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### Authors

Bird, Kevin A  
Niederhuth, Chad E  
Ou, Shujun  
et al.

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# Replaying the evolutionary tape to investigate subgenome dominance in allopolyploid *Brassica napus*

Kevin A. Bird<sup>1,2</sup> , Chad E. Niederhuth<sup>3</sup> , Shujun Ou<sup>4</sup> , Malia Gehan<sup>5</sup> , J. Chris Pires<sup>6</sup> , Zhiyong Xiong<sup>7</sup>, Robert VanBuren<sup>1,8</sup>  and Patrick P. Edger<sup>1,2</sup> 

<sup>1</sup>Department of Horticulture, Michigan State University, East Lansing, MI 48824, USA; <sup>2</sup>Ecology, Evolutionary Biology and Behavior, Michigan State University, East Lansing, MI 48824, USA; <sup>3</sup>Department of Plant Biology, Michigan State University, East Lansing, MI 48824, USA; <sup>4</sup>Department of Ecology, Evolution and Organismal Biology, Iowa State University, Ames, IA 50011, USA; <sup>5</sup>Donald Danforth Plant Science Center, St Louis, MO 63123, USA; <sup>6</sup>Division of Biological Sciences, University of Missouri, Columbia, MO 65211, USA; <sup>7</sup>Key Laboratory of Herbage and Endemic Crop Biotechnology, Inner Mongolia University, Hohhot 010070, China; <sup>8</sup>Plant Resilience Institute, Michigan State University, East Lansing, MI 48824, USA

## Summary

Authors for correspondence:

Zhiyong Xiong

Email: xiongyz2003@aliyun.com

Robert VanBuren

Email: vanbur31@msu.edu

Patrick P. Edger

Email: edgerpat@msu.edu

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**Key words:** *Brassica napus*, DNA methylation, genomic shock, hybridisation, polyploidy, rapeseed, subgenome dominance.

- Allopolyploidisation merges evolutionarily distinct parental genomes (subgenomes) into a single nucleus. A frequent observation is that one subgenome is 'dominant' over the other subgenome, often being more highly expressed.
- Here, we 'replayed the evolutionary tape' with six isogenic resynthesised *Brassica napus* allopolyploid lines and investigated subgenome dominance patterns over the first 10 generations postpolyploidisation.
- We found that the same subgenome was consistently more dominantly expressed in all lines and generations and that >70% of biased gene pairs showed the same dominance patterns across all lines and an *in silico* hybrid of the parents. Gene network analyses indicated an enrichment for network interactions and several biological functions for the *Brassica oleracea* subgenome biased pairs, but no enrichment was identified for *Brassica rapa* subgenome biased pairs. Furthermore, DNA methylation differences between subgenomes mirrored the observed gene expression bias towards the dominant subgenome in all lines and generations. Many of these differences in gene expression and methylation were also found when comparing the progenitor genomes, suggesting that subgenome dominance is partly related to parental genome differences rather than just a byproduct of allopolyploidisation.
- These findings demonstrate that 'replaying the evolutionary tape' in an allopolyploid results in largely repeatable and predictable subgenome expression dominance patterns.

## Introduction

Hybridisation among closely related species is a widespread and recurrent evolutionary process (Arnold & Meyer, 2006; Mallet, 2007; Soltis & Soltis, 2009). By merging the genomes of independently evolved species into a single nucleus, hybridisation creates a unique opportunity for immense variability that natural selection can act upon in subsequent generations (Anderson & Stebbins, 1954; Rieseberg *et al.*, 2003). Hybridisation is known to produce transgressive phenotypes, including heterosis and novel phenotypic variation not observed in the parents (Pires *et al.*, 2004; Dittrich-Reed & Fitzpatrick, 2013). However, the hybridisation of highly diverged genomes, particularly those with different base chromosome numbers, can also lead to chromosome pairing issues during meiosis, which greatly reduces fertility. Proper bivalent pairing of homologous chromosomes in such interspecific hybrids can be restored through whole genome duplication (i.e. polyploidisation), resulting in the formation of an allopolyploid species (Charron *et al.*, 2019). This may in part

explain the high prevalence of polyploidy across flowering plants (Leitch & Leitch, 2008; Van de Peer *et al.*, 2009e; Peer *et al.*, 2009) including the multiple origins of various naturally established allopolyploids (e.g. *Tragopogon mirus* and *Tragopogon miscellus*; Soltis *et al.*, 2012), *Mimulus peregrinus*; Vallejo-Marín *et al.*, 2015), and *Elymus caninus*, (Yan & Sun, 2012). Several cultivated species, including rapeseed (*Brassica napus*; Chalhoub *et al.*, 2014), cotton (*Gossypium hirsutum*; Wendel & Cronn, 2003), and wheat (*Triticum aestivum*; International Wheat Genome Sequencing Consortium, 2014), among many others (Leitch & Leitch, 2008), are also allopolyploids.

Genome-scale analyses of recent and ancient allopolyploids led to the discovery that one of the parental species' genomes (referred to as subgenomes) often exhibits greater gene retention (Thomas, 2006), more tandem gene duplications (Edger *et al.*, 2019), higher gene expression (Schnable *et al.*, 2011) and lower DNA methylation (Woodhouse *et al.*, 2014). Collectively this phenomenon is referred to as 'subgenome dominance'. The expression bias associated with subgenome dominance has the

potential to greatly impact the observed phenotypic variation within a crop or species. In maize, the dominant subgenome contributes more to trait heritability than the nondominant subgenome (Simon *et al.*, 2017) and in strawberry, the dominant subgenome largely controls several biological pathways related to agriculturally valuable traits like fruit flavour, colour, and aroma (Edger *et al.*, 2019). A previous study demonstrated that subgenome dominance at the gene expression level can occur at the moment of interspecific hybridisation and increases over subsequent generations in the allopolyploid (Edger *et al.*, 2017). This finding agrees with theoretical work of transcription factor binding and regulatory mismatch that predicts increasing subgenome dominance over generations in newly established allopolyploids (Bottani *et al.*, 2018). Pre-existing differences between parental genomes have been shown to influence these observed subgenome dynamics in allopolyploids (Buggs *et al.*, 2014; Kryvokhyzha *et al.*, 2019). For example, analyses of diverse allopolyploid species have revealed that gene expression differences among subgenomes mirrors differences in transposable element (TE) densities in flanking regions surrounding genes (Freeling *et al.*, 2012; Cheng *et al.*, 2016; Edger *et al.*, 2019). These findings collectively suggest that subgenome dominance may be largely predetermined based on subgenome differences in certain genomic features including TE densities. This tendency for subgenome expression dominance patterns to mirror parental differences in some species has been termed ‘parental legacy’ (Buggs *et al.*, 2014).

