





Phenotyping transpiration efficiency: Linking trait dissection to genetics

Erik van Oosterom, Karine Chenu, Greg McLean, Geetika, Kurt Deifel, Richard Sulman, Emma Mace, David Jordan, Graeme Hammer

> Working together with the Queensland Government



Outline

- What is Transpiration Efficiency (TE) and why it is important.
- Set up of phenotyping platform.
- Role of phenotyping within integrated approach to crop improvement.
- Concluding remarks.







What is Transpiration Efficiency (TE)?



Figure 10. Schematic of a cross-section of a leaf. The resistance not defined in the text is the mesophyll resis



Set up of phenotyping platform



- Large lysimeters
 - Pot size: 50+ L.
 - Capacity: 128 lysimeters.
 - Harvest: anthesis (maturity).
 - Use: Detailed trait studies on TE and its components.





Set up of phenotyping platform



Small lysimeters

- Pot size: 4 L.
- Capacity: 560 lysimeters.
- Harvest: mid-vegetative. Use: High throughput phenotyping for TE.











- Each lysimeter is located on a load cell.
- Weights are recorded ever 10 minutes.
- Rewatering of each lysimeter is fully automated.

Set up of phenotyping platform



Integrated approach to phenotyping







- Approach: High throughput phenotyping of mapping populations to detect QTL for TE.
- Platform: Small lysimeters

Detect (statistion strates)	Genetics						
Option) Connect(0.P m(del development)	(QTL, genes)	Population #	NRP	RP	N	Experiment	
		1	RTx7000	R931945-2-2	27	201601	-
		2	IS3541	R931945-2-2	30	201601	Low TE
		3	SC237-14E	R931945-2-2	22	201601	High TE
		4	QL12	R931945-2-2	58	201601	
		5	QL12	R986087-2-4-1	. 38	201601	
		6	SC237-14E	R986087-2-4-1	. 42	201601	
		7	Macia	R931945-2-2	47	201603	
		8	M35-1	R931945-2-2	36	201603	
		9	ICSV745	R931945-2-2	39	201603	
		10	SC56-14E	R931945-2-2	73	201603	
		11	Rio	R931945-2-2	45	201603	

Subset of 11 NAM populations based on 9 exotic non recurrent parents (and 2 elite recurrent parents)

- 457 individuals in total.
- 217 in experiment 1 (201601).
- 240 in experiment 2 (201603).



Physiology (Trait analysis)



Preliminary data only on QTL

rait analysis

Genetics

- QTL consistent across experimental runs.
- Most QTL identified across multiple populations.

Genetics to Physiology - Connect



Germplasm with contrasting alleles for TE QTL can be selected for detailed trait studies to determine the function of these QTL (eg. photosynthesis, transpiration).









Physiology to Modelling – dissect



Trait dissection studies:

- Approach: identify physiological processes that underpin genotypic differences in TE.
- Application: Science from trait dissection provides simulation models with biological functionality.
- Platform: Large lysimeters.



Physiology to Modelling – dissect

Can trait dissection meet the requirements for model development:

- Relationships need to be generic.
 - allows robust parameterisation.
- Plant-level data need to be representative of leaf-level data.
 - allows connection across levels of biological organisation.
- Trait dissection needs to capture genotypic differences in the complex trait (TE).
 - dynamic models simulate complex traits as an emergent consequence of component traits.

Example: transpiration rates.





Response of transpiration rates to VPD is consistent :

Time

- across experiments.
- across days within experiments. \rightarrow Results are generic.



TE =

Biomass

Transpiration



Plant-level measurements of transpiration per unit leaf area (T/LA) are representative of leaf-level measurements of conductance.

600



Modelling to Physiology – Predict



- Science from trait dissection provides simulation models with biological functionality.
- This gives models predictive capability for trait evaluation in the target environments to unravel G × E × M interactions.











Concluding remarks

Phenotyping for TE in our lysimetry platform is part of an integrated approach that exploits synergies of combining phenotyping and crop modelling with genotyping.

- Quantifies the role of TE in sorghum production.
- Informs decision making in the breeding program.

Two platforms we have are complementary and serve different purposes.

• Addition of imaging system would allow some trait dissection work to be done in the small lysimeters.





Acknowledgements



THE UNIVERSITY

OF OUEENSLAND

Australian Government

Australian Centre for International Agricultural Research

BILL& MELINDA GATES foundation





