Cytotoxic Evaluation of *Daphne pontica* L. Aerial Part Extracts on Three Cancerous Cell Lines by MTT Assay

Behshad Eskandari¹, Maliheh Safavi², Seyede Nargess Sadati Lamardi³, Mahdi Vazirian¹,⁴ *

¹Department of Pharmacognosy, Faculty of Pharmacy, Tehran University of Medical Sciences, Tehran, Iran  
²Department of Biotechnology, Iranian Research Organization for Science and Technology (IROST), Tehran, Iran  
³Department of Traditional Pharmacy, School of Persian Medicine, Tehran University of Medical Sciences, Tehran, Iran  
⁴Medicinal Plants Research Center, Tehran University of Medical Sciences, Tehran, Iran

Received: 3 May 2019  
Revised: 12 May 2019  
Accepted: 14 May 2019

Abstract

Nowadays, cancer is the second prevalent cause of mortality after cardiovascular diseases in developed and the third one in developing countries. Adverse effects of chemotherapeutic agents bring the necessity of investigating about new medications with fewer side effects. *Daphne* L. genus is one of the natural sources with valuable reported anticancer effects. This study aimed to assess the cytotoxic effect of some extracts from the aerial parts of *Daphne pontica* collected from North of Iran on cancer cell lines. Extraction of the plant material was performed by maceration (3×72 h) of 200 g of sample with petroleum ether, chloroform, ethyl acetate, and methanol, respectively. The total extract was also obtained by maceration of the sample with 80% ethanol. Different concentrations of the dried extracts were prepared to assess their cytotoxic effect by 24 h incubation of cell lines with different extracts and then MTT (dimethyl thiazolyl diphenyl tetrazolium) assay on three cancerous cell lines (MDA-MB-231, MCF-7 and T47D), performed in triplicate. IC₅₀ was then estimated from curves constructed by plotting cell survival (%) versus sample concentration (µg/ml). Results indicated that ethyl acetate fraction of *D. pontica* had the most potent cytotoxic effect in MTT assay with IC₅₀ = 977.46 µg/ml; while other fractions were weaker in toxicity (IC₅₀ >1000 µg/ml). By comparing to potent cytotoxic effects of other *Daphne* species, it seems that the cytotoxic properties of *D. pontica* is different from other species of this genus since according to this study, no significant antineoplastic properties against the three breast cancer cell lines were determined. Further studies on other pharmacological activities of this plant are recommended.

Keywords: Cytotoxic; MTT; *Daphne pontica*; Cancer cell line


*Corresponding Author: Mahdi Vazirian  
Department of Pharmacognosy, Faculty of Pharmacy, Tehran University of Medical Sciences, Tehran, Iran  
Tel/Fax: +98-2164121223  
Email: vazirian_m@sina.tums.ac.ir*
Cytotoxic evaluation of Daphne pontica L. B. Eskandari et al.

Introduction
Cancer is currently one of the most important challenges in medical sciences and despite all the advances made, no definitive treatment has been found. One of the approaches to find anticancer agents is the screening of plants to find natural compounds that have cytotoxic effects on human cancer cell lines, which has so far led to the discovery of remarkably effective therapeutic compounds in a variety of cancers including the alkaloids of Vinca rosea and Taxus brevifolia. Phytochemical studies of plants with a history of traditional or folk use for the treatment of cancer often results in the isolation of compounds with antitumor activity [1].

The genus Daphne belongs to the Thymelaeaceae family and has a history of therapeutic use in different countries. Various pharmacological effects have been reported from different species of this plant, such as analgesic and anti-inflammatory effects [2-4], wound healing properties [5], antimicrobial effects [6], and antioxidant effects [6,7].

Daphne pontica L., known with the local name of Bargebooei, is one of the Daphne species in Iran. The leaf infusion of D. pontica is used by native people in Mazandaran province to treat low back and abdominal pain, diarrhea, heartburn, and kidney infections [8]. In an in vivo study, anti-inflammatory and antinociceptive activity of ethyl acetate extract of the roots of D. pontica was reported [4]. Considering the cytotoxic effects of various Daphne species in different types of cancer cells, for example, D. gniidium and D. genkwa on lung cancer, and D. mucronata on breast and leukemia cell lines [9-18], the aim of this study was to investigate the cytotoxic effect of some extracts from the aerial part of D. pontica collected from North of Iran against T-47D, MDA-MB231, MCF-7 cancer cell lines.

