Cytotoxic Activity of Some Medicinal Plants from Hamedan District of Iran

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Abstract

Medicinal plants have been investigated for possible anti-cancer effects. The aim of the present study was to examine the cytotoxic activity of several medicinal plants on different tumor cell lines. 11 selected plant species which have been used in folkloric prescriptions were collected from different sites of Hamedan district of Iran. The methanolic extracts of the plants were prepared and their cytotoxic effects on four human cancer cell lines (A549, human lung adenocarcinoma; MCF7, human breast adenocarcinoma; HepG2, hepatocellular carcinoma and HT-29, human colon carcinoma) and one normal cell line (MDBK, bovine kidney) were examined using the MTT assay. Three of these were exhibited antiproliferative activity against one or more of the cell lines. The extract from \textit{Primula auriculata} demonstrated the highest cytotoxicity with IC\textsubscript{50} of 25.79, 35.79 and 43.34 \textmu g.mL\textsuperscript{-1} against MCF7, HepG2 and HT-29 cells, respectively. For some of the plants, their traditional use was correlated with the cytotoxic results, whereas for others the results may support the non-cytotoxicity of species used traditionally as natural remedies. The cytotoxic species could be considered as potential of anticancer compounds.

Keywords: Cytotoxic activity; MTT assay; Iranian medicinal plants; Hamedan.

Introduction

Cancer is one of the main causes of death all over the world. The world health organization (WHO) estimates that 84 million people would die of cancer between 2005 and 2015 (1). Accordingly, much effort has been made to develop various approaches to reduce the threat caused by cancer. Chemotherapy is an important option in modern cancer treatment, and many clinically available anticancer drugs are currently used to treat some types of leukemia, lymphoma and solid tumors (2).

The introduction of active agents derived from nature into the cancer armamentarium has changed the natural history of many types of human cancer (3, 4). Statistics indicated that a half part of anticancer drugs approved internationally between 1940(s) and 2006 was either natural products or their derivatives (5).

Iran’s unique meteorological conditions have contributed to the diversity of more than 8000 plant species (6), for this reason many botanists believe the flora of Iran is a green gold (7). Traditional records and ecological diversity indicate that Iranian plants represent an exciting resource for possible lead structures in drug design (6). Local communities in different parts of the country have developed a deep knowledge of various uses of plants during their old history (8). Hamedan district with a long medical
Preparation of extracts

The aerial part of each plant (100 g) was separated, shade dried and ground into powder using a mortar and pestle at room temperature. Then extracted by maceration with methanol for 72 h. The supernatants were filtered and evaporated under vacuum by means of a rotary evaporator to obtain crude methanolic extracts.

Cytotoxic assay

Cell lines and culture medium

The following cancer cell lines were used for this study: A549 (human lung adenocarcinoma), MCF7 (human breast adenocarcinoma), MDBK (bovine kidney cells), HepG2 (hepatocellular carcinoma) and HT-29 (human colon carcinoma). Cells were obtained from National Cell Bank of Iran (Pasteur Institute, Tehran, Iran). MCF7 and HT-29 were cultured in Dulbecco’s modified eagle medium (DMEM; Gibco) with respectively 5% and 10% bovine serum (FBS; Gibco) while other three cell lines were cultured in RPMI 1640 medium (Sigma) with 10% FBS to maintain the desired growth. All cell lines were treated with 1% penicillin-streptomycin (Sigma) in a humidified atmosphere with 5% CO$_2$ at 37 °C throughout the assay.

MTT assay

Cell viability was quantified by an MTT colorimetric assay (3-[4, 5-dimethylthiazol-2-yl]-2,5 diphenyl tetrazolium bromide assay) (11). The cells were seeded in 96-well plates at

