

THE HUNT FOR MODIFIERS OF THE Tcb1 LOCUS

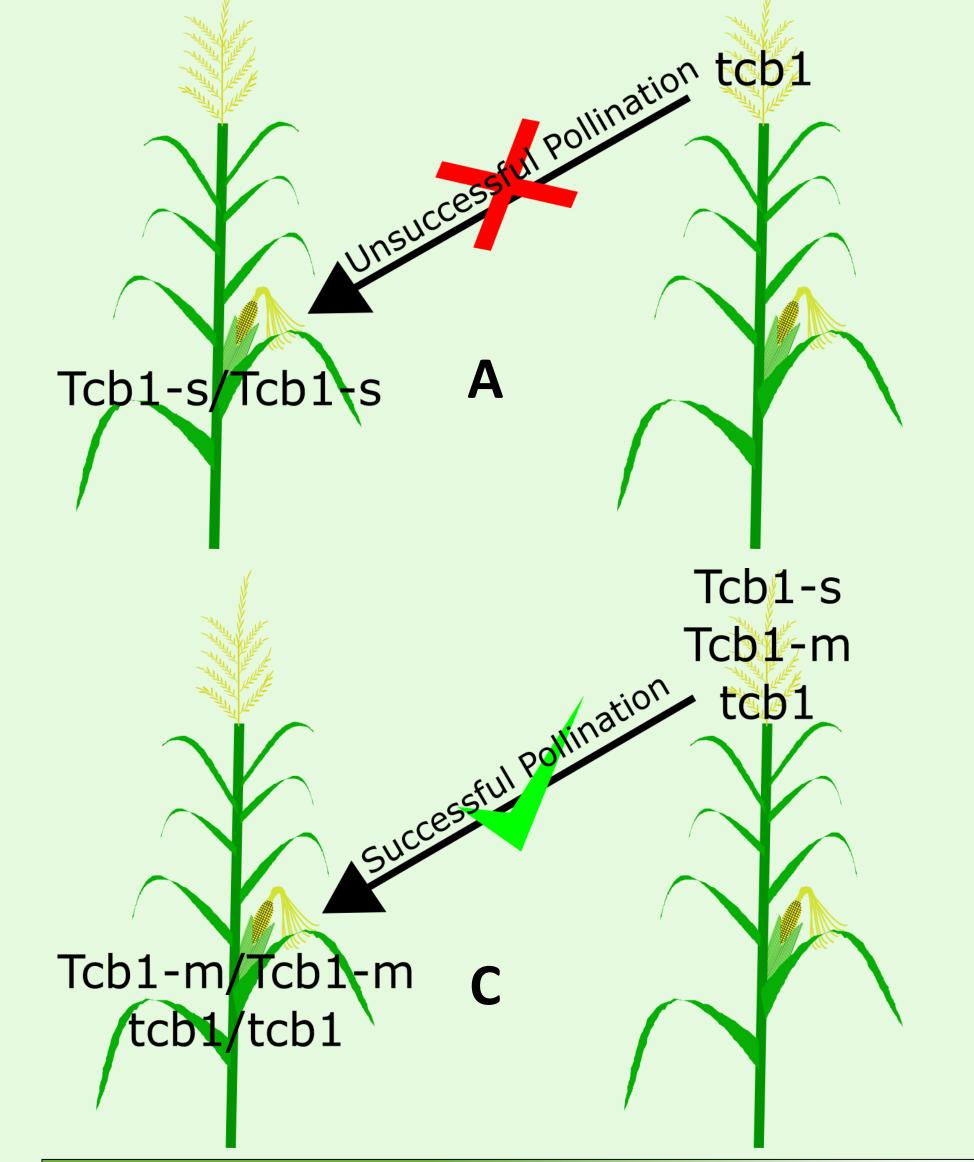
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Introduction & Background

- Windborne contamination of organic maize by genetically modified pollen can decrease profits. Currently, farmers place barrier crops, separate fields, or displace planting times to prevent cross contamination
- An easier alternative would be to introgress one of three gametophyte cross incompatibility (CI) systems: Gametophyte factor 1 (Ga1), Gametophyte factor 2 (Ga2), or Teosinte crossing barrier 1 (Tcb1)
- *Tcb1* was originally found in teosinte (*Zea mays spp. Mexicana*) and prevents fertilization from pollen with incompatible alleles (*tcb1*)
- Tcb1 has three alleles: strong allele (-s), male allele (-m), and a null allele (Fig 1)
- F1's of many maize inbreds (*e.g.* B73) x W22 *Tcb1-s* show high incompatibility with *tcb1*; Mo17 x W22 *Tcb1-s* shows a weakened incompatibility
- Using an Intermated B73 x Mo17 (IBM) recombinant inbred line (RIL) containing extensive SNP markers will help to fine map QTL of potential *Tcb1* modifiers



