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

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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 phtochemicals 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, ResearchGate 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.

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 elegeon 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
mortality. 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 protease) was reported. This family of enzymes comprises 11 members in humans and 13 in mammalians. 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 an 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.

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.

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, 1944,000 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, ResearchGate 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 tribes recommended in the review by Xavier et al. 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 likeness 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 likeness 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 likeness 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 1×10⁵ cells/mL were grown on 12 mm glass coverslips. When cell density reached 80–90% confluence, cells were treated with eugenol (500m/mL) and berberine (500m/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 synthentic 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

![Table 1: Expression of caspases in cancerous condition.](image-url)

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

↑ = activated/ improved activity/ upregulated
↓ = inhibited/ downregulated
Italized = gene; unitalicized = enzyme;
Table 2: Assays to detect active caspase induced by apoptotic stimuli.

<table>
<thead>
<tr>
<th>Method</th>
<th>Assays</th>
<th>Advantages</th>
<th>Disadvantages</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fluorogenic and chromogenic assay ELISA</td>
<td>• Estimation of cleaved substrate products and estimating fluorogenic or chromogenic leaving groups</td>
<td>• Substrates used: aminofluorocoumarins N-acetylaspartyl-glutamylvalylaspartyl-7-amino-4-trifluoromethyl coumarin (DEVD-AFC) for caspase-3 and -7; N-acetylvalinyl-glutamylisoleucylaspartyl-AFC (VEID-AFC) for caspase-6;</td>
<td>• Non-specificity as DEVD-AFC, preferred substrate for caspase-3 is cleaved by caspases-1, -2, -4, -6, -7, -8, -10, and -14</td>
</tr>
<tr>
<td>Fluorescence microscopy or flow cytometry</td>
<td>• Fluorescence microscopy or flow cytometry</td>
<td>• Advantages:</td>
<td></td>
</tr>
<tr>
<td>Immunoblotting</td>
<td>Using sera for caspase cleavage products, the anti-neoepitope antibodies</td>
<td>• Procaspe 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)</td>
<td>• The level of caspase cleavage might not reflect the quantity of activated caspase as it is dependent on the stability of cleavage products</td>
</tr>
<tr>
<td>Immunofluorescence or flow cytometry</td>
<td>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.</td>
<td>• Specificity of the reagents used aids in assessing with certainty activation of a particular caspase</td>
<td></td>
</tr>
<tr>
<td>Affinity labeling for immunoblotting, fluorescence microscopy and flow cytometry</td>
<td>Using covalent modification of the caspase active site with a substrate-like molecule with an inhibitory group and a reporter moiety</td>
<td>• Acyloxymethylketones (aomks) with reporter moiety (biont 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.</td>
<td>• The reagents are expensive and require custom synthesis</td>
</tr>
<tr>
<td>RT-PCR</td>
<td>Accession numberSequence</td>
<td>• Caspase 3U49930F5′-AATTCAAGGGACGGGTCATG-3′</td>
<td></td>
</tr>
</tbody>
</table>

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

<table>
<thead>
<tr>
<th>Bioactive compound</th>
<th>Cell line (s); cytotoxic concentrations (IC\textsubscript{50} values)</th>
<th>Therapeutic targets / Mode of action</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>Curcumin</td>
<td>PC-3; IC\textsubscript{50} 20 mm/L HepG2, [40 mg/mL]</td>
<td>TRAIL-induced apoptosis; ↓Δ\textsubscript{ψm}t</td>
<td>Deeb \textit{et al.}\textsuperscript{18}</td>
</tr>
<tr>
<td>Sulfuraphane</td>
<td>MDA-MB-231, [25 mm/L]</td>
<td>↑Caspase-8, -3; ↓Bcl-2</td>
<td>Pledgie-Tracy \textit{et al.}\textsuperscript{27}</td>
</tr>
<tr>
<td>EGC or EGCG</td>
<td>SH-SYSY, [50 mm/L]</td>
<td>↑Caspase-8,</td>
<td>Das \textit{et al.}\textsuperscript{67}</td>
</tr>
<tr>
<td>beta-carotene</td>
<td>HT-29 [55 mm/L]</td>
<td>↑Caspase-8, -9; ↓Bcl-2</td>
<td>Khan \textit{et al.}\textsuperscript{79}</td>
</tr>
<tr>
<td>Apigenin</td>
<td>22Rv1, [80 mm/L]</td>
<td>↑Caspase-9, -3; ↑Δ\textsubscript{ψm}t; ↑Caspase-9</td>
<td>Khan \textit{et al.}\textsuperscript{79}</td>
</tr>
<tr>
<td>Luteolin</td>
<td>SCC-4, [100 mm/L]</td>
<td>↑Caspase-3, -9; ↓PARP</td>
<td>Khan \textit{et al.}\textsuperscript{79}</td>
</tr>
<tr>
<td>Resveratrol</td>
<td>MD-MB-231, [50 mm/L]</td>
<td>↑Caspase-3; ↓PARP</td>
<td>Khan \textit{et al.}\textsuperscript{79}</td>
</tr>
<tr>
<td>Iberin</td>
<td>SK-N-AS, [50 mm/L]</td>
<td>↑Caspase-3, -9; ↓PARP</td>
<td>Khan \textit{et al.}\textsuperscript{79}</td>
</tr>
<tr>
<td>p-coumaric acid</td>
<td>N2a,[150 mm/L]</td>
<td>↑Caspase-8</td>
<td>Shailasree \textit{et al.}\textsuperscript{80}</td>
</tr>
<tr>
<td>α-mangostin</td>
<td>COLO, [205 9.74±0.85 mg/mL]</td>
<td>↑Caspase-3, -8, -9</td>
<td>Gavrilas \textit{et al.}\textsuperscript{18}</td>
</tr>
<tr>
<td>Omega-3 (PUFA)</td>
<td>LT97, [100 mM]</td>
<td>↑Caspase-3, -9</td>
<td>Khan \textit{et al.}\textsuperscript{79}</td>
</tr>
</tbody>
</table>

