Introduction

Cupressaceae is the largest conifer family with worldwide distribution. It consists of 32 genera, and the members of this family are highly valued for their timber, resin, fruit, and ornamentals [1, 2]. The genus Calocedrus belongs to the cypress family and is native to eastern Asia and North America. Calocedrus mainly includes 4 species viz., Calocedrus decurrens, Calocedrus macrolepis, Calocedrus formosana, and Calocedrus rupestris [3, 4]. These evergreen coniferous trees were primarily cultivated for ornamental purposes or lumber industry. Conifers are also renowned for their ability to produce an enormous array of bioactive molecules, leading to the discovery of new drugs. Calocedrus macrolepis and C. formosana are well-studied species in this genus and are reported to possess antioxidant, antiradical, antifungal, antimicrobial, anticancer, and immuno-activating properties [5–10]. The chemical constituents of these plants include lignans, sesquiterpenes, diterpenoids, terpenoids, abietane-O-abietane type dimers, labdane diterpenoids, and phaeophorbides [5, 8, 11–13]. Calocedrus rupestris is only recorded in Vietnam and is considered a rare and endemic species [4].
Calocedrus decurrens (Torr.) Florin, commonly known as the "Incense cedar tree," is indigenous to the United States’ Pacific region, such as Oregon and California [14]. The essential oil of cedarwood is highly aromatic and contains phenolic terpenes as major constituents. The heartwood was reported to contain carvacrol, thymoquinone, p-methoxythymol, p-methoxycarvacrol, libocedrol, hedyeriol, calocedrol A, and calocedrol B [15–17]. Von Rudloff identified the main constituents of leaf essential oil as limonene (31.3%), Δ-3-carene (21.0%), α-pinene (9.2%), myrcene (8.0%), and α-terpenyl acetate (5.7%) [18]. Velthour et al. reported that the essential oil of the heartwood contains thymoquinone (35.9%), carvacrol (29.2%), p-methoxythymol (11.0%), and p-methoxycarvacrol (3.2%) as the major constituents [19]. Garcia et al. reported 4 unusual pinane derivatives: pin-2-en-8-al, pin-2-en-8-ol, pin-2-en-8-yl acetate, and methyl pin-2-en-8-oate from the essential oil in young branches of C. decurrens[20]. The extracts and essential oil of the heartwood exhibited antifungal, antimicrobial, and biocidal activity, which supports the decay-resistant nature of the wood [21–23]. Native Americans used the decoction of cedar leaves for stomach trouble, as a vapor infusion for colds and as a food spice [24]. The wood’s durable and decay-resistant nature made it unique in the lumber industry, especially for making pencils and general building purposes. From the literature review, only a few reports were available regarding the phytochemistry and pharmacology of the species C. decurrens. Our study mainly focused on the phytochemical investigation of the heartwood to identify its constituents, with particular emphasis on the anticancer property of the extracts and compounds. To the best of our knowledge, this is the first report on the anticancer activity of C. decurrens.

Results and Discussion

About 1 kg of the heartwood of C. decurrens was extracted with hexane, acetone and ethanol. The extracts were assessed for their cytotoxicity against human lung adenocarcinoma (A549) cell lines using MTT assay. Doxorubicin was used as the standard, and the results are shown in ▶ Table 1. The effect of various concentrations of the extracts is depicted in ▶ Fig. 1. The hexane extract showed excellent activity with an IC50 value < 5 µg/mL, which was comparable with the control doxorubicin at 24 h (IC50 1.63 µg/mL) as well as 48 h (IC50 1.18 µg/mL). The acetone extract exhibited good activity with an IC50 11.26 µg/mL, while the ethanol extract was least toxic with an IC50 value > 100 µg/mL at 24 h. Since hexane and acetone extracts demonstrated good inhibitory activity, further phytochemical investigation was focused on these extracts. Hence, the present work describes the successive isolation and purification of phytoconstituents from the hexane and acetone extracts of the heartwood of C. decurrens and their antiproliferative study toward human lung adenocarcinoma (A549) and normal lung fibroblast (WI-38).

