## F323 RFLP Analysis of the Intergenic Spacer in the rDNA of Fusarium spp.

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The Intergenic Spacer (IGS), which separates rDNA repeat units, appears to be the most rapidly evolving spacer region and closely related species may show considerable divergence in IGS, often reflection both length and sequences variation. Because of the potential for intraand interspecific variation within the ICS, we compared this region of the rDNA from 9 formae speciales of Fusarium oxysporum (Section Elegans) and from eleven species of Fusarium belonging to Section Liseola. Primer CNL12 and CNS1 were used to amplified the IGS region of Fusarium. An amplification product was approximately 2.6 Kb that varied somewhat in size depending on the strain. When PCR products were digested with BamHI, EcoRV. Sphl, Xbal, Hindll, Stul, Sacl and Xhol, there were not observed distinct restriction fragments. Digestion of the amplified ICS with Smal and Hincll showed a very similar restriction patterns among F. oxysporum formae speciales, but different from those of species belonging to Section Liseola. When isolates were digested with Bg/II, KpnI and PstI, the former two enzymes (Bg/II, KpnI) appeared one restriction site in Elegans but no site in Liseola. The latter enzyme Pstl showed one restriction site in section Liseola but no site in section Elegans. In case of EcoRI and Nrul, all strains belong to two sections had similar band patterns which could not distinguish section Elegans from section Liseola. The all strains of Fusarium exhibited different Hinfl, Alul, Haelll, Mspl and Taql patterns. In general, IGS divergence within the section was more variablee than within the specise. Comparision of IGS restriction patterns will provide more phylogenetic information in Fusarium spp.

## F324 Redox-sensitive Modulation of an Anti-Sigma Factor RsrA from Streptomyces coelicolor (RsrA, an anti-sigma factor, regulated by redox change in Streptomyces coelicolor)

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SigR ( $\sigma^R$ ) is a sigma factor in *Streptomyces coelicolor* responsible for inducing *trxBA* gene encoding thioredoxin/thioredoxin reductase upon oxidative stress. We identified an anti-sigma factor for  $\sigma^R$  (RsrA) which binds and inhibits  $\sigma^R$ -directed transcription *in vitro* under reduced conditions. Exposure to  $H_2O_2$  or diamide resulted in dissociation of  $\sigma^R$  from  $\sigma^R$ -RsrA , allowing  $\sigma^R$ -dependent transcription. Thioredoxin was able to reduce RsrA, thus completing the feedback regulatory loop in response to oxidative stress. The result presents a novel example of modulating anti-sigma activity in a redox-dependent manner.