

Caspase Activators: Phytochemicals with Apoptotic Properties Targeting Cancer, a Health Care Strategy to Combat this Disease

Asma Saqib¹, Sharath Pattar², Chandrakant Shivappa Karigar³, Shailasree Sekhar^{4,*}

¹Department of Biochemistry, Maharani's Science College for Women, Palace Road, Bangalore, Karnataka, INDIA.

²Department of Bioinformatics, Maharani's Science College for Women, Palace Road, Bangalore, Karnataka, INDIA.

³Department of Biochemistry, Bangalore University, Bengaluru, Karnataka, INDIA.

⁴Scientist, Institution of Excellence, Vijnana Bhavana, University of Mysore, Mysuru, Karnataka, INDIA.

ABSTRACT

Context: Caspases, a family of cysteine-aspartic proteases have a pivotal role in apoptotic pathways. Their down-regulation is reported to induce inappropriate cell survival and enhanced carcinogenic potential. Screening of phytochemicals with a capacity to activate caspases enhancing apoptotic capacity has been proven to be effective anticancer agents. **Objectives:** This review consolidates data on phytochemicals traditionally used to treat cancerous conditions. The scientific validation of caspase-activated apoptosis for this traditional application has been compiled. **Methods:** Internet assisted scientific literature was collected from Google, Google Scholar, Research Gate and NCI, restricted to publications from 1997 to 2019. Search terms 'caspases and cancer', 'assay of caspases', 'traditionally used medicinal plants', 'Kani tribes', 'plant extracts activating caspase', 'cytotoxicity assay', 'docking phytochemicals to caspase', 'technological advancement for anticancer therapy', 'clinical studies of plant extracts and phytochemicals' and 'herbal drugs approved by FDA' was included. **Results:** The compilation revealed significance of multiple experimental strategies, traditional research laboratory practices and advanced *in silico* molecular docking techniques in anticancer therapy. Technological advancement such as MALDI-TOF assisted phytochemical mediated protein target identification and designing promoter for caspases activation and synthesizing functionalized nano carriers for clinical studies has been included for identification of hit molecule and lead optimization. Eugenol and berberine were identified as phytochemicals with potential drug characteristics by both *in silico* and *in vivo* studies. **Conclusion:** The phytochemicals from important Kani tribal medicinal plants via *in silico* docking and *in vivo* studies identified could be explored at clinical trials.

Key words: Anticancer, Apoptosis-associated caspase assay, Advanced *in silico* techniques, NCI, Kani tribes, HeLa cell lines.

Submission Date: 05-08-2019;

Revision Date: 18-09-2019;

Accepted Date: 14-04-2020

DOI: 10.5530/ijper.54.3.98

Correspondence:

Dr. Shailasree Sekhar

Scientist, Institution of Excellence, University of Mysore, Mysuru-570006, Karnataka, INDIA.

Phone: +91 9480476031

E-mail: shailasree@ioe.

uni-mysore.ac.in

INTRODUCTION

Caspases, cysteine-dependent aspartic-acid-directed endoproteases, participate in ordered processes such as inflammation and cell death, essential for the initiation and execution of apoptosis. Synthesized as catalytically inactive pro-caspases, they are activated upon apoptotic stimulus, such as death ligands or permeabilization of outer mitochondrial membrane. In 1992, the first member of this family was described in *Caenorhabditis elegans* var. Bergerac (Rhabditidae), as essential for developmental

cell death and named as CED-3 homologue interleukin-1beta-converting enzyme.¹ Genetic analysis identified two genes (*ced-3*, *ced-4*) for cell death execution (Figure 1). Cloning of *ced-3* identified it to encode a protease with homology to mammalian interleukin-1beta-converting enzyme. The cleavage site for this enzyme in interleukin 1beta was after aspartic acid residue 116. It provided the first indication of cysteine proteases as crucial components of the cell death



www.ijper.org

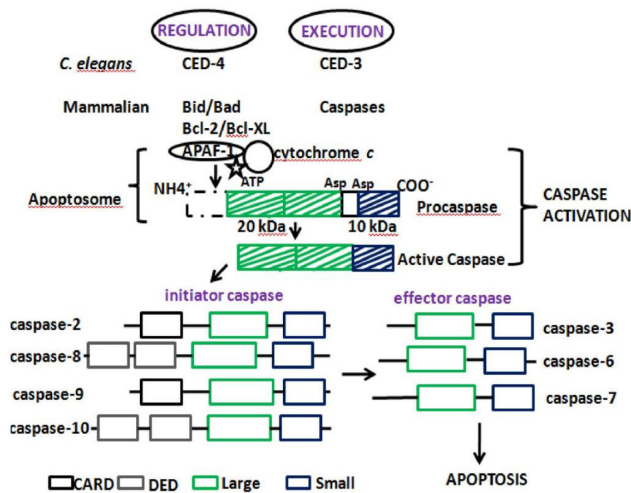


Figure 1: Caspase cascade in apoptosis.

CED-3 and CED-4 function to kill cells. APAF-1 protein contains (from the N terminal) a caspase recruitment domain. Upon binding cytochrome c and dATP, this protein forms an oligomeric apoptosome upon binding to procaspase. The apoptosome binds and cleaves procaspase protein, releasing its mature, activate caspase. Alternative splicing results in several transcript variants encoding different caspase isoforms; initiator caspase-2, -8, -9, -10 and effector caspase-3, -6, -7.

CARD - The caspase recruitment domain; DED - Death-effector domain;

machinery.² Use of caspase inhibitors resulted in cancer proliferation pointing to an essential role for caspases in execution of apoptosis. Subsequently, a growing family of cysteine proteases with homology to CED-3, designated as caspases (for cysteinyl aspartate-specific proteinase was reported. This family of enzymes comprises 11 members in humans and 13 in mammals.³ They are classified based on reported apoptotic inducing capacity (caspase-3, -6, -7, -8 and -9 in mammals). Caspases with a role in human inflammation are caspases-1, -4, -5, -12. Those with a inflammatory role in mice are caspase-1, -11 and -12. The functional role of caspase-2, -10 and -14 are less recognized. Apoptotic caspases are sub-classified by the site and mechanism of action as initiator caspases (caspase-2, -8, -9, -10) activated by dimerization on large multiprotein complexes or apoptotic executioner (caspase-3, -6, -7). They have various interaction with cellular proteins interacting with their prodomains leading to complex formation with adaptors. Caspase Recruitment Domain (CARD) is recognized in caspase-1, -2, -4, -5, -9. Death effector domain is in caspase-8 and -10.³

Western Ghats, a major biodiversity hotspot along the Western coast covers an area of 159,000 sq km with >15,000 plant species. Several of them are endemic with potential medicinal value. Tribals dwelling in these remote areas of the forest cover of Western Ghats have used its flora since antiquity as first aid remedies. With increased prominence to explore traditional knowledge and ethnobotanical studies, validating

traditional use by presenting scientific data has offered a route to develop novel drugs.

The Agasthyamalai hills, the mountain range in Kerala state of the Western Ghats, houses the indigenous Kani tribe. They are traditionally nomadic population of almost 25,000 individuals. This population meets their healthcare needs by preparations based on traditional knowledge using non-timber minor forest produces. A study of Xavier *et al.*⁴ identified and reported use of forest produces by these tribal documented through interviews of the Kani traditional healers. The non-timber produce could be used in fresh or dried state. Some of the preparations included a single herb or use of a combination of more than one in some of their preparations for a single or multiple ailments. A high degree of consensus among the tribal healers for use of a particular plant indicated presence of a consortium of phytochemicals that could be attributed to its medicinal property.

An increased global interest in identifying pharmacological potent compounds as preventive medicine without side effects has warranted attention. There are recommendations to increase consumption of foods rich in bioactive components for overall maintenance of health. Mankind, through trial and error, has found medicinal properties in seeds, barks, roots and leaves of certain plants and traditional knowledge has given clues to the discovery of these valuable drugs. High-throughput ADMET screening provides an effective paradigm for filtering compounds for drug discovery process. The technique employed is based on the prediction of binding modes and binding affinities of each compound in the dataset by means of docking to an X-ray crystallographic structure. Various studies reported in literature stated the importance of dataset size such as 10,000 compounds using Flex and X, 19 44,000 compounds using Surflex20 and several others. Therefore, an alternative approach to eliminate unpromising compounds before docking by restricting the dataset to drug-like compounds is by filtering the dataset based on appropriate property and sub-structural features and by performing diversity analysis. These approaches can be highly effective in reducing the dataset to be docked.

MATERIALS AND METHODS

The present review was intended to consolidate extensive collection of scientific literature on plants traditionally used to treat cancerous conditions. The scientific validation of their anticancer activity via caspase-activated apoptosis for traditional application has been compiled.

Internet assisted literature was collected from Google, Google Scholar, Research Gate and NCI, restricted to publications from 1997 to 2017. Search terms 'caspases and cancer', 'assay of caspases', 'traditionally used medicinal plants', 'Kani tribes', 'plant extracts activating caspase', 'cytotoxicity assay', 'docking phytochemicals to caspase', 'clinical studies of plant extracts and phytochemicals', 'herbal drugs approved by FDA' and 'technological advancement for anticancer therapy' was included.

