Fatima, Nishat; Baqri, Syed Shabihe Raza; Alsulimani, Ahmad; Fagoonee, Sharmila; Slama, Petr; Kesari, Kavindra Kumar; Roychoudhury, Shubhadeep; Haque, Shafiul

Phytochemicals from Indian ethnomedicines

Published in: Antioxidants

DOI: 10.3390/antiox10101606

Published: 01/10/2021

Document Version
Publisher's PDF, also known as Version of record

Published under the following license: CC BY

Please cite the original version: Fatima, N., Baqri, S. S. R., Alsulimani, A., Fagoonee, S., Slama, P., Kesari, K. K., Roychoudhury, S., & Haque, S. (2021). Phytochemicals from Indian ethnomedicines: Promising prospects for the management of oxidative stress and cancer. Antioxidants, 10(10), [1606]. https://doi.org/10.3390/antiox10101606
Abstract: Oxygen is indispensable for most organisms on the earth because of its role in respiration. However, it is also associated with several unwanted effects which may sometimes prove fatal in the long run. Such effects are more evident in cells exposed to strong oxidants containing reactive oxygen species (ROS). The adverse outcomes of oxidative metabolism are referred to as oxidative stress, which is a staple theme in contemporary medical research. Oxidative stress leads to plasma membrane disruption through lipid peroxidation and has several other deleterious effects. A large body of literature suggests the involvement of ROS in cancer, ageing, and several other health hazards of the modern world. Plant-based cures for these conditions are desperately sought after as supposedly safer alternatives to mainstream medicines. Phytochemicals, which constitute a diverse group of plant-based substances with varying roles in oxidative reactions of the body, are implicated in the treatment of cancer, ageing, and all other ROS-induced anomalies. This review presents a summary of important phytochemicals extracted from medicinal plants which are a part of Indian ethnomedicine and Ayurveda and describes their possible therapeutic significance.

Keywords: ethnomedicine; oxidative stress; ROS; cancer; phytochemicals

1. Introduction

Living cells are subjected to constant wear and tear throughout their lives, and oxidative metabolism constitutes an important mechanism involved in this process. The cellular milieu is characterized by a tricky equilibrium between pro- and antioxidants, which dictates the overall health of organisms. Any imbalance in the relative abundance of key components of oxidative metabolism leads to oxidative stress which forms the basis of a variety of diseases. Failure of the body to deal with oxidative damage is presumed to be the reason for cell senescence, cancer, and many immunological disorders. Thus, understanding oxidative metabolism is an important prerequisite for devising suitable therapeutic strategies against such diseases.

Cancer is a global health problem because of its high prevalence and poor prognosis. It is an outcome of uncontrolled cell division which results from the loss of regulation over
various cell cycle checkpoints. Cancer can involve almost every part of the body, a feature that accounts for the great diversity of known cancers. Many types of cancer have a clear genetic basis and arise from an altered expression of certain genes called oncogenes [1]. Important physiological roles are assigned to cellular oncogenes in a normal cell and most of them are involved in cell cycle regulation through their effects on molecular mechanisms of chromatin condensation, DNA damage, replication, transcription, and translation. Cellular oncogenes are homologous to viral oncogenes, which explains the viral origin of some well-known cancers. A different class of genes involved in cancer progression are tumor suppressor genes, such as p53 and retinoblastoma protein (Rb) that avert the development of cancer by inciting apoptosis in cancerous cells [2]. Any defect in tumor suppressor genes pushes the cancerous cell to escape apoptosis and continue proliferation leading to tumorigenesis [3].

Oxidative metabolism involving reactive oxygen species (ROS) has been shown to play a significant role in key cellular pathways of DNA repair and damage, cell cycle control, and apoptosis [4]. Since all these pathways are correlated with the incidence of cancer, it is obvious that oxidative stress has a huge bearing on cancer epidemiology. In fact, oxidative stress has been shown in causing cancer by inducing DNA damage and mutagenesis. As a result, antioxidant substances are seen as potential candidates in treating various types of cancer, and this has been a promising research area to explore further in cancer therapeutics. In some cases, the theoretical basis of anticarcinogenic effect of certain antioxidants is well known. In this connection, molecules involved in the oxidative metabolism of the cell (such as glutathione, superoxide dismutase - SOD, and peroxidase) appear to be the likely targets for managing cancer [4].

Extensive research has been performed on ways to treat cancer over the past few decades but an ultimate cure remains elusive. However, there has been a significant improvement in cancer prognosis due to the development of strategies such as radiotherapy, chemotherapy, and immunotherapy that aim at killing malignant cells through the use of radiation, powerful cytotoxic chemicals, or antibodies against tumor-specific antigens, respectively. A flip side of all these strategies is that they result in widespread cytotoxicity of healthy cells too and cause severe side effects such as hair loss and skin damage [5]. This has led to an intensified quest for alternative cures that are free of such side effects. Alternative medicine provides a ray of hope for a safe cancer cure exempt from undesirable side effects. Phytochemicals obtained from traditional herbs are known to have a wide chemical diversity, and several classes of such phytochemicals may have anticancer properties. In particular, chemicals such as terpenes, carotenoids, alkaloids, organosulfur compounds, and steroids are abundantly present in plants. Some of these have shown clinical efficacy in treating cancer [6,7]. We discuss these aspects in a detailed manner in the below mentioned sections.

2. Relation between Oxidative Stress and Cancer

Oxidative stress is described as a type of disturbance in the balance between the production of ROS [free radicals and non-free radical oxygen comprising chemical molecules such as hydrogen peroxide (H$_2$O$_2$), singlet oxygen, hydroxyl radical, and superoxide] and antioxidant defense [8]. Oxidative stress has numerous pro-tumorigenic effects such as genome instability, enhancing DNA mutation rate, or promoting DNA damage as well as cell proliferation. On the other hand, oxidative stress also exerts anti-tumorigenic activities and studies have shown that it is correlated with apoptosis and senescence - the two key mechanisms that neutralize cancer development [9]. Cancer cells exhibit enhanced ROS production that may increase cell proliferation. Undeniably, cancer initiation and its progression are correlated to oxidative stress-mediated DNA damage, increased DNA mutations or genome instability as well as cell proliferation [10]. In order to guard the organism against such detrimental pro-oxidants (endobiotic or xenobiotic), a systematically designed complex system of enzymatic antioxidants, for example, SOD, glutathione peroxidase (GPx), glutathione reductase, catalase and non-enzymatic antioxidants (which act by
interrupting free radical chain reactions) such as glutathione (GSH), vitamins C, D and E come into play [11]. Under regular physiological situations, these protective antioxidant molecules are present in quantities adequate to tackle the physiological rate of free-radical production. It is also well known that any further load of free radicals can disturb the antioxidant balance (anti-free radical) and pro-oxidant (free radical) balance. This imbalance leads to oxidative stress, when it goes beyond the capability of the oxidation-reduction system of the body, gene mutation may result leading to many chronic diseases including cancer [12].

3. Indian Ethnomedicinal Plants as a Potential Source of Anticancer Phytochemicals

Phytochemicals of remedial plants encompass a diverse chemical space for healthcare and management of common disorders. India is well known for its custom of traditional remedies and ethnopharmacology [13]. The Indian herbal medicines or the conventional medicaments have been derived from the prosperous customs of prehistoric civilizations and scientific inheritance. Herbal therapeutics, thus comprise a key stake of all the legitimate Indian organizations of medicine such as Ayurveda, Yoga, Unani, Siddha, and Homeopathy (AYUSH) [14]. These herbal remedies are used by more than 70% of the inhabitants for their primary health-related troubles [15]. It is believed that these plant-based herbal remedies are non-toxic against normal cells and are therefore well tolerated (or accepted) by the human body. This attribute is a great motivation to spark the interest of drug designers who are in the pursuit of cheap and safer alternatives to modern synthetic medicines. A brief list of some ayurvedic medicinal plants having potent anticancer activity and their main phytochemicals is provided in Table 1. A state-wise geographical distribution of some prominent Indian ethnomedicinal plants has been provided in Figure 1.

The currently used drug development program based on ayurvedic prescriptions has gained extensive reception in the existing healthcare system. Table 2 listed out several important Indian ethnomedicinal/Ayurvedic remedies and their uses in medical science. Although, validation of a combined therapeutic approach based on Ayurvedic prescriptions and modern medicine with better efficiency and safety is likely to be a big leap in overcoming some of the hurdles in the way of treating difficult disorders such as cancer [16].

Anticancer phytochemicals and their derived secondary metabolites present in leaf, root, flower, bark, and stem catalyze numerous pharmacological functions in human healthcare systems. Flavonoids, alkaloids, glycosides, phenolics, tannins, gums, oils, and resins are such representative compounds [17]. These phytochemicals or their modified derivatives have displayed significant anti-tumor potential.

There are four major classes of plant-derived or plant-based anti-cancer agents that are currently being used commercially: (1) the vinca alkaloids from *Catharanthus roseus* (vincristine, vinblastine, and vindesine), (2) the epipodophyllotoxins from *Podophyllum peltatum* (etoposide and teniposide), (3) the taxanes from the genus *Taxus* (docetaxel and paclitaxel) and (4) the camptothecin derivatives from *Camptotheca acuminata* (irinotecan and camptotecin) [18]. Moreover, the anti-cancer potential of a variety of plants is still being actively investigated and even some have displayed very encouraging results.
Table 1. A brief list of some ayurvedic medicinal plants having potent anticancer activity and their main phytochemicals.

