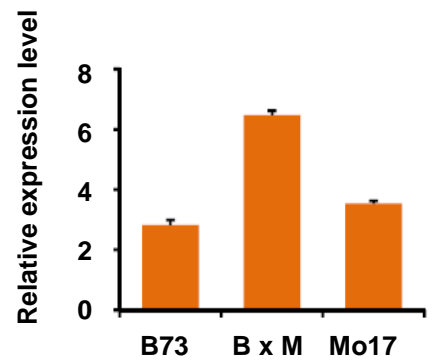
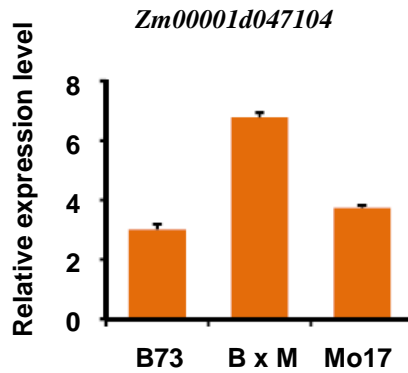
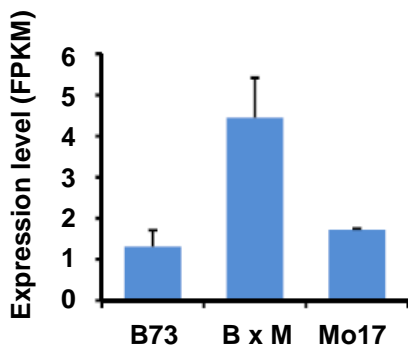
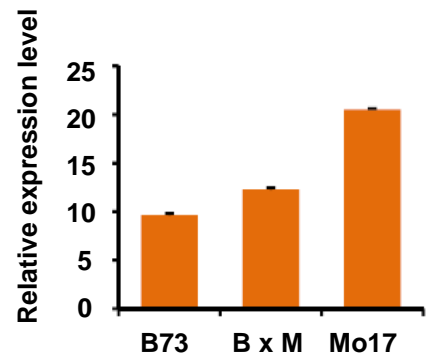
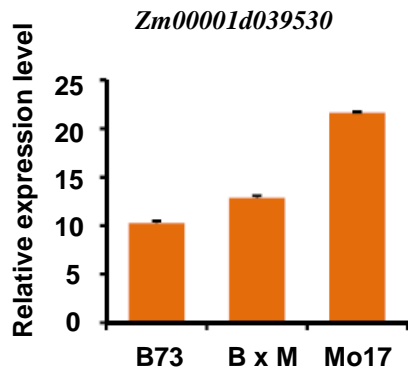
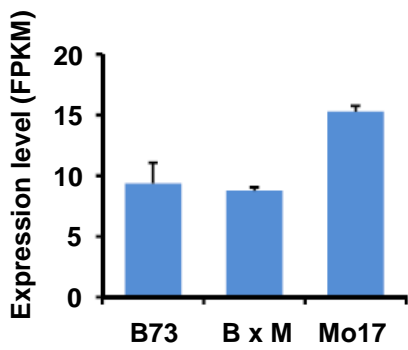
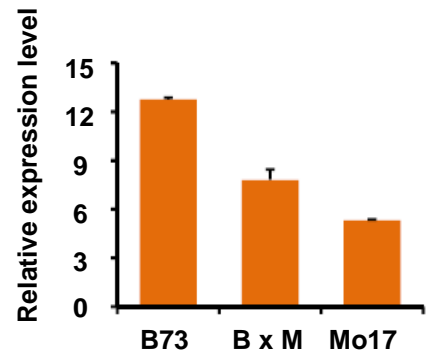
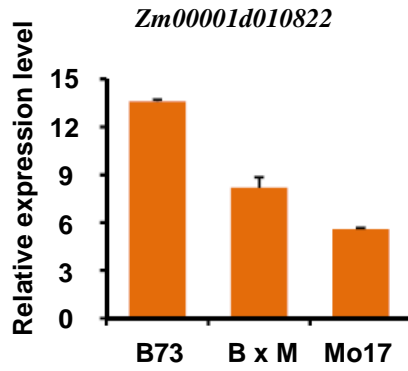
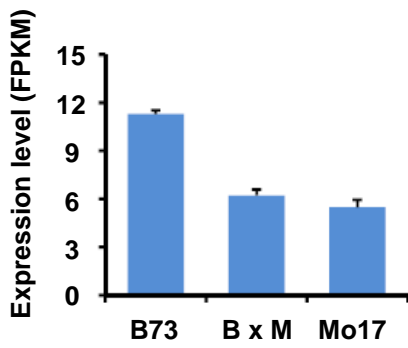
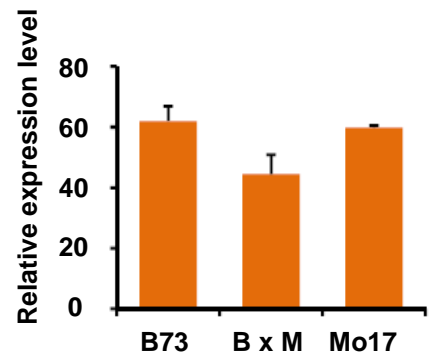
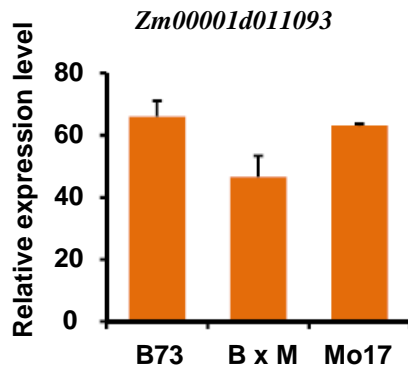
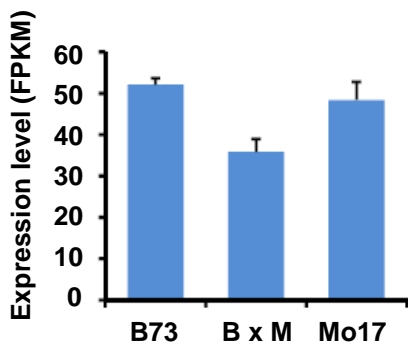


