Coffee Leaf Rust (CLR) in Hawaii: Fungal Attacker Revealed

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Coffee Leaf Rust (CLR)

- Pathogenicity
 - Disease Cycle
 - Biology/Development/ Genetics
- Management
 - Fungicides
 - Varieties
 - Natural Enemies



CLR Teamwork!









Growers & Producers



Lionel Sugiyama



Eva Brill



Karma Kissinger

And Many Others



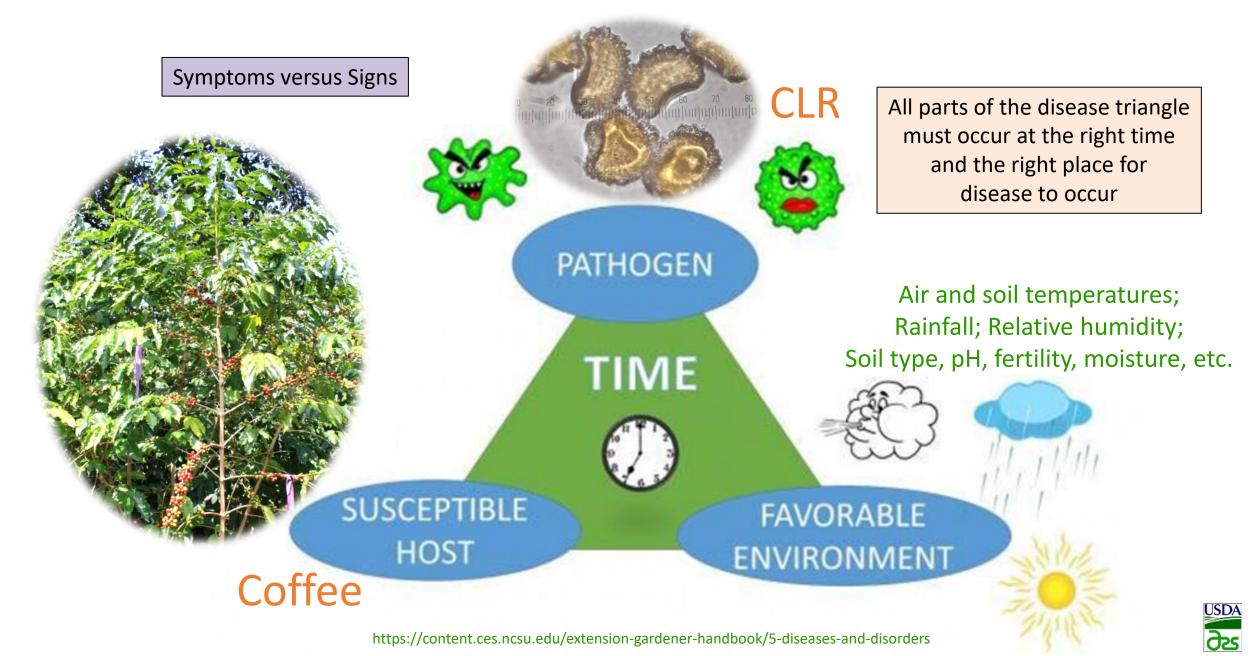
Pepe Miranda



Pathogenicity Testing



How is Disease Caused in Plants?



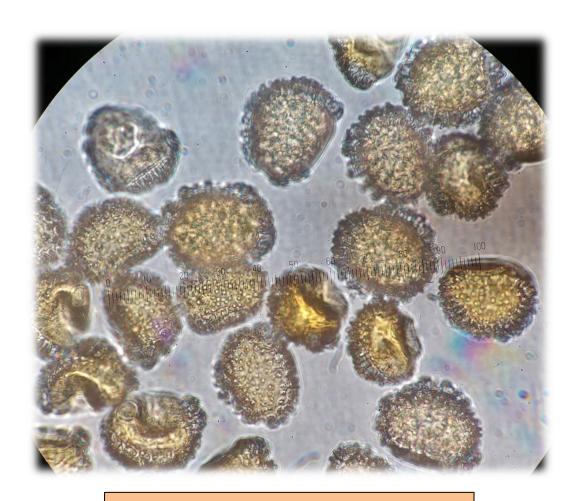
Pathogenicity Testing

- Collect fresh spores
- Adjust concentration to 1 x 10⁵ spores/ml
- Brush on/spray on inoculum (Water control); Typica
- Growth chamber
- 2 days in the dark/ high humidity
- 12 light/12 dark
- Observe symptoms/ signs