Molecular Docking of Phytochemicals from Important Kani Tribe Medicinal Plants Swiss Autodock

2Y1L

Caspase-8 of from *Homo sapiens* complex with DARP (dopamine releasing protein) in -8.4 was used for the study. 2Y1L has all the unique 4 protein chain and all the strict requirement for Asp at position P1 and has a preferred cleavage sequence of (Leu/Asp/Val)-Glu-Thr-Asp-1-(Gly/Ser/Ala).

Ligand preparation for docking

A set 82 phytochemicals were selected from literature survey reported from traditional medicinal plants of Kani tribes⁴ recommended in the review by Xavier *et al.*⁴ as a possible source of potential new drugs. They were screened for caspase-8 activation via *in silico* technology. A total of 16 of them exhibited drug likeliness and selected for docking. Drawing window of chemsketch was applied to obtain 3D-coordinates. Gross biological capacity of Lipinski rule 5 was evaluated. Further drug likeliness and drug score was estimated.

Molecular docking

In silico docking was performed using the Swiss Doc of the Lead IT software. Pubchem was used to download ligands. Addition of H-atoms as required was carried out. The molecular models were built. A strategy of steepest descent approximation by running 1000 cycles of energy was applied in this process. The Chimera tool UCSF 1.6.2 was used at a gradient of 0.02. The AMBERff99SB Force field procedure was also applied. The charges of Gastiger were added to the ligands. The Mol2 format was applied to save them. The resulting product was uploaded into the docking toll of Swiss Doc from the Lead IT software.

Target protein minimization

The 2Y1L protein was loaded into the prepared molecule module Bio solve IT software. For experimental initiation Lead IT chain A of the protein was prepared and selected for docking procedure. The binding pocket

of caspase-8 comprised site of interest. The binding energy of this site was minimized. The binding pocket amino acid atomic coordinates were converged. The 2Y1L protein was now ready for docking.

Active Site prediction

The binding pockets of 2Y1L were identified for the binding site analysis. The largest binding pockets were selected for docking studies. A search was carried out for possible binding residues of receptor. The results were applied in our study.

Swiss ADME was applied for drug likeliness evaluation

Swiss ADME analysis for the ligands was assessed.

In vivo evaluation of in silico observations

HeLa cell lines were procured from National Centre for Cell Science (NCCS), Cell Repository, Pune. The procured cell lines were approved by NCCS ethics committee. HeLa cells were maintained in Eagle's Minimum Essential Medium (EMEM). They were supplemented with 10% fetal bovine serum (FBS). The cells were incubated in 5% CO₂ at 37°C.

Confocal Laser Scanning microscopy

For fluorescence microscopy HeLa cells were incubated in 6-well plates at an initial cell density of 1×10^5 cells/mL were grown on 12 mm glass coverslips. When cell density reached 80–90% confluence, cells were treated with eugenol (500 microg/mL) and berberine (500 microg/mL) procured from Sigma Aldrich-Merck Pvt., Ltd, Bengaluru, India for the indicated times (24 and 48h). At the time intervals (24 and 48hr) cells were washed once with PBS (145 mM NaCl, 7.5 mM Na₂HPO₄, 2.8 mM NaH₂PO₄). Additional medium was then removed with filter paper. The CaspGLOW™ Fluorescein active Caspase-8 staining kit was procured for detection of active Caspase-8 in living cells and the manufacturers' protocol was followed. The FITC (fluorescent marker)-conjugated caspase-8 inhibitor, IETD-FMK permeabilized the cells. It bound to cytosolic active caspase-8 irreversibly. FITC on the IETD-FMK was detected by Confocal Laser Scanning Microscope, LSM710 (Carl Zeiss, Germany).

RESULTS

The search identified an essential role for caspase-activated apoptosis and its up-regulation leading to cancer suppression property. The compilation revealed significance of multiple experimental strategies of

both traditional research laboratories practice and advanced *in silico* molecular docking techniques in anticancer therapy. Technological advancement applied in screening of a set of phytochemicals via MALDI-TOF assisted phytochemical mediated protein target identification with designing promoter for caspases activation and synthesizing functionalized nano carriers for clinical studies has been included. Construction and maintenance of natural product library, albeit, compiled information of microbial, plant, marine or sources composed of crude extracts, semi pure mixtures or single purified natural product has been reported to carry a distinctive advantage.^{5,6} Additionally they could provide links to available literature on phytochemicals reported from various research laboratories with potentially more chemical interactive sites for computational studies. This could facilitate their introduction into clinical trials for hit identification and lead optimization leading to developing anticancer drugs with lower effective dosage.⁷

Caspases and Cancer

A note on evidence for down-regulation of caspases resulting in cancerous conditions has been detailed (Table 1). Caspase-8, an initiator caspase enzyme, has a role in extrinsic death receptor pathway, with obligatory role in apoptosis have a putative role in cancer suppression.⁸ Silencing of *CASP8* gene by deletion or methylation of its promoter was reported in tumors and cell lines pediatric cancer. An amplified *MYCN*

oncogene transcript with elevated levels of its protein was reported.^{9,10} A *CASP8* gene mutation due to stop codon modification and Alu repeat to its site was observed in oral cavity cytolytic T lymphocytes of human squamous cell carcinomas. A relatively low frequency of *CASP8* mutations have been reported in colorectal and gastric cancers. In patients suffering from hepatocellular carcinoma, a somatic frame-shift mutation (1225-1226delTG) was reported in nine of the 69 patients.^{11,12} Thus, caspase-8 enzyme alterations resulted in decreased capacity to initiate apoptosis in these cancerous conditions.

Caspase activity assays include a variety of techniques with specific reagents by ELISA using fluorescent or luminescent groups for measurement of their activity. They provide the vital information on its cascade in apoptotic pathway. As each of these techniques has their own advantages and disadvantages, it has been recommended to use at least two complementary assay methods during such studies (Table 2).¹³

Plants and Phytochemicals against Cancer

The World Health Organization (WHO) estimates cancer deaths (to the tune of 30%) to dietary habits and lifestyles (physical inactivity, smoking and alcohol consumption). In the year 2012 alone, deaths related to cancer were estimated to be 8.2 million. The condition of lung cancer at 19% was leading the statistics. Harmful compounds such as polycyclic amines, polycyclic aromatic hydrocarbons, *N*-nitroso compounds from nitrite and reactive oxygen species from heme iron induce increased cell proliferation and DNA damage.¹⁴ The proof is the decades of observation and epidemiological studies suggesting dietary interventions in preventing and decreasing the onset or even reversing chronic disease conditions.¹⁵ Plant diet finds application due their ability to re-sensitize treatment of resistant cancer cells.¹⁶ Dietary bioactive compounds inducing apoptosis via extrinsic / death receptor pathway of apoptosis reviewed¹⁷ are listed (Table 3, Figure 2a, 2b). Questioning the safety of anticancer drugs like synthetic cisplatin and cyclophosphamide with several undesirable side effects, the search for anticancer agents from plant sources, modern science has turned to it for exploitation of the vast phytochemicals to therapeutic targets.¹⁸⁻²¹ In many instances, the cancerous condition was reported as 'hard swellings', 'abscesses', 'calluses', 'corns', 'warts', 'polyps', or tumors.²² The United States initiated extensive plant collection program through National Cancer Institute (NCI) as a part of the National Institute of Health (NIH) in 1957. A total of 1,14,000 extracts from around 35,000 plant samples for anticancer activity

Table 1: Expression of caspases in cancerous condition.

Caspases	Cancer	Reference
↓ <i>CASP8</i>	Neuroblastomas (children)	Teitz <i>et al.</i> ⁹
↓Caspase-8	Advanced gastric invasive colorectal Head and neck Hepatocellular Lung and breast cancer cell lines	Soung <i>et al.</i> ¹² Kim <i>et al.</i> ¹¹ Soung <i>et al.</i> ¹² Soung <i>et al.</i> ¹²
↓Caspase-10	Lung and breast cancer cell lines	Kischkel <i>et al.</i> ⁷³
↓Caspase-2	Gastric	Yoo <i>et al.</i> ⁷⁴
↓Caspase-9	Colorectal	Palmerini <i>et al.</i> ⁷⁵
↓Caspase -6	Gastric	Yoo <i>et al.</i> ⁷⁴
↓ <i>CASP7</i>	Colorectal	Soung <i>et al.</i> ¹²
↓Caspase-7	Head and neck Colorectal Gastric	Soung <i>et al.</i> ¹² Palmerini <i>et al.</i> ⁷⁵ Yoo <i>et al.</i> ⁷⁴
↓Caspase-3	Breast	Devarajan <i>et al.</i> ⁷⁶
↑Caspase-3		Nakopoulou <i>et al.</i> ⁷⁷

↑ = activated/ improved activity/ upregulated

↓ = inhibited/ downregulated

Italicized = gene; unitalicized = enzyme;

Table 2: Assays to detect active caspase induced by apoptotic stimuli.

