

Mapping and analysis of loci controlling quality traits in broccoli

Peter Glen Walley

Aims of the project

1. Identify and map additional QTL linked to a range of quality traits
 - a. Shelf life (turgor); QTL for yellowing already identified
 - b. Nutrient content (different components)
 - c. Stability of nutrients during storage
 - d. Head morphology

2. Analyse nutrients and potential flavour attributes
 - a. Global metabolome analysis
 - b. Key nutrients (antioxidants, glucosinolates, flavanoids, vitamins)

3. Fine scale mapping of QTL, potential gene identification
 - a. Backcross selected lines for fine scale mapping
 - b. Microarray analysis
 - c. Candidate gene nomination

Introduction – Broccoli shelf life

'short' and unpredictable shelf life

Determined by Genotype



Head remains green
For longer

Yellowing of head

Firm and attractive
Increased storage potential
Slower deterioration of nutrients
Flavour
Crunch



Soft - unattractive
Prone to impact damage
Reduced nutrients
Reduced Flavour
Unfit for retail



Broccoli x Broccoli Double Haploid (DH) lines

P1 Mar 34



P2 GD33



Mar 34 x GD33



F₁

Microspore culture



72 unique DH lines

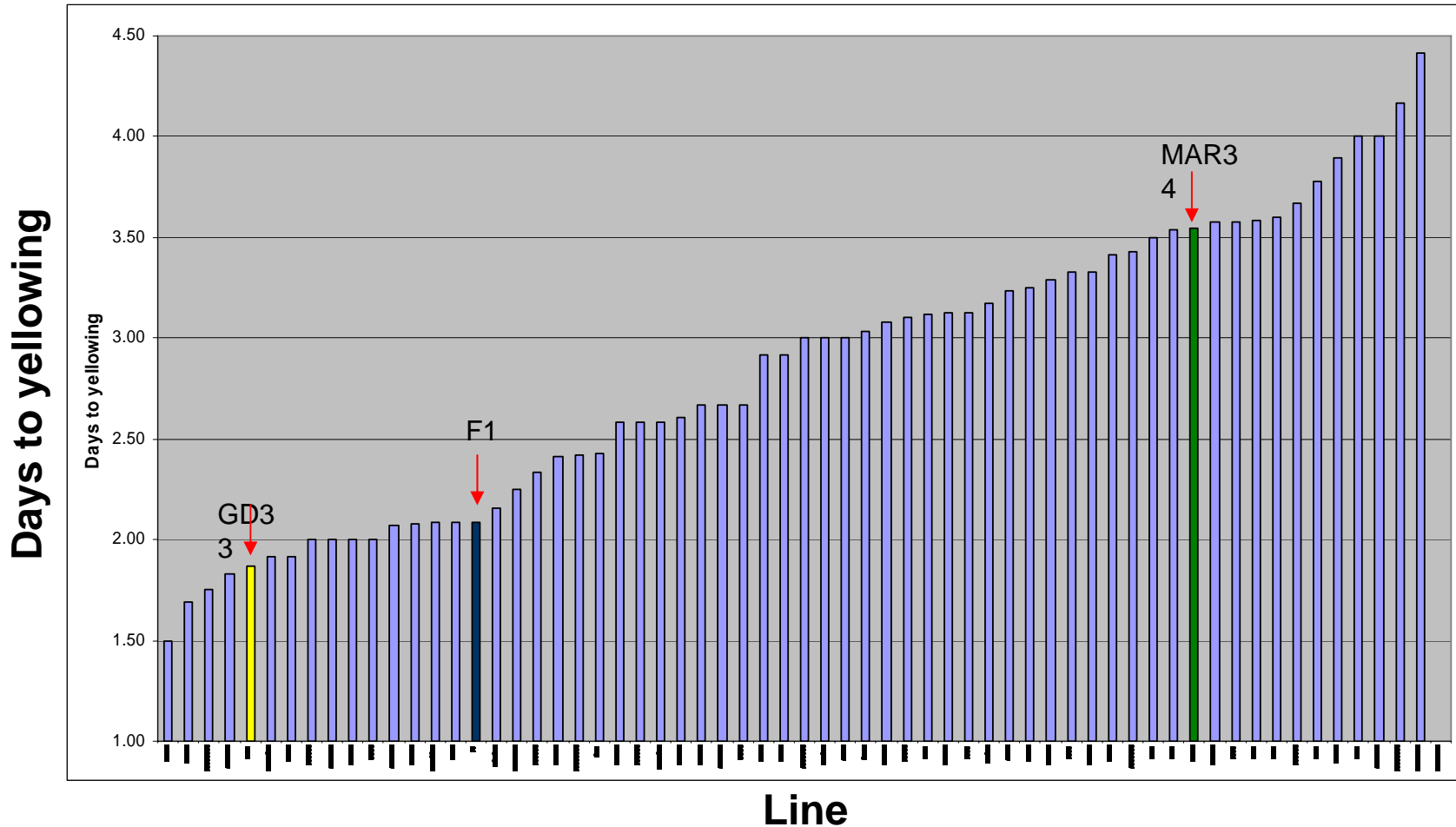
Fixed mapping resource

DH line from the F₁ cv Marathon.
Yellowing >4 days at RT

DH line from Green Duke.
Yellowing >2 days at RT

Variation in shelf life

Distribution of head yellowing for a broccoli x broccoli DH population



DH line head morphology



Head size and colour variation: (a, c)

