

## PŮVODNÍ PRÁCE

**Biotic elicitation of the *Trifolium pratense* L. suspension culture**

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Received 17 March 2008 / Accepted 21 April 2008

## SUMMARY

**Biotic elicitation of the *Trifolium pratense* L. suspension culture**

The important substances of secondary metabolism include flavonoids and isoflavonoids. The *Trifolium pratense* L. (*Fabaceae*) suspension culture's yield of these metabolites is low, thus an attempt was made to increase the production by elicitation. The endogenous signal substances of the plants' defensive responses include jasmonic acid that also functions as a biotic elicitor in the case of exogenous application. In the experiment the authors monitored the impact of 4 different concentrations of jasmonic acid on the *Trifolium pratense* L. (variety DO-8 and variety DO-9) suspension culture's yield. The culture was cultivated in Gamborg medium to which 2 mg.l<sup>-1</sup> of 2,4-dichlorophenoxyacetic acid and 2 mg.l<sup>-1</sup> of 6-benzylaminopurine were added. The maximum increase in the production of flavonoids was achieved, when compared with the control samples, with both varieties of the 500 μmol concentration (DO-9, 6-hour application by 140%; DO-8, 24-hour application by 65%). The production of isoflavonoids (genistin, daidzein, genistein, and formononetine) was best stimulated in both varieties by the 50 μmol concentration (in the case of DO-9 variety, the 48-hour application increased the content of genistin by up to 845%).

**Key words:** *Trifolium pratense* L. suspension culture – flavonoids and isoflavonoids – biotic elicitation – jasmonic acid

Čes. slov. Farm., 2008; 57, 107–110

## SOUHRN

**Biotická elicítace suspenzní kultury *Trifolium pratense* L.**

Mezi významné látky sekundárního metabolismu patří flavonoidy a isoflavonoidy. Suspenzní kultura *Trifolium pratense* L. (*Fabaceae*) má však nízkou produkci těchto metabolitů, a proto byla snaha zvýšit jí elicítací. K endogenním signálním látkám rostlinných obraných reakcí patří kyselina jasmínová, která v případě exogenní aplikace působí také jako biotický elicitor. V práci byl sledován vliv čtyř koncentrací kyseliny jasmínové na produkci suspenzní kultury *Trifolium pratense* L. (variety DO-8 a varieta DO-9). Kultura byla kultivovaná na mediu podle Gamborga s přísadkou 2 mg.l<sup>-1</sup> 2,4-dichlorfenoxyoctové kyseliny a 2 mg.l<sup>-1</sup> 6-benzylaminopurinu. Maximální zvýšení produkce flavonoidů oproti kontrole vyvolala u obou variet koncentrace 500 μmol (6hodinová aplikace o 140 % u variety DO-9 a 24hodinová aplikace o 65 % u variety DO-8). Produkci isoflavonoidů (genistinu, daidzeinu, genisteinu a formononetinu) stimulovala u obou variet nejvíce koncentrace 50 μmol (u variety DO-9 zvýšila 48hodinová aplikace obsah genistinu až o 845 %).

**Klíčová slova:** suspenzní kultura *Trifolium pratense* L. – flavonoidy a isoflavonoidy – biotická elicítace – kyselina jasmínová

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Čes. slov. Farm., 2008; 57, 107–110

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The elicitation method is one of the means to increase the production of secondary metabolites in explant cultures of plants. The elicitation utilizes substances possessing the signal effect properties—the elicitors. These function as stress factors, induce increased synthesis of phytoalexins and of other secondary metabolites that contribute to the defense potential of the plant or of the explant.

Jasmonic acid and its precursors and derivatives are some of the endogenous signal substances of the plants' defensive responses<sup>1)</sup>. Jasmonic acid and its methyl ester are natural hormonal regulators controlling the plant's aging and inducing processes that take place after the plant is injured. In the exogenic application case, they may also function as elicitors<sup>2)</sup>.

Important secondary metabolites of intact plants and of explant cultures include flavonoids and isoflavonoids which possess a wide range of biological effects<sup>3–12)</sup>. In the observed suspension culture, *Trifolium pratense* L. (*Fabaceae*), the production rate of these metabolites is low, thus there was an attempt to increase their production with the jasmonic acid biotic elicitor.