| Ayurvedic Name                      | Scientific Name | Family               | Active/Antioxidant/Anticancer Phytochemicals                                                                                                                                                                                                 | References |
|-------------------------------------|-----------------|----------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------|
| Amrita/Guduchi/Giloya              | Tinospora cordifolia | Menispermaceae       | Palmative, new clerodane furanoditerene glycoside, arabinogalactan, berberine, phenolic compounds, and epoxy cleodane diterpene                                                                                                                                                      | [19]       |
| Amritavallari                      | Basella rubra    | Basellaceae           | p-Coumaric acid, caffeic acid, diosmetin                                                                                                                                                                                                   | [20]       |
| Amruta, Kalgu, or Narkya            | Mappia foetida/Nothapodytes foetida | Icacinaceae          | Camptothecin                                                                                                                                                                                                                                      | [21]       |
| Anikol                              | Alangium salviifolium | Cornaceae            | Deoxytubulosine, alangimarcine, dehydropseudoemetine, salviifosides A-C, kaempferol, salicin, and kaempferol 3-O-b-D-glucopyranoside                                                                                                                                 | [22]       |
| Ashwagandha                         | Withania somnifera | Solanaceae           | Withanolides, sitoindosides                                                                                                                                                                                                                 | [23]       |
| Ashwatha                            | Ficus religiosa  | Moraceae             | Quercetin, myricetin, kaempferol, β-sitosteryl-D-glucoside                                                                                                                                                                                    | [24]       |
| Bhukamtaka suksharanphala          | Ziziphus nummularia | Rhamnaceae           | Rutin, chlorogenic acid, quercetin, pyrogallol, mandelic acid, morin                                                                                                                                                                        | [25]       |
| Bhumyamalaki                        | Phyllanthus amarus | Phyllanthaceae       | Nirtetralin (NIRT), nirantrhin (NIRA), phyllanthin (PHYLLA), phyltetralin                                                                                                                                                                    | [26]       |
| Bhunimba and kalmegha              | Andrographis paniculata | Acanthaceae         | Andrographolide                                                                                                                                                                                                                                 | [27]       |
| Bhurjapatra                         | Betula utilis    | Betulaceae           | Betulinic acid                                                                                                                                                                                                                                   | [28]       |
| Bilva                               | Aegle marmelos  | Rutaceae             | Skimmianine, marmesin, imperatorin, ciminaldehyde, xanthotoxol                                                                                                                                                                                 | [29]       |
| Brahmamanduki                       | Centella asiatica | Acanthaceae          | Asiaticoside                                                                                                                                                                                                                                    | [30]       |
| Brahmi                              | Bacopa monnieri  | Scrophulariaceae     | Brahmine, herpestine, nicotinipe, bacosides A and B, betulinic acid, wogonin, and oroxindin                                                                                                                                                  | [31,32]    |
| Daruharidra                         | Berberis aristata | Berberidaceae        | Berberine, berbamine, oxyberberine, aromoline, a protoberberine alkald karachine, palmatine, taxilamine, and oxycanthine                                                                                                                                                       | [33]       |
| Datura                              | Datura metel     | Solanaceae           | Pterodontriol B, scopoline, adenosine, disciferitriol, thymidine, dioscoroside D, and ilekudinoside C                                                                                                                                            | [34]       |
| Gunja                               | Abrus precatorius | Leguminosae          | Abrin, Cycloartenol, Luteolin, Isoorientin, Trigonelline                                                                                                                                                                                       | [35]       |
| Haridra                             | Curcuma longa    | Zingiberaceae        | Curcumin, Curcuminoid, Desmethylcurcurmin, Bisdemethylcurcumin, Curdione, Bisacurone                                                                                                                                                          | [36]       |
| Ayurvedic Name                  | Scientific Name          | Family         | Active/Antioxidant/Anticancer Phytochemicals                                                                 | References |
|--------------------------------|--------------------------|----------------|----------------------------------------------------------------------------------------------------------|------------|
| Haritaki                        | Terminalia chebula       | Combretaceae   | Arjunglucoside I, chebulosides I and II, arjungenin, chebulin, 2,4-chebulyl-ß-D-glucopyranose, chebulic acid, chebulinic acid, terchebin | [37]       |
| Kadamba, Vrattapuspa, Sisupala  | Anthocephalus cadamba   | Rubiaceae      | Cadamine, isocadambine, isocadambine                                                                     | [38]       |
| Kakamachi                       | Solanum nigrum          | Solanaceae     | Solamargine, Solasonine, Solasodine, Solanidine                                                          | [39]       |
| Kumbhini                        | Croton tiglium           | Euphorbiaceae  | Corydine and salutaridine                                                                                 | [40]       |
| Kumkuma/Ghusrun/Agneeshekhar    | Crocus sativus           | Iridaceae      | Crocin, crocetin, picrocrocin, and safranal                                                              | [41]       |
| Lakshamanap hala                | Annona muricata          | Annonaceae     | Annonacin, Isoquinoline, Anonaine, Bullatacin, Annonamine, Gentisic acid                                 | [42]       |
| Lavanga                         | Syzygium aromaticum     | Myrtaceae      | Eugenol, Bicornin, Caryophyllene, Tellimagrandin II                                                      | [43]       |
| Neem                            | Azadirachta indica      | Meliaceae      | Azadirachtin, Nimbin, Gedunin, epoxyazadiradione, Oleic acid                                             | [44]       |
| Nimbuka                         | Citrus medica            | Rutaceae       | Citronellal, Undecanal, Bisabolene                                                                         | [45]       |
| Pandalu                         | Allium cepa              | Liliaceae      | Quercetin, Diallyl disulfide, Allicin                                                                       | [46]       |
| Pippali                         | Piper longum             | Piperaceae     | Piperine, Chavicine, Piperlongumine, Piperlongumunine                                                      | [47]       |
| Rasna                           | Alpinia galangal         | Zingiberaceae  | Galangin, Kaempferide, Cadinene, fenchyl acetate                                                         | [48]       |
| Rasona                          | Allium sativum           | Amaryllidaceae | Allii, allicin allii, alliinase                                                                           | [49]       |
| Revandachini                    | Rheum emodi              | Polygonaceae   | Emodin, aloë-edomin, chrysophanol, physcion, rhein, emodin glycoside, and chrysophanol glycoside          | [50]       |
| Shatavari                       | Asparagus racemosus      | Liliaceae      | Shatavaroside A and B, shatavarins, filiasparoside C, immunoside, and schidigerasaponin                  | [51]       |
| Suthi                           | Zingiber officinalis     | Zingiberaceae  | Gingerol, Zingiberene, Shogaol, Paradol, Zingerone                                                        | [52]       |
| Tulsi                           | Ocimum sanctum           | Labiatae       | Eugenol, Caryophyllene, Linalool, Methyl eugenol, Estragole, Carvacrol, Cadinene, Farnesene              | [53]       |
| Vacha, Ugargandha, Chhadgrantha | Acorus calamus           | Acoraceae      | Asarone, Cadinene, Methyl isoeugenol                                                                      | [54]       |
Table 2. Some popular Indian ethnomedicinal/Ayurvedic remedies and their uses.

| Ethnomedicinal/Ayurvedic Remedy | Plant Species | Description | Benefits and Medicinal Uses | References |
|---------------------------------|---------------|-------------|-----------------------------|------------|
| Jatamansi                        | *Nardostachys jatamansi* DC | Ayurvedic Powder | Use as brain tonic that help to improve memory and brain functions by preventing cell damage due to its antioxidant property | [55] |
| Bhootkeshi                       | *Selenium vaginatum* (Edgew) Cl | Ayurvedic Powder | Sleep and mental disorders, high blood pressure | [56] |
| Haridra/Daruharidra              | *Curcuma longa* | Ayurvedic Powder | Chronic anterior uveitis, rheumatoid arthritis, conjunctivitis, small pox, skin cancer, chicken pox, urinary tract infections, wound healing, and liver ailments | [57] |
| Drakshasava                      | *Vitis Vinifera, Woodwardia Fruticosa, Piper Cubeba, Cinnamomum Tamala, Mesua Ferrea, Syzygium Aromaticum Myristica Fragrans, Piper Nigrum, Plumbago Zeylanica, Piper Retrofractum Piper Longum, Vitex Negundo* | Ayurvedic rejuvenators | Weakness, lethargy and heat exhaustion, curing haemorrhoids and cardiac disorders | [58] |
| Kanchanar Guggulu                | *Bauhinia variegata* L. (BV), *Zingiber officinale, Piper nigrum, Piper longum, Terminalia chebula, Terminalia bellirica, Embelia officinalis, Crataeva nurvala, Cinnamomum tamala, Elletaria cardemomun, Cinnamomum zeylanicun, Comniphora muku* | Ayurvedic Formulation | Hormonal imbalance, PCOS, hypothyroidism, and joint pains. weight loss, lipoma, tumor, cysts, cancer, goiter, fistula, boils and skin related diseases | [59] |
| Hingvastaka                      | *Zingiber officinale, Piper nigrum, Piper longum, Trachyspermum amni, Carum carvi, Cuminum Cyminum, Ferrula asafetida* | Ayurvedic churna | Indigestion, Anorexia and all Vata Disorders, relieve flatulence | [60] |
| Triphala                         | *Phyllanthus emblica,Terminalia bellirica, Terminalia chebula* | Polyherbal Ayurvedic medicine | Indigestion, weight loss, Reducing inflammation, regulate blood sugar levels, Lower cholesterol, reduce tumors, prevents cancer, inhibit HIV | [61] |
| Ethnomedicinal/Ayurvedic Remedy | Plant Species | Description | Benefits and Medicinal Uses | References |
|-------------------------------|--------------|-------------|----------------------------|------------|
| Aswagandharishtam             | *Withania somnifera, Rubia cordifolia, Chlorophytum tuberosum, Terminalia chebula, Curcuma longa, Berberis aristata, Glycyrrhiza glabra, Pluchea lanceolata, Pueraria tuberosa, Terminalia arjuna, Cyperus rotundus, Ipomoea turpethum, Hemidesmus indicus, Santalum album, Acorus calamus, Plumbago indica* | Liquid Ayurveda medicine | Anti-anxiety, anti-stress, antidepressant | [62] |
| Kasisadi Tailam               | *Leucaena Leucocephala, Ficus Religiosa, Dry Ginger, and Plumbago Zeylanica* | Polyherbal Ayurvedic oil | External application on corns, piles and warts | [63] |
| Pusyanuga curna              | *Cissampelaos pareira, Syzygium cumini, Mangifera indica, Bergenia lingulata, Berberis aristata, Ambastha (Patha)- Cissampelos pareira, Mochrara- Salalina malabarica, Mimosa pudica, Holarrhena antidysentrica, Crocus sativus, Aconitum heterophyllum, Aegle marmelos, Cyperus rotundus, Symplcos racemosa, Ochre or Haematite, Alaththus excels, Piper nigrum, Zingiber officinalis, Vitis vinifera, Pterocarpus santalinus, Myrica esculenta, Holarrhena antidysentrica, Hemidesmus indicus, Woodfordia fruticosa, Glycyrrhiza glabra, Terminalia arjuna* | Herbo-mineral formulation | Conditions involving menstrual irregularities such as menorrhagia, metrorrhagia, dysmenorrhoea and endometriosis, piles, diarrhoea, bloody stools and different types of discharges from vaginal tract | [64] |
| Lashuna taila                | *Allium Sativum linn, Sesamum indicum linn* | Ayurvedic oil | Fungal infections, warts, and corns, hair loss and thrush | [65] |
4. Antioxidant Phytochemicals in the Management of Cancer

It is well established that the incidence of cancer and its progression has been associated with oxidative stress owing to the role played by oxidative agents in mutagenesis and induction of DNA damage which lead to cell proliferation and genome instability. This gives credence to the speculation that antioxidant agents could intercept carcinogenesis [66]. A plethora of studies has reported that phytochemicals from ethnomedicinal plants can yield numerous bioactive compounds with promising anti-cancer potential. Besides, in
addition to their renowned antioxidant properties, several phytochemicals target epige-
netic processes participating in cancer progression by modulating oxidative stress [67,68].
In treating cancer, an efficient strategy is to induce senescence and cytostasis by using
cytotoxic agents. Several anticancer polyphenols arrest cellular growth via the induction
of ROS-dependent premature senescence (RDPS), and are contemplated as promising
antitumor therapeutic options [69]. In the light of this mechanism, pro-oxidant therapy
which involves drugs that trigger oxidation-induced cell death has been emerging as an
effective strategy for treating cancer [70].

Several recent epidemiological researches have demonstrated that nutritional antioxi-
dant phytochemicals such as phenolic compounds, carotenoids, flavonoids, and alkaloids
can act as defensive agents against many chronic diseases including cancer [71]. In the re-
cent past, numerous studies have been conducted to elucidate the mechanisms that impart
anticarcinogenic properties to phytochemicals, such as vitamin C, vitamin E, carotenoids,
flavonoids, and phenolic acids [72]. Curcumin, which is a bright yellow polyphenolic
phytochemical produced by Curcuma longa (Turmeric) plants and is widely used in In-
dian ethnomedicine, is in fact a strong antioxidant agent. It is an exceptionally potent,
lipid-soluble compound owing its healing properties to its pro-oxidant/antioxidant effects
since the generation of ROS by curcuminoids and curcumin correlates with their apoptotic
activity on cancer cells [73]. Antioxidant or pro-oxidant capacity of any prospective can-
idate drug against cancer thus appears heavily dependent on the concentrations of active
phytochemicals in it. A number of studies using different cell models have highlighted the
pro-oxidative properties of polyphenols, which are also known as antioxidants, such as
genistein, quercetin, catechins, epicatechin, and resveratrol [74,75]. Figure 2 presents the
structure of published articles indexed in PubMed online database pertaining to the most
studied antioxidant phytochemicals associated with oxidative stress and cancer.

Quercetin can be used for averting cancer via the modulation of oxidative stress factors
and several antioxidant enzymes for preventing the progression of a variety of cancers,
such as lung, liver, prostate, colon, breast, and cervical cancers [75,76]. In an in vivo
research study performed on rats by Sharmila et al., the antioxidant activity of quercetin
was matched with that of a carcinogen and testosterone by histological evaluation and
measuring oxidative stress markers, such as lipid peroxidation (LPO), H₂O₂ and reduced
GSH. They observed that carcinogen and testosterone-treated rats had higher levels of LPO
and H₂O₂ and lower levels of GSH compared with rats with quercetin treatment. They also
reported that quercetin improved the levels of antioxidant enzymes and apoptotic proteins
in animals having prostate cancer. Additionally, the authors observed that quercetin
synchronized the expression of androgen receptors (AR), insulin-like growth factor receptor
1 (IGFR1), protein kinase B (AKT), cell proliferation factors, and anti-apoptotic proteins
which are usually elevated in cancer [77]. Quercetin also trims down the overproduction of
ROS, inhibits the expression of tumor necrosis factor (TNF)-α gene, corrects the damage
inflicted by trauma and averts myocardial cell injury caused due to Ca²⁺ overload [78].
Thus, quercetin can efficiently prevent injuries caused by oxidative stress [79].