Coffee Leaf Rust: Prove Pathogenicity

- Re-isolate spores and confirm by morphology and molecular ID
 - Obligate parasitic fungus
 - can't be cultured
 - Infection through stomata
 - Dark
 - High humidity
 - 22°C / ~72°F



Magnification of the urediniospores of coffee leaf rust (x400)





CLR Biology & Development



Coffee Leaf Rust: Symptoms & Signs



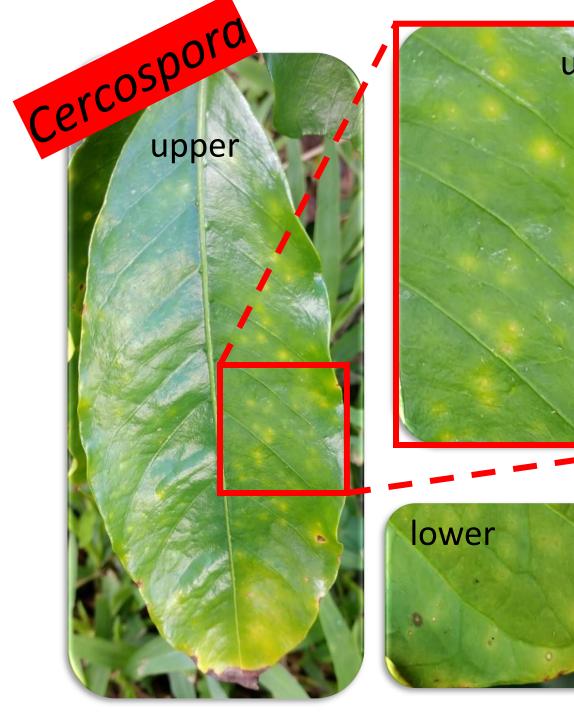




- Under optimum conditions with susceptible variety
- First observable symptoms: small, pale yellow spots on the upper surfaces of the leaves (~21 days)
- Spots gradually increase in size; spores appear on the lower leaf surface
- Lesion centers become necrotic (~35 days)
- Infected leaves fall off (~70 days)



