
#### Abstract

Nowadays, humankind is facing the global climate crisis with new challenges. One of them is a threat of vast crop yield losses caused by growing populations of fungal plant pathogens such as Fusarium proliferatum. This hemibiotrophic organism infects various host plants, including asparagus, maize, garlic and pineapple. Moreover, this fungus is able to produce a number of harmful mycotoxins including fumonisin $\mathrm{B}_{1}$, moniliformin, bauvericin and fusaproliferin. The lack of effective plant protection products widens an issue of F. proliferatym management in an environment. Therefore, in order to evaluate the F. proliferatum mechanisms triggered during plant-pathogen interaction, three experiments with extracts, fractions and bioactive compounds derived from host plants were conducted. Experiments were carried out in vitro in liquid cultures of $F$. proliferatum strains treated with plant stressors. Then, a series of analyses were performed, including dry biomass measurements, selected genes' expression analysis as well as quantification of type B ( $\mathrm{FB}_{1-3}$ ) fumonisins. The results obtained have confirmed the significant effect of tested extracts, fractions and plant bioactive compounds on primary and secondary metabolism of F. proliferatum. Stress factors influenced the expression of primary target genes - especially genes encoding heat shock proteins, and also the expression of the FUM gene cluster involved in fumonisins biosynthesis. Extracts, fractions and plant bioactive compounds might act as stress factors or signal molecules, depending on their concentrations. Applied stress factors caused a significant reduction of fumonisins content, while low concentration of chlorogenic acid was an exception and led to $\mathrm{FB}_{1}$ induction. Further research on the defense mechanisms of $F$. proliferatum and its interaction with host plant is needed. There is a chance that extracts, fractions or plant bioactive compounds will be used as plant protection products in the future.