Furthermore, silymarin from milk thistle, green tea polyphenols, and proanthocyani-
dins from grape seeds all have the capability to defend the skin from the unfavourable
effects of UV radiation including skin cancers. The protection against UV damage was
achieved mainly through four mechanisms, namely inhibition of DNA damage, induction
of inflammation, suppression of immune responses, and oxidative stress [80].

Resveratrol is a stilbene-type aromatic phytoalexin, mainly available in red wine,
grape skin, peanuts, berries, purple grape juice, turmeric, and other dietary foodstuffs.
Resveratrol has shown potential anticancer activity primarily because of induction of apop-
tosis through several pathways, all leading to a reduction in tumor initiation, promotion,
and progression, and alteration in gene expression [81]. Lately, it is reported that the an-
tioxidant potential of hydroxyl stilbenes (trans- and cis-resveratrol and their hydroxylated
analogs) is associated with the regulation of eicosanoid synthesis, and the basic stilbene
structure of two benzene rings bonded by a central ethylene is chiefly accountable for its
properties on Caco-2 cell growth, DNA synthesis, and cell cycle independently of redox state and modulation of eicosanoid synthesis [82].

Figure 2. Illustration of the most reviewed antioxidant phytochemicals associated with oxidative stress and cancer. The structure of published literature is based on the citations in PubMed (April 2021). The numbers of published manuscripts are provided in parentheses along with the name of each phytochemical and “oxidative stress and cancer”, within title/abstract, is stated.

Lycopene is a non-provitamin A carotenoid which has shown potential anticancer activity against advanced and aggressive conditions of prostate cancer [83]. Lycopene and its auto-oxidant products may stimulate apoptosis in HL-60 cells. It is involved in modulating Bcl-2, thereby influencing apoptosis, which may explain the observed antioxidant properties of lycopene. It also restrains carcinogenesis by saving indispensable biomolecules such as proteins, lipids, LDL, and DNA [84]. Besides, lycopene also possesses the quenching capability against singlet oxygen that can be accredited to its conjugated double bonds [85].

5. Classification of Dietary Antioxidant Phytochemicals in the Management of Oxidative Stress and Cancer

5.1. Phenolic Compounds

Phenolic compounds represent one of the largest and most extensive groups of secondary metabolites in the plant kingdom and are characterized by an aromatic ring holding one or more hydroxyl groups. More than 8000 natural phenolic compounds have been identified which are divided into different subgroups such as phenolic acids, tannins, flavonoids, coumarins, quinones, lignans, curcuminoids, and stilbens [86].

Flavonoids are further classified as flavonols, flavones, flavanones, isoflavonoids, and anthocyanidins. Polyphenols are explored as promising medicinal agents for the treatment of various diseases including ulcer, bacterial infections, hypertension, allergies, hypercholesterolemia, and vascular fragility, especially in various types of cancer. Table 3 lists the
There is a popular faith that nutritional polyphenols have anti-cancer properties because of their anti-oxidative characteristics. Their conjugated structures give rise to wonderful electron delocalization characteristics that bestow the ability to quench free radicals and react with a large number of ROS, including singlet oxygen, superoxide radical, nitric oxide, peroxyl radical, hydroxyl radical, peroxynitrite and nitrogen dioxide. The occurrence of numerous hydroxyl groups in their structures makes polyphenols exceptional hydrogen bond donors. Their ability to bind hydrogen is supposed to be responsible for their elevated affinity for nucleic acids and proteins. Polyphenols have a molecular basis to induce cancer cell death by the down-regulation of oncogenic survival kinases such as PI3K and Akt, D-type cyclins, cyclin-dependent kinases (CDKs), cell proliferation regulators that include Erk1/2, angiogenic factors such as VEGF, FGFR1, and MIC-1 and transcription factors such as NF-κB, NRF2, and STATs etc. [87].

Among all the polyphenols, flavonoids are the most abundant in nature, having a basic skeleton of phenylbenzopyrone consisting of two aromatic rings. They can occur in nature in both free and conjugated forms. They have a wide range of anti-cancer and anti-mutagenic properties. However, whether a particular polyphenol is antimutagenic or not solely depends upon its chemical structure, the concerned gene, and the mutagenic factor (alcohol, ultraviolet radiation, tobacco consumption etc.).

Resveratrol (RE; 3,4′,5 trihydroxystilbene) is a stilbenoid natural polyphenol, found in many Indian ethnomedicinal plants such as Drakshasava (*Vitis vinifera* L.), Dadima (*Punica granatum* Linn. fruits), and Shahoot (*Morus indica*) [88]. It demonstrated potential inhibitory activity in different cancer types, such as for the initiation, promotion, and progression stages.
The apoptotic signals are originated through two main pathways: (i) the intrinsic pathway and (ii) the extrinsic pathway. The intrinsic pathway operates by stimulating the mitochondrial membrane to hinder the expression of anti-apoptotic proteins Bcl-2 and Bcl-XL [89]. Curcumin perturbs the balance in the mitochondrial membrane potential, leading to enhanced suppression of the Bcl-xL protein [90].

Ginger has served as a principal drug in Ayurveda, Unani, Siddha, Chinese, Homeopathy, and Tibetan traditional medicines for more than 2000 years [91]. It contains a variety of active phenolic compounds, e.g., [6]-gingerol, [8]-gingerol, [10]-gingerol, and [6]-shogaol. Among these, [6]-gingerol is one of the most significant pungent components of ginger and was found to have varied pharmacological actions, such as antioxidant, antiemetic, cardiotonic, anti-tumor, anti-inflammatory, and anti-platelet properties [92]. de Lima et al. (2018) found that ginger extract and [6]-gingerol exerted their action through imperative mediators and pathways of cell signaling, including Bax/Bcl2, Nrf2, p38/MAPK, p65/NF-κB, ERK1/2, SAPK/JNK, ROS/NF-κB/COX-2, TNF-α, caspases-3, -9, and p53 [93].

The vitamin E family, tocopherols (α-, β-, γ-, δ-), and tocotrienols (α-, β-, γ-, δ-) are an important group of fat-soluble phenolic compounds having strong antioxidant and anticancer properties. The major Indian ethnomedicinal dietary sources of tocopherols (vitamin E) are Vatada (Prunus dulcis), Surajmukhi seeds (Helianthus annuus), and Gehu jawara (wheatgrass oil). There are several reports demonstrating that tocopherol mixture rich in γ-tocopherol inhibits lung, mammary, prostate, and colon tumorigenesis in animal models [94]. It is also evident by numerous in vitro and in vivo reports that γ-tocopherol, δ-tocopherol, and vitamin E metabolite 13’-carboxychromanol have potent anti-cancer activities via modulating key signaling and several mediators that regulate tumor progression and cell death for example PI3K, NF-κB, STAT3, sphingolipid and eicosanoids metabolism [95].

Flavonoid riboflavin (vitamin B2) is a water-soluble vitamin found in milk, dairy products, green leafy vegetables, fortified cereals, and grain products. This vitamin is necessary for normal cell growth and function, is required to process fats and amino acids, and helps convert carbohydrates into fuel. Along with several medicinal properties, this vitamin also has antioxidant nature and can protect the body against oxidative stress, reperfusion oxidative injury, and especially LPO [96].

| Bioactive Molecules | Indian Ethnomedicinal Plant Source | Chemical Classification | Associated Condition | Cell Line Tested | Biological Approach (In Vitro/In Vivo) | Mechanism of Action | References |
|---------------------|-----------------------------------|-------------------------|----------------------|-----------------|----------------------------------------|---------------------|------------|
| Chlorogenic acid    | Jjatamansi (Nardostachys jatamansi) | Ester of caffeic acid and (−)-quinic acid | Osteosarcoma         | U2OS, Saos-2, and MG-63 OS | In vitro | Activates extracellular-signal-regulated kinase1/2 (ERK1/2) | [97] |
| Epigallocatechin-3-gallate | Syamaparni (Camellia sinensis) | Ester of epigallocatechin and gallic acid | Oral squamous cell carcinoma | KBV200 | In vivo | Inhibition of angiogenesis via VEGF down-regulation | [98] |
| Genistein | Rajapatha (Stephania glabra) | Isoflavones | Kidney cancer | In vitro | | Induces apoptosis and inhibit the proliferation via regulating CDKN2a methylation | [99] |
| Resveratrol | Drakhasava (Vitis vinifera L.) | Stilbenoid | Renal cell carcinoma | In vitro/in vivo | | Depressing activity of NLRP3, and NLRP3 | [100] |
### Table 3. Cont.

| Bioactive Molecules | Indian Ethnomedicinal Plant Source | Chemical Classification | Associated Condition | Cell Line Tested | Biological Approach (In Vitro/In Vivo) | Mechanism of Action | References |
|---------------------|-------------------------------------|-------------------------|----------------------|-----------------|----------------------------------------|---------------------|------------|
| Quercetin           | Tulsi (Ocimum sanctum)              | Flavonol                | Malignant melanoma   | B16             | In vitro                              | Reduced the proportion of cells in the S and G2/M stages of the cell cycle | [101] |
| Daidzein            | Vidari (Pueraria tuberosa)          | Isoflavones             | Colorectal cancer    | In vivo         | Lessened the protein expression of p-ERK/ERK and p-AKT/AKT | [102] |
| Gallic acid         | Amla (Emblica officinalis)          | Phenolic acid           | Prostate cancer      | In vitro/In vivo| Inhibits HDAC1 and 2 expression        | [103] |

### 5.2. Alkaloids

Alkaloids are amongst the highly copious plant secondary metabolites. They create a large conglomerate of fundamental heterocyclic nitrogen encompassing natural phytochemicals that are usually produced by plants as toxic substances [17]. Out of 27,000 diverse alkaloids known, nearly 17,000 have shown varied pharmacological properties comprising anticancer activities [104]. Among the many pharmacological features of alkaloids are their prominent antioxidant properties which make them capable of averting a range of degenerative ailments either through their binding to catalysts involved in diverse oxidation processes happening within an individual’s body or through capturing free radicals.

Alkaloids can be classified into several groups such as pyrrolizidines, quinolizidines, pyrrolidines, indoles, piperidines, tropanes, purines, isoquinolines, and imidazoles. Among these groups, the leading anticancer alkaloids isolated from the plants are compounds such as taxol, vinblastine, vincristine, topotecan, vinflunine, and camptothecins while other important isolated alkaloids include harringtonine, rohitukine, thalicarpine, acronycine, usambaresine, matrines, and ellipticine. Several studies have proved the efficacy of plant-based alkaloids in oncogenesis inhibition. Alkaloids have been shown to modulate key signaling pathways participating in proliferation, metastasis, and cell cycle regulation, and these very properties form the basis of their use as the principal components of numerous clinical anti-cancer agents. Table 4 lists the anticancer activity of some recently reported potent antioxidant alkaloids, whereas Figure 4 shows the chemical structures of some lead alkaloids.

Vinca alkaloids, colchicine, and paclitaxel are amongst the most primitive plant-derived anticarcinogenic compounds. These are microtubule-targeting agents which can be used to abort spindle formation during the cell division of rapidly dividing cancer cells. Thus, they are of clinical significance against many types of cancer such as ovarian, breast, prostate, and non-small cell lung cancer.

Berberine is an isoquinoline derivative extracted from *Berberis aristata* also known as “Daruharidra” in Ayurveda. It is a wild shrub found at altitudes of 5000 feet and above in the Nepal and Himalayas region [105]. It is an isoquinoline derivative extracted from *Coptis chinensis* [Family; Berberidaceae], which possesses diverse pharmacological actions. It is equipped with anti-bacterial, anti-diabetic, anti-inflammatory, and anti-cancer properties and is also beneficial for the cardiovascular system [106].

Quite recently, Wang ZC et al. synthesized some novel berberine derivatives with di-substituents on positions C9 and C13 and assessed their anticarcinogenic activity against breast cancer cell line (MDA-MB-231), human prostate cancer cell lines (PC3 and DU145), and human colon cancer cell lines (HT29 and HCT116). Out of these, a specific compound designated as 18e showed the highest cytotoxicity against PC3 cells having an IC50 value of 0.19 µM. Additional studies revealed that 18e can arrest the cell cycle at the G1 phase and can extensively stop tumor cell colony formation and migration even at very low concen-
trations. Remarkably, 18e could notably bring about cytoplasmic vacuolation, suggestive of a different mechanistic action from berberine. It is well evidenced that potential molecular targets, as well as mechanistic action of berberine, are rather complex. It interacts with RNA or DNA to form a berberine-RNA or a berberine-DNA complex, respectively.

