

Figure S1. The enriched KEGG pathways between 20 and 80 DAFB (A), between 80 and 140 DAFB (B) and between 20 and 140 DAFB (C).

The X-axis represents the percentage of genes identified in this study. The same gene can be included in more than one category.

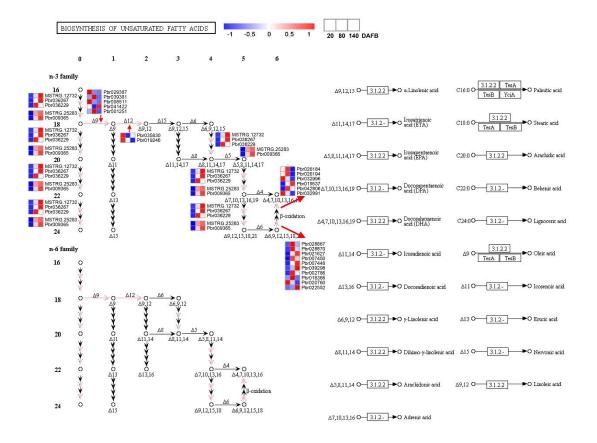


Figure S2. Unsaturated fatty acid biosynthesis pathway from the KEGG database.

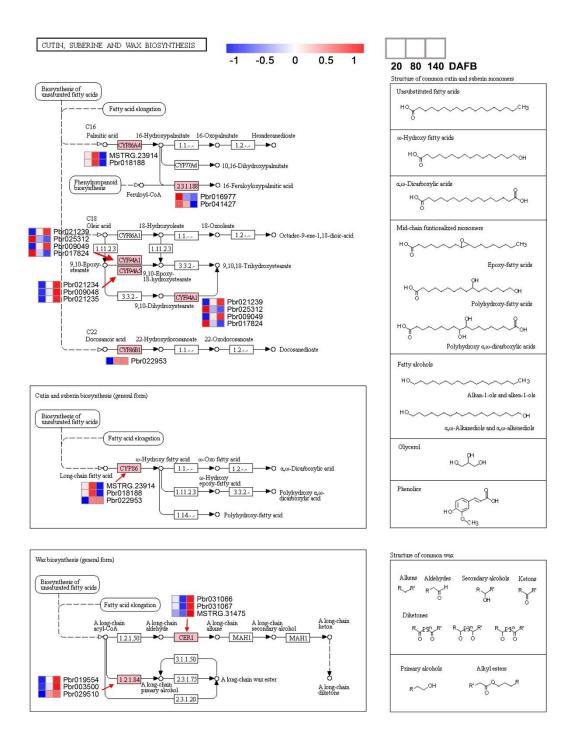


Figure S3. Cutin, suberin and wax biosynthesis from the KEGG database.

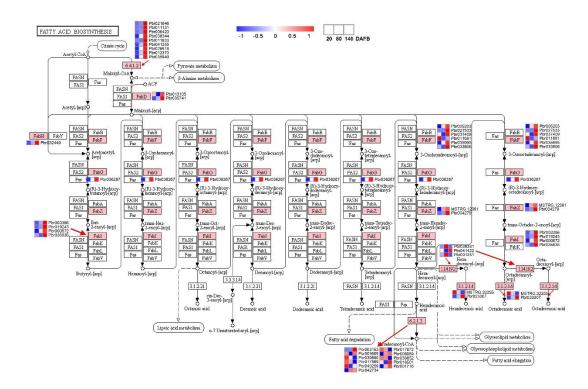


Figure S4. Fatty acid biosynthesis from the KEGG database.

