



# Flavonoids, Breast Cancer Prevention, and Its Treatment: A Growing Evidence

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## Authors' contributions

This work was carried out in collaboration among all authors. Author MRA reviewed the literature and wrote the manuscript. Authors SJ, SH, MMI and NI cooperated in writing the manuscript. Authors SAI and HBS reviewed the manuscript. All authors read and approved the final manuscript.

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## ABSTRACT

There is mounting evidence linking certain lifestyle factors such as food, weight, and physical activity to an increased risk of breast cancer. Flavonoids are commonly used in traditional medicine. Plenty of studies have investigated the relationship between flavonoid consumption and breast cancer in humans. This review aimed to examine the association between flavonoids, each flavonoid subclass and the risk of breast cancer besides therapeutic use of flavonoids to break the multidrug resistance in breast cancer. Prospective cohort, case-control, and laboratory-based

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studies published between around 1990's to date and referred to the impact of flavonoids on breast cancer prevention, treatment, or other roles were included. Odd Ratio (OR)/ Risk Ratio (RR) and Hazard ratio (HR) along 95% confidence interval (CI) were carefully reviewed to reveal the association between different subgroups of flavonoids and breast cancer risk. Other adjustments (e.g. age, menopausal status, food habits, race, BMI, etc.) were also considered. The antioxidant properties of flavonoids, as well as their ability to inhibit apoptosis, suppress estrogen activity, and limit the proliferation of breast cancer cells, all contribute to a significant reduction in the risk of developing breast cancer. There are linings enough that some polyphenolic compounds are effective as drugs for treating breast cancer.

**Keywords:** Polyphenolic compounds; mammary carcinogenesis; antioxidants; multi-drug resistance (MDR); phytoestrogen; CYP1A1 inhibitors; ATP-binding cassette.

## 1. INTRODUCTION

Breast cancer (or mammary carcinogenesis) is the most frequently diagnosed cancer in women worldwide. It has emerged as a growing global health concern in recent decades (Ganesan *et al.*, 2022). According to GLOBOCAN (2020), the prevalence of new instances of breast cancer among women is expected to be 24.5% (Sung *et al.*, 2021). The number of women who were diagnosed with breast cancer worldwide reached 2.3 million in the year 2020, leading to 685,000 confirmed deaths. With 7.8 million confirmed cases in the previous five years, breast cancer exceeded all other cancers in terms of prevalence by the end of 2020 (WHO, 2023). Breast cancer is the consequence of cumulative exposure of the mammary cells to endogenous estrogens (Colditz, 1998). A wide variety of biological processes, including cell proliferation, cancer progression, apoptosis, and others, are made easier as a result of the interaction between estrogen and the estrogen receptor that is present in breast cells. This phenomenon is the result of this interaction (Thomas and Gustafsson, 2011).

Diet is the most pronounced modifiable factor for breast cancer occurrence, recurrence, and mortality (Hamer and Warner, 2017). Several studies in the last few decades have investigated the relationship between breast cancer and certain foods, such as beans, tomatoes, strawberries, grapes, meat, soy products, and a variety of vegetables, fruits, and teas (Peterson and Dwyer, 1998). Jang *et al.* in 2018 reported an association between anti-inflammatory diets and cancer recurrence as well as overall mortality in breast cancer patients. Experimental evidence suggests that flavonoids can both inhibit and stimulate a wide array of enzyme systems found in mammals. Enzymes like these

have an impact on processes including cell proliferation and division, detoxification, platelet aggregation, inflammation, and immunological responses (Franke *et al.*, 1998). Flavonoids have gone under numbers of investigations for their anti-carcinogenic mechanisms and antiproliferative effects on human lymphocytes as well as breast cancer cells and have also been reported (Peterson *et al.*, 2003). Laboratory and animal studies suggest that dietary flavonoids may be protective against breast cancer risk. An increased consumption of fruits and vegetables rich in flavonoids has been linked to a decreased risk of cancer, according to data from epidemiological studies. Which bioactive chemicals, if any, are responsible for this correlation remains unknown. This review aims at a clear idea about the 1990-to-date study pattern, interest in flavonoid subclasses, and major findings in both epidemiological and laboratory-based studies.

### 1.1 What is Breast Cancer?

Cancer is a cluster of abnormally grown/growing cells in a tissue. This anomaly is caused by a genetic mutation or other environmental factors that let them divide/grow out of control. These out-of-control cells form ball-shaped lumps of mutated tissue which is termed as tumor(s). When these forms of tumors are from breasts, they're classified as breast cancer. Most breast cancers begin in the milk glands (lobules) or the tubes (ducts) by which the nipple is connected to the milk glands. According to WHO (2023), the female gender has come up as the strongest risk factor for the occurrence of breast cancer. But in males, 1% or less of breast cancer cases are occurred. The main variables that increase the likelihood of breast cancer, both in terms of incidence and mortality, include being older (40 or older), having a higher body mass index

(BMI), smoking, not being physically active, eating a diet rich in fat, experiencing menstruation at a young age, having late in first full-term pregnancy, breastfeeding for shorter periods or not at all, using oral contraceptives, using menopausal hormone therapy, breast density, and genetic predisposition (Zhang *et al.*, 2020). A woman's chance of getting breast cancer is increased by both behavioral and genetic factors. Signs and symptoms of breast cancer include; thickening or lump in the breast, sometimes without pain; change in overall appearance (size and shape) of the breast; creek, reddish appearance, pit, or other changes in the skin; change in the physiognomy of nipples or areola (the darkish area around the breast); and abnormal fluid (may be bloody) from the nipple (WHO, 2023).

### 1.2 Molecular Subtypes of Breast Cancer

Physiologically, the human female mammary gland is under the primary control of different hormones. It is established that among them, estrogen appears to play the central role. There are mainly four molecular subtypes of breast cancer, characterized by hormone receptors (HR) in addition to protein involvement (or not involvement) in each cancer: (I) Luminal A or HR+/HER2- (HR-positive/HER2-negative), (II) Luminal B or HR+/HER2+ (HR-positive/HER2-positive), (III) HER2 enriched and (IV) Triple-negative (TNBC) or basal-like (HR/HER2-negative) (Fragomeni *et al.*, 2018). Each of the subtypes of breast cancer is described in Table 1, which includes their characteristics.

### 1.3 Cancer Growth and Clinical Stages

The female breast consists of 15 to 20 lobes of tissue, each of which contains lobules that house milk-producing glands and ducts. The nipple receives milk from the lobules via the ducts. Cancers can begin anywhere. However, the lymph nodes are often the sites of spreading after beginning in the ducts or lobules. Lymph nodes are like bus stations for cancer cells; they transport the disease throughout the body. There are many lymph nodes close to the breasts-around the chest, neck, and armpits. Cancer can spread regionally to these nodes from the breasts and referred as regionally spreading. Metastatic breast cancer spreads beyond those nodes. During diagnosis with breast cancer, cancer is categorized termed as "staged". Staging lets healthcare providers to make decision on treatment mechanisms and conditions after treatment. There are three distinct categories used to classify breast cancer stages: clinical prognostic, pathologic, and anatomic.

