



# Sclerotinia

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Character	Species	Country
Sclerotinia	<i>B. napus</i> <i>B. juncea</i>	China, Australia India, Australia, China



# Objectives

- 1. Develop disease screening protocols, especially for Australia**
- 2. Screen *B. napus* and *B. juncea* germplasm for resistance (India, China and Australia)**

# Key findings

## DEVELOPMENT OF SCREENING PROTOCOLS

1. Evaluated published leaf, petiole and stem as inoculation sites

*In certain tests: such as petiole, detached leaves*

Varietal differentiation occurs

BUT

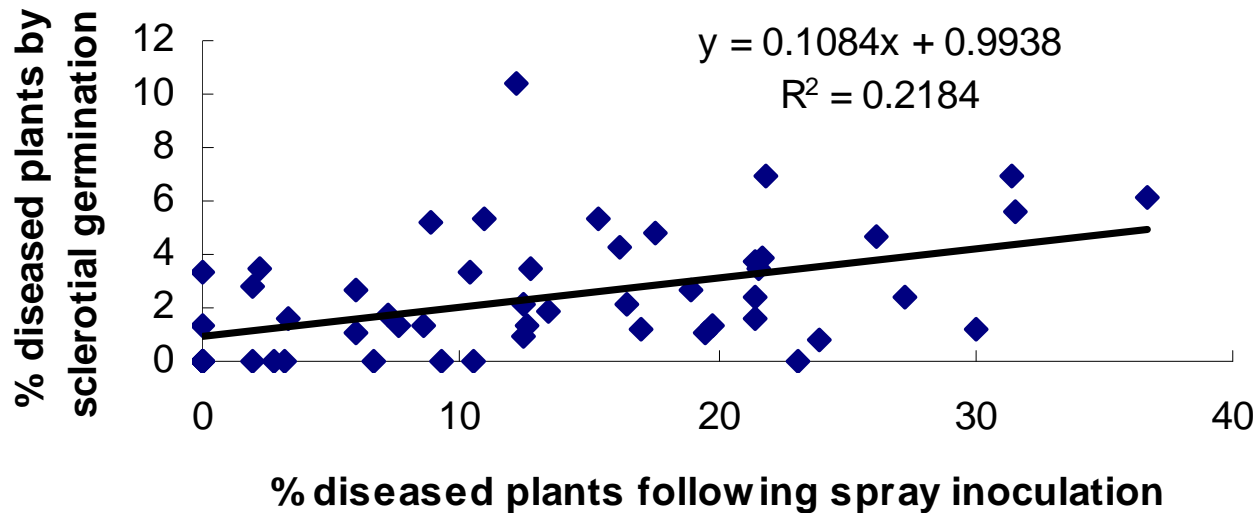
can sometimes/frequently correlates poorly with infection from artificial stem inoculations or natural inoculations in the field

# Key findings

## DEVELOPMENT OF SCREENING PROTOCOLS

### 2. Evaluated different field inoculation types

- (i). Application of a spray of mycelial suspension
- (ii). Myceliogenic germination originating from sclerotia resident in soil