Given that gene expression level dominance may occur instantly following the initial hybridisation event (Edger *et al.*, 2017), resynthesised allopolyploids are the ideal system to investigate the establishment and escalation of subgenome dominance. Few studies have used multiple independently derived resynthesised allopolyploids to investigate subgenome dominance (Chagué *et al.*, 2010; Combes *et al.*, 2015; Hao *et al.*, 2017; Wu *et al.*, 2018; Gaebelein *et al.*, 2019; Li *et al.*, 2019). It remains unclear the extent to which the emergence of subgenome expression dominance is a result of pre-existing characteristics of the diploid progenitors or independent and nonrecurrent events during polyploid formation. In other words, if we were to replay the tape of life (Gould, 1991) will multiple independently established allopolyploids consistently exhibit the same patterns of subgenome expression dominance (e.g. towards the same subgenome)?

Here we analysed subgenome dominance in six independent resynthesised allopolyploid *Brassica napus* ( $2n=4x=38$ ) lines formed by hybridising two doubled haploid parents from the progenitor species *Brassica rapa* (AA;  $2n=2x=20$ ) and *Brassica oleracea* (CC;  $2n=2x=18$ ) (Song *et al.*, 1995). The crop *B. napus* was formed between 7500 and 12 500 yr ago and is widely grown present-day as an oilseed crop (rapeseed), vegetable fodder crop (rutabaga) and vegetable crop (Siberian kale) (Chalhoub *et al.*, 2014; An *et al.*, 2019). The strengths of the *B. napus* polyploid system include not only having high-quality reference genomes for both diploid progenitors and *B. napus*, but also being closely related to the model plant *Arabidopsis thaliana*, allowing for the integration of diverse genomic and bioinformatic

resources (Cheng *et al.*, 2012; Chalhoub *et al.*, 2014; Parkin *et al.*, 2014; Koenig & Weigel, 2015). Furthermore, a previous analysis of the *B. napus* reference genome identified a greater number of retained genes in the *B. oleracea* (BnC) subgenome compared to the *B. rapa* (BnA) subgenome, though the overall number of genes lost was small (Chalhoub *et al.*, 2014). This is consistent with patterns observed in older allopolyploids that exhibit subgenome dominance, where the dominant subgenome retains more genes (Bird *et al.*, 2018). Lastly, because the resynthesised *B. napus* lines were made with doubled haploids, each of the independent lines started out genetically identical (Ren *et al.*, 2017). This permitted us to examine and compare the establishment of subgenome dominance across independently derived polyploid lines without the added influence of allelic variation segregating between different lines. We surveyed gene expression and methylation dynamics in each of the six resynthesised polyploid lines over 10 generations, which allowed with RNA-seq and Bisulfite-seq data to characterise gene expression and methylation differences between high confidence homoeologs over the first 10 generations. This permitted us to assess the variability of subgenome dominance during the earliest stages after polyploid formation.

## Materials and Methods

### Plant growth, tissue collection, library prep

The resynthesised *B. napus* allopolyploid lines (CCAA) were obtained from a previous study (Xiong *et al.*, 2011). Plants were grown under 23°C:20°C, 16 h:8 h, day:night cycles in a growth chamber. True leaf three was collected from all plants within 1 h, starting at 10 am (4 h into the day) and immediately flash-frozen in liquid nitrogen. Leaves were split in half for RNA and DNA isolation. Total RNA and DNA was isolated using the respective KingFisher Pure Plant kits (Thermo Fisher Scientific, Waltham, MA, USA) and quantified using a Qubit 3 fluorometer (Thermo Fisher Scientific). DNA and RNA libraries were prepared using the KAPA HyperPrep and mRNA HyperPrep kit protocols, respectively (KAPA Biosystems, Roche, USA). Bisulfite conversion was performed using the EZ DNA Methylation Kit (Zymo, Irvine, CA, USA). All libraries were submitted to a genomics facility (Beijing Nuohe Zhiyuan Technology Corp., Beijing, China) and sequenced with paired-end 150-bp reads on an Illumina HiSeq 4000 system.

### *In silico* reference genome construction

Paired-end 150-bp Illumina reads for the doubled haploid *Brassica rapa* accession IMB-218, were aligned to the *Brassica rapa* R500 reference genome using BOWTIE2 v.2.3.4.1 (Langmead & Salzberg, 2012) on default settings with the flag ‘--very-sensitive-local’. The resulting alignment files were sorted and had read groups added with PICARDTOOLS v.2.8.1. Single nucleotide polymorphisms (SNPs) were called between the R500 reference and the IMB-218 alignment using GATK v.3.5.0 Unified Genotyper, filtered to only include homozygous SNPs, and a new fasta

reference was made using GATK v.3.5.0 FastaAlternativeReferenceMaker. This IMB-218 reference genome was concatenated to the *B. oleracea* TO1000 reference genome to create an *in silico* reference genome for *B. napus* matching the two progenitor used in our study.

### Homoeologous exchange analysis

Paired-end 150-bp genomic Illumina reads were filtered with TRIMMOMATIC v.0.33 (Bolger *et al.*, 2014) to remove Illumina TruSeq3 adapters. Trimmed reads were aligned to the *in silico* *B. napus* reference genome with BOWTIE2 v.2.3.4.1 (Langmead & Salzberg, 2012) on default settings with the flag '--very-sensitive-local'. BAM files were sorted using BAMTOOLS (Barnett *et al.*, 2011) for use in downstream analyses.

The MCSscan toolkit (Tang *et al.*, 2008) was used to identify syntenic, homologous gene pairs (syntelogs) between *Brassica rapa* (reference genome R500) and *Brassica oleracea* (reference genome TO1000; Parkin *et al.*, 2014). In the synthetic polyploid these can be thought of as syntenic homoeologs. BED files based on the chromosome and start/stop position information for each subgenome were generated. For all 18 samples (6 individuals  $\times$  3 generations) read depths for the A subgenome (BnA) syntenic homoeologs were determined in BEDTOOLS (Quinlan & Hall, 2010) with BEDCOV using the R500 syntelog BED file and for the C subgenome (BnC) using the TO1000 syntelog BED file. In R v.3.4.1, read depths for each syntenic homoeolog were normalised to reads per million for subgenome of origin and the ratio of reads mapping to a syntenic homoeolog compared to the overall read mapping for a syntenic homoeolog pair was averaged over a window of 50 genes with a step of one gene.

Homoeologous exchanged regions were identified by calculating average read depth for the BnC subgenome along a sliding window of 170 (85 upstream and downstream) genes and step size of one. If 10 or more consecutive genes had read depths within a preselected range it was called a homoeologous exchange. Regions  $0 \leq \text{read depth} < 0.2$  were predicted to be in a 0BnC-to-4BnA ratio, 1BnC-to-3BnA was predicted for  $0.2 \leq \text{read depth} < 0.4$ , 2BnC-to-2BnA was predicted for  $0.4 \leq \text{read depth} < 0.6$ , 3BnC-to-1BnA for read depth between  $0.6 \leq \text{read depth} < 0.8$  and 4BnC-to-0BnA for read depth between  $0.8 \leq \text{read depth} < 1$ .