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## EXPERIMENTAL PHASE

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### Instruments

A 200S analytical scales manufactured by Sartorius, Göttingen; a PS 20A autoclave by Chirana, Brno; an HS 31A hot-air sterilizer by Chirana, Brno; a laminar flow workbench by Fatran LF, Žilina; a roller by Vývojové Dílny AV ČR, Praha; a 2010 shaker by Unimax, Heidolph; a CE 1010 spectrophotometer by Cecil Instruments, Cambridge; a liquid chromatograph (a PU-2089 pump, a MD-2015 detector, an AS-2055 automatic sample injector) by Jasco, Tokyo.

### *Trifolium pratense* L. suspension culture

The suspension cultures (variety DO-8, DO-9) were derived from callus cultures. First by mechanical loosening on the shaker followed by the cultivation in Gamborg nutrient media<sup>13)</sup> with the addition of 2 mg.l<sup>-1</sup> of 2,4-dichlorophenoxyacetic acid and 2 mg.l<sup>-1</sup> of 6-benzylaminopurine, at the temperature of 25 °C, 16-hr light/8-hr dark period. The subcultivation interval lasted 14 days<sup>14)</sup>.

### Elicitation

To elicitate, we used jasmonic acid dilutions in 96% ethanol in the concentrations of 5 µmol, 50 µmol, 500 µmol, and 5000 µmol. During the elicitation process, we added 1.0 ml of the elicitor to a two-year-old suspension culture on the 21<sup>st</sup> day of cultivation<sup>14)</sup>. The control cultures received 1.0 ml of distilled water. The elicitor application durations were 6, 24, 48, and 168 hours. The cells were separated from the media by vacuuming, rinsed in distilled water, and dried at the laboratory room temperature.

### Determining the flavonoids and isoflavonoids

The elicited and the inspection samples underwent photometric determination of flavonoids in accordance with the Czech Pharmacopoeia 2005<sup>15)</sup> and the determination of isoflavonoids using the HPLC method<sup>16)</sup>. The HPLC determination process conditions were as follows: a RP-18 Lichrospher column (250 × 4 mm, particles size 5 µm) with a precolumn made of the same material; elution: linear gradient of the mobile phase A (methanol) in phase B (water containing 0.15% of phosphoric acid) 30–80% from 0 to 9 minutes was followed by the isocratic elution mobile phase in the composition of 80% of phase A in phase B from 9 to 15 minutes; the elution speed was 1.1 ml/min; the detection was carried out at the 260 nm wave length. The obtained results were statistically evaluated in the t-test for at least 3 members of the team and at the p = 0.05 level of statistical significance.

Tab. 1. Production of flavonoids in suspension culture of *Trifolium pratense* L. (variety DO-8, DO-9) elicited with jasmonic acid

Elicitor concentration (µmol)	Time after elicitation (hours)	Flavonoid contents (%) variety DO-8	Flavonoid contents (%) variety DO-9
control	6	0.068 ± 0.007	0.031 ± 0.008
	24	0.068 ± 0.007	0.031 ± 0.008
	48	0.068 ± 0.007	0.031 ± 0.008
	168	0.057 ± 0.009	0.053 ± 0.006
5	6	0.044 ± 0.011	0.048 ± 0.009
	24	0.085 ± 0.004	0.032 ± 0.008
	48	0.059 ± 0.003	0.033 ± 0.010
	168	0.024 ± 0.004	0.058 ± 0.007
50	6	0.083 ± 0.012	0.060 ± 0.011
	24	0.091 ± 0.005	0.057 ± 0.008
	48	0.057 ± 0.002	0.030 ± 0.006
	168	0.042 ± 0.009	0.056 ± 0.010
500	6	0.070 ± 0.003	0.074 ± 0.005
	24	0.112 ± 0.005	0.059 ± 0.013
	48	0.056 ± 0.005	0.044 ± 0.009
	168	0.024 ± 0.002	0.060 ± 0.007
5000	6	0.079 ± 0.011	0.010 ± 0.009
	24	0.092 ± 0.003	0.019 ± 0.007
	48	0.089 ± 0.006	0.025 ± 0.011
	168	0.023 ± 0.003	0.049 ± 0.008

Tab. 2. Production of isoflavonoids in suspension culture of *Trifolium pratense* L. (variety DO-8) elicited with jasmonic acid