Phytochemical investigation of the heartwood of C. decurrens resulted in the isolation of 5 compounds viz: libocedrol (1) [17], thymoquinone (2) [25], libocedroquinone (3) [17], diethylphthalate (4) [26], and (1 R, 2 R, 4 R)-p-methane-1,2,4-triol (5) [27]. The structures of the compounds (▶ Fig. 2) was established by extensive spectroscopic analysis and on comparison of their spectral data with those in the literature (see Supporting Information). Libocedroqui-

none was isolated as a reddish crystalline solid, and the structure was further supported with a single-crystal XRD (▶ Fig. 3). Crystal data for 3 (CCDC 2086733): C21H26O4, M= 342.42, Triclinic, space group P-1, a = 8.666(5) Å, b = 9.634(6) (3) Å, c = 13.266(8) Å, α = 108.260(1)°, β = 90.060(4)°, γ = 110.285(10)°, cell formula units Z = 2, crystal density = 1.162 mg/m3, T = 302(2), 3503 reflections collected, 1505 reflections with I > 2σ(I) R factor = 0.0872. The single crystal X-ray structure of this compound is being reported for the first time. Compounds 1–3 were previously reported by Zavarin et al. from this species [14–16]. This is the first report on the presence of compounds 4 and 5 in the Calocedrus genus. (1 R, 2 R, 4 R)-p-methane-1,2,4-triol was previously reported from Artemisia suksdorfii (Asteraceae family) and Daucus carota (Apiaceae family) [27, 28].

There are no reports available regarding the cytotoxicity of the 2 compounds, libocedrol and libocedroquinone. Therefore, compounds 1–3 were examined for their growth inhibitory properties against A549 and WI-38 cell lines. The standard drug used was doxorubicin (DOX), with similar concentrations as the compounds. Treatment of each compound resulted in a concentration-dependent reduction of cell viability. The cytotoxicity results in terms of IC50 values are depicted in ▶ Table 2, and the effect of various concentrations is represented in ▶ Fig. 4. Compounds 1 and 3 showed an IC50 value of 7 and 4.6 µM for 24 h and 7.2 and 3 µM for 48 h, respectively. Compound 2 did not produce any IC50 effect (> 20 µM) in 24 h of incubation, whereas an IC50 value of 16.5 µM was obtained in 48 h. Doxorubicin showed IC50 at 1.8 µM in 24 h and 0.8 µM in 48 h. Compounds 1 and 3 exhibited similar cytotoxic effects of doxorubicin upon 48 h of treatment (p < 0.05, non-significant when compared to doxorubicin, compound 1 and 3). Whereas in terms of IC50 value, compound 3 is found to be more efficient when compared to other compounds. Upon 24 h treatment, the IC50 value of compound 3 in the A549 cell line was 4.6 ± 0.9 µM, whereas, for WI-38, the IC50 value was not achieved even at a concentration of 20 µM, emphasizing its minimal cytotoxic effects in normal cells. Even though these compounds are not as effective as doxorubicin against the human lung adenocarcinoma cell line, they exhibited low toxicity against the normal cell line, which was comparable with doxorubicin (16.5 µM) at 24 h.

Out of the selected compounds, compound 3 exhibited appreciable apoptotic effects toward the lung adenocarcinoma cells (A549) and was found to be least toxic to normal lung fibroblast cells (WI-38) at the specified concentration. The apoptotic ability of compound 3 was further demonstrated by using ethidium bromide-acridine orange live dead assay. In the dual staining proce-

<table>
<thead>
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<th>Table 1 Cytotoxicity data for different extracts in A549 over a time of 24 and 48 h.</th>
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<tr>
<td><strong>Extracts</strong></td>
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<tr>
<td>Hexane</td>
</tr>
<tr>
<td>Acetone</td>
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<tr>
<td>Ethanol</td>
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<tr>
<td>Doxorubicin*</td>
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* Standard drug; b statistical significance: doxorubicin vs. extracts p < 0.001; c IC50 not achieved. A549: human lung adenocarcinoma.
dure, ethidium bromide is impermeant to the cell membrane and only enters the dead cells with damaged cell membrane; it indicates red fluorescence upon intercalation with DNA. Acriflavin orange is a nonfluorescent membrane-permeable dye that becomes fluorescent upon conversion with the presence of cellular esterase enzymes, causing the live cells to appear green in color. Fluorescent images showed a reduced number of cells in compound 3 treated group in the live dead assay, with a high proportion of cells exhibiting yellow/red fluorescence compared with the untreated control cells (Fig. 5). Thus, the results demonstrated the apoptotic-inducing ability of compound 3 even at 4.6µM concentration.

