National Plant Disease Recovery System (NPDRS)

2020 Research Progress Reports
March 1, 2021

Tim Widmer, USDA-ARS (tim.widmer@usda.gov)



The NPDRS Program

 Called for in Homeland Security Presidential Directive Number 9 (HSPD-9) which was issued in February of 2004

- Oversight by ARS
- Consists of two parts:
 - Recovery Plans:
 www.ars.usda.gov/crop-production-and-protection/plant-diseases/docs/npdrs/
 - Research



Dr. Jo Anne Crouch

Mycology and Nematology Genetic Diversity and Biology Laboratory:

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<u>Problem</u>: Tools & resources to accurately identify Downy Mildews of cereals/grasses very limited, especially the foreign pathogens**





^{**}The PI has requested a hold on viewing their presentation at this time. If you would like further information, please contact them directly.

Dr. Christopher Dunlap

Crop Bioprotection Research Laboratory: Peoria, IL christopher.dunlap@usda.gov





Novel approaches to controlling laurel wilt and Fusarium dieback in avocados.





Dr. Christopher Dunlap,

Crop Bioprotection Research Unit National Center for Agricultural Utilization Research,

In collaboration with our state partners and local growers



Dr. Daniel Carrillo, TREC- University of Florida







Research and Accomplishments

• Evaluated and developed entomopathogens for use against ambrosia beetles, which are now routinely used by growers during phytosanitation procedures.



- Identified antifungal antagonists against the plant pathogens. Identified two new Paenibacillus species and B. velezensis.
- Developed experimental foam formulations to improve delivery of biocontrol agents to the target.

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Dr. William Rutter

U.S. Vegetable Laboratory: Charleston, SC william.rutter@usda.gov



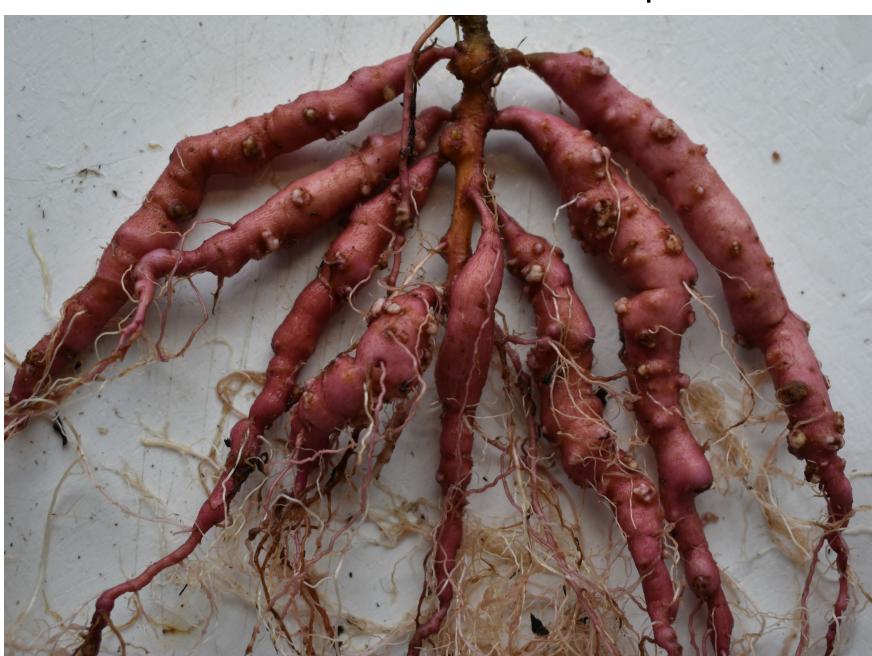


Developing resources to manage Guava Root-Knot Nematode (*Meloidogyne enterolobii*)