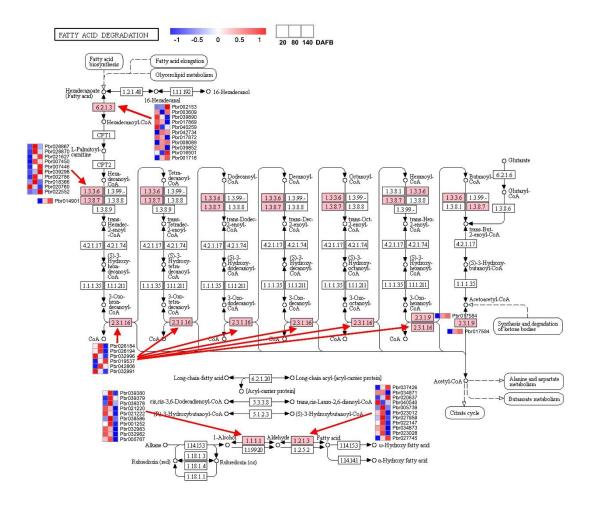


Figure S5. Fatty acid degradation from the KEGG database.

The three columns represent developmental stages at 20, 80 and 140 DAFB, respectively.

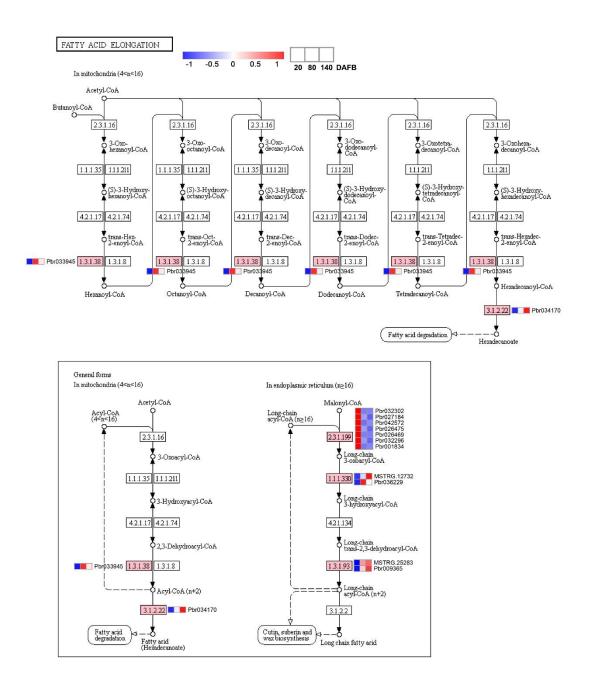


Figure S6. Fatty acid elongation from the KEGG database.

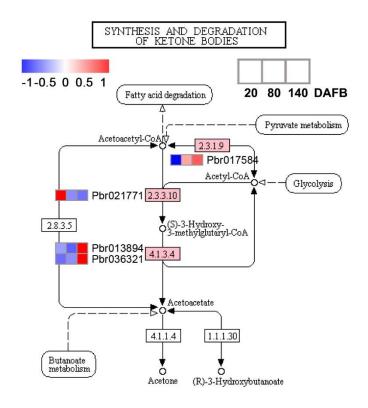


Figure S7. Synthesis and degradation of ketone bodies from the KEGG database.

The three columns represent developmental stages at 20, 80 and 140 DAFB, respectively.

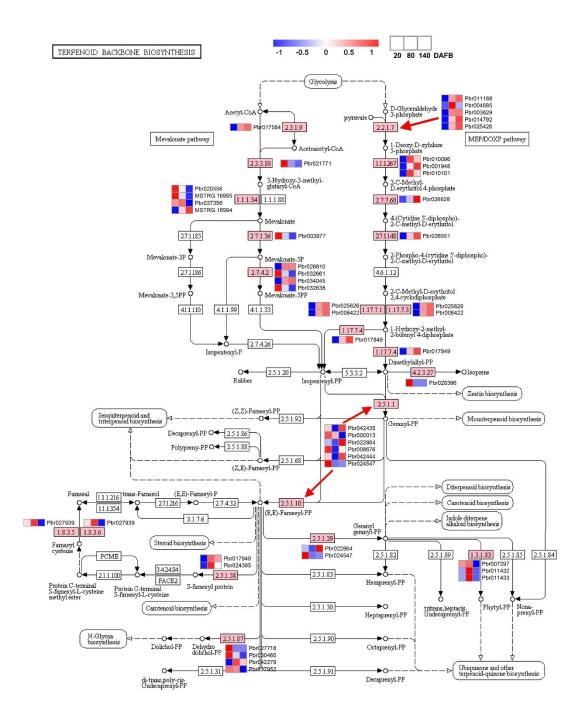


Figure S8. Terpenoid backbone biosynthesis from the KEGG database.