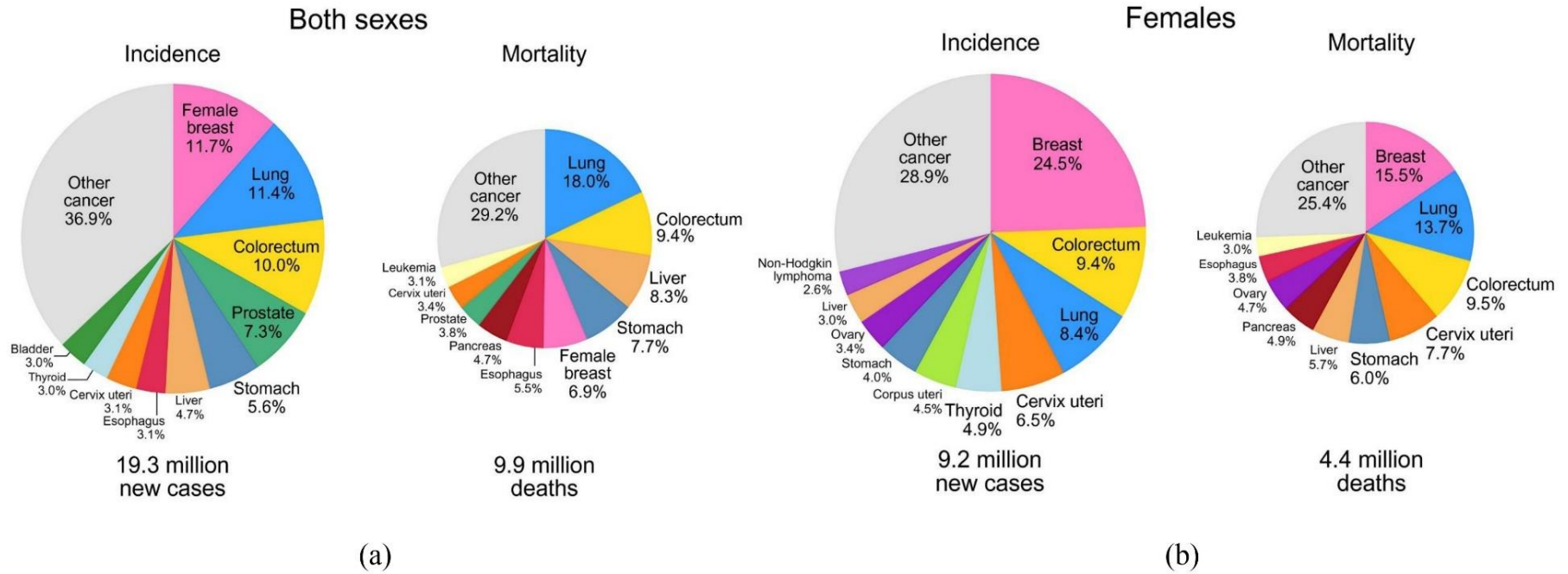
### 1.4 Breast Cancer Scenario Worldwide

Taking aside lung cancer from the title of "most commonly diagnosed cancer globally", breast cancer has emerged as the most prevalent cancer type. The year 2020 had an estimated 2.3 million cases and 685,000 fatalities due to breast cancer and among eight cancer diagnoses one cancer case was diagnosed as breast cancer case (Sung *et al.*, 2021). Over 2.3 million new instances of breast cancer are added annually. As more and more instances have been reported, breast cancer has outpaced all others among adults.

**Table 1. Distinctive characteristics of the various subtypes of breast cancer**

Attribute	Luminal A	Luminal B	HER2	TNBC	References
Frequency (%)	50	15	20	15	Barzaman <i>et al.</i> , 2020
ER	Yes	Yes	Some cases	No	Gao and Swain, 2018
PR	Yes	Some cases	Some cases	No	Barzaman <i>et al.</i> , 2020
HER	No	No	Yes	No	Engel and Kaklamani, 2007
Mutations	No	BRCA2	p53	p53 and BRCA1	Duffy <i>et al.</i> , 2018
Prognosis	Good	Middle	Middle/Bad	Bad	Ahern <i>et al.</i> , 2014
Therapy	Hormonal	Hormonal /Chemo	Hormonal/ Chemo/ Herceptin	Chemo/ Experimental	Loibl and Gianni, 2017

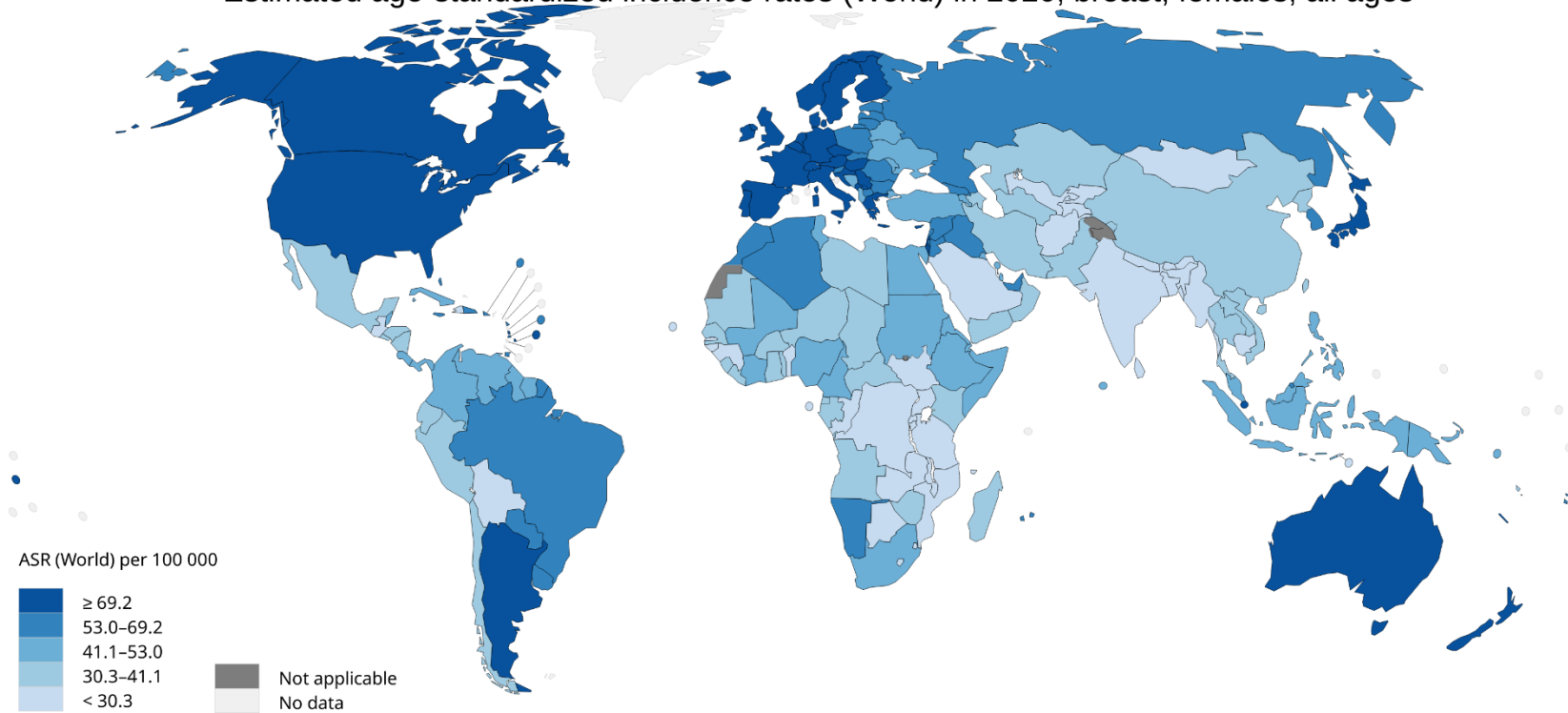
ER: estrogen receptor, PR: progesterone receptor, HER: human epidermal receptor, HER2: human epidermal growth factor receptor 2, and TNBC: triple-negative breast cancer



