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# CYTOKININ ACTION IS LINKED TO OXIDATIVE STRESS AS SHOWN BY PROTEOME ANALYSIS IN ARABIDOPSIS



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#### I. Introduction

Cytokinins (CKs) regulate diverse developmental processes in plants. The molecular mechanisms of CK action are currently the subject of intense research. We analyzed changes in the *Arabidopsis* proteome caused by increase in endogenous CKs which was achieved by the employment of an improved pOp-*iptl*/LhGR transgenic system in which the production of CKs is inducible by dexamethasone (DEX) (Craft et al., 2005).

#### II. Results

## 1. Plant material



**Fig. 1**: Wild-type Columbia 0 (**Col**; control) and a transgenic pOp-*iptl*LhGR line (**ipt**) grown for 15 days on MS medium in the absence and presence of 2.5 uM DEX, at 16h light/8h dark; 21 °C day/19 °C night; light intensity 117 umol.m<sup>-2</sup>.s<sup>-1</sup>.

#### 2. Quantitative analysis of CKs

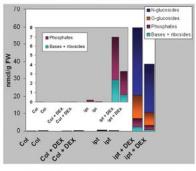


Fig. 2: CKs extracted by methanol/ chloroform/acetic acid and purified on RP-C18 columns were quantified by LC/MS using deuterated CKs as internal standards. Samples were analysed in duplicates.

# 3. Two-dimensional (2-D) electrophoresis

2-D electrophoresis was performed as follows: IPG strips pH 4-7; 500 ug protein/strip; 11% SDS-PA gel; colloidal Coomassie G250 staining.

Analysis of gel images in triplicates and subsequent statistical evaluation revealed significant differences between protein maps from DEX treated and non-treated pOp-ipt/LhGR seedlings (Fig. 3). No significant effects of DEX on wild-type protein maps were observed.

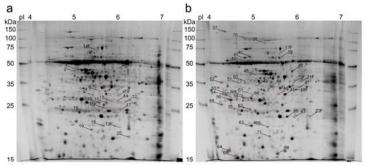


Fig. 3: Protein maps of pOp-ipt/LhGR seedlings grown in the absence (a) and presence (b) of 2.5 uM DEX; numbers indicate the significantly overexpressed spots.

#### 4. Protein identification and functional classification

Total of 62 differentially expressed spots were subjected to MALDI-TOF MS analysis; 58 of them were characterized: 2 specific, 35 upregulated, 19 downregulated and 2 absent in gels from DEX treated pOp-ipt/LhGR plants (Table 1). The identified proteins were classified into functional categories (Fig. 4).

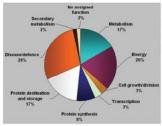


Table 1: Abbreviated list of differentially expressed proteins.

**Fig. 4:** Assignment of the differentially expressed proteins to functional categories according to Bevan et al. (1998).

Regulation	Spot	Protein
Up	45	Jasmonate inducible protein isolog
	18F	2'-hydroxyisoflavone reductase
	37	Cor78 protein
	20F	Glutathione-S-transferase
	43	S-adenosylmethionine synthase
	66	GSH-dependent dehydroascorbate reductase
Down	13	L-ascorbate peroxidase
	10	Probable ethylene-inducible protein

Validity of 2-D analysis results, particularly the high proportion of defence and oxidative stress related proteins, was examined by the following experiments:

#### 5. Quantitative analysis of jasmonic acid (JA)

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Fig. 5: Content of JA as quantified by LC/MS.

#### 6. Localization of hydrogen peroxide

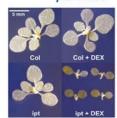
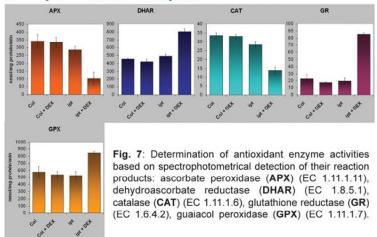
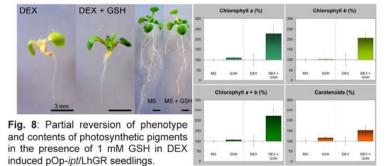


Fig. 6: Chromogenic detection of  $H_2O_2$  in plant tissues by diaminobenzidine (DAB) staining (2.5 mM/2 h).

#### 7. Assays for antioxidant enzymes



#### 8. Reduced glutathion (GSH) treatment



#### III. Summary

- □ 2-D electrophoresis as a top-down approach is a suitable tool for identifying candidates related to CK action on the protein level.
- ☐ High proportion of the differentially expressed proteins were related to oxidative stress and defense mechanisms including JA inducible proteins.
- □ Subsequent analyses revealed increased levels of hydrogen peroxide and JA in seedlings with elevated CKs together with significant changes in activity of several antioxidant enzymes.
- □ Application of reduced glutathion, an antioxidant, partially reversed the adverse effects caused by high CK levels (i.e. retarded development and contents of photosynthetic pigments).
- ☐ The data demonstrate a link between CKs and oxidative stress at the proteome level, implying cross-talk between CKs and JA *in planta*.

References: Bevan M, et al. (1998) Nature 391:485-488 Craft J, et al. (2005) Plant J 41:899-918

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