Methods
Plant material
The aerial part (leave and stem) of the Daphne pontica was collected from the altitudes of Sari in Mazandaran province on June 2015 and a voucher specimen deposited in Herbarium Ministrii Iranici Agriculturae (Iran) (Code No. IRAN70100/1).

Extraction
The air-dried aerial parts of D. pontica (200 g) were powdered and extracted by maceration (3×72 h) with petroleum ether, chloroform, ethyl acetate and methanol, respectively. The total extract was also obtained by maceration of the plant with 80% ethanol. Removal of the solvents with a rotary evaporator resulted in the production of petroleum ether (PEE), chloroform (ChE), ethyl acetate (EAE) and methanol (MEE) extracts.

Cell lines and cell culture
Human breast cancer cell lines comprising MDA-MB231(ATCC® HTB-26), MCF-7 (ATCC® HTB-22) and T47D (ATCC® HTB-133) cells were purchased from the National Cell Bank of Iran, Pasteur Institute, Tehran, Iran.
Determination of cell viability by MTT assay
The in vitro cytotoxic activity of all samples was assessed against three human breast cancer cell lines by dimethyl thiazolyl diphenyl tetrazolium (MTT; Merck, Germany) colorimetric test (19). Cancer cell lines were grown in RPMI-1640 medium supplemented with 10% FBS (fetal bovine serum protein) (Gibco BRL), 100 μg/ml streptomycin and 100 U/ml penicillin (Roche, Germany). All cell lines were cultured at 37° C in air/carbon dioxide (95:5) atmosphere.

Samples were dissolved in DMSO (dimethyl sulfoxide) and were further diluted with cell culture medium. The final DMSO concentration used was 1% of total volume of medium in all treatments, including the control group. For the MTT assay, 5×10⁴ cells /wells were plated into 96-well plates (Nunc, Denmark) and incubated for 24 h before addition of the extracts. The plates were incubated overnight at 37°C with 5% CO₂. Then 5 μl of the media having various concentrations of the samples was added per well in triplicate. The plates were then incubated for 72 h. In each plate, there were three control wells (cells without test samples) and three blank wells (the medium with 0.1% DMSO) for cell viability. After treatment, the medium was removed and 200 μl phenol red-free medium containing MTT (1 mg/ml), was added to the wells, followed by 4 h incubation at 37°C. After incubation, the culture medium was then replaced with 100 μl of DMSO and the absorbance of each well was measured by using a microplate reader at 570 nm. The cytotoxicity value for each sample was presented as IC₅₀ (the median growth inhibitory concentration) of the reagents compared with the control. IC₅₀ values were calculated by Sigmaplot 9.0 software [20].

Results
The in vitro cytotoxic activity of total extract (TE) and other extracts including PEE, ChE, EAE, and MEE were assessed against three human breast cancer cell lines including MCF-7, T47D, and MDA-MB-231. The IC₅₀ for the samples are represented in Table 1-3.

Table 1. In vitro effect of Daphne pontica different extracts on viability (Mean ± SD) of MDA-MB 231 cell line.

<table>
<thead>
<tr>
<th>Samples a</th>
<th>(Concentration (µg/ml))</th>
<th>IC₅₀ (µg/ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>1000</td>
<td>100</td>
</tr>
<tr>
<td>PEE</td>
<td>107.19±11.19</td>
<td>82.19±7.75</td>
</tr>
<tr>
<td>ChE</td>
<td>59.69±12.03</td>
<td>75.59±10.00</td>
</tr>
<tr>
<td>EAE</td>
<td>48.85±7.43</td>
<td>87.64±14.14</td>
</tr>
<tr>
<td>MEE</td>
<td>111.21±4.68</td>
<td>88.51±15.05</td>
</tr>
<tr>
<td>TE</td>
<td>104.70±12.36</td>
<td>102.89±4.80</td>
</tr>
<tr>
<td>CON+DMSO</td>
<td>94.18±5.71</td>
<td></td>
</tr>
<tr>
<td>DMSO</td>
<td>3.83±0.16</td>
<td></td>
</tr>
</tbody>
</table>

a Samples; PEE Petroleum ether extract, ChE: Chloroform extract, EAE: Ethyl acetate extract, MEE: Methanol extract, TE: Total extract, CON+DMSO: Control+DMSO
Table 2. In vitro effect of Daphne pontica different extracts on viability (Mean±SD) of T47D cell line.