Figure S1. The repeatability between two biological replicates for RNA-seq data (A), m⁶A-seq data (B) and polysome profiling data (C) in hybrid. The Pearson correlation coefficient (r) and p -values are shown in the top left corner.



Continued

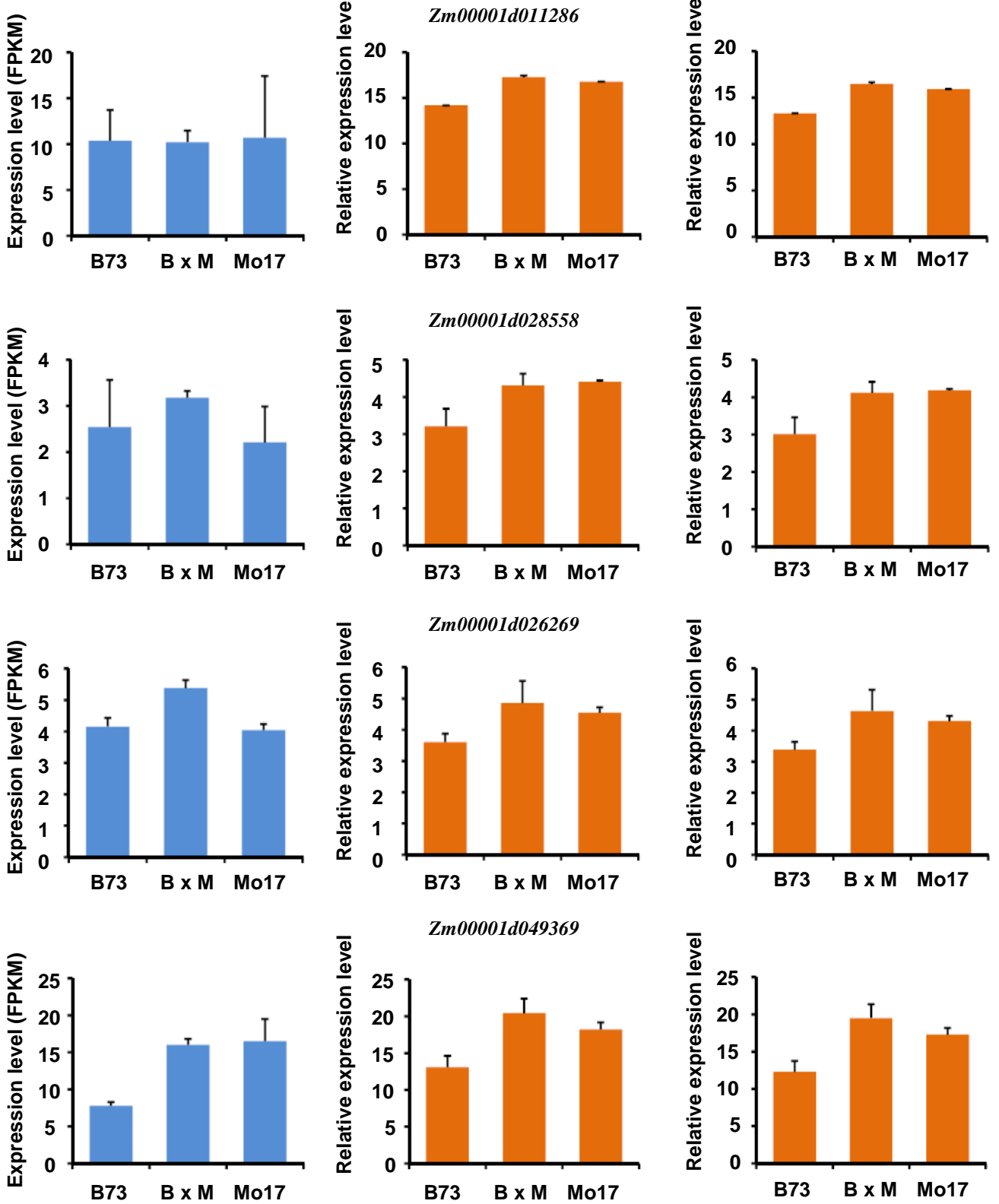
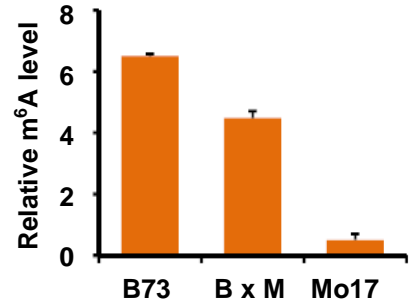
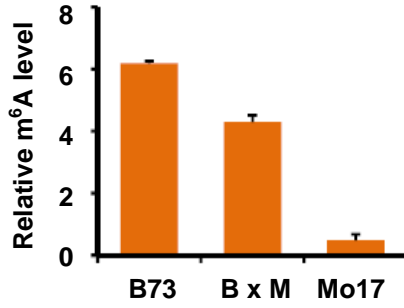
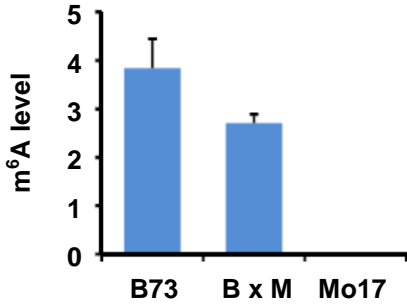
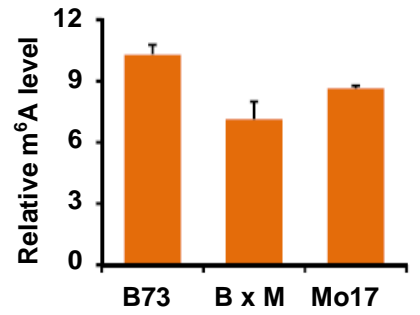
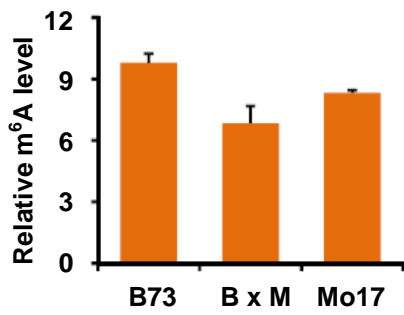
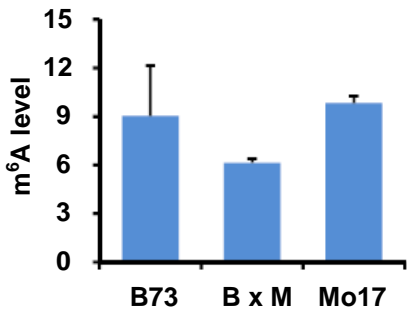