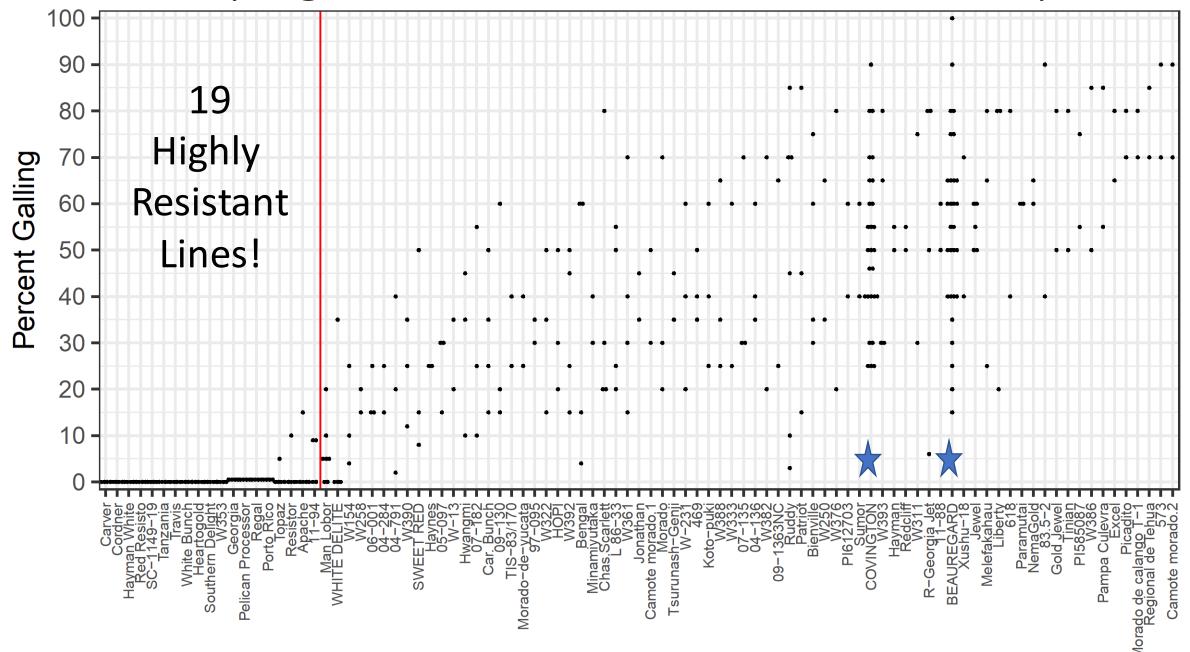
William Rutter
USDA-ARS
Charleston, SC

Identifying M. enterolobii resistant Sweetpotato

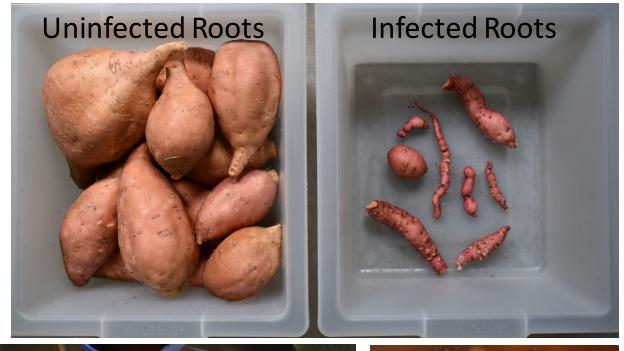




Identifying M. enterolobii resistant Sweetpotato



A new Method for Detecting M. enterolobii in Sweetpotato



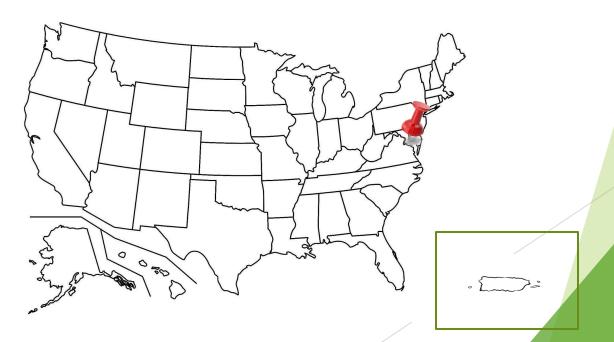




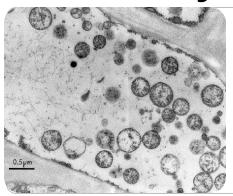


Dr. Yan Zhao and Dr. Wei Wei Molecular Plant Pathology Laboratory: Beltsville, MD yan.zhao@usda.gov wei.wei@usda.gov



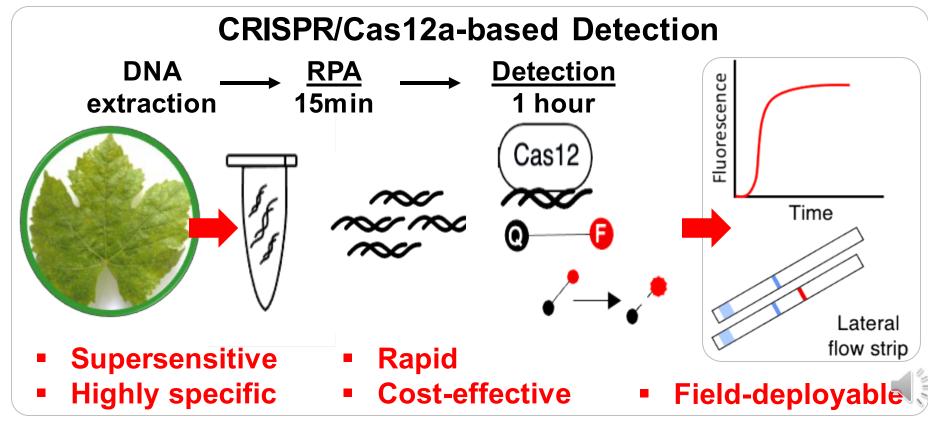


Developing CRISPR/Cas12a-based Technologies for Early Detection of Exotic Phytoplasmas



Phytoplasmas

- Phloem-inhabiting bacteria without a cell wall
- Cause numerous diseases in agricultural crops
- Six species are quarantine pests



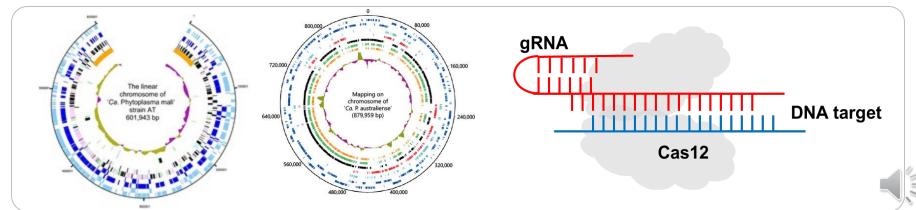
Developing CRISPR/Cas12a-based Technologies for Early Detection of Exotic Phytoplasmas



Apple proliferation disease caused by 'Ca. Phytoplasma mali' (Photo credit: Osler/DBADP, Univ. of Udine, Italy)



Grapevine yellows disease caused by 'Ca. Phytoplasma solani' (Photo credit: Fabio Quaglino, Univ. of Milan, Italy)



Dr. Scott Adkins U.S. Horticultural Research Laboratory: Ft. Pierce, FL scott.adkins@usda.gov





Tomato Resistance-Breaking Tospoviruses

Scott Adkins, USDA-ARS, Fort Pierce, FL Ozgur Batuman & Salih Yilmaz, UF, SWFREC, Immokalee, FL Tong Geon Lee & Samuel Hutton, UF, GCREC, Wimauma, FL



TSWV and **TCSV** management by resistance



TSWV R gene:

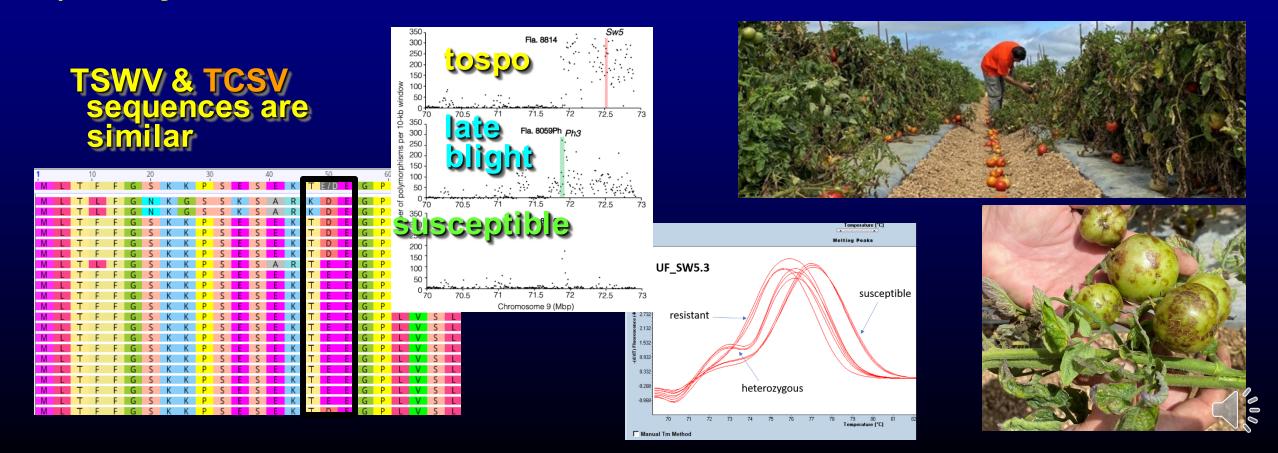
identified in 2016 for the first time in California



Tomato Resistance-Breaking Tospoviruses

Planting Sw-5 tomato in Florida and Puerto Rico for TSWV & TCSV:

- 1) Tools/protocols for detecting resistance-breaking TSWV & TCSV
- 2) Tools/protocols for detecting tomato tospovirus resistance genes
- 3) Analyze resistant tomato cultivars in Florida and Puerto Rico



Dr. Kai Ling

U.S. Vegetable Laboratory: Charleston, SC kai.ling@usda.gov





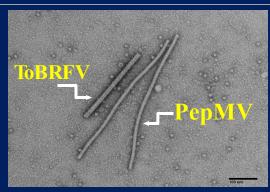
Managing emerging Tomato brown rugose fruit virus through sensitive detection, effective disinfection and improved immunity

-Kai-Shu Ling, USDA-ARS, Charleston, SC-











Objectives:

- ➤ To evaluate selected disinfectants for field application and seed treatment.
- To optimize real-time PCR useful for seed health assays.
- ➤ To conduct genetic and genomic analysis of novel resistance to ToBRFV in tomato.