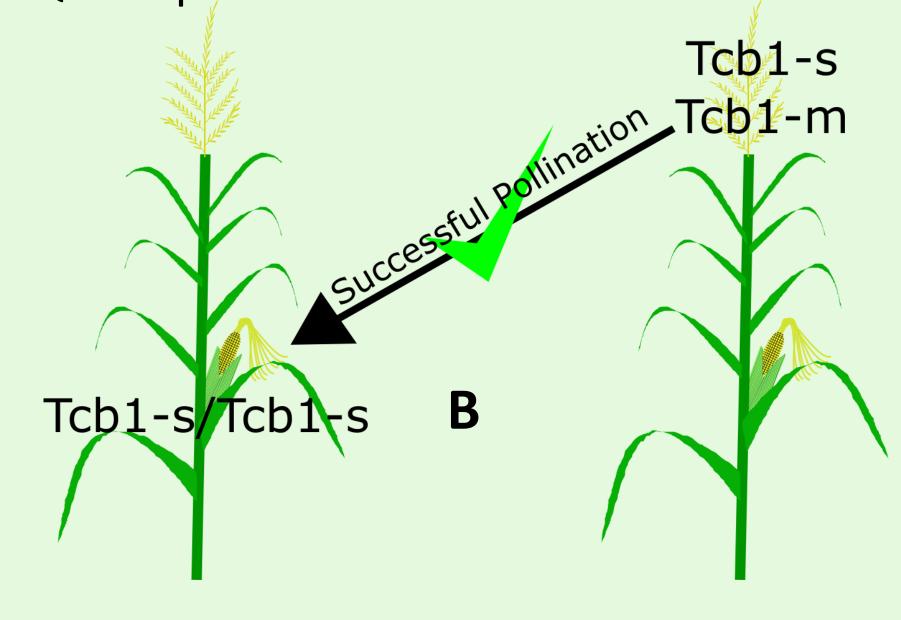


Fig 1: Teosinte crossing barrier 1 (Tcb1) cross incompatibility system. **A:** Unsuccessful pollination between Tcb1-s silks and tcb1 pollen. **B:** Successful pollination between Tcb1-s silks and Tcb1-s OR Tcb1-m pollen. **C:** Successful pollination between Tcb1-m or tcb1 silks and Tcb1-s, Tcb1-m, OR tcb1 pollen

Objectives

- Find QTL from *Tcb1* modifying loci
- From discovered QTL regions identify and isolate candidate genes
- Determine the molecular mechanism is for Tcb1 cross incompatibility

