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PR3.26

The black truffle of perigord responds to cold stress with an extensive reprogramming of its transcriptional activity

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Free-living fungi often encounter different kinds of environmental stresses, including changes in temperature, osmolarity, pH, humidity, availability of O2 and nutrients, exposure to toxins, UV or heavy metals, as well as competition with other organisms. To understand the cell adaptation and the survivor in non-ideal conditions, a better comprehension of many basic events is required. Tuber melanosporum can be subjected to different stress conditions, considering its life cycle. In this work, the genome sequence of the ectomycorrhizal ascomycete Tuber melanosporum was analysed with the aim to identify and characterize genes involved in environmental stress response. As a second step whole genome arrays were used to verify the transcriptional profiling in the presence of a cold shock (4°C for 7 days). In a whole genome microarray (7496 genes/probe), 423 genes resulted significantly differentially expressed (> 2.5 fold; p-value < 0.05) in stressed mycelia compared to the control ones. After 4°C exposure for 7 days the number of up-regulated genes was 187; the down-regulated genes were 236. The 50-60% of the up- or down-regulated transcripts had no KOG classification and were clustered as unclassified proteins, which represent the most abundant category both in up- and down-regulated genes. A gene subset, concerning a range of biological functions, was chosen to validate the microarray experiment using qRT-PCR. Sixteen out of 22 considered genes confirmed the array data. At our knowledge this is the first work, which considers the global gene expression profiling in a filamentous fungus under cold stress condition.

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Screening and sequence analysis of zhd101 (zearalenone lactonohydrolase) homologues in *Trichoderma/Clonostachys* sp.

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Zearalenones are economically important group of *Fusarium sp.* mycotoxins, exhibiting estrogenic activity and chemical strucure consisting of a resorcinol moiety fused to a 14-membered macrocyclic lactone. These compounds are converted into a far less estrogenic product by incubation with *Clonostachys rosea* IFO 7063 expressing *zhd101* zearalenone lactonohydrolase gene. In the present study we described screening of *Trichoderma/Clonostachys* combined collection for new strains with functional lactonohydrolase homologues. In the screened samples, we observed degradation reactions in 10 of 79 total *Trichoderma sp.* and *Clonostachys sp.* isolates and have been able to determine new lactonohydrolase homologue sequences with average sequence identity of 90%.

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