As per our study, libocedrol and libocedroquinone exhibited significant anticancer potency compared to compound 2 (i.e., thymoquinone [TQ]). TQ is the main constituent in black cumin (Nigella sativa, Ranunculaceae family) seed oil and is well known for its anticancer properties. TQ exerts excellent anticancer effects on various cancer cell lines, including pancreatic, uterine sarcoma, cervical, colon, lung, prostate, breast, myeloblastic leukemia, etc.[29, 30]. Previous reports revealed that TQ stimulated apoptosis in A549 lung cancer cells via p53 and caspase cascade-dependent pathway activation [31, 32]. Yang et al. have shown that TQ significantly inhibited proliferation, migration, and invasion of A549 cells in a time- and dose-dependent manner by suppressing the ERK1/2 pathway in A549 cells [33]. Banerjee et al. reported that the synthetic analogs of TQ exhibited more potent anticancer potential than the parent TQ in terms of inhibition of cell growth and induction of apoptosis [34]. Considering the biosynthetic pathway, libocedroquinone is derived from the oxidative coupling of thymoquinone and p-methoxythymol [22]. As a structural derivative of thymoquinone, it exhibits excellent activity than thymoquinone toward cancer cells and is less toxic to normal cells. Our findings suggested that libocedroquinone could be developed as an effective chemotherapeutic agent for lung cancer. More studies are needed to uncover the exact molecular targets and mechanism of the action of libocedroquinone.

In summary, the current study was designed to evaluate the anticancer potency of various extracts and isolated compounds from the heartwood of Calocedrus decurrens. Five compounds were isolated from the heartwood extracts of C. decurrens, and compounds 4 & 5 were reported for the first time from this genus. In addition, the in vitro cytotoxic analysis of compounds 1–3 in human lung adenocarcinoma and normal lung fibroblast cell lines was conducted. Even though not up to the standard drug, libocedroquinone (3) showed satisfactory antiproliferative activity with minimal toxicity toward normal cells at 24 h incubation. To the best of our knowledge, this is the first anticancer study of the species C. decurrens. Our results suggested that libocedroquinone could be a potential candidate as an anticancer agent against lung cancer, and further investigations are necessary to validate its activity.
Materials and Methods

General experimental procedure

The NMR spectra were acquired with Bruker Avance II AMX 500 MHz spectrometer; NMR solvents used were CDCl3 and acetone-d6 with 0.03 % tetramethylsilane (TMS) as the internal standard. Mass spectra were measured using Thermo Scientific Exactive mass spectrometer under ESI/HRMS at 60,000 resolutions. Silica gel (230 – 400 mesh and 100–200 mesh) and Sephadex LH-20 were used for column chromatography. All the chemicals and solvents used were purchased from Sigma Aldrich and Merck. The diffraction data of single crystals were collected on a Rigaku Saturn 724+ diffractometer using graphite monochromated Mo-Kα radiation.

Plant material

Incense cedar (Calocedrus decurrens) trees were identified and collected by Dr. Richard Halse, OSU herbarium director, from a logging and sawmill operations near Warm Springs, Jefferson, Oregon, USA. A voucher specimen was deposited in the Oregon State University Herbarium (Voucher number OSC228848). The heartwood was removed and shavings made and stored in vacuum-sealed containers at −20 °C.

Table 2  IC50 Values of different compounds in A549 and WI-38 cell lines at 24 h and 48 h. Average of more than three independent experiments.

<table>
<thead>
<tr>
<th>Compounds</th>
<th>A549</th>
<th>WI-38</th>
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<tr>
<td></td>
<td>24 h</td>
<td>48 h</td>
</tr>
<tr>
<td>1</td>
<td>7 ± 1.5b</td>
<td>7.2 ± 0.3b</td>
</tr>
<tr>
<td>2</td>
<td>&gt; 20b</td>
<td>16.5 ± 1.2b</td>
</tr>
<tr>
<td>3</td>
<td>4.6 ± 0.9b</td>
<td>3 ± 0.2b, d</td>
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<tr>
<td>Doxorubicina</td>
<td>1.8 ± 0.2</td>
<td>0.8 ± 0.1</td>
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a Standard drug; b statistical significance: doxorubicin vs. compounds p < 0.001; c IC50 not achieved; d compound 1 vs. compound 3 p < 0.001; A549: human lung adenocarcinoma; WI-38: normal lung fibroblast.