Elicitor concentration (μmol)	Time after elicitation (hours)	Isoflavonoid contents (%)			
		genistin	daidzein	genistein	formononetin
control	6	0.112 ± 0.007	0.042 ± 0.005	0.013 ± 0.004	0.033 ± 0.006
	24	0.111 ± 0.004	0.042 ± 0.008	0.013 ± 0.002	0.033 ± 0.004
	48	0.111 ± 0.006	0.041 ± 0.003	0.012 ± 0.006	0.032 ± 0.002
	168	0.054 ± 0.003	0.033 ± 0.006	0.012 ± 0.003	0.021 ± 0.005
5	6	0.112 ± 0.009	0.042 ± 0.008	0.011 ± 0.001	0.022 ± 0.003
	24	0.112 ± 0.006	0.041 ± 0.003	0.011 ± 0.001	0.024 ± 0.006
	48	0.093 ± 0.008	0.041 ± 0.005	0.010 ± 0.003	0.031 ± 0.004
	168	0.011 ± 0.007	0.022 ± 0.003	–	0.023 ± 0.006
50	6	0.280 ± 0.003	0.094 ± 0.008	0.050 ± 0.010	0.043 ± 0.005
	24	0.182 ± 0.007	0.032 ± 0.006	0.013 ± 0.008	0.023 ± 0.006
	48	0.063 ± 0.008	0.030 ± 0.004	0.014 ± 0.002	0.022 ± 0.009
	168	–	0.023 ± 0.003	0.032 ± 0.006	0.012 ± 0.008
500	6	0.252 ± 0.009	0.087 ± 0.006	0.013 ± 0.004	0.023 ± 0.003
	24	0.160 ± 0.004	0.062 ± 0.008	0.013 ± 0.005	0.024 ± 0.002
	48	0.061 ± 0.007	0.053 ± 0.004	0.021 ± 0.003	0.031 ± 0.004
	168	–	0.012 ± 0.008	–	0.044 ± 0.011
5000	6	0.092 ± 0.008	0.032 ± 0.006	0.017 ± 0.012	0.010 ± 0.004
	24	0.101 ± 0.003	0.024 ± 0.004	0.022 ± 0.006	0.010 ± 0.010
	48	0.110 ± 0.006	0.032 ± 0.002	0.021 ± 0.004	0.031 ± 0.008
	168	0.012 ± 0.004	0.030 ± 0.003	–	0.042 ± 0.006

Tab. 3. Production of isoflavonoids in suspension culture of *Trifolium pratense* L. (variety DO-9) elicited with jasmonic acid

Elicitor concentration (μmol)	Time after elicitation (hours)	Isoflavonoid contents (%)			
		genistin	daidzein	genistein	formononetin
control	6	0.052 ± 0.003	0.023 ± 0.006	0.020 ± 0.000	0.011 ± 0.003
	24	0.052 ± 0.006	0.023 ± 0.002	0.022 ± 0.006	0.011 ± 0.001
	48	0.051 ± 0.004	0.020 ± 0.010	0.024 ± 0.002	0.012 ± 0.000
	168	0.024 ± 0.006	0.013 ± 0.006	0.010 ± 0.001	0.011 ± 0.003
5	6	0.062 ± 0.010	0.013 ± 0.002	0.011 ± 0.000	0.010 ± 0.000
	24	0.053 ± 0.006	0.023 ± 0.006	0.011 ± 0.001	0.011 ± 0.003
	48	0.101 ± 0.008	0.024 ± 0.008	0.024 ± 0.006	0.012 ± 0.006
	168	0.053 ± 0.004	0.022 ± 0.002	0.010 ± 0.000	0.011 ± 0.000
50	6	0.097 ± 0.008	0.011 ± 0.006	0.011 ± 0.001	0.011 ± 0.002
	24	0.112 ± 0.009	0.013 ± 0.004	0.023 ± 0.006	0.031 ± 0.004
	48	0.482 ± 0.007	0.081 ± 0.006	0.032 ± 0.000	0.042 ± 0.006
	168	0.081 ± 0.004	0.013 ± 0.001	0.010 ± 0.002	0.020 ± 0.010
500	6	0.062 ± 0.004	0.010 ± 0.006	0.011 ± 0.001	0.021 ± 0.006
	24	0.087 ± 0.006	0.012 ± 0.004	0.023 ± 0.006	0.032 ± 0.004
	48	0.123 ± 0.010	0.024 ± 0.002	0.023 ± 0.001	0.033 ± 0.002
	168	0.075 ± 0.008	0.013 ± 0.006	0.011 ± 0.001	0.024 ± 0.011
5000	6	0.061 ± 0.005	0.013 ± 0.006	0.011 ± 0.002	0.010 ± 0.006
	24	0.084 ± 0.007	0.010 ± 0.002	0.012 ± 0.002	0.012 ± 0.004
	48	0.102 ± 0.008	0.023 ± 0.008	0.023 ± 0.006	0.032 ± 0.006
	168	0.073 ± 0.004	0.013 ± 0.006	0.011 ± 0.002	0.031 ± 0.002