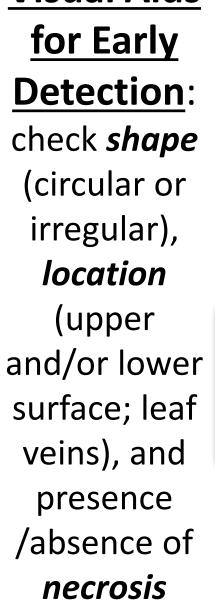




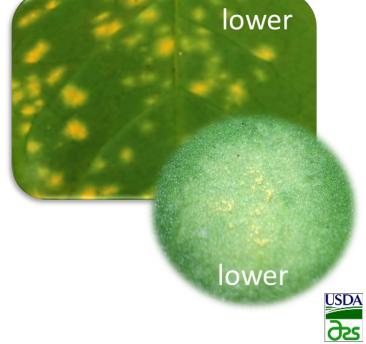




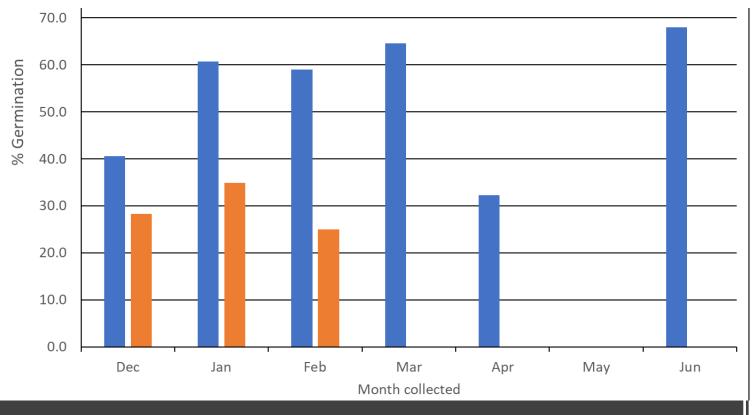
CLR Visual Aids











Stage of Infection/Environment Affects Spore Viability



Coffee Leaf Rust: Symptoms & Signs

CLR impact on wild coffee in Kona

- Lesions can develop anywhere on the leaf
- First spots can appear on any leaf in the canopy
 - Initial stages of the epidemic; windblown spores
- Multiple infections visible on the same leaf
- Typica cotyledons can become infected





 Wild: leaves drop prematurely, leaving twigs with no leaves; regrowth observed







Characterization of CLR in Hawaii

- ARS PBARC Lab: Basic molecular ID
 - All Hawaii isolates are identical
- Genotyping (Dr. Aime, Purdue U.) PURD

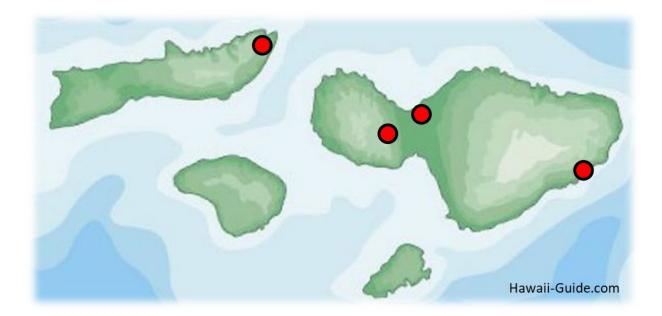


- Molecular analysis/Geographical location
 - Central America and the Caribbean: Jamaica, Panama, Guatemala, El Salvador, and Honduras
- Single introduction
- Race identification (Dr. Vítor Várzea & Dr. Maria do Céu Silva, University of Lisbon, Portugal)
 - Screening a set of coffee plant differentials that contain various resistance genes



CLR Early Detection (DLNR/MISC)

Active spore traps: battery powered; on timers; can be used directly for molecular ID



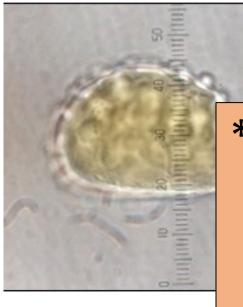
Maui: DLNR Baseyard; Koali; Hanaula: Oct 2019 to Dec 2020 (Feb/Aug 2020)

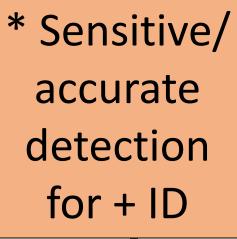
Molokai: Puu O Hoku: June 2020 to Mar 2021

CLR was not detected in any spore trap using PCR Increased sensitivity using nested PCR approach = 2 CLR +