Figure S2. RT-qPCR validation of eight genes at the mRNA level in F_1 hybrid B73 \times Mo17 and its two parental lines B73 and Mo17. The mRNA levels were determined by RNA-seq (blue) or RT-qPCR (red), respectively. The relative mRNA level of each gene was compared to the internal control genes *Zm00001d034600* (middle) and *Zm00001d042939* (right). Data are represented as means + s.d. of two independent biological replicates.

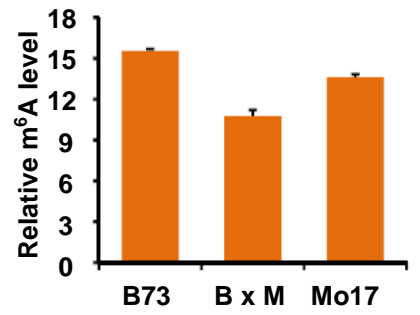
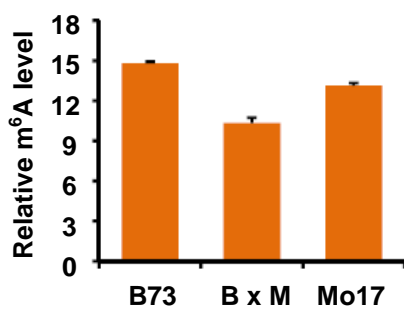
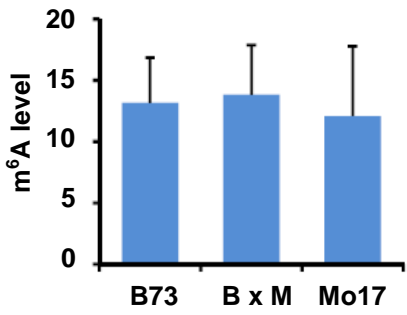
Zm00001d011093



