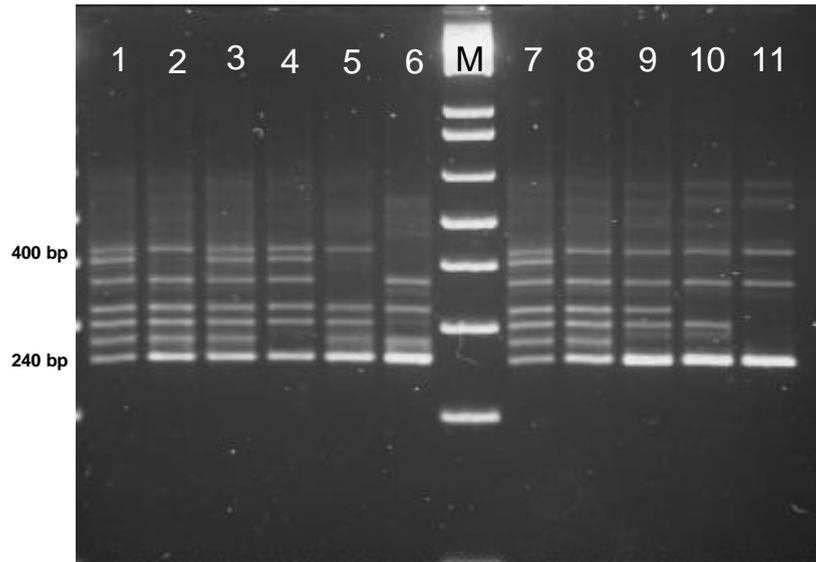


DNA gel-electrophoresis of samples obtained by the PCR-based “DNA-flag test” on *A. niger* ISO strains having different compositions of the *DglaA* loci as a result of gene-conversion or amplification events



Lanes 1-6: DNA-patterns of production strains having still all 7 marked $\Delta glaA$ loci single in lane 1; a conversion of *Bam*HI marked $\Delta glaA$ loci towards the *Pst*I- $\Delta glaA$ locus (*Bam*HI²⁺, *Pst*I⁻ convertant) in lane 2; two *Bam*HI⁺ loci (*Bam*HI²⁺) as a result of an amplification in lane 3, a *Bam*HI²⁺/*Sal*I⁻ convertant in lane 4, a *Bam*HI³⁺/*Pst*I/*Mlu*I⁻ convertant in lane 5; a *Bam*HI⁴⁺/*Sac*I/*Pst*I/*Kpn*I⁻ convertant in lane 6.

M = Marker DNA

Lanes 7 - 11: DNA-patterns of the *A. niger* ISO-500 host strains having still all marked $\Delta glaA$ loci single in lane 7; a conversion of *Bam*HI marked $\Delta glaA$ locus towards the *Pst*I- $\Delta glaA$ locus (*Bam*HI²⁺, *Pst*I⁻ convertant) in lane 8; a *Bam*HI³⁺/*Pst*I/*Sal*I⁻ convertant in lane 9; a *Bam*HI⁴⁺/*Pst*I/*Sal*I/*Bgl*II⁻ convertant in lane 10; a *Bam*HI⁵⁺/*Pst*I/*Sal*I/*Bgl*II/*Kpn*I⁻ convertant in lane 11.