# Key findings

## DEVELOPMENT OF SCREENING PROTOCOLS

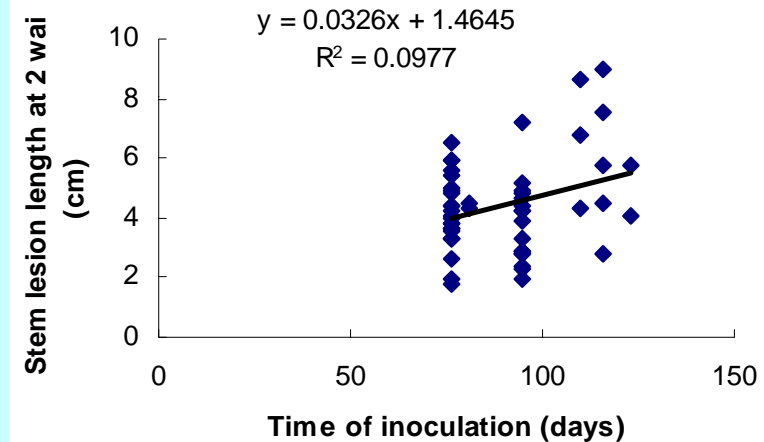
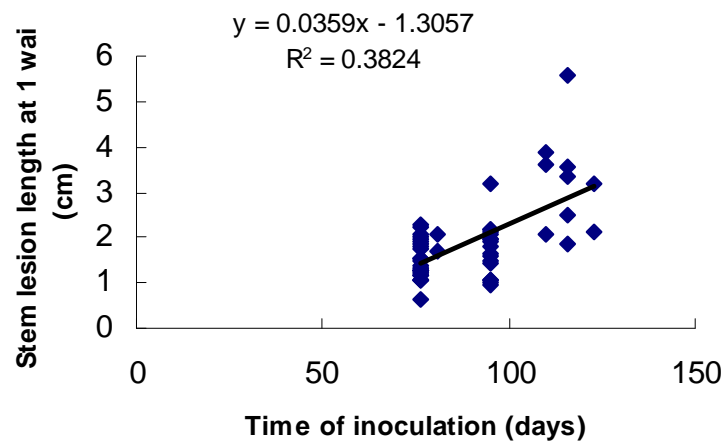
2(iii). Stem inoculation: Chosen for screening genotypes under field conditions at the flowering stage [single agar plug disc bearing actively growing mycelium according to Buchwaldt et al. (2005)]

# Key findings

## DEVELOPMENT OF SCREENING PROTOCOLS

### *Stem inoculation*

*Time of disease assessment resolves challenge of different genotype maturities*



BUT - No effect of flowering time if wait for 3 weeks post-inoculation to assess disease

i.e., The impact of different flowering times rendered insignificant when assessment of stem inoculation is delayed until 3 wks post-inoculation