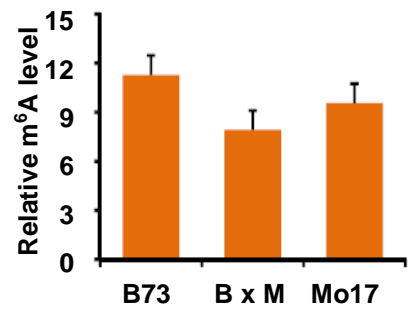
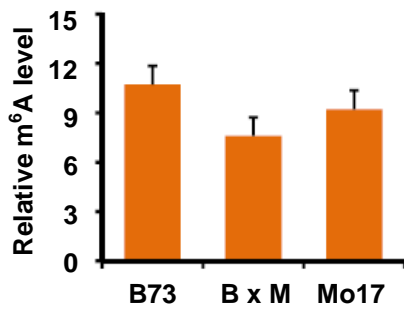
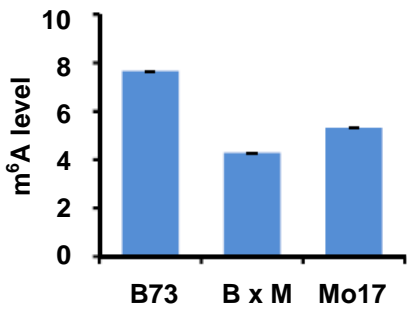
Zm00001d010822



