

Natural Populations of Chickpea Rhizobia Evaluated by Antibiotic Resistance Profiles and Molecular Methods

Journal	Microbial Ecology
Publisher	Springer New York
ISSN	0095-3628 (Print) 1432-184X (Online)
Issue	Volume 51, Number 1 / January, 2006
DOI	10.1007/s00248-005-0085-3
Pages	128-136
Subject Collection	Biomedical and Life Sciences
SpringerLink Date	Tuesday, January 03, 2006

PDF (366.1 KB) HTML

Ana Alexandre^{1, 2}, Marta Laranjo^{1, 2} and Solange Oliveira^{1, 2} 

- (1) Departamento de Biologia, Universidade de Évora, Apartado 94, 7002-554 Évora, Portugal
- (2) Laboratório de Microbiologia do Solo, Instituto de Ciências Agrárias Mediterrânicas, Universidade de Évora, Évora, Portugal

Received: 28 July 2005 **Accepted:** 11 August 2005 **Published online:** 3 January 2006

Abstract The aims of this study were to investigate the hypothesis that intrinsic antibiotic resistance (IAR) profiles of chickpea rhizobia are correlated with the isolates site of origin, and to compare the discriminating power of IAR profiles with molecular approaches in rhizobial strain identification and differentiation. Rhizobial diversity from five Portuguese soils was assessed by IAR profiles and molecular methods [16S rDNA restriction fragment length polymorphism (RFLP) analysis, direct amplified polymorphic DNA (DAPD) fingerprinting, and SDS-PAGE analysis of protein profiles]. For each analysis, a dendrogram was generated using the software BioNumerics. All three molecular methods generated analogous clustering of the isolates, supporting previous results on 16S rDNA sequence-based phylogeny. Clusters obtained with IAR profile are similar to the species groups generated with the molecular methods used. IAR groups do not correlate significantly with the geographic origin of the isolates. These results may indicate a chromosomal location of antibiotic resistance genes, and suggest that IAR is species related. DAPD and IAR profiles proved to be the most discriminating approaches in strain differentiation and can be used as fast methods to screen diversity in new isolates.