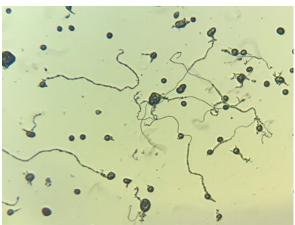






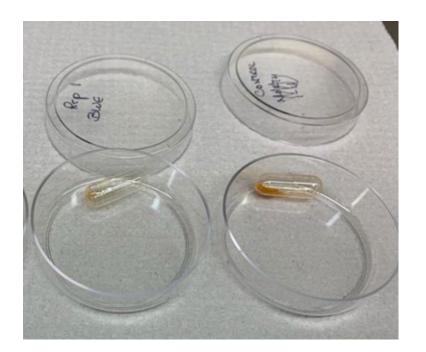








USDA



Laboratory and Field Fungicide Testing

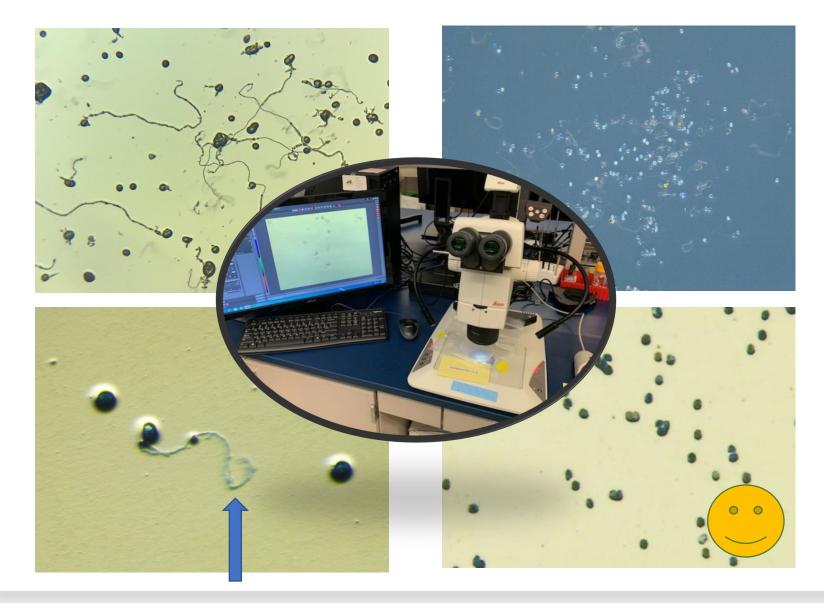
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In Vitro Fungicide Assays

Method

Reg. for use: 7
Julie/UH IR-4: 7
Additional: 10+

*Direct contact/timing











Sanitation Protocols Developed

- 70%, 80%, 95% Ethanol
- 70%, 91% IPA
- 10% Freshly Prepared Bleach Solution

Field Fungicide Trial

- Badge X2
- Kocide 3000-O
- Cueva
- DoubleNickel
- Serenade
- Oxidate 2.0







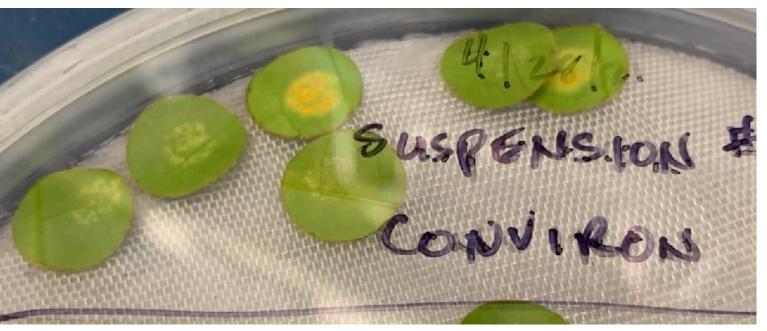


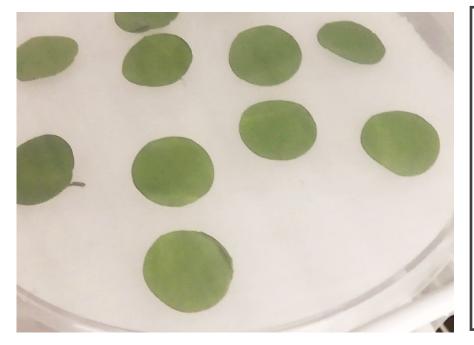


Variety Testing (HARC)







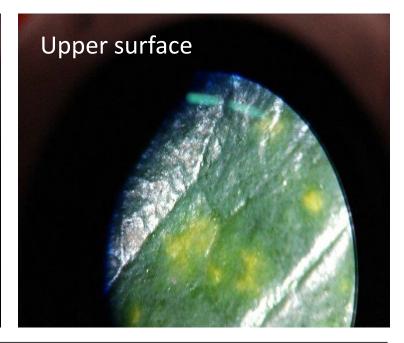


Typica v. Obata











Potted Tupi & Obata





Natural Enemies of CLR



Mycoparasites









Combating CLR in HI

- Reduce % incidence and severity
- Improve/maintain plant and soil health
- Scout regularly for disease symptoms
 - Early and accurate detection
- Weed control and pruning
- Practice good hygiene and sanitation
- Manage as a continuous epidemic
- Short- & Long-term strategies
- No silver bullet/Use integrated approach