Zm00001d039530



Zm00001d047104



Continued

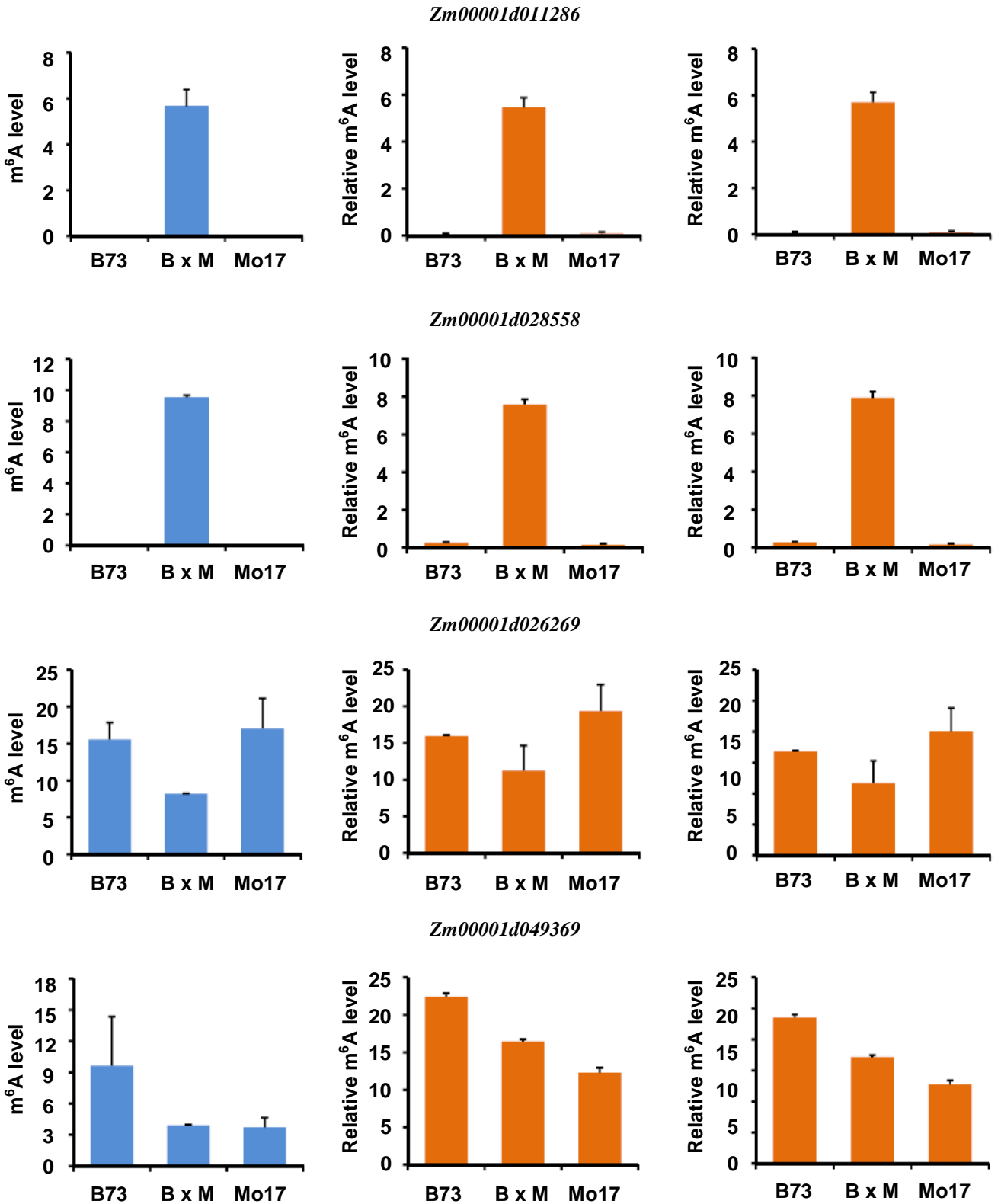
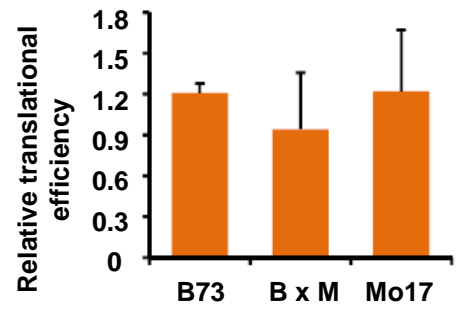
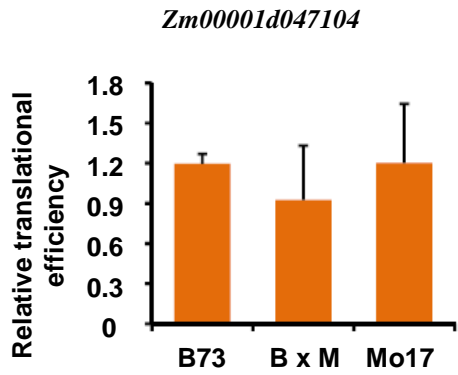
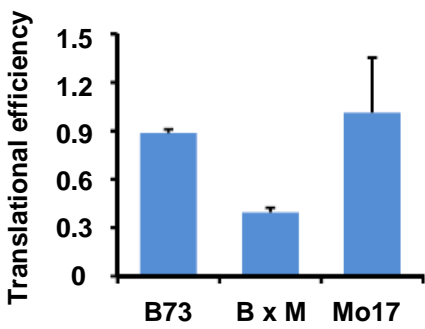
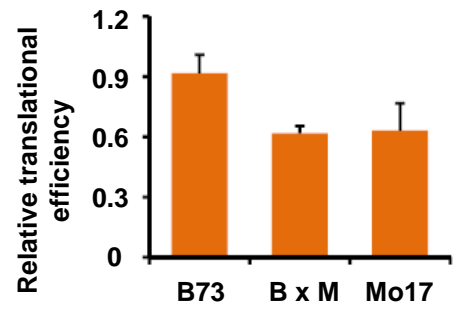
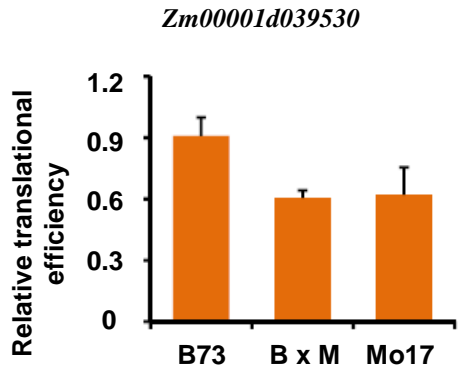
