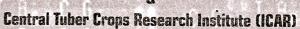


# Abstract of papers

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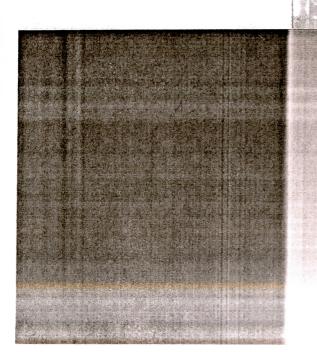


Indian Phytopathological Society (South Zone)



Venue: Central Tuber Crops Research Institute (ICAR)

Thiruvananthapuram 695 017, Kerala



ally distinct isolates of BBrMV.

## morphic DNA analysis among the various urica causing powdery mildew diseases in

#### m and J. Rajalakshmi

ımil Nadu Agricultural University, Coimbatore- 625 104

leaves showing symptoms of powdery mildew disrmer's fields in different chilli growing areas of Ta-I pathogens were isolated from the infected leaves. l as Leveillula taurica based on morphological and ulence of all the fungal isolates was determined in n on detached chilli leaf assay. The measurement of aves revealed differences in the virulence between riability among isolates of L.taurica that differed by means of random amplified polymorphic DNA primers. Analysis of the genetic coefficient matrix RAPD profile showed that minimum and maximum the L.taurica were in the range of 12 to 100% reusing unweighted pair-group method with arithmetic separated the isolates into three clusters (I, II and III) ersity among the isolates of L.taurica from chilli. e isolates namely Lv 4, Lv 10 and Lv 6 and Cluster Lv 2 and the cluster I have Lv1, Lv 3, Lv 5 and Lv 9

#### PG-P3

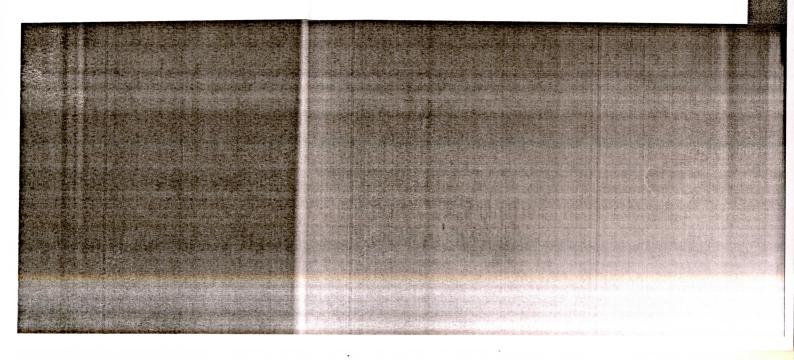
Recombination and negative selection pressure in the P1 gene of Indian Sugarcane streak mosaic virus

Abstracts

### K. Bagyalakshmi, B. Parameswari, C. Chinnaraja, R. Viswanathan

Plant Pathology Section, Division of Crop Protection, Sugarcane Breeding Institute (ICAR), Coimbatore Email: rasaviswanathan@yahoo.co.in

Sugarcane streak mosaic virus (SCSMV), a member of the genus Poacevirus, of this family, an important viral pathogen affects the sugarcane production in India. The genome has a single open reading frame that is translated in to a large polypeptide and consequently cleaved into functional proteins. This virus causes mosaic of sugarcane along with the Sugarcane mosaic virus (SCMV) which is a serious disease causing varietal degeneration reported from India in 1999 and later has been reported from geographically different Asian countries. The coding region for P1 peptidase is located at the very beginning of the viral genome of the family Potyviridae. P1 was thought of as serine peptidase with RNA-binding activity and with possible influence in cell-to-cell viral spreading. In order to unveil its mechanism of evolution we initiated the study by characterizing 10 P1 gene of Indian isolates and the sequences were compared with previously reported SCSMV isolates from different countries. Comparison of all of the sequenced virus isolates revealed a high level of diversity in the P1 gene (83-98% nt sequence identity; 87-100% aa sequence identity), and the Indian isolates were found to be the most divergent (up to 9% variation at the amino acid level). Phylogenetic analysis revealed clustering of 17 SCSMV isolates into two groups. Group I included isolates from India (except SCSMV-TPT) and Pakistan, and group II consisted of isolates from Japan, Indonesia, Thailand and SCSMV-TPT. The results obtained from phylogenetic study were further supported with the SNPs (single nucleotide polymorphism), INDELs (insertion and deletion) and evolutionary distance analysis. A significant proportion of recombination sites were found at the N terminal region of I'l gene of Indian isolates. Analysis of selection pressure indicated that the P1



Abstracts

gene of Indian SCSMV isolates is under strong negative selection. It is likely that recombination, along with strong negative selection, enhances the speed of elimination of lethal mutations in the P1 gene of Indian SCSMV isolates.

#### PG-P4

Assessing the incidence of grain discolouration and seed mycoflora load in rice varieties

C. Jeyalakshmi, R. Renuka and C. Rettinassababady

Department of Plant Pathology, Pandit Jawaharlal Nehru College of Agriculture & Research Institute, Karaikal – 609 603, U.T. of Puducherry E-mail: csjayal@yahoo.co.in

Experiments were conducted to assess the grain discolouration (%) and seed mycoflora load in the popularly grown rice varieties of the Cauvery delta region of U.T. of Puducherry. Ten representative panicles of ADT 39, ADT 43 ADT 46, CR 1009, KKLR1, Samba Mahsuri and White ponni collected ran domly at harvest from Pandit Jawaharlal Nehru College of Agriculture and Research Institute farm during February 2013 and per cent grain discoloura tion was assessed as per IRRI, (1998). Seed mycoflora assay was carried ou and observations on fungal growth were recorded and frequency occurrenc of mycoflora was calculated. The results revealed that the variety White Por ni was statistically different from other varieties by recording the maximur grain discoloration of 40 per cent. The variety CR 1009 recorded the lower incidence (13.00%). Eight mycoflora viz., Aspergillus niger, A. flavus, Curvi laria lunata, Fusarium moniliforme, Helminthosporium oryzae, Microdoch um oryzae, Penicillium sp. and Rhizopus'sp were found to be associated wit the grain discoloration. Of which, C. lunata and H. oryzae were present all test varieties. The presence of the aflatoxicosis fungi Aspergillus sp. wa noticed in all test varieties except ADT 39, however it harboured Rhizopus s The study on the distribution of mycoflora in the grain discolouration revea that Penicillium sp. is the most frequently isolated fungus with 30.57 per ce distribution followed by C. lunata (18.85%), H. oryzae (17.14%) and F. mon. iforme (10.00%). The A. niger and A. flavus were less frequently distribute

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