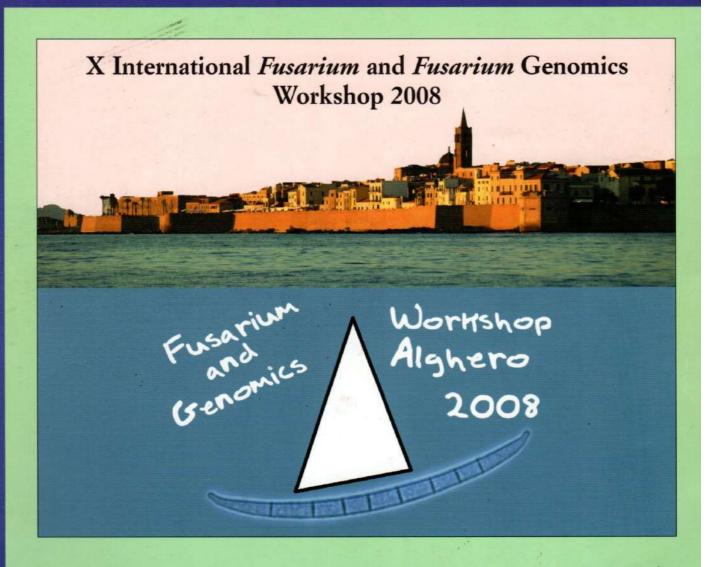
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Abstracts of presentations

Alghero, Sardinia (Italy), August 30 - September 2, 2008



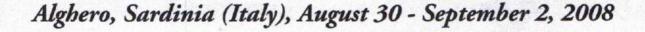
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X International Fusarium and Fusarium Genomics Workshop 2008





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X International *Fusarium* Workshop and *Fusarium* Genomics Workshop 2008

Hotel Carlos V, Alghero, Sardinia (Italy), August 30 - September 2, 2008

Fusarium Committee, ISPP:

H. Corby Kistler, USDA-ARS Cereal Disease Laboratory, St. Paul, USA Ulf Thrane, Technical University of Denmark, Kgs. Lyngby, Denmark

Scientific Committee:

Marie-Josèe Daboussi, Université Paris-Sud, Orsay, France
Harold Corby Kistler, USDA-ARS Cereal Disease Laboratory, St. Paul, USA
Antonio Logrieco, CNR - ISPA, Bari, Italy
Brett Summerell, Royal Botanic Gardens, Sydney, Australia
Ulf Thrane, Technical University of Denmark, Kgs. Lyngby, Denmark

Local organizers:

Virgilio Balmas, Quirico Migheli, University of Sassari, Sassari and Istituto Nazionale di Biostrutture e Biosistemi, Italy





Program schedule: .

Saturday 30 August

13.00-14.30: registration of participants

14.30-15.30: Keynote address (Marie-Josèe Daboussi and Lester W. Burgess)

15.30-17.30: Systematics and phylogenetics (session leaders: Kerry O'Donnell, Keith Seifert)

17.30-17.45: ISPP Fusarium Committee Meeting

17.30-19.30: poster session and Welcome party

Sunday 31 August

08.30-10.00: Genomics (session leaders: Li-Jun Ma, Corby Kistler)

10.00-10.15: coffee break

10.15-13.15: Host-fungal interactions (session leaders: Willie Schäfer, Antonio Di Pietro)

13.15-14.30: lunch

14.30-16.00: Biological control (session leaders: Claude Alabouvette, Angelo Garibaldi)

16.00-17.30: Ecology and biogeography (session leaders: David Geiser, Brett Summerell)

17.30-17.45: coffee break

17.45-18.00: ISPP Fusarium Committee Meeting

18.00-18.30: Fusarium Genomes Research Policy Group Meeting

18.00-19.00: poster session

Monday 1 September

08.30-10.45: Mycotoxins and metabolism (session leaders: Naresh Magan, Ulf Thrane)

10.45-11.00: coffee break

11.00-13.15: Diagnostics (session leaders: Kamel Abd-El Salam, Theo van der Lee)