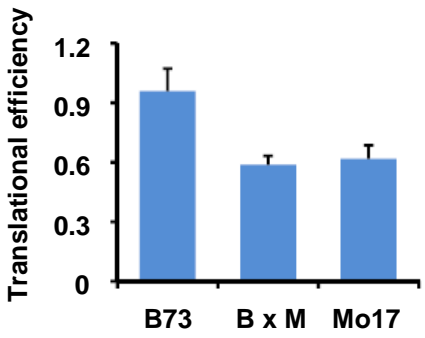
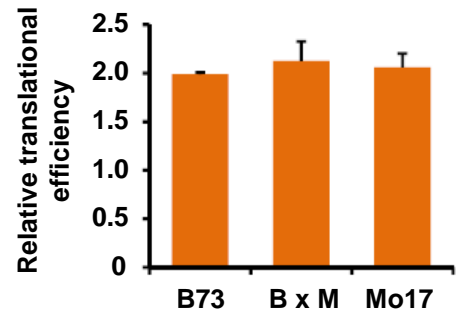
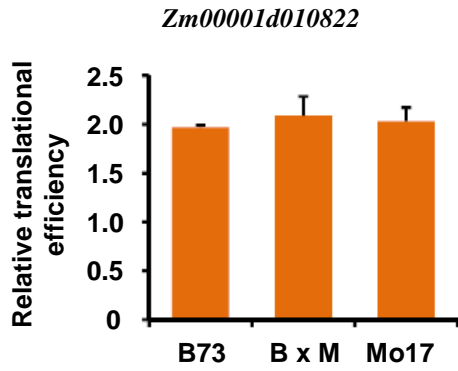
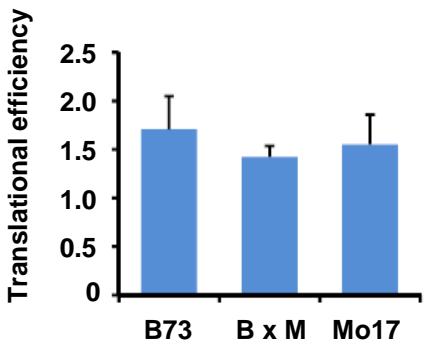
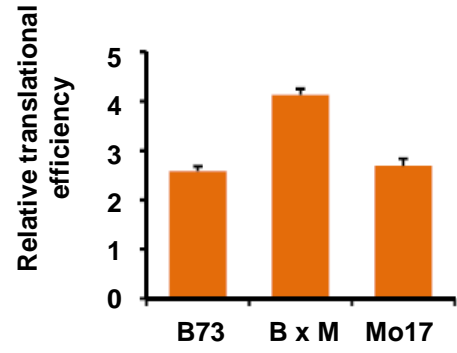
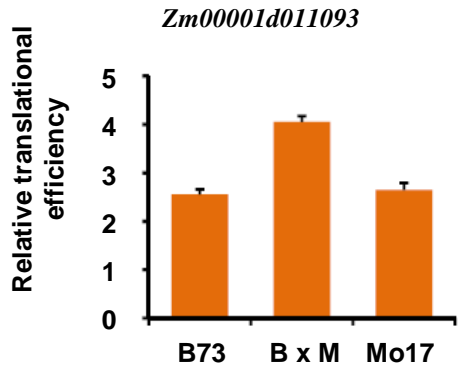
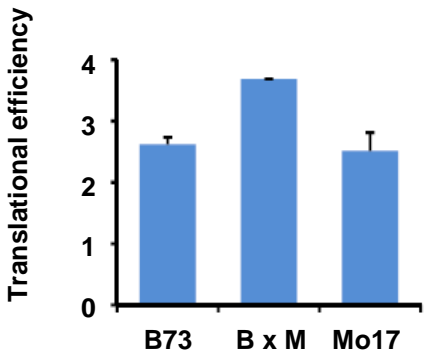


