Abstract

Ayurveda, “life knowledge”, is an ancient Indian traditional medical system, which has been practiced for >5000 years and still in use with the same importance among many cultural tribes in Indian sub-continents. Ayurvedic medicine is a unique holistic approach to treatment where herbal medicines, special diets, yoga and relaxation methods, and lifestyle management are key strategies for curing various chronic diseases like diabetes, cancer, cardiovascular, neurological disorders and many other. Cancer is biologically complex disease, causing local damage and inflammation of cells due to lack of full response in a specific tissue that is responsible to maintain cellular differentiation, survival, proliferation and death. Studies have suggested that bioactive phytochemicals present in Ayurvedic plants (fruits, vegetables and herbs) mediate their effects by inhibiting some of the recently identified inflammatory pathways. Nevertheless, for successful implementation of Ayurvedic principles to drug discovery, major concerns relating to proportion, practicality, safety and drug interactions and possible side effects of newly developed Ayurvedic drugs and formulations need to be addressed. Systematic investigations of Ayurvedic drugs, employing contemporary scientific tools and methods, are expected to lead to the discoveries of major significance. Explorations involving studies of bioactive phytochemicals, their side effects, the specific potential targets and the mechanisms of action would transform this traditional therapeutic approach to standard Ayurvedic drug discovery which would be accepted and embraced by global populations. Current review focuses on the importance of anti-cancer drug discovery from Ayurvedic medicines in the modern context, ongoing improvements and major achievements in the field over the past decade or two.

Keywords: Ayurveda; Harmony; Balance

Introduction

“Ayurveda” – ayus (life) and veda (science/knowledge) means “the science of life”, has unbroken long history as ancient Indian medicinal system for five thousand years. Ayurveda is often known as the “Mother of all healing” [1] that brings harmony and balance in all areas of life including mind, body, spiritual well-being and social welfare of mankind [2].

According to Ayurveda, the primary components of human beings and universe is one or combination of the five basic elements (Panchamahabhutas) [3], namely: Pritivi (earth - the principle of inertia), Ap (water - the principle of cohesion), Teja (fire - the principle of radiance), Vayu (air - the principle of vibration), Akasha (ether - the principle of pervasiveness). These five basic principles form three vital forces called Tridoshas: Vata (maintaining nervous system responsible for cell growth, differentiation, cell death, movement of cells, molecules, nutrients and waste), Pitta (maintaining venous system for chemical reactions in the body including digestion, metabolic process, immunity and temperature control) and Kapha (stream nutrition to the arterial system responsible for structure, growth, and protection) [3].

Physical, mental and physiological features (constitution or Prakriti) of every individual are distinctly different from one another. In Ayurvedic medical practice, the combination of Tridosha and panchamahabhutas are used to determine the nature (Prakriti) of an individual, accordingly, the treatments are individualized [3]. For example, alterations in “Pitta dosha” is allied with inflammatory diseases [4] resulting from weakened organs (Dhatu), and hence “Rasayanaprayoga” (immunotherapy) is commonly practiced for rejuvenating the body’s support system [5]. Ayurvedic medicine is a holistic and multidimensional medical system with built capacity to diagnose and treat various illnesses including chronic diseases such as cardiovascular, neurological, diabetes, cancer and many other diseases.

In the context of Ayurveda, a good health is defined as balanced situation of Tridoshas. Imbalance in these vital forces causes diseases and the specific treatment plan for any particular imbalance is defined and practiced in Ayurveda from vedic era. As illustrated in (Figure 1A). The treatment modality involves a combination of different approaches including yoga, dietary control, meditation, prayers and other means. However, medicinal herbs received greater prominence than other mode of treatments due to their efficacy and prompt curative properties. Globally, herbs and their derivatives became major component of different traditional medicinal systems to treat diseases. Modern scientific and technological advancements have the potential to transform the traditional knowledge of medicinal plants into modern drugs, as described in (Figure 1 B). Many medicinal plant derived drugs such as aspirin, reserpine, and digitalis are a few
examples of therapeutic agents discovered from medicinal plants [6]. However, more systematic and robust research is required to derive full benefits of traditional herbs and to discover effective therapeutic agents for life threatening diseases.

Ayurvedic formulations have been the first line of treatment in Indian subcontinent for thousands of years due to their lesser toxicity and wide acceptability [7]. The emerging chemotherapeutic approaches have hampered the practice of Ayurvedic medicine system during the past couple of decades. This is primarily due to the effectiveness of medicinal herbs on disease symptoms is relatively slower than chemotherapeutic drugs. Apparent success of chemotherapeutic drugs, in the treatment of cancer and other serious diseases, has been masked by their side effects and toxicity which have become major downside. In addition, chemotherapy approaches often are not successful for the treatment of major death causing diseases including cancer. This situation has primarily lead to the renewed interest in complementary therapies as a pathway for finding cure for cancer.

In Ayurveda, Charaksha and Sushruta Samhitas define cancer as inflammatory or non-inflammatory swelling often known as Granthi (minor neoplasm) or Arbuda (major neoplasm) [8]. Minor neoplasm is mainly caused by weak Shukra Dhatu (tissue regeneration and cell division) interrelated with imbalanced Kapha and/or Vata Dosha. However, missing coordination amongst all these three vital forces is responsible for unhealthy and maline tridoshic major tumor [9]. In principle, it is caused due to unhealthy lifestyle, such as consumption of unhealthy foods, poor hygiene, or bad habits/addiction, or from physical trauma, causing major discrepancies in Tridoshas (Vata, Pitta, and Kapha) [10]. Inhibition of the tumoral growth and their destruction is accomplished by making changes in metabolic defects and restoring normal tissue functions by improving lifestyle, known as “Sama Dhatu Parampara” [11].