<p>Fluorogenic and chromogenic assay ELISA</p> <p>Fluorescence microscopy or flow cytometry</p>	<ul style="list-style-type: none"> • Estimation of cleaved substrate products and estimating fluorogenic or chromogenic leaving groups • Substrates used: aminofluorocoumarins <i>N</i>-acetylaspartyl-glutamylvalinylaspartyl-7-amino-4-trifluoromethyl coumarin (DEVD-AFC) for caspase-3 and -7; <i>N</i>-acetylvalinyl-glutamylisoleucylaspartyl-AFC (VEID-AFC) for caspase-6; • Advantages: <ul style="list-style-type: none"> • Obtaining temporal and spatial information about caspase activation • Allows estimation of multiple activities in replicates • Disadvantages: <ul style="list-style-type: none"> • Non-specificity as DEVD-AFC, preferred substrate for caspase-3 is cleaved by caspases-1, -2, -4, -6, -7, -8, -10 and -14 • Apo-ONE™ (Promega, Madison, WI) homogenous caspase-3/7 kit –estimation based on release of rhodamine 110 from peptide bound substrate • PhiPhiLux™ (OncoImmunin, Gaithersburg, MD), peptide with caspase cleavage site tagged with fluorescent moiety, rhodamine 110 on one end and a quencher on the other, generating a fluorescent signal upon cleavage of the peptide that separates the fluorophore and quencher • Disadvantage: <ul style="list-style-type: none"> • The requirement to load the peptide into cells prior applying an apoptotic stimulus
<p>Immunoblotting</p>	<ul style="list-style-type: none"> • Using sera for caspase cleavage products, the anti-neoepitope antibodies • Procaspase 3 and 6 rabbit polyclonal antibodies from Cell Signaling Technology (Beverly, MA); monoclonal antibodies to poly(ADP)-ribose polymerase (PARP) and cleaved caspase-3 antibodies from Sigma (St. Louis, MO) • Advantages <ul style="list-style-type: none"> • The high quality antibodies recognize caspase cleavage products making it possible to detect procaspase cleavage during apoptosis • It provides important evidence for caspase activation during apoptosis • Disadvantages <ul style="list-style-type: none"> • Some of the initiator caspases gain enzymatic activity even without cleavage. Thus it is possible for caspases-8 and/or 9 detecting in immunoblots. • The level of caspase cleavage might not reflect the quantity of activated caspase as it is dependent on the stability of cleavage products • Assessing the appearance of a band at a molecular weight or disappearance of a band at a molecular weight is difficult is the immunological reagent recognizes 10 bands on a blot. • The cleavage of a particular substrate does not necessarily indicate the activated caspase. The cleavage of a particular substrate, PARP, a classically caspase-3 substrate, is cleaved by other caspases, caspase-3, -7, -8, -9 and -10 <i>in vitro</i> and by caspases 7and or/-9 in cells deficient of caspase-3.
<p>Immunofluorescence or flow cytometry</p>	<ul style="list-style-type: none"> • Using antibodies that recognize the active conformation of caspase-3, IETD (Cell Signaling Technologies), the sequence that becomes the C-terminal of the caspase-3 large subunit when procaspase 3 is cleaved by initiator caspase. • Using anti-neo-epitope antibodies that recognizes C- or N-terminal amino acids • Advantages <ul style="list-style-type: none"> • Quantitative data regarding the percentage of cells stained could be obtained • Specificity of the reagents used aids in assessing with certainty activation of a particular caspase • Disadvantages <ul style="list-style-type: none"> • There is a lack of availability of conformation-sensitive antibodies to most of the active caspases • A potential cross-reactivity of anti-caspase neoepitope antibodies with other polypeptides cleaved during apoptosis containing similar sequences
<p>Affinity labeling for immunoblotting, fluorescence microscopy and flow cytometry</p>	<ul style="list-style-type: none"> • Using covalent modification of the caspase active site with a substrate-like molecule with an inhibitory group and a reporter moiety • Acyloxymethylketones (aomks) with reporter moiety (biotin labeled streptavidin; fluorescein; 2,4-dinitrophenol) attached to the amino acid immediately upstream of the scissile aspartate (the P2 amino acid) exposed to solvent and with the capacity to accommodate a bulky substituent upon inhibition binding the caspase or at the N-terminus of the peptide. The biotin-X-VAD(OMe)-fmk and zEK(bio)D-aomk from Calbiochem and the Osaka Peptide Institute (Osaka, Japan) respectively are used. • Advantages <ul style="list-style-type: none"> • Uniquely capable of identification of the caspases activated in the cells at the onset of apoptosis • Can be used for studying caspases that lack anti-caspase antibodies • Disadvantages <ul style="list-style-type: none"> • The reagents are expensive and require custom synthesis • Sometime endogenously biotinylated polypeptides could be detected or co-purified • Not all caspases react with affinity ligands.zEK(bio)D-amok readily labels caspases 2, 3, 6, 7and -9 but not activated caspase-8
<p>RT-PCR</p>	<p>Accession numberSequence</p> <ul style="list-style-type: none"> • Caspase 3U49930F5'-AATCAAGGGACGGGTCATG-3' • R5'-GCTTGTGCGGTACAGTTTC -3' • Caspase 8AF288372F 5'-CTGGAAGGATCGACGATTA-3'R 5'-CATGTCCTGCATTTTGATGG-3' • Caspase 9NM031632F 5'-AGCCAGATGCTGTCCCATA-3'R 5'-CAGGAGACAAAACCTGGGA-3'

Table 3: Selective dietary bioactive compounds inducing extrinsic and intrinsic pathways of apoptosis via activation of caspases.

Bioactive compound	Cell line (s); cytotoxic concentrations(IC ₅₀ values)	Therapeutic targets / Mode of action	References
Curcumin	PC-3; IC ₅₀ 20 micromol/L HepG2, [40 micromol/L]	TRAIL-induced apoptosis; ↓Δψmt	Deeb et al. ⁷⁸
Sulforaphane	MDA-MB-231, [25 micromol/L]	↑Caspase-8, -3; ↓Bcl-2	Pledgie-Tracy et al. ²⁷
EGC or EGCG	SH-SY5Y, [50 micromol/L]	↑Caspase-8,	Das et al. ⁶⁷
beta-carotene	HT-29 [55 micromol/L]	↓Δψmt; ↑Caspase-9	Khan et al. ⁷⁹
Apigenin	22Rv1, [80 micromol/L]	↑Caspase-9, -3; ↓Bcl-2	Khan et al. ⁷⁹
Luteolin	SCC-4, [100 micromol/L]	↑Caspase-3, -9; ↓PARP	Khan et al. ⁷⁹
Resveratrol	MD-MB-231, [50 micromol/L]	↑Caspase-3; ↓PARP	Khan et al. ⁷⁹
Iberin	SK-N-AS, [50 micromol/L]	↑Caspase-3, -9; ↓PARP	Khan et al. ⁷⁹
p-coumaric acid	N2a, [150 micromol/L]	↑Caspase-8	Shailasree et al. ⁸⁰
α-mangostin	COLO, [205.974±0.85 micromol/L] MIP-101, [11.35 ±1.12 micromol/L] SW 620, [19.6 ± 1.53 micromol/L]	↑Caspase-3, -8, -9	Gavrilas et al. ¹⁸
Omega-3 (polyunsaturated fatty acids) PUFA	LT97, [100 microM]	↑Caspase-3, -9	Khan et al. ⁷⁹

Δψmt – mitochondrial membrane potential; PARP: poly ADP ribose polymerase; TRAIL:TNF-related apoptosis inducing ligand

↑= activated/ improved activity/ unregulated

↓= inhibited/ downregulated

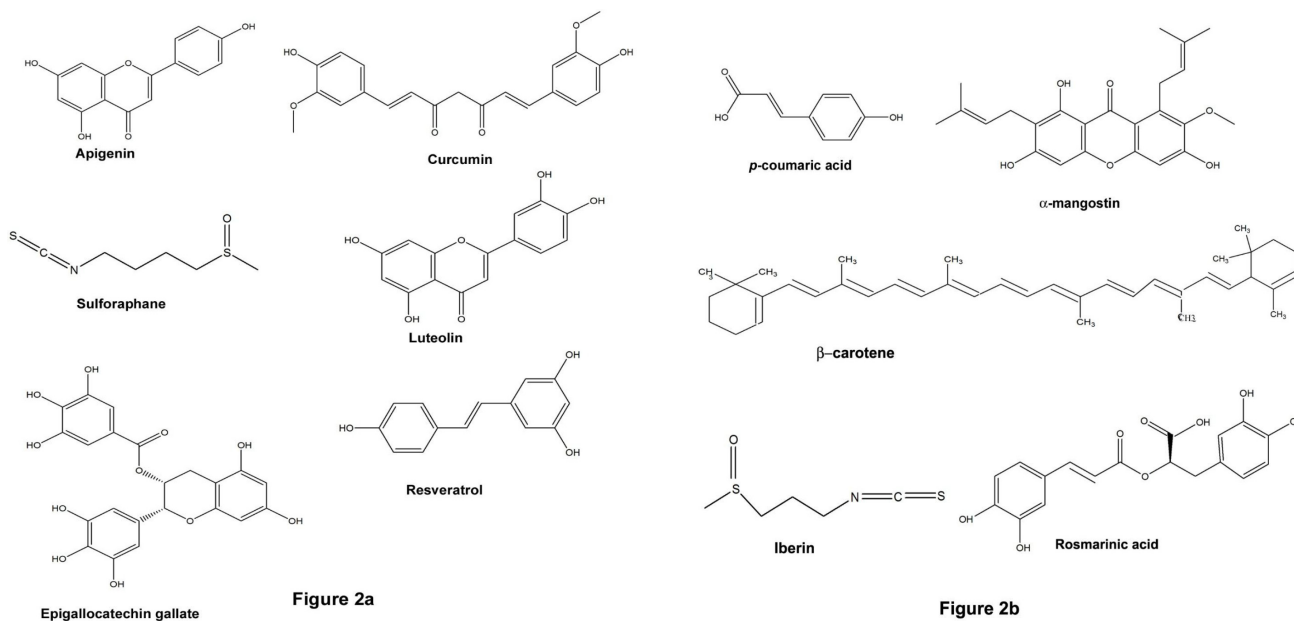


Figure 2a, 2b: Dietary phytochemicals inducing caspase-activated apoptosis.