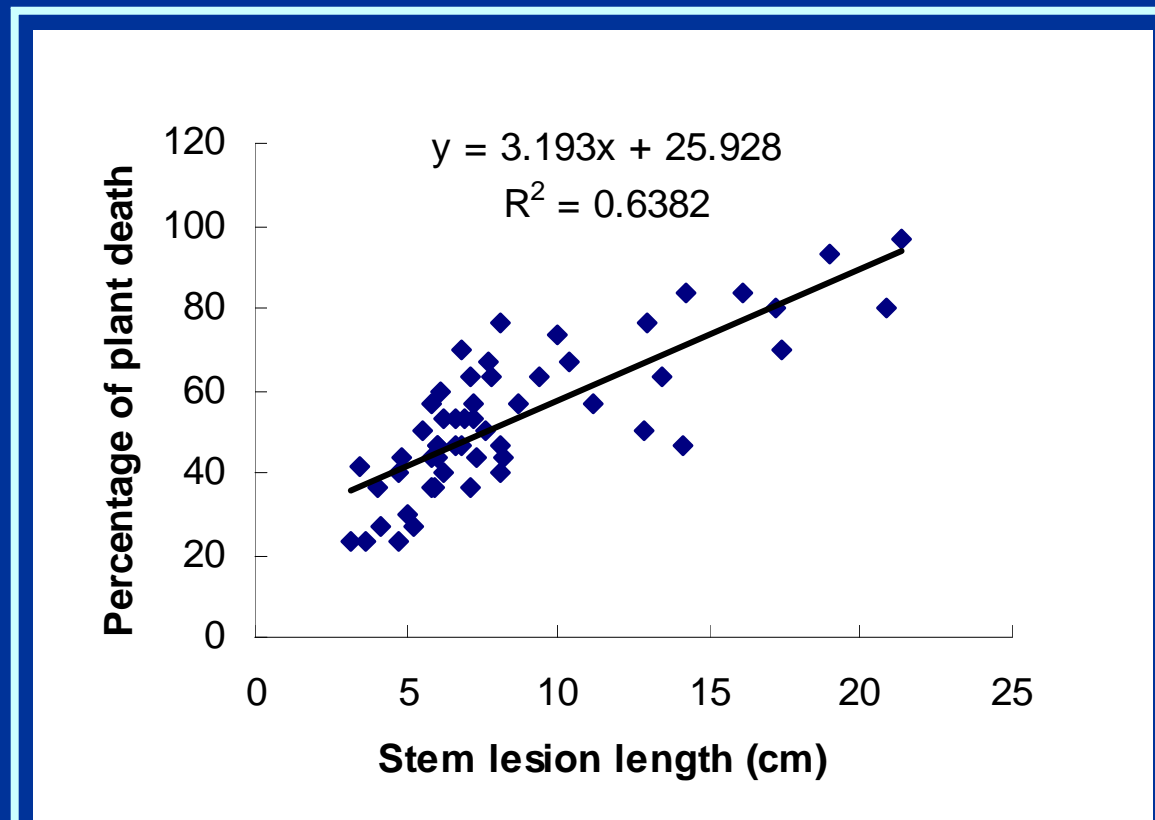
# Key findings

## DEVELOPMENT OF SCREENING PROTOCOLS

*Stem inoculation*

*Other advantages of field stem test*

Stem lesion length relates well to plant death



# Key findings

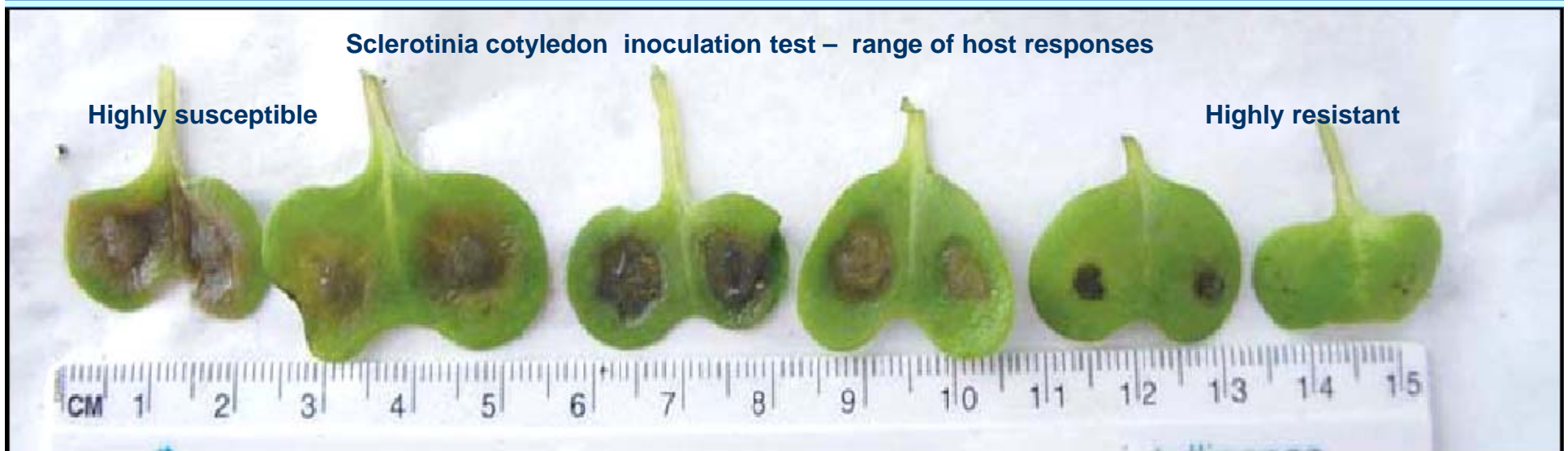
Garg, H., Hua Li, Sivasithamparam, K. and Barbetti, M.J. (2008). Cotyledon assay as a rapid and reliable method of screening for resistance against *Sclerotinia sclerotiorum* in *Brassica napus* genotypes. *Australasian Plant Pathology* 37: 106-111

## DEVELOPMENT OF SCREENING PROTOCOLS

3. **Cotyledon test** already used for *Sclerotinia* disease on legumes (Grau and Bissonette, 1974) refined for *B. napus* [cotyledons drop-inoculated using macerated mycelium under controlled environmental conditions]