Figure 4. Chemical structures of some lead alkaloids.

Rohitukine is a chromone alkaloid, that was initially isolated from the leaves and stems of *Amoora rohituka* widely used in the Ayurvedic system of medicines [107]. It has been shown to possess many important biological functions including anti-inflammatory, anti-fertility, and immunomodulatory effects [108]. The unique chemical structure of rohitukine presents a framework for derivatization and chemical synthesis of some novel molecules [109]. P-276-00 and flavopiridol are two rohitukine analogs presently evaluated in the advanced phase II clinical trials for a potential anticancer drug [110]. Flavopiridol demonstrated efficient action against most cancer cell lines and tumorous growth suppression in animals [111]. It is a pan-CDK inhibitor that arrests the cell cycle in G1/S or G2/M phase and has been used in the treatment of chronic lymphocytic leukemia [112]. In a recent study, it has been observed that rohitukine induces cytotoxic effects in lung cancer (A549) cells and stimulates the production of ROS following the exposure of 24 hrs. It was also found that rohitukine activated apoptosis in A549 cell line through the upregulation of p53 and caspase 9 and the downregulation of Bcl-2 protein [113].
Pain is an important point of consideration in dealing with cancer patients especially when its severity increases too much in advanced stages of the disease. There are two types of pain depending on its origin. The nociceptive pain which is sensed by nociceptors (pain receptors) is of peripheral origin, arising in bodily organs due to the invasive growth of the tumor or due to invasive surgical procedures. The second type of pain which is neuropathic and is either central or peripheral in origin occurs when the growing tumor in its advanced stages invades into the nervous tissue. Very often, the pain is iatrogenic and is caused by the drugs used in the treatment of cancer. Vinca alkaloids used as chemotherapeutic agents have been found to cause such iatrogenic pain. One estimate suggests that pain is not properly managed in more than half of cancer patients and a quarter of all cancer patients eventually die of pain. There exists a three-step guideline from the WHO to manage pain in cancer patients and it recommends the use of opiate alkaloids in the second and third stages of cancer. However, the use of opiate drugs is discouraged because of the fear of dependency and addiction, yet a study has found that such apprehensions are not true.

### Table 4. Anticancer activity of some potent antioxidant alkaloids (recent data).

| Bioactive Molecules | Indian Ethnomedicinal Plant Source | Chemical Classification | Associated Condition | Cell Line Tested | Biological Approach (In Vitro/In Vivo) | Mechanism of Action | References |
|--------------------|-----------------------------------|-------------------------|----------------------|------------------|----------------------------------------|---------------------|------------|
| Berberine          | Daruharidra (Berberis aristata)    | Isoquinoline alkaloid   | Colon cancer         | In vitro         | Targets SCAP/SREBP-1 pathway driving lipogenesis |                     | [114] |
|                    |                                   |                         | Chronic lymphocytic leukemia | Clinical trial/human patient | Induces apoptosis by decreasing ROR1, Bcl-2, and mir-21 |                     | [115] |
| Capsaicin          | Twak (Cinnamomum verum)           | Capsaicinoids           | Prostate cancer       | PC-3, DU145      | In vitro                              | Suppresses prostate cancer stem cells via inhibition of Wnt/β-catenin pathway | [116] |
| Piperine           | Pipali (Piper nigrum)             | N-acylpiperidine        | Human melanoma cells  | A375P, A375SM    | In vitro and in vivo                   | Increased the expression of BCL2-associated X, apoptosis regulator (BAX), cleaved poly (ADP-ribose) polymerase, cleaved caspase-9, phospho-c-Jun N-terminal kinase and phospho-p38, and reduced that of B-cell lymphoma 2 (BCL2) | [117] |
| Sanguinarine       | Svarnakshiri (Argemone mexicana)  | Benzophenanthridine alkaloid | Multiple myeloma | U266, IM9, MM1S, and RPMI-8226 | In vitro | Induces mitochondrial/caspase-dependent apoptosis, produces oxidative stress, and suppresses proliferation cancer cell lines | [118] |
| Tetrandrine        | Patha (Cissampelos pareira)       | Bisbenzylisoquinoline alkaloid | Breast cancer | MDA-MB-231     | In vivo | TET-upregulated Caspase-3, Bid, Bax, and, downregulated by Bcl-2, Survivin, and PARP | [119,120] |
and addiction has not been reported in cancer patients taking opiates [121]. The standard strategy of using opiates as analgesics involves the use of alkaloids such as codeine in mild pain and morphine or other members of the class in later stages of severe pain [122]. It has long been known that morphine and other opiate alkaloids work by binding to $\mu$-opiate receptors in the central nervous system to suppress pain [123].

5.3. Organosulfur Compounds

Organosulfur compounds (OSCs) are organic molecules that contain sulfur principally responsible for the characteristic odour and flavour of onion and garlic. They are also abundant in cruciferous vegetables such as cabbage and broccoli [124]. There are two major groups of natural resources that contain OSCs with unique properties. Onion, garlic, shallot chives, and leeks are famous representatives of the Allium genus that have S-alk(en)yl-l-cysteine sulfoxides [125]. Cauliflower, cabbage, and kale are representatives of the genus Brassica, while rucola belongs to the genus Eruca of cruciferous family that contain S-methyl cysteine-l-sulfoxide [126]. Earlier research studies have revealed the anti-cancer mechanisms of extracts, preparations, and organosulfur products of alliaceous vegetables. These mechanisms presumably include their antimicrobial activity, redox modification, and reduced bioactivation of cancer-causing agents. Hence, Allium vegetables and their active phytochemicals have assumed significance in efficiently regulating the biological processing of carcinogenesis so as to alter or reduce the hazard of cancer. OSCs inhibit the development of cancer cells by several pathways including inhibition of metabolism, inhibition of angiogenesis, and cell cycle arrest. These compounds are classified into water-soluble OSCs such as S-Allyl Mercaptocysteine and S-Allyl Cysteine and oil-soluble OSCs, such as diallyl disulfide, diallyl sulfide, diallyl trisulfide, ajoene, and dithiins [127]. Numerous organosulfur compounds are declared to have potent antioxidant capacity. Their potential to scavenge ROS plays conceivably a significant role in their anti-senescence activity in vitro. Table 5 reports the anticancer activity of some recently reported potent antioxidant organosulfur phytochemicals and Figure 5 shows chemical structures of some lead organosulfur phytochemicals.

![Chemical structures of some lead organosulfur phytochemicals.](image)

**Figure 5.** Chemical structures of some lead organosulfur phytochemicals.

Alliin, allicin, S-alllylcysteine, and S-allylmercaptocysteine are some major phytochemicals present in garlic, which is commonly known as ‘Lasuna’ in Sanskrit, considered to be one of the best Ayurvedic medicine and termed as Mahaauashadha [128]. Recently, Rosas-González et al. appraised the effects of alliin and allicin on cell death and senescence, and their senolytic potential in MCF-7 (luminal A) and HCC-70 (triple-negative breast
cancer cells). Their results revealed that allicin has anti-proliferative, anti-clonogenic, and senolytic properties. In addition, allicin reduced cell viability and induced apoptosis by triggering the loss of ∆Ψm, caspase-3, caspase-8, and caspase-9 activation, downregulation of BCL-XL expression as well as upregulation of NOXA, BAK, and P21. On the other hand, alliin endorsed clonogenicity, induced senescence, and did not demonstrate pro-apoptotic effects in breast cancer cells [129]. Allicin also suppressed proliferation and induced glioma cell apoptosis in vitro by intrinsic mitochondrial and extrinsic Fas/FasL mediated pathways [130].

Allicin is extremely unstable and is easily transformed into lipid-soluble sulfides such as diallyl sulfide (DAS), diallyl disulfide (DADS), and diallyl trisulfide (DATS) [131]. DAS, DADS, and DATS were revealed to play a significant role in inducing histone acetylation thereby resulting in the suppression of the oncogenic protein expression in human glioblastoma cells [132]. It has also been shown that DATS and DAS have the ability to induce apoptosis in the carcinogen-induced two-stage mouse model of skin tumour via targeting of multiple cancer signaling pathways [133].

Some garlic-based water-soluble phytochemicals such as S-allylmercaptocysteine (SAMC) and S-allyl cysteine (SAC) also exhibited potent anticancer activity [134]. SAC could also exhibit stronger antioxidant properties than fresh garlic, and has powerful anti-cancer activity not only at the early stage but also at the late stages of the ailment.

| Table 5. Anticancer activity of some potent antioxidant organsulfur phytochemicals (recent data). |
|---------------------------------------------------------------|
| **Bioactive Molecules** | **Ethnomedicinal Plant Source** | **Chemical Classification** | **Associated Condition** | **Cell Line Tested** | **Biological Approach (In Vitro/In Vivo)** | **Mechanism of Action** | **References** |
|--------------------|-----------------------------|--------------------------|--------------------------|---------------------|------------------------------------------|---------------------------|----------------|
| S-allylcysteine    | Lasuna (Fresh garlic)       | Derivative of the amino acid cysteine | Lung cancer               | A549                | In vitro/in silico                       | Reduced expression of PD-L1 and HIF-1α, showed efficient docking with PD-L1 immune checkpoint target | [135]          |
| Allicin             | Lasuna (Crushed or chopped garlic) | Sulfoxide              | Skin cancer/melanoma       | A375                | In vitro                                 | Reduction of MMP-9 mRNA expression | [136]          |
|                    |                             |                          | Cholangiocarcinoma         | MCF-7, HCC-70       | In vitro/in vivo                         | Overpowers cell proliferation and invasion through STAT3 signaling | [137]          |
| S-allylmercaptocysteine | Lasuna (Aged garlic)      | Thioallyl compound      | Lung cancer               | A549                | In vitro                                 | Suppression of cell proliferation, regulation of cell cycle, attenuation of ROS formation, inhibition of DNA damage, increase of SOD activity and inhibition of nuclear factor-kappa B (NF-kB) activity | [138]          |
| Sulfuraphane       | Shatavari (Asparagus racemosus) | Isothiocyanate         | Small-cell lung cancer    | SCLC cell lines NCI-H69, NCI-H69AR and NCI-H82 | In vitro                                 | Induced cell death was mediated via ferroptosis and Inhibition of the mRNA and protein expression levels | [139]          |
5.4. Carotenoids

Carotenoids are tetraterpenes mainly synthesized in plants with a parent skeleton of hydrocarbon, which consists of \( \text{C}_{40}\text{H}_{56} \) with definite rotation in single and double bonds. As of now, more than seven hundred carotenoids have been identified and can be categorized as carotenes or xanthophylls [140]. Furthermore, carotenoids can be oxidatively sliced by dioxygenase derivatives, called apocarotenoids [141].

A polyene backbone consisting of a sequence of conjugated \( \text{C}=\text{C} \) bonds forms the “core” structural element of carotenoids. This special feature is mainly accredited for both, their pigmenting properties and the capability of many of these phytochemicals to interrelate with singlet oxygen and free radicals, and consequently act as efficient antioxidants [142]. Carotenoids are the primary line protection mechanism that extinguishes oxygen either physically by energy relocation mechanism or chemically through direct reaction with the radicals [143]. Several carotenoids work as powerful antioxidants and anticancer agents. Various epidemiologic and investigational studies have reported that eating of carotenoids is inversely proportional with the incidence of cancer [144,145].

Nutritionally, carotenoids can be divided into two groups, pro-vitamin A and non-provitamin A. Out of thousands of known phytochemicals, only three of them (\( \alpha \)-carotene, \( \beta \)-carotene, and \( \beta \)-cryptoxanthin) act as pro-vitamin A. They can be converted in vivo into vitamin A (retinol). Major dietary sources of non-provitamin A are lutein, lycopene, and zeaxanthin [146]. There are several epidemiologic studies suggesting that carotenoids reduced the hazard of different chronic ailments, especially cancers of gastrointestinal tract, pancreas, lung, and breast with dietary consumption of different carotenoids [147]. On the other hand, randomized clinical trials have not shown any reliable decrease in the frequency of cancers or cancer mortality [144]. Worst of all, in a study on asbestos workers and smokers, an elevated risk of lung cancer had been reported, whether they were given high doses of beta-carotene only or in combination with other antioxidants [148].