### RNA-seq analysis

Raw RNA-seq reads were filtered using TRIMMOMATIC v.0.33 (Bolger *et al.*, 2014) to remove Illumina TruSeq3 adapters and mapped to the *in silico* reference using STAR v.2.6.0 (Dobin *et al.*, 2013) on default settings. Transcripts were quantified in transcripts per million (TPM) from RNA-seq alignments using STRINGTIE v.1.3.5 (Pertea *et al.*, 2015). Because the syntelogs in the progenitor genomes are in the subgenomes of the synthetic polyploids, they can be thought of as syntenic homoeologs. To avoid dosage imbalance, only syntenic homoeologs determined to be at a 2:2 dosage balance were analysed for homoeolog

expression bias. Additionally, to remove lowly expressed genes that might be noise, syntenic homoeologs were only kept if the total TPM of the pair was  $> 10$ . Syntenic homoeolog pairs with  $\log_2$  fold change (FC)  $> 3.5$  were called BnC biased, and  $< 3.5$  were called BnA biased. This cutoff follows the practice of Woodhouse *et al.* (2014) who used a log FC cutoff of 2 to determine homoeolog expression bias, however to more confidently reduce false positives a higher FC cutoff of 3.5 was used. Because the lack of subgenome dominance would follow a normal distribution in which deviations from 0 FC are equal in either direction, a chi-squared goodness of fit test was carried out to test for normality. The R package UPSETR was used to identify and plot syntenic homoeologs shared by all lines for a given generation. Intermediate files and code to reproduce plots can be found at <https://doi.org/10.5061/dryad.h18931zjr>. For each generation, *Arabidopsis thaliana* orthologues were identified for genes showing the same subgenome bias in all six lines and the progenitors and were investigated for Gene Ontology (GO) and Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway enrichment (Ashburner *et al.*, 2000; Kanehisa & Goto, 2000) in the STRING PPI network (Szklarczyk *et al.*, 2017) using the online STRING network search application. STRING also calculated and reported average node degree, clustering coefficients and enrichment for network interactions.

### DNA methylation analysis

Whole genome bisulfite sequencing (WGBS) data was mapped to the combined *in silico* reference genome using METHYLPLY v.1.3.8 (Schultz *et al.*, 2015); using CUTADAPT v.2.3 (Martin, 2011) for adaptor trimming, BOWTIE2 v.2.3.5 (Langmead & Salzberg, 2012) for alignment, and PICARD tools v.2.20.2 for marking duplicates. The chloroplast genome is unmethylated in plants and was used as an internal control for calculating the non-conversion rate of bisulfite treatment, the percentage of unmethylated sites that fail to be converted to uracil (Lister *et al.*, 2008). Methylpy accounts for this nonconversion in calling methylated sites.

When the parental WGBS data were mapped to the combined genome (TO1000 + R500), a small fraction of reads of each sample mapped to the other subgenome, *c.* 1.3% TO1000 to *B. rapa* and *c.* 6.1% IMB218 to *B. oleracea*. We compared results from mapping of the parental data to either the combined genome or their own respective genome. There was little difference in DNA methylation levels or patterns for either parent and we therefore concluded that the impact of this mismapping was insignificant. As a further control, we created a 'mock' allopolyploid *in silico*. The TO1000 data were randomly downsampled to an equal number of read pairs as for IMB218. The two datasets were combined and mapped to the combined genome to mimic an *in silico* allopolyploid. DNA methylation levels in this 'mock' allopolyploid were either about half-way between the two parents for the whole genome or nearly identical to their respective parent at a subgenome level. If DNA methylation in the resynthesised lines is simply a combination of both parent's methylomes, then we expected global DNA methylation in the resynthesised lines to be

similar to this combined mock dataset. Deviation from this pattern would indicate global remodelling of DNA methylation.

Genome-wide levels of DNA methylation and DNA methylation metaplots were analysed as previously described (Niederhuth *et al.*, 2016) using PYTHON v.3.7.3, PYBEDTOOLS v.0.8 (Dale *et al.*, 2011), and BEDTOOLS v.2.25.0 (Quinlan & Hall, 2010). Genome-wide DNA methylation levels were calculated for each sequence context (CG, CHG, and CHH) using the weighted methylation level (Schultz *et al.*, 2012), which accounts for sequencing coverage. For gene metaplots, cytosines from 2 kilobase (kb) upstream, 2 kb downstream and within the gene/TE body were extracted. For gene bodies, only cytosines in coding sequences were used, as the presence of TEs in introns and problems of proper UTR annotation can obscure DNA methylation at start/stop sites and introduce misleadingly high levels of DNA methylation (Niederhuth *et al.*, 2016). Each of these three regions (upstream, gene body, and downstream) were then divided into 20 windows and the weighted methylation level for each window calculated and averaged for all genes. For long-terminal repeat (LTR) metaplots, the same analysis was performed, except all cytosines within the LTR body were included. Plots were made in R v.3.6.0 (R Core team, 2018) using GGPLOT2 (Wickham, 2009). All code and original analysed data and plots are available on github (<https://github.com/niederhuth/replaying-the-evolutionary-tape-to-investigate-subgenome-dominance>).

## Results

### Homoeolog expression bias

This population of resynthesised polyploids provided a unique opportunity to examine if the same subgenome would repeatedly exhibit subgenome expression dominance. Gene expression in leaves was surveyed using RNA-seq for 16 of the 18 resequenced individuals (six lines and three generations). Library construction failed for two individuals, therefore these were not able to be included in this analysis. However, representative generations from all six lines were included in these sets of analyses. Samples were aligned to an *in silico* polyploid reference genome. We restricted gene expression analyses to genomic regions with balanced gene dosage (2 : 2; AA : CC) identified using genome resequencing data for each individual to reduce the confounding factor of dosage changes in regions that have undergone homoeologous exchange. Expression patterns of the six lines were also compared with the parental *B. rapa* and *B. oleracea* genotypes to test if expression differences may exist among the diploid progenitors.

The mean expression bias ( $\log_2$  FC BnC expression/BnA expression) for homoeologs in balanced (2 : 2) regions ranged from 0.12 to 1.16 (median  $-0.50$  to  $0.96$ ), with 15 of 16 individuals having mean expression bias significantly  $> 0$  (one-way Wilcoxon–Mann–Whitney test,  $P < 2.2e^{-16}$ ; Fig. 1, Supporting Information Figs S1–S5). These results suggested a transcriptome-wide bias toward the BnC subgenome, however the magnitude of the expression bias was smaller than that observed previously in other allopolyploids (Edger *et al.*, 2017). The homoeolog expression bias between the parents was also significantly  $> 0$  in these balanced regions.

