QUEENSLAND: Mung bean powdery mildew, wheat powdery mildew, barley net blotches

AUSTRALIAN FUNGICIDE RESISTANCE EXTENSION NETWORK





Monday 14 August 2023

AUSTRALIAN FUNGICIDE RESISTANCE EXTENSION NETWORK



Regionally specific resources and training to help growers and advisors understand the status, risks and management of fungicide resistance in Australian grains.

Develop and deliver:

- Fungicide resistance management guide
- > Workshops, info sessions & webinars
- > Factsheets, updates & email alerts





SARDI

RESEARCH AND

Department of

adcommunicators.















Government of South Australi

Department of Priman lustries and Reni

> Department of Primary Industries and Regional Development





AUSTRALIAN FUNGICIDE RESISTANCE EXTENSION NETWORK



Fungicide resistance in Queensland grains crops: Introduction and a case study

Prof Levente Kiss



Fungicide resistance terminology

AUSTRALIAN FUNGICIDE RESISTANCE EXTENSION NETWORK

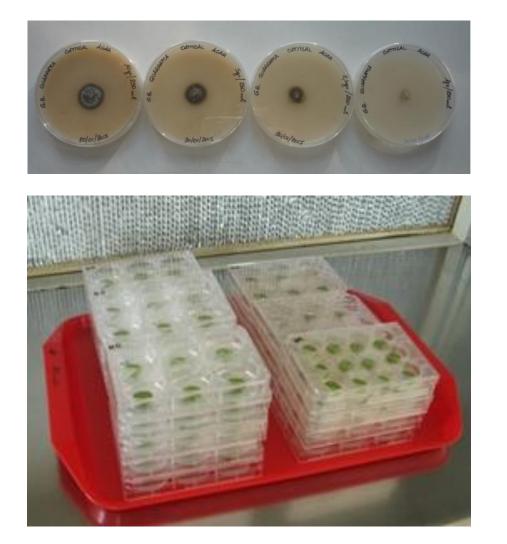


Term	Impact on fungicide use
Sensitive	Still works
Reduced sensitivity Lab confirmation	Might still work okay May need to use higher rates Higher risk of developing resistance
required Resistant	Doesn't work – avoid use • Field failure detected

Lab detection	Measurable decrease in sensitivity when				
	fungus cultured in the lab ± mutation detection				

AUSTRALIAN FUNGICIDE RESISTANCE EXTENSION NETWORK

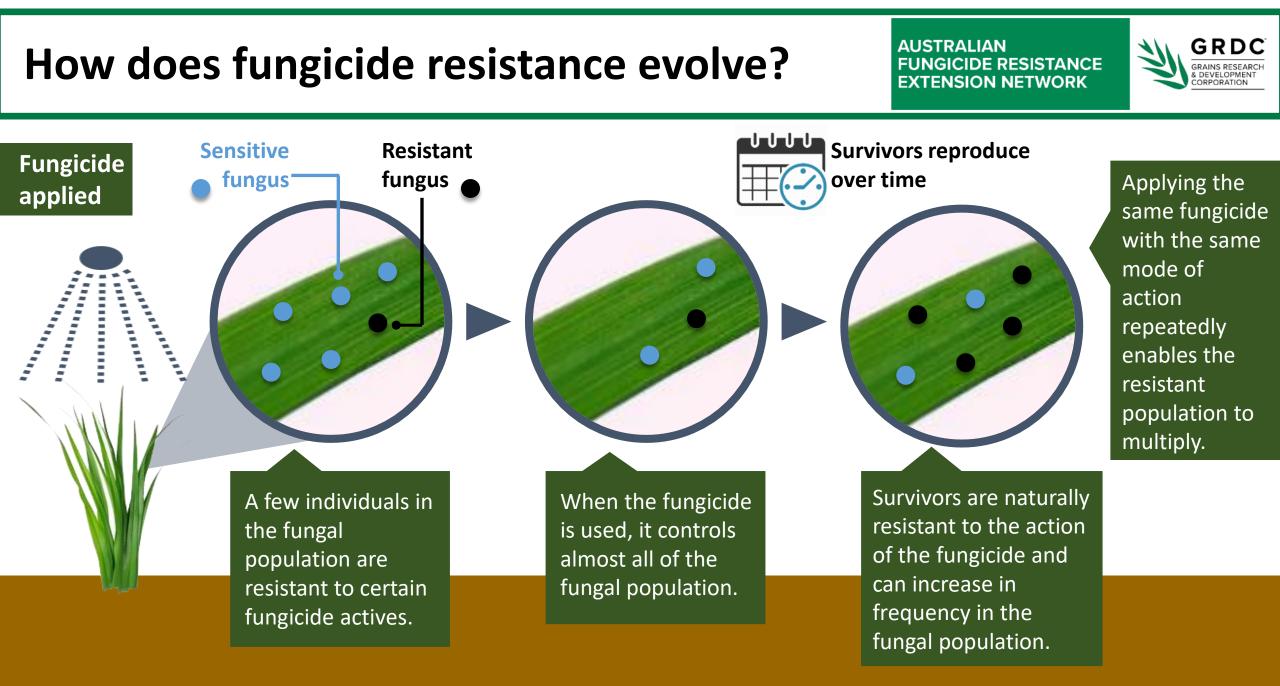




To detect reduced sensitivity or resistance to a fungicide:

- 1. Field failure
- 2. Lab detection of reduced sensitivity of pathogenic strains isolated from the field baseline sensitivity!
- 3. DNA-level detection of one or more mutations in the pathogen's gene(s) associated with the mode of action of the fungicide





Modified from CropLife Australia Fungicide Resistance Management Fact Sheet - https://www.croplife.org.au/resources/programs/resistance-management/fact-sheet-fungicide-resistance/

Fungicide resistance in Australian crops

AUSTRALIAN FUNGICIDE RESISTANCE EXTENSION NETWORK



As of July 2023

NB. Dots point to state only, not area where resistance was discovered.

