Breakthrough in the Identification of the Causal Virus of Pigeonpea Sterility Mosaic Disease

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Sterility mosaic disease (SMD), a potential threat to pigeonpea production in the Indian subcontinent was first reported in 1931. SMD results in annual losses exceeding 205,000 tones of grain (over \$100 million) in India alone. The eriophyid mite Aceria cajani transmits the causal agent under natural conditions. Several attempts to identify the SMD causal agent were not successful and this problem remained enigmatic and posed a big challenge to the scientific community (Reddy et al., 1994; Singh et al., 1999). Recently, using a newly developed protocol a tenui-like virus of asymmetric morphology was isolated from the SMD-affected plants. The virus particles are highly flexuous and branched filaments measuring 3-8 nm in diameter (Plate 1). The virus contains a 32 kDa coat protein and 4-5 segmented RNA of size 0.8 - 3.5 kilo bases. It has been named as pigeonpea sterility mosaic virus (PPSMV) (Kumar et al., 1999). Polyclonal antibodies were produced to PPSMV in rabbit and mice that are now being used to detect the virus in SMD-affected plants by ELISA. This virus was detected consistently from all the SMD-affected samples analysed from several different

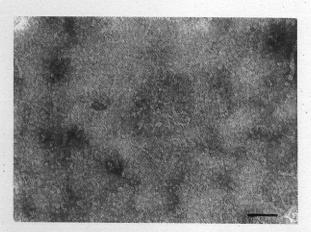


Plate 1. Transmission electron micrograph of the PPSMV particles. Bar represents 100 nm.

locations of the Indian peninsula, and from every one of the pigeonpea plants inoculated with infectious mites (A. cajani) under laboratory conditions. Leaves from over 300 healthy or uninoculated pigeonpea plants gave negative ELISA results. Additionally none of the healthy samples processed by the same protocol as that for virus isolation, contained any virus particles. This is the first confirmatory report of the identification of the SMD causal agent ending seven decades long search for the elusive pathogen. Further this is the first report of isolation of a tenui-like virus from dicotyledonous plants, transmitted by eriophyid mites.

With the availability of data on virus identity and diagnostic tests, it should be possible to screen pigenopea genotypes for virus resistance precisely and device molecular tools for distinguishing rapidly various biotypes of the PPSMV.

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