

# AFLP Analysis and Improved Phytoextraction Capacity of Transgenic *gshI*-Poplar Clones (*Populus* × *canescens* L.) for Copper *in vitro*

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Z. Naturforsch. **60c**, 300–306 (2005)

Clone stability and *in vitro* phytoextraction capacity of vegetative clones of *P. × canescens* (2n = 4x = 38) including two transgenic clones (*ggs11* and *lgl6*) were studied as *in vitro* leaf disc cultures. Presence of the *gshI*-transgene in the transformed clones was detected in PCR reactions using *gshI*-specific primers. Clone stability was determined by fAFLP (fluorescent amplified DNA fragment length polymorphism) analysis. In total, 682 AFLP fragments were identified generated by twelve selective primer pairs after *EcoRI*–*MseI* digestion. Four fragments generated by *Eco*AGT–*Mse*CCC were different (99.4% genetic similarity) which proves an unexpectedly low bud mutation frequency in *P. × canescens*. For the study of phytoextraction capacity leaf discs (8 mm) were exposed to a concentration series of ZnSO<sub>4</sub> (10<sup>–1</sup> to 10<sup>–5</sup> M) incubated for 21 days on aseptic tissue culture media WPM containing 1 μM Cu. Zn<sup>2+</sup> caused phytotoxicity only at high concentrations (10<sup>–1</sup> to 10<sup>–2</sup> M). The transgenic poplar cyt-ECS (*ggs11*) clone, as stimulated by the presence of Zn, showed elevated heavy metal (Cu) uptake as compared to the non-transformed clone. These results suggest that *gshI*-transgenic poplars may be suitable for phytoremediation of soils contaminated with zinc and copper.

**Key words:** Phytoextraction, cyt-ECS (*ggs11*), chl-ECS (*lgl6*)