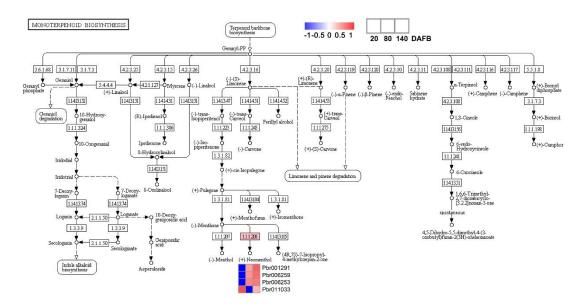


Figure S9. Monoterpenoid biosynthesis from the KEGG database.

The three columns represent developmental stages at 20, 80 and 140 DAFB, respectively.

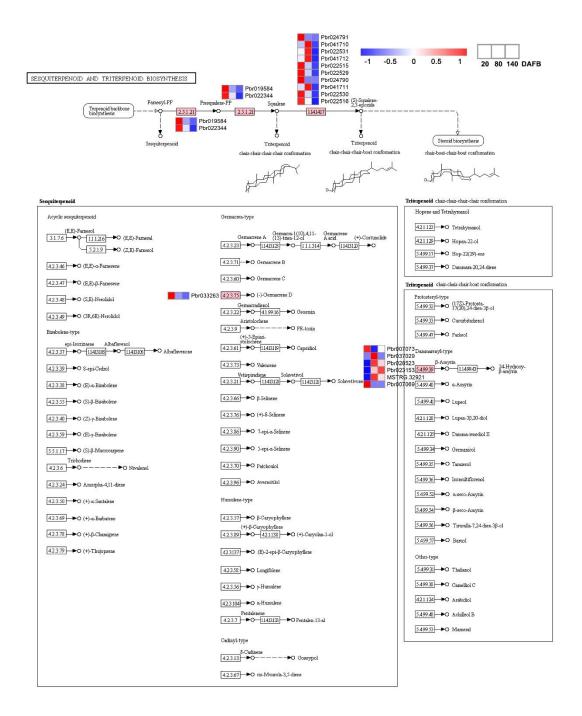


Figure S10. Sesquiterpenoid and triterpenoid biosynthesis from the KEGG database.

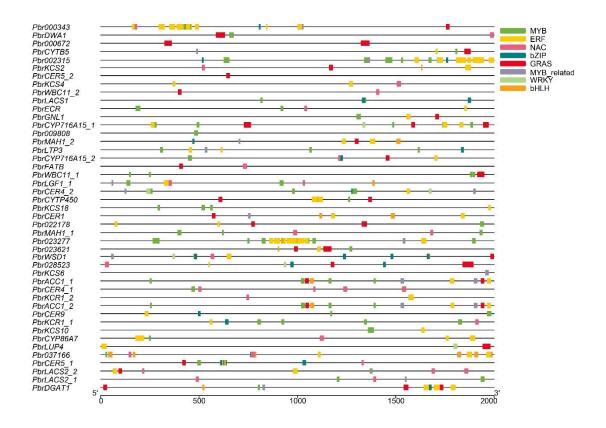


Figure S11. Prediction of the binding sites of TFs in the 2-kb upstream regulatory regions of genes

involved in cuticular wax. The different binding sites are represented by different colored boxes.

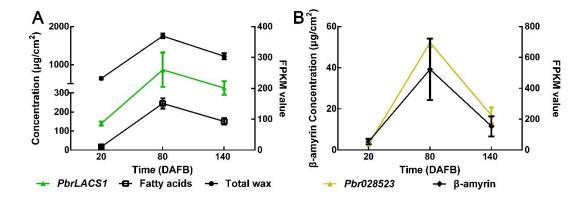


Figure S12. Wax concentrations and FPKM values of genes.

A. The total wax, fatty acid concentrations and FPKM values of *PbrLACS1* genes. B. The β -amyrin concentrations and FPKM values of *Pbr028523* genes.