Progress and Accomplishments

- 1. Effective disinfectants identified and recommended to growers.
- 2. Developed a sensitive Real-time PCR useful for seed health testing.
- 3. Screened tomato germplasm and identified new sources of resistance to ToBRFV.

Need for research

- 1. Improving methods of application for effective disinfectants to achieve disease management in tomato productions.
- 2. Understanding the genetic basis of resistance and molecular marker development.
- 3. Developing genetic materials that are useful in tomato breeding for resistant to ToBRFV.

Publications:

Chanda, B., Shamimuzzaman, M., Gilliard, A., and Ling, K.-S. 2021. Virology Journal 18:7.

Chanda, B., Rivera, Y., Nunziata, S.O., Galvez, M.E., Gilliard, A.C., Ling, K.-S. 2020. Microbiology Resource Announcements. 9(29):e00630-20.

Dr. Bob Jarret

Plant Genetic Resources Conservation Unit: Griffin, GA bob.jarret@usda.gov

Screening the ARS Capsicum
Germplasm Collection for
Tobamoviruses**





^{**}The PI has requested a hold on viewing their presentation at this time. If you would like further information, please contact them directly.

Dr. Ricardo Goenaga

Tropical Crops and Germplasm Research Laboratory: Mayaguez, P.R. ricardo.goenaga@usda.gov





Highly sensitive qPCR screening of the U.S. cacao germplasm collection in Puerto Rico for presence of the emergent Cacao mild mosaic badnavirus (CaMMV) previously restricted to the Trinidad Gene Bank collection.

INTRODUCTION:

- The NPGS living cacao germplasm collection comprises strategically representative accessions belonging to the 10 genetic cacao groups, and is the only world collection planted in an RCBD.
- During 1943, two virus-like symptom phenotypes were observed for the first time in cacao trees in Trinidad and Tobago and symptoms attributed to two strains of a suspect viral disease, referred to as Trinidad cacao virus (TCV) disease.
- These strains represent two new badnaviruses, named *Cacao mild mosaic virus* (CaMMV) and *Cacao yellow vein banding virus* (CYVBV), making them the first cacao-infecting badnaviruses known in the Western Hemisphere.
- During 2019-2020, symptoms reminiscent of TCV, now known to be associated with CaMMV or CYVBV, were observed in a commercial field in Puerto Rico, and several clones housed in the USDA germplasm collections in Mayaguez and Miami quarantine greenhouse.
- Quantitative PCR (qPCR) and PCR assays that target the viral movement protein (MP) have been developed based on CaMMV and CYVBV sequences (1 genome each), the only available isolates. A SNP genotyping tool has been developed to identify accessions by genetic group affiliation, and genotype analysis of virus-infected trees is underway to aid in identifying germplasm with apparent resistance or tolerance, and potentially, link historical origins of CaMMV with genetic group.

OBJECTIVES:

- Screen the USDA cacao germplasm collection in Mayaguez, PR, and the cacao quarantine greenhouse in Miami, FL and any symptomatic trees observed in the Trinidad and CATIE-Costa Rica collections for CaMMV/CYVBV using qPCR to determine badnavirus distribution and simultaneously guide replacement of infected clones with uninfected trees, and removal of infected commercial trees;
- 2. Confirm identity of badnaviral species in representative samples found positive for CaMMV or CYVBV presence by PCR amplification, cloning, and DNA sequencing of MP;
- 3. Genotype cacao samples using cacao SNPs analysis found to be CaMMV or CYVBV-infected



Highly sensitive qPCR screening of the U.S. cacao germplasm collection in Puerto Rico for presence of the emergent Cacao mild mosaic badnavirus (CaMMV) previously restricted to the Trinidad Gene Bank collection.

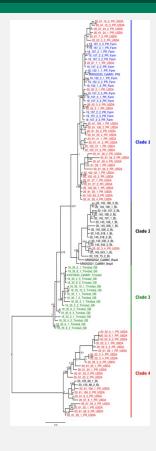
RESULTS AND DISCUSSION

- 1. Screen USDA cacao germplasm collection in Mayaguez, PR, Miami cacao quarantine greenhouse, and Trinidad cacao genebank for CaMMV/CYVBV.
- a. Amplification with qPCR primers designed to detect CaMMV-Trinidad (only known isolates);
- b. Positive samples verified CaMMV by PCR amplification with CaMMV-T original MP primers; **Results**: Phylogenetic analysis revealed not one but three CaMMV clades (*Fig. 1*). **Conclusion**: There is genomic variability among Mayaguez, Brazil and Trinidad CaMMV isolates.
- c. Re-designed and validated modified MP-PCR primers to account for all CaMMV variants: **Results**: Re-designed primers amplified all CaMMV variants. **Conclusion**: Development of a much more powerful test for CaMMV.
- d. Re-run Miami & Mayaguez samples again with previously re-design primer combination (to account for false negatives in 1st run). **Results:** Positive detection was greater with new re-designed primer combination. Is the virus endemic? TBD.
- e. Validation of CYVBV-T with 8 isolates from Trinidad (2020). Primer testing was halted due to budget constraints.
- 2. In the Quarantine Miami greenhouse: selected samples from plants showing virus-like symptoms were subject to Illumina DNA sequencing and RNAseq: Results: No CaMMV /badnavirus detected. In DNA reads, no DNA-virus found; in RNA reads: polerovirus-like sequences detected. Whether representative of a complete polerovirus genome will require further study. New virus??
- 3. SNP discovery analysis using the raw sequencing data from collection genotypes is pending additional funding. Do cacao genetic groups differ in their susceptibility to badnaviruses?

IMPACT OF DELIVERABLES

- 1) Developed/validated improved qPCR and PCR detection of CaMMV and three groups of variants, are now recognized through this research (Fig. 2).
- 2) Development of highly useful assay for virus-indexing and managing USDA cacao germplasm collections.
- 3) Positive impact on improving current regulatory testing capacity to aid APHIS-PPQ decision-making.

Figure 1. Phyloge netic analysis of CaMMV movement protein (MP) resolved three additional groups of clades of the virus, a result that agreed with the results of pairwise distance analysis (see Clades 1-4). Sequences from the current studyare highlighted in blue = commercial farms, Puerto Rico), red = USDA germplasm collection, Mayaguez, Puerto Rico), or green = Gene Bankin Trinidad.



