

ABSTRACT

The study is aimed to enhance *in vitro* production of secondary metabolites in *Genista tinctoria* L. via elicitor treatment. The different levels of elicitor concentration – selenium dioxide were utilized to affect the quantity of isoflavonoids occurred in cultures.

Experiment was performed in callus and suspension cultures on MS nutrient media supplemented with 10 g l^{-1} of NAA (α -naphthylacetic acid) as growth regulator. The elicitor was added in the form of solution in concentrations of $9,012 \cdot 10^{-3} \text{ mol l}^{-1}$; $9,012 \cdot 10^{-4} \text{ mol l}^{-1}$ and $9,012 \cdot 10^{-5} \text{ mol l}^{-1}$. It was exposed for 6, 12, 24, 48, 72 and 168 hours. The content of isoflavonoids was determined by HPLC in dry weight (DW) and medium.

The most effective production of genistin ($6,20 \text{ mg g}^{-1} \text{ DW}$, $8,30 \text{ mg g}^{-1} \text{ DW}$) in callus culture was measured. It was reached in concentrations of $9,012 \cdot 10^{-4} \text{ mol l}^{-1}$ and $9,012 \cdot 10^{-5} \text{ mol l}^{-1}$ after 168 h elicitor treatment. The second most satisfactory genistin level $5,20 \text{ mg g}^{-1} \text{ DW}$ was detected after elicitor application in concentration of $9,012 \cdot 10^{-4} \text{ mol l}^{-1}$ after 6 h. The content of genistein, daidzein and formononetin in callus culture was low and in the most cases equal zero compared to control samples. The content of biochanin A was equal zero compared to control samples.

The most efficient daidzein production ($37,10 \text{ mg g}^{-1} \text{ DW}$) in suspension culture was detected after elicitor treatment in concentrations of $9,012 \cdot 10^{-3} \text{ mol l}^{-1}$ and $9,012 \cdot 10^{-5} \text{ mol l}^{-1}$ after 24 h. The second most abundant content $11,30 \text{ mg g}^{-1} \text{ DW}$ of daidzein was reached after selenium dioxide treatment in concentrations of $9,012 \cdot 10^{-3} \text{ mol l}^{-1}$ and $9,012 \cdot 10^{-5} \text{ mol l}^{-1}$ after 12 h. High production of genistin was observed in concentration of elicitor $9,012 \cdot 10^{-3} \text{ mol l}^{-1}$ ($15,2 \text{ mg g}^{-1} \text{ DW}$) after 24 h, in concentration $9,012 \cdot 10^{-4} \text{ mol l}^{-1}$ ($9,5 \text{ mg g}^{-1} \text{ DW}$) after 168 h and in concentration $9,012 \cdot 10^{-5} \text{ mol l}^{-1}$ ($10,7 \text{ mg g}^{-1} \text{ DW}$) after 48 h. The content of genistein and biochanin A was equal zero after elicitor application. Production of formononetin was very low.

Isoflavonoids were not released into nutrient media. Selenium dioxide can be recommended to increase efficiently isoflavonoids production – genistin in callus and suspension cultures and daidzein in suspension cultures.