		Disease and fungicide group
L, RS, R		Barley Powdery Mildew – Group 3 (DMI)
L, RS, R	٠	Barley Net Form Net Blotch – Group 3
RS, R	•	Barley Net Form Net Blotch – Group 7 (SDHIs)
RS, R	•	Barley Spot Form of Net Blotch – Group 3
L, RS, R	•	Barley Spot Form of Net Blotch – Group 7
L, R	•	Wheat Powdery Mildew – Group 3
L, R	0	Wheat Powdery Mildew – Group II (strobilurins)
RS	•	Wheat Septoria tritici blotch – Group 3
L, R	0	Wheat Septoria tritici blotch – Group II
L	0	Canola Blackleg – Group 2 (MAP-kinase)
RS	٠	Canola Blackleg – Group 3
L	•	Ascochyta Blight of Lentil – Group I (MBC)
L	•	Botrytis Grey Mould of Chickpea – Group I

L = Lab detection RS = Reduced sensitivity R = Resistant

AUSTRALIAN FUNGICIDE RESISTANCE EXTENSION NETWORK



Case study #1: FR in mungbean powdery mildew (PM) in Queensland

Research on mungbean PM was supported by:



AUSTRALIAN FUNGICIDE RESISTANCE EXTENSION NETWORK





Mungbean PM

- Common disease
- Serious yield losses if appears before flowering and the environmental conditions are conducive afterwards.
- May have an impact on desiccation efficacy.
 - Available fungicides: Tebuconazole (Group 3) Veritas Opti (Teb & azoxystrobin, Groups 3 & 11)
- Current recommendation: spray at first sign of disease and 2 weeks later, if needed.
- And when is it needed? \rightarrow use an app!

AUSTRALIAN FUNGICIDE RESISTANCE **EXTENSION NETWORK**



- Industry practice: a "preventive" spray with insecticide applications, followed by one (or two) more fungicide sprays
- How many sprays are economical?
- App (Decision Support Tool): free, easy to use, always reliable





PowderyMildew MBM - Powdery Mildew management app for mungbean

PowderyMildewMBM uses a forecasting model to assist munabean growers with fungicide application decisions, on a paddock by paddock basis, and the likely economic returns from those decisions.

The user can specify individual paddock data as well as expected weather conditions so that the output relates to their own cropping circumstances.

To download the PowderyMildewMBM App, click on the App store link below from your iPad, or the Google play link below from your Android tablet.





Department of **Primary Industries and** Regional Development



GOVERNMENT OF WESTERN AUSTRALIA



2022 & 2023: a number of field trials to validate and demonstrate the value of the app – conclusions:

- "Preventive" sprays are not needed.
- App recommendations have always resulted in the most economic disease management.
- Some of the trials were the first experiments that compared the efficacy of the two available fungicide products, Tebuconazole and Veritas Opti[®], against mungbean PM.

Improving Powdery Mildew Management in Mungbean (USQ2202-001RTX)

Australian Fungicide Resistance Extension Network

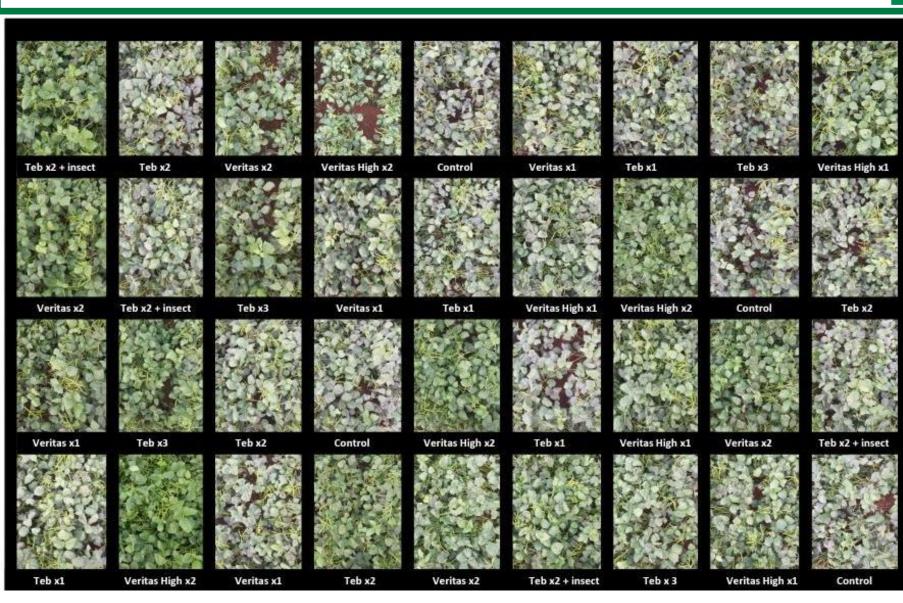
AUSTRALIAN FUNGICIDE RESISTANCE EXTENSION NETWORK





AUSTRALIAN FUNGICIDE RESISTANCE EXTENSION NETWORK



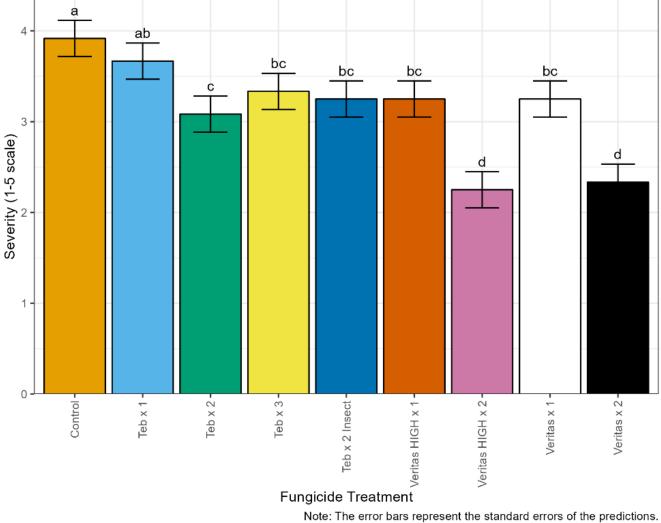


An aerial image of a mungbean PM fungicide trial in 2023. (Courtesy Neil Robinson)

AUSTRALIAN FUNGICIDE RESISTANCE EXTENSION NETWORK



USQ CCH Predictions for severity - Treatment main effect



The severity of PM infection was significantly reduced by all fungicide treatments except Tebuconazole applied once, compared to the control (P<0.001).

No significant effect of fungicide treatments on yield (P=0.453) – PM appeared late in the season, at late flowering/green pod stages.

→ The app did NOT recommend any sprays!

The LSD letters denote treatment differences between Treatments, averaged over Time (days after sowing).

AUSTRALIAN FUNGICIDE RESISTANCE EXTENSION NETWORK





100% PM control is never achieved with foliar sprays – why?

We sequenced the DNA markers for resistance to Group 3 (tebuconazole) and Group 11 (azoxystrobin) fungicides in mungbean PM samples collected from diverse paddocks & experiments since 2019 – results: * DNA marker for Group 3 resistance (G461S) detected in a single sample from a glasshouse. * DNA marker for Group 11 resistance (G143A) detected in three paddocks.