Figure S3. RT-qPCR validation of eight genes at the m⁶A level in F₁ hybrid B73 × Mo17 and its two parental lines B73 and Mo17. The m⁶A levels determined by m⁶A-seq (blue) or RT-qPCR (red), respectively. The relative m⁶A level of each gene was compared to the internal control genes *Zm00001d034600* (middle) and *Zm00001d042939* (right). Data are represented as means + s.d of two independent biological replicates.



Continued

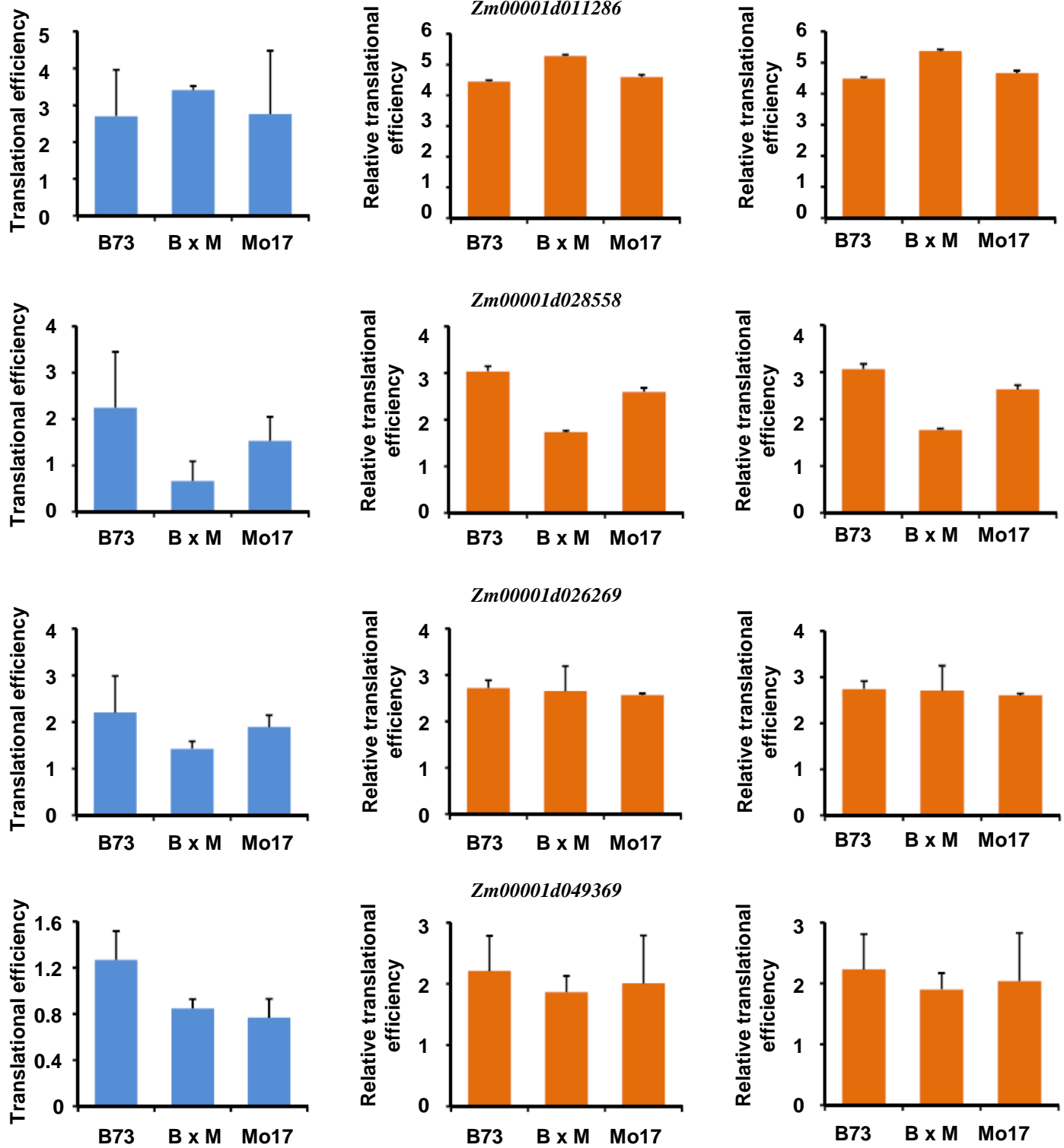


Figure S4. RT-qPCR validation of selected eight genes at the level of translational efficiency in F₁ hybrid B73 × Mo17 and its two parental lines B73 and Mo17. The translational efficiency was determined by polysome profiling sequencing (blue) or RT-qPCR (red), respectively. The relative translational efficiency of each gene was compared to the internal control genes *Zm00001d034600* (middle) and *Zm00001d042939* (right). Data are represented as means + s.d. of two independent biological replicates.

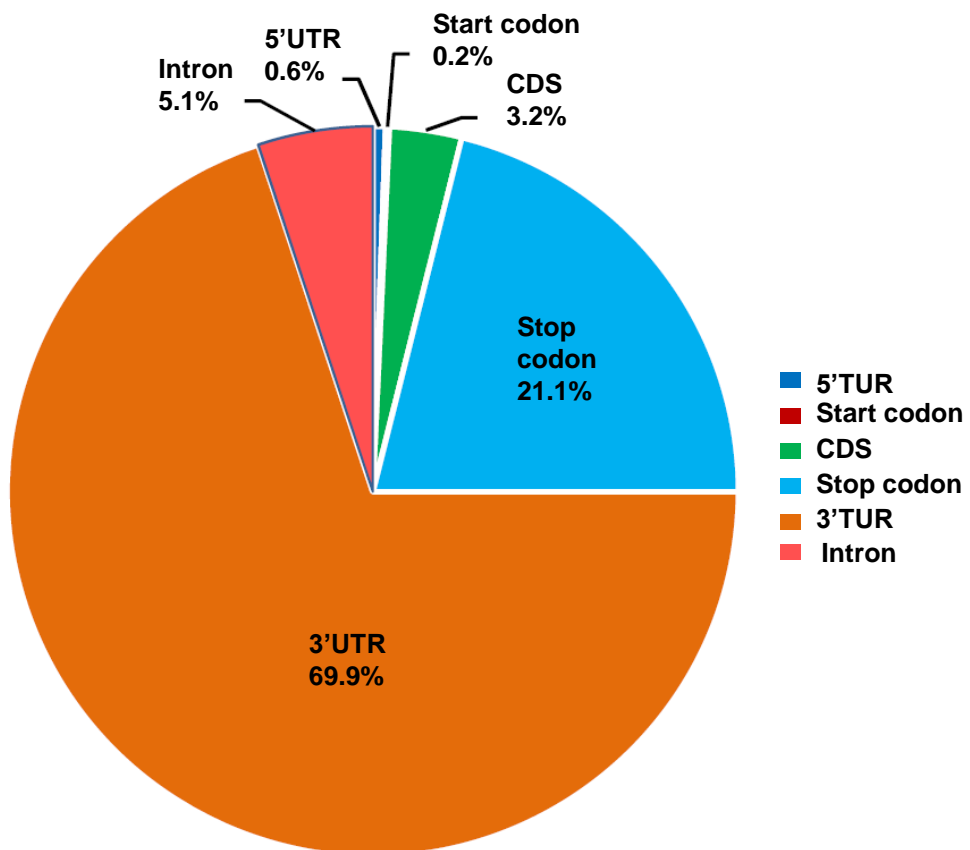


Figure S5. Pie-chart depicting the percentage of m⁶A peaks within six transcript segments in hybrid.