Δ\textsubscript{ψm}t – mitochondrial membrane potential; PARP: poly ADP ribose polymerase; TRAIL:TNF-related apoptosis inducing ligand

\textsuperscript{↑} = activated/improved activity/unregulated

\textsuperscript{↓} = inhibited/downregulated

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

from around 35,000 plant samples for anticancer activity were screened. By 1991, 28,800 plant samples from over 20 countries were catalogued for chemotherapeutic capacity initially screened \textit{in vitro} against human cancer cell lines.\textsuperscript{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 \textit{Taxus brevifolia}, vinblastine and vincristine from \textit{Catharanthus roseus}, etoposide and teniposide from \textit{Podophyllum peltatum}, topotecan from \textit{Camptotheca acuminate} and irinotecan from \textit{Camptotheca acuminate} (Table 4). The most successful discoveries are the vinca alkaloids (vinblastin and vincristine) from \textit{Catharanthus roseus}.\textsuperscript{23}
Table 4: Plant extracts and compounds (selective) that were screened against NCI-60 Cancer cell panel, information referring NCI website.

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

*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.

Clinical Trials
Paclitaxel (taxol/ taxane) from Taxus brevifolia L. with reported ability to block mitosis, disrupt microtubule and spindle formation and inhibit translational 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 nanoliposomes are being developed as alternate form of administration of this drug. 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. 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. Vinca alkaloids from Catharanthus roseus G. Don with apoptotic capacity, bind beta-tubulin and destabilize microtubule, induce cell cycle arrest have been incorporated in clinical trials. Vinca alkaloids from Catharanthus roseus G. Don with apoptotic capacity, bind beta-tubulin and destabilize microtubule, induce cell cycle arrest have been incorporated in clinical trials. Vinca alkaloids from Catharanthus roseus G. Don with apoptotic capacity, bind beta-tubulin and destabilize microtubule, induce cell cycle arrest have been incorporated in clinical trials. Vinca alkaloids from Catharanthus roseus G. Don with apoptotic capacity, bind beta-tubulin and destabilize microtubule, induce cell cycle arrest have been incorporated in clinical trials. Vinca alkaloids from Catharanthus roseus G. Don with apoptotic capacity, bind beta-tubulin and destabilize microtubule, induce cell cycle arrest have been incorporated in clinical trials. Vinca alkaloids from Catharanthus roseus G. Don with apoptotic capacity, bind beta-tubulin and destabilize microtubule, induce cell cycle arrest have been incorporated in clinical trials. Vinca alkaloids from Catharanthus roseus G. Don with apoptotic capacity, bind beta-tubulin and destabilize microtubule, induce cell cycle arrest have been incorporated in clinical trials. Vinca alkaloids from Catharanthus roseus G. Don with apoptotic capacity, bind beta-tubulin and destabilize microtubule, induce cell cycle arrest have been incorporated in clinical trials. Vinca alkaloids from Catharanthus roseus G. Don with apoptotic capacity, bind beta-tubulin and destabilize microtubule, induce cell cycle arrest have been incorporated in clinical trials. Vinca alkaloids from Catharanthus roseus G. Don with apoptotic capacity, bind beta-tubulin and destabilize microtubule, induce cell cycle arrest have been incorporated in clinical trials. Vinca alkaloids from Catharanthus roseus G. Don with apoptotic capacity, bind beta-tubulin and destabilize microtubule, induce cell cycle arrest have been incorporated in clinical trials.