Conclusions:

- The app supports spray decisions against mungbean PM in a reliable way. The app's recommendations are useful for FR management, in addition to calculating the immediate monetary value returns on sprays – if any.
- DNA mutations conferring resistance to both MoA groups that are available for mungbean PM control were **detected in the lab** in Qld samples, but their incidence appears to be low (monitoring needed).
- Mungbean PM control can be achieved in the field in an economic way with both MoA group fungicides.



AUSTRALIAN FUNGICIDE RESISTANCE EXTENSION NETWORK The Fungicide Resistance Five – for Mungbean PM

AUSTRALIAN FUNGICIDE RESISTANCE EXTENSION NETWORK



1. Avoid susceptible crop varieties

All mungbean varieties are susceptible to PM to some extent

- 2. Rotate crops use time and distance to reduce disease carry-over Inoculum is airborne, difficult to control by rotation
- 3. Use non-chemical control methods to reduce disease pressure Plant early in the summer season!
- 4. Spray only if necessary and apply strategically Use the app!
- 5. Rotate & mix fungicides / MoA groups
- Only two MoA groups are available

AUSTRALIAN FUNGICIDE RESISTANCE EXTENSION NETWORK



Acknowledgements:

Ayesha Senanayake, MSCN Student, UniSQ SoAES Kirsty Owen, UniSQ CCH Lisa Kelly, QDAF Neil Robinson, UniSQ CCH Niloofar Vaghefi, Uni Melbourne Noel Knight, UniSQ CCH Sadegh Balotf, UniSQ CCH

Fungicide resistance frequencies in Queensland net blotch

Noel Knight Senior Research Fellow – Centre for Crop Health

noel.knight@usq.edu.au



University of **Southern Queensland**

Fungicide resistance in net blotch

Pathogen: Pyrenophora teres f. teres (net form net blotch) Pyrenophora teres f. maculata (spot form net blotch)

Host: Barley

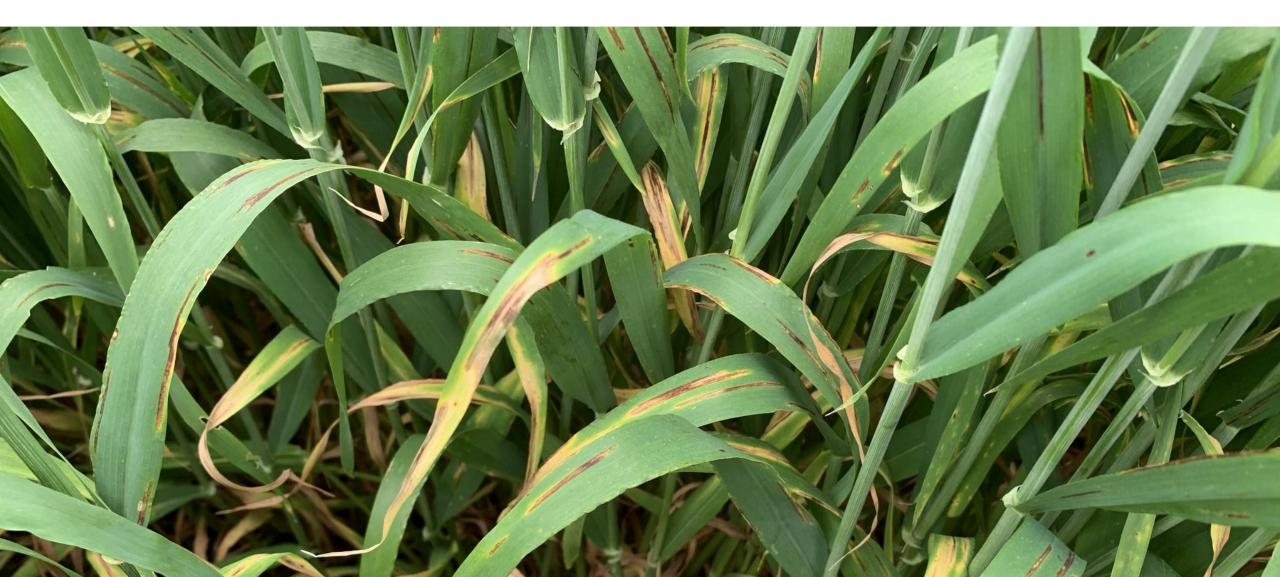
- Fungicides important disease control option
- Reduced sensitivity and resistance reported for:
 - Demethylation inhibitors (DMI) Group 3
 - Succinate dehydrogenase inhibitors (SDHI) Group 7
- Significant implications for disease management



Fungicide resistance management

 Use fungicides only when necessary & apply strategically Rotate between and within modes of action Use mixtures (if available) Stay within label rates 	Reduce populations on the plant
Integrated Disease ManagementSupport with IDM to reduce disease pressureOutputStubble managementSow later & not into previous year's stubbleOutputCrop rotationprevious year's stubbleOutputGood hygieneManage the green bridge	Reduce pathogens in the environment
Start with a solid foundationVariety SelectionSelect less susceptible varieties to reduce reliance on fungicides throughout the season	Reduce infection potential

Know Your Field!



Fungicide resistance terminology

Sensitive

• Recommended label rate controls disease

Reduced sensitivity

- Fungi can persist at low fungicide rates
 - Reduction in product performance
 - May not be obvious in the field

Resistant

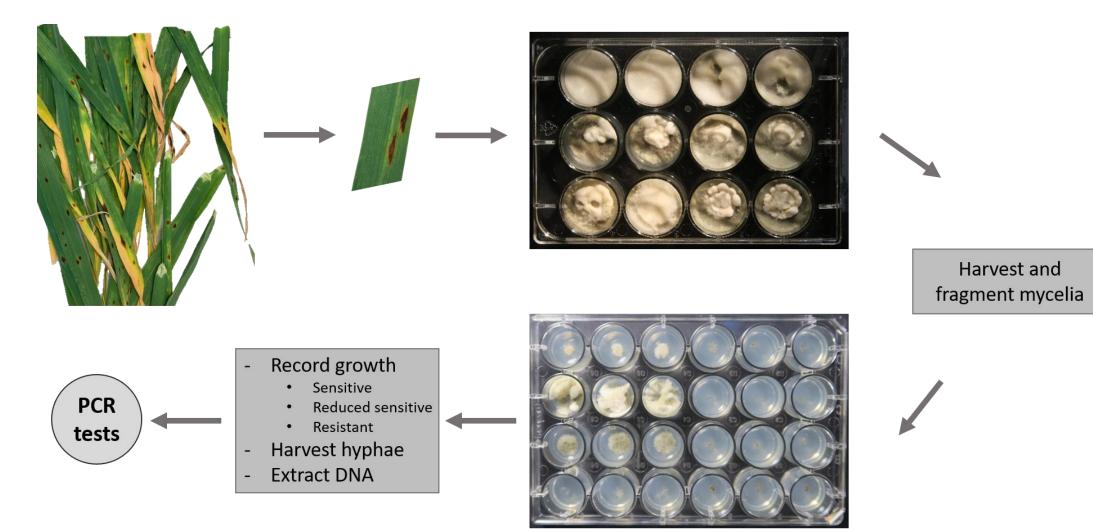
- Fungi survive at maximum fungicide rates
 - Fungicide fails to provide acceptable disease control