In modern science, cancer is defined as hyper proliferative disorder that involves damage and inflammation of cells in a specific tissue that is responsible to maintain cellular differentiation, survival, proliferation and death [12]. After cardiovascular diseases, cancer is still the second leading cause of death worldwide worse than AIDS, tuberculosis and malaria. According to World Health Organization (WHO), in the year 2012, total cancer deaths were 8.2 million (~22,000 cancer deaths a day), which means that around 15% of total deaths are caused by cancer alone [13]. It is estimated by 2030, globally 21.7 million new cancer cases may exist and 13 million deaths simply due to the growth and aging of the population [14].

In modern medicine, chemotherapy is the first line treatment method for many types of cancer (especially for metastatic disease) [15]. After a huge investment, intensive and systematic research in drug discovery, there are some useful therapeutic agents in the market that include penicillin [16], temozolomide, carmustin, carboplatin and thalidomide [17]. Biologically, once normal cells get mutated and/or become cancer cells and lose most of the normal regulatory functions and continue to multiply cancer cells which are not seen in normal cells [18]. Thus chemotherapeutic drugs do not work properly on cancer as they are mainly targeted to kill cancer cells but not repair the regulatory function. As a result, the degradation takes place continuously. The disappointment in the use of chemotherapeutic treatments, as expected, is also due to many toxic side effects that occur during and after the treatment [19] which will affect the quality of life of the patient due to the very treatment that is intended to provide relief. As a result, the global community is still looking for an alternative and natural medicine that could cure diseases with least side effects and issues.

Towards these goals, traditional medical systems including Ayurveda, Chinese medicine and European medicine have attracted attention as complementary approaches. Ayurvedic medicinal plants have a special place in the treatment of different types of cancer with established outcomes for thousands of years and hence received particular interest. This review is focused on the discovery of bioactive molecules, ongoing improvements, major achievements, current issues and the areas to be improved in the field of anti-cancer drug development from the vast knowledge of Ayurvedic medicine.

Bioactive molecules from Ayurvedic medicine

Plants produce armoury of secondary metabolites that play a vital role in their biochemical pathways and defence against pests. Taking the advantage of bioactive properties, humans started consuming plant products as neutraceuticals. In Ayurveda treatment practices, diet rich in fruits, vegetables, grains and herbal extracts [20] rich in neutraceuticals are consumed to bring health benefits [21,22]. Traditionally, Ayurvedic medicinal plants are consumed in different forms [7]: chewing, swallowing, applying over the skin, teas, consuming with milk, ghee, honey or other solvents commonly known as “Anupana” in Ayurveda [23].

To date, more than 25,000 plant-based formulations and individual herbs are used in folk and traditional medicine in India by more than 387 thousand qualified practitioners of traditional medicinal system [24]. Over 75 formulations are readily available in the Indian market for health and vitality; all contain Withania somnifera (100%), Asparagus racemosus (81.5%), A. adscendens (48%) and Curcullago orchoides (15 %), with other plants in small proportions [25]. The plants Glycrrhiza glabra, Piper longum, Adhatoda vasica, W. somnifera, Ciprus rotundus, Tinospora cordifolia, Berberis aristata, Tribulus terrestris, Holarrhena antidysenterica and Boerhavia diffusa are used in approximattely 140 herbal formulations. Triphala containing potent anticancer agents (Terminalia chebula, T. bellerrica and Embelia officinalis) has been found in more than 219 formulations [26]. Some of the formulations that are widely used in treatment of cancer are listed in (Table 1).
Modern drug discovery process is primarily driven by tracing the traditional medicinal use followed by bioactivity guided fractionation to identify major active ingredients.

| S.No | Name of formulation | Name of herbal plants |
|------|---------------------|-----------------------|
| 1.   | United States Patent 6780441 Sahajanand Biotech Private Limited (Gujarat, IN) | Withania somnifera, Chlorophytum borivilianum, Boerhavia diffusa, Elephantopus scaber, Moringa oleifera, Tecoma undulata, Bauhinia purpurea, Ficus racemosa, Cyperus rotundus, Sphaeranthus acmella and Tinospora cordifolia. |
| 2.   | Immunozone Sino-Vedic Cancer Research Centre | Immunozone herbal formulation contains active principles isolated from selected antimutagenic and immunoenhancing herbs such as Aegle marmelos, Aloe vera, Alpinia galanga, Andrographis paniculata, Azadirachta indica, Berberis vulgaris, Curcuma domestica, Emblica officinalis, Glycine max, Morinda citrifolia, Ocimum sanctum, Tinospora cordifolia, Trigonella foenum-graecum, Viscum album, Withania somnifera and Zingiber officinale. |
| 3.   | Cancertame Sino-Vedic Cancer Research Centre | Cancertame herbal formulation contains active principles isolated from Bauhinia variegata, Catharanthus roseus, Curcuma longa, Glycine max, Glycyrrhiza glabra, Gossypium hirsutum, Nigella sativa, Phyllanthus emblica, Plumbago zeylanica, Rubia cordifolia, Solanum indicum, Zingiber officinale, Aloe vera, Amoora rohiikula and Azadirachta indica. |
| 4.   | Oncotame Sino-Vedic Cancer Research Centre | Oncotame herbal formulation contains active principles isolated from Ginkgo biloba, Oldenlandia diffusa, Podophyllum emodi, Punella vulgaris, Psoralea corylifolia, Saussurea lappa, Solanum nigrum, Withania somnifera, Panax ginseng, Catharanthus alba, Curcuma aromatica, Andrographis paniculata, Aloe barbadensis, Emblica officinalis and Viscum album. |
| 5.   | Chen Pi Ageless Herbs | Citrus reticulate, Astragalus membranaceus, Arctium lappa, Ganoednera lucidum, Trifolium pretense, Eleutherococcus senticosus, Rumex crispus, Atractylodes macrocephala, Spatholobus sub erectus, Ascorbyum nodosum, Punella Vulgaris, Salvia milifiorrhiza, Carthamus tinctorius, Rumex acetosella, Foeniculum vulgare, Althaea officinalis. |
| 6.   | Caractol | Blepharis Edulis, Piper Cubea Linn, Similia Chin Linn, Hemidesmus Indicus, Tribulus Terrestris, Ammania Vesieatoria, Lepidium Sativum Linn, Rheum emodi wall. |
| 7.   | Triphala | Terminalia chebula, Emblica officinalis, Terminalia bellericia. |
| 8.   | Vidaikan choornam | Embelia ribes, Migna oleifera, piper longum |
| 9.   | Liv 52 | Capparis spinosa, Cichorium intybus,Mandur bhasma, Solanum nigrum, Terminalia arjuna, Cassia occidentalis, Achilea milefolium, Tamarix gallica |
| 10.  | Immu-21 | Ocimum sanctum, Withania somnifera, Emblica officinalis and Tinospora cordifolia |