**Fig. 1. Death and case distributions for the 10 most frequent malignancies in 2020;(a) Both sexes and (b) Females. The percentage of cases or deaths is shown by the area of the pie chart. The "other" group includes nonmelanoma skin cancers, which do not include basal cell carcinoma for incidence**

Data source: GLOBOCAN 2020. Pie chart: Sung et al., 2021

### Estimated age-standardized incidence rates (World) in 2020, breast, females, all ages



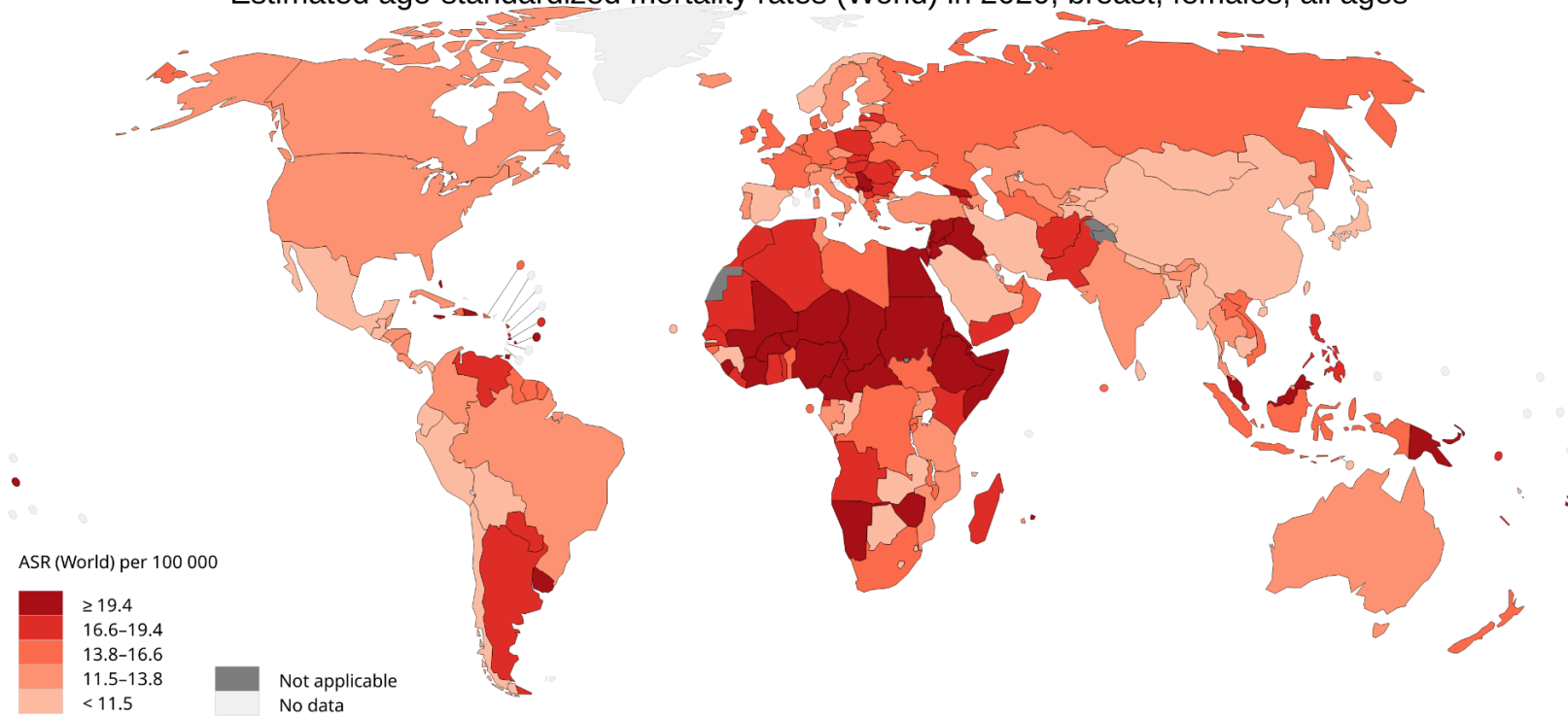
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Data source: GLOBOCAN 2020  
Map production: IARC  
(<http://gco.iarc.fr/today>)  
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**Fig. 2. Estimated age-standard female breast cancer incidence rates in 2020**  
*Data source: GLOBOCAN 2020, Map production: IARC, World Health Organization*

### Estimated age-standardized mortality rates (World) in 2020, breast, females, all ages



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Data source: GLOBOCAN 2020  
Map production: IARC  
(<http://gco.iarc.fr/today>)  
World Health Organization

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**Fig. 3. Estimated age-standard female breast cancer mortality rates in 2020**  
*Data source: GLOBOCAN 2020, Map production: IARC, World Health Organization*

More than 95% of countries have identified breast cancer as either the first or second major cause of mortality among females due to cancer (WHO, 2023). The majority of countries throughout the world have breast cancer as their primary cause of mortality and morbidity in the year 2020. Breast cancer was responsible for more than a quarter of all cancer diagnoses in women and approximately 15% of all deaths that were caused by cancer (Sung et al., 2021). Low and middle-income nations account for almost 80% of breast cancer and cervical cancer deaths because of their larger populations (WHO, 2023). By 2070, the anticipated number of breast cancer cases will have risen to 4.4 million (Soerjomataram and Bray, 2021).

### 1.5 Flavonoids, Classification, and Distribution

Flavonoids, whose name derives from the Latin word "flavus" signifying yellow, are ubiquitous secondary metabolites found in plants. They are most recognizable as the anthocyanin pigments,

which are red, blue, and purple in color, that cover plant tissues (Winkel-Shirley, 2001). Flavonoids generally consist of a 15-carbon skeleton comprising two benzene rings joined by a 3-carbon linkage chain (Nabavi et al., 2018). They are therefore shown as C6-C3-C6 compounds. The production of these compounds is accomplished by two distinct pathways: the shikimate pathway and the acetate pathway (Nabavi et al., 2018). Over six thousand distinct flavonoids have been identified. These chemicals are found in many different types of plants and serve multiple purposes, including protecting plants from various biotic and abiotic challenges, acting as signal molecules, detoxifying the body, and fighting microbes (Liu et al., 2021). The degree of oxidation, the connecting location of ring B, the annularity of ring C, and the position of hydroxy groups in the carbon skeleton all contribute to the classification of flavonoids into several subgroups, which include isoflavones, flavones, flavonols, flavanones, flavanonols, and flavones (Wang et al., 2018). Because of their extraordinary health advantages, foods rich in