Lab detection

- Phenotype fungal growth on media
- Genotype fungal DNA sequence associated with resistance

Steps of characterising fungicide resistance

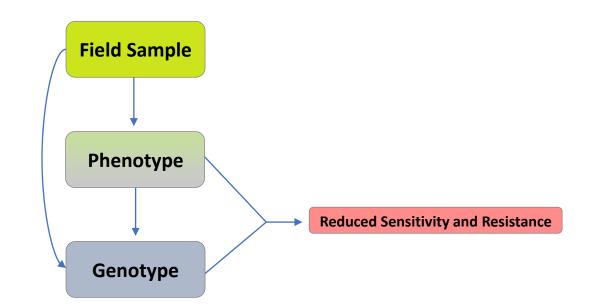
• First detection – the ground work



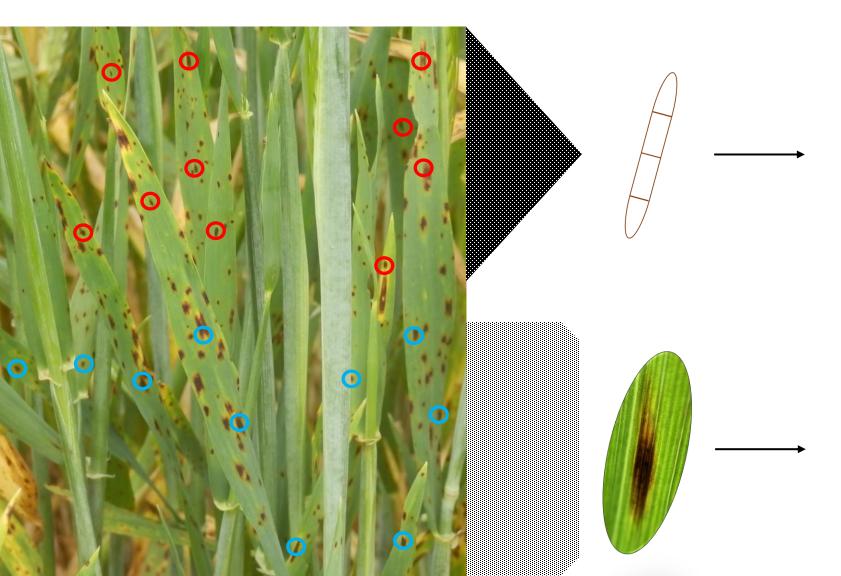
Tebuconazole: 50µg/mL

Steps of characterising fungicide resistance

- Characterisation
 - Phenotype
 - Growth with fungicides
 - Genotype
 - Link DNA changes to phenotype
- Informed Detection
 - Monitor DNA changes in field samples
 - Fungal isolates pure
 - Leaf lesions mixtures



It's all about sampling



Fungal isolation

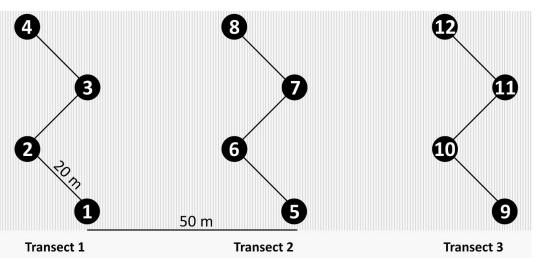
- 10 lesions total
- 1 conidia from 1 lesion
- That 'represents' the field
- Requires sporulation
- One type from one lesion
- Living fungi

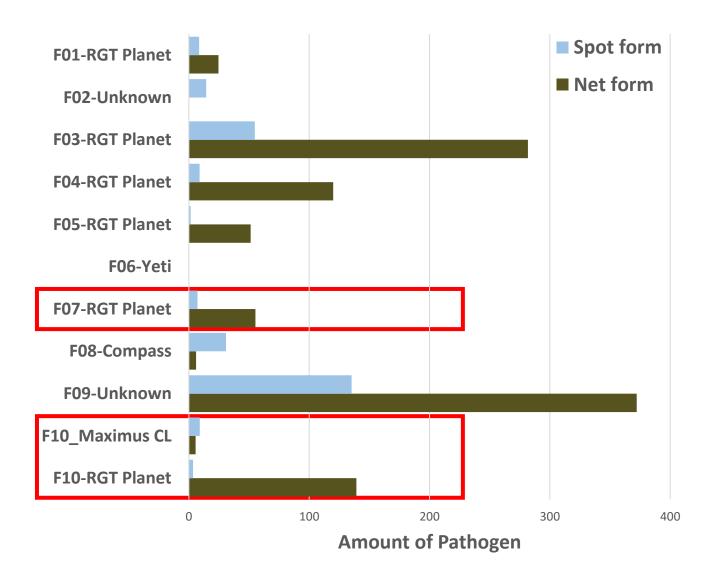
DNA isolation

- 10 lesions total
- Mixed DNA from lesions
- That 'represents' the field
- Variation within lesion kept
- Amount of each type
- DNA only
- Frequencies

Net blotch sampling and severity - 2022

- 10 barley fields sampled
- Leaves collected at 12 points in each field
 - targeting upper 3 leaves
- Combined 60 lesions
- Genotype monitoring