**Cotyledon test** provided *B. napus* genotype responses that were:

- repeatable between experiments
- proved to be a relatively reliable indicator of field performance





# Key findings

## EXCELLENT RESISTANCE FOUND IN ACIAR PROJECT

*Best =*

*B. napus* ZY006 (China)  
(stem lesion length <0.45cm)

*Others excellent =*

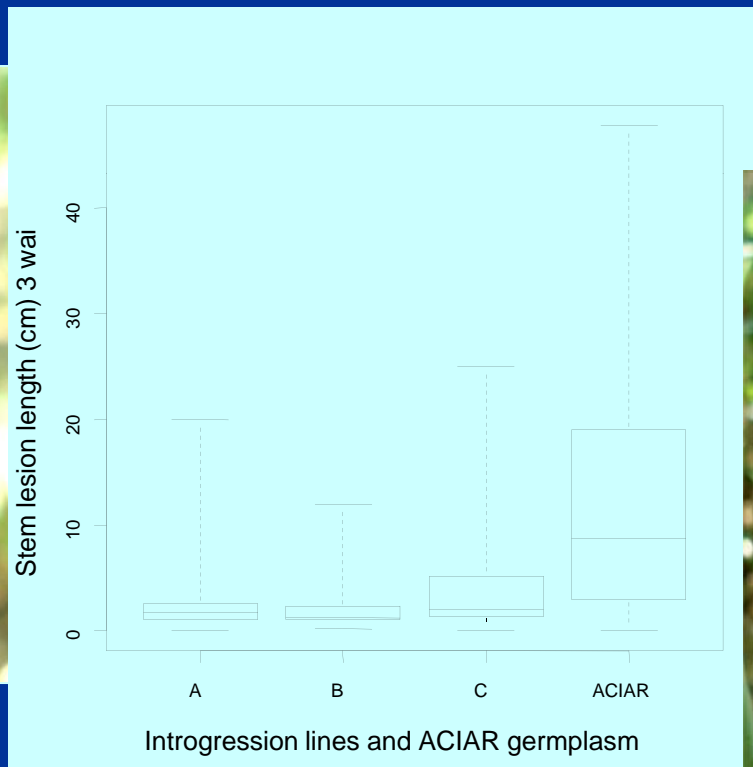
*B. napus*  
06-6-3792 & ZY004 (China)  
RT108 (Australia)

*B. juncea*  
JM06018 & JM06006 (Australia)  
*B. juncea-2* (China)

# Key findings

## FANTASTIC RESISTANCE FOUND IN PAU COLLABORATION (INDIA)

Introgression lines developed following hybridization of three wild crucifers (viz. *Erucastrum cardaminoides*, *Diplotaxis tenuisiliqua* and *E. abyssinicum*) with *B. napus* or *B. juncea*



*A = Erucastrum cardaminoides*  
*B = Diplotaxis tenuisiliqua*  
*C = Erucastrum abyssinicum*

Garg, H., Atri, C., Sandhu, P.S., Kaur, B., Renton, M., Banga, S.K., Singh, H., Singh, C., Barbetti, M.J., Banga, S.S. (2010). High level of resistance to *Sclerotinia sclerotiorum* in introgression lines derived from hybridization between wild crucifers and the crop *Brassica* species *B. napus* and *B. juncea*. *Field Crops Research* (Online at [http://www.elsevier.com/wps/find/journaldescription.cws\\_home/503308/description#description](http://www.elsevier.com/wps/find/journaldescription.cws_home/503308/description#description))

# Impact of project for Australia

1. Now have a reliable field stem inoculation test
  - one that differentiates host resistance across germplasm from Australia, China and India under Western Australian field conditions
2. Now have high level host resistance is now available for oilseed *Brassica* breeding programs in Australia
3. Now have a cotyledon test developed for rapid growth room screenings for *B. napus* genotypes
4. Now have substantially better understanding of this pathogen and *Sclerotinia-Brassica* pathosystem, especially in terms of identifying host resistance
5. Now understand need for screening for *Sclerotinia* resistance to be undertaken in each country using regional pathogen isolates and that host resistances identified may not be applicable across countries