Table 1: List of Herbal formulations available in the market for cancer treatment [73-75]
Curcumin is a hydrophobic polyphenol compound derived from *Curcuma longa*, commonly known as turmeric, consumed in regular culinary [39]. Traditionally, turmeric is commonly used in the treatment of microbial diseases, stomach and liver ailments. Phytochemical evaluation of turmeric revealed two major constituents demethoxycurcumin (DMC) and bisdemethoxycurcumin (BDMC), apart from curcumin [40]. In vitro and in vivo studies have demonstrated that curcumin is a potent anti-tumorigenic agent [41,42] against different cancer types including colon, skin, duodenum and stomach. Curcumin is considered as a noble chemopreventive and antitumor agent with its multiple targeting property without side effects even at high dose [41-43].

On the contrary, poor bioavailability, solubility and absorption hindered its approval as chemotherapeutic agent [39]. Recent studies showed that curcumin possesses in vitro anti-angiogenic and in vivo anti-tumor properties through combined phosphodiesterases-PDE2 and PDE3 inhibition [44]. Furthermore, DMC induced the expression of promoter of methylated genes more than cumin in leukemia cells and on combination could give optimum re-expression of epigenetically silenced genes [45,46]. The ongoing research outcomes provide strong indication that curcumin will be a potential anticancer chemotherapy agent derived from Ayurvedic herb.

Andrographolide is an active compound derived from *Andrographis paniculata Nees* commonly known as 'King of Bitters' (*Kalmeg*) in Ayurveda [47]. *Kalmeg* is distributed throughout South India and Srilanka displayed broad spectrum of activities including anticarcinogenic, anti HIV and several other pharmacological activities [47]. Pharmaceutical constituents including andrographolide, neandrographolide and dehydroandrographolide have been shown to possess anti-cancer properties [48]. Andrographolide being the most active compound from *kalmeg* has shown anticancer properties in several in vitro and in vivo studies against B16F0 melanoma syngeneic and HT-29 xenograft models [47], 2-cell line panel containing MCF-7 (breast cancer cell line) and HCT-116 (colon cancer cell line)[49], multiple myeloma [50,51], rats induced with gastric ulcer [52]. In one of the human study commenced by Darryl and co-authors, ingestion of *A. paniculata* at a dose of 500 mg twice daily along with other nutraceuticals by 20 patients with late stage cancer in different parts showed significant improvement in the patients [53]. However, the poor oral bioavailability has remained a major setback [54].

*Withania somnifera* is commonly known as *"Ashwagandha"*, widely used in Ayurveda for treatment of arthritis, menstrual cycle and uterus related disorders, cancer and other inflammatory diseases [55]. Previous in vivo studies demonstrated antitumor properties of roots of *Aswagandha* against Sarcoma-180 solid tumor with low toxicity [56]. *Withaferin A* is a most abundant bioactive constituent of *W. somnifera* and received a marked interest globally for its exceptional anticancer and anti-inflammatory properties [57].