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, baikal skullcap (Scutellaria baicalensis), baikalins, glycyrrhizin (licorice root), saikosaponins, ginsenosides, wogonin and gingerol. It has been in use for nearly 3000 years. Tanaka et al. 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\textsuperscript{39,40} has kept alive natural product discovery.\textsuperscript{41} There is also reported use of various standardized extracts or fractions of single or mixed herbs with anticancer effects as dietary supplements.\textsuperscript{12} 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.\textsuperscript{43-46}

Advanced \textit{in silico} Techniques

Comparative \textit{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.

\textbf{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.\textsuperscript{47} Compound from \textit{Alkanna tinctor}, 5-methoxyangenylalkannin, 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.\textsuperscript{48} Binding of eucalyptol, phytochemical from cardamom to caspase-3 analyzed by Auto Dock Pyrx was reported with poly-pharmacological anticancer property.\textsuperscript{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.\textsuperscript{50} Quercetin docked with caspase-3 exhibiting a strong interaction and docking score of \(-4.09\text{ kcal/mol}\) validates its anticancer activity.\textsuperscript{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.\textsuperscript{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 triaspartic acid (DDD), blocking the isoleucine-glutamine-threone-aspartate, ile-glu-thr-asp (IETD) site of proteolysis. A small promotor molecule, procaspase-3 activating compound, PAC-1, discovered by Putt \textit{et al.}\textsuperscript{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 piperazone nitrogen in PAC-1,\textsuperscript{54} were screened for their capacity to directly interact with safety catch DDD in procaspase-3. \textit{In silico} using CDOCKER-a CHARMM-based MD docking algorithm tool of Discovery Studio was used.\textsuperscript{55} This study identified above cited molecules with a capacity to promote caspase-activated apoptosis in cancer cells.\textsuperscript{54}

\textbf{Phytochemical mediated differential expression of protein targets}

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

\textbf{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/Adrxenografts of nude mice. A caspase-9 and -3 dependent apoptosis via activation of Bax, Bid and inhibition of Bel-2 were reported. A six-fold intracellular uptake of DOX and a decreased drug efflux in comparison to free drug was reported.\textsuperscript{60}

ii. A KLA peptide, D[KLAKLAK]\textsubscript{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 extra-
cellular 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.\textsuperscript{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.\textsuperscript{62} miRNAs are involved in all stages of colorectal cancer.\textsuperscript{63} Curcumin, resveratrol, quercetin, alpha-mangostin, Vitamin D, omega-3 PUFA have been reported with miRNA modulatory activity.\textsuperscript{18} Their anticancer capacity could be enhanced when entrapped in nanostructures detailed\textsuperscript{65} improving their antineoplastic effect.

Synergism of phytochemical, cannabinoids from Cannabis sativa L. with cancer treatment drugs, temozolomide (chemotheraphy drug) has been commonly used in patients with glioblastoma. Cannabinoids could enhance efficacy of bicalutamide or docetaxel, standard drugs used in prostate cancer treatment.\textsuperscript{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.\textsuperscript{3} 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.\textsuperscript{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 (500mg/ml)\textsuperscript{67} and berberine (500 mg/ml)\textsuperscript{68} at said time intervals of 24 and 48h (Figure 4) support...
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 -7.90; (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) Caffoylqunic 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) Arocarpin interaction with 2Y1L and estimated ΔG (kcal/mol) was -9.22.

Figure 4: CaspGlow™ Fluorescein detection of caspase-8 activated in HeLa cells upon treatment with eugenol (500 mg/mL) and berberine (500 mg/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 10 microm. Fluorescence was quantified (g) as detailed in materials and methods.

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

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 (500 m/mL) and berberine (500 mg/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 m/mL eugenol and 500 μg/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. Consumption of plants and their parts has been advocated since time immemorial. As the elicitation of oxidative stress by generation of reactive oxygen species (ROS), a product of mitochondrial oxidative phosphorylation in human body has been considered to be involved in initiation of cancer onset. Ample literature presently available, advocates on the merits of medicinal plants extracts and phytochemicals to alleviate cancer. A few of the reported plant extracts and phytochemicals 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. 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.

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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, Surflex 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

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Saqib, et al.: Photochemicals with Apoptotic Properties Targeting Cancer

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PICTORIAL ABSTRACT

SUMMARY

- Phytochemicals with a capacity to activate caspases enhancing apoptotic capacity has been proven to be effective anticancer agents.
- Data on photochemicals 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.

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