# Opportunities-Challenges-Future

1. **Opportunity** to introgress resistance into Australian cultivars
2. **Opportunity** to screen the final ACIAR 'trait-cross' materials
3. **Opportunity** to identify wider range of sources of resistance
1. **Challenge** to define the pathotype-host interactions for Australia
2. **Challenge** to define/monitor Sclerotinia pathotypes in Australia
3. **Challenge** to find resistance that is independent of pathotype

**Future** prospect for using host resistance as a critical component of Sclerotinia management is, for the first time, a real possibility



# White rust or blister

*Caixia Li*  
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Character	Species	Country
White rust	<i>B. juncea</i>	Australia

**GRDC**

Grains  
Research &  
Development  
Corporation

National Canola Pathology Workshop 23 February 2010



Australian Government

Australian Centre for  
International Agricultural Research

# Objectives

**1. Develop disease screening protocols for Australia**

**2. Screen in Australia *B. juncea* germplasm for resistance**

# Key findings

## DEVELOPMENT OF SCREENING PROTOCOLS

1. Evaluated under Glasshouse conditions, the disease development on:
  - Cotyledons
  - Seedling plant leaves
  - Mature plant leaves
  - Leaves and flowers at flowering
2. Evaluated under Field conditions
  - Leaf incidence over time
  - Leaf severity over time
  - Stagheads
3. Compared Glasshouse and Field evaluations

# Key findings

## DEVELOPMENT OF SCREENING PROTOCOLS

**Glasshouse testing: identifies most resistant genotypes irrespective of point of inoculation – e.g.**

**Cotyledon test:** The most resistant genotypes: CBJ-001, CBJ-002, CBJ-003, CBJ-004 from China and JR049 from Australia

**Seedling stage test:** The most resistant genotypes were CBJ-001 CBJ-002, CBJ-003, CBJ-004 from China and JR049 from Australia

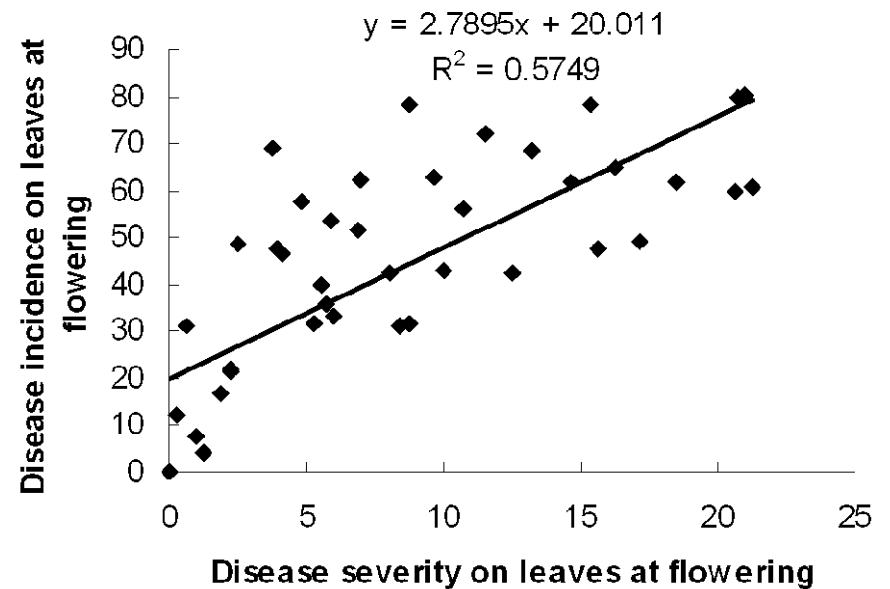
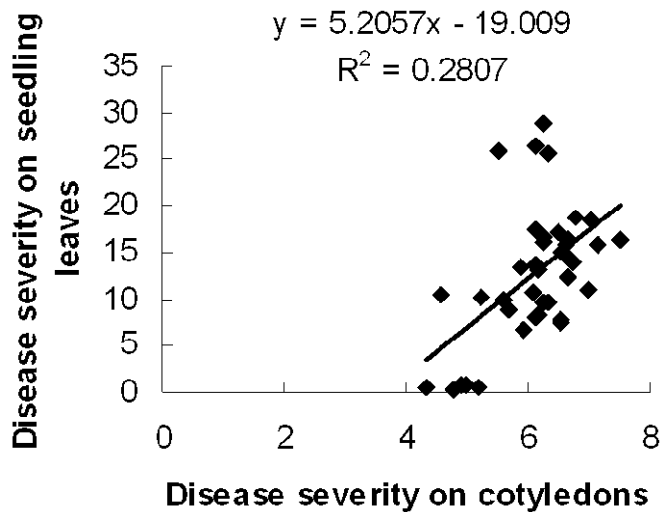
**Flowering stage test:** The most resistant genotypes: CBJ-001, CBJ-002, CBJ003 and CBJ004 from China and JR049 from Australia