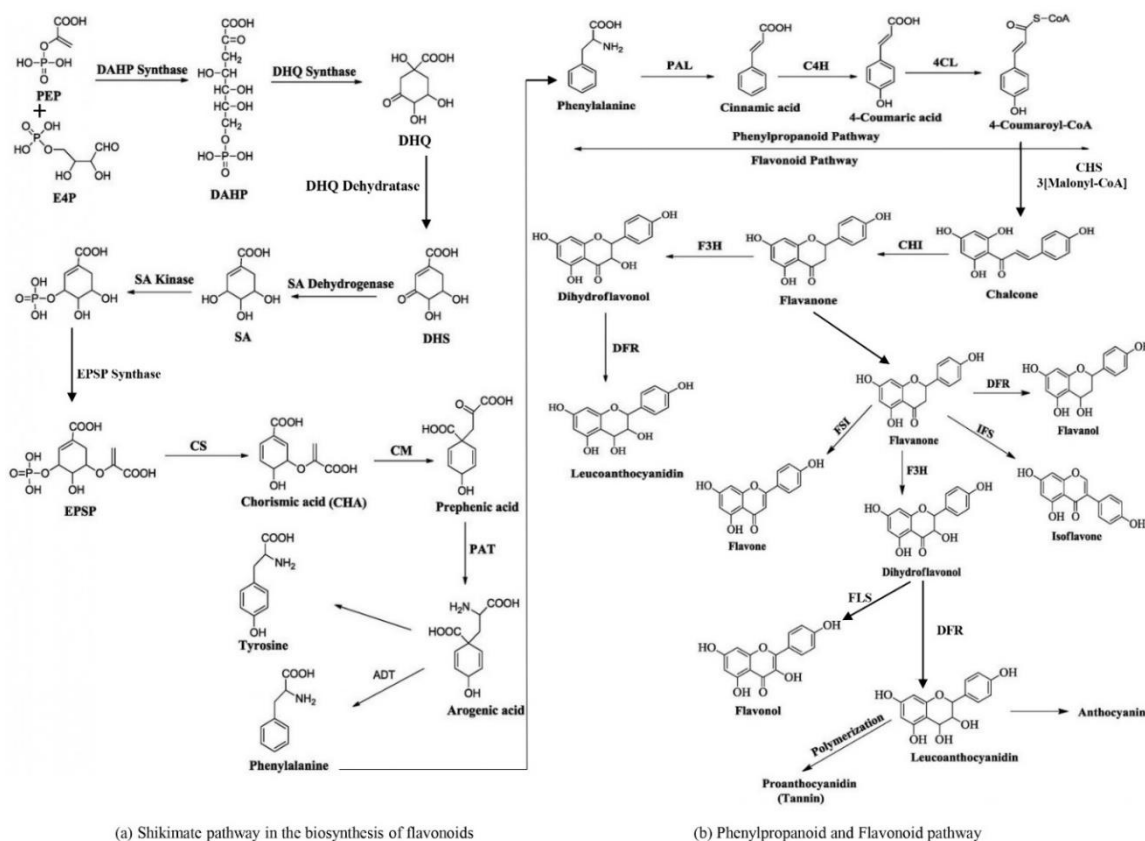


Fig. 4. Biosynthesis of various classes of flavonoids through the phenylpropanoid and shikimate pathways (Rehan, 2021)



flavonoids are called superfoods. Beverages, fruits, vegetables, cereals, beans, nuts, and tea make up the vast majority of these plant-based items (Harborne and Williams, 2000). Westerners get most of their flavonoids from wine, while Easterners get most of theirs from tea. Green vegetables, onions, apples, berries, cherries, soybeans, and citrus fruits are also believed to have a high concentration of dietary flavonoids (Butt et al., 2014).

## 1.6 Flavonoid Biosynthesis in Plants

The shikimate, phenylpropanoid, and polyketide pathway is the main pathway for higher plant species to synthesize flavonoids. These processes lead to the formation of numerous different types of flavonoids, with chalcones and flavanones serving as intermediates (Rehan, 2021). Plant species and sets of enzymes including hydroxylases, reductases, and isomerases are essential for this principal process to synthesis different forms of flavonoid skeletons. The flavanone is the progenitor of many flavonoid groups, including anthocyanins, isoflavonoids, and proanthocyanidins, which are condensed tannins (Fig. 4). Typically, the phenylpropanoid pathway is where flavonoid synthesis gets started. Following the completion of the shikimate pathway, the phenylpropanoid pathway eventually starts. Erythrose 4-phosphate and phosphoenol pyruvate condense, which is the first step in the shikimate pathway.

## 2. METHODOLOGY

**Selection of articles:** Article search was performed in PubMed, Google Scholar and NCBI using combinations of keywords relating to “breast cancer” AND (“dietary flavonoids” OR “flavonoids rich foods” OR “flavonols” OR “flavonones” OR “anthocyanidins” OR “flavones” OR “isoflavones” OR “soy flavonoids” OR “quercetin” OR “flavonoid subclasses” OR “polyphenolic compounds” OR “flavonoids rich foods” OR “flavonoids rich fruits and vegetables”) AND (“preventive role” OR “treatment” OR “flavonoids in drug” OR “flavonoids as chemoprotective agent”) etc. Only original articles both in epidemiological and laboratory-based studies were sorted out for this review.

**Inclusion criteria:** The eligible criteria included studies in the English language published between around 1990's to date. Prospective cohort, case-control, and laboratory-based studies that referred to the impact of flavonoids

on breast cancer prevention, treatment, or other roles were included. Titles and abstracts were independently screened. Hence, articles that linked flavonoids to (a) breast cancer incidence, (b) breast cancer-specific mortality, (c) the role of flavonoids, (d) therapeutic use of flavonoids, (e) multidrug resistance, etc. are included. The menopausal phase, cancer subtype, and type of anti-cancer medication received by patients were not subject to any restrictions.

**Data Extraction:** Study characteristics were recorded in Excel spreadsheet as follows: (1) name of the first author and publication year; (2) country or origin; (3) study design; (4) length of follow-up; (5) total cases and controls; (7) exposures to flavonoids; (8) way of flavonoids intakes; (9) Method and cell lines used and (10) RR/HR/OR from the most fully adjusted model for the highest versus the lowest flavonoids exposure and their 95% CI; (10) potential confounders and (11) limitations of studies.

**Retrieval of information:** Odd Ratio (OR)/ Risk Ratio (RR)/ Hazard ratio (HR) along 95% CI and p-value were carefully reviewed to reveal the association between different subgroups of flavonoids and breast cancer risk. Other adjustments (e.g. age, menopausal status, food habits, education level, race, BMI, multi-vitamin use, smoking habit, energy intake, age at first live birth, physical activity level, lifetime lactation, etc.) were also considered.

**How flavonoids fight against breast cancer:** Flavonoids, a plant source of polyphenolic compounds have grown its evidence from the 1990's to date that can fight against breast cancer.

### 1. Flavonoids as antioxidants

Flavonoids demonstrate their antioxidant action in a variety of ways, including activating antioxidant enzymes, scavenging reactive oxygen species (ROS), metal-chelating activity etc. Enzymes including catalase, glutathione peroxidase (GPX), and superoxide dismutase (SOD) constitute the backbone of this antioxidant defense system and deactivate free radicals generated by metabolic processes. (Jeeva et al., 2015). In 1999, Nacaia et al. (1999) demonstrated that flavonoids' intracellular antioxidative effect is dependent on their interaction with GSH-PO, at least in cells expressing the enzyme, and concluded that quercetin and catechin-activate glutathione peroxidase (GSH-PO) clearly. Superoxide



dismutase (SOD) levels were increased and glutathione peroxidase (GPx) levels were potentially decreased upon intraperitoneal dose of catechins.

Flavonoids, being polyphenols, have a high reactivity due to the hydroxyl substituents, which makes them effective at scavenging free radicals through a process called hydrogen atom abstraction (Korkina, and Afanas'Ev, 1997). Flavonoids possess the ability to effectively capture the highest levels of active solar wavelengths (namely, UV-B and UV-A). Not only that, but they are equally adept at halting the creation of ROS and removing them once they've formed (Agati *et al.*, 2012). Troxerutin, a flavonoid, exhibited its cytoprotective efficacy by protecting various cell types (intestinal epithelial cells, fibroblasts, and lymphocytes) against oxidative stress (Panat *et al.*, 2016). The antioxidant properties of silibinin and quercetin stem from their capacity to bind to iron or copper ions and inhibit oxidases, in addition to scavenging free radicals and other oxidizing chemicals (de Groot and Rauen, 1998). Diniz *et al.* in 2015 suggested that flavonoids have the potential for neuroprotection in epilepsy.