Carotenoids such as \( \beta \)-carotene, \( \alpha \)-carotene, lycopene, \( \beta \)-cryptoxanthin, violaxanthin, lutein, fucoxanthin, neoxanthain, canthaxanthin, zeaxanthin, astaxanthin, and siphonaxanthin have been proven for bearing anticancer activity in different cancer cells such as breast, colon, prostate, cervix, leukemia, and liver [149,150]. The proposed mechanisms of cancer chemoprevention using carotenoids probably involve modulations in pathways leading to cell development or cell demise. These mechanisms comprise hormone and growth factor signaling, immune modulation, regulatory mechanisms of cell cycle development, cell differentiation, and apoptosis [151]. The anticancer activities of some recently reported potent antioxidant carotenoids are listed in Table 6, whereas, chemical structures of some leading carotenoids are given in Figure 6.
Table 6. Anticancer activity of some potent antioxidant carotenoids (recent data).

| Bioactive Molecules | Indian Ethnomedicinal Plant Source | Chemical Classification | Associated Condition | Cell Line Tested | Biological Approach (In Vitro/In Vivo) | Mode of Action | References |
|---------------------|-----------------------------------|-------------------------|----------------------|-----------------|----------------------------------------|---------------|------------|
| α-carotene          | Tulsi (Ocimum tenuiflorum)        | Carotenes               | Lung carcinoma       | LLC             | In vitro, In vivo                      | Inhibits Lewis lung carcinoma metastasis and suppresses lung metastasis | [152]       |
| β-carotene          | Gajara (Daucus carota subsp. sativus) | Carotenes               | Gastric cancer       | gastric epithelial cells | In vitro | β-catenin signaling and oncogene expression. | [153]       |
| Lycopene            | Devataruni (Rose hips-Rosa canina L.) | Carotenes               | Human cervical cancer | HeLa cells      | In vitro                              | Inhibition of cell viability, upregulation of Bax expression, and downregulation of Bcl-2 expression | [154]       |
| Lutein              | Sthulapushpa-Marigold (Tagetes erecta) petals | Xanthophylls           | Human cervical cancer | HeLa cells      | In vitro                              | Induce apoptosis by increasing ROS production, interaction with mitochondrial factors, and upregulation of caspase-3-mediated pathway resulting in fragmentation of nuclei DNA | [155]       |
| Zeaxanthin          | Kumkuma (Crocus sativus)          | Xanthophylls            | Gastric cancer       | AGS, KATO-3, MKN-45, MKN-28, NCI-N87, YCC-1, YCC-6, SUN-5, SUN-216, YCC-16, SUN-668, SUN-484 | In vitro | Upregulating intracellular ROS levels, and regulating AKT, MAPK, NF-KB, and STAT3 signaling pathways | [157]       |
| β-cryptoxanthin     | Tulsi (Ocimum tenuiflorum)        | Xanthophylls            | Human cervical cancer | HeLa            | In vitro                              | Activated nuclear condensation and disruption of the integrity of the mitochondrial membrane, upregulation of caspase-3, -7, and -9 mRNA, and increased activation of caspase-3 proteins leading to apoptosis and nuclei DNA damage | [158]       |
| Bioactive Molecules | Indian Ethnomedicinal Plant Source | Chemical Classification | Associated Condition | Cell Line Tested | Biological Approach (In Vitro/In Vivo) | Mode of Action | References |
|---------------------|-----------------------------------|-------------------------|----------------------|-----------------|----------------------------------------|---------------|------------|
| Fucoxanthin         | Kelp (seaweed)                    | Xanthophylls            | Brain and spinal cord cancer | U251-human-glioma-cell | In vitro | Apoptosis by triggering ROS-mediated oxidative damage and dysfunction of MAPKs and PI3K-AKT pathways | [159] |
| Fucoxanthinol       | Kelp (seaweed)                    | Xanthophylls            | Human cervical cancer | HeLa, HepG2, and Jurkat cells | In vitro | Induces apoptosis by cleavage of DNA, LDH release, activation of caspase-3, and decrease in cell count | [160] |
| β-ionone            | Shatapatri (Rosa Centifolia)      | Apocarotenoids/cyclic isoprenoid | Hepatocellular carcinoma |                | In vivo | Apoptogenic signal induction mediated by downregulation of Bcl-2 and upregulation of Bax, PPAR-γ, and FOXO-1 expressions | [161] |
| Crocin              | Kumkuma (Crocus sativus-flower)   | Apocarotenoids          | Esophageal squamous cell carcinoma | KYSE-150 | In vitro | Activated mitochondrial-mediated apoptosis pathway with an eventual disruption of MMP, increased levels of Bax and cleaved caspase-3, and decreased levels of Bcl-2 | [162] |
| Crocin              | Kumkuma (Crocus sativus-flower)   | Apocarotenoids          | Breast cancer         | HCC70, HCC1806, HeLa and CCD1059sk | In vitro | Inhibited cell proliferation mainly by disrupting the microtubule network | [163] |
| Picrocrocin         | Kumkuma (Crocus sativus-flower)   | Apocarotenoids          | Skin cancer           | SK-MEL-2         | In vitro | Targeting signaling pathway of JAK/STAT5, cell cycle arrest and mitochondrial assisted apoptosis | [164] |
| Bixin               | Sinduri (Bixa orellana L.)        | Apocarotenoids          | Cutaneous melanoma    | A2058            | In vitro | ROS-mediated cytotoxicity | [165] |
Figure 6. Chemical structures of some lead carotenoids.

6. Conclusions and Future Prospects

The importance of oxidative metabolism in the context of health and disease cannot be overlooked. All the major cellular and molecular pathways rely on redox reactions involving oxidizing and reducing equivalents such as NADH.H⁺, NADPH.H⁺, and FADH₂. For their normal functioning, cells need to maintain an equilibrium between pro-oxidants and antioxidants, failing which they stand to face consequences of oxidative stress. Overproduction of ROS is detrimental and interferes with cellular processes of DNA synthesis and repair, cell-cycle control, protein synthesis, and regulation of gene expression. These are critical mechanisms that decide the fate of a cell by influencing its survival and longevity. A cell exposed to oxidative stress is likely to undergo gene defects, apoptosis, and premature senescence. Therefore, cells are naturally equipped with mechanisms to cope with oxidative stress. In humans, glutathione, SOD, vitamin C (ascorbate), and vitamin E (tocopherol) are specifically concerned with defense of the body against oxidative stress. Herbal and ethnomedicinal drugs are usually effective against stress owing to the presence of some bioactive compounds.

The results of studies presented here are quite reassuring considering the enormous range of compounds that can be obtained from the herbs and their therapeutic potential. A number of findings presented in this review point to the fact that herbal extracts are particularly rich in compounds implicated in oxidative metabolism. In the case of many classes of phytochemicals, the basis of their therapeutic effects is known at the cellular and molecular level. However, there are many ethnomedicinal prescriptions which are considered effective against certain diseases but the exact nature of their effects is not clearly understood. In such cases, it is advisable to exercise caution because the claim that
herbal preparations are always safe is misleading in nature. Toxicity associated with many herbal medicines has been reported in numerous studies and the adverse effects of herbal drugs have recently been reviewed [166].

Therefore, considering the growing body of experimental evidence about the medicinal efficacy of herbs, it is necessary to identify their active principles responsible for such effects. This should be followed by studies to explore the mechanisms of action of active principles of interest. It is hoped that the progress in the refinement of our analytical tools will pave the way for the identification of more chemicals of therapeutic value from herbal and ethnomedicinal prescriptions. There is a hidden wealth of knowledge about naturally occurring medicinal compounds, which is yet to be explored in the years to come.

Author Contributions: N.F., S.S.R.B., A.A., S.F., P.S., K.K.K., S.R., S.H., conceptualization, methodology, investigation, formal analysis, writing—original draft preparation; N.F., S.S.R.B., A.A., S.F., S.H., methodology, investigation, formal analysis, validation; N.F., S.S.R.B., A.A., S.F., S.H., investigation, formal analysis, N.F., S.S.R.B., A.A., S.F., S.H., formal analysis, data curation; N.F., S.S.R.B., A.A., S.F., S.H., investigation, formal analysis; N.F., S.S.R.B., A.A., S.F., P.S., K.K.K., S.R., S.H., supervision, project administration, writing—review and editing. All authors have read and agreed to the published version of the manuscript.

Funding: This research received no external funding.

Acknowledgments: The authors, S.H. & A.A. sincerely acknowledge Jazan University, Saudi Arabia for providing the access of the Saudi Digital Library for this research work.

Conflicts of Interest: The authors declare no conflict of interest.