# Key findings

## DEVELOPMENT OF SCREENING PROTOCOLS

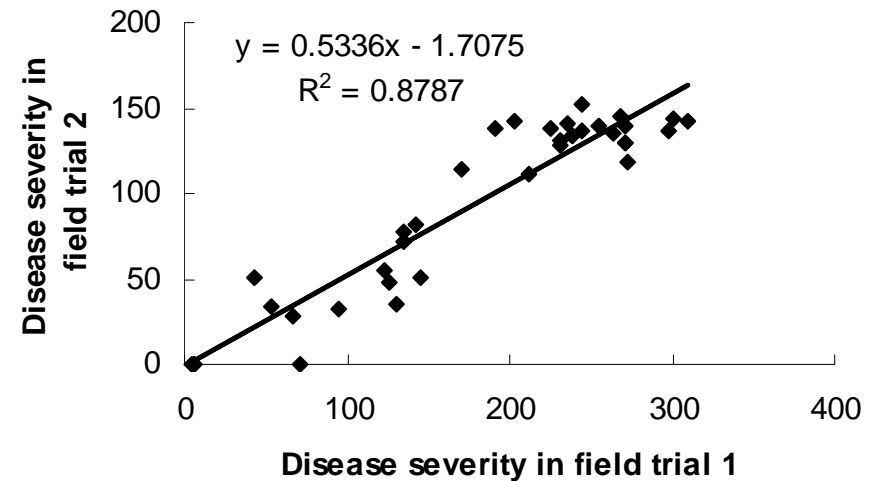
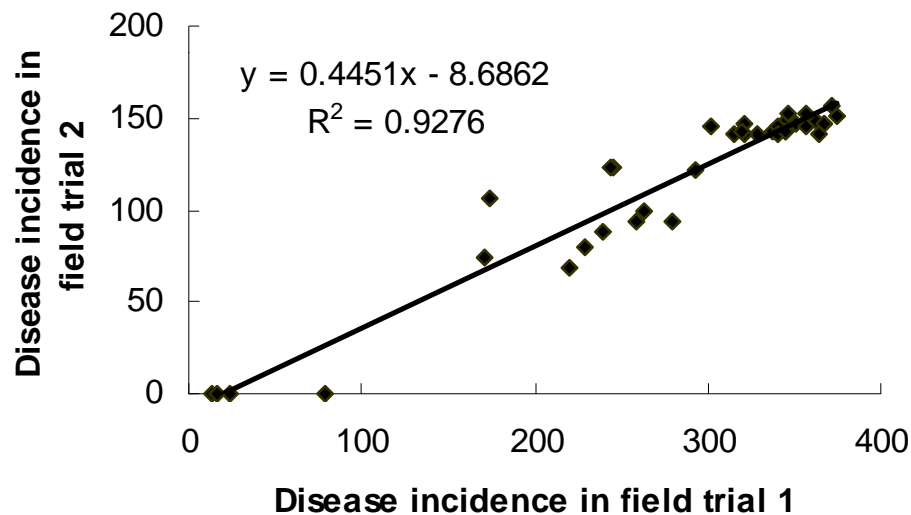
Glasshouse testing: often good overall correlation but some individual genotype exceptions, e.g.