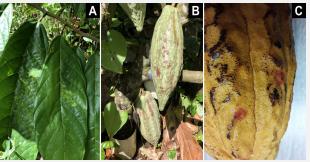


Figure 2. Ca MMV-like symptoms in Puerto Rico.

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National and international collaborations to proactively combat threats to U.S. cotton. Partnered with Pakistani scientists to mitigate the effects of Cotton Leaf Curl Virus (CLCuV) a

virus transmitted by whiteflies.



Infected plants (left) and resistant (right)

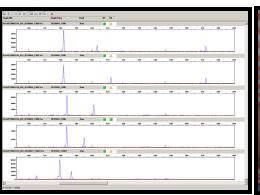
Developed advanced cotton lines.
Provided seed to U.S. public
breeders and breeding companies
and breeders in China, India, Brazil
and Australia. Seed is available on
request.

In 2021, more improved resistant elite cotton lines will be available.



Created diagnostic tests to detect then identify and track the virus.

Have already used the tests on new viruses found in the U.S., but so far not found.

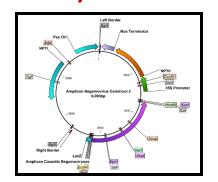




DNA markers for marker assisted selection. Can breed without having the disease present and can also decrease the number of years for development from ten years to six!







Using RNAi, CRISPR and other emerging technologies to develop plants with their own self protection mechanisms.

Emerging Threat Cotton Leaf Roll Dwarf Virus (CLRDV). In 2017, a variant of this virus, which causes Cotton Blue Disease in Brazil, was detected in the U.S. Took advantage of Brazil's experience to get a jump on tackling CLRDV in the U.S.







By 2019 the virus was found in ten states across the cottonbelt.



Phylogenetic study of CLRDV samples collected across the cottonbelt, indicates a single main point of origin that spread into other states.

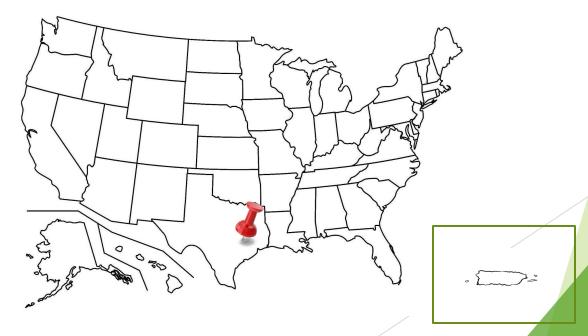


Cotton Inc. organized a Task Force to tackle CLRDV

- Research to understand the best ways to mitigate the effects of CLRDV
- Develop cheap and effective diagnostic tests
- Identify resistant cotton germplasm, so far best candidates have come from the USDA Cotton Germplasm Collection and originated in Africa
- Breed resistant cotton varieties

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Resistance to Fusarium wilt Race 4 (FOV4) identified from the U.S. National Cotton Germplasm Collection



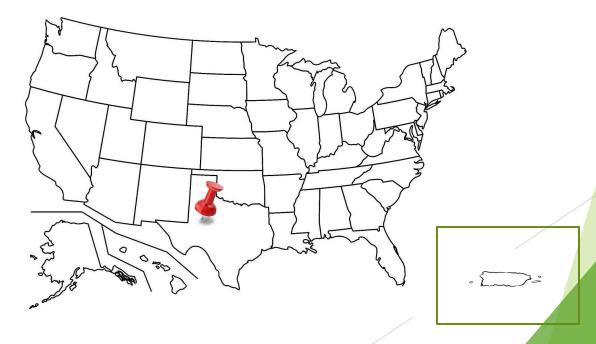
Figure 1. A. FOV4 in San Joaquin Valley (photo credit: Dr. Bob Hutmacher, University of California). B. FOV1 in Georgia (photo credit Dr. Peng Chee, University of Georgia). C. FOV4 in El Paso County Tx (photo credit Dr. Jason Woodward)

- ~10,000 accessions are available in the Collection
- ~2,000 accessions screened by multiple collaborators using differing methods to search for FOV4 tolerance/resistance
- ONE (1) accession identified as tolerant/resistant to FOV4 in highly replicated field studies

Dr. Mauricio Ulloa

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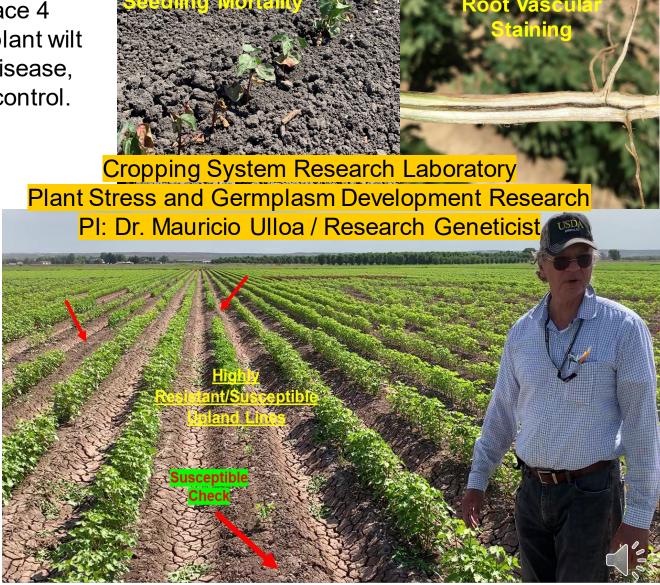


Unraveling population shift of Fusarium wilt race 4 (FOV4) fungi-strains and their interaction for overcoming tolerance/resistance in cotton

➤ The Fungus Fusarium oxysporum f. sp. vasinfectum race 4 (FOV4) adversely impacts cotton production causing plant wilt and death. FOV4 is an inoculum density-dependent disease, and, so far, resistant cultivar is the only management control.

Previous and Ongoing Research

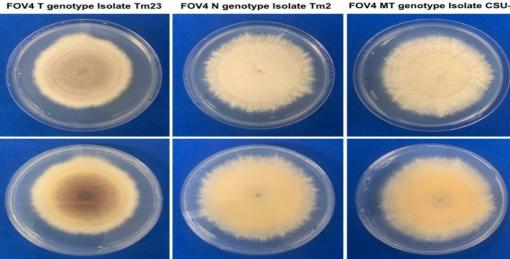
- Identified new sources of FOV4 resistance in Upland and Pima cotton
- ➤ Developed new Upland and Pima cotton progeny or breeding lines with high resistance to FOV4.
- ➤ Publicly release Upland and Pima germplasm lines with high resistance to FOV4.
- Preliminary characterization of current Fusarium oxysporum f. sp. vasinfectum isolates from cotton in the San Joaquin Valley of California and El Paso Texas Valley.