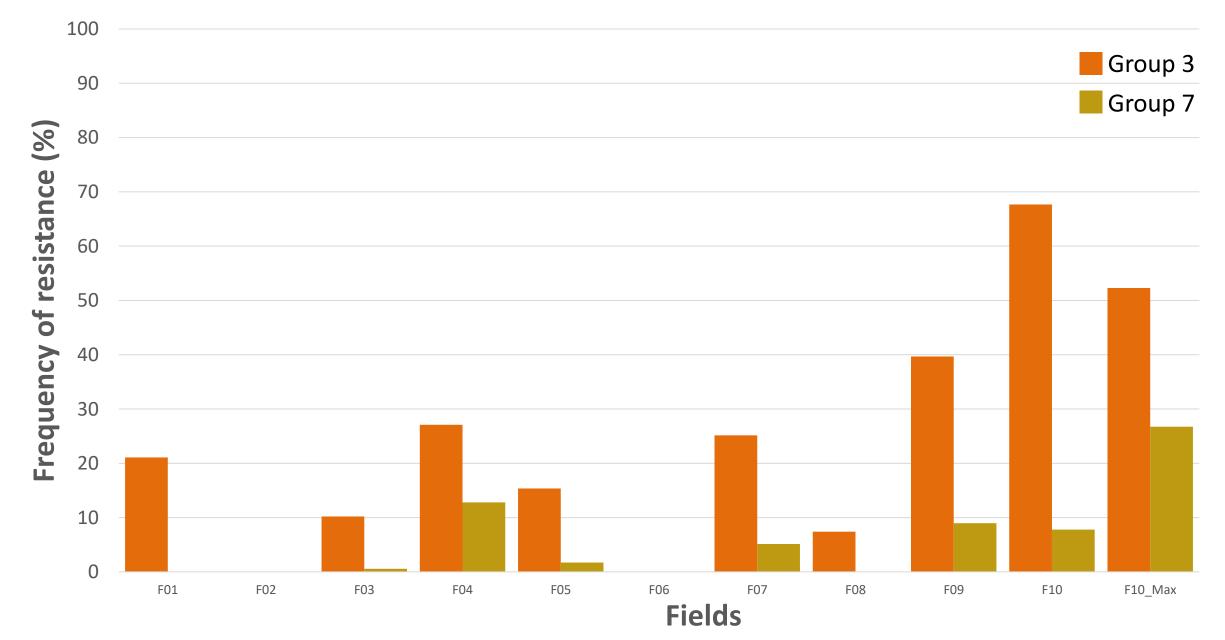




Locations and genotype frequencies



Genotype frequencies



Group 3 - Demethylation Inhibitors (DMI)

• Genotype results

- 80% of fields had reduced sensitivity and/or resistance genes
- Across all fields, 34% of the population had reduced sensitivity or resistance genes
 - Within fields ranged from 0 to 68%

Group 7 - Succinate Dehydrogenase Inhibitors (SDHI)

• Genotype results

– 60% of fields had reduced sensitivity and/or resistance genes

- Across all fields, 6% of the population had reduced sensitivity or resistance genes
 - Within fields ranged from 0 to 13%

Distribution

- Reduced sensitivity and resistance to Group 3 fungicides widespread across sampled region
- Reduced sensitivity and resistance to Group 7 fungicides less frequent
- However ...
 - Detection of Group 7 reduced sensitivity genotypes unexpected



Effects of paddock management

- Complex and diverse systems
 - No clear 'simple' impact due to fungicide history
 - Multiple seasons of any single mode of action may increase frequency of resistance
- Need to manage inoculum
 - Variety
 - Stubble
 - Crop rotations, tillage, height
 - Seed



Questions to consider

- What is the perceived risk to disease control?
 - Does a greater frequency of resistance mean more disease/loss?
 - Impact of fungicide treatments or rotations?
 - Are fungicide applications effective?

- If you have concerns, contact your local expert
 - AFREN Guide



• I am interested in sampling 2023 net blotch in Qld fields



Benefits

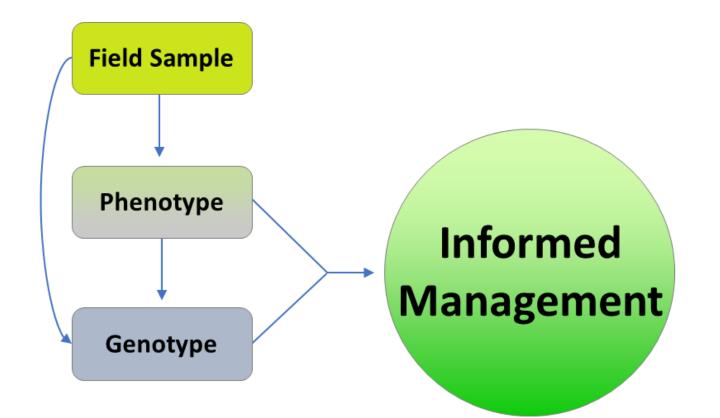
- Aim to provide a summary to growers
 - Frequencies
 - General overview

- Can these values be used to guide management now?
 - Is more information on effect of frequency required?