Rutin and quercetin were found to be more effective in suppressing iron ion-dependent lipid peroxidation systems due to their ability to form stable compounds with iron ions, which are incapable of initiating lipid peroxidation (Afanas'Ev *et al.*, 1989). Apigenin, luteolin, kaempferol, quercetin, myricetin, and rutin, isoflavones (daidzein and genistein), flavanones (taxifolin, naringenin, and naringin; and catechin) were found to have a greater ability to reduce copper ions compared to iron ions (Mira *et al.*, 2002).

## 2. Flavonoids as phytoestrogens:

High consumption of phytoestrogens can result in increased amounts of phytoestrogens in the bloodstream, which have been empirically shown to exhibit estrogen-like actions. Breast cancer risk is lower in women who regularly consume phytoestrogens, especially throughout childhood (Lee *et al.*, 2009). Korde *et al.* in 2009 also concluded a correlation between breast cancer risk reduction with higher soy consumption during childhood, adolescence, and adulthood where the most potent protective effect was shown in childhood. According to Rice and Whitehead (2006), phytoestrogens have the ability to change estrogen production by inhibiting aromatase (known as cytochrome P450

19 (Cyp19)) as well as 17 $\beta$ -hydroxysteroid dehydrogenases (HSD), estrone sulfatase, and sulfotransferase. In breast tissue, these enzymes play a role in the generation of estradiol, and when they are overexpressed, they are associated with an increased risk of breast cancer. Research has shown that a number of flavonoids can significantly lower estrogen levels in the blood. They may be useful in treating malignancies that are resistant to other treatments, even though their efficacy is lower than that of steroidal aromatase inhibitors often employed in clinical practice.

## 3. Flavonoids as CYP1A1 inhibitors/substrates

Inhibiting CYP1A is a plausible approach to preventing the development of cancer. The selective metabolism of dietary flavonoids by CYP1 enzymes, resulting in the production of conversion products that limit the proliferation of cancer cells, is the key mechanism behind cancer prevention. Cytochrome P450 enzymes (CYP450s) are the primary enzyme group accountable for the metabolism and detoxification of foreign substances, which are commonly referred to as xenobiotics. They are also involved in the biosynthesis as well as metabolism of endogenous compounds like fatty acids and steroids. CYP450s in the liver where they are highly expressed catalyze reactions that are responsible for converting lipophilic compounds into more hydrophilic derivatives. Most drugs follow this biotransformation pathway to be catabolized and eliminated from the body (Gonzalez, 1988). But other xenobiotic substances, such as dietary flavonoids, can modulate the CYP450 enzyme function, either increasing or decreasing the catalytic activity. As a result, drug detoxification or functionality could be interrupted (Hodek *et al.*, 2002).

It has been established that the CYP1 family is involved in the process of carcinogenesis and the advancement of cancer, and demonstrated through several research that flavonoids, such as resveratrol, have the ability to effectively inhibit the activation of CYP1 in both in-vivo and ex-vivo environments because of their function as competitive antagonists for the aryl hydrocarbon receptor AhR, which plays a role in the activation of CYP1 expression (Ciolino *et al.*, 1999). For instance, quercetin and myricetin are potent CYP1B1 inhibitors, but they appear to have a lesser impact on the CYP1A1 and CYP1A2 variants (Arroo *et al.*, 2009).

#### 4. Flavonoids as ABC (ATP-binding cassette) transporter regulators

ATP-binding cassette superfamily comprises P-glycoprotein (P-gp), ABCG2, multidrug resistance proteins 1 and 2 (MRP1 and 2) and breast cancer resistance protein (BCRP). Several xenobiotic-related compounds rely on these membrane proteins for absorption, distribution, and excretion. These transporters are essential for the distribution of pharmaceuticals often used in medical therapy, and they are the principal family responsible for removing drugs from cells (Tan *et al.*, 2013). In recent years, there has been a significant amount of focus placed on the role that ATP-binding cassette (ABC) transporters play role in the biology of cancer cells (Nobili *et al.*, 2020) and for this reason they are the subject of substantial investigation. The development of multidrug resistance, which is predominantly driven by factors related to ABC transporter genes, is the primary obstacle that stands in the way of effective cancer treatment. There are several examples of ATP-binding cassette (ABC) transporters that impart resistance to particular chemotherapy medications. Some of these transporters include breast cancer resistance protein (BCRP), P-glycoprotein (P-gp), and multidrug resistance-associated protein 1 (MRP1). Intestinal efflux ABC (ATP binding cassette) transporters, some of which include P-glycoprotein, breast cancer resistance protein, and multidrug resistance-related proteins, perform the function of "pumping doors" to regulate the elimination of flavonoids from intestinal epithelial cells into either the intestinal cavity or the systemic circulation. Breast cancer resistance protein (BCRP) is an ATP-binding cassette transporter that has recently come to light due to the report that it can regulate drug disposition and induce multidrug resistance (MDR) to certain important anticancer treatments. Flavonoids serve as inhibitors of transporters and restrict the ATP hydrolysis activity that leads to the elimination of anticancer drugs from tumor tissues (Zhang *et al.*, 2004). Presumably, this occurs through the interaction with the nucleotide-binding domain (NBD), which is the specific site of action for flavonoids in both BCRP and P-gp (Morris and Zhang, 2006). The complete understanding of the interaction between flavonoids and BCRP, despite their ability to effectively modify medication pharmacokinetics, remains incomplete.

#### 5. Flavonoids' impact on cell death, cell cycle reversal, and other signaling mechanisms

Flavonoids are well acknowledged for their significance as anti-carcinogenic substances and have shown remarkable efficacy as cytotoxic anti-cancer medicines, triggering apoptosis in most cancer cells. A number of flavonoids have shown promise in inducing cell death in cancers, including breast cancer (Huang *et al.*, 2012) via means of the two primary apoptotic pathways: internal apoptosis involving caspase-9 and mitochondria, and extrinsic apoptosis involving caspase-8 and death receptors (Kamsteeg *et al.*, 2003). Brusselmans *et al.* (2005) proposed the ability of flavonoids to inhibit fatty acid synthase (FAS) is closely linked to their potential to cause apoptosis in cancer cells.

#### 6. Effect of flavonoids on multi-drug resistance

The primary factor contributing to the failure of the majority of cancer chemotherapy is drug resistance. The fundamental cause of chemoresistance is believed to be the dysregulation of the epigenetic machinery. It was reported that the dysregulation of DNA methylation in resistant cancer cells leads to the abnormal expression of genes involved in cancer proliferation, apoptosis, DNA repair, and drug efflux. The dysregulation of many enzymes that catalyze histone post-translational modifications can also lead to changes in chromatin structure. Moreover, expression of numerous drug-resistance genes is regulated by this imbalance. Alterations in the patterns of microRNA play a role in the development of drug resistance in cancer cells (To Kenneth and Cho, 2021). Curcumin, stilbenes, ellagitannins, phenol carboxylic acids, and flavones are all examples of phytochemicals that possess antioxidant, anti-inflammatory, and anti-cell-growth activities. These features make them useful chemopreventive agents. In addition to this, they prevent the development of new blood vessels and the spread of cancer cells, regulate immune and inflammatory responses, and deactivate substances that can cause cancer. Furthermore, preclinical and clinical investigations have revealed that these drugs successfully change several pathways, which prevents cancer from developing multidrug resistance (Costea *et al.*, 2020). Researchers are now interested in the effect of flavonoids on drug resistance in chemotherapies. To overcome treatment

resistance in breast cancer, flavonoids have been suggested as a potential medication by many studies. According to Rao *et al.* (2012), luteolin effectively reverses multidrug resistance (MDR) in breast cancer cells that are resistant to mitoxantrone.