References
1. Croce, C.M. Oncogenes and cancer. N. Engl. J. Med. 2008, 358, 502–511. [CrossRef]
2. Wang, L.H.; Wu, C.F.; Rajasekaran, N.; Shin, Y.K. Loss of Tumor Suppressor Gene Function in Human Cancer: An Overview. Cell Physiol. Biochem. 2018, 51, 2647–2693. [CrossRef]
3. Bhattacharya, S.; Ghosh, M.K. Cell death and deubiquitinases: Perspectives in cancer. BioMed Res. Int. 2014, 2014, 435197. [CrossRef] [PubMed]
4. Redza-Dutordoir, M.; Averill-Bates, D.A. Activation of apoptosis signalling pathways by reactive oxygen species. Biochim. Biophys. Acta 2016, 1863, 2977–2992. [CrossRef]
5. Goodman, M. Managing the side effects of chemotherapy. Semin. Oncol. Nurs. 1989, 5 (Suppl. 2), 29–52. [CrossRef]
6. Di Gioia, F.; Tzortzakis, N.; Rouphael, Y.; Kyriacou, M.C.; Sampaio, S.L.; CFR Ferreira, I.; Petropoulos, S.A. Grown to Be Blue—Antioxidant Properties and Health Effects of Colored Vegetables. Part II: Leafy, Fruit, and Other Vegetables. Antioxidants 2020, 9, 97. [CrossRef] [PubMed]
7. Shin, S.A.; Moon, S.Y.; Kim, W.Y.; Paek, S.M.; Park, H.H.; Lee, C.S. Structure-Based Classification and Anti-Cancer Effects of Plant Metabolites. Int. J. Mol. Sci. 2018, 19, 2651. [CrossRef]
8. Reuter, S.; Gupta, S.C.; Chaturvedi, M.M.; Aggarwal, B.B. Oxidative stress, inflammation, and cancer: How are they linked? Free Radic. Biol. Med. 2010, 49, 1603–1616. [CrossRef]
9. Visconti, R.; Greco, D. New insights on oxidative stress in cancer. Curr. Opin. Drug Discov. Dev. 2009, 12, 240–245.
10. Duračková, Z. Some current insights into oxidative stress. Physiol. Res. 2010, 59, 459–469. [CrossRef] [PubMed]
11. Kurutas, E.B. The importance of antioxidants which play the role in cellular response against oxidative/nitrosative stress: Current state. Nutr. J. 2016, 15, 71. [CrossRef] [PubMed]
12. Noda, N.; Wakasugi, H. Cancer and oxidative stress. Jpn. Med. Assoc. J. 2001, 44, 535–539.
13. Sen, S.; Chakraborty, R. Revival, modernization and integration of Indian traditional herbal medicine in clinical practice: Importance, challenges and future. J. Tradit. Complement. Med. 2017, 7, 234–244. [CrossRef]
14. Vaidya, A.D.; Devasagayam, T.P. Current status of herbal drugs in India: An overview. J. Clin. Biochem. Nutr. 2007, 41, 1–11. [CrossRef] [PubMed]
15. Samal, J. Medicinal plants and related developments in India: A peep into 5-year plans of India. Indian J. Health Sci. Biomed. Res. (KLEU) 2016, 9, 14–19. [CrossRef]
16. Balachandran, P.; Govindarajan, R. Cancer—An ayurvedic perspective. Pharmacol. Res. 2005, 51, 19–30. [CrossRef]
17. Hussein, R.; El-Anssary, A. Plants Secondary Metabolites: The Key Drivers of the Pharmacological Actions of Medicinal Plants. Herb. Med. 2019, 1, 13. [CrossRef]
18. Pan, L.; Chai, H.; Kinghorn, A.D. The continuing search for antitumor agents from higher plants. Phytochem. Lett. 2010, 3, 1–8. [CrossRef]
19. Ansari, J.A.; Rastogi, N.; Ahmad, M.K.; Mahdi, A.A.; Khan, A.R.; Thakur, R.; Srivastava, V.K.; Mishra, D.P.; Fatima, N.; Khan, H.J.; et al. ROS mediated pro-apoptotic effects of Tinospora cordifolia on breast cancer cells. Front. Biosci. (Elite Ed.) 2017, 9, 89–100.
20. Kumar, B.R.; Anupam, A.; Manchikanti, P.; Rameshbabu, A.P.; Dasgupta, S.; Dhara, S. Identification and characterization of bioactive phenolic constituents, anti-proliferative, and anti-angiogenic activity of stem extracts of Basella alba and rubra. J. Food Sci. Technol. 2018, 55, 1675–1684. [CrossRef]
21. Khan, N.; Tamboli, E.; Sharma, V.K.; Kumar, S. Phytochemical and pharmacological aspects of Nothapodytes nimmoniana. An overview. Herba Pol. 2015, 59, 1. [CrossRef]
22. Tran, M.H.; Nguyen, H.D.; Kim, J.C.; Choi, J.S.; Lee, H.K.; Min, B.S. Phenolic glycosides from Alangium salviifolium leaves with inhibitory activity on LPS-induced NO, PGE(2), and TNF-alpha production. Bioorg. Med. Chem. Lett. 2009, 19, 4389–4393. [PubMed]
23. Gavande, K.; Jain, K.; Jain, B.; Mehta, R. Comprehensive Review on Phytochemistry and Pharmacological Prominence of Withania somnifera. UK J. Pharm. Biosci. 2015, 3, 15. [CrossRef]
24. Devanesan, E.B.; Anand, A.V.; Kumar, P.S.; Vinayagamoorthy, P.; Basavaraju, P. Phytochemistry and Pharmacology of Ficus religiosa. Syst. Rev. Pharm. 2018, 9, 45–48. [CrossRef]
25. Uddin, N.; Ali, N.; Uddin, Z.; Nazir, N.; Zahoor, M.; Rashid, U.; Ullah, R.; Alqahtani, A.S.; Alqahtani, A.M.; Nasr, F.A.; et al. Evaluation of Cholinesterase Inhibitory Potential of Different Genotypes of Ziziphus nummularia, Their HPLC-UV, and Molecular Docking Analysis. Molecules 2020, 25, 5011. [CrossRef]
26. Devi, S.; Rashid, R.; Kumar, M. Phytochemistry and pharmacological properties of Phyllanthus amarus Schum: A review. Pharma Innov. J. 2017, 6, 169–172. [PubMed]
27. Jayakumar, T.; Hsieh, C.Y.; Lee, J.J.; Sheu, J.R. Experimental and Clinical Pharmacology of Andrographis paniculata and Its Major Bioactive Phytoconstituent Andrographolide. Evid.-Based Complement. Altern. Med. eCAM 2013, 2013, 846740. [CrossRef] [PubMed]
28. Pandey, S.; Phulara, S.C.; Mishra, S.K.; Baipai, R.; Kumar, A.; Niranjani, A.; Lehri, A.; Upreti, D.K.; Chauhan, P.S. Betula utilis extract prolongs life expectancy, protects against amyloid-β toxicity and reduces Alpha Synuclein in Caenorhabditis elegans via DAF-16 and SKN-1. Comp. Biochem. Physiol. Toxicol. Pharmacol. CBP 2020, 228, 108647. [CrossRef]
29. Seenamisyi, R.; Faruck, L.H.; Gattu, S.; Neelamegam, R.; Bakshi, H.A.; Rashan, L.; Al-Buloshi, M.; Hasson, S.S.A.A.; Nagarajan, K. Anti-Microbial and Anti-Cancer Activity of Aegle Marmelos and Gas Chromatography Coupled Spectrometry Analysis of Their Chemical Constituents. Int. J. Pharm. Sci. Res. 2019, 10, 373–380. [PubMed]
30. Prakash, V.; Jaiswal, N.; Srivastava, M. A Review on Medicinal Properties of Centella asiatica. Asian J. Pharm. Clin. Res. 2017, 10, 69–74. [CrossRef]
31. Jeyasri, R.; Muthuramalingam, P.; Suba, V.; Ramesh, M.; Chen, J.T. Bacopa monnieri and Their Bioactive Compounds Inferred Multi-Target Treatment Strategy for Neurological Diseases: A Cheminformatics and System Pharmacology Approach. Int. J. Pharm. Tech. 2020, 10, 536. [CrossRef]
32. Koczurkiewicz, P.; Łojewski, M.; Piska, K.; Michalik, M.; Wójcik-Pszczola, K.; Szewczyk, A.; Hałaszuk, P.; Pękala, E.; Muszyńska, B. Chemopreventive and Anticancer Activities of Bacopa monnieri Extracted from Artificial Digestive Juices. Nat. Prod. Commun. 2017, 12, 337–342. [CrossRef]
33. Aswal, J.; Dobhal, R.; Uniyal, D.P.; Nautiyal, V. A review on Pharmacological potential of Berberine; an active component of Himalayan Berberis aristata. J. Phytopharmacol. 2017, 6, 53–58. [CrossRef]
34. Al-Snafi, A. Medical importance of Datura fastuosa (syn: Datura metel) and Datura stramonium—A review. IOSR J. Pharm. 2017, 7, 43–58. [CrossRef]
35. Okoro, E.E.; Osoniyi, O.R.; Jabeen, A.; Shams, S.; Choudhary, M.I.; Onajobi, F.D. Anti-proliferative and immunomodulatory activities of fractions from root extract of Abrus precatorius L. Clin. Phytoscience 2019, 5, 45. [CrossRef]
36. Tomeh, M.A.; Hadianamrei, R.; Zhao, X. A Review of Curcumin and Its Derivatives as Anticancer Agents. Int. J. Mol. Sci. 2019, 20, 1033. [CrossRef] [PubMed]
37. Rathinamourthy, R.; Thilagavathi, G. Terminalia chebula-review on pharmacological and biochemical studies. Int. J. Pharm. Tech. Res. 2014, 6, 97–116. [PubMed]
38. Fatima, N.; Ahmad, M.; Ansari, J.; Khan, H.; Rastogi, N.; Srivastava, S.; Ahmad, S.; Ali, Z. Antiproliferative and Antioxidant Studies of Anthocephalus cadamba Rox. Miq. Bark. Indian J. Pharm. Sci. 2016, 78, 525–531. [CrossRef]
39. Khan, H.J.; Ahmad, M.K.; Khan, A.R.; Rastogi, N.; Mahdi, A.A.; Ansari, J.A.; Fatima, N.; Satyanarayan, G.N.V. Identification of Anticancer and Antioxidant phytoconstituents from chloroform fraction of Solanum nigrum L. berries using GC-MS/MS analysis. Indian J. Exp. Biol. 2016, 54, 774–782. [PubMed]
40. Pal, P.K.; Nandi, M.K.; Singh, N.K. Detoxification of Croton tiglium L. seeds by Ayurvedic process of Šodhana. Anc. Sci. Life 2014, 33, 157. [PubMed]
41. Bukhari, S.I.; Manzoor, M.; Dhar, M.K. A comprehensive review of the pharmacological potential of Crocus sativus and its bioactive apocarotenoids. Biomed. Pharmacother. 2018, 98, 733–745. [CrossRef]
42. Coria-Téllez, A.V.; Montalvo-Gonzalez, E.; Yahia, E.M.; Obledo-Vázquez, E.N. Annona muricata: A comprehensive review on its traditional medicinal uses, phytochemicals, pharmacological activities, mechanisms of action and toxicity. Arab. J. Chem. 2018, 11, 662–691. [CrossRef]
43. El-Saber Batia, G.; Alakzmi, L.M.; Wasef, L.G.; Beshbishy, A.M.; Nadwa, E.H.; Rashwan, E.K. Syzygium aromaticum L. (Myrtaceae): Traditional Uses, Bioactive Chemical Constituents, Pharmacological and Toxicological Activities. Biomolecules 2020, 10, 202. [CrossRef] [PubMed]
44. Braga, T.M.; Rocha, L.; Chung, T.Y.; Oliveira, R.F.; Pinho, C.; Oliveira, A.L.; Morgado, J.; Cruz, A. Azadirachta indica A. Juss In Vivo Toxicity—An Updated Review. Molecules 2021, 26, 252. [CrossRef]

45. Chihikara, N.; Kour, R.; Jaglan, S.; Gupta, P.; Gai, Y.; Panghal, A. Citrus medica: Nutritional, phytochemical composition and health benefits—A review. Food Funct. 2018, 9, 1978–1992. [CrossRef]

46. Marrelli, M.; Amodeo, V.; Statti, G.; Conforti, F. Biological Properties and Bioactive Components of Allium cepa L.: Focus on Potential Benefits in the Treatment of Obesity and Related Comorbidities. Molecules 2019, 24, 119. [CrossRef]

47. Yadav, V.; Krishnan, A.; Vohora, D. A systematic review on Piper longum L.: Bridging traditional knowledge and pharmacological evidence for future translational research. J. Ethnopharmacol. 2020, 247, 112255. [CrossRef]

48. Anirban, C.; Santanu, F. A Review on Phytochemical and Pharmacological Potential of Alpinia galanga. Pharmacogn. J. 2018, 10, 9–15.

49. Alam, K.; Hoq, O.; Uddin, S. Medicinal plant Allium sativum: A Review. J. Med. Plants Stud. 2016, 4, 72–79.

50. Zargar, B.A.; Masoodi, M.H.; Ahmed, B.; Ganie, S.A. Phytoconstituents and therapeutic uses of Rheum emodi wall. ex Meissn. Food Chem. 2011, 128, 585–589. [CrossRef]

51. Dixit, V.; Jain, P.; Joshi, S.; Joshi, S. Hypolipidaemic effects of Curcuma longa Linn., and Nardostachys jatamansi DC and Its Phytochemical Analysis by RP-HPLC and GC-MS. Antioxidants 2015, 4, 185–203. [CrossRef]

52. Ansari, J.A.; Ahmad, M.K.; Khan, A.R.; Fatima, N.; Khan, H.J.; Rastogi, N.; Mishra, D.P.; Mahdi, A.A. Anticancer and Antioxidant activity of Zingiber officinale Roscoe rhizome. Indian J. Exp. Biol. 2016, 54, 767–773. [PubMed]

53. Joshi, R.K.; Setzer, W.N.; Da Silva, J.K. Phytoconstituents, traditional medicinal uses and bioactivities of Tulsi (Ocimum sanctum Linn.): A review. Am. J. Essent. 2017, 5, 18–21.

54. Kami, S.S.; Hameed, I.H.; Hamza, L.F. Acorus calamus: Parts used, insecticidal, anti-fungal, antitumour and anti-inflammatory activity: A review. Int. J. Pharm. Clin. Res. 2018, 10, 153–157.

55. Razack, S.; Kumar, K.H.; Nallamuthu, I.; Naika, M.; Khanum, F. Antioxidant, Biomolecule Oxidation Protective Activities of Nardostachys jatamansi DC and Its Phytochemical Analysis by RP-HPLC and GC-MS. Antioxidants 2015, 4, 185–203. [CrossRef]

56. Pandey, M.M.; Katara, A.; Pandey, G.; Rastogi, S.; Rawat, A.K.S. An Important Indian Traditional Drug of Ayurveda Jatamansi and Its Substitutes Bhootkeshi: Chemical Profiling and Antioxidant Activity. Evid.-Based Complement. Altern. Med. 2013, 2013, 142517. [CrossRef]

57. Dixit, V.; Jain, P.; Joshi, S. Hypolipidaemic effects of Curcuma longa Linn., and Nardostachys jatamansi DC, in triton-induced hyperlipidaemic rats. Indian J. Pharm. Pharmacol. 1988, 32, 299–304.

58. Pillai, D.; Pandita, N. Determination of Quality Standards for Drakasharisha, a Polyherbal Ayurvedic Formulation. Indian J. Pharm. Sci. 2016, 78, 129–135.

59. Sathiya, M. Scientific Evaluation of Antioxidant and Anti Cancer Activity of Kanchanara Guggulu Vati by Invitro Methods. Ph.D. Thesis, Madras Medical College, Chennai, India, 2017.

60. Sarawathy, A.; Sundaresan, R.; Joy, S.; Gopal, R.H. Effect of Container on Ayurvedic Drugs—A Select Study. Anc. Sci. Life 2004, 24, 11.