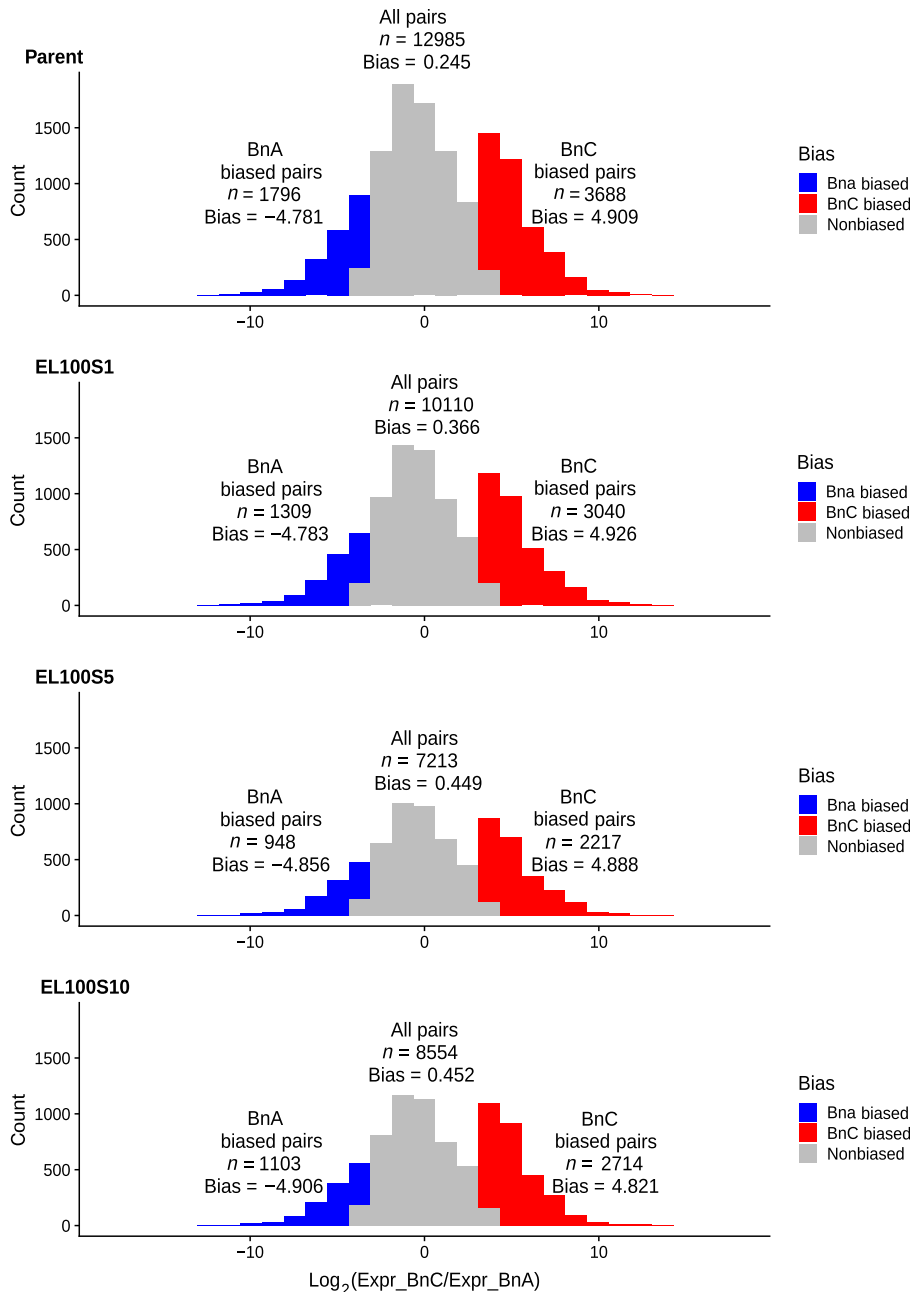
Comparing the bias difference between the parental lines and the synthetic polyploids revealed that only 5 of the 16 were significantly different from the parents (two-way Wilcoxon–Mann–Whitney test,  $P < 0.001$ ).

We examined homoeolog pairs that were biased toward either subgenome ( $\log_2$  FC  $> |3.5|$ ) and found that there were significantly more BnC-biased homoeolog pairs than expected in all individuals ( $\chi^2$ -test,  $\chi^2 > 170$ ,  $df = 1$ ,  $P < 2.2e^{-16}$ ; Figs 1, S1–S5; Table S1). This indicated that there was a bias in homoeolog expression on a gene-by-gene basis towards the BnC subgenome. This BnC homoeolog bias was consistent across generations. We also found that homoeolog bias in the synthetic polyploids was significantly different than existing expression bias in the parents for 7 of 16 individuals ( $\chi^2$  test,  $\chi^2 > 12.459$ ,  $df = 2$ ,  $P < 0.003125$ ; Figs 1, S1–S5; Table S2). In five of those seven individuals, there were more BnC-biased homoeologs than expected based on expression biases of the progenitor genomes. A bimodal distribution was observed when comparing BnA and BnC subgenome expression, the rightmost distribution being largely due to homoeolog pairs with a lack of BnA expression.

Next, we investigated whether individual gene pairs were biased in the same direction across the six lines. Due to the stochastic nature of homoeologous exchanges, dosage of a gene pair may differ between lines. To adjust for this, we first looked only at genes found in 2 : 2 dosage for all lines in a generation, resulting in 6917, 3574 and 2252 homoeologous pairs for generations S1, S5 and S10, respectively. The reduction in 2 : 2 homeologs over time suggested that homoeologous exchange is a dynamic and variable process across independent lines and that it reaches some stability or equilibrium over successive generations. The frequency of homoeologous exchanges across each chromosome of this synthetic *B. napus* population was previously reported based on cytogenetic analyses (Xiong *et al.*, 2011). A majority of BnC-biased gene pairs was biased toward the BnC subgenome in all synthetic lines for each generation and in the parents (S1 = 1806 (75%), S5 = 772 (71%), and S10 = 602 (70%); Fig. 2a–c). In the 1<sup>st</sup> generation, 36 gene pairs (1.5%) were uniquely dominant in only the parents and 32 (1.3%) were dominant in all six synthetic lines but not the parent (Fig. 2a). In the 5<sup>th</sup> and 10<sup>th</sup> generation, there was a similar number of BnC dominant homoeologs that were dominant in only the parents (S5 = 17 (1.5%), S10 = 14 (1.6%)) and dominant in all six lines but not the parent (S5 = 17 (1.5%), S10 = 13 (1.5%); Fig. 2c). Similar patterns were observed for BnA-biased homoeologs with most genes showing similar bias in all lines across generations and the parents (S1 = 698 (61%), S5 = 401 (58%), S10 = 221 (51%), respectively, Fig. S6a–c), and a consistently low number of genes biased in all six lines but not the parents in each generation (S1 = 28 (2.5%), S5 = 15 (2.2%), S10 = 10 (2.3%)) and only biased in the parents (S1 = 41 (3.6%), S5 = 22 (3.2%), S10 = 18 (4.1%); Fig. S6a–c).