*Withaferin A* inhibits Notch-1 signalling and down regulates pro survival pathways such as Akt/NF-kB/Bcl-2, in HCT-116, SW-480 and SW-620 cancer cell lines [58]. Also other in vivo studies suggest antiproliferative activity on NCL-H460 (lung), HCT-116 (colon), SF-268 (central nervous system; CNS and MCF-7 (breast) human tumor cell lines [59]. Recent studies conducted by Li Xu and coauthors [60] suggest the synergistic antitumor activity of withaferin A combined with oxaliplatin.
| S.No | Plant species | Active molecules/plant extracts | Biological anti-cancer activities against |
|------|--------------|---------------------------------|-------------------------------------------|
| 1    | Andrographis paniculata Nees (Green chirayta) | Andrographolide, neoandrographolide and dehydroandrographolide | B16F0 melanoma syngenic and HT-29 xenograft models[47], 2-cell line panel containing MCF-7 (breast cancer cell line) and HCT-116 (colon cancer cell line)[49], multiple myeloma [50,51], rats induced with gastric ulcer [52] |
| 2    | Alpinia galanga Wild (Greater galangal) | 1’S-1’-acetoxychavicol acetate, 1,8-cineole, beta-bisabolone and beta- seline, Galangin and Camphor | COR L23 cells (lung cancer cell line) and MCF7 cells (breast cancer cell line) with IC50 7.8µM and 23.9µM[76] |
| 4    | Aristolochia indica (Birthwort) | Aristolochic Acid | Cytotoxic against Chinese hamster ovary-K1 (CHO-K1) cells [77] |
| 5    | Agati grandiflora (Humming bird tree) | Whole plant material ethanol extract | Anticancer effect against Ehrlich Ascites Carcinoma (EAC) in mice cell line [78] |
| 6    | Albizia lebbeck (Rain tree) | Triterpenoid saponins [79] | Antitumor activity against gall tumors [80] |
| 7    | Allium sativum (Garlic) | Abstract Diallyl sulfide [81], S-allylcysteine, S-allymercaptol-cysteine | Cytotoxic against colorectal adenocarcinoma [81] |
| 8    | Aloe vera (Aloe) | Barbabin (aloe-emodin, aloesin, aloin), acemannan | Selectively inhibited the growth of human neuroectodermal tumors in mice [82], caused tumor shrinkage, tumor necrosis and lymphocytic infiltration in dogs [83] |
| 9    | Alstonia scholaris Linn (Milky pine, sapthapama) | Ethanol extract (Echitamine[84]) | Selectively inhibited the forestomach carcinogenesis in mice [85] |
| 10   | Amorphophallus campanulatus (Elephant foot yam) | Methanolic extract | Cytotoxic and apoptotic activities against human hepatoma cell line [86], against human colon carcinoma cell line HCT-15 [86] |
| 11   | Amoora rohituka (Rohituka tree) | Petroleum ether extract | Cytotoxic effects on breast and pancreatic cancer cells [87] |
| 12   | Azadirachta indica (The wonder tree) | Nimbidolde[88] | Antiproliferative effect on human canccl lines [88] |
| 13   | Bacopa monnieri (Indian pennywort) | Bacoside A and B[89] Stigmasterol [90] | Antitumor against Ehrlich Ascites carcinoma in mice [90] |
| 14   | Berberis aristate(Indian ophthalmic barberry) | Methanolic extract | Cytotoxic activity in MCF7 cell line [91] |
| 15   | Calotropis gigantea R. Br(Gigantic swallow wort) | Alcoholic extract, Lupeol | Anti-cervical cancer activity [92] |
| 16   | Calotropis procera R. Br(Crown floer) | Asclepin, Alcoholic extract | Antiproliferative activity [93] |
| 17   | Cassia fistula (Golden shower tree) | Methanolic extract | Antitumor activities against Ehrlich ascites carcinoma [94] |
| 18   | Carum roxburghianum Kurz(Azowan) | Monoterpen hydrocarbon | Cytotoxic in the brine shrimp lethality assays [95] |
| 19   | Curcuma longa L. | Methanolic extract | Cytotoxic activity against human breast adenocarcinoma (MCF7) and human colon carcinoma (HT-29) [96] |
| 20   | Cucumis sativus (Three leaces caper) | Methanol extract, Lupeol[97] | Cytotoxic activity in brine shrimp lethality bioassay [98] |
| 21   | Cucumis sativus var. longa (Tumeric) | Curcumin I-III [43,99] | Potent inhibitor of Leukotriene B4 formation in rat peritoneal polymorphonuclear neutrophils (PMNL)[43,99] |
| 22   | Cucumis sativus (White turmeric) | Curcuminoids [100] Isocurcumenol [101] | Cytotoxic against human ovarian cancer OVCAR-3 cells [100] |
Bhandari M, Ravipti AS, Reddy N, Koyyalamudi SR (2015) Traditional Ayurvedic medicines: Pathway to develop anti-cancer drugs. J Mol Pharm Org Process Res 3: 130. doi:10.4172/2329-9053.1000130

23 Cymbopogon citrates (Lemongrass) Citral [102] Induced apoptosis in caspase-3 enzymatic activity
24 Cinnamomum cassia (Cinnamon bark) Aqueous extract, 2’-Hydroxycinnamaldehyde (HCA)[103] Induced apoptosis in human cervical cancer line (SiHa) [104]
25 Datura metel L/ (Angel’s trumpet) Withanolides [105] Daturametelins H–J [106] Cytotoxic activities again S549 (lung), BGC-823 (gastric) and K562 (leukemia) cancer cell lines [105]
26 Euphorbia nerifolia L (India spurge tree) Hydro-ethanolic extract Chemopreventive effect against DENA-induced renal carcinogenesis in mice [107]
27 Elephantopus scaber (Elephant’s foot) Deoxyelephantopin (ESD) [108] Elescaberin, isodeoxyelephantopin, Deoxyelephantopin [109,110] Antineoplastic effects of a sesquiterpene lactone on lung adenocarcinoma (A549) cells [109,110]
28 Ficus glomerata (Goolar fig) Ethanolic extract Anticancer activities in mammary carcinoma DMBA induced rats [111]
29 Madhuca indica (Butter tree) Ethanolic extract Cytotoxicity against ling (A-549), colon (502713 HT-29) and neuroblastoma (IMR-32) cell line [112]
30 Mallotus philippinensis Muell. Arg (Kamala tree) 3alpha-hydroxy-D:A-friedooleanan-2-one [113], Ethanolic extract with Rottlerin [114,115] Cytotoxic activity on human cancer cell lines [116]
31 Saraca indica (Ashoka tree) 7,12-dimethyl benz(a)anthracene [117] Chemopreventive property in skin cancer in mice models [117]
32 Emblica officinalis, Gaerth, pyrogallol [118] Cyclophosphamide (CP) [120] Chemopreventive property on DMBA induced skin tumorigenesis in Swiss albino mice [119]
33 Urginea indica (Indian squill/ wild onion) 29kDa glycoprotein[121] Antitumor activity against ascites tumor, mouse mammary carcinoma [121]
34 Taxus buccata (Himalayan yew) Taxol[122], taxotere Cytotoxic against breast, lung, ovarian, colorectal cancer and melanoma tumor colony [123]
35 Withania somnifera (Indian ginseng)[124] Withaferin A[55] Viscosalactone B[59] Antitumor activity on S-180 in BALB/c mice [124], antiproliferative activity on NCL-H460 (lung), HCT-116 (colons), SF-268 (central nervous system;CNS and MCF-7(breast) human tumor cell lines [59]
36 Xantium strumarium(Burdock datura)[125] Xanthatin and xanthinosin[126] Cytotoxicity in the human cancer cell lines WiDr ATCC (colon), MDA-MB-231 ATCC (breast) and NCI-417 (lung) [127], Cytotoxic effects on the human cell line, HL-60 [127]