Head shape variation: b (bracted and loose); e (domed and flat); f (flat)

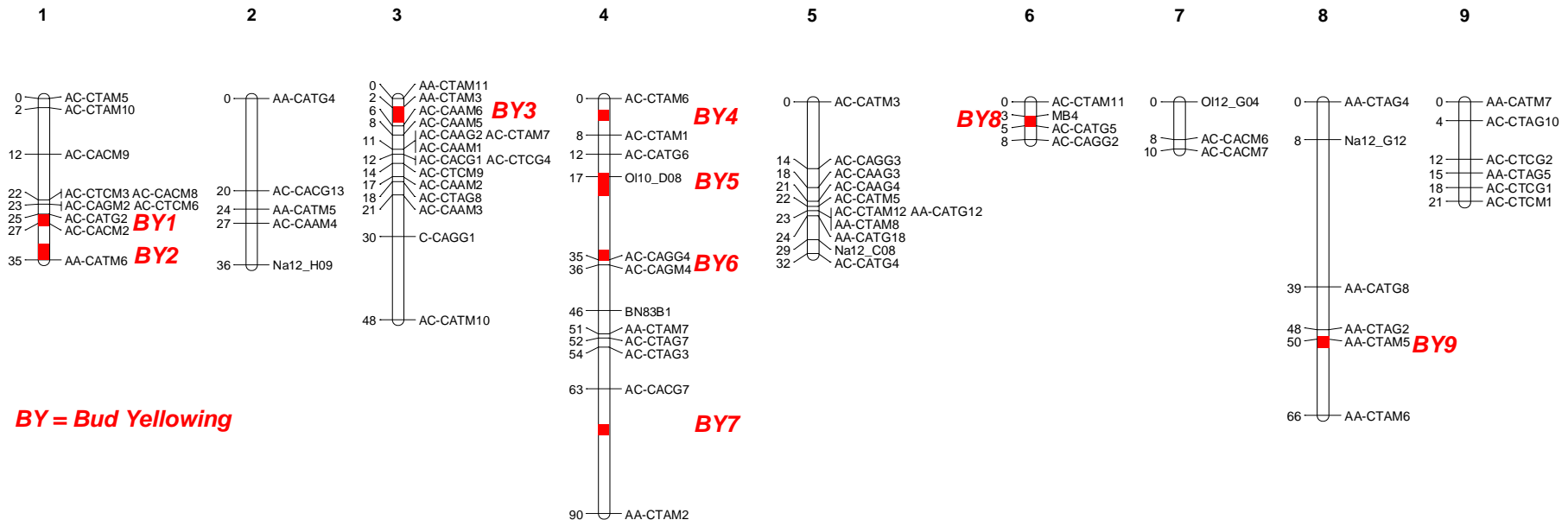
Leaf shape and size (a, c, e)

Phenotype data

- **Field trial 2007 – All lines grown together**
- **Improved assessment of shelf life quality components**
- **Metabolite analysis - QTL linked to quantity and relative composition**
 - **Metabolite loss during storage**
- **Global metabolome variation**
- **Lines identified for backcross programme**

Linkage map

- Linkage analysis resulted in a *B. oleracea* map
- Based on 94 loci derived from AFLP and SSR marker data.
- The nine linkage groups identified cover 346 cM (total length)
- Corresponds to 38.3 % of the *B. oleracea* genome



Improved Linkage map

- Skeleton linkage map

- New linkage map
- Greater AFLP, SSR coverage

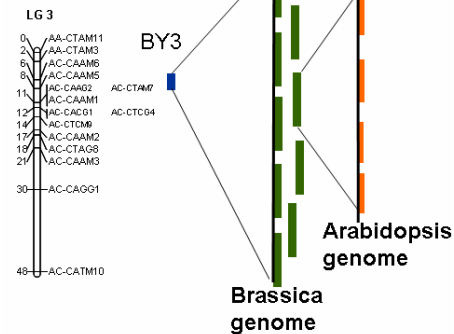
Mapped SSRs – anchor map to N x G; A x G maps

Improved AFLP marker coverage – comparative mapping

- Syngenta SSR platform

- Significant QTLs – Postharvest Yellowing

- Improved QTL analyses
Multi –trait analyses



Future direction

- BC F₁ DH lines – fine mapping of QTL
- Metabolome variation
- Microarray analysis
- Functional role of candidate gene (s)
- Markers developed for breeding programme
- QTL transferred to elite breeding lines
- Happy consumers

Acknowledgements

Team Members

WHRI

Peter Glen Walley
Vicky Buchanan-Wollaston
Dave Pink
Paul Hand
John Carder
Emma Skipper

WHRI Horticultural Services
WHRI Genome Centre

Evy Mathas

Syngenta

Ian Puddephat
Sjaak Ploeg
Charles Baxter
Martin Hill