Lastly, we performed GO (Ashburner *et al.*, 2000), KEGG pathway (Kanehisa & Goto, 2000) and network enrichment analyses in the *Arabidopsis thaliana* protein–protein interaction

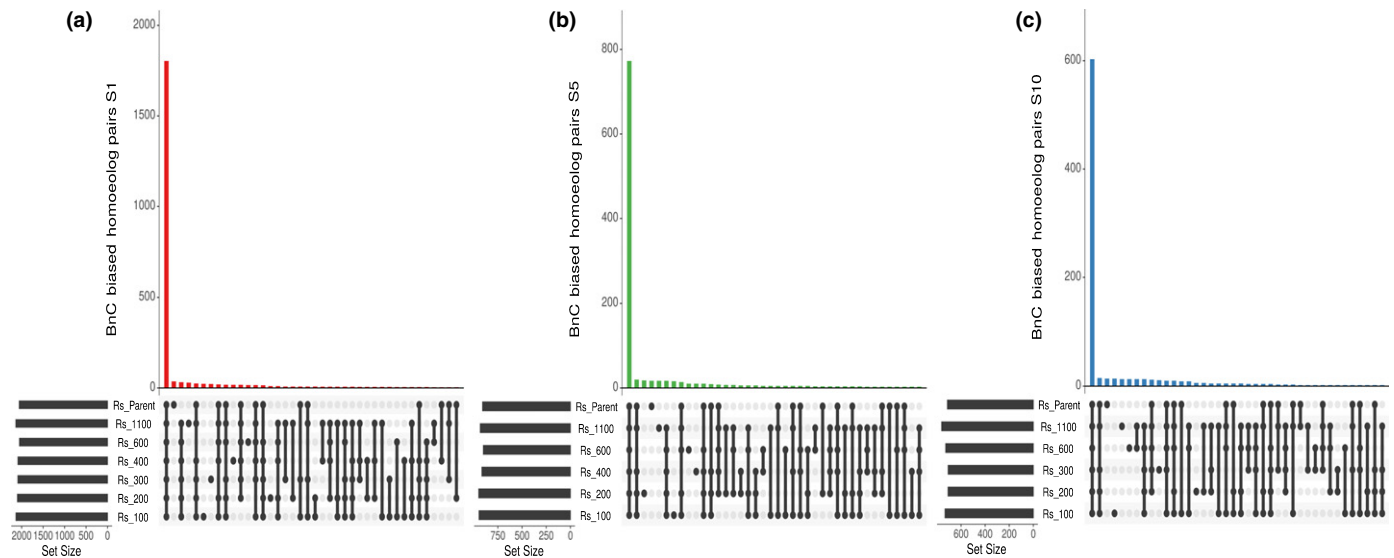




**Fig. 1** Homoeolog expression bias in resynthesised *Brassica napus*. Distribution of homoeolog bias in the parent and three generations of line EL100. Red regions indicate BnC-biased homoeologs with  $\log_2$  expression fold change  $> 3.5$  and blue regions indicate BnA-biased homoeologs with  $\log_2$  expression fold change  $< -3.5$ .

network (Szkłarczyk *et al.*, 2017) to determine if *Arabidopsis thaliana* homologues of BnC and BnA-biased gene pairs were enriched with certain biological functions. Placing biased homoeologs in a network context allowed us to determine the extent to which the genes interact with each other, the average number of connections a gene has with other genes (average node degree), and the extent to which genes are clustered together in the network. In all generations, BnC-biased gene pairs showed enrichment for interactions in the PPI network in the 1<sup>st</sup>, 5<sup>th</sup> and 10<sup>th</sup> generations (all three generations;  $P < e^{-16}$ ) with high average node degree (S1 = 20.6, S5 = 12.1, S10 = 7.99) and clustering coefficient (S1 = 0.386, S5 = 0.385, S10 = 0.385). GO terms for BnC-biased gene pairs were highly enriched with core metabolic processes, including photosynthesis and organellar and ribosomal

functions. For example, in the 1<sup>st</sup> generation, 263 genes were annotated as being active in the chloroplast (Table S3). BnC-biased gene pairs were also significantly overrepresented in KEGG pathways annotated for amino acid biosynthesis, MAPK signalling and photosynthesis (Tables S3–S5). The enrichment for organellar functions for BnC-biased pairs may be expected, given that the BnC subgenome is the maternal progenitor of the resynthesised lines. This suggests that subgenome dominance may, in part, be related to maintenance of balanced nuclear–organellar interactions. By contrast, BnA-biased gene pairs showed no enrichment for any GO terms or KEGG pathways. However, there was an observed enrichment for interactions in the PPI network for BnA-biased gene pairs in generations 1 ( $P = 9.75e^{-07}$ ) and 5 ( $P = 0.00681$ ), but not generation 10 ( $P = 0.16$ ). Lower



**Fig. 2** Common shared biased homoeolog pairs in resynthesised *Brassica napus*. UpSet plot showing BnC-biased homoeologs pairs for generations 1 (a), 5 (b), and 10 (c) that are shared or unique among comparisons across all six lines for the three sampled generations. This analysis was restricted only to homoeolog pairs in 2 : 2 balance in all six lines.

network statistics were also observed for average node degree ( $S1 = 1.13$ ,  $S5 = 0.912$ ,  $S10 = 0.338$ ) and average clustering coefficient ( $S1 = 0.270$ ,  $S5 = 0.235$ ,  $S10 = 0.209$ ) for BnA-biased gene pairs. In summary, subgenome expression dominance in our resynthesised *Brassica napus* was biased towards a set of highly interconnected BnC genes that were enriched for a wide variety of biological processes.

