P A T H O L O G Y

SECONDARY NUTRIENT APPLICATIONS TO MANAGE CHARCOAL ROT IN SOYBEAN

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Charcoal rot of soybean is an important disease that occurs annually and can limit profitable soybean production. The estimated yield losses as a result of charcoal rot are comparable to other soybean root-associated diseases including sudden death syndrome and seedling disease. Nationally, approximately 400

million bushels of soybean were lost to charcoal rot over the past 20 years. Macrophomina phaseolina (Tassi) Goid, the fungus responsible for causing charcoal rot, is a ubiquitous soilborne pathogen that affects over 500 hosts including rotational hosts such as corn, cotton and grain sorghum. Charcoal rot is typically more devastating in hot dry years in non-irrigated situations, but can also be problematic in irrigated fields when stressed. Symptoms of charcoal rot can be visible at vegetative stages, but are predominately observed during reproductive stages especially once plants set pods (> R3). the season. The symptoms associated with charcoal rot include interveinal chlorosis, stunting, wilting, and premature plant death. Currently, management practices suggest reducing stress and planting resistant germplasm; however, a general lack of complete resistance is available within the commercially available germplasm

so the management options for charcoal rot remain extremely limited.

In prior research, a relationship between the concentrations of calcium and magnesium in plant tissue and the incidence of charcoal rot was observed. Between 2014 and 2016, non-irrigated field trials were conducted at the Delta Research and Extension Center with a susceptible cultivar, Pioneer P46T21R, and a moderately resistant cultivar, Pioneer P49T80. Treatment applications consisted of 1,000 pounds per acre of calcium and magnesium alone and in combination at pre-plant, at-plant, and

"ANNUALLY, CHARCOAL ROT ACCOUNTS FOR SIGNIFICANT YIELD LOSSES THROUGHOUT MISSISSIPPI, ESPECIALLY DURING YEARS WHERE THE ENVIRONMENT IS DOMINATED BY HOT AND DRY CONDITIONS. HOWEVER, MANAGING CHARCOAL ROT IS MADE DIFFICULT SINCE RESISTANT VARIETIES ARE RARELY AVAILABLE AND FUNGICIDE TREATMENT IS INEFFECTIVE."

Tom Allen

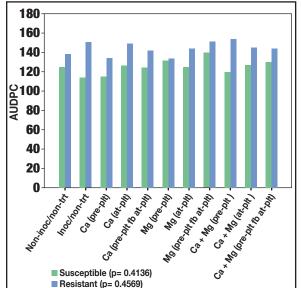
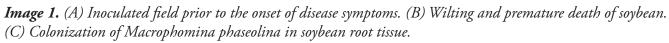


Figure 1. AUDPC calculated from disease severity assessments from root tissue throughout the season.

pre-plant followed	Ave
by at-plant compared	
1 1	Ave Treat Non Inoc Calc Calc Calc Calc Calc Mag Mag Mag Calc
secondary nutrients). In-season evalu- ations, including stand count, vigor, and plant height were taken at vegetative	(Calc (a Calc (r P-va

Moderately Resistant Varie	ties	tible and	tures the multiple ratings conducted
Treatment (timing) Non-Inoculated untreated Inoculated untreated Calcium 1,000 lb/ac (pre-plant) Calcium 1,000 lb/ac (at-plant) Calcium 1,000 lb/ac (pre-plant fb at-plant) Magnesium 1,000 lb/ac (pre-plant) Magnesium 1,000 lb/ac (at-plant) Calcium 1,000 lb/ac + Magnesium 1,000 lb/ac (pre-plant) Calcium 1,000 lb/ac + Magnesium 1,000 lb/ac (at-plant) Calcium 1,000 lb/ac + Magnesium 1,000 lb/ac (pre-plant) P-value for F-statistic	Susceptible 56.3 c 64.4 ab 63.8 ab 64.2 ab 63.5 ab 65.4 a 64.3 a 62.7 ab 60.2bc 61.5 ab	Moderately Resistant 59.1 58.8 55.9 58.1 58.2 59.0 59.5 59.1 57.6 56.7 56.8 0.7732	during the season. Yield was collected from all plots and post-harvest evalua- tions were conduct- ed to determine 100 kernel weight and % germination. Colony forming units (cfu) were used to quan- tify <i>Macrophomina</i> <i>phaseolina</i> coloni- zation from ground root tissue. Collec- tively, environmental

and reproductive stages. Root samples were taken beginning at pod initiation (R3) through maturity (R8). Disease was visually assessed in root tissue using a 1-5 scale (1= no visible microsclerotia and 5= completely colonized). These values were used to calculate the area under the disease progress curve (AUDPC) which capconducive for charcoal rot development in any of the three years. Although no significant benefit was observed between treatments when compared to control treatments, minimal numerical differences occurred in disease severity (Figure 1.) and yield (Table 1.).





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