### 3. RESULTS AND DISCUSSION

Flavonoid consumption may have an association with a lower risk of cancer development, according to the findings of the cohort and case-control studies that were reviewed in this article (Table 2). The fact that flavonoids have been shown to have an inverse correlation with cancer risk may be explained by the fact that flavonoids have an effect on a number of critical biological activities. The capacity of flavonoids to scavenge free radicals has been characterized in experimental settings to a satisfactory degree.

More recently, flavonoids have been shown to influence the signal transduction pathway (Melnik *et al.*, 2002), accelerate apoptosis (Choi *et al.*, 2001), suppress inflammation and impede proliferation in human cancer cell lines (Manthey *et al.*, 2002). In vitro and animal model systems were used to publish these findings. Some flavonoids can also enhance the transcription of phase II detoxifying enzymes, which would help eliminate pro-carcinogenic substances. This would reinforce the idea that flavonoids prevent cancer. Flavonoids were shown to dramatically reduce the number of focal areas of dysplasia that were caused by exposure to azoxymethane (AOM). According to the findings of a study that employs mice that were subjected to treatment with azoxymethane (AOM) and fed either a standard diet, a standard diet plus rutin, or a standard diet plus quercetin (Yang *et al.*, 2000). This kind of research is noteworthy because it suggests that flavonoids could be related to a number of early-stage processes in the cascade that result in cancer development.

The development of multidrug resistance (MDR) in cancer cells through transporter-mediated active efflux of cytotoxic drugs is one of the processes that has been studied and characterized the most. Previous studies have demonstrated that BCRP (Breast Cancer resistance protein) bestows resistance upon several significant anticancer agents, including doxorubicin, topotecan, mitoxantrone, SN38, methotrexate, and flavopiridol (Doyle *et al.*, 1998). Therefore, to potentially reverse multidrug resistance (MDR), it is necessary to identify

inhibitors of BCRP that are both potent and nontoxic. Flavonoids, which have a lengthy history of human ingestion and a stellar safety record, are an essential component of the average diet (Havsteen, 2002). Furthermore, these compounds have been linked to numerous anticancer mechanisms (Havsteen, 2002), as well as synthetic compounds that are analogous to them, including flavone acetic acid and flavopiridol (Senderowicz *et al.*, 1998).

In addition to its role in regulating drug distribution, the ATP-binding cassette transporter breast cancer resistance protein (BCRP), which was only recently discovered, has been shown to impart multidrug resistance (MDR) to many important anticancer drugs. Flavonoids, a group of polyphenolic chemicals, are widely present in plant products and dietary sources. It has been reported that flavonoids interact with P-glycoprotein and multidrug resistance-associated protein 1; nevertheless, the nature of their interaction with BCRP remains unknown. The objective of this study was to evaluate the effects of naturally derived flavonoids on the cytotoxicity and cellular accumulation of mitoxantrone in human cell lines that exhibit BCRP expression but lack BCRP. Most of the reviewed investigations in this study utilized human breast cancer cells (MCF-7) that over expressed BCRP and large-cell lung carcinoma cells (NCI-H460) that lacked BCRP.

A considerable number of the flavonoids that were assessed (50  $\mu\text{M}$ ) exhibited an augmentation in mitoxantrone accumulation in cells that over expressed BCRP, while also effectively reversing the development of mitoxantrone resistance. It is important to note that these flavonoids did not have any effect on the BCRP-negative cells that corresponded to them, which suggests that they act as antioxidants.

The impact of these flavonoids on the cellular accumulation and cytotoxicity of mitoxantrone was found to be concentration-dependent, with the majority of the flavonoids exhibiting significant modifications at concentrations below 10  $\mu\text{M}$  (Hung *et al.*, 2015). Additional research is required to optimize the compounds' bioavailability, bio-distribution, and safety in preparation for future human studies. In particular, it is imperative to elucidate how every flavonoid, in accordance with its distinct chemical composition, can traverse the membrane.

**Table 2. Summary of epidemiological studies investigating the association of flavonoid intake with breast incidence and mortality**

Author(s)	Method	Subject(s)	Cancer type	Exposer	Conclusion
Zheng <i>et al.</i> , 1999	<ul style="list-style-type: none"> <li>• Case-control study</li> <li>• Structured questionnaire</li> <li>• Urinary excretion rates</li> </ul>	Sixty (60) breast cancer cases and individually matched controls examined for rates of excretion in urine. Face-to-face interviews conducted for 746 urbane Chinese women in Shanghai	Breast cancer	Total phenols and five major isoflavonoids (glycitein, daidzein, genistein, equol and O-desmethylangole nsin)	<ul style="list-style-type: none"> <li>• Breast cancer cases had lower urinary total phenols and all isoflavonoids.</li> <li>• The disparity between the case and control groups became more apparent when comparing the median levels of these compounds.</li> <li>• Individuals with the greatest levels of daidzein, glycitein, and total isoflavonoids had around 50% lower risk of cancer compared to those with the lowest levels.</li> </ul>
Horn-Ross <i>et al.</i> , 2001	<ul style="list-style-type: none"> <li>• Case-control study</li> <li>• Food Frequency Questionnaire (FFQ)</li> <li>• Ontario phytoestrogen database</li> <li>• Multivariate logistic regression</li> </ul>	1,326 cases and 1,657 controls non-Asian US women (35–79 years) residing in the San Francisco Bay Area	Breast cancer	Isoflavones, total isoflavones, lignans, total lignans, total phytoestrogens	<ul style="list-style-type: none"> <li>• Phytoestrogen was found not to be protective against breast cancer risk (odds ratio = 1.0 and 95% CI = 0.80, 1.3 for the highest versus lowest quartile).</li> </ul>
Peterson <i>et al.</i> , 2003	<ul style="list-style-type: none"> <li>• Case-control study in Greece</li> <li>• Semi-quantitative food frequency questionnaire</li> <li>• US Department of Agriculture-Iowa State University Database</li> </ul>	820 women diagnosed with breast cancer against 1548 control women	Breast cancer	Flavones, flavonols, flavanones, flavan-3-ols, isoflavones and anthocyanidins	<ul style="list-style-type: none"> <li>• A significant negative correlation between flavone consumption and breast cancer was observed. The odd ratio for an increment in daily flavone consumption equal to one standard deviation (0.5 mg/day) was 0.87 (95% CI = 0.77, 0.97).</li> </ul>
Silva <i>et al.</i> , 2004	<ul style="list-style-type: none"> <li>• Case-control study</li> <li>• Face-to-face interview with food frequency questionnaire (FFQ)</li> <li>• Conditional logistic regression models</li> </ul>	240 South Asians diagnosed with breast cancer residing in England against 477 population-based controls matched for age	Breast Cancer	Total isoflavones, (genistein, daidzein) as well as total lignans (secoisolariciresinol, matairesinol)	<ul style="list-style-type: none"> <li>• After controlling for known and established risk factors for breast cancer, there is moderate evidence of a dose-response relationship between isoflavone consumption and the likelihood of developing breast cancer (P value for trend 0.08).</li> </ul>