Table 2: List of pharmaceutical drug leads derived from Ayurvedic medicinal plants.

Withaferin A enhanced oxaliplatin induced growth suppression and apoptosis in PC cells via mitochondrial dysfunction and inactivation of the P13K/AKT pathway [60]. Withaferin A is of particular interest due to its ability to interact and inhibit different cancer-specific pathways with least toxic effects.

Market share of Ayurvedic medicine in the pharmaceutical industry

Natural products derived from plants, animals and other sources have been pivotal in treatment of human diseases. Ayurvedic medicine has significant history in this respect and there is an ongoing demand due to their efficacy, low toxicity and fewer side effects. These positive attributes have led to mushrooming of Ayurvedic medical practices and related herbal based drug manufacturing companies worldwide.

The market value for Ayurvedic herbs and their derivatives is growing annually at 20% and the sales have increased by 25% in the last 10 years. To date, China and India are major sources of herbal products. In the global market, China shares 13% of total market whereas India shares only 2.5% [61]. Remarkably, Traditional Chinese medicinal (TCM) system comprises 5000 plant species while Ayurvedic medicinal system uses 7000 plant species [62]. According to Export / Import Bank, TCM plants have a global trade value of US $6 billion, while Ayurvedic medicine have only a share of US $1 billion. The trade of Ayurvedic products are expected to increase rapidly in the near future, due to the fact that a number of drugs with high efficacy are primarily derived by the utilization of realm of traditional Ayurvedic knowledge.

The trade of Ayurvedic medicine is mainly being affected by poor quality control practices, lack of efficient agricultural practices, lack of knowledge on the global marketing strategies and standardization of processes and services. The projection made by WHO states that the global herbal market would grow to $5 trillion by 2050 from the current market level of $62 billion [61].
associations like National Medicinal Plants Board (NMPB), Department of Ayurveda, Yoga and Naturopathy, Unani, Siddha and Homeopathy (AYUSH), World Health organization (WHO) in India, USA, UK, Russia, Germany, Hungary and South Africa are now yielding good results in promoting Ayurveda which has opened-up new opportunities [63].

Implications of using Ayurvedic medicine

Ayurvedic medicine system is a very ancient and is now practiced throughout the world. The Ayurvedic medicine preparations are formulated by a combination of medicinal plants, animal products and minerals and metals including gold, copper, lead, mercury, iron and zinc [64]. These metals are primarily added to herbal products as ashes or bhasmas, with a belief that they act as catalyst and improves the target specificity. However, Ayurvedic practitioners often prescribe medicine which do not meet the criteria of modern GMP or manufactured by practitioners themselves. According to Centre for Disease Control, during the years between 2000 – 2003, 12 cases were reported with lead toxicity after consumption of Ayurvedic products [65]. Strict regulations for Ayurvedic preparations are therefore expected to minimize adverse effects and improve the acceptance of these medicine by global communities.

Many herbal supplements that are available in the market are subjected to limited regulations. Individuals consume these herbal products as dietary supplements without proper knowledge of pros and cons and hence often affected by toxic effects. In a case study of a 69-year-old Caucasian male retired lawyer consumed ‘Bhasma’ an ayurvedic herbal medication on family advice while traveling in India. He reported illness of being lethargy, fatigue, memory impairment, generalized weakness, severe constipation, anorexia and weight loss of 18 kg over preceding eight months. After thorough clinical investigations, discontinuation of Ayurvedic Bhasmas and chelation therapy treatment helped him to recover from these issues [66].

Chemical constituents of plants can be influenced by several factors including habitat, cultivation site, seasonal variations, temperature, soil composition, water availability and pollutant levels in the atmosphere. Furthermore, physicians prescribe different plant species for specific symptoms due to morphological similarities. For instance, Shankhpushpi is herbal drug widely used in the treatment of epilepsy, leucoderma, bronchitis and teething problems. This drug is sourced from three plant species, namely, Clitoria ternatea, Convolvulus pluricaulis and Evolvulus alsinoides in different regions of the country [67]. In fact, the chemical constituents of these plants are significantly distinct from one another. Due to ever increasing demand for Shankhpushpi and other important herbal products, the preparations are often altered with other plant material. Phytochemical investigations suggest that over 60 % herbal products available in the market are adulterated [68].