were screened. By 1991, 28,800 plant samples from over 20 countries were catalogued for chemotherapeutic capacity initially screened *in vitro* against human cancer cell lines.²² The survey carried out showed that above 60% of clinically viable anticancer drug reported were of natural origin or were modeled on natural product parents. The seven plant derived anticancer drugs with US Food and Drug Administration (FDA) approval for commercial production were taxol from *Taxus brevifolia*, vinblastine and vincristine from *Catharanthus*

roseus, etoposide and teniposide from *Podophyllum peltatum*, topotecan from *Camptotheca acuminata* and irinotecan from *Camptotheca acuminata* (Table 4). The most successful discoveries are the vinca alkaloids (vinblastine and vincristine) from *Catharanthus roseus*.²³

Clinical Trials

Paclitaxel (taxol/ taxane) from *Taxus brevifolia* L. with reported ability to block mitosis, disrupt microtubule and spindle formation and inhibit translational

Table 4: Plant extracts and compounds (selective) that were screened against NCI-60 Cancer cell panel, information referring NCI website.

Plant extracts/compounds	Reported properties in NCI Website
Mistletoe extracts (PDQ®)	Cancerous conditions
Eribulin mesylate (Halaven)	Breast and lung cancer
Crizotinib (Xalkori®)	Non-small cell lung cancers
Taxol	Breast and ovarian cancers
Ibrutinib	Effective against Graft-Versus-Host Disease
Gardasil 9	HPV Types (HPV types 6, 11, 16, 18, 31, 33, 45, 52 and 58)*
Velcade®	Multiple Myeloma, blood cancers and solid tumors
Sonidegib	Advanced Basal Cell Carcinoma
Vinblastine and vincristine	Leukemia, bladder and testicular cancers
Etoposide and teniposide	Small-cell lung cancer, testicular cancer, lymphomas
Topotecan	Ovarian and small-cell lung cancers
Irinotecan	Metastatic colorectal cancer

*HPV types 6 and 11 are low-risk types that do not cause cancer but can cause warts on or around the genitals, anus, mouth, or throat. 16 and 18 are responsible for approximately 70 percent of all cervical cancers and HPV types 31, 33, 45, 52 and 58 are responsible for another 20 percent of cervical cancers.

machinery is in Phase I-III clinical use. It is tested against breast-, non-small lung cancer, ovarian- cancer and Kaposi sarcoma. Nano-particles, nanochealtes and nano-liposomes are being developed as alternate form of administration of this drug.^{14,24,25} Sulphoraphane in cruciferous vegetables inhibits growth in breast cancer has been taken up for clinical trials. Administration of cruciferous vegetable preparations orally with sulphoraphane is underway.²⁶⁻²⁸ Podophyllotoxin isomer, epidophyllotoxin from *Podophyllum peltatum* L. with pro-apoptotic capacity interfering with cell cycle has been tested for clinical trials against lymphomas and testicular cancer.^{29,30} Vinca alkaloids from *Catbaranthus roseus* G. Don with apoptotic capacity, bind beta-tubulin and destabilize microtubule, induce cell cycle arrest have been incorporated in clinical trials. Vincristine for lymphomas, sarcomas and leukaemias; vinblastine for testicular cancer, Hodgkins disease and lymphomas;³⁰⁻³³ vinorelbine for Phase I-III against non-small cell lung cancer;^{31,33} vindesine against acute lymphocytic leukaemia;³³ vinflunine in Phase III clinical trials against solid tumors and Vinflunine in Phase III clinical trials against solid tumors.²⁹ Isoflavonoid, pomiferin isolated from *Maclura pomifera* and *Dereeis malaccensis* exhibit antioxidant activity. It inhibits histone-deacetylase. It is reported to be cytotoxic to MDA-MB-231-breast, NCI-H23- lung, ACHN - kidney, PC-3 - prostate, HCT-15 -colon human cancer cell lines, LOX-IMVI – Melanoma and MDA-MB-231-breast cancer cells.^{32,33} Epigallocatechin-3-gallate derived from catechin rich in green tea is in Phase I trial for oral administration related to prostate cancer.^{33,34} Combretastatin A-4 phosphate is a water soluble analogue of combretastatin from *Combretum caffrum*

exhibited anti-angiogenic capacity and has been in early trails against cancer development.¹⁴ Roscovitine derived from olomucine isolated from *Raphanussativus* L. has been tested for in Phase II clinical trials in Europe due to its ability to inhibit cylin dependent kinases and inhibit cell cycle progress.³⁵ Flavopiridol based on rohitukine (isolated from *Dysoxylum binectariferum* Hook. f.) structure, a synthetic flavonoid is being tested in clinical trials of Phase I and Phase II for leukaemias, solid tumors and lymphomas.³⁵ Opium (isolated from *Papaver somniferum*) derived noscapine is being tested in Phase I and Phase II trials and nanotechnology assisted drug administration is being experimented due to its limited water-solubility.²⁴

Herbal Mixtures

The anticancer effect of popular Japanese and Chinese herbal formulation (Minor *Bupleurum* Combination), also known as Xiao Chai Hu Tang (China) or Sho-saiko-to (Japan) is attributed to herbal components, baical skullcap (*Scutellaria baicalensis*), baicalein, glycyrrhizin (licorice root), saikosaponins, ginsenosides, wogonin and gingerol. It has been in use for nearly 3000 years.^{36,37} Tanaka *et al.*³⁸ reported promising results in patients with colorectal adenomas.

Medicinal Plants Inducing Caspase Activated Apoptosis

The search for novel natural compounds by the natural product research^{39,40} has kept alive natural product discovery.⁴¹ There is also reported use of various standardized extracts or fractions of single or mixed herbs with anticancer effects as dietary supplements.⁴² Evaluation of potential cytotoxic anticancer agents

leading to apoptosis of cancer cells, theoretically arresting their growth and spread of neoplasms has been adapted in many manuscripts.⁴³⁻⁴⁶

Advanced *in silico* Techniques

Comparative *in silico* and biotechnological techniques have been applied to develop optimal screening strategies of medicinal plants and phytochemicals for revealing their elusive mode of caspase-activated apoptotic activity.

In silico docking

Molecular assay via computational methodology of docking process finds application in drug discovery program involves ligand (small molecule) - protein X-ray crystallography structure.⁴⁷ Compound from *Alkanna tinctoria*, 5-methoxyangerylalkannin, exhibited arrest of cancer cell in the S and G2/M phases inducing apoptosis. Docking (Surflex-Dock, Tripos, St. Louis, MO, USA) of its structure to caspase-9 (PDB code 2AR9) suggested formation of hydrogen bonds with hydroxyl groups of Asp-340 and Ser-339 at the active site.⁴⁸ Binding of eucalyptol, phytochemical from cardamom to caspase-3 analyzed by Auto Dock Pyrx was reported with poly-pharmacological anticancer property.⁴⁹ In another study, the 3D structures of neophytadine, nitrocyclohexane, octadecane and tetadecanoic acid were docked to caspase-3 receptor (PDB code 2X70). Nitrocyclohexane and neophytadiene with lower energy values specifically interact with the receptor supporting caspase-activated apoptosis.⁵⁰ Quercetin docked with caspase-3 exhibiting a strong interaction and docking score of -4.09 kcal/mol validates its anticancer activity.⁵¹ A set of 32 anticancer 3-aryl-5-aryl-1,2,4-oxadiazoles docked to caspase-3 with precise correlations validating apoptosis via caspase-3 activation.⁵² This combined assessment showed a good correlation between predicted and observed activity.

Designing molecules of smaller size for activation of caspase-3 from procaspase-3

Caspase-3, the executioner protein, is stored in its inactive procaspase-3 form with zinc inhibiting it at low concentrations and has a safety catch, a tripartite acid (DDD), blocking the isoleucine-glutamine-threonine-aspartate, ile-glu-thr-asp (IETD) site of proteolysis. A small promoter molecule, procaspase-3 activating compound, PAC-1, discovered by Putt *et al.*⁵³ could directly interact with and cleave this catch promoting the cancer cell to apoptosis. A set 38 molecules with positive charge at physiological pH, similar to piperazine nitrogen in PAC-1,⁵⁴ were screened for their capacity to directly interact with safety catch DDD in procaspase-3.