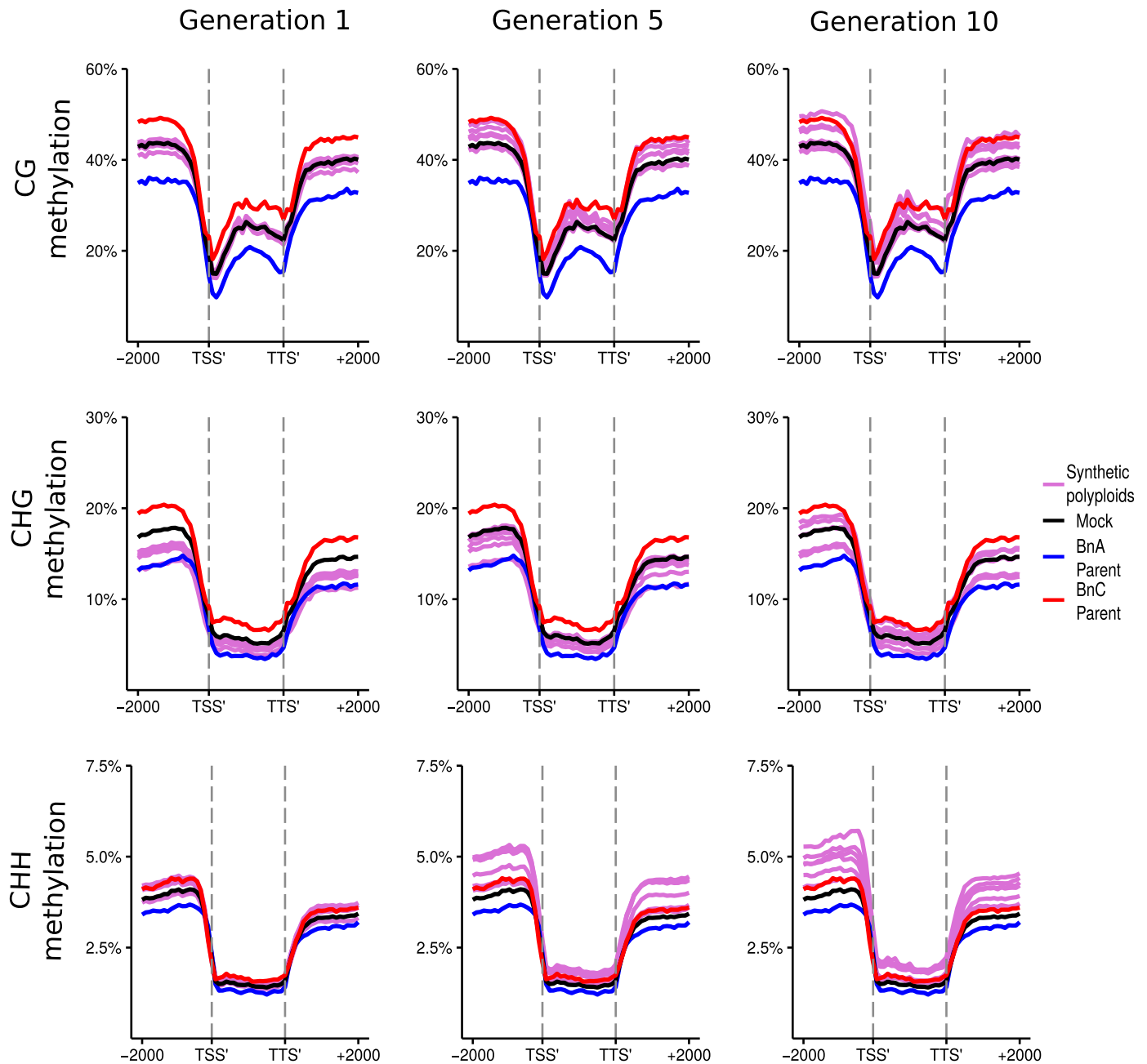
## DNA methylation

Cytosine methylation is involved in defining regions of chromatin, silencing TEs, maintaining genome integrity and can affect gene expression. DNA methylation itself is shaped by factors such as gene expression and the underlying DNA sequence (Niederhuth & Schmitz, 2017). To understand how DNA methylation evolves following polyploidy and may contribute to subgenome dominance, DNA methylation was assessed using WGBS in the parents and allopolyploids (Cokus *et al.*, 2008; Lister *et al.*, 2008). Three sequence contexts are generally recognised for DNA methylation in plants: dinucleotide CG (or CpG), and trinucleotide CHG and CHH (where H = A, T or C). Methylation of these contexts is established and maintained by different molecular pathways and their associations with gene expression differs based on the pattern of DNA methylation (Niederhuth & Schmitz, 2017). Coding sequences methylated in all three contexts (CG, CHG and CHH) are associated with transcriptional silencing. By contrast, so-called gene body methylated (gbM) genes exhibited relatively high and broad expression patterns. GbM is a specific pattern of CG-only DNA methylation in gene bodies, with depletion at the transcriptional start site (TSS) and termination site. Finally, unmethylated genes showed variable expression patterns and were often developmentally and environmentally responsive genes. Regardless of context, depletion of methylation around the TSS appears to be important for

transcription as even CG methylation in this region is associated with lower expression.

In line with current conceptual models of subgenome dominance (Woodhouse *et al.*, 2014; Garsmeur *et al.*, 2014; Bird *et al.*, 2018; Alger and Edger, 2020), we might expect to see generally lower methylation, especially in the CHH context, upstream of genes or in TEs in the dominant subgenome (Edger *et al.*, 2017, 2019; Renny-Byfield *et al.*, 2017; Zhao *et al.*, 2017). The predicted mechanism at play is that silencing of TEs through the RNA-directed DNA methylation pathway affects expression of nearby genes, as part of the trade-off described by Hollister and Gaut (2009). Although as noted above, these are not the only methylation patterns that can explain expression differences among homoeolog pairs.

In all methylation contexts, the BnC subgenome progenitor showed higher methylation levels than the BnA subgenome progenitor (Figs 3, 4). For genes, CG methylation levels are predominantly at mid-parent levels, but appear to increase over time, most notably upstream and downstream of gene bodies. By the 10<sup>th</sup> generation, some lines showed upstream and downstream CG methylation closer to the BnC progenitor, while others showed methylation levels in line with the mid-parent level. CG methylation of LTR retrotransposons showed a similar trend as the genes, with 1<sup>st</sup> generation methylation levels matching mid-parent levels and increasing over time, particularly in the LTR body (Figs 3, 4). In the 1<sup>st</sup> generation, CHG methylation fell between the mid-parent value and the BnA progenitor for both genes and LTRs. In the 5<sup>th</sup> and 10<sup>th</sup> generations, CHG methylation progressively increased to more reflect the mid-parent value. As with CG methylation, at the 10<sup>th</sup> generation, we saw two clusters, one with CHG methylation levels above the mid-parent value and one with methylation below the mid-parent value in the upstream and downstream regions of genes (Figs 3, 4). CHH methylation patterns were even more striking. In genes and



