

"HDIR-6: Targeting Cancer"

The 6th Meeting of the Croatian Association for Cancer Research with International Participation

November 10-12, 2022

Hotel International, Zagreb, Croatia

BOOK OF ABSTRACTS

Hrvatsko društvo za istraživanje raka (HDIR) Croatian Association for Cancer Research (CACR)

"HDIR-6: Targeting Cancer" The 6th Meeting of the Croatian Association for Cancer Research with International Participation – Book of Abstracts

Impressum: Editor: Petar Ozretić, HDIR Secretary

Publisher: Croatian Association for Cancer Research, Zagreb, Croatia ISBN 978-953-48672-1-1

URL: http://stari.hdir.hr/docs/HDIR6_BoA.pdf

Place and year of publication: Zagreb, November 2022

Copyright @ 2022, Croatian Association for Cancer Research. All rights reserved.

"HDIR-6: Targeting Cancer" The 6th Meeting of the Croatian Association for Cancer Research with International Participation

Organizer: Croatian Association for Cancer Research, Bijenička 54, HR-10000 Zagreb, Croatia

Venue: Hotel International, Miramarska cesta 24, HR-10000 Zagreb, Croatia

Dates: November 10-12, 2022

Scientific Committee:

Sonja Levanat, Andreja Ambriović Ristov, Paola Defilippi, Katja Ester, Maja Herak Bosnar, Dinko Leović, Vesna Musani, Petar Ozretić, Maja Sabol, Maja Sirotković-Skerlev, Neda Slade, Sandra Sobočanec, Ivan Šamija, Engin Ulukaya

Organizing Committee:

Sonja Levanat, Josipa Čonkaš, Anđela Horvat, Maja Jazvinšćak Jembrek, Matea Kurtović, Vesna Musani, Petar Ozretić, Tina Petrić, Nikolina Piteša, Bastien Proust, Maja Sabol, Iva Škrinjar, Ana Tadijan, Ignacija Vlašić

Supported by: European Association for Cancer Research (EACR), Foundation of the Croatian Academy of Sciences and Arts (HAZU)

Sponsors:

Alphachrom, Avantor / VWR, Biosistemi, Biovit, Diagnostica Skalpeli, Gorea plus, Jasika, KEFO, Labena, Medic, Vita Lab Nova



"HDIR-6: Targeting Cancer"

The 6th Meeting of the Croatian Association for Cancer Research with International Participation

November 10-12, 2022

Hotel International, Zagreb, Croatia

BOOK OF ABSTRACTS

Hrvatsko društvo za istraživanje raka (HDIR) Croatian Association for Cancer Research (CACR)

P10: Cytotoxic Effects of *Lavandula angustifolia* Mill. and *Laurus nobilis* L. Essential Oils on Human Cervical Adenocarcinoma Cells

<u>Ivana Pašić</u>¹, Ivana Srbljak Ćuk¹, <u>Nina Petrović</u>^{1,2}, Ivana Matić¹, Hurija Džudžević-Čančar³, Alema Dedić³, Amra Alispahić³, Emina Boškailo⁴, Tatjana Stanojković¹

¹Department of Experimental Oncology, Institute for Oncology and Radiology of Serbia, Belgrade, Serbia; ²Laboratory for Radiobiology and Molecular Genetics, "VINČA" Institute of Nuclear Sciences-National Institute of the Republic of Serbia, University of Belgrade, Serbia; ³Department of Chemistry in Pharmacy, Faculty of Pharmacy, University of Sarajevo, Sarajevo, Bosnia and Herzegovina; ⁴Department of Pharmacy, Faculty of Health Studies, University of Modern Sciences-CKM, Mostar, Bosnia and Herzegovina (e-mail: ivannapasic@gmail.com, dragoninspiration@yahoo.com)

Lavandula angustifolia Mill. (lavender) is an aromatic and medicinal herb whose flower essential oils (EO) are widely used for the treatment of gastrointestinal, nervous, and rheumatic disorders, and in the perfume industry. Laurus nobilis L. (laurel bay) is an evergreen tree whose EOs have antimicrobial and anti-inflammatory effects. Lavender and bay were collected from Sarajevo and Mostar in Bosnia and Herzegovina. The extraction was performed by hydrodistillation in Clevenger-type apparatus. Phytochemical composition was analyzed by gas chromatography coupled with mass spectrometry. Cytotoxic activities of lavender EO and bay leaf, fruit and seed EOs were investigated against human cervical adenocarcinoma HeLa cells and non-transformed human lung fibroblasts MRC-5 by MTT cell survival assay. Cell cycle phase distribution was examined by flow cytometry. In bay EOs the most abundant component was 1,8-cineole, followed by linalool, bicyclic monoterpenes sabinene, α pinene, and β -pinene. Components identified in the fruit and seed, but not in the leaf were (E)- β -ocymene, camphene, β-elemene, bornyl acetate and trans-caryophyllene. The major component of lavender extract was linalool accompanied by linalyl acetate, lavandulyl acetate, camphor, 1,8-cneole, borneol, a-terpineol, and terpinene-4-ol. The four tested EOs showed concentration-dependent cytotoxic effects on HeLa and MRC-5 cells. Among examined EOs, lavender EO exerted the strongest cytotoxic activity on HeLa cells with IC50 value of 0.11 µL/mL. Bay seed and fruit EOs exerted stronger cytotoxicity on HeLa cells than bay leaf EO (IC50 values: 0.17, 0.21, and 3.35 µL/mL, respectively). When compared with sensitivity of HeLa cells, normal MRC-5 cells showed similar sensitivity to the cytotoxic activity of the four tested EOs. Lavender EO applied at IC50 concentration, during 24 h caused remarkable increase in the percentage of HeLa cells within the subG1 cell cycle phase, in comparison with control cells (64.69% vs 2.47%). Pretreatment with caspase-3, caspase-8 or caspase-9 inhibitor before 24 h treatment with lavender EO did not cause changes in the percentage of cells in the subG1 phase in comparison with HeLa cells exposed only to lavender oil. Our results showed that lavender and bay EOs exerted potent cytotoxic activity against HeLa cells. Additional investigations are necessary to explore cytotoxic effects of these EOs against various cancer cell lines and mechanisms underlying anticancer effects.