In silico using CDOCKER-a CHARMM-based MD docking algorithm tool of Discovery Studio was used.⁵⁵ This study identified above cited molecules with a capacity to promote caspase-activated apoptosis in cancer cells.⁵⁴

Phytochemical mediated differential expression of protein targets

Proteomics of apoptotic maslinic acid, a natural pentacyclic triterpene subjected to comparative proteomics and biotechnological studies revealed its capacity to down-regulate dUTP and stathmin. These are two proteins involved in induction of early S and G2 cell cycle arrest.^{55,56} A set of fourteen cytoskeleton proteins that were differentially expressed were reported.⁵⁷ Apoptosis induced by maslinic acid has been correlated with caspase and c-Jun N-terminal kinase activation.⁵⁸ These combinatorial studies could provide opportunities for phytochemicals to be developed into site- and target-specific anticancer agents.⁵⁹

Nano formulations for optimal activity

Targeting phytochemicals and drugs to dedicated sites via functionalized nanoparticles has evolved as new standard in anticancer methods.

- i. Functionalized nano carriers, folate-terminated polyrotaxanes (drug carrier) with dequalinium (compound for selective delivery of drug into mitochondria) with 101 nm size containing 18% classic cytotoxic anticancer drug, doxorubicin (DOX) was tested against DOX resistant cancerous cells of breast - MCF-7 and MCF-7/Adr xenografts of nude mice. A caspase-9 and -3 dependent apoptosis via activation of Bax, Bid and inhibition of Bcl-2 were reported. A six-fold intracellular uptake of DOX and a decreased drug efflux in comparison to free drug was reported.⁶⁰
- ii. A KLA peptide, D[KLAKLAK]2 modified with 2,3-dimethylmaleic anhydride (DMA) was combined with 1,2-distearoyl-*sn*-glycero-3-phosphoethanolamine (DSPE) and paclitaxel. It resulted in apoptotic paclitaxel nanodrug, DSPE-KLA-DMA (DKD) lipid. They could facilitate liposome internalization by reversing surface charge (-ve to +ve) at extracellular pH of ~ 6.8 into lung cancer, A549 cells. Apoptosis (cytotoxicity, 86.7%) via mitochondrial signaling pathway was evidenced by release of cytochrome c, increased caspase-9 and -3 activities.⁶¹
- iii. Liposomal assisted internalization of Anti-RNAs (antisense oligonucleotides, asOs) asOs initiates its binding and silencing of miRNAs with impaired invasion and proliferation of cancer. Overexpression

of phosphatase and tensin homolog (PTEN), programmed cell death protein 4 (PDCD4) with activation of caspase 3/7, an apoptotic mode of cytotoxicity was reported.⁶² miRNAs are involved in all stages of colorectal cancer.⁶³ Curcumin, resveratrol, quercetin, alpha-mangostin, Vitamin D, omega-3 PUFA have been reported with miRNA modulatory activity.¹⁸ Their anticancer capacity could be enhanced when entrapped in nanostructures detailed²⁵ improving their antineoplastic effect.

Synergism of phytochemical, cannabinoids from *Cannabis sativa* L. with cancer treatment drugs, temozolomide (chemotherapy drug) has been commonly used in patients with glioblastoma. Cannabinoids could enhance efficacy of bicalutamide or docetaxel, standard drugs used in prostate cancer treatment.⁶⁴

Phytochemicals of Important Kani Tribe Medicinal Plants with Apoptotic Properties Activating Caspase-8: an *in silico* Study

Kani tribal communities in Thodu hills of Kerala use non-timber minor forest products and preparations to meet their healthcare like for first aid remedies, to treat cough, cold, fever, headache, poisonous bites among others based on traditional knowledge.⁴ The present study was also initiated with an aim to identify phytochemicals reported in these plants with a capacity to activate caspases directing cancer cells towards apoptosis via *in silico* technology (Table 5). The data compiled and presented in this review includes phytochemicals (Figure 3) reported with potential anticancer activities find rare mention.^{65,66}

In vivo evaluation of *in silico* observations

Eugenol and berberine were identified to exhibit a relatively higher caspase-8 activating capacity (Table 5). The other phytochemicals did not exhibit this property. Hence Eugenol and berberine were identified for further studies for *in vivo* analysis. The activation of caspase-8 has been reported to be a pre-requisite for apoptosis. Data generated by Confocal Laser Scanning Microscopy (CLSM) on activation of apoptosis by eugenol (500 microg/ml)⁶⁷ and berberine (500 microg/ml)⁶⁸ at said time intervals of 24 and 48h (Figure 4) support their apoptotic property as visualized by blebbing of the cells visualized (indicated by arrows) as early as at 24hr of treatment.

HeLa cells were visualized under CLSM for morphological changes induced by exposure to eugenol and berberine. In contrast to clear body observed in HeLa cells of the control group, cells exposed to eugenol

Table 5: Swiss ADME Results.

Phytochemicals	Lipophilicity Log P o/w iLOGP)	Water solubility Log S (ESOL)	GI absorption	Lipinski	Ghose	Muegge	BA
Eugenol MPN: <i>Ocimum tenuiflorum</i> Linn. ⁸¹	2.37	-2.46	High	Yes; 0 violation	Yes	No; 1 violation: MW<200	0.55
Berberine MPN: <i>Coscinium fenestratum</i> (Gaertner) Colebr ⁸²	4.10	-4.20	High	Yes; 0 violation	yes	yes	0.55
Isorientine MPN: <i>Biophytumsensitivum</i> (Linn.) ⁸³	2.12	-2.70	Low	No; 2 violations: NorO>10, NHorOH>5	No; 1 violation: WLOGP<-0.4	No; 3 violations: TPSA>150, H-acc>10, H-don>5	0.15
methoxy Luteoline MPN: <i>Biophytumsensitivum</i> (Linn.) ⁸⁴	2.40	-3.73	Low	No; 2 violations: NorO>10, NHorOH>5	Yes	No; 3 violations: TPSA>150, H-acc>10, H-don>5	0.15
4-caffeoylquinic acid <i>Biophytumsensitivum</i> (Linn.) ⁸⁵	0.96	-1.62	low	Yes; 1 violation: NHorOH>5	No; 1 violation:	No; 2 violations: TPSA>150, H-don>5	0.11
Brucine MPN: <i>Strychnos nux-vomica</i> ⁸⁶	3.16	-2.92	High	Yes; 0 violation	yes	yes	0.55
Artocarpine MPN: <i>Artocarpus heterophyllus</i> (Linn.) ⁸⁴	4.14	-6.12	High	Yes; 0 violation	No; 1 violation: WLOGP>5.6	No; 1 violation: XLOGP3>5	0.55

The lipophilicity, water solubility, Lipinski, Ghose and Muegge rules for drug-like molecules have also approved the ligand. Gastro-intestinal (GI) tract absorption is high and bioavailability (BA) of the ligand resulted in the partition coefficient (QPlogPo/w) ranges from -0.59 to -1.62 and water solubility (QPlogS) i.e., critical for estimation of absorption and distribution of drugs within the body, ranged between -6.12 and 0.5.

Topological Polar Surface Area of the ligand is also appreciable.

All these pharmacokinetic parameters are within the acceptable range signifies the ligand (s) to be a typical drug molecule with <5 hydrogen bond donors, <10 hydrogen bond acceptors and QPlogPo/w <5.

The n-octanol/water partition coefficient (log Po/w) is a key physicochemical parameter for drug discovery depicting lipophilicity indices of the ligand as within the range.

The parameters measured for the ligand's solubility in water is to propose the ligand (s) to be an ideal drug.

MPN: Medicinal plant name

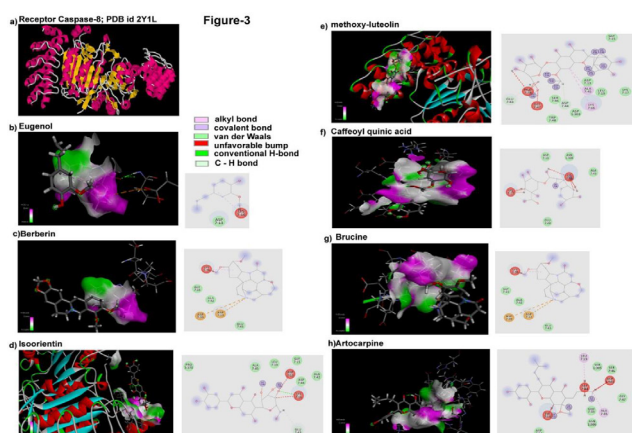


Figure 3: *In silico* docking of phytochemicals from important Kani tribe medicinal plants docked to receptor caspase-8 (PDB id 2Y1L).

(a) 2Y1L receptor caspase obtained from RCSB PDB; (b) Eugenol interaction with 2Y1L and estimated ΔG (kcal/mol) was -6.03; (c) Berberine interaction with 2Y1L and estimated ΔG (kcal/mol) was -6.28; (d) Isoorientin interaction with 2Y1L and estimated ΔG (kcal/mol) was -7.89; (e) methoxy Luteolin interaction with 2Y1L and estimated ΔG (kcal/mol) was -7.90; (f) Caffeoyl quinic acid interaction with 2Y1L and estimated ΔG (kcal/mol) was -8.19; (g) Brucine interaction with 2Y1L and estimated ΔG (kcal/mol) was -8.94 and (h) Artocarpine interaction with 2Y1L and estimated ΔG (kcal/mol) was -9.22.

(500 microg/mL) and berberine (500 microg/mL) exhibited the loss of these structural characteristics.