In order to supply quality herbal products for better health, there is a requirement of standardized authentication of herbs. For instance, rigorous morphological analysis and quantification of molecular markers can be used as an effective means of identification of plants. It is therefore essential that the regulations on authentication of plant material must be strictly implemented in order to minimize adverse effects due to adulteration of herbal medicine and also to gain the desired health benefits.

Interface between Ayurveda and modern medicine

In the realm of modern drug discovery, traditional knowledge of Ayurvedic medicinal plants is playing pivotal role. There has been significant correlation between the use of herbs for particular symptoms and the targets of derived drugs. Anticancer drugs derived from Ayurvedic medicinal plants exert their activity by means of activation or inactivation of signaling pathways [69]. Oncogenes, cancer growth factors, cancer promoting enzymes, protein kinases have been identified as major targets in drug discovery and development process [70]. Thousands of years ago, Ayurvedic medicinal system used many medicinal plants in the treatment of cancer (when there was no modern medicinal knowledge existed).

Drugs derived from many of these traditional plants are now well established inhibitors of the targets responsible for different cancers. For instance, Vinca rosea and Taxus species were widely used in the treatment of different cancer types [71]. Vinristine and paclitaxel isolated from these plants showed strong anticancer properties. [71]. These compounds are now available in the drug market.

Modern cancer treatment typically involves administration of a single therapeutic agents into human system. These agents usually lack specificity to the target or multiple targets, leads to apoptosis of healthy cells and causes adverse side effects. Toxicity and side effects remained to be major setbacks in chemotherapeutic approaches to cancer treatment. For example, anticancer drugs, including vincristine and paclitaxel, are responsible for peripheral neuropathy during cancer treatment [71].

Typical Ayurveda treatment involves administration of an individual or mixture of different plant extracts. Ayurveda recommends formulation (multiple plants at different proportions) to obtain synergism and diminish drug associated side effects such as cachexia (including nausea, wait loss and anorexia). For instance, W. somnifera, Sida cordifolia, Vitis vinifera, Plumbago zeylanica, Tinospora cordifolia Asparagus racemosa and Zingiber officinale are often used to improve appetite, malnutrition, body resistance and fatigue. In addition, Ayurvedic treatment incorporates yoga and relaxation therapies which provide spiritual well-being of the patient. The effectiveness of these holistic approaches goes beyond the symptom relief and improves physiological and behavioral aspects of the patient.

Owing to these benefits, modern cancer treatments are gradually adapting combinatorial approaches including yoga, meditation, mind-positive thinking and cognitive practices along with chemotherapy. The case of Anita Moorjani’s experience with lymphoma evidences role of mind and positive thinking in curing diseases[72]. Anita underwent many conventional chemotherapy cancer treatments. During the cancer treatment she fell into deep coma for several days and claimed to have had near death experience and out-of-body experience [72]. As the lymphoma had metastasized throughout her body and her death was confirmed. Subsequently, Anita came out of coma surviving from cancer, claiming that it was only possible through spontaneous healing developed from positive thinking attitude [33]. Meditation, yoga and prayers have been practiced in Ayurvedic treatment to develop mind-body control [33]. Furthermore, a study conducted by Özlem Ülger and co-authors evidenced improved health conditions in cancer patients after yoga practices [65]. These observations suggest that a revolutionized approach can be devised through integration of Ayurvedic practices with modern day treatments for enhanced efficacy of cancer therapies.
Conclusion

The quest for pharmaceutical drug leads from Ayurvedic medicinal plants has been rapidly growing. The prime reason for this interest is: medicinal plants are cost effective, reliable and possess least side effects. However, ever growing requirement of medicinal plants often leads to cultivation of malpractices. In this respect, educating the communities about correct procedures of collecting, extracting, handling and storing of herbal material, and strict implementation of regulations on quality control would minimize wrong practices and improve the grade of the products and their efficiency. Furthermore, the toxicity and safety assessment procedures for these herbal medicines should be conducted using modern analytical techniques and tools for their improved efficacy. Many international authorities and agencies including the WHO, European Agency for the Evaluation of Medicinal Products and European Scientific Cooperation of Phytomedicine, US Agency for Health Care Policy and Research, European Pharmacopeia Commission and the Department of Indian System of Medicine have initiated developing and implementing new and effective strategies for regulating quality control and standardization of herbal medicine. In addition, appropriate global marketing strategies would boost the usage and economic value of Ayurvedic medicinal plants.