# Key findings

## DEVELOPMENT OF SCREENING PROTOCOLS

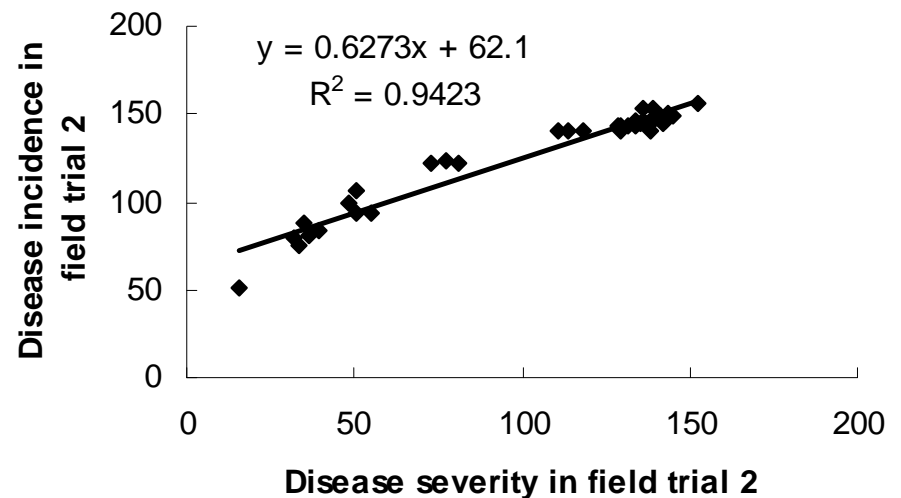
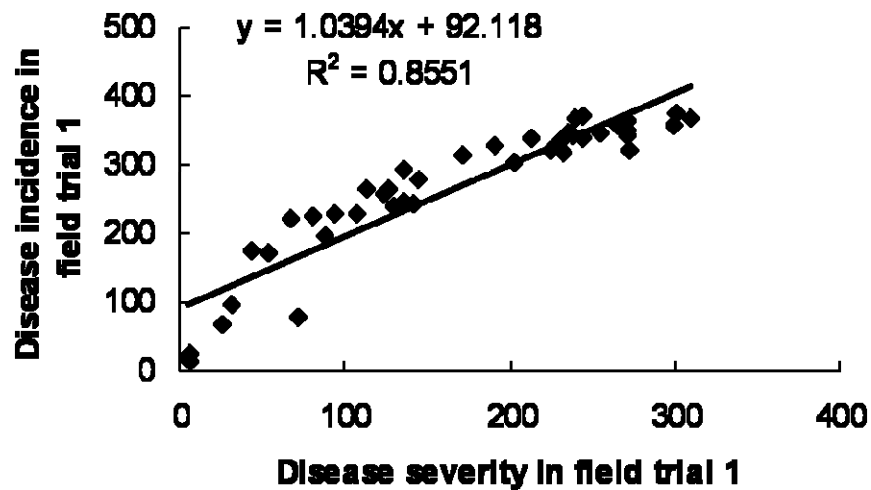
Field testing: excellent overall correlation between experiments but some individual genotype exceptions, e.g.



# Key findings

## DEVELOPMENT OF SCREENING PROTOCOLS

Field testing: excellent overall correlation between different disease parameters but some individual genotype exceptions, e.g.