Eugenol and berberine treatment Caused caspase-8 activation in HeLa Cells

We tested whether eugenol and berberine-induced apoptosis was caspase-8 dependent after staining with CaspGLOW™ Fluorescein active Caspase-8 staining kit was procured for detection of active Caspase-8 in living cells. The FITC (fluorescent marker)-conjugated caspase-8 inhibitor, IETD-FMK permeabilized the cells. It bound to cytosolic active caspase-8 irreversibly. FITC on the IETD-FMK was detected. Eugenol and berberine-treated HeLa cells exhibited a time-dependent increase in mean caspase-8 fluorescence compared to control cells (Figure 4). For example, the caspase-8 fluorescence in HeLa cells treated with 500 microg/ml eugenol and 500 $\mu\text{g}/\text{mL}$ berberine was increased by approximately 7.9- and 6.5-fold, respectively compared to the control group (Figure 4g). The higher the capacity to activate caspase-8 supports the apoptotic potential of the phytochemical.

CONCLUSION

Cancer onset could be attributed to several factors. Apart from epigenetic, environmental factors, dietary habits could not be excluded.^{15,69} Consumption of plants and their parts has been advocated since time immemorial.⁶ As the elicitation of oxidative stress by generation of reactive oxygen species (ROS), a product

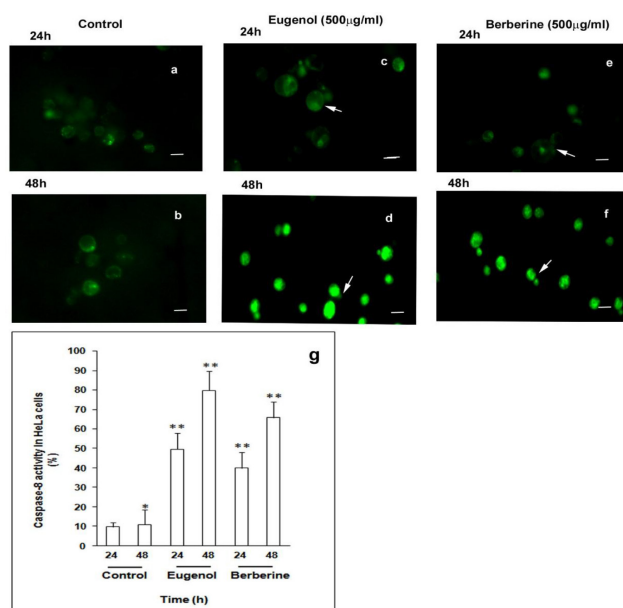


Figure 4: CaspGLOW™ Fluorescein detection of caspase-8 activated in HeLa cells upon treatment with eugenol (500mg/mL) and berberine (500mg/mL) for 24h and 48h.

Intracellular fluorescence of FITC on the IETD-FMK for all the samples was detected by Confocal Laser Scanning Microscope, LSM710 (Carl Zeiss, Germany). Control HeLa cells (a, b); eugenol treatment for 24h (c, d) and 48h (e, f); berberine treatment for 24h (c) and 48h (e). Arrows indicate blebbing of the cells. Bar corresponds to 10microm. Fluorescence was quantified (g) as detailed in materials and methods.

*Significant ($p < 0.05$) versus control cells.

of mitochondrial oxidative phosphorylation in human body has been considered to be involved in initiation of cancer onset.⁷⁰ Ample literature presently available, advocates on the merits of medicinal plants extracts and phytochemicals to alleviate cancer.³⁹ A few of the reported plant extracts^{36,38} and phytochemicals⁷¹ have already shown anticancer activity at clinical trials. Phytochemicals at academic level identified from the present compilation at pre-clinical stage should be introduced for clinical studies. They could be used alone or augmented with chemotherapeutic drugs. Phytochemicals identified in the present compilation could act as proactive preventive anticancer dietary regime even in genetically predisposed individuals to cancer.⁷² Thus, collaboration, guidance and support for herbal medicine use with prior systematic evaluation for cancer patients are needed as the dialogue between mainstream medicine and herbal medicine practitioners. An integration of herbal medicine into conventional cancer management strategies will aid in full realization of its value in cancer therapeutics. It could be a medico driven in comparison to a patient driven strategy.

ACKNOWLEDGEMENT

SS acknowledges the recognition of University of Mysore as an Institution of Excellence. The financial

support from the Ministry of Human Resource Development, New Delhi, Govt. of India through UGC under UOM/IOE/RESEARCH/1/2010-11, dt 22-04-2010 project. The computational facility at the Bioinformatics Centre, Maharani Science College, Palace Road, Bengaluru, India was used during this work and it is gratefully acknowledged.

CONFLICT OF INTEREST

There is no conflict of interest and the authors declare the same.

ABBREVIATIONS

2Y1L: Caspase-8 of from *Homo sapiens* complex with DARP (dopamine releasing protein) in -8.4 from protein data bank; **ADMET**: Absorption, Distribution, Metabolism, Excretion, Toxicity; **Alu elements**: An Alu element is a short stretch of DNA originally characterized by the action of the *Arthrobacter luteus* (*Alu*) restriction endonuclease; **CARD**: Caspase Recruitment Domain; **CASP-8**: *CASP8* gene encodes Caspase-8 protein; **CED-3**: Ced-3 is an executioner caspase (cysteine-dependent aspartate-directed protease); **DARP**: Dopamine releasing protein; **FDA**: US Food and Drug Administration; Flex and X, Surflex20: Docking tools and programs have been developed for both academic and commercial applications; **MALDI-TOF**: Matrix-Assisted Laser Desorption/Ionization-Time of Flight (MALDI-TOF); **NCI**: The National Cancer Institute; **WHO**: The World Health Organization

REFERENCES

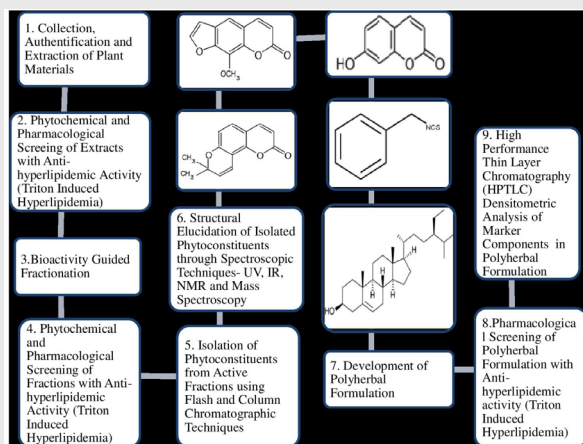
- Nicholson DW, Thornberry NA. Caspases: Killer proteases. *Trends Biochem Sci.* 1997;22(8):299-306.
- Parrish AB, Freel CD, Kombluth S. Cellular mechanism controlling caspase activation and function. *Cold Spring Harb Perspect Biol.* 2013;5(6):1-5.
- Kumar S. Caspase function in programmed cell death. *Cell Death Differ.* 2007;14(1):32-43.
- Xavier TF, Kannan M, Lija L, Auxillia A, Kanthi A, Rose F, et al. Ethnobotanical study of Kani tribes in Thoduhills of Kerala, South India. *J Ethnopharmacol.* 2014;152(1):78-90.
- Koehn FE. High impact of technologies for natural products screening. *Prog Drug Res.* 2008;65:177-210.
- Atanasov AG, Waltenberger B, Wenzing EMP, Linder T, Wawrosch C, Uhrin P, et al. Discovery and resupply of pharmacologically active plant-derived natural products: A review. *Biotechnol Adv.* 2015;33(8):1582-614.
- Rodrigues T, Reker D, Schneider P, Schneider G. Counting on natural products for drug design. *Nat Chem.* 2016;8(6):531.
- Olsson M, Zhivotovsky B. Caspases and cancer. *Cell Death Differ.* 2011;18(9):1441-9.
- Teitz T, Wei T, Valentine MB, Vanin EF, Grenet J, Valentine VA, et al. Caspase 8 is deleted or silenced preferentially in childhood neuroblastomas with amplification of MYCN. *Nat Med.* 2000;6(5):529-35.
- Ashley DM, Riffkin CD, Muscat AM, Knight MJ, Kaye AH, Novak U, et al. Caspase 8 is absent or low in many ex vivo gliomas. *Cancer.* 2005;104(7):1487-96.