Extraction and isolation

About 1 kg of the heartwood shavings of the plant material was soaked in 5 L hexane for 3 days at room temperature. The supernatant liquid was decanted and filtered. The procedure was repeated three times, giving 11 g of the crude hexane extract. The residue obtained after extraction was further extracted with acetone and ethanol, which yielded 28 g of acetone and 17 g of ethanol extract. All the crude extracts were evaluated for cytotoxicity using an MTT assay. Further, the crude hexane extract was then fractionated on a silica gel (100–200 mesh) column and eluted with ethyl acetate in hexane gradient, increasing the amount of ethyl acetate to rise polarity. The elution was started with hexane and gradually increased the polarity (between 5 to 30 %), finally with 30 % ethyl acetate/hexane. A total of 25 fractions of approximately 150 mL each were collected. Fractions collected were pooled into 3 fraction pools after analyzing the TLC. Fraction pool F2 (3.04 g) showed highly UV active spots. These fractions were purified successively with silica gel column chromatography in 5 % ethyl acetate/hexane gradient to afford compound 1 (500 mg, purity > 95 %, from NMR) and 2 (563 mg, purity > 95 %, from NMR). Another subfraction (100 mg) of F2 was rechromatographed on silica gel (230–400 mesh), a reddish compound with some minor impurities was obtained. It was further purified with Sephadex LH-20 in methanol yielded compound 3 (27 mg, purity > 95 %, from NMR). Next, we carried out the column chromatographic separation of the acetone extract. About 28 g of the acetone extract was subjected to silica gel CC (100–200 mesh) afforded two more compounds along with compounds 1 and 2. The elution was started with hexane, and an increase in polarity was carried out by increasing the amount of ethyl acetate (between 5 to 100 %). Final elution was carried out with ethyl acetate. As in the previous procedure, a total of 52 fractions of approximately 250 mL each were collected. From their similarity in TLC, collected fractions were combined into 5 fraction pools (F1–F5). Compound 4 was obtained from the fraction pool F2 as a colorless oily substance (17 mg) by eluting the column with...
5% ethyl acetate-hexane polarity. Compound 5 isolated from fraction pool F4 as a white crystalline solid (30 mg, 90% EtOAc-hexane). The chemical structures of the compounds were characterized using various spectroscopic techniques.

**Cell lines and cultural conditions**

A549 (human lung adenocarcinoma) and WI-38 (human normal lung fibroblast) were obtained from National Centre for Cell Science (NCCS), Pune, India, after short tandem repeat (STR) profiling. The cell lines were cultured in DMEM supplemented with 10% heat-inactivated FBS and 1% antibiotics antifungal solution (10,000 U Penicillin, 10 mg streptomycin and 25 µg amphotericin B per mL in 0.9% normal saline) in a humidified CO₂ incubator at 37°C with 5% CO₂.

**Cytotoxicity assay**

The cytotoxicity effects of the extracts and the isolated compounds from Calocedrus decurrens were evaluated using the MTT assay [35]. Doxorubicin hydrochloride was used as the standard (D1515, Sigma-Aldrich, 98.0–102.0%, HPLC purified). Mitochondrial dehydrogenase enzyme in the live cell will convert the tetrazolium salt into soluble formazan crystals whose absorbance can be measured at 570 nm. The cells were seeded at a density of 10000 cells per well in 100 µL volume with complete DMEM medium for 24 h for cells to adhere. Cells were then treated with different compounds having concentrations ranging from 0.5 to 20 µM followed by further incubation for 24 h and 48 h separately. 100 µL MTT solution (0.5 mg/mL in HBSS) was added to each well and incubated for 3 h. The solution was subsequently removed from wells, and resulting formazan crystals were solubilized in 100 µL DMSO. Culture plates were rocked gently for 20 min to solubilize the crystals, and optical density was recorded spectrophotometrically using a microplate reader (BioTek) at 570 nm. The relative cell viability in percentage was calculated using the formula: \( \text{Absorbance of treated/Absorbance of control} \times 100 \). As a result of cytotoxicity, the inhibition rate is calculated as Inhibition % = 100 − Viability %.

Also, concentrations providing inhibition of 50% on cell growth (IC₅₀) were calculated.

**Live dead assay to determine apoptotic cells**

The most effective compound with the minimal IC₅₀ value at 24 h from the MTT assay was carefully chosen for further screening. Staining with acridine orange and ethidium bromide dual dye is one of the most frequently used method for detecting apoptotic cells, where the varying uptake of 2 dyes by live and dead cells are observed [36]. In order to perform the assay, A549 cells were cultured at a seeding density of 5000 cells/well and treated with and without compound 3. A dual-staining dye cocktail was prepared by adding 1 µL each acridine orange and ethidium bromide from 5 mg/mL and 3 mg/mL stock, respectively, to 1 mL PBS buffer. One hundred µL of the as-prepared reagent was added to the compound 3 treated and untreated control cells. After incubation for 5 min, the cells were then washed thrice with PBS buffer and observed under a fluorescence microscope (Nikon Eclipse TS100, Japan).

**Statistical analysis**

All data were acquired from 3 independent experiments. The results were provided as mean ± standard deviation. The statistical analysis was performed by the software GraphPad Instat 2. Concentrations inducing a 50% inhibition of cell growth (IC₅₀) were determined graphically using Easy Plot software for each experiment.

**Supporting Information**

The 1H and 13C NMR spectra of compounds 1–5 are available in the supporting information.

**Acknowledgments**

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Conflicts of Interest

The authors declare that they have no conflict of interest.

References