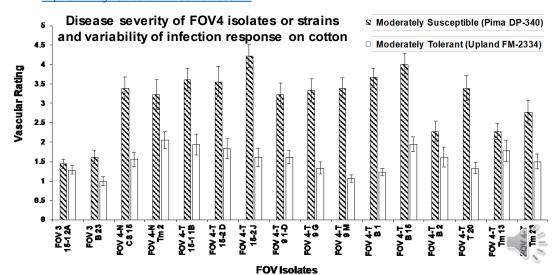
- With the support of the NPDRS program, we are going to further identify and characterize emerging devastating *Fusarium* strains in different soil types and infected cotton plants from evaluation sites in California and Texas

 Simple variability can be observed on growth and pigmentation
- ➤ So far, a preliminary FOV molecular characterization study identified diversity/variability within FOV4 strains and confirmed the presence of two (T and N) strains of FOV4 (VCG 0014i) in California cotton fields and possibly population shift in some fields due to selective pressure towards the N strains
- ➤ Seedling assays to assess pathogen infection reveals differences in isolate's virulence or aggressiveness and the complexity of FOV4, wilt disease and plant resistance mechanisms.
- ➤ Overall, results from this and previous studies suggest that the FOV4 populations identified to date in far-west Texas vary from those found in California.
 - DNA sequencing or blueprint-information generated from this project will help us to identify and distinguish these new FOV4 variants and to develop new diagnostic tools and assays for Soil FOV Quantification and Breeding Resistant Varieties.

Simple variability can be observed on growth and pigmentation when fungus grows on media

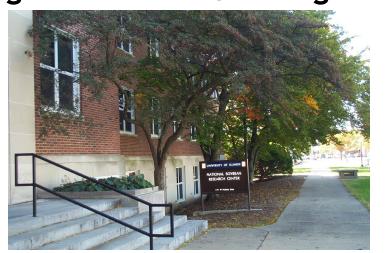


Diaz, J., Garcia, J., Lara, C., Hutmacher, R. B., Ulloa, M, Nichols, R. L, Ellis M. L. 2020. Characterization of current *Fusarium oxysporum* f. sp. *vasinfectum* isolates from cotton in the San Joaquin Valley of California and Lower Valley B Paso Texas. Plant Disease Journal. https://doi.org/10.1094/PDIS-05-20-1038-RE



Dr. Glen Hartman

Soybean/Maize Germplasm, Pathology, and Genetics Research Unit: Urbana, IL glen.hartman@usda.gov





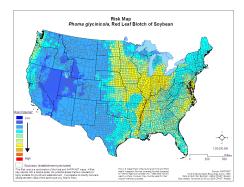
An Exotic Foliar Fungal Disease - Red Leaf Blotch of Soybean

Background

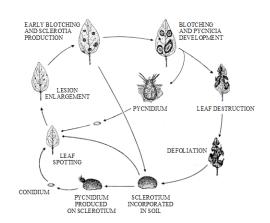
- The disease occurs only in Sub-Saharan
 African countries on soybean and a native legume plant, Neonotonia wightii
- Based on the APHIS Recovery Plan developed in 2009, distribution in the USA if introduced could become widespread from the south to the northern areas of production

Symptoms and signs

- Leaf lesions are most noticeable (lesions on petioles, pods, and stems also occur; not known to be seed borne)
- Lesions progress from small red-brown spots to larger leaf blotches
- Older lesions contain fungal structures including sclerotia (survival structures) and pycnidia (spore-producing structures)







Issues/problems

- Yield losses of 50% reported in several African countries
- With more recent information about the disease and conducive epidemic environments, the potential risk to soybeans in the USA needs updating even though there is a lack of detailed research on distribution, spread, fungal variability, and management in Africa

Disease management

- Fungicides may be effective for controlling RLB, but questions about efficacy, application timing, and registrations remain unknown
- Biological and cultural management may aid in alleviating disease pressure, but research in this area is lacking

Evaluation of Soybean Genotypes for Resistance to RLB

- Until now, there has been little concerted effort to evaluate soybean genotypes for resistance to RLB
- One report screened soybean germplasm in the USA (ARS Foreign Disease-Weed Science Research Unit) and concluded that all 23 soybean genotypes tested (representing 90% of the genes present in public U.S. materials) were susceptible

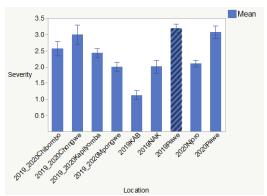
Research Objective

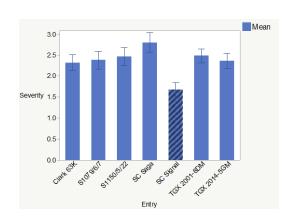
 Evaluate red leaf blotch resistance in the field in breeding lines and cultivars developed in Africa

Acknowledgements

- ARS National Plant Disease Recovery System
- Soybean Innovation Laboratory (SIL)
- Field cooperators with SIL







Materials and Methods

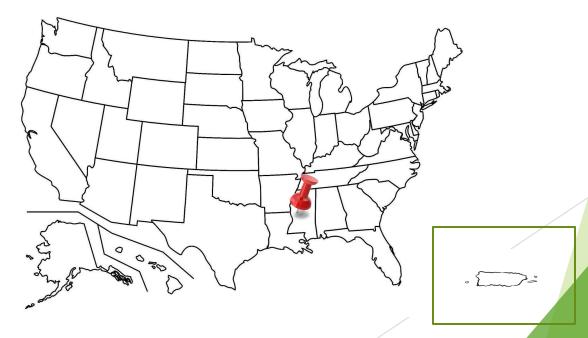
- Nine locations were evaluated for RLB resistance in the Pan African Trials (PAT); the PAT is part of a USAID Feed the Future Initiative funded project to the Soybean Innovation Lab
- A total of 69 entries were evaluated; all locations had a minimum of 30 entries
- Seven entries were common among locations

Summary

- Disease severity differed by local and entry
- Top bar graph shows Uganda with the lowest level of disease; Ethiopia with the highest
- Lower bar graph shows cultivar SC Signal with the lowest severity (commercially available cultivar developed in Zimbabwe)
- Our mandate is to continue to screen more germplasm to find sources of resistance that are more complete

Dr. Shuxian (Susan) Li Crops Genetics Research Unit: Stoneville, MS shuxian.li@usda.gov





Recovery and Preservation of *Phakopsora pachyrhizi C*ollection from Southern US: Meeting the Needs for Soybean Rust Research

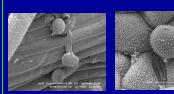


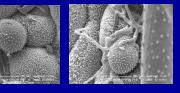
Shuxian Li

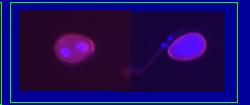
USDA, ARS, Crop Genetics Research Unit Stoneville, Mississippi 38776, USA

- Soybean rust (SBR) is one of the most economically important soybean foliar diseases caused by the fungal pathogen *Phakopsora pachyrhizi*.
- Yield losses from 13% to 90% due to SBR have been reported.
- **SBR** has the potential to cause major economic damage to U.S. soybean production.
- The causal agent has a broad host range, an airborne nature, and diverse and complex virulence pattern.
- Developing soybean cultivars with resistance to different isolates of *P. pachyrhizi* are needed to combat the emerging potential threat of soybean rust in the U.S.