13.15-14.30: lunch

14.30-16.45: Disease management (session leaders: Rafael M. Jiménez-Diaz, Wade Elmer)

16.45-17.00: ISPP Fusarium Committee Meeting

17.00-18.00: poster session and coffee break

18.00: sightseeing tour and evening banquet

Tuesday 2 September

09.30-11.45: Pathogenicity and disease epidemiology (session leaders: Paul Nicholson,

Sukumar Chakraborty)

11.45: coffee break and departure

12.00-13.00: Meeting of the International Society of Plant PathologyInternational Mycological Association Subcommittee on Fusarium Systematics (chair David M. Geiser)

14.00-17.00: NordForsk Network Meeting (chair Tapani Yli Mattila)

yields. Field and laboratory pathogenicity tests with the two species showed that they were pathogenic, that more than one fungus is associated with this disease, their prevalence varying with location and season, and that these *Fusarium* species reproduced similar symptoms on the same rice hosts. Generally dwarf cultivars were more susceptible than the taller ones.

FUSARIUM WILT OF CHICKPEA AND PIGEONPEA AND THEIR MANAGEMENT IN SEMI-ARID TROPICS. S. Pande¹, M. Sharma¹, P.M. Gaur¹, K.B. Saxena¹, C.L.L. Gowda¹, O. Gupta², L. Kaur³, M.S. Sangwan⁴, R.N. Chaudhary⁵, B.M. Jamadagni⁶, D.R. Saxena² and H.K. Ramappa. ¹International Crops Research Institute for the Semi-Arid Tropics (ICRISAT), Patancheru, Andhra Pradesh. ²Department of Plant Breeding & Genetics, JNKVV, Jabalpur. ³Department of Plant Breeding, Genetics and Biotechnology, PAU, Ludhiana, Punjab. ⁴Department of Plant Pathology, CCSHAU, Hisar. ⁵Indian Institute of Pulses Research, Kanpur. ⁶MPKV, Rahuri, Maharashtra. ¬RAK College of Agriculture, Sehore, Madbya Prades. ®UAS, GKVK, Bangalore, India. E-mail: s.pande@cgian.org

Fusarium wilt of chickpea (Cicer arietinum L.) caused by Fusarium oxysporum f. sp. ciceris (FOC) and pigeonpea (Cajanus cajan (L.) Millsp. caused by F. udum (FU) are the major production constraints to these legumes worldwide, specifically in the Indian subcontinent and Eastern Africa (chickpea and pigeonpea) and the Mediterranean region (chickpea). In India alone these wilts cause up to 100% crop losses on susceptible cultivars under favorable environmental conditions. Host plant resistance offers the most sustainable and effective wilt management option either alone or as a major component of integrated disease management. Effective greenhouse and field screening techniques have been developed at the International Crops Research Institute for the Semi-Arid tropics (ICRISAT), Patancheru, India to identify resistance in the germplasm and to incorporate resistance in breeding lines of chickpea and pigeonpea. More than 17,000 lines of chickpea and 12,000 lines of pigeonpea have been screened using these techniques at ICRISAT. Significant positive correlation was found between seedling screening in greenhouse and adult plant screening in the wilt-sick fields. Chickpea and pigeonpea lines found resistant to Fusarium wilt at ICRISAT were further evaluated through International Chickpea Wilt and Root Rot Nursery (ICWRRN) and International Pigeonpea Wilt Nursery (IPWN) to identify stable and broad based resistance at hot spot locations in Asia and Africa. Several breeding lines have shown stable and durable resistance to Fusarium wilt of chickpea and pigeonpea, and many improved wilt resistance varieties have been adopted by farmers. The differential wilt disease reaction in few lines in multilocation testing indicated variability in FOC and FU populations.