61. Albert, J.; Albert, L.; Albert, B. Therapeutic uses of triphala in ayurvedic medicine. J. Altern. Complement. Med. 2017, 23, 607–614. [CrossRef]

62. Kotteswari, M.; Rao, M.; Kumar, S.; Prabhu, K.; Sundaram, R.L.; Dinakar, S. GC MS Analysis of One Ayurvedic Preparation Azadirachta indica DC, in triton-induced hyperlipidaemic rats. Indian J. Pharm. Clin. Res. 2015, 9, 76–81. [PubMed]

63. Patil, V.; Parmar, N. A Comparative Clinical Study of Kasisadi Taila and Jatyadi Taila in the Management of Arsha. J. Ayurveda Holist. Med. 2017, 5, 16–24.

64. Jadhav, A.N.; Bhutani, K. Ayurveda and gynecological disorders. J. Ethnopharmacol. 2005, 97, 151–159. [CrossRef]

65. Govardhan, B.; Manjunatha, A.; Kumar, S.S. Standardization of Lashuna taila: An ayurvedic oil based medicine. J Pharmacogn. Phytochem. 2018, 7, 28338.

66. Ferguson, L.R.; Chen, H.; Collins, A.R.; Connell, M.; Damia, G.; Dasgupta, S.; Malhotra, M.; Meeker, A.K.; Ameedi, A.; Amin, A.; et al. Genomic instability in human cancer: Molecular insights and opportunities for therapeutic attack and prevention through diet and nutrition. Semin. Cancer Biol. 2015, 35, S5–S24. [CrossRef] [PubMed]

67. Forni, C.; Facchiano, F.; Bartoli, M.; Pieretti, S.; Facchiano, A.; D’Arcangelo, D.; Norelli, S.; Valle, G.; Nisini, R.; Beninati, S.; et al. Beneficial Role of Phytochemicals on Oxidative Stress and Age-Related Diseases. BioMed Res. Int. 2019, 2019, 8748253. [CrossRef] [PubMed]

68. Zhang, Y.-J.; Gan, R.-Y.; Li, S.; Zhou, Y.; Li, A.-N.; Xu, D.-P.; Li, H.-B. Antioxidant Phytochemicals for the Prevention and Treatment of Chronic Diseases. Molecules 2015, 20, 21138–21156. [CrossRef] [PubMed]

69. Mileo, A.M.; Miccadi, S. Polyphenols as Modulator of Oxidative Stress in Cancer Disease: New Therapeutic Strategies. Oxidative Med. Cell. Longev. 2016, 2016, 6475624. [CrossRef] [PubMed]

70. Choi, D.G.; Venkatesan, J.; Shim, M.S. Selective Anticancer Therapy Using Pro-Oxidant Drug-Loaded Chitosan–Fucoidan Nanoparticles. Int. J. Mol. Sci. 2019, 20, 3220. [CrossRef] [PubMed]

71. Kocyigit, A.; Guler, E.M.; Dikilitas, M. Role of antioxidant phytochemicals in prevention, formation and treatment of cancer. In Reactive Oxygen Species (ROS) in Living Cells; InterchOpen: London, UK, 2018; pp. 21–45.

72. Swallah, M.S.; Sun, H.; Affoh, R.; Fu, H.; Yu, H. Antioxidant potential overviews of secondary metabolites (polyphenols) in fruits. Int. J. Food Sci. 2020, 2020, 9081686. [CrossRef]
100. Tian, X.; Zhang, S.; Zhang, Q.; Kang, L.; Ma, C.; Feng, L.; Li, S.; Li, J.; Yang, L.; Liu, J.; et al. Resveratrol inhibits tumor progression by down-regulation of NLRP3 in renal cell carcinoma. J. Nutr. Biochem. 2020, 85, 108489. [CrossRef]

101. Soli, F.; Ternet, C.; Berry, I.M.; Kumari, D.; Moore, T.C. Quercetin Inhibits Proliferation and Induces Apoptosis of B16 Melanoma Cells In Vitro. Assay Drug Dev. Technol. 2020, 18, 261–268. [CrossRef]

102. Salama, A.A.A.; Allam, R.M. Promising targets of chrysins and daidzein in colorectal cancer: Amphiregulin, CXCL1, and MMP-9. Eur. J. Pharmacol. 2021, 892, 173763. [CrossRef]

103. Jang, Y.G.; Ko, E.B.; Choi, K.C. Gallic acid, a phenolic acid, hinders the progression of prostate cancer by inhibition of histone deacetylase 1 and 2 expression. J. Nutr. Biochem. 2020, 84, 108444. [CrossRef]

104. Habli, Z.; Toumiez, G.; Fatfat, M.; Rahal, O.N.; Gali-Muhtasib, H. Emerging Cytotoxic Alkaloids in the Battle against Cancer: Overview of Molecular Mechanisms. Molecules 2017, 22, 250. [CrossRef] [PubMed]

105. Mohammadlou, M.; Abdollahi, M.; Hemati, M.; Baharlou, R.; Doulabi, E.M.; Pashaei, M.; Ghahremanfard, F.; Faranoush, M.; Safia; Kamil, M.; Jadiya, P.; Sheikh, S.; Haque, E.; Nazir, A.; Lakshmi, V.; Mir, S.S. The Chromone Alkaloid, Rohitukine, Affords apoptotic inducing activity of Amoora rohituka leaf extracts in human breast cancer cells. J. Ayurveda Integr. Med. 2020, 11, 383–390. [CrossRef] [PubMed]

106. Singh, R.K.; Ranjan, A.; Srivastava, A.K.; Singh, M.; Shukla, A.K.; Atri, N.; Mishra, A.; Singh, A.K.; Singh, S.K. Cytotoxic and apoptotic inducing activity of Amoora rohituka leaf extracts in human breast cancer cells. J. Ayurveda Integr. Med. 2020, 11, 383–390. [CrossRef] [PubMed]

107. Wiernik, P.H. Alvocidib (flavopiridol) for the treatment of chronic lymphocytic leukemia. Expert Opin. Investig. Drugs 2016, 25, 729–734. [CrossRef]

108. Safia; Kamil, M.; Jadiya, P.; Sheikh, S.; Haque, E.; Nazir, A.; Lakshmi, V.; Mir, S.S. The Chromone Alkaloid, Rohitukine, Affords Anti-Cancer Activity via Modulating Apoptosis Pathways in A549 Cell Line and Yeast Mitogen Activated Protein Kinase (MAPK) Pathway. PLoS ONE 2015, 10, e0137991. [CrossRef] [PubMed]

109. Liu, Y.; Hua, W.; Li, Y.; Xian, X.; Zhao, Z.; Liu, C.; Zou, J.; Li, J.; Fang, X.; Zhu, Y. Berberine suppresses colon cancer cell proliferation by inhibiting the SCAP/SREBP-1 signaling pathway-mediated lipogenesis. Biochem. Pharmacol. 2020, 174, 113776. [CrossRef]

110. Harmon, A.D.; Weiss, U.; Silverton, J. The structure of rohitukine, the main alkaloid of Amoora rohituka (syn. Aphanamixis polystachya)(Meliaceae). Tetrahedron Lett. 1979, 20, 721–724. [CrossRef]

111. Isah, T. Anticancer Alkaloids from Trees: Development into Drugs. Phytother. Res. PTR 2010, 24, 892–900. [CrossRef] [PubMed]

112. Saisomboon, S.; Kariya, R.; Vaeteewoottacharn, K.; Wongkham, S.; Sawanyawisuth, K.; Okada, S. Antitumor effects of flavopiridol, a cyclin-dependent kinase inhibitor, on human cholangiocarcinoma in vitro and in an in vivo xenograft model. Heliotrop 2019, 5, e01675. [CrossRef]

113. Alviocidib (flavopiridol) for the treatment of chronic lymphocytic leukemia. Expert Opin. Investig. Drugs 2016, 25, 729–734. [CrossRef]

114. Srivastava, S.; Rawat, A.K. Quality evaluation of ayurvedic crude drug daruharidra, its allied species, and commercial samples from herbal drug markets of India. Evid.-Based Complement. Altern. Med. eCAM 2013, 2013, 472973. [CrossRef] [PubMed]

115. Akhtar, S.; Achkar, I.W.; Siveen, K.S.; Kuttikrishnan, S.; Prabhu, K.S.; Khan, A.Q.; Ahmed, E.I.; Sahir, F.; Jerobin, J.; Raza, A.; et al. Anticancer Activity via Modulating Apoptosis Pathways in A549 Cell Line and Yeast Mitogen Activated Protein Kinase (MAPK) Pathway. J. Nutr. Biochem. 2020, 84, 108444. [CrossRef]

116. Zhu, M.; Yu, X.; Zheng, Z.; Huang, J.; Yang, X.; Shi, H. Capsaicin suppressed activity of prostate cancer stem cells by inhibition of Wnt/β-catenin pathway. Phytother. Res. PTRY 2021, 35, 2025–2033. [CrossRef]

117. Akhtar, S.; Achkar, I.W.; Siveen, K.S.; Kuttikrishnan, S.; Prabhu, K.S.; Khan, A.Q.; Ahmed, E.I.; Sahir, F.; Jerobin, J.; Raza, A.; et al. Sanguinarine Induces Apoptosis Pathway in Multiple Myeloma Cell Lines via Inhibition of the JaK2/STAT3 Signaling. Front. Oncol. 2019, 9, 285. [CrossRef] [PubMed]

118. Wang, C.H.; Yang, J.M.; Guo, Y.B.; Shen, J.; Pei, X.H. Anticancer Activity of Tetrandrine by Inducing Apoptosis in Human Breast Cancer Cell Line MDA-MB-231 In Vivo. Evid.-Based Complement. Altern. Med. eCAM 2020, 2020, 6823520. [CrossRef]

119. Bhagya, N.; Chandrashekar, K.R.; Prabhu, A.; Rekha, P.D. Tetrandrine isolated from Cyclea peltata induces cytotoxicity and caspase pathways in breast and pancreatic cancer cells. In Vitro Cell. Dev. Biol. Anim. 2019, 55, 331–340. [CrossRef] [PubMed]

120. Portenoy, R.K. Opioid therapy for chronic nonmalignant pain: A review of the critical issues. J. Pain Symptom Manag. 1996, 11, 203–217. [CrossRef]

121. Practice guidelines for chronic pain management. A report by the American Society of Anesthesiologists Task Force on Pain Management, Chronic Pain Section. Anesthesiology 1997, 86, 995–1004.

122. Gutstein, H.; Akil, H. Opioid Analgesics in the Pharmacological Basis of Therapeutics; McGraw-Hill: New York, NY, USA, 2001.

123. Boeing, H.; Bechthold, A.; Bub, A.; Ellinger, S.; Haller, D.; Kroke, A.; Leschik-Bonnet, E.; Müller, M.J.; Oberritter, H.; Schulze, M.; et al. Critical review: Vegetables and fruit in the prevention of chronic diseases. Eur. J. Nutr. 2012, 51, 637–663. [CrossRef]

124. Corzo-Martinez, M.; Corzo, N.; Villamiel, M. Biological properties of onions and garlic. Trends Food Sci. Technol. 2007, 18, 609–625. [CrossRef]
126. Goncharov, N.; Belinskaia, D.; Ukolov, A.; Jenkins, R.; Avdonin, P. *Organosulfur Compounds as Nutraceuticals*; Academic Press: New York, NY, USA, 2021; pp. 911–924.