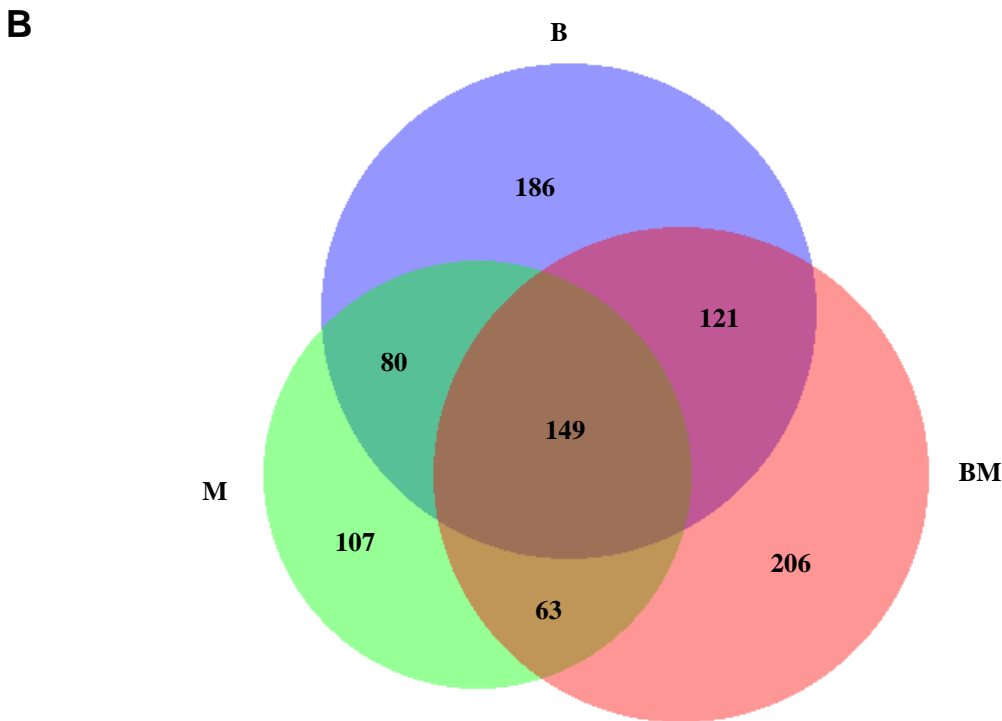
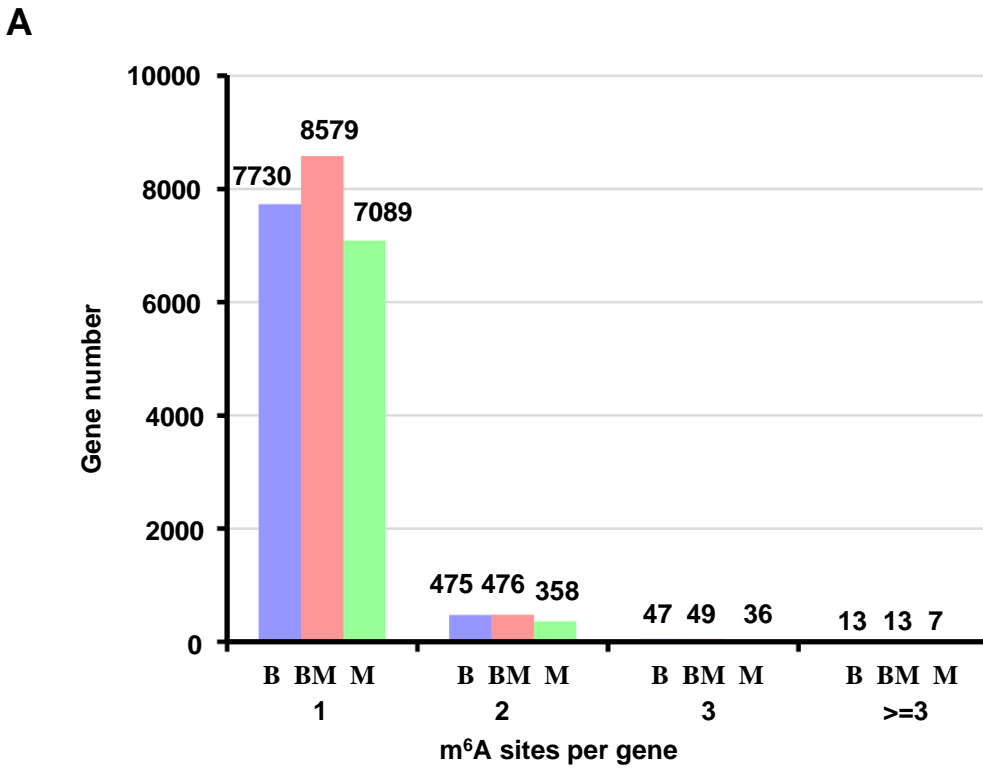


Figure S6. Comparison of gene numbers containing multiple m⁶A peaks between hybrid and parents.

(A) No. of genes containing different number of m⁶A peak between hybrid and parents.

(B) Venn diagram showing the overlap of genes containing multiple m⁶A peak per gene between hybrid and parents. B, B73; M, Mo17; BM, B73 x Mo17.