<table>
<thead>
<tr>
<th>Samples</th>
<th>(Concentration (µg/ml))</th>
<th>IC_{50} (µg/ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>1000</td>
<td>100</td>
</tr>
<tr>
<td>PEE</td>
<td>54.06±4.35</td>
<td>79.85±6.87</td>
</tr>
<tr>
<td>ChE</td>
<td>63.07±1.86</td>
<td>115.60±4.97</td>
</tr>
<tr>
<td>EAE</td>
<td>59.19±2.21</td>
<td>120.54±12.27</td>
</tr>
<tr>
<td>MEE</td>
<td>94.26±8.59</td>
<td>107.03±12.74</td>
</tr>
<tr>
<td>TE</td>
<td>84.61±3.61</td>
<td>93.69±5.82</td>
</tr>
<tr>
<td>CON+DMSO</td>
<td></td>
<td></td>
</tr>
<tr>
<td>DMSO</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Table 3. In vitro effect of Daphne pontica different extracts on viability (Mean±SD) of MCF-7 cell line.

<table>
<thead>
<tr>
<th>Samples</th>
<th>(Concentration (µg/ml))</th>
<th>IC_{50} (µg/ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>1000</td>
<td>100</td>
</tr>
<tr>
<td>PEE</td>
<td>70.41±6.17</td>
<td>113.88±4.49</td>
</tr>
<tr>
<td>ChE</td>
<td>81.85±1.53</td>
<td>129.67±23.22</td>
</tr>
<tr>
<td>EAE</td>
<td>77.58±0.12</td>
<td>171.23±18.60</td>
</tr>
<tr>
<td>MEE</td>
<td>120.32±18.47</td>
<td>135.69±7.12</td>
</tr>
<tr>
<td>TE</td>
<td>183.18±19.59</td>
<td>125.95±14.88</td>
</tr>
<tr>
<td>CON+DMSO</td>
<td></td>
<td></td>
</tr>
<tr>
<td>DMSO</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Discussion

According to the results, among all samples, EAE showed the highest cytotoxic effect on MDA-MB231 cell line with IC_{50} value 977.46 µg/ml and the remaining samples did not show significant cytotoxicity (IC_{50} > 1000 µg /ml). A review of previous studies showed that the compounds in ethyl acetate, hydroalcoholic, and chloroform extracts of other Daphne species had significant cytotoxic effects. Ethyl acetate extract of D. gnidiunm inhibited the growth of lung cancer cell lines (A549), liver cancer cell line (SMMC-7721) and breast cancer cell line (MCF-7) by inducing apopto-
sis after 48 hours [18]. Also, daphnane diterpenoids from *D. genkwa* have been shown to inhibit proliferation of the A549 lung cancer cell line [9]. In other studies, diterpenoids with cytotoxic effects were isolated from *D. genkwa*, *D. mucronata*, *D. mezereum* and *D. linearifolia*, such as yuanhuahine, yuanhuadine, yuanhuagine [9], gnidilatimonecin [12,13], mezerein [11], and bioflavonoids including 4-methylgenkwanol A, and 2-hydroxygenkwanol A [21]. Several compounds with diterpene ester and secolignans structure which had a moderate cytotoxic effect were isolated from the ethyl acetate fraction of *D. acutiloba* and *D. feddei* [10, 22]. The results of this study, which was performed on three breast cancer cell lines including T47D cell line which was evidently susceptible to progesterone [23], an estrogen-positive (MCF-7) and an estrogen-negative (MDA-MB-231) human breast cancer cell line [24], showed that the ethyl acetate extract had a poor effect on the MDA-MB-231.

**Conclusion**

Based on the results, the most potent effect of *D. pontica* ethyl acetate extract on MDA-MB231 cancer cell line was found, and the rest of the extracts did not show cytotoxicity on breast cancer cell lines. Future studies on pure compounds of the ethyl acetate extract as well as other cancer cell lines can be done to elucidate further biological effects of this plant.

**Conflicts of Interest**

The authors confirm that this article content has no conflicts of interest.

**Acknowledgments**

None.

**References**


[10] Huang SZ, Zhang XJ, Li XY, Kong LM, Jiang HZ, Ma QY. Daphnane-type diterpene esters with cytotoxic and
Cytotoxic evaluation of *Daphne pontica* L.  

B. Eskandari et al.


