

## CYCLIN-DEPENDENT KINASE INHIBITORS INDUCE APOPTOSIS IN PLANT CELLS

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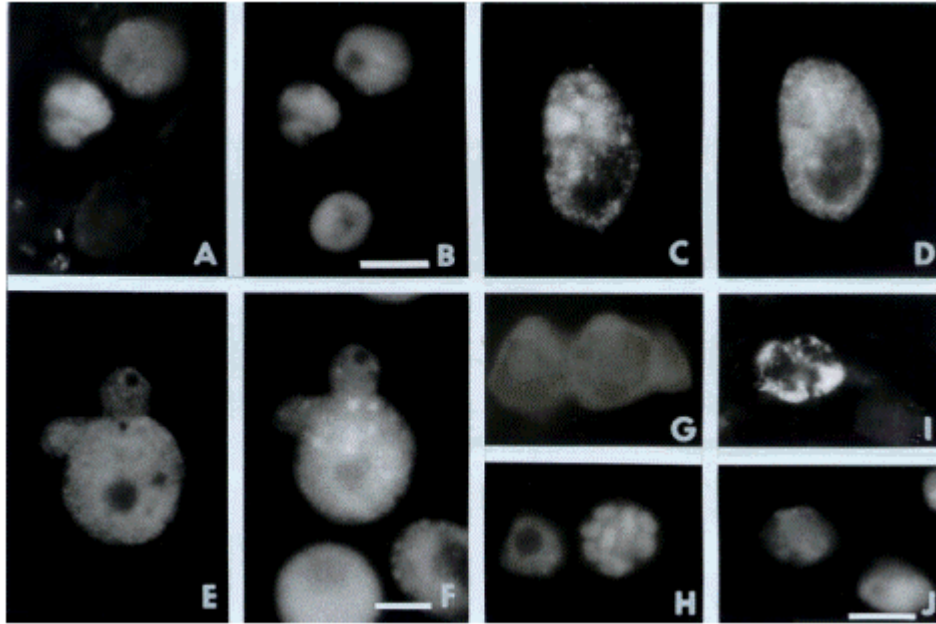
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**INTRODUCTION.** Programmed cell death (PCD) appears to be a ubiquitous process which allows a multicellular organism to eliminate unwanted or defective cells by the process of cellular disintegration without inducing an inflammatory response. As compare to animal cells molecular and cellular mechanisms underlying PCD (or apoptosis) in plants were not characterised yet, although there are many data describing PCD during plant development or resulting form an interactions with environment (hypoxia, plant pathogen) (1). Here we collected the evidence that programmed cell death is induced with cdk inhibitors treatment also in plant cells.

**METHODS.** Seeds of field bean (*Vicia faba* L.) were germinated at 25°C in Hoagland solution. Seedlings with about 2-cm long main roots were treated with cdk inhibitor bohemin or roscovitin at concentration from 20-300 µM for various time periods of 2.612,24 and 48h. CDC-2 was detected with rabbit polyclonal Ab prepared against to a peptide (RITARGALEHEYFKDIK). Microtubular structures were detected with mouse monoclonal antibody DmlA (Sigma) against α-tubulin, or with rabbit affinity purified antibody against α,β-tubulin heterodimer. The γ-tubulin was detected with mouse monoclonal antibody TU-31, or with affinity purified rabbit polyclonal antibody (2).

**RESULTS.** The purine analogues bohemin and roscovitin were used to study the role on CDKs in cell cycle progression and microtubule organisation in *Vicia faba* root tip cells. Both drugs inhibited the activity of immunopurified *Vicia faba* and alfalfa CDC2-kinase. The transcript levels of an A- and B-type cyclin, as well as of the *cdc2* genes, declined in treated root tips, while the mRNA level of a D-type cyclin gene was not affected. An observed transient arrest at the G1/S and G2/M regulatory points indicated that inhibition of the CDC2-kinase had an effect on both transitions (see Fig. 1). In contrast to the regular bipolar spindle in untreated cells, in drug treated metaphase cells abnormally short and dense kinetochore microtubule fibres were observed. These microtubules were randomly arranged in the vicinity of the kinetochores and connected the chromosomes. Thus, the chromosomes were not aligned on the metaphase plate but were arranged in a circle, with kinetochores pointing inwards and chromosome arms pointing outwards. γ-Tubulin, which plays a role in microtubule nucleation, also localised to the centre on the monopolar spindle. The observed abnormalities in mitosis, after inhibition of CDC2-kinase by specific CDK drugs, suggest a role for this enzyme in regulating some of the steps leading to a bipolar spindle structure. These compounds also induce apoptosis of different plant cells *in vivo*.



**Fig. 1.** Immunofluorescence in situ detection of DNA double strands breaks in cells of *V. faba*.

**DISCUSSION.** The doses of roscovitine used here to induce PCD in plant cells were several times higher than that reported as apoptosis inducing in animal cells (3). Lowered sensitivity of plant cells to roscovitine and bohemin treatment might be explained by different penetration or different metabolisms of the drugs, which are analogues of naturally occurring plant growth regulators from the group of cytokinins (4).

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