



## A STUDY ON THE ANTIBACTERIAL ACTIVITY OF ETHANOLIC AND METHANOLIC EXTRACTS OF *SATUREJA HORTENSIS* L.

Seyyed Mansour Seyyednejad,<sup>1,\*</sup> Zahra Hasannejad,<sup>1</sup> Hossein Motamedi,<sup>1</sup> Fariba Dehghani<sup>1</sup>

<sup>1</sup>Department of Biology, Faculty of Science, Shahid Chamran University, Ahvaz, Iran  
E-mail: author1@institution.edu

Aromatic plants are known as one of important sources of antimicrobial agent. The genus *Satureja* L. (savory, saterei) includes more than 30 species belonging to the family Lamiaceae. *Satureja hortensis* L. produce antimicrobial secondary metabolites and essential oils as a part of their normal program of growth or in response to pathogens attack or stress condition. The aim of this study was evaluation and comparing the antibacterial activity of methanolic and ethanolic extracts of *S. hortensis* L. Against some bacterial pathogens [1, 2]. Plants were collected from farmland in Ahvaz, Khouzestan. Then ethanolic and methanolic extracts were prepared using 1g of dried plant powder and 10 mL of 80% ethanol or methanol. Sterile blank discs were saturated with (100, 200, 400 and 600 mg/ml) concentrations of these extracts and Antibacterial activity of extracts was assessed using standard disc diffusion method against some pathogenic bacteria. A lawn culture of test bacteria with 0.5 McFarland turbidity prepared on Muller-Hinton agar and discs were placed on lawn cultures and incubated at 37°C for 24h. Simultaneously discs containing standard antibiotics were placed on cultures. After incubation the inhibition zone diameter around each disc was measured in millimeter [3, 4]. Different concentrations of both ethanolic and methanolic extracts of this plant showed considerable antibacterial activity against some gram positive bacteria *Staphylococcus aureus* (14mm), *Bacillus cereus* (19mm) and also gram negative bacteria *Proteus mirabilis* (14mm) and *Pseudomonas aeruginosa* (11mm). Based on these results it can be suggested that *S. hortensis* L. is an medicinal plant that have this potential to be used as a new source for antibiotic discovery against bacterial pathogens especially pathogens such as *Staphylococcus*, *Bacillus* and also for treatment of *Proteus mirabilis* infection.

### References

- [1] Mihajilov-krstev T. et al. *Arch. Biol Sci Belgrade*. 2010, 63(1), 109-116.  
[2] Mihajilov-krstev T. et al. *Bulletin USAMV-CN*. 2007, 62(1-2), 146-150.  
[3] Seyyednejad, S. M.; Koochak, H.; Darabpour, E.; Motamedi, H. *Asian P J of Tro Med*. 2010, 351-355.  
[4] Motamedi, H.; Darabpour, E.; Gholipour, M.; Seyyednejad, S. M. *Inter J of Pharma*. 2010, 6(1), 117-122.

## ZINC AND IRON NANO OXIDE INDUCED PRODUCTION OF HYPERICIN AND HYPERFORIN IN CELL SUSPENSION CULTURE OF *HYPERICUM PERFORATUM* (L.)

Ebrahim Sharafi,<sup>1,2</sup> Tahereh Hasanloo,<sup>3,\*</sup> Seyyed Mojtaba Khayyam Nkoei,<sup>4</sup> Mohamad Hossin Fotokian,<sup>1</sup> Daruosh Davoodi,<sup>2</sup> Hossin Hadavand Mirzae<sup>1</sup>

<sup>1</sup>Department of Molecular Physiology, Agricultural Biotechnology Research Institute of Iran, Karaj, Iran

<sup>2</sup>Department of Agricultural Biotechnology, Shahed University, Tehran, Iran

<sup>3</sup>Department of Microbial Biotechnology, Agricultural Biotechnology Research Institute of Iran, Karaj, Iran

<sup>4</sup>Department of Nanotechnology, Agricultural Biotechnology Research Institute of Iran, Karaj, Iran

E-mail : [hasanloo@abrii.ac.ir](mailto:hasanloo@abrii.ac.ir)

*Hypericum perforatum* L. (St. John's wort) is an herbal remedy widely used in treatment of mild to moderate depression [1]. Hypericin and hyperforin, a photosensitive naphodianthrone, are believed to be the compounds responsible for reversing the depression symptoms [2]. Hypericin and hyperforin production have been induced using elicitors in cell cultures of *H. perforatum* [3]. In the present study the effect of zinc and iron nano oxides at different concentrations (0, 50, 100, 150 ppb/20 ml culture) were investigated for hypericin and hyperforin production in *H. perforatum* cell suspension culture. Detection and identification of hypericin and hyperforin was carried out by high performance liquid chromatograph method. The highest hypericin and hyperforin production were observed in media supplemented with 100ppb zinc and iron nano oxide. Zinc and iron nano oxide enhanced hypericin production (11.18 and 7.87  $\mu\text{g g}^{-1}$  DW, respectively) 3 and 13-fold, respectively, higher than that of the control. A dramatic increase in hyperforin production was achieved after exposure to zinc and iron nano oxide (195.62 and 217.45  $\mu\text{g g}^{-1}$  DW, respectively) 5 and 12-fold, respectively, higher than that of the control. These observations suggested that nanoparticles could be introduced as an appropriate candidate for elicitation studies of *in vitro* secondary metabolites production.

### References

- [1] Cui, X. H.; Murthy, H. N.; Wu, C. H.; Paek, K. Y. *In Vitro Cell Dev Biol*. 2010, 48, 437-444.  
[2] Gadzovska, S.; Maury, S.; Delaunay, A.; Spasenoski, M.; Joseph, J.; Hagege, D. *Plant Cell Tiss Organ Cult*. 2007, 19, 1-12.  
[3] Liu, X.; Zhang, X.; Zhang, S.; Sun, J. *Plant Cell Tiss Organ Cult*. 2007, 91, 1-7.