## RESULTS AND DISCUSSION

Increasing the production of therapeutically important secondary metabolites in *in vitro* cultures is one of the closely watched spheres in pharmaceutical biotechnology. Tangibly speaking, the elicitation of explant cultures is a very promising method.

The *Trifolium pratense* L. suspension culture (variety DO-8, DO-9) was observed for the effect of jasmonic acid's biotic elicitor on the production of flavonoids and isoflavonoids.

The findings of previous experiments imply<sup>14)</sup> that the elicitation of the *Trifolium pratense* L. suspension culture is best on day 21 of subcultivation; our culture's elicitation was therefore carried out on that day of cultivation.

Four concentrations of jasmonic acid were tested (5 μmol, 50 μmol, 500 μmol, and 5000 μmol); these concentrations had been chosen from within the range that is usually applied with this type of elicitor<sup>17–19)</sup>. The time periods of the elicitor's effect that were tested (6, 24, 48 and 168 hours) had been based on the results of the already performed tests<sup>20)</sup> and on different pieces of published information that deal with an increase in the

production of the secondary metabolites after the addition of an elicitor in the range of several hours to several days<sup>17,18,21,22</sup>. Inspection cultures were collected after 6 and 168 hours since their production does not change notably in such short time intervals.

The results of the biotic elicitation of the *Trifolium pratense* L. suspension culture (variety DO-8) with jasmonic acid (Table 1) clearly indicate that flavonoid production was influenced in a positive way. The maximum content (0.112%) was recorded after a 24-hour exposure to a 500  $\mu\text{mol}$  concentration and, after comparing this result with the control culture result, we can state that a statistically important increase in the production by 65% was achieved. Statistically important results were also achieved with the remaining concentrations of the elicitor after a 24-hour exposure when compared with the control samples.

The *Trifolium pratense* L. suspension culture (variety DO-9) showed a better elicitation effect with a 500  $\mu\text{mol}$  concentration, too (Table 1). The maximum content of flavonoids (0.074%) and a statistically important increase in the production by 140%, compared with the control samples, were recorded after a 6-hour application.

In the *Trifolium pratense* L. suspension culture (variety DO-8, DO-9) elicited with jasmonic acid, we also checked the production of isoflavonoids. In the control culture we recorded the isoflavonoids genistin, daidzein, genistein, and formononetin. In the case of the DO-8 variety (Table 2), we can value namely the 6-hour application of the 50  $\mu\text{mol}$  concentration which increased, compared with the control culture, the production of all isoflavonoids under observation – genistin by 150%, daidzein by 123%, genistein by 285%, and formononetin by 30%. The increase in the production of these isoflavonoids was also induced by the 500  $\mu\text{mol}$  concentration.

In the case of the variety DO-9 (Table 3), the best elicitation effect of jasmonic acid was also recorded at the 50  $\mu\text{mol}$  concentration. The maximum increase in the content of all isoflavonoids under observation, when compared with the control culture, was brought about during the 48-hour application of the 50  $\mu\text{mol}$  concentration when the increase was as follows: genistin by 845%, daidzein by 300%, genistein by 30%, and formononetin by 250%. The production of genistin was also increased by the other concentrations of the elicitor and the increase values were again highest after the 48-hour application.

The 48-hour application of the 50  $\mu\text{mol}$  concentration also stimulated best the production of anthracene derivatives in the *Rheum palmatum* L. suspension culture<sup>20</sup>. Other studies confirm the fact that jasmonic acid is one of the important signal substances of the plants' defensive responses that result in an increased production of secondary metabolites<sup>23–26</sup>.

This work was sponsored under the Research Project No. MSM 0021620822.

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