Oxidative stress response in Arabidopsis thaliana after exposure to uranium: the role of glutathione

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Introduction

Uranium is a heavy metal and radionuclide with greater risk for chemical toxicity than radiological. Due to anthropogenic pollution 2-5 mg/kg uranium occurs in the earths crust. It is know that uranium induces oxidative stress in plants. This is a condition with an **imbalance** in the production and elimination of reactive oxygen species (ROS), causing cellular damage such as oxidized DNA bases and lipid peroxidation. The plant developed a defence mechanism to cope with oxidative stress of which glutathione is an important element. However, the role of GSH under uranium stress is not fully understood. Therefore, the objective of this work is to further analyse the role of GSH in *Arabidopsis* 0 thaliana plants under uranium stress.

Materials & methods

Arabidospis thaliana plants were cultivated in a hydroponic set-up during 18 days after which they were exposed during 3 days to different uranium concentrations (0, 3, 6.25, 12.5, 25 or 50 µM). After harvesting the plants, the transcriptome was sequenced at the University of Antwerp (Ilumina HiSeq2000 + TruseqTM RNA sample prep kit) and the **Differentially expressed** genes were identified (cut-off values: |Log2 Fold Changes| > 1 & False discovery rate < 0.05). The differentially expressed genes were analysed in a **GO enrichment analaysis** using the web-based tool Metascape [2]. The analyses were done separately for up-and downregulated genes.



Results & discussion



Figure 4: Results of Arabidopsis thaliana after exposure to different uranium concentrations: (A) Differential expressed genes; (B) Venn diagram upregulated genes; (C) Venn diagram downregulated genes.



Conclusion

- No differentially expressed genes related to: GSH synthesis, AsA-GSH and phythochelatins
- Important role for GSTs and peroxides in both detoxification and redox homeostasis
- Uranium may be detoxified by direct quenching with a role for GSTU3/4/12

Perspectives

- Cad2-1 mutant (GSH deficient) with analyses such as uranium uptake, lipid peroxidation, enzymatic activity and growth
- Cad1-3 mutant (phythochelatin deficient)
- Investigate specific genes with the (in this work) already developed primers
- Measure GST enzymatic activity

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