

## Short notes

### A NEW RACE OF *MELOIDOGYNE CHITWOODI* GOLDEN, O'BANNON, SANTO & FINLEY, 1980?

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Species identification of *Meloidogyne* can be reliably carried out on the basis of the isozyme band patterns of esterase (Est) and malate dehydrogenase (Mdh) in combination with morphological characterization. To obtain these band patterns on a routine scale a fast and accurate method is described by Esbenshade and Triantaphyllou (1985, 1990) using just one single *Meloidogyne* female. This method uses prefabricated mini polyacrylamide gels (4.3 x 5 cm slabs, gradient 10-15 %) on the Phastsystem (Pharmacia LKB), a completely standardized and computerized electrophoresis system.

For screening for resistance and for diagnostic purposes seventeen populations of *Meloidogyne hapla* and *M. chitwoodi* were simultaneously identified with this system at the Plant Protection Service (PD) and the DLO-Centre of Plant Breeding and Reproduction Research (CPRO-DLO). It became evident that five populations of *M. chitwoodi* from locations in the south-eastern part of the Netherlands deviated from the originally described pattern for the Mdh band, while the Est band pattern was absent (Fig. 1). This null allele has been observed in a few occasions within the *Meloidogyne* species (Dalmaso & Bergé, 1983; Esbenshade & Triantaphyllou, 1987). The different band patterns for Est and Mdh were consistent and reproducible and occurred in many of the investigated populations of *M. chitwoodi*. Preliminary morphological studies on these populations indicated that several characteristics can be used for identification of this "new race".

Within *M. chitwoodi* two races are distinguished on the bases of differential host ranges (Santo & Pinkerton, 1985). Field and greenhouse trials showed that maize could be described as a non-host for these deviating populations, whereas maize is a good host for *M. chitwoodi*, type race (Golden *et al.*, 1980). Further host range experiments, morphological studies and biochemical analyses of other isozymes will help to designate the status of these populations in relation to the earlier described races.

#### References

DALMASSO, A. & BERGÉ, J. B. (1983) Enzyme polymorphism and the concept of parthenogenetic species, exemplified by

*Meloidogyne*. In: Stone, A. R., Platt, H. M. & Khalil, L. F. (Eds). *Concepts in nematode systematics*. London, Academic Press : 187-196.

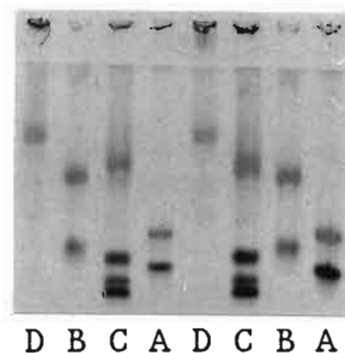
ESBENSHADE, P. R. & TRIANTAPHYLLOU, A. C. (1985). Use of enzyme phenotypes for identification of *Meloidogyne* species. *J. Nematol.*, 17 : 6-20.

ESBENSHADE, P. R. & TRIANTAPHYLLOU, A. C. (1987). Enzymatic relationships and evolution in the genus *Meloidogyne* (Nematoda : Tylenchida). *J. Nematol.*, 19 : 8-18.

ESBENSHADE, P. R. & TRIANTAPHYLLOU, A. C. (1990). Isozyme phenotypes for the identification of *Meloidogyne* species. *J. Nematol.*, 22 : 10-15.

GOLDEN, A. M., O'BANNON, J. H., SANTO, G. S. & FINLEY, A. M. (1980). Description and SEM observations of *Meloidogyne chitwoodi* n. sp. (Meloidogynidae), a root-knot nematode on potato in the Pacific Northwest. *J. Nematol.*, 12 : 319-327.

SANTO, G. S. & PINKERTON, J. N. (1985). A second race of *Meloidogyne chitwoodi* discovered in Washington. *Pl. Dis.*, 69 : 361.



**Fig. 1.** Gradient 8-25 % mini gel showing the original esterase and malate dehydrogenase phenotype of a single *Meloidogyne* female of *Meloidogyne hapla* (A), *M. chitwoodi* (B) and *M. javanica* (C), and the new deviating band pattern of *M. chitwoodi* (D).