| Species             | Family     | Local name | parts Traditionally used       | Voucher number |
|---------------------|------------|------------|--------------------------------|---------------|
| *Alhagi camelorum* | Fabaceae   | Taranjabin | Whole plant parts              | 3258 (TMRC)   |
| *Centaurea aucheri*| Astraecae  | Gole gandom | -                              | 3235 (TMRC)   |
| *Centaurea pseudoscabiosa* | Astraecae | Gole gandom | Aerial parts                    | 3233 (TMRC) |
| *Cerasus microcarpa* | Rosaceae   | Albalooye valshi | Fruits, flowers, core, wood | 3259 (TMRC) |
| *Primula auriculata* | Primulaceae | Tootia     | Flower                         | 3224 (TMRC)   |
| *Silene amputa*     | Caryophyllaceae | silene     | Aerial parts                   | 3238 (TMRC)   |
| *Silene peduncularis* | Caryophyllaceae | silene     | Aerial parts                   | 3229 (TMRC)   |
| *Smyrniposis aucheri* | Apiaceae  | peakal     | -                              | 2261 (TMRC)   |
| *Stachys lavandifolia* | Lamiaceae | Toklijeh   | Leaves, and flowers            | 2835 (TMRC)   |
| *Thymus pubescens*  | Lamiaceae  | Azarbeh    | Aerial parts                   | 1593 (TMRC)   |
| *Tripleurospermum disforme* | Astraceae | Babooneh   | Aerial parts                   | 3245 (TMRC)   |

Table 1. Selected medicinal plants of Hamedan district, Iran.
8.5× 10^3 for MDBK cells, 7.5 × 10^3 for MCF7, 15 × 10^3 for HepG2, 9 × 10^3 for A549 cells, and 5 × 10^3 for HT-29 cells and incubated at 37 °C. After 24 h of incubation, when cells reached more than 80% confluence, the medium was removed and the cells were treated with fresh medium containing various concentrations of plant extracts to be tested. Control cells were supplemented with 0.05% DMSO (v/v) vehicle. After 24 h incubation, the supernatants were removed and fresh medium containing MTT (0.5 mg mL^{-1}) was added to each well at the time of incubation. After 4 h incubation, the supplement was carefully removed, and the remaining formazan crystals were dissolved in DMSO. The plates were shaken for 20 min. The absorbance of each well was measured on an enzyme-linked immunosorbent assay reader (TECAN) at the wavelength of 570 nm.

The dose–response curves of the compounds were fitted by means of the computer program GraphPad Prism 6.0 (GraphPad Software, USA), and IC50 values (the concentration at which the cell proliferation is 50% of the untreated control) were calculated. All in-vitro experiments were carried out on two microplates with at least three parallel wells. The antitumor agent 5-FU was used as a positive control in all cell lines.

Results

In order to evaluate the cytotoxic effect of 11 plant extracts that are used in Hamedan district of Iran, an antiproliferative assay on four human cancer cell lines (A549, human lung adenocarcinoma; MCF7, human breast adenocarcinoma; HepG2, hepatocellular carcinoma and HT-29, human colon carcinoma) and one normal cell line (MDBK, bovine kidney) was performed. Table 2 presents the list of the investigated plants with traditional uses, chemical constituent and biological activities.

Cytotoxicity activity (IC_{50}) of the eleven plant extracts was shown in Table 3.

Extracts with IC_{50} > 100 µg.mL^{-1} in MTT assay were considered inactive.

Discussion

Our study describes investigations into the anticancer potential of 11 so far not studied Iranian medicinal plants by screening for cytotoxic activity against normal bovine kidney and four human cancer cell lines. All plant extracts showed no toxicity against normal bovine kidney, but methanolic extract of Centaurea aucheri, Centaurea pseudoscabiosa subsp pseudoscabiosa and Primula auriculata was expected to show more cytotoxic activity against the tumor cells, whereas others were not cytotoxic against any of the cell lines tested.

The large genus Centaurea comprises about 500 species, which are predominantly distributed around the Mediterranean area and in west Asia (22). In Iran this genus has 74 annual to perennial herbaceous species that are widespread around the country (62). Several species of the genus Centaurea are well known for their traditional medicinal uses for the treatment of a number of ailments including bacterial infections, cancers, diabetes, diarrhea, fever, hypertension, malaria, rheumatism and tumors (63). Many reports showing the existence of various cytotoxic compounds in different Centaurea species, including alkaloids, flavonoids, lignans, sesquiterpenes and simple phenolics (64-66).