Author(s)	Method	Subject(s)	Cancer type	Exposer	Conclusion
Bosetti <i>et al.</i> , 2005	<ul style="list-style-type: none"> <li>• Case-control study</li> <li>• Standard structured questionnaire</li> <li>• Food Frequency Questionnaire (FFQ)</li> </ul>	2,569 women with histologically confirmed breast cancer and 2,588 hospital controls (age range 23-74 years, median age 55) from six Italian areas	Breast Cancer	Flavanones, flavones, anthocyanidins, flavan-3-ols, flavonols, and isoflavones	<ul style="list-style-type: none"> <li>• Increased intake of flavones reduced the risk of breast cancer (OR = 0.81 for the highest versus the lowest quintile with a P-trend of 0.02).</li> <li>• Significant association of risk reduction of breast cancer was not found for other flavonoids.</li> </ul>
Ha <i>et al.</i> , 2006	<ul style="list-style-type: none"> <li>• Cross-sectional study</li> <li>• Dietary questionnaire</li> </ul>	128 women aged 40–79 year and newly diagnosed with invasive breast cancer.	Breast cancer	Isoflavonoids and lignans	<ul style="list-style-type: none"> <li>• The chance of getting a cancer diagnosis at any stage other than stage 1 was 32% lower in women who consumed a higher level of phytoestrogens.</li> </ul>
Fink <i>et al.</i> , 2007	<ul style="list-style-type: none"> <li>• Case-control study</li> <li>• Interviewed about known as well as suspected risk factors</li> <li>• Food Frequency Questionnaire (FFQ)</li> </ul>	1,434 female cases (English-speaking and newly diagnosed with breast cancer) and 1,440 control. Both are from Long Island, New York, USA	Breast Cancer	Flavonols, flavan-3-ols, flavones, isoflavones, lignans anthocyanidins,	<ul style="list-style-type: none"> <li>• Flavonoid intake was associated with breast cancer risk reduction.</li> <li>• The reduction was most pronounced among postmenopausal women for flavonols (odds ratio (OR) ¼ 0.54 with 95% CI = 0.40, 0.73).</li> </ul>
Cotterchio <i>et al.</i> , 2008	<ul style="list-style-type: none"> <li>• Case-control study</li> <li>• Block food frequency questionnaire (FFQ) expanded to include foods containing phytoestrogen</li> </ul>	3,063 breast cancer cases (age range 25-74 years and diagnosed in 2002 and 2003) and 3430 controls were identified by using the Ontario Cancer Registry	Breast cancer	Lignans and isoflavones	<ul style="list-style-type: none"> <li>• Lignan intake was found to be effective in reducing breast cancer risk.</li> <li>• A significant reduction in breast cancer risk was observed in overweight pre-menopausal women who had a high consumption of phytoestrogens.</li> </ul>
Cutler <i>et al.</i> , 2008	<ul style="list-style-type: none"> <li>• Prospective cohort</li> <li>• Food frequency questionnaire</li> <li>• Food composition databases</li> <li>• Baseline questionnaire (1985) and 5 follow-up questionnaires (1987,</li> </ul>	34,708 postmenopausal women (55-69 years) in the Iowa Women's Health Study	Pancrease, lung, breast, colorectal and upper aerodigestive cancer	Seven flavonoid subclasses along with total flavonoids	<ul style="list-style-type: none"> <li>• Isoflavone intake and overall cancer incidence were inversely associated (HR= 0.93; 95% CI = 0.86–1.00).</li> </ul>

Author(s)	Method	Subject(s)	Cancer type	Exposer	Conclusion
	<ul style="list-style-type: none"> <li>1989, 1992, 1997, 2004)</li> <li>Followed for cancer incidence from 1986 through 2004</li> </ul>				
Torres-Sanchez et al., 2009	<ul style="list-style-type: none"> <li>Case-control study</li> <li>Semi quantitative food frequency questionnaire (FFQ)</li> <li>Logistic regression models</li> </ul>	A total of 198 women with breast cancer (age range 21–79 years) and age-matched control ( $\pm 3$ yr) for each case	Breast cancer	Onion, lettuce, tea, apple, and spinach	<ul style="list-style-type: none"> <li>The adjusted odds ratio was 0.27 (95% CI = 0.16, 0.47) with a statistically significant trend (<math>P &lt; 0.001</math>) for the consumption of more than one slice of onion per day.</li> <li>Consumption of lettuce and spinach had a substantial preventive impact.</li> </ul>
Pantavos et al., 2015	<ul style="list-style-type: none"> <li>Prospective cohort study</li> <li>Semi quantitative FFQ</li> <li>Ferric reducing antioxidant potential (FRAP)</li> <li>Follow-up of 17 years</li> <li>Crude and multivariate Cox proportional hazard models</li> </ul>	Women aged 55 years and older (n = 3,209) and 199 breast cancer cases, identified from the Rotterdam study	Breast cancer	Vitamin A, C and E, selenium, flavonoids and carotenoids	<ul style="list-style-type: none"> <li>No correlation between total antioxidant intake and the risk of breast cancer was found.</li> <li>A reduced consumption of flavonoids was linked to an increased risk of breast cancer in women aged 70 and above (HR = 1.80; 95% CI = 1.09, 2.99).</li> </ul>
Wang et al., 2014	<ul style="list-style-type: none"> <li>Prospective cohort study</li> <li>Self-administered questionnaires</li> <li>Modified Willett FFQ</li> <li>Cox proportional hazards regression</li> </ul>	56,630 postmenopausal women from the CPS-II Nutrition Cohort were included in the analytical cohort	Breast cancer	Total flavonoids, Anthocyanidins, flavan-3-ols, flavanones, flavones etc	<ul style="list-style-type: none"> <li>Total flavonoid intake was not associated with breast cancer risk.</li> <li>Despite this, the total risk of breast cancer was somewhat inversely related to flavone consumption.</li> </ul>
Feng et al., 2020	<ul style="list-style-type: none"> <li>Case-control study</li> <li>Food frequency questionnaire (Face-to-face)</li> <li>Multivariable logistic regression models.</li> </ul>	1522 breast cancer cases and 1547 frequency-matched controls (both cases and controls are selected Chinese population)	Breast cancer	Total flavonoids, Anthocyanidins, Flavanols, flavan-3-ol etc.	<ul style="list-style-type: none"> <li>After considering possible influencing factors, a clear inverse relationship was shown between the risk of developing breast cancer and the intake of various types of flavonoids,</li> </ul>

**Table 3. Summary of laboratory-based studies investigating the association of flavonoid intake with breast incidence and mortality**