			DMI Resistance	e Frequency:	Medium			SDHI R	esistance Frequency:	Medium		
Field Number:	9		Demthylation	Inhibitor Resista	ance (DML F	RAC Gro	oup 3)	3) Succinate Dehydrogenase Inhib		tor Resistance (SDHL FRAC G	Gre
Location:		Demanyiation	initiation neolote	ince (Dirii, I		up 3/	Juccinia	te benyarogenuse minst	tor nesistance (Sonn, mare c	_	
Property:			Phenot	pe Assessment ¹					Phenotype Assessment ⁴			
GPS Coordinates:			Filehot	pe Assessment					Phenotype Assessment			
or 5 coordinates.					Isolates	%				Isolates	%	
Cultivar:	Spartacus CL		Sensitive	, ,	5	63			Sensitive	3	38	
Date sampled:	17/09/2021			I Sensitive	0	0			Reduced Sensitive	5	63	
outo sumpreur			Resistan		3	38			Resistant	0	0	
Net blotch form:	Spot form net blotch			-	-					-		1
			Isolate	iotal ²	8				Isolate Total ²	8		
			100/010	otar					isolate rotal			
In-season fungicide tre	estments			Phenotype	s: DMI				Phenot	types: SDHI		
	d. Propiconazole (435gai) 250ml/ha	737										
575110 1154/10 OII SEEL	a oprovidzore (+55gar) z50111/11a											
Previous season fungio	cide treatments											
Systiva FB Propiconazo												
Field Resistance Profile	e Rating (resistance frequencies) ^a											
	× 750/											
High				•								
High Medjum	25 to 75%											
High Medium Low	25 to 75%											
High Medium Low	25 to 75% < 25%		Sensiti	ve Reduced S	ensitive •	Resistan	+		Sensitive Reduc	ced Sensitive	Resistant	
Low	< 25%		Sensiti	ve Reduced S	ensitive ■	Resistan	t		Sensitive Reduc	ced Sensitive = I	Resistant	
Low ^a The greater of the phe	< 25% enoptyic or genotypic reduced		Sensiti	ve Reduced S	ensitive 🔳	Resistan	t		Sensitive Reduc	ced Sensitive	Resistant	
Low ^a The greater of the phe sensitive and resistant	< 25% enoptyic or genotypic reduced frequencies was used to indicate				ensitive •	Resistan	t			ced Sensitive 🗖	Resistant	
Low ^a The greater of the phe	< 25% enoptyic or genotypic reduced frequencies was used to indicate			ve Reduced S	ensitive •	Resistan	t		Sensitive Reduction Genotype Assessment ³	ced Sensitive	Resistant	
Low ^a The greater of the phe sensitive and resistant	< 25% enoptyic or genotypic reduced frequencies was used to indicate				ensitive Copies	Resistan %	t			ced Sensitive	Resistant %	
Low ^a The greater of the phe sensitive and resistant i the Field Resistance Pro	< 25% enoptyic or genotypic reduced frequencies was used to indicate ofile Rating			be Assessment ³			t					
Low ^a The greater of the phe sensitive and resistant the Field Resistance Pro ¹ Phenotypes represent the	< 25% enoptyic or genotypic reduced frequencies was used to indicate		Genoty	be Assessment ³	Copies 196	%	t		Genotype Assessment ³	Copies	%	
Low ^a The greater of the phe sensitive and resistant the field Resistance Pro ⁱ Phenotypes represent the cultures. Growth on DMI for	< 25% enoptyic or genotypic reduced frequencies was used to indicate ofile Rating e growth of the fungus in laboratory		Genoty	pe Assessment ³	Copies 196	%	t		Genotype Assessment ³ Sensitive	Copies 85	% 33	
Low ^a The greater of the phe sensitive and resistant the field Resistance Pro ⁱ Phenotypes represent the cultures. Growth on DMI for	< 25% enoptyic or genotypic reduced frequencies was used to indicate ofile Rating e growth of the fungus in laboratory unglcides was assessed using		Genoty	e Assessment ³ 9 I Sensitive/Resistant	Copies 196	% 77 23	t		Genotype Assessment ³ Sensitive Reduced Sensitive/Resistant	Copies 85	% 33	
Low The greater of the phe sensitive and resistant of the Field Resistance Pro Phenotypes represent the cultures. Growth on DMI fit tebuconazole at 0 (sensiti	< 25% enoptyic or genotypic reduced frequencies was used to indicate ofile Rating e growth of the fungus in laboratory unglcides was assessed using		Genoty	pe Assessment ³	Copies 196 60	% 77 23	t		Genotype Assessment ³ Sensitive	Copies 85 172	% 33	
Low ^a The greater of the phe sensitive and resistant the the Field Resistance Pro- ^a Phenotypes represent the cultures. Growth on DMI fi tebuconaziot at 0 (sensiti (resistant) µg/mL.	< 25% enoptyic or genotypic reduced frequencies was used to indicate ofile Rating e growth of the fungus in laboratory unglcides was assessed using		Genoty	e Assessment ³ e I Sensitive/Resistant be count (copies)	Copies 196 60 256	% 77 23	t		Genotype Assessment ³ Sensitive Reduced Sensitive/Resistant Genotype count (copies)	Copies 85 172 257	% 33	
Low ^a The greater of the phe sensitive and resistant the the Field Resistance Pro- ^a Phenotypes represent the cultures. Growth on DMI fi tebuconaziot at 0 (sensiti (resistant) µg/mL.	< 25% enoptyic or genotypic reduced frequencies was used to indicate ofile Rating e growth of the fungus in laboratory ungicides was assessed using ve), 15 (reduced sensitive) and 50 et from 12 lesions. The 'Isolate Total'		Genoty	e Assessment ³ 9 I Sensitive/Resistant	Copies 196 60 256	% 77 23	t		Genotype Assessment ³ Sensitive Reduced Sensitive/Resistant Genotype count (copies)	Copies 85 172	% 33	
Low ^a The greater of the phe sensitive and resistant the field Resistance Pro- ¹ Phenotypes represent the cultures. Growth on DMI fi tebuconazole at 0 (sensiti (resistant) µg/mL. ² Isolations were attempte	< 25% enoptyic or genotypic reduced frequencies was used to indicate ofile Rating e growth of the fungus in laboratory ungicides was assessed using ve), 15 (reduced sensitive) and 50 et from 12 lesions. The 'Isolate Total'		Genoty	e Assessment ³ e I Sensitive/Resistant be count (copies)	Copies 196 60 256	% 77 23	t		Genotype Assessment ³ Sensitive Reduced Sensitive/Resistant Genotype count (copies)	Copies 85 172 257	% 33	
Low ^a The greater of the phe sensitive and resistant the field Resistance Pro- ¹ Phenotypes represent the cultures. Growth on DMI fi tebuconazole at 0 (sensiti (resistant) µg/mL. ² Isolations were attempte	< 25% enoptyic or genotypic reduced frequencies was used to indicate ofile Rating e growth of the fungus in laboratory ungicides was assessed using ve), 15 (reduced sensitive) and 50 et from 12 lesions. The 'Isolate Total'		Genoty	e Assessment ³ e I Sensitive/Resistant be count (copies)	Copies 196 60 256	% 77 23	t		Genotype Assessment ³ Sensitive Reduced Sensitive/Resistant Genotype count (copies)	Copies 85 172 257	% 33	
Low ^a The greater of the phe sensitive and resistant of the Field Resistance Pro- ^a Phenotypes represent the cultures. Growth on DMI fi tebuconazole at 0 (sensiti (resistant) µg/mL. ² Isolations were attempter represents the number of	< 25% enoptyic or genotypic reduced frequencies was used to indicate ofile Rating e growth of the fungus in laboratory ungicides was assessed using ve), 15 (reduced sensitive) and 50 et from 12 lesions. The 'Isolate Total'		Genoty	e Assessment ³ e I Sensitive/Resistant be count (copies)	Copies 196 60 256	% 77 23	t		Genotype Assessment ³ Sensitive Reduced Sensitive/Resistant Genotype count (copies)	Copies 85 172 257	% 33	
Low ^a The greater of the phe sensitive and resistant the the Field Resistance Pro- ^b Phenotypes represent the cultures. Growth on DMI fi- tebuconazole at 0 (sensiti- resistant) µg/mL. ^b Isolations were attempter represents the number of ^b Genotypes (DNA sequence)	< 25% enoptyic or genotypic reduced frequencies was used to indicate ofile Rating e growth of the fungus in laboratory fungicides was assessed using ve), 15 (reduced sensitive) and 50 ed from 12 lesions. The 'Isolate Total' successful isolations.		Genoty	e Assessment ³ e I Sensitive/Resistant be count (copies)	Copies 196 60 256	% 77 23	t		Genotype Assessment ³ Sensitive Reduced Sensitive/Resistant Genotype count (copies)	Copies 85 172 257	% 33	