References

1. Lad V (1984) Ayurveda: The science of self-healing: A practical guide. Lotus press.
2. Verhoef MJ, Lewith G, Ritenbaugh C, Roen H, Fleishman S, et al. (2005) Complementary and alternative medicine whole systems research: beyond identification of inadequacies of the RCT. Complement Ther Med 13: 206-212.
3. Ganeshrao BN, Somaji LD, Shirikrishna PA (2014) International Journal of Ayurveda and Pharma Research. Int J Ayur Pharma Research 2: 101-110.
4. Lokhande P, Jagdale S, Chabukswar A (2006) Natural remedies for heart diseases. Indian J Tradit Know 5: 420-427.
5. Rastogi S (2014) Ayurvedic Principles of Food and Nutrition: Translating theory into Evidence-Based Practice. Ayurvedic Science of Food and Medicinal plants: Springer. pp. 325-347.
6. Schmidt LM (2004) Herbal remedies: the other drugs your patients take. Home Healthc Nurse 22: 169-175.
7. Ramawat KG, Goyal S (2008) The Indian herbal drugs scenario in global perspectives. Bioactive Molecules and Medicinal Plants: Springer. pp. 325-347.
8. Balachandran P, Govindarajan R (2005) Cancer—an ayurvedic perspective. Pharmacol Res 51: 19-30.
9. Singh RH (2002) An assessment of the ayurvedic concept of cancer and a new paradigm of anticancer treatment in Ayurveda. J Altern Complement Med 8: 609-614.
10. Rastogi S (2014) Ayurvedic Principles of Food and Nutrition: Translating Theory into Evidence-Based Practice. Ayurvedic Science of Food and Nutrition: Springer. pp. 3-14.
11. Sumantran VN, Tiliu G (2012) Cancer, inflammation, and insights from ayurveda. Evid Based Complement Alternat Med 2012: 306346.
12. Aggarwal BB, Sung B, Gupta SC (2014) Inflammation and Cancer: Springer.
13. Lindsey Torre M, Rebecca Siegel M, Ahmedn Jemal D (2015) American Cancer Society. Global Cancer Facts & Figures. Atlanta: American Cancer Society 3.
14. Torre LA, Bray F, Siegel RL, Ferlay J, Lortet-Tieulent J, et al. (2015) Global cancer statistics, 2012. CA Cancer J Clin 65: 87-108.
15. Van Cutsen E, Kohn C-H, Hité E, Zalúsji J, Chang Chien C-R, et al. (2009) Cetuximab and chemotherapy as initial treatment for metastatic colorectal cancer. N Engl J Med 360: 1408-1417.
16. Chain E, Florey HW, Gardner AD, Heatley NG, Jennings MA, et al. (1940) Penicillin as a chemotherapeutic agent. The lancet 236: 226-228.
17. Guzmán M (2003) Cannabinoids: potential anticancer agents. Nat Rev Cancer 3: 745-755.
18. Reya T, Morrison SJ, Clarke MF, Weissman IL (2001) Stem cells, cancer, and cancer stem cells. Nature 414: 105-111.
19. Partridge AH, Burstin HJ, Winer EP (2000) Side effects of chemotherapy and combined chemohormonal therapy in women with early-stage breast cancer. J Natl Cancer Inst Monogr 30: 135-142.
20. Sanjivani GV, Raina MK, Sharma M (1976) Medicinal plants of India: Indian council of medical research New Delhi.
21. Sinha R, Anderson DE, McDonald SS, Greenwald P (2003) Cancer risk and diet in India. J Postgrad Med 49: 222-228.
22. Park EJ, Pezutto JM (2002) Botanicals in cancer chemoprevention. Cancer Metastasis Rev 21: 231-255.
23. Tavidi K, Galib PB, Patgiri BJ, Prajapati PK Metal toxicity due to Ayurvedic drugs-Facts and Myths.
24. Jain AK, Sharma BK Developments in the field of ayurveda-past to present.
25. Ramawat KG, Jain S, Suri SS, Arora DK (1998) Aphrodisiac plants of Aravalli Hills with special reference to safed musli. Role of biotechnology in medicinal and aromatic plants 1: 210-223.
26. Kamboj VP (2000) Herbal medicine. Current science-Bangalore- 78: 35-38.
27. Dutta S (2015) Natural sources as potential anti-cancer agents: A review.
28. Caltagirone S, Rossi C, Poggi A, Ranellieto FO, Natali PG, et al. (2000) Flavonoids apigenin and quercetin inhibit melanoma growth and metastatic potential. Int J Cancer 87: 595-600.
29. Dragsted LO, Strube M, Larsen JC (1993) Cancer-protective factors in fruits and vegetables: biochemical and biological background. Pharmacol Toxicol 72 Suppl 1: 116-135.
30. Salim AA, Chin Y-W, Kinghorn AD (2008) Drug discovery from plants. Bioactive molecules and Medicinal plants: Springer. pp. 1-24.
31. Wheeler NC, Jech K, Masters S, Brobst SW, Alvarado AB, et al. (1992) Effects of genetic, epigenetic, and environmental factors on taxol content in Taxus brevifolia and related species. J Nat Prod 55: 432-440.
32. Colegate SM, Molyneux RJ (2007) Bioactive natural products: detection, isolation, and structural determination.(2nd edn), CRC press.
33. Ebers GC1 (1994) Treatment of multiple sclerosis. Lancet 343: 275-279.
34. Mohanakumara P1, Sreejayan N, Priti V, Ramesha BT, RaviKanth G, et al. (2010) Dysoxylum binectariferum Hook.f (Melaceae), a rich source of rebaudioside A. J Ethnopharmacol 127: 145-148.
35. Ali S, Heathcote DA, Kroll SHB, Jogalekar AS, Scheiper B, et al. (2009) The development of a selective cyclin-dependent kinase inhibitor that shows antitumor activity. Cancer research 69: 6208-6215.
36. Phelps MA, Lin TS, Johnson AJ, Hurh E, Rozewski DM, et al. (2009) Clinical response and pharmacokinetics from a phase 1 study of an active dosing schedule of flavopiridol in relapsed chronic lymphocytic leukemia. Blood 113: 2637-2645.
37. Shindalipina P, Brown JR, Danilov AV (2014) A new hope: novel therapeutic approaches to treatment of chronic lymphocytic leukaemia with defects in TP53. Br J Haematol 167: 149-161.
38. Asghar U, Witkiewicz AK2, Turner NC3, Knudsen ES2 (2015) The history and future of targeting cyclin-dependent kinases in cancer therapy. Nat Rev Drug Discov 14: 130-146.
39. Anand P, Kunnunakkarra AB, Newman RA, Aggarwal BB (2007) Bioavailability of curcumin: problems and promises. Mol Pharm 4: 807-818.
40. Himesh S, Sharan PS, Mishra K, Govind N, Singhai A (2011) Qualitative and quantitative profile of curcumin from ethanolic extract of Curcuma longa. Int Res J Pharm 2: 180-184.
41. Shankar S, Ganapathy S, Chen Q, Srivastava RK (2008) Curcumin sensitizes TRAIL-resistant xenografts: molecular mechanisms of apoptosis, metastasis and angiogenesis. Mol Cancer 7: 7-16.
42. Shankar S, Chen Q, Sarva K, Siddiqui I, Srivastava RK (2007) Curcumin enhances the apoptosis-inducing potential of TRAIL in prostate cancer cells: molecular mechanisms of apoptosis, migration and angiogenesis. J Mol Signal 2: 10.