- Kim HS, Lee JW, Soung YH, Park WS, Kim SY, Lee JH, et al. Inactivating mutations of caspase-8 gene in colorectal carcinomas. *Gastroenterol.* 2003;125(3):708-15.
- Soung YH, Lee JW, Kim HS, Park WS, Kim SY, Lee JH, et al. Inactivating mutations of CASPASE-7 gene in human cancers. *Oncogene.* 2003;22(39):8048-52.
- Kaufmann SH, Lee SH, Meng XW, Loegering DA, Kottke TJ, Henzing AJ, et al. Apoptosis-associated caspase activation assays. *Methods.* 2008;44(3):262-72.
- Cragg GM, Newman DJ. Plants as a source of anticancer agents. *J Ethnopharmacol.* 2005;100(1-2):72-9.
- Bhaskaran K, Douglas I, Forbes H, Dos-Santos-Silva I, Leon DA, Smeeth L. Body-mass index and risk of 22 specific cancers: A population-based cohort study of 5-24 million UK adults. *Lancet.* 2014;384(9945):755-65.
- Willett WC, Koplan JP, Nugent R, Dusenbury C, Puska P, Gaziano TA. Prevention of chronic disease by means of diet and lifestyle changes. In *Disease control priorities in developing countries* chapter 44, 2nd edition. Oxford University Press. 2006;833-50.
- Wang H, Khor TO, Shu L, Su Z, Fuentes F, Lee JH, et al. Plants against cancer: A review on natural phytochemicals in preventing and treating cancers and their drug ability. *Anticancer Agents Med Chem.* 2012;12(10):1281-305.
- Gavrilas LI, Ionescu C, Tudoran O, Lisencu C, Balacescu O, Miere D. The role of bioactive dietary components in modulating miRNA expression in colorectal cancer. *Nutrients.* 2016;8(10):590.
- Pezzuto JM. Plant-derived anticancer agents. *Biochem Pharmacol.* 1997;53(2):121-33.
- Cassileth BR. Evaluating complementary and alternative therapies for cancer patients. *CA Cancer J Clin.* 1999;49(6):362-75.
- Aggarwal BB, Ichikawa H, Garodia P, Weerasinghe P, Sethi G, Bhatt ID, et al. From traditional Ayurvedic medicine to modern medicine: Identification of therapeutic targets for suppression of inflammation and cancer. *Expert Opin Ther Targets.* 2006;10(1):87-118.
- Newman DJ, Cragg GM. Natural products as sources of new drugs over the 30 years from 1981 to 2010. *J Nat Prod.* 2012;75(3):311-35.
- Wong FC, Tan ST, Chai TT. Phytochemical mediated protein expression profiling and the potential application in therapeutic drug target identification. *Critical Rev Food Sci Nutr.* 2016;56(Sup 1):S162-70.
- Che E, Gao Y, Wan L, Zhang Y, Han N, et al. Paclitaxel/gelatin coated magnetic mesoporous silica nanoparticles: Preparation and antitumor efficacy *in vivo*. *Microporous Mesoporous Mater.* 2015;204:226-34.
- Piktel E, Niemirowicz K, Watek M, Wolny T, Deptula P, Bucki R. Recent insights into nanotechnology-based drugs and formulations designed for effective anticancer therapy. *J Nanobiotechnol.* 2016;14(1):39.
- Heiss E, Herhaus C, Klimo K, Bartsch H, Gerhäuser C. Nuclear Factor κB is a molecular target for sulforaphane-mediated anti-inflammatory mechanisms. *J Biol Chem.* 2001;276(34):32008-15.
- Pledge-Tracy A, Sobolewski MD, Davidson NE. Sulforaphane induces cell type-specific apoptosis in human breast cancer cell lines. *Mol Cancer Ther.* 2007;6(3):1013-21.
- Comblatt BS, Ye L, Dinkova-Kostova AT, Erb M, Fahey JW, Singh K, et al. Preclinical and clinical evaluation of sulforaphane for chemoprevention in the breast. *Carcinogenesis.* 2007;28(7):1485-90.
- Unnati S, Ripal S, Sanjeev A, Niyati A. Novel anticancer agents from plant sources. *Chinese J Nat Med.* 2013;11(1):16-23.
- Soloway E, Lichtenstein M, Sallo S, Paavilainen H, Soloway E, Lorberbaum-Galski H. Evaluating medicinal plants for anticancer activity. *Sci World J.* 2014;2014:1-2.
- Jordan MA, Wilson L. Microtubules as a target for anticancer drugs. *Nat Rev Cancer.* 2004;4(4):253-65.
- Risinger AL, Giles FJ, Mooberry SL. Microtubule dynamics as a target in oncology. *Cancer Treat Rev.* 2009;35(3):255-61.
- Amin A, Muhtasib GH, Ocker M, Schneider-Stock R. Overview of major classes of plant-derived anticancer drugs. In *J Biomed Sci.* 2009;5(1):1-1.
- Raza H, John A. Green tea polyphenol epigallocatechin-3-gallate differentially modulates oxidative stress in PC12 cell compartments. *Toxicol Appl Pharmacol.* 2005;207(3):212-20.
- Newcomb EW. Flavopiridol: Pleiotropic biological effects enhance its anticancer activity. *Anti Cancer Drug.* 2004;15(5):411-9.