- Collections of plant pathogens are crucial for phytopathological research and breeding for host resistance to diseases.
- Objective for phase I research: Recover stored P. pachyrhizi isolates
- □ Accomplishments during the pandemic in 2020:
- Fifteen isolates were recovered using the heat-shock method (Bonde et al. (2006) or Tween-water soaking technique.
- Isolate MS 06-1 was used to identify the *Rpp6* gene in 2012.
- A Florida isolate was reported to defeat *Rpp 1* and *Rpp6* genes previously.
- Ongoing research:
- Confirm the pathogenicity of recovered isolates by inoculating a panel of differentials with known resistance (*Rpp*) genes and susceptible plants.
- Set up long-term storage.







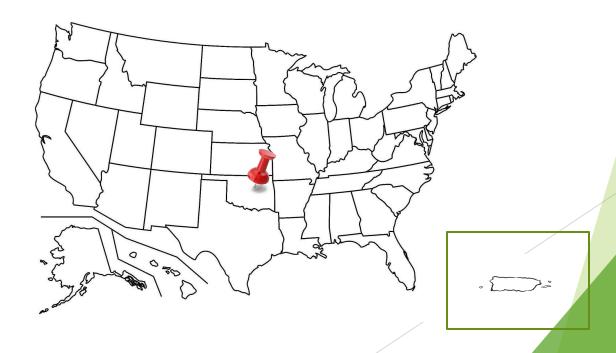


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Dr. Kelly Chamberlin

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Breeding for Resistance to Peanut Smut

What is peanut smut?

- Fungal disease of peanuts
- No above ground symptoms; Nuts (seed) replaced with fungal spores
- No control method; Highly dispersive and easily spread
- Not found outside of Argentina and Brazil

Why are we worried about peanut smut?

- 100% incidence in Argentina, up to 50% yield loss
- Threat to global peanut production (\$47 B in 2020) and thus U.S. peanut production (\$1.13 B in 2019)
- U.S. stakeholder requested action

What is the plan?

- USDAARS scientists are working with INTA, Argentina scientists to:
 - Identify sources of smut resistance through screening
 - Incorporate smut resistance into peanut for the U.S.
 - Identify and deploy markers for selecting resistance
 - Develop high-throughput methods for phenotyping and detection
 - Determine the variability of pathogen population

Accomplishments to Date:

- Identified 13 sources with near immunity over 3 years of phenotyping U.S. germplasm in Argentina
- Conducting crosses to incorporate smut resistance into cultivars for production in the U.S.
- Advancing breeding populations for resistance screening and cultivar development
- Resistance marker development nearly complete

Impact

- Protection of the U.S. peanut crop, valued at \$1.13 B in 2019
- Resistant germplasm available to peanut breeding programs
- Molecular markers for use in peanut breeding programs





Dr. Shyam Tallury

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Development and validation of a molecular diagnostic assay for the detection of peanut clump virus (PCV) and Indian peanut clump virus (IPCV)**





^{**}The PI has requested a hold on viewing their presentation at this time. If you would like further information, please contact them directly.

Dr. Shahryar Kianian Cereal Disease Lab: St. Paul, MN shahryar.kianian@usda.gov





Understanding and combating increased virulence to oat crown rust (*Puccinia coronate* f. sp. *avenae*)

Shahryar F. Kianian

USDA-ARS Cereal Disease Lab



- ✓ Oat crown rust is the most widespread and damaging disease of oat worldwide.
- ✓ Moderate to severe epidemics can reduce grain yield by 10 to 40%.
- ✓ In 2014 yield losses due to this disease in Minnesota was reported at 50% and in South Dakota as 35.

Oat crown rust isolates are becoming substantially more virulent

"Resistance is futile"?

- ➤ 100+ seedling resistance genes identified in oat
- 3 years effective lifespan
- innumerable races
- Low apparent fitness cost to virulence
- No negative associations between virulences
- Abundant sexual recombination on common buckthorn

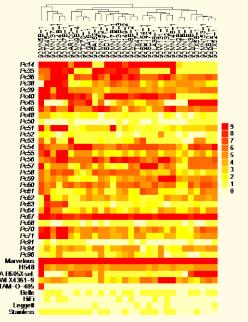
Entry 191- BT1021-1-1 X O/CI1712-5-131



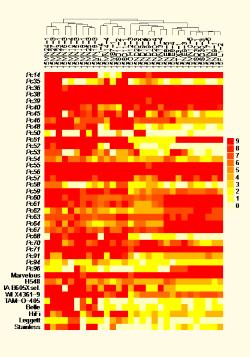
Entry 147- BT1020-1-1 X O/CI8000-4-66



1990 Pca Isolates



2015 Pca Isolates



Dr. Les Szabo
Cereal Disease Lab: St. Paul, MN
les.szabo@usda.gov





Genotyping of Critical Wheat Stem Rust Pathogen Stains. Les J. Szabo, St. Paul, MN

Wheat Stem Rust

- Historically the most devasting disease of wheat.
- Caused by the fungal pathogen *Puccinia* graminis.
- Recently re-emerged as serious threat to wheat production in Africa, Western Asia and Southern Europe.
 - Ug99 race group (Africa)
 - TKTTF race group
 - TTRTF race group
- These new strains of *P. graminis* are a threat wheat production in the U.S.

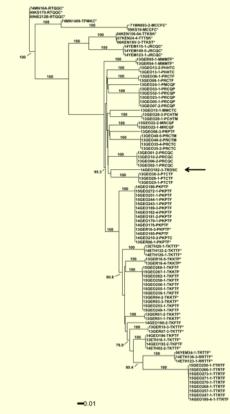




Infected wheat field in Kenya

Genotyping of Critical Wheat Stem Rust Pathogen Stains. Les J. Szabo, St. Paul, MN

- Objective: Develop genotyping tools and characterize U.S. and global population.
 - Developed high-throughput SNP chips.
 - Genotyped over 2,500 samples.
 - Defined the major genetic groups in the U.S. and global populations.
- Objective: Develop rapid molecular diagnostic tools.
 - Developed rapid molecular assays for the U.S. and global populations.
 - Assays are being used as part of an international surveillance program for wheat stem rust pathogen.