Materials and Methods

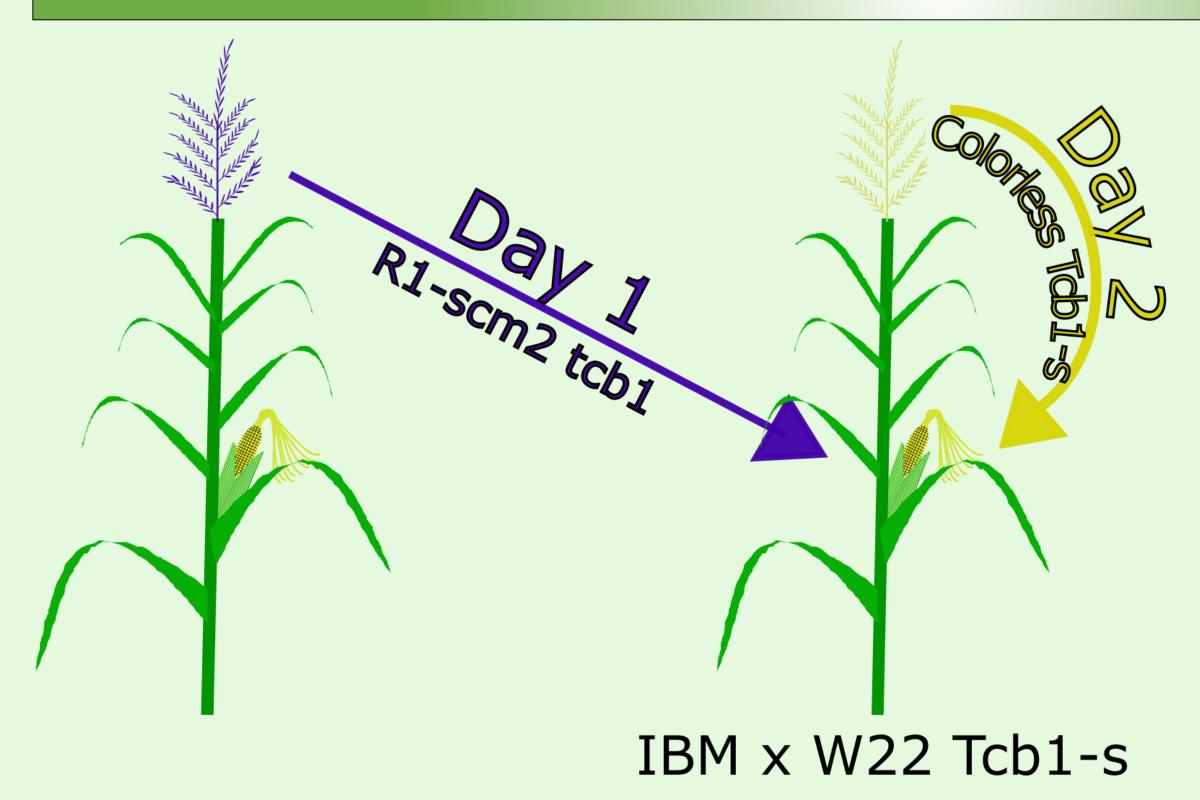


Fig 2: Pollination of IBM *Tcb1-s* silks with colored *R1-scm2 tcb1* pollen on day one (to test IBM x W22 *Tcb1-s* efficiency) and self-pollination with colorless *Tcb1-s* on day two

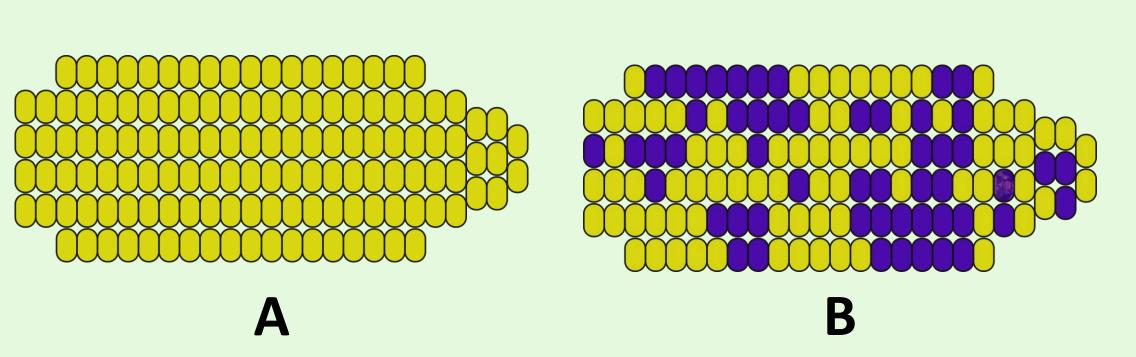


Fig 3: Predicted ear phenotypes of Fig 2 crosses. **A:** Strong effect of IBM x W22 *Tcb1-s* silks at rejecting colored *tcb1* pollen. **B:** Weak effect of IBM x W22 *Tcb1-s* silks at rejecting colored *tcb1* pollen

- Preform experiment in summer 2017 in Brookings, South Dakota
- Five plants from each IBM RIL will be crossed with *R1-scm2 tcb1* pollen and colorless self *Tcb1-s* pollen
- After harvest each RIL will be scored for the degree of tcb1 contamination
- IBM contamination data will be associated with RIL markers to determine if QTL for *Tcb1* modifiers are present

Expected Results

• Find QTL that cause *tcb1* to have lower efficacy in Mo17 F1s and identify those regions of the maize genome that support *tcb1* activity

Future Work

- Isolate and sequence candidate genes to shed light on Tcb1 incompatibility
- Quantify pollen tube growth in compatible and incompatible silks

Acknowledgements & References

Reproduction, 27(1), 19-29. doi:10.1007/s00497-013-0236-5



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