127. Farhat, Z.; Hershberger, P.A.; Freemanheim, J.L.; Mammen, M.J.; Hageman Blair, R.; Aga, D.S.; Mu, L. Types of garlic and their antioxidant activity: A review of the epidemiologic and experimental evidence. *Eur. J. Nutr.* 2021, 60, 3585–3609. [CrossRef]

128. Joshi, V.; Joshi, A. *Garlic in Traditional Indian Medicine (Ayuurveda) for Health and Healing*; IntechOpen: London, UK, 2021. [CrossRef]

129. Rosas-González, V.C.; Téllez-Bañuelos, M.C.; Hernández-Flores, G.; Bravo-Cuellar, A.; Aguilar-Lemarroy, A.; Jave-Suárez, L.F.; Haramati, J.; Solorzano-Ibarra, F.; Ortiz-Lazareno, P.C. Differential effects of alliin and allicin on apoptosis and senescence in luminal A and triple-negative breast cancer: Caspase, ΔΨm, and pro-apoptotic gene involvement. *Fundam. Clin. Pharmacol.* 2020, 34, 671–686. [CrossRef]

130. Li, C.; Jing, H.; Ma, G.; Liang, P. Allicin induces apoptosis through activation of both intrinsic and extrinsic pathways in glioma cells. *Mol. Med. Rep.* 2018, 17, 5976–5981. [CrossRef]

131. Miękus, N.; Marszałek, K.; Podlacha, M.; Iqbal, A.; Puchalski, C.; Świergiel, A.H. Health Benefits of Plant-Derived Sulfur Compounds, Glucosinolates, and Organosulfur Compounds. *Molecules* 2020, 25, 3804. [CrossRef]

132. Das, A.; Banik, N.L.; Ray, S.K. Garlic compounds generate reactive oxygen species leading to activation of stress kinases and cysteine proteases for apoptosis in human glioblastoma T98G and U87MG cells. *Cancer 2007, 110*, 1083–1095. [CrossRef]

133. Wang, H.C.; Yang, J.H.; Hsieh, S.C.; Sheen, L.Y. Allyl sulfides inhibit cell growth of skin cancer cells through induction of DNA damage mediated G2/M arrest and apoptosis. *J. Agric. Food Chem.* 2010, 58, 7096–7103. [CrossRef]

134. Kanamori, Y.; Via, L.D.; Macone, A.; Canettieri, G.; Greco, A.; Toninello, A.; Agostinelli, E. Aged garlic extract and its constituent, S-allyl-L-cysteine, induce the apoptosis of neuroblastoma cancer cells due to mitochondrial membrane depolarization. *Exp. Ther. Med.* 2020, 19, 1511–1521. [CrossRef]

135. Khan, F.; Pandey, P.; Mishra, R.; Arif, M.; Kumar, A.; Jafri, A.; Mazumder, R. Elucidation of S-Allylcysteine Role in Inducing Apoptosis by Inhibiting PD-L1 Expression in Human Lung Cancer Cells. *Anti-Cancer Agents Med. Chem.* 2021, 21, 532–541. [CrossRef]

136. Jobani, B.M.; Najafzadeh, N.; Mazani, M.; Arzanlou, M.; Vardin, M.M. Molecular mechanism and cytotoxicity of allillic and all-trans retinoic acid against CD44(+) versus CD117(+) melanoma cells. *Phytochem. Int. J. Phytother. Phytopharm.* 2018, 48, 161–169. [CrossRef] [PubMed]

137. Chen, H.; Zhu, B.; Zhao, L.; Liu, Y.; Zhao, F.; Feng, J.; Jin, Y.; Sun, J.; Geng, R.; Wei, Y. Allicin Inhibits Proliferation and Invasion in Vitro and in Vivo via SHP-1-Mediated STAT3 Signaling in Cholangiocarcinoma. *Cell. Physiol. Biochem.* 2018, 47, 641–653. [CrossRef] [PubMed]

138. Wang, K.; Wang, Y.; Qi, Q.; Zhang, F.; Zhang, Y.; Zhu, X.; Liu, G.; Luan, Y.; Zhao, Z.; Cai, J.; et al. Inhibitory effects of S-allylmercaptocysteine against benzo(a)pyrene-induced precancerous carcinogenesis in human lung cells. *Int. Immunopharmacol.* 2016, 34, 37–43. [CrossRef] [PubMed]

139. Iida, Y.; Okamoto-Katsuyama, M.; Maruoka, S.; Mizumura, K.; Shimizu, T.; Shikano, S.; Hikichi, M.; Takahashi, M.; Tsuya, K.; Okamoto, S.; et al. Effective ferroptotic small-cell lung cancer cell death from SLC7A11 inhibition by sulforaphane. *Oncol. Lett.* 2021, 21, 71. [CrossRef] [PubMed]

140. Fernandes, A.S.; do Nascimento, T.C.; Jacob-Lopes, E.; De Rosso, V.V.; Zepka, L.Q. Carotenoids: A brief overview on its structure, biosynthesis, synthesis, and applications. In *Progress in Carotenoid Research*; IntechOpen: London, UK, 2018; pp. 1–15. [CrossRef]

141. Giuliano, G.; Al-Babili, S.; von Lintig, J. Carotenoid oxygenases: Cleave it or leave it. *Trends Plant Sci.* 2015, 20, 968–985. [CrossRef] [PubMed]

142. Ramel, F.; Birtic, S.; Cuin, S.; Triantaphylidès, C.; Ravanat, J.-L.; Havaux, M. Chemical Quenching of Singlet Oxygen by Carotenoids in Plants. *Plant Physiol.* 2012, 158, 1267–1278. [CrossRef]

143. Black, H.S.; Boehm, F.; Edge, R.; Truscott, T.G. The Benefits and Risks of Certain Dietary Carotenoids that Exhibit both Anti- and Pro-Oxidative Mechanisms—A Comprehensive Review. *Antioxidants 2020*, 9, 264. [CrossRef]

144. Niranjana, R.; Gayathri, R.; Nimish Mol, S.; Sugawara, T.; Hirata, T.; Miyashita, K.; Ganesan, P. Carotenoids modulate the hallmarks of cancer cells. *J. Funct. Foods* 2015, 18, 968–985. [CrossRef]

145. Toti, E.; Chen, C.O.; Palmyre, M.; Villan, D.; Peluso, I. Non-Provitamin A and Provitamin A Carotenoids as Immunomodulators: Recommended Dietary Allowance, Therapeutic Index, or Personalized Nutrition? *Oxidative Med. Cell. Longev.* 2018, 2018, 4637861. [CrossRef]

146. IARC Working Group on the Evaluation of Cancer-Preventive Agents. *IARC Handbooks of Cancer Prevention: Carotenoids*; IARC: Lyon, France, 1998.

147. Omenn, G.S.; Goodman, G.E.; Thornquist, M.D.; Balmes, J.; Cullen, M.R.; Glass, A.; Keogh, J.P.; Meykens, F.L.; Valanis, B.; Williams, J.H.; et al. Effects of a combination of beta carotene and vitamin A on lung cancer and cardiovascular disease. *N. Engl. J. Med.* 1996, 334, 1150–1155. [CrossRef]

148. Sathasivam, R.; Ki, J.-S. A Review of the Biological Activities of Microalgal Carotenoids and Their Potential Use in Healthcare and Cosmetic Industries. *Marine Drugs* 2018, 16, 26. [CrossRef]
150. Takata, Y.; Xiang, Y.-B.; Yang, G.; Li, H.; Gao, J.; Cai, H.; Gao, Y.-T.; Zheng, W.; Shu, X.-O. Intakes of fruits, vegetables, and related vitamins and lung cancer risk: Results from the Shanghai Men's Health Study (2002–2009). *Nutr. Cancer* 2013, 65, 51–61. [CrossRef] [PubMed]

151. Tanaka, T.; Shinmizu, M.; Moriwaki, H. Cancer chemoprevention by carotenoids. *Molecules* 2012, 17, 3202–3242. [CrossRef] [PubMed]

152. Liu, Y.Z.; Yang, C.M.; Chen, J.Y.; Liao, J.W.; Hu, M.L. Alpha-carotene inhibits metastasis in Lewis lung carcinoma in vitro, and suppresses lung metastasis and tumor growth in combination with taxol in tumor xenografted C57BL/6 mice. *J. Nutr. Biochem.* 2015, 26, 607–615. [CrossRef] [PubMed]

153. Kim, D.; Lim, J.W.; Kim, H. β-carotene Inhibits Expression of c-Myc and Cyclin E in Helicobacter pylori-infected Gastric Epithelial Cells. *J. Cancer Prev.* 2019, 24, 192–196. [CrossRef] [PubMed]

154. Cui, L.; Xu, F.; Wu, K.; Li, L.; Qiao, T.; Li, Z.; Chen, T.; Sun, C. Anticancer effects and possible mechanisms of lycopene intervention on N-methylbenzylnitrosamine induced esophageal cancer in F344 rats based on PPARγ. *Eur. J. Pharmacol.* 2020, 881, 173230. [CrossRef] [PubMed]

155. Aktepe, O.H.; Şahin, T.K.; Güner, G.; Arik, Z.; Yalçın, Ş. Lycopene sensitizes the cervical cancer cells to cisplatin via targeting nuclear-factor- kappa B (NF-kB) pathway. *Turk. J. Med. Sci.* 2021, 51, 368–374.

156. Gansukh, E.; Mya, K.K.; Jung, M.; Keum, Y.-S.; Kim, D.H.; Saini, R.K. Lutein derived from marigold (*Tagetes erecta*) petals triggers ROS generation and activates Bax and caspase-3 mediated apoptosis of human cervical carcinoma (HeLa) cells. *Food Chem. Toxicol.* 2019, 127, 11–18. [CrossRef] [PubMed]

157. Sheng, Y.-N.; Luo, Y.-H.; Liu, S.-B.; Xu, W.-T.; Zhang, Y.; Zhang, T.; Xue, H.; Zuo, W.-B.; Li, Y.-N.; Wang, C.-Y.; et al. Zeaxanthin Induces Apoptosis via ROS-Regulated MAPK and AKT Signaling Pathway in Human Gastric Cancer Cells. *Onco Targets Ther.* 2020, 13, 10995–11006. [CrossRef]

158. Gansukh, E.; Nile, A.; Sivanesan, I.; Rengasamy, K.R.R.; Kim, D.-H.; Keum, Y.-S.; Saini, R.K. Chemopreventive Effect of β-Cryptoxanthin on Human Cervical Carcinoma (HeLa) Cells Is Modulated through Oxidative Stress-Induced Apoptosis. *Antioxidants* 2020, 9, 28. [CrossRef] [PubMed]

159. Wu, H.L.; Fu, X.Y.; Cao, W.Q.; Xiang, W.Z.; Hou, Y.J.; Ma, J.K.; Wang, Y.; Fan, C.D. Induction of Apoptosis in Human Glioma Cells by Fucoxanthin via Triggering of ROS-Mediated Oxidative Damage and Regulation of MAPKs and PI3K-AKT Pathways. *J. Agric. Food Chem.* 2019, 67, 2212–2219. [CrossRef] [PubMed]

160. Shukla, M.; Varalakshmi, K.N. Apoptosis induction in cancer cell lines by the carotenoid Fucoxanthinol from Pseudomonas stutzeri JGI 52. *Indian J. Pharmacol.* 2018, 50, 116. [PubMed]

161. Abd-Elbaset, M.; Mansour, A.M.; Ahmed, O.M.; Abo-Youssef, A.M. The potential chemotherapeutic effect of β-ionone and/or sorafenib against hepatocellular carcinoma via its antioxidant effect, PPAR-γ, FOXO-1, Ki-67, Bax, and Bcl-2 signaling pathways. *Naunyn-Schmiedeberg's Arch. Pharmacol.* 2020, 393, 1611–1624. [CrossRef] [PubMed]

162. Li, S.; Qu, Y.; Shen, X.Y.; Ouyang, T.; Fu, W.B.; Luo, T.; Wang, H.Q. Multiple Signal Pathways Involved in Crocetin-Induced Apoptosis in KYSE-150 Cells. *Pharmacology 2019*, 103, 263–272. [CrossRef]

163. Hire, R.R.; Srivastava, S.; Davis, M.B.; Kumar Konreddy, A.; Panda, D. Antiproliferative Activity of Crocin Involves Targeting of Microtubules in Breast Cancer Cells. *Sci. Rep.* 2017, 7, 44984. [CrossRef] [PubMed]

164. Yu, L.; Li, J.; Xiao, M. Picrocrocin exhibits growth inhibitory effects against SKMEL-2 human malignant melanoma cells by targeting JAK/ STAT5 signaling pathway, cell cycle arrest and mitochondrial mediated apoptosis. *J. B.U.ON. Off. J. Balk. Union Oncol.* 2018, 23, 1163–1168.

165. De Oliveira Júnior, R.G.; Bonnet, A.; Bracconier, E.; Grout, H.; Prunier, G.; Beaugeste, L.; Grumet, R.; da Silva Almeida, J.R.G.; Ferraz, C.A.A.; Picot, L. Bixin, an apocarotenoid isolated from *Bixa orellana L.*, sensitizes human melanoma cells to dacarbazine-induced apoptosis through ROS-mediated cytotoxicity. *Food Chem. Toxicol.* 2019, 125, 549–561. [CrossRef]

166. Posadzki, P.; Watson, L.K.; Ernst, E. Adverse effects of herbal medicines: An overview of systematic reviews. *Clin. Med.* 2013, 13, 7–12. [CrossRef]