										Pg	t Core	SNP	Marke	ers						
				A101	A105	A106	A107	A109	A111	A119	A121	A126	A127	A128	A129	A130	A131	A133	A134	A13
olate	RACE	Clade	MLG	C/A	C/T	T/G	T/C	G/C	A/G	T/C	A/G	G/A	G/A	C/G	T/C	T/C	G/A	G/T	C/A	C/1
GEF R1 56/04	THESE	1	C=-AB1	CA	СТ	TG	TC	GC	AG	cc	AG	GA	AA	CG	cc	TC	GA	GT	CA	TT
XET108-3	JMCQC		Co-AB2	CA	CC	GG	CC	GC	GG	cc	AG	AA	AA	CG	cc	TC	GG	GT	CA	TT
TEMB4-1	TRUTTE	■ A	Cu-AD3	CA	TT	TG	TC	GC	GG	TC	AA	GA	GA	GG	TC	TC	GA	П	CA	TT
U Œ0129-1	TIMIF		C=-AD4	CA	CT	TG	CC	CC	GG	CC	AA	AA	AA	GG	CC	TC	GG	GG	CA	TT
SETH18-1	TETTE	N-A.1	Cu-ADS	CC	CC	π	TT	GG	GG	TC	AG	GA	AA	GG	TC	TT	GG	GG	CC	CT
KE1107-3	TRITE	N-A.2	Co-ADC	CA	CT	TG	TC	GC	GG	TC	GG	GG	GA	CC	π	TT	GG	GG	CC	CT
ETHD4-1	TETTE	N-E	Co-AD7	AA	CT	TG	TC	CC	AA	π	AG	GA	GA	GG	TC	TT	GG	GG	CC	CT
HE 106-1	PKPTF	N-C	C=-ADS	CA	CT	TG	TC	GC	AG	П	AG	GG	AA	CG	TC	TT	GA	GG	cc	CC
GEII01-1	TKKTF	IV-D	O=-AD9	CA	CT	TG	TC	GC	AG	TC	AG	GA	GA	CG	TC	TT	GA	GT	CA	CT
HEERLE_5	THEFT	N-E.1	Co-ALD	AA	CC	π	TC	CC	AG	TC	AA	GA	GA	GG	cc	TC	GA	GG	cc	CT
HEE 1003-1	THEFT	N-E.2	O= A11	AA	CT	TG	TC	CC	AG	TC	AA	GA	AA	CG	cc	TC	AA	GG	CA	CT
GE 0129-1	TOTE	₩-F	C=-A14	CC	CT	TG	TC	CC	GG	TC	AG	GA	AA	CG	TC	TC	GA	GG	cc	CT
GE104-1	MAME	Clarke V-A	C=-T1D	AA	TT	TG	CC	CC	AA	cc	AG	GG	AA	GG	CC	TC	GG	GT	CC	CT
GE 015-2	MMCTC	Clarke V-B	O=-T11	CA	TT	П	CC	CC	AG	TC	AG	GA	AA	CG	TC	TC	GA	GT	CC	CT
SD 19a	HFLG	WA	O⇒-A15	CA	TT	TG	CC	GC	GG	CC	AA	GA	AA	GG	TC	TC	AA	TT	AA	TT
THD SZA	ROUS	W-E	On-A16	AA	CT	GG	TC	CC	GG	cc	AG	AA	AA	CG	cc	CC	GA	GT	AA	TT
185 60 a	QFCS	W-C	C=-A17.d	AA	TT	GG	CC	CC	GG	cc	AA	AA	GA	GG	cc	CC	AA	TT	AA	TT

Dr. Bob Bowden

Hard Winter Wheat Genetics Research Unit: Manhatta

Hard Winter Wheat Genetics Research Unit: Manhattan, KS robert.Bowden@usda.gov





Breeding Resistant Wheat Varieties to Stem Rust Using Marker-Assisted Selection

Robert Bowden and Mary Guttieri, USDA-ARS, Manhattan, KS Amir Ibrahim, Texas A&M University, College Station, TX



Summary of Problem: Wheat crops in the US are threatened by new exotic virulent races of stem rust, such as the Ug99 group. New durably resistant germplasm is needed that also has elite agronomic characteristics.

Research Objectives:

- 1. Determine rust resistance reactions to stem rust, stripe rust, and leaf rust diseases for advanced breeding lines in the Southern Plains at the Castroville, TX hot-spot for wheat rusts.
- 2. Develop high-yielding, high-quality winter wheat germplasm lines adapted to the Great Plains region with durable resistance to stem rust, stripe rust, and leaf rust diseases.

Relevance of the Objectives to NPDRS Program Priorities

Wheat is the most widely grown crop in the Great Plains region. New races of all three rust species threaten the sustainable production of wheat crops. Durably resistant varieties are within reach through cooperation of USDA-ARS, Texas A&M University, and others.



PROGRESS REPORT

Castroville Screening Nursery, 2021



Lines Screened

• Winter Wheat: 16,334

Spring Wheat: 2,421

• Oats: 1,548

• Barley: 114

Germplasm Development



Durable, Quantitative Resistance Breeding

- Four lines with durable SR resistance from 'Kingbird'
- Sr2, Sr9b, Sr12, Sr57, Sr58

Gene Pyramid Breeding

- Elite backgrounds: TAM114, TAM205, TX13M5625, Smith's Gold, Showdown, KS090387K-20
- Sr22, Sr26, Sr35, Sr38, Sr57, Sr58

Dr. David Marshall

Plant Science Research Unit: Raleigh, NC david.marshall@usda.gov





Stacking Ug99 Stem Rust Resistance with Other Resistances in Wheat – David Marshall, Raleigh, NC

IMPORTANCE:

Rusts and other wheat diseases are global problems.

Genetic resistance is the most cost-effective disease management for farmers.



- Identify effective, long-lasting resistance to three wheat rusts.
- Combine those resistances with resistance to powdery mildew and Fusarium head blight.
- Breed those combined resistances into diverse wheat backgrounds and market classes.





















RESULTS

APPROACH:

- National & International
- Controlled screening
- Molecular markers
- Phenotyping & genotyping
- Breeding & selection

Dr. Matthew Rouse
Cereal Disease Lab: St. Paul, MN
matthew.rouse@usda.gov





Rouse: Wheat resistance to Ug99

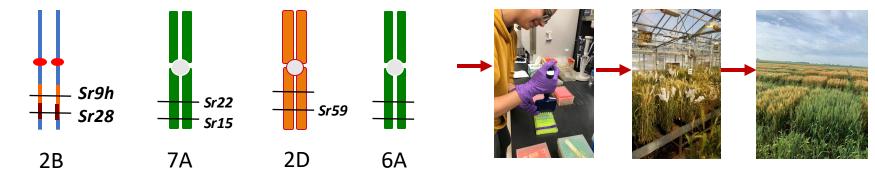
2007 Ug99 Resistance Loci

- 19 resistance genes,
- 5 with markers

2020 Ug99 Resistance Loci

- 43 resistance genes
- 39 with markers

Linkage Blocks of Ug99 resistance genes



- Marker-assisted backcrossing
- BC₄-derived fixed +/- lines
- Yield trials in collaboration with breeders

- Bolles
 - Advance

Linkert

- Forefront
- Lang-MN Prevail

Linkert

- UMN **SDSU**
 - **SDSU**

7 lines

NDSU

6 lines

UMN

Field Screening of Spring Wheat and Barley in Africa



- 1500+ spring wheat lines
- 200+ spring barley lines
- 7 US universities
- 4 private companies
- **Recent resistant varieties** 'MN-Torgy', 'SY-Longmire'