Low The greater of the phe sensitive and resistant of the Field Resistance Pro- Phenotypes represent the cultures. Growth on DMI for tebuconazole at 0 (sensiti- resistant) µg/mL. Isolations were attempte represents the number of Genotypes (DNA sequence) reduced sensitivity or resisted	< 25% enoptyic or genotypic reduced frequencies was used to indicate ofile Rating e growth of the fungus in laboratory ungicides was assessed using ve), 15 (reduced sensitive) and 50 ed from 12 lesions. The 'Isolate Total' successful isolations.		Genoty	e Assessment ³ e I Sensitive/Resistant be count (copies)	Copies 196 60 256	% 77 23	t		Genotype Assessment ³ Sensitive Reduced Sensitive/Resistant Genotype count (copies)	Copies 85 172 257	% 33	
Low The greater of the phesessesses and resistant the Field Resistance Presentitive and resistant of the Field Resistance Present the Field Resistance Present the cultures. Growth on DMI freebuconazole at 0 (sensitifreesistant) µg/mL. Solations were attempte represents the number of Genotypes (DNA sequence densitivty or resis Reduced sensitivty on resis Reduced sensitivty and resogether due to the potential of	< 25% enoptyic or genotypic reduced frequencies was used to indicate ofile Rating e growth of the fungus in laboratory ungicides was assessed using ve), 15 (reduced sensitive) and 50 ed from 12 lesions. The 'Isolate Total' successful isolations. etces) associated with sensitivity, stance were detected using PCR. sistant genotypes were grouped tail for phenotypic growth to		Genoty	e Assessment ³ e I Sensitive/Resistant be count (copies)	Copies 196 60 256	% 77 23	t		Genotype Assessment ³ Sensitive Reduced Sensitive/Resistant Genotype count (copies)	Copies 85 172 257	% 33	
Low The greater of the phe sensitive and resistant of the Field Resistance Pro phenotypes represent the cultures. Growth on DMI fi tebuconazole at 0 (sensiti resistant) µg/mL. Isolations were attempte represents the number of Genotypes (DNA sequence reduced sensitivity or resis Reduced sensitive and resi	< 25% enoptyic or genotypic reduced frequencies was used to indicate ofile Rating e growth of the fungus in laboratory ungicides was assessed using ve), 15 (reduced sensitive) and 50 ed from 12 lesions. The 'Isolate Total' successful isolations. etces) associated with sensitivity, stance were detected using PCR. sistant genotypes were grouped tail for phenotypic growth to		Genoty	e Assessment ³ e I Sensitive/Resistant be count (copies)	Copies 196 60 256	% 77 23	t		Genotype Assessment ³ Sensitive Reduced Sensitive/Resistant Genotype count (copies)	Copies 85 172 257	% 33	
Low The greater of the phesessitive and resistant the Field Resistance Presentitive and resistant of the Field Resistance Present the Field Resistance Presents are consistent on DMI feebuconazole at 0 (sensiti resistant) µg/mL. Isolations were attempte epresents the number of Genotypes (DNA sequence deuced sensitive and resisted use to the potent	< 25% enoptyic or genotypic reduced frequencies was used to indicate ofile Rating e growth of the fungus in laboratory ungicides was assessed using ve), 15 (reduced sensitive) and 50 ed from 12 lesions. The 'Isolate Total' successful isolations. etces) associated with sensitivity, stance were detected using PCR. sistant genotypes were grouped tail for phenotypic growth to		Genoty	e Assessment ³ e I Sensitive/Resistant be count (copies)	Copies 196 60 256	% 77 23	t		Genotype Assessment ³ Sensitive Reduced Sensitive/Resistant Genotype count (copies)	Copies 85 172 257	% 33	
Low The greater of the phesessitive and resistant the Field Resistance Presentitive and resistant of the Field Resistance Present the Field Resistance Presents are consistent on DMI feebuconazole at 0 (sensiti resistant) µg/mL. Isolations were attempte epresents the number of Genotypes (DNA sequence deuced sensitive and resisted use to the potent	< 25% enoptyic or genotypic reduced frequencies was used to indicate ofile Rating e growth of the fungus in laboratory ungicides was assessed using ve), 15 (reduced sensitive) and 50 ed from 12 lesions. The 'Isolate Total' successful isolations. etces) associated with sensitivity, stance were detected using PCR. sistant genotypes were grouped tail for phenotypic growth to		Genoty	e Assessment ³ e I Sensitive/Resistant be count (copies)	Copies 196 60 256	% 77 23	t		Genotype Assessment ³ Sensitive Reduced Sensitive/Resistant Genotype count (copies)	Copies 85 172 257	% 33	
Low The greater of the phesensitive and resistant of the Field Resistance Present the Field Resistance Present the cultures. Growth on DMI frebuconazole at 0 (sensitiresistant) µg/mL. Isolations were attempte expresents the number of Genotypes (DNA sequence educed sensitive and resistend use to the potent encompass each of these	< 25% enoptyic or genotypic reduced frequencies was used to indicate ofile Rating e growth of the fungus in laboratory unglcides was assessed using we), 15 (reduced sensitive) and 50 ed from 12 lesions. The 'Isolate Total' successful isolations. etces) associated with sensitivty, stance were detected using PCR. sistant genotypes were grouped tial for phenotypic growth to classes.		Genoty	e Assessment ³ e I Sensitive/Resistant be count (copies)	Copies 196 60 256	% 77 23	t		Genotype Assessment ³ Sensitive Reduced Sensitive/Resistant Genotype count (copies)	Copies 85 172 257	% 33	
Low The greater of the phe sensitive and resistant of the Field Resistance Pre Phenotypes represent the ultures. Growth on DMI fi ebuconazole at 0 (sensitiv resistant) µg/mL. Isolations were attempte epresents the number of Genotypes (DNA sequence deuced sensitivity or resis Reduced sensitivity and resis Reduced sensitive and resi Reduced sensitive and resis Reduc	< 25% enoptyic or genotypic reduced frequencies was used to indicate ofile Rating e growth of the fungus in laboratory ungicides was assessed using ve), 15 (reduced sensitive) and 50 ed from 12 lesions. The 'Isolate Total' successful isolations. etces) associated with sensitivity, stance were detected using PCR. sistant genotypes were grouped tail for phenotypic growth to		Genoty	e Assessment ³ e I Sensitive/Resistant be count (copies)	Copies 196 60 256	% 77 23	t		Genotype Assessment ³ Sensitive Reduced Sensitive/Resistant Genotype count (copies)	Copies 85 172 257	% 33	
Low The greater of the phe- sensitive and resistant the resisting represent the ultures. Growth on DMI f ebuconazole at 0 (sensiti- resistant) µg/mL. Isolations were attempte epresents the number of Genotypes (DNA sequence educed sensitive or resis reduced sensitive or resis reduced sensitive or the potent incompass each of these Phenotypes represent the ultures. Growth on SDMI	< 25% enoptyic or genotypic reduced frequencies was used to indicate ofile Rating e growth of the fungus in laboratory ungicides was assessed using ve), 15 (reduced sensitive) and 50 ed from 12 lesions. The 'Isolate Total' successful Isolations. edited with sensitivity, stance were detected using PCR, sistant genotypic growth to classes. e growth of the fungus in laboratory		Genoty	e Assessment ³ e I Sensitive/Resistant be count (copies)	Copies 196 60 256 :DMI	% 77 23	t		Genotype Assessment ³ Sensitive Reduced Sensitive/Resistant Genotype count (copies)	Copies 85 172 257 Cypes:SDHI	% 33 67	

