 23-29. <https://doi.org/10.3892/mmr.2018.9665>
- Song, Y., Feng, T., Zhang, Q., 2020. Hesperetin derivatives: synthesis and anticancer activities. European Journal of Medicinal Chemistry. 188, 112020. <https://doi.org/10.1016/j.ejmech.2019.112020>
- Sudhakaran, M., Navarrete, T.G., Mejía-Guerra, K., Mukundi, E., Eubank, T.D., Grotewold, E., Arango, D., Doseff, A.I., 2023. Transcriptome reprogramming through alternative splicing triggered by apigenin drives cell death in triple-negative breast cancer. Cell death & disease. 14(12), 824. <https://doi.org/10.1038/s41419-023-06342-6>
- Ting, Y., Jiang, Y., 2020. Challenges and limitations of silibinin in breast cancer therapy: Insights from molecular mechanisms and clinical trials. Pharmaceuticals. 13(10), 297. <https://doi.org/10.3390/ph13100297>
- Tuli, H.S., Tuorkey, M.J., Thakral, F., Sak, K., Kumar, M., Sharma, A.K., Sharma, U., Jain, A., Aggarwal, V., Bishayee, A., 2019. Molecular mechanisms of action of genistein in cancer: Recent advances. Frontiers in pharmacology. 10, 1336. <https://doi.org/10.3389/fphar.2019.01336>
- Ubaid, M., Salauddin, Shadani, M.A., Kawish, S.M., Albratty, M., Makeen, H.A., Alhazmi, H.A., Najmi, A., Zoghebi, K., Halawi, M.A., Ali, A., Alam, M.S., Iqbal, Z., Mirza, M.A., 2023. Daidzein from dietary supplement to a drug candidate: An evaluation of potential. ACS omega. 8(36), 32271-32293. <https://doi.org/10.1021/acsomega.3c03741>
- van Breemen, R.B., 2019. Development of effective cancer chemopreventive agents: A perspective from the nci's cancer prevention research program. Pharmaceuticals. 12(1), 36. <https://doi.org/10.3390/ph12010036>
- Vargas-Castro, R., García-Becerra, R., Díaz, L., Avila, E., Ordaz-Rosado, D., Bernadez-Vallejo, S.V., Cano-Colín, S., Camacho, J., Larrea, F., García-Quiroz, J., 2023. Enhancing tamoxifen therapy with α-mangostin: Synergistic antiproliferative effects on breast cancer cells and potential reduced endometrial impact. Pharmaceuticals. 16(11), 1576. <https://doi.org/10.3390/ph16111576>
- Vasconcelos, A.G., Valim, M.O., Amorim, A.G.N., do Amaral, C.P., de Almeida, M.P., Borges, T.K.S., Socodato, R., Portugal, C.C., Brand, G.D., Mattos, J.S.C., Relvas, J., Plácido, A., Eaton, P., Ramos, D.A.R., Kückelhaus, S.A.S., Leite, J.R.S.A., 2020. Cytotoxic activity of poly-ε-caprolactone lipid-core nanocapsules loaded with lycopene-rich extract from red guava (psidium guajava l.) on breast cancer cells. Food research international. 136, 109548. <https://doi.org/10.1016/j.foodres.2020.109548>

- [.org/10.1016/j.foodres.2020.109548](https://doi.org/10.1016/j.foodres.2020.109548)
- Wang, X., Yang, Y., An, Y., Fang, G., 2019. The mechanism of anticancer action and potential clinical use of kaempferol in the treatment of breast cancer. *Biomedicine & pharmacotherapy*. 117, 109086. <https://doi.org/10.1016/j.biopha.2019.109086>
- Wang, Z., Yu, B.W., Rahman, K.M., Ahmad, F., Sarkar, F.H., 2008. Induction of growth arrest and apoptosis in human breast cancer cells by 3,3-diindolylmethane is associated with induction and nuclear localization of p27kip. *Molecular cancer therapeutics*. 7(2), 341-349. <https://doi.org/10.1158/1535-7163.MCT-07-0476>
- Wink, M., 2015. Modes of action of herbal medicines and plant secondary metabolites. *Medicines (Basel, Switzerland)*. 2(3), 251-286. <https://doi.org/10.3390/medicines2030251>
- Wulandari, A.A., Choiri, A.A., Fitria, Widiandani, T., 2021. Thymoquinone and its derivatives against breast cancer with her2 positive: in silico studies of admet, docking and qspr. *Journal of basic and clinical physiology and pharmacology*. 32(4), 393-401. <https://doi.org/10.1515/jbcpp-2020-0431>
- Yap, K.M., Sekar, M., Wu, Y.S., Gan, S.H., Rani, N.N.I.M., Seow, L.J., Subramaniam, V., Fuloria, N.K., Fuloria, S., Lum, P.T., 2021. Hesperidin and its aglycone hesperetin in breast cancer therapy: A review of recent developments and future prospects. *Saudi journal of biological sciences*. 28(12), 6730-6747. <https://doi.org/10.1016/j.sjbs.2021.07.046>
- Yi, X., Zuo, J., Tan, C., Xian, S., Luo, C., Chen, S., Yu, L., Luo, Y., 2016. Kaempferol, a flavonoid compound from *Gynura medica* induced apoptosis and growth inhibition in mcf-7 breast cancer cell. *African journal of traditional, complementary, and alternative medicines : AJTCAM*. 13(4), 210-215. <https://doi.org/10.21010/ajtcam.v13i4.27>
- Younas, M., Hano, C., Giglioli-Guivarc'h, N., Abbasi, B.H., 2018. Mechanistic evaluation of phytochemicals in breast cancer remedy: current understanding and future perspectives. *RSC advances*. 8(52), 29714-29744. <https://doi.org/10.1039/c8ra04879g>
- Zhang, H., Zhang, M., Yu, L., Zhao, Y., He, N., Yang, X., 2012. Antitumor activities of quercetin and quercetin-5',8-disulfonate in human colon and breast cancer cell lines. *Food and chemical toxicology*. 50(5), 1589-1599. <https://doi.org/10.1016/j.fct.2012.01.025>
- Zhang, M., Holman, C.D., Huang, J.P., Xie, X., 2007. Green tea and the prevention of breast cancer: a case-control study in southeast china. *Carcinogenesis*. 28(5), 1074-1078. <https://doi.org/10.1093/carcin/bgl252>
- Zhu, J., Li, Q., Wu, Z., Xu, Y., Jiang, R., 2024. Curcumin for treating breast cancer: A review of molecular mechanisms, combinations with anticancer drugs, and nanosystems. *Pharmaceutics*. 16(1), 79. <https://doi.org/10.3390/pharmaceutics16010079>
- Zhu, L., Xue, L., 2019. Kaempferol suppresses proliferation and induces cell cycle arrest, apoptosis, and dna damage in breast cancer cells. *Oncology research*. 27(6), 629-634. <https://doi.org/10.3727/096504018X15228018559434>