FUSARIUM LANGSETHIAE INFECTION AND MYCOTOXIN CONTENTS OF OATS AND BARLEY. P. Parikka¹, V. Hietaniemi², S. Rämö² and H. Jalli¹. ¹MTT Agrifood Research Finland, Plant Production Research, Finland. ²MTT Agrifood Research Finland, Laboratories, 31600 Jokioinen, Finland. E-mail: paivi.parikka @mtt.fi

In the field trial of 2004-2006, the first Fusarium species, detected at panicle emergence of oats, was F. langsethiae. It was found also on barley at ear emergence, but the amount of infected kernels was not so high as on oats. F. langsethiae was the most common Fusarium species on oats during the early development of kernels. The other species detected at early stages in flowers

and kernels of oats was F. poae. Later in the season, other Fusarium species infected kernels and the prevalence of F. langsethiae in harvested grain was low. In 2006, however, infection by species like F. avenaceum and F. culmorum was inhibited in dry conditions and F. langsethiae was fairly abundant in harvested, dried grain. It was present both in oats and barley grown in traditionally tilled and direct drilled areas. As a whole, more infection was seen on oats than on barley and more in late cultivars than in early ones. The prevalence of F. langsethiae varied: in 2004 it was slightly more prevalent in tilled than in direct drilled areas but in 2006 in dry conditions direct drilling seemed to produce more infected kernels and grain than tillage. F. langsethiae seems to be a more important producer of T2 and HT-2 toxins than F. sporotrichioides in Finland. The analysed contents of these mycotoxins were higher on oats (max 950 µg/kg) than on barley (max. 710 µg/kg). In 2006, the T2+HT-2 contents of oats were higher in direct drilled than in tilled plots.

FUSARIUM SPECIES ASSOCIATED WITH VANILLA STEM ROT IN INDONESIA. A. Pinaria¹, L.W. Burgess¹ and E.C.Y. Liew². ¹Faculty of Agriculture, Food and Natural Resources, The University of Sydney, NSW 2006, Australia. ²Botanic Gardens Trust, Mrs Macquaries Road, Sydney, NSW 2000, Australia; Email: apin3761@mail.usyd.edu.au



Indonesia is one of the world's leading producers of vanilla, an important crop offering high economic returns to small-holder farmers. A major constraint in vanilla production in Indonesia is stem rot disease, which has caused significant economic losses over the last decade. Previous reports of vanilla stem rots in the Asia-pacific region include those caused by Fusarium, Colletotrichum, and Phytophthora species. In this paper, we report Fusarium species associated with the disease. Seven major vanillaproducing provinces were surveyed for disease incidence. Isolates were obtained from diseased stem tissues using selective media. Pure cultures were subcultured onto CLA and PDA for species identification. A total of 542 Fusarium isolates were recovered, comprising 7 species, namely F. decemcellulare, F. oxysporum, F. proliferatum, F. pseudograminearum, F. semitectum, F. solani, F. subglutinans, and 14 isolates of undescribed species. F. oxysporum was most commonly isolated from all the areas surveyed, followed by F. solani and F. semitectum. Of the species tested in pathogenicity studies, only F. oxysporum was shown to be pathogenic to vanilla. Further studies to investigate genetic diversity of the pathogen and host resistance are underway.

ASSESSMENT OF ENDOPHYTIC COLONIZATION OF SORGHUM BICOLOR SEEDLINGS BY GIBBERELLA ZEAE. S.A.J. Quazi¹, L.W. Burgess² and J. Smith-White². ¹Plant Pathology Division, Bangladesh Rice Research Institute, Gazipur-1701, Bangladesh. ²Faculty of Agriculture, Food and Natural Resources, The University of Sydney, NSW 2006, Australia. E-mail: q.shireen@bdonline.com

The susceptibility of grain sorghum (Sorghum bicolor) to colonization by Gibberella zeae was assessed by isolation studies involving plants grown in the glasshouse. The glasshouse studies on infection of sorghum seedlings by G. zeae indicated that this pathogen can infect sorghum at early growth stages and gradually colonize adjacent tissues as an endophyte. The results also showed that roots as well as stalk tissues are susceptible to infection. The results suggested that the fungus could infect and colonize the proximal parts of roots more aggressively than the leaf sheaths and