Author	Cell line	Model	Flavonoids exposer	Conclusion
Miksicek, 1993	MCF7, HeLa, COS-7	Transfection studies, competition binding studies	Chalcones, flavanones, flavones, and isoflavones	<ul style="list-style-type: none"> <li>Selected hydroxylated flavonoids interacted directly with the estrogen receptor based on their ability to compete for binding <math>17\beta</math>-[3H]estradiol to the receptor in the free cell extracts.</li> <li>These substances exhibited lower activity, when measured on a molar basis, compared to <math>17\beta</math>-[3H]estradiol or the synthetic dihydroxystilbene estrogens.</li> </ul>
So <i>et al.</i> , 1996	Human breast carcinoma cell line, MDA-MB-435.	Proliferation assay, growth curve at $ic_{50}$ , MTT viability assay, tumorigenicity experiments	Two citrus flavonoids along with four noncitrus flavonoids	<ul style="list-style-type: none"> <li>Among the six individual flavonoids tested, baicalein had the highest potency to effectively suppress the proliferation of the cells.</li> <li>Hesperetin and naringenin (found in citrus fruits) exhibited moderate inhibitory effects.</li> </ul>
Zand <i>et al.</i> , 2000	BT-474 human breast cancer cells	ELISA-type immune fluorometric assays,	72 flavonoids and structurally related compounds.	<ul style="list-style-type: none"> <li>Flavonoids exhibited significant steroid hormone activity. Based on the findings they concluded that flavonoids may have an effect on cancer risk, prevention and cancer therapeutics.</li> </ul>
Imai <i>et al.</i> , (2004)	K562 (K562/BCRP), LLC/BCRP, K562/MDR, and KB/MRP Cell Lines	Cell growth inhibition assay, cellular [3H]genistein accumulation in K562/BCRP Cells, transcellular transport assay of [3H]genistein	Estrone, kaempferol, naringenin, naringenin-7-glucoside	<ul style="list-style-type: none"> <li>Phytoestrogens/flavonoids, including genistein, naringenin, acacetin, and kaempferol, boosted the toxicity of SN-38 and mitoxantrone in BCRP-transduced K562 (K562/BCRP) cells.</li> </ul>
Zhang <i>et al.</i> , 2004	MCF-7, NCI-H460	Western blot analysis of BCRP, P-gp, and MRP1, mitoxantrone accumulation studies, mitoxantrone cytotoxicity studies.	Apigenin, biochanin a, chrysin, daidzein, fisetin, morin, myricetin, genistein etc.	<ul style="list-style-type: none"> <li>Many of the tested flavonoids (50 M) are BCRP inhibitors.</li> <li>Chrysin and biochanin A exhibited the highest level of potency as inhibitors of BCRP.</li> </ul>
Katayama <i>et al.</i> , 2007	Human leukemia K562 cells and human epidermoid	Growth inhibition assay, topotecan uptake	Flavone, flavanol, flavanone, isoflavone, chalcone	<ul style="list-style-type: none"> <li>3,4,7-trimethoxyXavone demonstrated the most potent anti-BCRP activity for SN-38 and mitoxantrone respectively.</li> <li>3,4,7-trimethoxyXavone and acacetin, exhibited</li> </ul>



Author	Cell line	Model	Flavonoids exposer	Conclusion
	carcinoma KB-3-1 cells			minimal anti-P-gp activity, while the others did not demonstrate any inhibitory effects on P-gp.
Chen <i>et al.</i> , 2007	MDA-MB-231 breast cancer cell cultures and xenografts	Proteasomal chymotrypsin-like and caspase-3/caspase-7 activity assays, MTT assay, western blot analysis and human breast tumor xenograft experiments	Apigenin	<ul style="list-style-type: none"> <li>● Apigenin suppresses the proteasomal chymotrypsin-like activity and induces apoptosis in both cultured MDA-MB-231 cells and MDA-MB-231 xenografts.</li> </ul>
Yang <i>et al.</i> , 2012	MCF-7, MDA-MB-231, MCF-10A cells	MTT colorimetric assay, Luminescence ATP detection assay, Western blot analysis	Fisetin (3,3',4',7-tetrahydroxyflavone)	<ul style="list-style-type: none"> <li>● Fisetin exhibited greater cytotoxicity in MCF-7 human breast cancer cells compared to MDA-MB-231 cells.</li> <li>● Fisetin can trigger a novel atypical apoptosis in caspase-3-deficient MCF-7 cells.</li> </ul>
Shan <i>et al.</i> , 2013	MCF-7 and MDA-MB-231	MTT, clonogenicity assay, apoptosis assay, western blot analysis	Triticuside A from wheat bran	<ul style="list-style-type: none"> <li>● Triticuside A dose-dependently suppressed the growth of human breast cancer cells (MCF-7 and MDA-MB-231).</li> </ul>
Pradhan <i>et al.</i> , 2015	MCF-7 cell lines and MCF-7 xenografts	Proteasomal chymotrypsin-like and caspase-3/ caspase-7 activity assays, cell viability assay, xenograft experiments	Quercetin	<ul style="list-style-type: none"> <li>● Quercetin suppresses the proteasomal chymotrypsin-like activity and triggers apoptosis in both cultivated MCF-7 cell lines and MCF-7 xenografts.</li> </ul>
Vrhovac Madunic <i>et al.</i> , 2018	ER-positive MCF-7, triple-negative MDA MB-231	MTT assay, cell viability, comet and lipid peroxidation assays,	Apigenin	<ul style="list-style-type: none"> <li>● Apigenin caused cell death in both cell lines resulting in significant toxicity and apoptosis.</li> <li>● No significant cytogenotoxic effects were detected in normal cells.</li> </ul>
Santes-Palacios <i>et al.</i> , 2020	<i>Escherichia coli</i> DH5 $\alpha$ cells	Ethoxyresorufin O-deethylase activity (EROD) assay, bacterial mutagenicity assay	Quercetin, myricetin, luteolin, fisetin, kaempferol, 5-hydroxyflavone, flavone etc.	<ul style="list-style-type: none"> <li>● 5-hydroxyflavone, 3 hydroxy-flavone and flavone exerted the most potent inhibitory activity with IC50 values of 0.07, 0.10 and 0.08 <math>\mu</math>M respectively.</li> </ul>

Based on the available research, it may be concluded that flavonoids have a negative correlation with the chance of developing cancer. Further research utilizing up-to-date dietary databases with HPLC results for flavonoid estimations from food will offer conclusive information about the correlation between plant chemicals and the risk of cancer. Furthermore, conducting feeding experiments that can accurately describe the process of how plant flavonoids are absorbed, metabolized, and eliminated from the body, as well as their interaction with important enzyme systems, would be crucial in comprehending the underlying biological reasons for the negative relationship between flavonoid consumption and the risk of developing cancer. Meanwhile, it is strongly recommended that the public adhere to dietary recommendations for cancer prevention, which involve consuming five or more servings of fruits and vegetables daily. Moreover, the findings of the researcher might offer significant backing for additional exploration into evaluating the effectiveness and safety of flavonoid chemicals whether included in a nutritious diet or used as a supplementary treatment for human malignancies.

#### 4. LIMITATIONS

This article only reviewed some of open access articles available in online. This narrowed down the number of articles reviewed. Some articles reviewed in this study may not adequately account for all confounding factors that could influence cancer risk. Few studies showed negative finding in the context of breast cancer prevention and treatment and need more investigation.

#### 5. CONCLUSION

There is evidence that dietary flavonoids can lower the risk of cancer, especially breast cancer. This review briefly discussed about that the different sources of flavonoid and their subclasses which are the polyphenolic compounds mainly present in plant sources like fruits, vegetables, and cereals. They mainly reduce the risk of breast cancer by antioxidant activity, anti-carcinogenic effect, inhibiting apoptosis, reducing estrogen activity, and inhibiting the growth of breast cancer cells. Researchers are enthusiastic about revealing the therapeutic properties of flavonoids to treat breast cancer cases. There are linings enough that some polyphenolic compounds are found as effective drugs for treating breast cancer.

#### CONSENT AND ETHICAL APPROVAL

It is not applicable.

#### DISCLAIMER (ARTIFICIAL INTELLIGENCE)

Author(s) hereby declare that NO generative AI technologies such as Large Language Models (ChatGPT, COPILOT, etc.) and text-to-image generators have been used during the writing or editing of this manuscript.

#### COMPETING INTERESTS

Authors have declared that no competing interests exist.

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