

ANTIMICROBIAL ACTIVITY OF *CHELIDONIUM ALKALOIDS* INVESTIGATED BY BIOARENA SYSTEM

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BioArena system, which integrates the up-to-date methodology and biological results of bioautography with TLC, is especially suitable for investigating biochemical interactions in a sorbent bed after chromatographic separation. The antimicrobial effect of *Chelidonium alkaloids* has been investigated in this system.

Greater celandine (*Chelidonium majus* L.), a member of Papaveraceae family, is widely distributed in Europe and Western Asia. This plant is a well known source of benzophenanthridine (chelidonine, sanguinarine and chelerythrine), protoberberine (berberine) and protopine (coptisine) alkaloids. These alkaloids exhibit choleric, colagogue, spasmolytic, antitumor, antiinflammatory and antimicrobial actions.

Although the antimicrobial activity of *Chelidonium alkaloids* against human bacteria has been reported the mechanism of this action is almost unknown. In this case for bacterial biotest, the phytopathogen *Pseudomonas savastanoi* pv. *phaseolicola* race 6, causing halo blight on bean, was used. The antibacterial activity of *Chelidonium alkaloids* was visualised by staining of bioautograms with aqueous solution of MTT (3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide). To study the mechanism of action of *Chelidonium alkaloids* co-factor substances, as L-arginine, glutathione and copper(II)-sulphate were added separately to the cell suspension in different concentration before inoculation [1].

An important fact is that there is a relationship between the amount of HCHO-capturing agent and antimicrobial activity of *Chelidonium alkaloids*, that is, larger quantities of L-arginine and glutathione always resulted in a greater decrease in antimicrobial activity. These alkaloids exhibit an antibacterial activity, while their methylated derivatives promote cell proliferations under the same conditions. The Cu(II), the oxidized form of copper, an essential trace metal found in all biological systems which enhances the antibacterial activity of HCHO.

It is obvious that BioArena provides more information than conventional bioautography. We can change the incubation time, and may observe the changes on the sorbent layer during 5-6 or more days, using endogenous and/or exogenous substances in culture medium on the antimicrobial action of the separated components.

[1] Tyihák E, Botz L, Ott P, Nagy S, Kocsis B, Király-Véghely Zs and Mincsovcics E. The combination of the overpressured chromatography and bioautography and its applications to the analysis of molecules influencing cell proliferation. *Chemical Analysis (Warsaw)* 2003; 48: 543-553.