# Key findings

## DEVELOPMENT OF SCREENING PROTOCOLS

### Across four field trials:

- Found that both incidence and severity of white rust disease reflected host resistance in *B. juncea* germplasm from Australia, China and India

### Conclusions from glasshouse and field testings:

- Differentiation of high levels of resistance among genotypes is similar in field as for artificially-inoculated seedlings or adult plants under glasshouse conditions – BUT, field is preferable (at least to confirm critical resistances)
- Leaf disease vs staghead disease relationship still needs further investigation – generally little or no correlation across genotypes

# Key findings

Kaur, P., Sivasithamparam, K. and Barbetti, M.J. (2008). Pathogenic behaviour of strains of *Albugo candida* from *Brassica juncea* (Indian mustard) and *Raphanus raphanistrum* (wild radish) in Western Australia. *Australasian Plant Pathology* 37: 353-356

## DEVELOPMENT OF SCREENING PROTOCOLS – must know the pathotypes present

Reactions of different cruciferous host differentials to Western Australian isolates of *Albugo candida*

Host differential	Disease reaction	
	<i>B. juncea</i> isolate	<i>R. raphanistrum</i> isolate
<i>Brassica carinata</i> 94024.2	-	-
<i>Brassica juncea</i> cv. Vulcan	+	-
<i>Brassica juncea</i> cv. Commercial Brown	+	+
<i>Brassica napus</i> cv. FAN 189 (China)	+	+
<i>Brassica napus</i> cv. Surpass 501TT	-	-
<i>Brassica nigra</i> 90745	+	+
<i>Brassica oleracea</i> var. <i>italica</i> cv. B sprouts	-	-
<i>Brassica rapa</i> cv. Torch	-	-
<i>Brassica rapa</i> cv. Reward	-	-
<i>Raphanus raphanistrum</i> WARR25	-	+
<i>Raphanus sativus</i> cv. White Icicle	+	+
<i>Brassica tournefortii</i> BTO2	+	-
<i>Eruca vesicaria</i> MJB1-06	-	-

***B. juncea* pathotype 2V is in Australia and infects:**

*B. napus* from China (FAN 189)  
*B. tournefortii* (wild turnip)  
*B. nigra*  
*Raphanus sativus*

***R. raphanistrum* pathotype infects:**

*B. juncea*  
*B. napus* from China  
*B. nigra*  
*R. sativus*

Warning: breeders to take care if:

- (i) sourcing white rust resistance from *B. napus*
- (ii) using China *B. napus* for breeding
- (iii) if these species are to be utilized commercially in Australia

Currently testing common host differentials to characterise WR races in India with PAU collaboration

# Key findings

## EXCELLENT RESISTANCE FOUND

### After glasshouse trials and then four field trials over four seasons:

- Most resistant genotypes were JM06011, JM06010, JM06021, JM06004 and JM06013 from Australia and CBJ-001, CBJ-003, CBJ-004 from China
- The very best resistance was on JM06011 that was similar to that of CBJ-003 and CBJ-004 from China, with incidence and severity scored zero
- JM06010, JM06021, JM06004 and JM06013 were more resistant than JR049 which was the best of the Australian genotypes from series 1 germplasm

# Impact of project for Australia

- Have reliable means to differentiate levels of resistance to white rust in germplasm under glasshouse or field tests  
(can utilise glasshouse screening initially and then confirm with field screening)
- Now have first high levels of resistance (foliage and stagheads) to pathotype 2V available for Australian oilseed Brassica breeding programs
- Now have substantially better understanding of this pathogen and *Albugo-Brassica* pathosystem, especially in terms of identifying host resistance (both foliage and stagheads)
- Now understand need for screening for White Rust resistance to be undertaken in each country using regional pathogen isolates and that host resistances identified may not be applicable across countries
- Now developing a clearer picture of the pathogen race status in Australia and the implications of this for disease screening and *Brassica* breeding and cultivation

# Opportunities-Challenges-Future

1. **Opportunity** to introduce resistance to pathotype 2V into all new Australian *B. juncea* cultivars
  2. **Opportunity** to screen the final ACIAR 'trait-cross' materials
  3. **Opportunity** to identify wider range of sources of resistance
1. **Challenge** to define the pathotype-host interactions for Australia
  2. **Challenge** to fully define/monitor White Rust pathotypes in Australia (need set of standard host differentials to characterise races worldwide)
  3. **Challenge** to manage White Rust if many different susceptible *Brassica* crops
- Future** prospects for using host resistance as a critical component of effective White Rust management in Australia are promising