Methanol extract from Centaurea aucheri displayed selective cancer cell line cytotoxicity with IC50 values of 53.31 µg.mL^{-1} against hepatocellular carcinoma. This plant has not previously been used as anticancer treatment in traditional Iranian medicine and was selected because of cytotoxic effect of this genus. Some flavonoids and their glycoside isolated from Centaurea pseudoscabiosa subsp pseudoscabiosa such as chrysin, hispidulin and luteolin (24) these compounds have been shown Significant cytotoxic and apoptotic effects of on various cancer cell lines (MCF7, Hela, HL-60 and KYSE-510)(67-69) and it may be involved in cytotoxic effect of methanolic extract of this species in this study. Interestingly, the plant has been used traditionally in skin ailments (23), however, no anticancer or cytotoxic activities have been reported to date. Methanol extract of Primula auriculata showed significant cytotoxic activity against breast, liver and colon cancer cell lines with IC50 values ranging from 25.79 to 43.34 µg.mL^{-1}.
### Table 2. Traditional uses, chemical constituents and biological activities of medicinal plants from Hamedan district, Iran.

| Species                | Biological activities                                                                 | Previously isolated compounds                                                                 | Traditional uses                                                                 |
|------------------------|---------------------------------------------------------------------------------------|---------------------------------------------------------------------------------------------|-----------------------------------------------------------------------------------|
| *Alhagi camelorum*     | Antidiarrheal (12), antinoceptive (13), antitulcerogenic (14), gastro protective (15), ureteral stone expulsion (16) | Kampferol, chrysoeriol, isohamnetin, chrysoeriol-7-o-xyloside, kaempferol-3-galacto rhamnoside, isohamnetin 3-o-B-D-apio-faranonosyl (1-2) B-D-galactopyranoside (14) alhagitan, alhagitan (17), ferulic acid, isohamnetin, 5-hydroxymalol (18), ß-phenethyllamine, N-methyl-ß-phenethyllamine, N-methyl-tyramine, hordenine, 3,4-dihydroxy-ß-phenethyltrimethylammonium hydroxide, 3-methoxy-4-hydroxy-ß-phenethyltrimethyl ammonium hydroxide, N-methyl mescaline, solsolidine (19) | Gastrointestinal disorders (8, 9), diuretic, wound healing, fever with rash, rheumatic pain (20) |
| *Centaurea aucheri*    | Antioxidant (21)                                                                      | Carnosine, cyclophellandrene, germacrine-D (22)                                                |                                                                                   |
| *Centaurea pseudoscabiosa* | Antibacterial (23)                                                                 | Chrysin, baicaline 6-methyl ether, protocatechuic acid, 5-cafeoyl quininc acid, hispidulin, chrysin 7-O-glucuronide, chrysin 7-O-glucuronide methyl ester, chrysin 6-C-glucoside, chrysin 8-C-glucoside, luteolin 7-glucoside, pinocembrin 7-O-a-arabinopyranosyl (1-2) B-gluopyranoside, chrysin 7-O-ß-galactopyranuronoside, baicaline 6-methyl-ß-7-O-ß-galactopyranuronoside, scopoletin (24) germacrene D, ß-caryophyllene, biocyclogermacrene, ß-sesquiphellandrene (25) | Skin ailments (23) |
| *Cerasus microcarpa*   | -                                                                                     | Alkaloid, tannin (26)                                                                         | Carminative, cure for pains of digestive system(8)                                 |
| *Primula auriculata*  | Antibacterial, antioxidant (27)                                                       | Saponin, flavonoid (27)                                                                      | Flu and sneezing (28), eye diseases, anti-infection, cataract, trachoma (29)       |
| *Silene ampulata*     | -                                                                                     | -                                                                                           | Insect repellent(8)                                                               |
| *Silene peduncularis* | -                                                                                     | -                                                                                           | Insect repellent(8)                                                               |
| *Smyrnigopsis aucheri*| antibacterial, antifungal (30)                                                        | ß-bisabolol (31), peyone, carophyllene oxide, spathulenol (32), smyrnilol, smyrnilol, smyrniloloside (33), smyrnilolidin (34), ß-pinene, ß-pirane, nachmyrin (30) | -                                                                                 |
| *Stachys lavandifolia*| Gastroprotective (35), wound healing (36), analgesic and antiinflammation (37), antiinflammatory (38), antinfectious (39), abortive effect (40) | ß-thujene, ß-pinene, ß-myrcene, ß-phellandrene, germacrene-D, cadinane, 1,4-methano-1 H-indene, γ-elemene benzaldehyde (41), apigenin, luteolin (42), lavandulifolioside A, lavandulifolioside B, verbascoside, leucosceptoside A, 5-O-ß-allopyranosyl-oxycucin (43) | Skin infection, monorragia, antibacterial (44, 45), gastrointestinal and respiratory disorder (46-48), wound healing, cardiac disorders, fever and malaria (49) |
| *Thymus pubescens*    | Antioxidant (50), antibacterial (51) analgesic and anti-inflammatory (52)             | Carvacoer, thymol, γ-terpinene, ß-cymene (53)                                                | Gastrointestinal disorder (54), herpes, lung infection and skin problem (55)       |
| *Tripleurospermum disforme* | Anti-ulcer (56), antibacterial (57), anti-inflammatory, analgesic (58), antinfectious (59), antifungal (60) | Flavonoid (61), ß-farcsene, ß-sesquiphellandrene, ß-methoxy-ß-cyclopropylstyrene, heptadecane, ß-methoxy-humulene oxide and benzene acetaldehyde (57) | Antispasmodic, antimflammatory, Acne and itching (61)                                 |