43. Shankar S, Srivastava RK (2007) Sax and Bak genes are essential for maximum apoptotic response by curcumin, a polyphenolic compound and cancer chemopreventive agent derived from turmeric, Curcuma longa. Carcinogenesis 28: 1277-1286.

44. Abusnina A, Keravis T, Zhou Q, Justiniano H, Lobstein A, et al. (2015) Tumour growth inhibition and antiangiogenic effects using curcumin correspond to combined PDE2 and PDE4 inhibition. Thromb Haemost 113: 319-328.

45. Yu J, Peng Y, Wu LC, Xie Z, Deng Y, et al. (2013) Curcumin down regulates DNA methyltransferase 1 and plays an anti-angiogenic role in acute myeloid leukemia. PLoS One 8: e59394.

46. Hassan HE, Carlson S, Abdallah I, Buttolph T, Glass KC, et al. (2015) Curcumin and dimethylycurcumin induced epigenetic changes in leukemia cells. Pharm Res 32: 863-875.

47. Rajagopal S, Kumar RA, Devi DS, Satyanarayana C, Razagopalan R (2003) Andrographolide, a potential cancer therapeutic agent isolated from Andrographis paniculata. J Exp Oncol 3: 147-158.

48. Lim JC, Chan TK, Ng DS, Sagineedu SR, Stanslas J, et al. (2012) Andrographolide and its analogues: versatile bioactive molecules for combating inflammation and cancer. Clin Exp Pharmacol Physiol 39: 300-310.

49. Iada SR, Matthews C, Saad MS, Hamzah AS, Lajis NH, et al. (2008) Benzylidine derivatives of andrographolide inhibit growth of breast and colon cancer cells in vitro by inducing G arrest and apoptosis. Br J Pharmacol 155: 641-654.

50. Matsuda T, Kurayogami M, Sugiyama S, Umehara K, Ueno A, et al. (1994) Cell differentiation-inducing diterpenes from Andrographis paniculata Nees. Chem Pharm Bull (Tokyo) 42: 1216-1225.

51. Gunn EL, Williams JT, Huynh DT, Iannotti MJ, Han C, et al. (2011) The natural products withanolide and andrographolide exhibit anti-cancer stem cell activity in multiple myeloma. Leuk Lymphoma 52: 1085-1097.

52. Saranya P, Geetha A, Selvamathy SM (2011) A biochemical study on the gastroprotective effect of andrographolide in rats induced with gastric ulcer. Indian J Pharm Sci 73: 550-557.

53. See D, Mason S, Roshan R (2002) Increased tumor necrosis factor alpha (TNF-alpha) and natural killer cell (NK) function using an integrative approach in late stage cancers. Immunol Invest 31: 137-153.

54. Je L, Wang T, Tang L, Liu W, Yang Z, et al. (2011) Poor oral bioavailability of a promising anticancer agent andrographolide is due to extensive metabolism and efflux by P-glycoprotein. J Pharm Sci 100: 5007-5017.

55. Devi PU (1996) Withania somnifera Dunal (Ashwagandha): potential plant source of a promising drug for cancer chemotherapy and radiosensitization. Indian J Exp Biol 34: 927-932.

56. Devi PU, Akagi K, Ostapenko V, Tanaka Y, Sugahara T (1996) Withaferin A: a new radiosensitizer from the Indian medicinal plant Withania somnifera. Int J Radiat Biol 69: 193-197.

57. Devi PU, Akagi K, Ostapenko V, Tanaka Y, Sugahara T (1996) Withaferin A: a new radiosensitizer from the Indian medicinal plant Withania somnifera. Int J Radiat Biol 69: 193-197.

58. Koduru S, Kumar R, Srinivasan S, Evers MB, Damodaran C (2010) Notch-1 inhibition by Withaferin-A: a therapeutic target against colon carcinogenesis. Mol Cancer Ther 9: 202-210.

59. Jayaprakash B1, Zhang Y, Seeram NP, Nair MG (2003) Growth inhibition of human tumor cell lines by withanolides from Withania somnifera leaves. Life Sci 74: 125-132.

60. Li X, Zhu F, Jiang J, Sun C, Wang X, et al. (2015) Synergistic antitumor activity of withaferin A combined with oxaliplatin triggers reactive oxygen species-mediated inactivation of the PI3K/AKT pathway in human pancreatic cancer cells. Cancer letters 357: 219-230.

61. Bhattacharya R, Reddy KRC, Mishra AK (2014) Export strategy of Ayurvedic Products from India. Intern J Ayurvedic Med 5.
published in this special issue (part 2 of 2) on the Safety and Efficacy of Natural Health Products. Can J Physiol Pharmacol 85: 1160-1172.

127. Nibret E, Youns M, Krauth-Siegel RL, Wink M (2011) Biological activities of xanthatin from Xanthium strumarium leaves. Phytother Res 25: 1883-1890.