


# Role of *Rhizoctonia solani* AG 3-PT inoculum source on disease development on potatoes in South Africa

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(A. Pitman, New Zealand)



# *Rhizoctonia* *solani*



White stem collar can develop early in season if sexual stage



## BACKGROUND

- Inoculum: seed tuber- and/or soil-borne **sclerotia or mycelium**
  - **Basidiospores** during crop closure >> sexual reproduction >> polycyclic nature of disease
- Role of **soil- vs seed tuber-borne inoculum** in disease development: Contentious issue
- **Mark-release-recapture experiments** can provide answers
  - Allow tracking of inoculum sources during disease development >> determine origin of inoculum

Infected seed and  
volunteer potatoes

Pathogen overwinters as  
sclerotia and mycelia

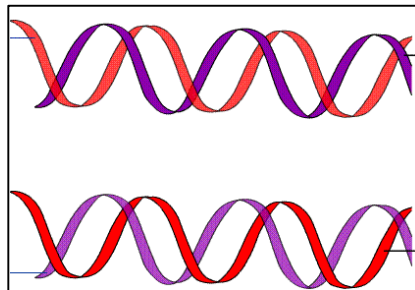
# Objectives

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1. To investigate the contribution of *R. solani* AG 3-PT **seed tuber- and soil-borne inoculum** to disease development



2. To investigate the **evolution** of an experimental population of *R. solani* AG 3-PT under field conditions



## MOTIVATION

The role of each source of inoculum is a contentious issue

Understanding the contribution of each source of inoculum is an important step in implementing effective **disease management strategies** for *R. solani*