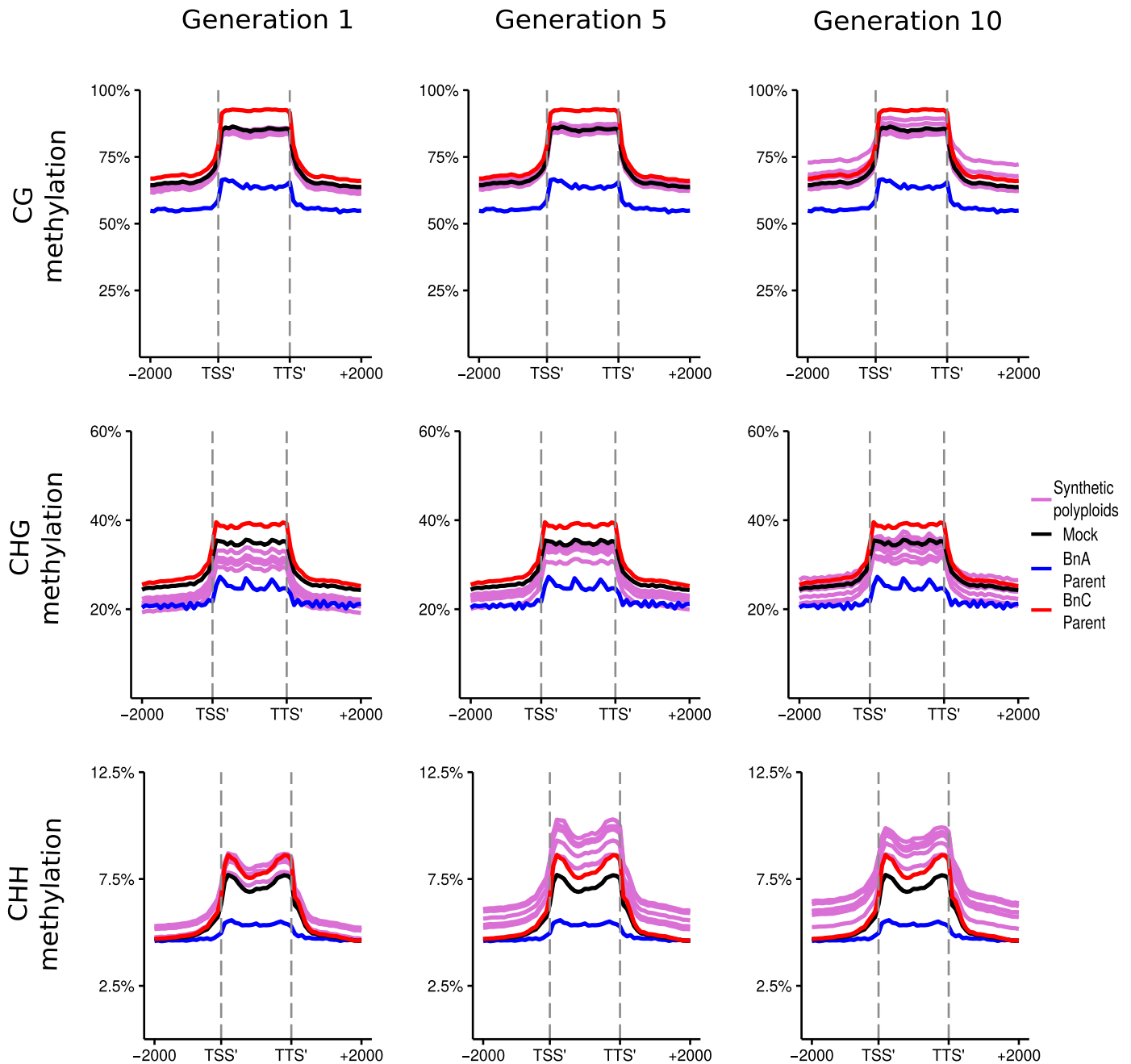
**Fig. 3** DNA methylation of genes in resynthesised *Brassica napus*. Metaplots of CG, CHG and CHH mean weighted methylation of all annotated gene models for generations 1, 5 and 10 assessed using Bisulfite-seq. The six resynthesised lines are shown in purple, the *Brassica oleracea* progenitor is shown in red, *Brassica rapa* progenitor in blue, and the *in silico* mock polyploid in black. Methylation levels are shown for the transcription start site (TSS'), gene body, transcription termination site (TTS') and 2 kb upstream and downstream of the TSS and TTS.

LTRs, the 1<sup>st</sup> generation shows methylation levels between the mid-parent and the BnC progenitor, with methylation increasing and surpassing the BnC progenitor levels in the 5<sup>th</sup> and 10<sup>th</sup> generation (Figs 3, 4). As with CG and CHG methylation, increased variation in the flanking regions by the 10<sup>th</sup> generation is observed, however all lines still showed a consistently higher methylation than the BnC progenitor.

Analysed by subgenome, methylation levels in the resynthesised lines were on average lower for the BnA subgenome than the BnC subgenome for every methylation context and

generation observed (Figs 5, 6). Additionally, there was a more pronounced increase in CG, CHG, and CHH methylation levels at the boundaries of flanking regions and the LTR body for the BnC subgenome compared with the BnA subgenome (Figs 7, 8). Syntenic genes are those that have remained in identical positions in the respective subgenomes, identified based on the collinearity of annotated genes across a region in the genome. Comparing syntenic genes to nonsyntenic genes, CG, CHG and CHH methylation levels in gene bodies were both higher and more variable for nonsyntenic genes. Differences in the changes in gene

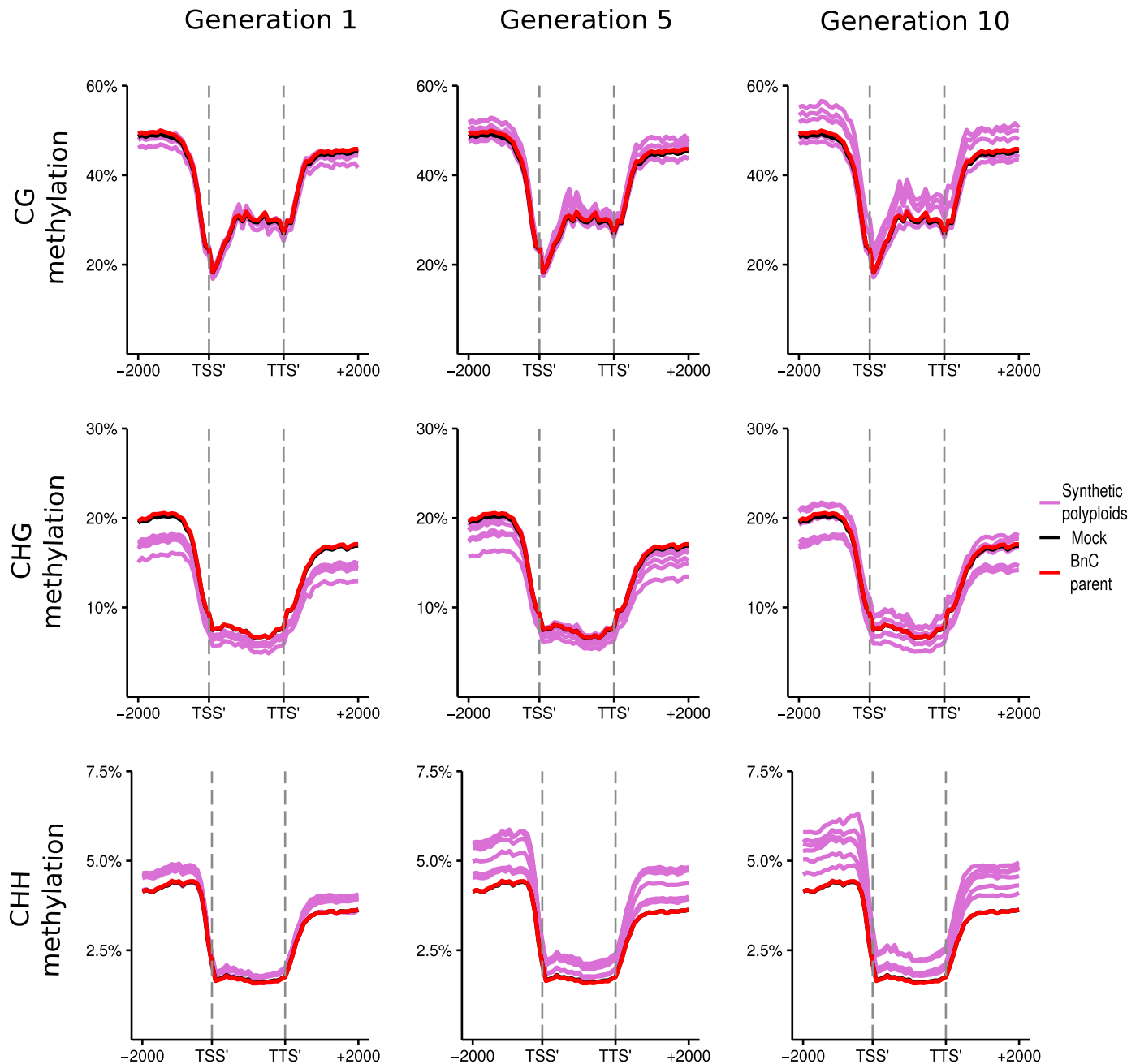




**Fig. 4** DNA methylation of long-terminal repeat (LTR) TEs in resynthesised *Brassica napus*. Metaplots of CG, CHG and CHH mean weighted methylation of all annotated LTR TEs for generations 1, 5 and 10 assessed using Bisulfite-seq. The six resynthesised lines are shown in purple, the *Brassica oleracea* progenitor is shown in red *Brassica rapa* progenitor in blue, and the *in silico* mock polyploid in black. Methylation levels are shown for the transcription start site (TSS'), gene body, transcription termination site (TTS') and 2 kb upstream and downstream of the TSS and TTS.