is one of the most important local medicinal plants in Hamedan district (locally named Tootia). White powders that were produced by plant inflorescences named Tootia have been used traditionally for eye infectious diseases (29). In turkey dried herb was sniffed into nose for sneezing to ease respiration in flu (28). The aerial part of this genus are rich in flavonoid (70) and they may be related to cytotoxic effect of *Primula auriculata* methanolic extract.

8 of 11 selected plants showed no cytotoxic activity against normal and cancer cell lines that
has a great significance for their traditional use in the treatment of various disorders other than cancer.

This is the first time that methanolic extracts from the 11 listed Iranian plants (*Alhagi camelorum*, *Centaurea aucheri*, *Centaurea pseudoscabiosa* subsp *pseudoscabiosa*, *Cerasus microcarpa*, *Primula auriculata*, *Silene ampulata*, *Silen peduncularis*, *Smyrniopsis aucheri*, *Stachys lavandifolia*, *Thymus pubescens* and *Tripleurospermum disiforme*) have been screened against human lung, liver, colon and breast cancer cell lines and one normal cell line. This study provides an important basis for further investigation into the isolation, characterization and mechanism of cytotoxic compounds from some of the screened Iranian medicinal plants. Thus, these plants could be used as a source for new lead structures in drug design to combat cancer.

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| Species                      | Yields (%) | A549   | MCF7 | HepG2 | HT-29 | MDBC |
|------------------------------|------------|--------|------|-------|-------|------|
| *Alhagi camelorum*           | 11.3       | >100   | >100 | >100  | >100  | >100 |
| *Centaurea aucheri*          | 9.6        | >100   | >100 | 53.31 | >100  | >100 |
| *Centaurea pseudoscabiosa*   | 11.77      | 54.82  | >100 | >100  | 98.15 | >100 |
| *Cerasus microcarpa*         | 10.86      | >100   | >100 | >100  | >100  | >100 |
| *Primula auriculata*         | 12         | >100   | 25.79| 35.79 | 43.34 | >100 |
| *Silene ampulata*            | 7.24       | >100   | >100 | >100  | >100  | >100 |
| *Silene peduncularis*        | 9.7        | >100   | >100 | >100  | >100  | >100 |
| *Smyrniopsis aucheri*        | 19.03      | >100   | >100 | >100  | >100  | >100 |
| *Stachys lavandifolia*       | 15.53      | >100   | >100 | >100  | >100  | >100 |
| *Thymus pubescens*           | 8.6        | >100   | >100 | >100  | >100  | >100 |
| *Tripleurospermum disiforme* | 11.25      | >100   | >100 | >100  | >100  | >100 |
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