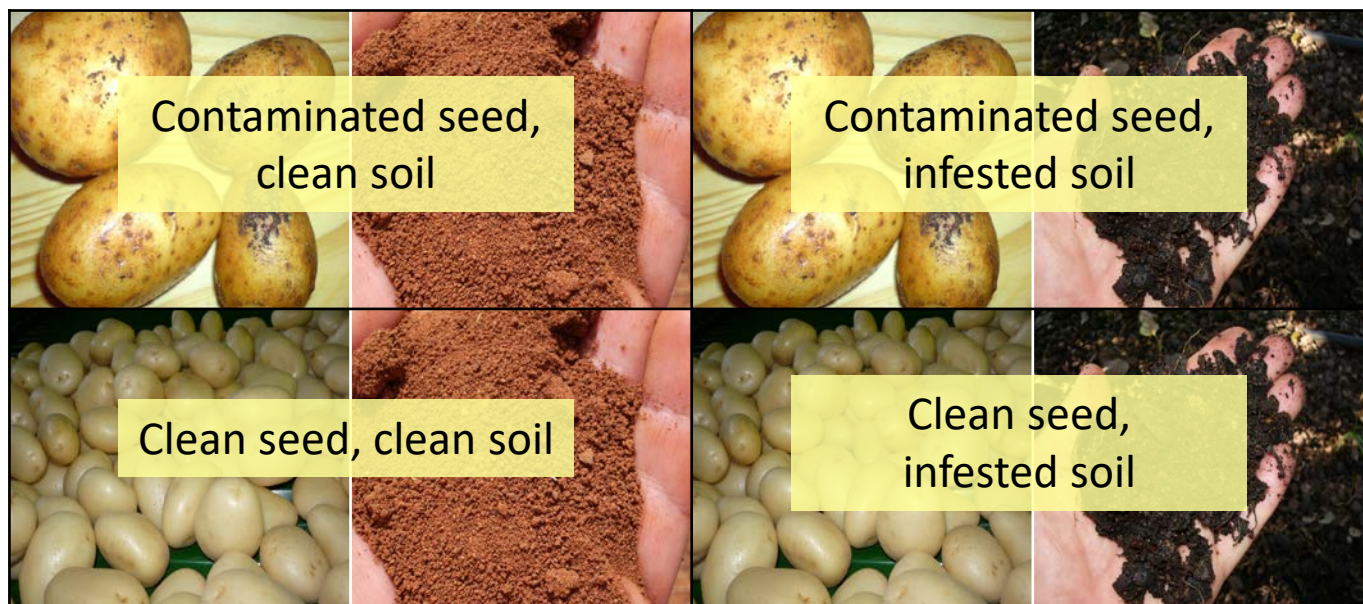


VS

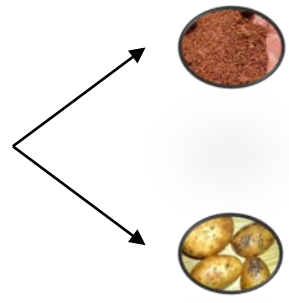


# Materials and Methods

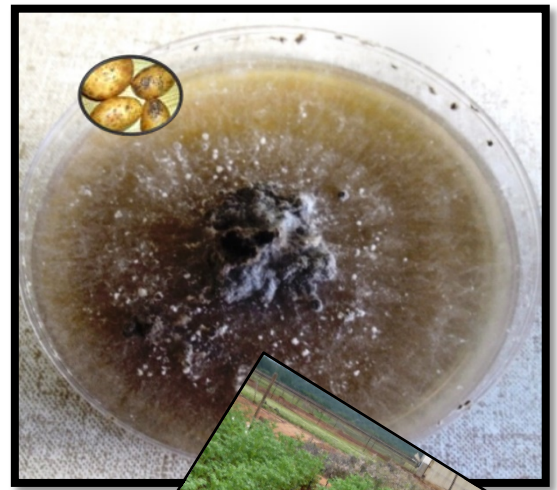
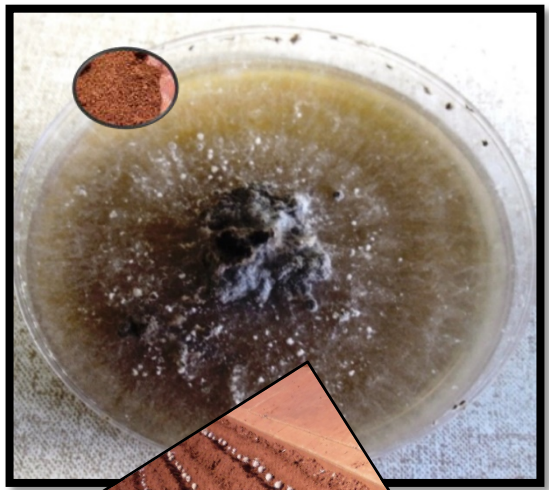
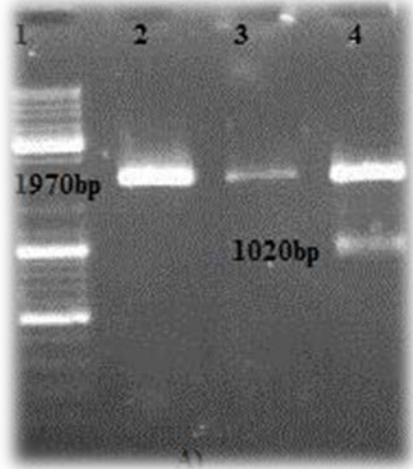
- University of Pretoria experimental plots
- RCBD
- Treatments: (4 treatments x 4 reps = 16 plots)



# Two distinct MLG groups of *R. solani* AG 3-PT isolates (five isolates per group)



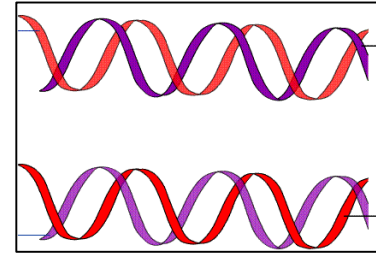
**PCR-RFLP profiling**  
(Ceresini et al. 2002)



**Disease assessments:**  
21 & 60 days after planting;  
At harvest  
**Isolations** from diseased material



# Materials and Methods



Contaminated seed,  
infested soil  
From previous trial –  
not reinoculated



**Disease assessments:**  
60 days after planting;  
At harvest  
**Isolations** from diseased  
material  
**PCR-RFLP**  
**Microsatellite analysis**  
(Ferrucho et al. 2009)





# Results: Disease index



VS



Inoculum source	Root infection		Stolon canker	
	2013	2014	2013	2014
Control	0.0 c*	0.0 c	0.0 d	0.0 d
Soil	14.5 b	10.5 b	8.5 b	15.0 a
Soil and tuber	17.5 a	12.5 a	10.2 a	12.0 b
Tuber	13.0 b	6.0 c	5.0 c	8.0 c