Impact



UMN variety 'Linkert' with Ug99 resistance – 833,900 acres in Minnesota and North Dakota in 2018

Dr. Yue Jin
Cereal Disease Lab: St. Paul, MN
yue.jin@usda.gov





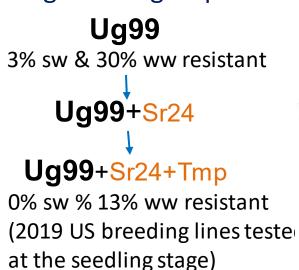
Stem rust surveillance to monitor Ug99 and other significant virulence

Y. Jin, USDA-ARS Cereal Disease Lab, St. Paul, MN

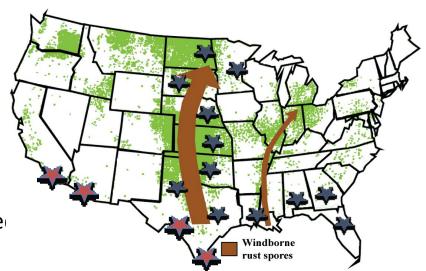
Stem rust of wheat:



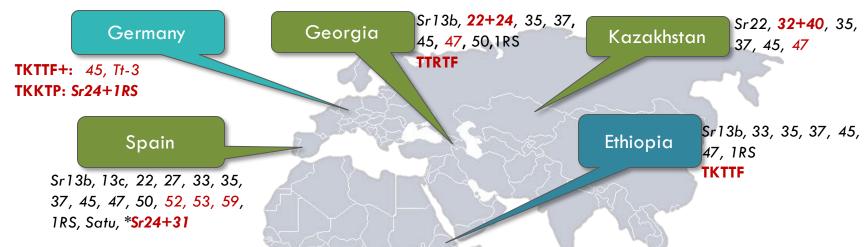
Ug99 race group:



Ug99 sentinel plots in US:



Detection of non-Ug99 races that have caused epidemics/outbreaks: (in close collaborations with D. Luster and P. Olivera)



Dr. Doug Luster

Foreign Disease - Weed Science Research Unit: Fort Detrick, MD doug.luster@usda.gov





Preliminary screening and storage of exotic cereal rust accessions in BSL3 containment

Douglas G. Luster Foreign Disease-Weed Science Research Unit, Fort Detrick, MD

Cereal rust pathogens are evolving new forms and threatening the world's food security

Stem rust of wheat:



- Foreign rust collections are received during the growing season at the ARS Ft. Detrick MD BSL-3 Plant Pathogen Containment Facility, recovered, screened and archived.
- Selected accessions are shipped during the Winter Containment season to the ARS Cereal Disease Laboratory, St. Paul MN for intensive screening, pathotyping and genotyping
- 2019 to 2020: over 320 sample receipts from 9 countries in Africa, Europe, and Asia recovered, screened and archived.

NPDRS FY 2018 Funding: "Rapid detection of *Phytophthora ramorum* (Sudden Oak Death) in nursery irrigation water"

Douglas G. Luster Foreign Disease-Weed Science Research Unit, Fort Detrick, MD

Problem: Detection of Sudden Oak Death in infested nurseries takes 10 days to 2 weeks

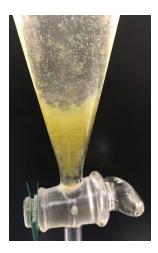
Mock Nursery Irrigation Retention Pond

National Ornamental Research Site Dominican University, San Raphael CA

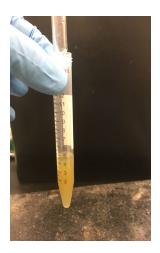


Rapid Sampling Strategy:

Flocculation of *P. ramorum* propagules from water samples







Objective: Rapid, efficient recovery and detection of *Phytophthora ramorum* from nursery irrigation water in 24 hrs

Mr. Gary Peterson

Foreign Disease - Weed Science Research Unit: Fort Detrick, MD gary.peterson@usda.gov





Wheat blast

Wheat blast is an emerging fungal disease once limited to South America but recently reported in Bangladesh and Zambia with disease Losses of 30-100%.

Wheat blast has not been reported in North America

USDAARS is currently screening U.S. wheat germplasm for disease resistance ARS Biosafety Level-3 Plant Containment Facility at Fort Detrick Maryland.

ARS Foreign Disease Weed Science Research Unit (FDWSRU) in Cooperation with Kansas State University and Asociación de Productores de Oleaginosas y Trigo (ANAPO) established a wheat blast screening nursery in La Paz, Bolivia

NPDRS is providing funding for the testing of U.S. wheat lines, identified at FDWSRU as potentially resistant in both the field and a greenhouse in Bolivia where new strains of the pathogen may exist.



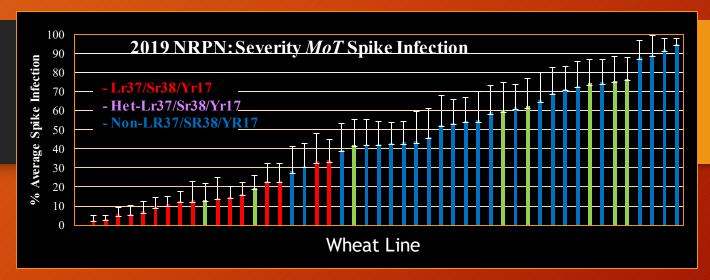


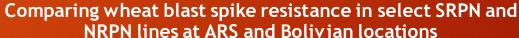
NPDRS: Wheat Blast

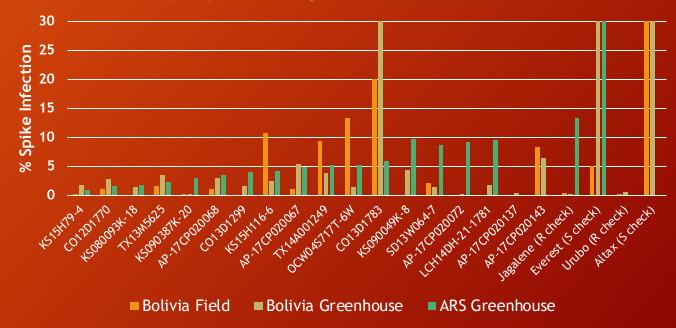
Of 93 lines from the U.S. SRPN and 102 NRPN lines tested, a total of 46 lines were shown to be resistant (<10%) to wheat blast at FDWSRU.

Of 38 potentially resistant lines sent to Bolivia, 20 were also resistant in Bolivian greenhouse and field studies.

Seed of 20 lines from SRPN, NRPN, Uniform Southern and Eastern Soft Red Wheat and several BASF lines are currently in Bolivia being prepared for planting this season







National Plant Disease Recovery System (NPDRS)

Tim Widmer, USDA-ARS (tim.widmer@usda.gov)