Key messages

- Fungicide resistance present in Queensland
 - Fungicide application strategies should include mixed modes of action and can be informed by testing field samples



Acknowledgements

Fungicide Resistance Group (CCDM)

Kul Chandra Adhikari Wesley Mair Kejal Dodhia **Leader:** Fran Lopez-Ruiz Lincoln Harper Steven Chang







Centre for Crop Health

Ahmed Saad Anke Martin Levente Kiss Lisle Snyman (DAFQ)

UniSQ Capacity Building Grant - 2022

Queensland Agronomists & Growers

Russell Wood (Wood Ag) Matthew Skerman (Nutrien Ag Solutions)

AUSTRALIAN FUNGICIDE RESISTANCE EXTENSION NETWORK









University of **Southern Queensland**

Questions

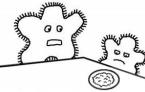
Wheat powdery mildew: DMI & Qol resistance

Field resistance to DMIs?

AUSTRALIAN FUNGICIDE RESISTANCE EXTENSION NETWORK







"See Timmy what happens when you finish all your fungicides?"



Courtesy of Steven Simpfendorfer, NSW DPI



Albury, NSW Bindaroi S-VS Wheat on wheat

- Tebuconazole 145mL/ha @ 4 August
- Propiconazole 250 mL/ha @ 6 October

AUSTRALIAN What happens under controlled conditions? FUNGICIDE RESISTANCE **EXTENSION NETWORK** Wild type Resistant Propiconazole (1x) Propiconazole (1x) Propiconazole (2x)

GRDC

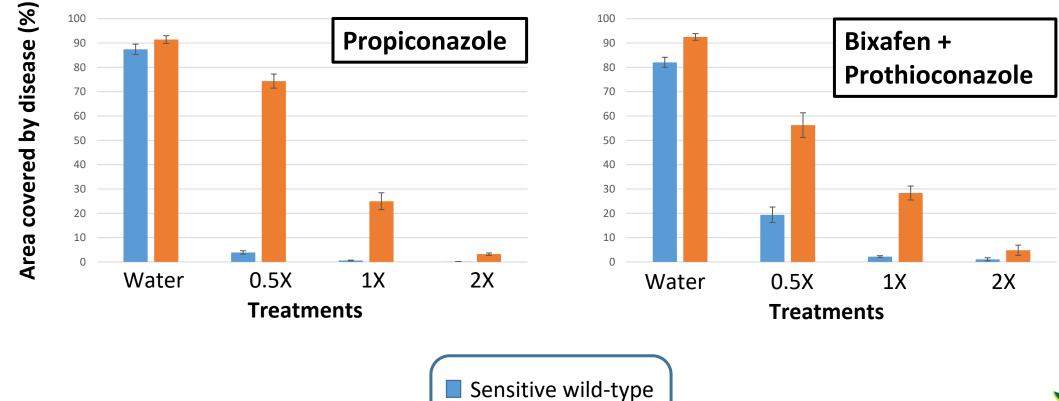
GRAINS RESEARCH & DEVELOPMENT

CORPORATION







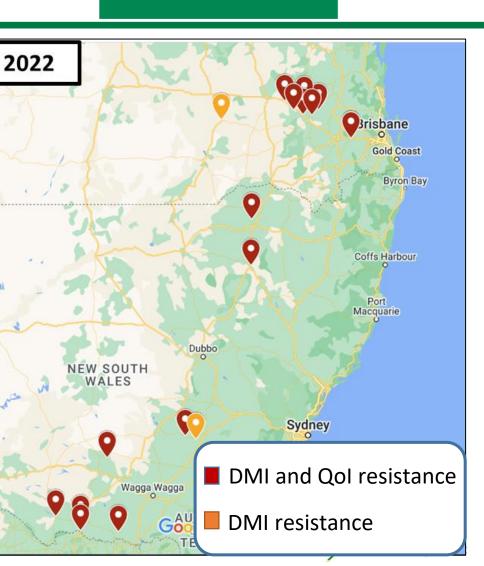


DMI resistant



QoI and DMI resistance detected in Qld

- Emergency permits for:
 - Quinoxyfen (Group 13, medium resistance risk)
 - Proquinazid (Group 13)
 - Metrafenone (Group 50, medium resistance risk)



AUSTRALIAN

FUNGICIDE RESISTANCE

EXTENSION NETWORK



GRDC

GRAINS RESEARCH & DEVELOPMENT

CORPORATION

Acknowledgements

The Fungicide Resistance Group Centre for Crop and Disease Management



Steven Chang

Kejal Dodhia

Centre for Crop and Disease Management Mark Gibberd Josh Mylne

The grains industry for

The myriad of samples Management information Feedback Advice



NSW Department of Primary Industries (NSW-DPI) Steven Simpfendorfer



Trengove Consulting Sam Trengrove

Trengove Consulting



Connect with AFREN

AUSTRALIAN FUNGICIDE RESISTANCE EXTENSION NETWORK



@theGRDC #AFREN



afren.com.au

afren@curtin.edu.au

- Fungicide resistance management guide
- Workshops, info sessions & webinars
- Factsheets, updates & email alerts



If you suspect fungicide resistance, let us know what's happening & send us a sample!