\*Duncan's Multiple Range test P = 0.05

# Results: Disease index



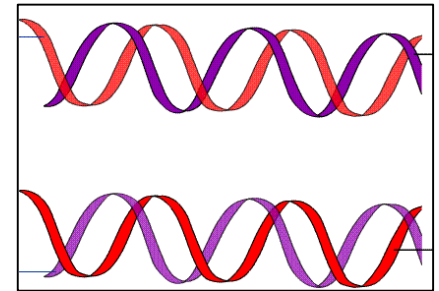
VS



	Stem canker		Black scurf	
<b>Inoculum source</b>	<b>2013</b>	<b>2014</b>	<b>2013</b>	<b>2014</b>
<b>Control</b>	0.0 d*	0.0	0.0 c	0.0 d
<b>Soil</b>	22.5 c	14.5 c	13.4 b	5.8 c
<b>Soil and tuber</b>	37.0 a	22.5 a	17.6 a	11.3 a
<b>Tuber</b>	29.0 b	17.4 b	12.6 b	7.8 b

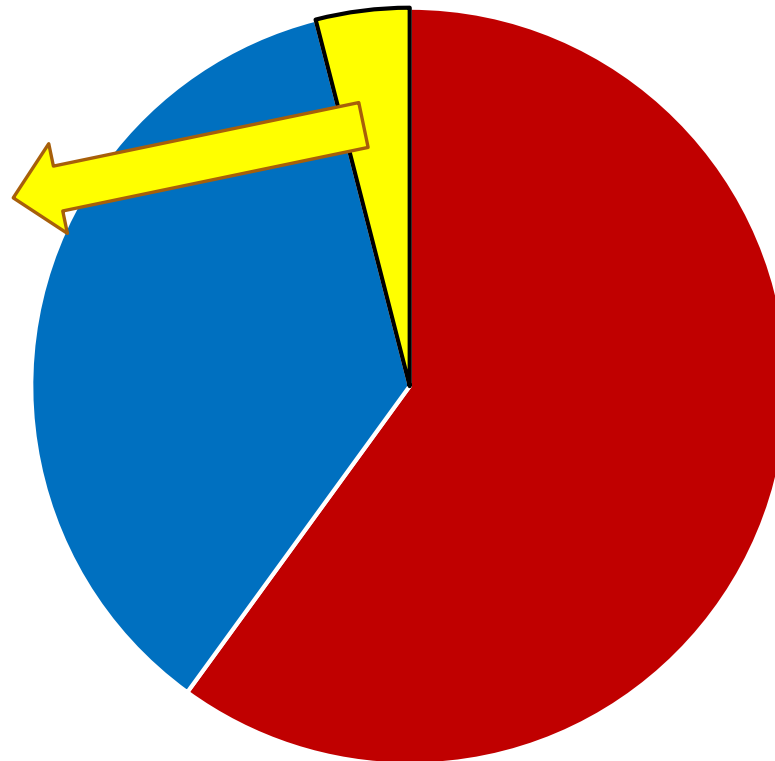
\*Duncan's Multiple Range test P = 0.05

# Results



Isolated from plants in plots inoculated with both seed and soil-borne inoculum

**Suggests recombination in field**

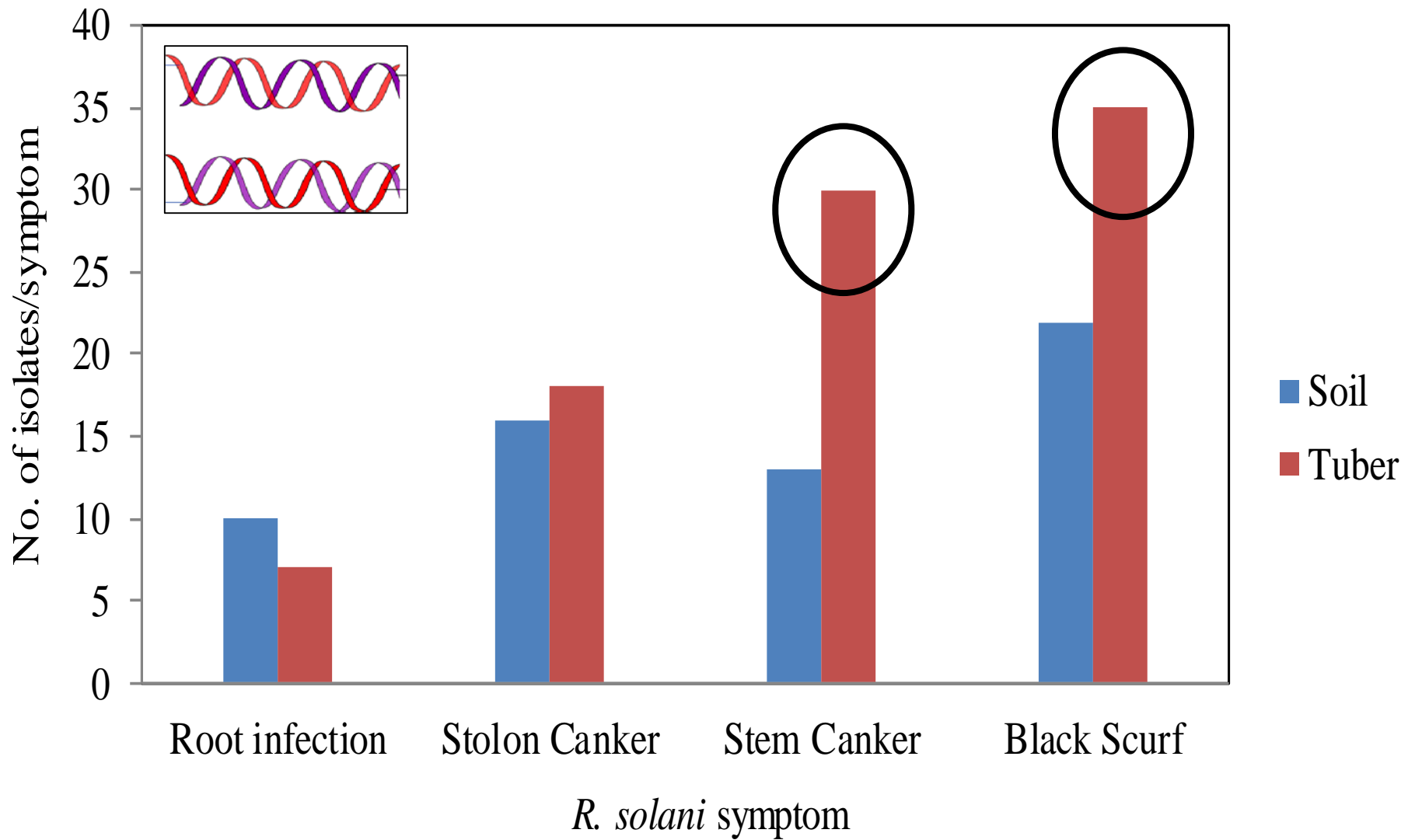


■ Seed (60%)

■ Soil (36%)

■ Non Released (4%)

Genotypes of isolates from symptomatic tissue (n = 151)



PCR-RFLP genotyping to correlate isolates obtained from symptoms to inoculum source

# Conclusions



VS



Black scurf: **Seed-borne** inoculum more important

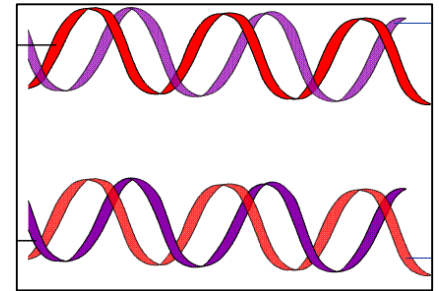


Root & stolon infection: **Soil- and seed-borne** inoculum equally important



Stem canker: **Seed-borne** inoculum more important

# Conclusions



**Recombination** may occur in field under conducive environmental conditions  
>> gradual **increase in genetic diversity**

Production of **basidiospores** >> initial inoculum

May gradually result in **changes in genetic structure** of field populations from year to year

Ensure the use of **pathogen-free seed tubers** to eliminate seed-borne inoculum and the introduction of new genotypes of *R. solani*

Apply to control measures to **limit development of sexual phase** in field

# Acknowledgements

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