36. Zheng N, Dai J, Cao H, Sun S, Fang J, Li Q, et al. Current understanding on anti-hepatocarcinoma effects of xiao chai hu tang and its constituents. *Evid Based Complement Alternat Med*. 2013;2013:1-14.
37. Steuer-Vogt MK. The effect of an adjuvant mistletoe treatment program in resected head and neck cancer patients: A randomized controlled clinical trial. *Eur J Cancer*. 2001;37(1):23-31.
38. Tanaka S, Haruma K, Yoshihara M, Kajiyama G, Kira K, Amagase H, et al. Aged garlic extract has potential suppressive effect on colorectal adenomas in humans. *J Nutr*. 2006;136(3):821S-6S.
39. Atanasov AG, Waltenberger B, Wenzing EMP, Linder T, Wawrosch C, Uhrin P, et al. Discovery and resupply of pharmacologically active plant-derived natural products: A review. *Biotechnol Adv*. 2015;33(8):1582-614.
40. Muhtasib GH. Emerging cytotoxic alkaloids in the battle against cancer: Overview of molecular mechanisms. *Molecules*. 2007;250:1-22.
41. Davis CD. Cellular cancer processes and their molecular targets for nutritional pre-emption of cancer. In: 'Bioactive compounds and cancer'. Humana Press. 2010;50-6.
42. Gunasinghe S, Arambewela L. The role played by plant sources in cancer therapy in Sri Lanka. *J Indigenous Med*. 2014;1-4.
43. Pan MH, Ghai G, Ho CT. Food bioactives, apoptosis and cancer. *Mol Nutr Food Res*. 2008;52(1):43-52.
44. Wang H, Khor TO, Shu L, Su Z, Fuentes F, Lee JH, et al. Plants against cancer: A review on natural phytochemicals in preventing and treating cancers and their drug ability. *Anticancer Agents Med Chem*. 2012;12(10):1281-305.
45. Baliga MS, Venkatesh S, Mrinal S, Bala N, Palatty PL. Bioactive dietary factors and plant extracts in dermatology Part III. In 'Plant and Plant components and skin care. Humana Press. 2013;93-136.
46. Mund MD, Alam S, Khan UH, Tahir U, Zubair MS, Younas T, et al. Phytochemicals as complementary and alternative therapeutic formulations with potential proapoptotic effects on various cancerous cell lines: A literature survey. *Focus Sci*. 2016;2(2):1-5.
47. Sliwoski G, Kothiwale S, Meiler J, JrLowe EW. Computational methods in drug discovery. *Pharmacol Rev*. 2014;66(1):334-95.
48. Tung NH, Du GJ, Yuan CS, Shoyama Y, Wang CZ. Isolation and chemopreventive evaluation of novel naphthoquinone compounds from *Alkanna tinctoria*. *Anticancer Drugs* 2013;24(10):1058-68.
49. Bhattacharjee B, Chatterjee J. Identification of proapoptotic, anti-inflammatory, anti-proliferative, anti-invasive and anti-angiogenic targets of essential oils in cardamom by dual reverse virtual screening an binding pose analysis. *Asian Pac J Cancer Prev*. 2013;14(6):3735-42.
50. Selvamangai G, Bhaskar A. Analysis of phytochemicals in the methanolic extract of *Eupatorium triplinerve* by GC-MS method. *Int J Drug Dev Res*. 2013;5:384-91.
51. Muthukala B, Sivakumari K, Ashok K. *In silico* docking of quercetin compound against the HeLa cell line proteins. *Int J Curr Pharma Res*. 2015;7(1):13-6.
52. Vaidya A, Jain AK, Kumar BRP, Sastry GN, Kashaw SK, Agarawal RK. CoMFA, CoMSIA, kNN, MFA and docking studies of 1,2,4-oxadiazole derivatives as potent caspase-3 activators. *Arabian J Chem*. 2017;10:S3936-46.
53. Putt KS, Chen GW, Pearson JM, Sandhorst JS, Hoagland MS, Kwon JT, et al. Small molecule activation of procaspase-3 to caspase-3 as a personalized anticancer strategy. *Nat Chem Biol*. 2010;2(10):543-50.
54. Kumar MS, Lainu KL, Aghila V, Purushothaman D, Gopal KV, Namboori PKK, et al. Designing a promoter for a novel target site identified in caspases for initiating apoptosis in cancer cells. In *Inf Commun Tech*: Springer-Verlag Berlin Heidelberg. 2010;62-7.
55. Wu G, Robertson DH, Brooks CL. Detailed analysis of Grid-based molecular docking: A case study of CDOCKER-A CHARMM-based MD docking algorithm. *J Comp Chem*. 2003;24(13):1549-62.
56. Yap WH, Khoo KS, Lim SH, Yeo CC, Lim YM. Proteomic analysis of the molecular response of Raji cells to maslinic acid treatment. *Phytomedicine*. 2012;19(2):183-91.
57. Rufino-Palomares EE, Reyes-Zurita FJ, Garcia-Salgureo L, Mokhtari K, Medina PP, Lupianez JA, et al. Maslinic acid, a triterpenic anti-tumoral agent, interferes with cytoskeleton protein expression in HT29 human colon-cancer cells. *J Proteomics*. 2013;83:15-25.
58. Reyes-Zurita FJ, Rufino-Palomares EE, Medina PP, Garcia-Salgureo L, Peragon J, Cascante M, et al. Antitumor activity on extrinsic apoptotic targets of the triterpenoid maslinic acid in p53 deficient Caco-2 adenocarcinoma cells. *Biochimie*. 2013;95(11):2157-67.
59. Wong FC, Tan ST, Chai TT. Phytochemical mediated protein expression profiling and the potential application in therapeutic drug target identification. *Critical Rev Food Sci Nutr*. 2016;56(Suppl 1):S162-70.
60. Wang H, Yin H, Yan F, Sun M, Du L, Peng W, et al. Folate-mediated mitochondrial targeting with doxorubicin-polyrotaxane nanoparticles overcomes multidrug resistance. *Oncotarget*. 2015;6(5):2827.
61. Jiang L, Li L, He X, Yi Q, He B, Cao J, et al. Overcoming drug resistant lung cancer by paclitaxel loaded dual-functional liposomes with mitochondria targeting and pH-response. *Biomater*. 2015;52:126-39.
62. Costa PM, Cardoso AL, Mendonça LS, Serani A, Custódia C, Conceição M, et al. Tumor-targeted chlorotoxin-coupled nanoparticles for nucleic acid delivery to glioblastoma cells: A promising system for glioblastoma treatment. *Mol Ther Nucleic Acids*. 2013;1-4.
63. Aslam MI, Patel M, Singh B, Jameson JS, Pringle JH. MicroRNA manipulation in colorectal cancer cells: From laboratory to clinical application. *J Transl Med*. 2012;10(1):128.
64. Velasco G, Sanchez C, Guzman M. Anticancer mechanisms of cannabinoids. *Curr Oncol*. 2016;23(Suppl 2):S23.
65. Dubois MAL. Bioactive saponins with cancer related and immunomodulatory activity: Recent developments. *Stud Nat Prod Chem*. 2005;32:209-46.
66. Dubois MAL. Bioactive saponins with cancer related and immunomodulatory activity: Recent developments. In: *Studies in Natural Products Chemistry: Bioactive natural products, Part I*, Elsevier, Netherlands. 2005;32:209-46.
67. Das A, Harshadha K, Kannan DSK, Hari RK, Jayaprakash B. Evaluation of therapeutic Potential of eugenol-a natural derivative of *Syzygium aromaticum* on cervical cancer. *Asian Pac J Cancer Prev*. 2018;19(7):1977.
68. LuB, Hu M, Liu K, Peng J. Cytotoxicity of berberine on human cervical carcinoma HeLa cells through mitochondria, death receptor and MAPK pathways and *in-silico* drug-target prediction. *Toxicol in vitro*. 2010;24(6):1482-90.
69. Habli Z, Toumeh G, Fattat M, Rahal ON, Muhtasib GH. Emerging cytotoxic alkaloids in the battle against cancer: Overview of molecular mechanisms. *Molecules*. 2017;22(2):250.
70. Aykan NF. Red meat and colorectal cancer. *Oncol Rev*. 2015;9(1):288.
71. Panieri E, Santoro MM. ROS homeostasis and metabolism: A dangerous liason in cancer cells. *Cell Death Dis*. 2016;7(6):e2253.
72. Jiang L, Li L, He X, Yi Q, He B, Cao J, et al. Overcoming drug resistant lung cancer by paclitaxel loaded dual-functional liposomes with mitochondria targeting and pH-response. *Biomater*. 2015;52:126-39.
73. Kischkel FC, Lawrence DA, Tinel A, LeBlanc H, Virmani A, Schow P, et al. Death receptor recruitment of endogenous caspase-10 and apoptosis initiation in the absence of caspase-8. *J Biol Chem*. 2001;276(49):46639-46.
74. Yoo NJ, Lee JW, Kim YJ, Soung YH, Kim SY, Nam SW, et al. Loss of caspase-2, -6 and -7 expression in gastric cancers. *APMIS* 2004;112:330-5.
75. Palmerini F, Devillard E, Jarry A, Birg F, Xerri L. Caspase 7 downregulation as an immunohistochemical marker of colonic carcinoma. *Hum Pathol*. 2001;32(5):461-7.
76. Devarajan E, Sahin AA, Chen JS, Krishnamurthy RR, Aggarwal N, Brun AM, et al. Downregulation of caspase 3 in breast cancer: A possible mechanism for chemoresistance. *Oncogene*. 2002;21(57):8843-51.
77. Nakopoulou L, Alexandrou P, Stefanaki K, Panayotopoulou E, Lazaris AC, Davaris PS. Immunohistochemical expression of caspase-3 as an adverse indicator of the clinical outcome in human breast cancer. *Pathobiology*. 2001;69(5):266-73.
78. Deeb D, Jiang H, Gao X, Al-Holou S, Danyluk AL, Dulchavsky SA, et al. Curcumin [1,7-bis(4-hydroxy-3-methoxyphenyl)-1,6-heptadine-3,5-dione; C21H20O6] sensitizes human prostatecancer cells to tumor necrosis factor-related apoptosis-inducing ligand/Apo2L-induced apoptosis by suppressing nuclear factor-kappaB via inhibition of the prosurvival Akt signaling pathway. *J Pharmacol Exp Ther*. 2007;321(2):616-25.
79. Khan N, Adhami VM, Mukhtar H. Apoptosis by dietary agents for prevention and treatment of prostate cancer. *Endocr Relat Cancer*. 2011;17(1):R39-52.
80. Shailasree S, Venkataramana M, Niranjana SR, Prakash HS. Cytotoxic effect of *p*-coumaric acid on neuroblastoma, N2a cell via generation of reactive oxygen species leading to dysfunction of mitochondria inducing apoptosis and autophagy. *Mol Neurobiol*. 2015;51(1):119-30.

81. Tuso PJ. Nutritional update for physicians: Plant based diets. *Perm J*. 2013;17(2):61.
82. Pattanayak P, Behera P, Das D, Panda SK. *Ocimum sanctum* Linn. A reservoir plant for therapeutic applications: An overview. *Pharmacogn Rev*. 2010;4(7):95.
83. Singh A, Bajpai V, Kumar S, Kumar KBR, Kumar B. Simultaneous quantification of protoberberine and aporphine alkaloids in different plant parts of *Cosciniium fenestratum* (Gaertner) Colebr. By liquid chromatography-hybrid triple quadrupole/liner ion trap mass spectrometer. *J Med Plants Studies*. 2016;4(3):144-8.
84. Sakhivel KM, Guruvayoorappan C. *Biophytum sensitivum*: Ancient medicine, modern targets. *J Adv Pharm Technol Res*. 2012;3(2):83.
85. Yin W, Deng XK, Yin FZ, Zhang XC, Cai BC. The cytotoxicity induced by brucine from the seed of *Strychnos nux-vomica* proceeds via apoptosis and is mediated by cyclooxygenase 2 and caspase 3 in SMMC 7221 cells. *Food Chem Toxicol*. 2007;45(9):1700-8.
86. Lee CW, Hsu LF, Lee MH, Lee I, Liu JF, Chiang YC, et al. Extracts of *Artocarpus communis* induce mitochondria-associated apoptosis via pro-oxidative activity in human glioblastoma cells. *Front Pharmacol*. 2018;9:411.

PICTORIAL ABSTRACT



SUMMARY

- Phytochemicals with a capacity to activate caspases enhancing apoptotic capacity has been proven to be effective anticancer agents.
- Data on phtochemicals traditionally used to treat cancerous conditions and the scientific validation of caspase-activated apoptosis for this traditional application has been compiled. Internet assisted scientific literature was collected.
- Technological advancement has been included for identification of hit molecule and lead optimization.
- Eugenol and berberine were identified as phytochemicals with potential drug characteristics by both *in silico* and *in vivo* studies.
- The phytochemicals from important Kani tribal medicinal plants via *in silico* docking and *in vivo* studies identified could be explored at clinical trials.

About Authors



Dr. Shailasree Sekhar is recipient of UGC-CSIR NET JRF-SRF grants for her doctoral studies from CSIR-CFTRI, 2000. She is "Young Scientist, SERB, Department of Science and Technology, Government of India" awardee. Currently, as Scientist at The Institution of Excellence, University of Mysore with the thrust area identified as the biodiversity of Western Ghats medicinal plants (MP), with immunological affections and cancer prevention properties, due to location advantage of this hot spot to the University, she has been actively involved in compilation of their scientific data as reviews. Recently, she has brought out a database on medicinal plants of Western Ghats in an efficient way. Screening of MP with immunological affections used by Western Ghats tribal population has resulted in identification of several of them with inflammation/ cancer inhibiting property. Fingerprinting their metabolites has been her priority. She has published several scientific reports/ papers in peer-reviewed journals, has 2 patents and is an adhoc reviewer of various journals. She has to her credit grants from National scientific agencies, DST and UGC under Government of India.

Cite this article: Saqib A, Pattar S, Karigar CS, Sekhar S. Caspase Activators: Phytochemicals with Apoptotic Properties Targeting Cancer, a Health Care Strategy to Combat this Disease. *Indian J of Pharmaceutical Education and Research*. 2020;54(3):496-508.