methylation levels across generations were also observed. From generations 1–10, individuals showed increased CG methylation in gene bodies and flanking regions for BnC genes, with some exceeding progenitor methylation levels (Fig. 5), but little increase was observed for BnA genes (Fig. 6). In genic regions on the BnC subgenome, CG methylation increase occurred primarily for nonsynthetic genes (Fig. S8), however in the flanking regions, CG methylation increases occurred for both synthetic and nonsynthetic genes (Figs S7–S10). For CHG sites, average

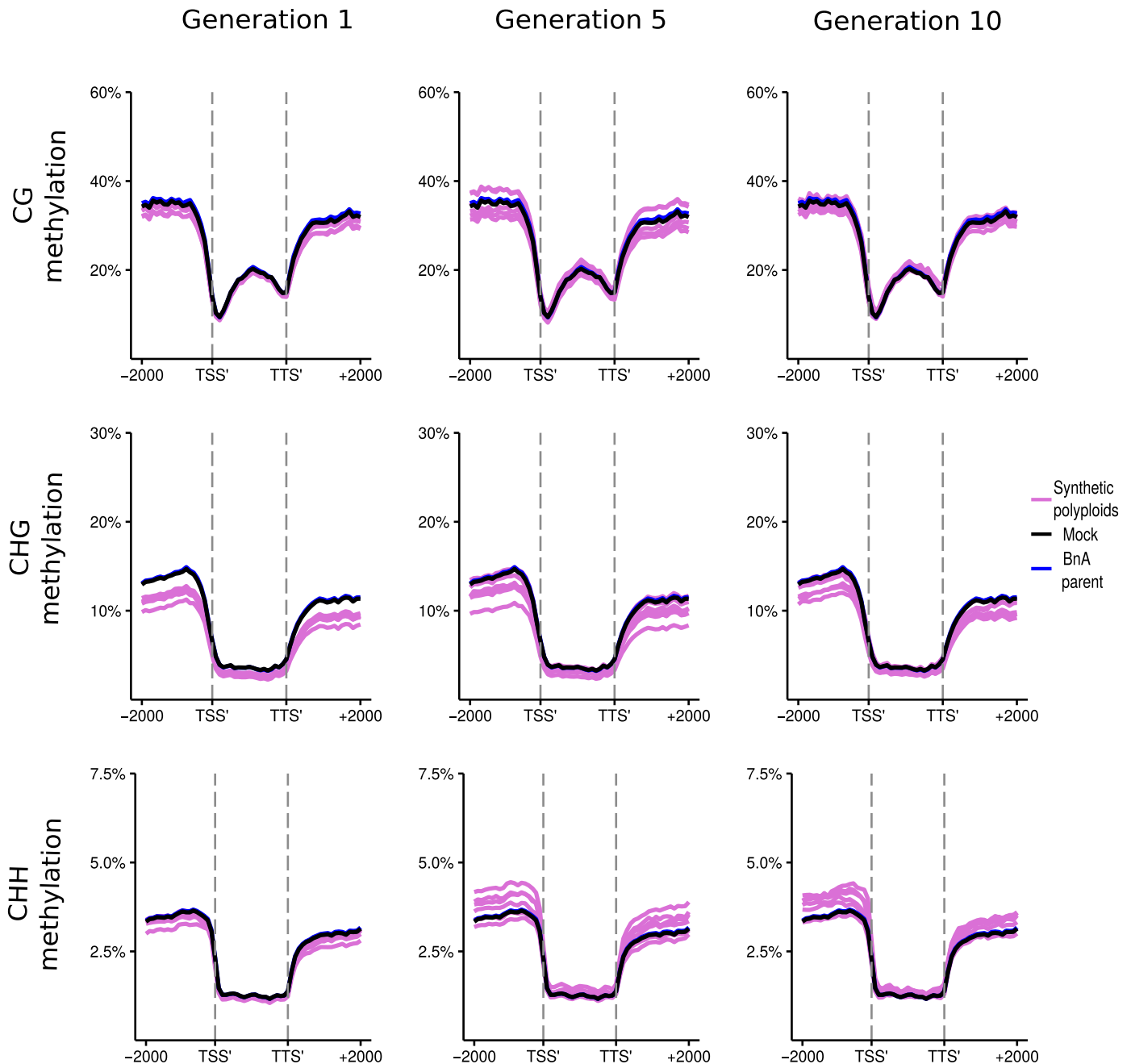
methylation levels in the 1<sup>st</sup> generation, samples were below progenitor levels, both in the bodies and in flanking regions of genes, and increased in subsequent generations with some lines equal to the progenitor level on the BnA subgenome and some exceeding the progenitor level on the BnC subgenome (Figs 5, 6). Syntenic genes showed both less variability and lack of increase across generations in CHG gene body methylation compared with nonsynthetic genes for both the BnA and BnC subgenomes, while patterns for flanking regions of genes were largely the same for



**Fig. 5** DNA methylation of BnC subgenome genes in resynthesised *Brassica napus*. Metaplots of CG, CHG and CHH mean weighted methylation of all gene models on the BnC subgenome for generations 1, 5 and 10 assessed using Bisulfite-seq. The six resynthesised lines are shown in purple, the *Brassica oleracea* progenitor is shown in red and *in silico* mock polyploid in black as a visual control for cross-mapping errors. Methylation levels are shown for the transcription start site (TSS'), gene body, transcription termination site (TTS') and 2 kb upstream and downstream of the TSS and TTS.

syntenic and nonsyntenic genes (Figs S7–S10). For LTRs, the same pattern of methylation increase over generations was observed, however LTRs on the BnC subgenome do not exceed progenitor methylation levels, while some lines showed BnA subgenome methylation exceeding the progenitor (Figs 7, 8). CHH methylation showed the most striking differences across generations. First generation allopolyploids showed flanking and gene body CHH methylation on the BnC subgenome that was slightly higher than the BnC progenitor methylation levels and

BnA subgenome methylation equal to or below the BnA progenitor (Figs 5, 6). In subsequent generations, CHH methylation in the flanking regions and gene bodies increased to surpass the progenitors on both BnA and BnC subgenomes, with the BnC subgenome showing the largest increase in methylation levels over time (Figs 5, 6). As with CG and CHG methylation, syntenic genes did not show an increase in gene body CHH methylation, while nonsyntenic genes did. CHH methylation of LTRs started around or below progenitor levels and increased to exceed

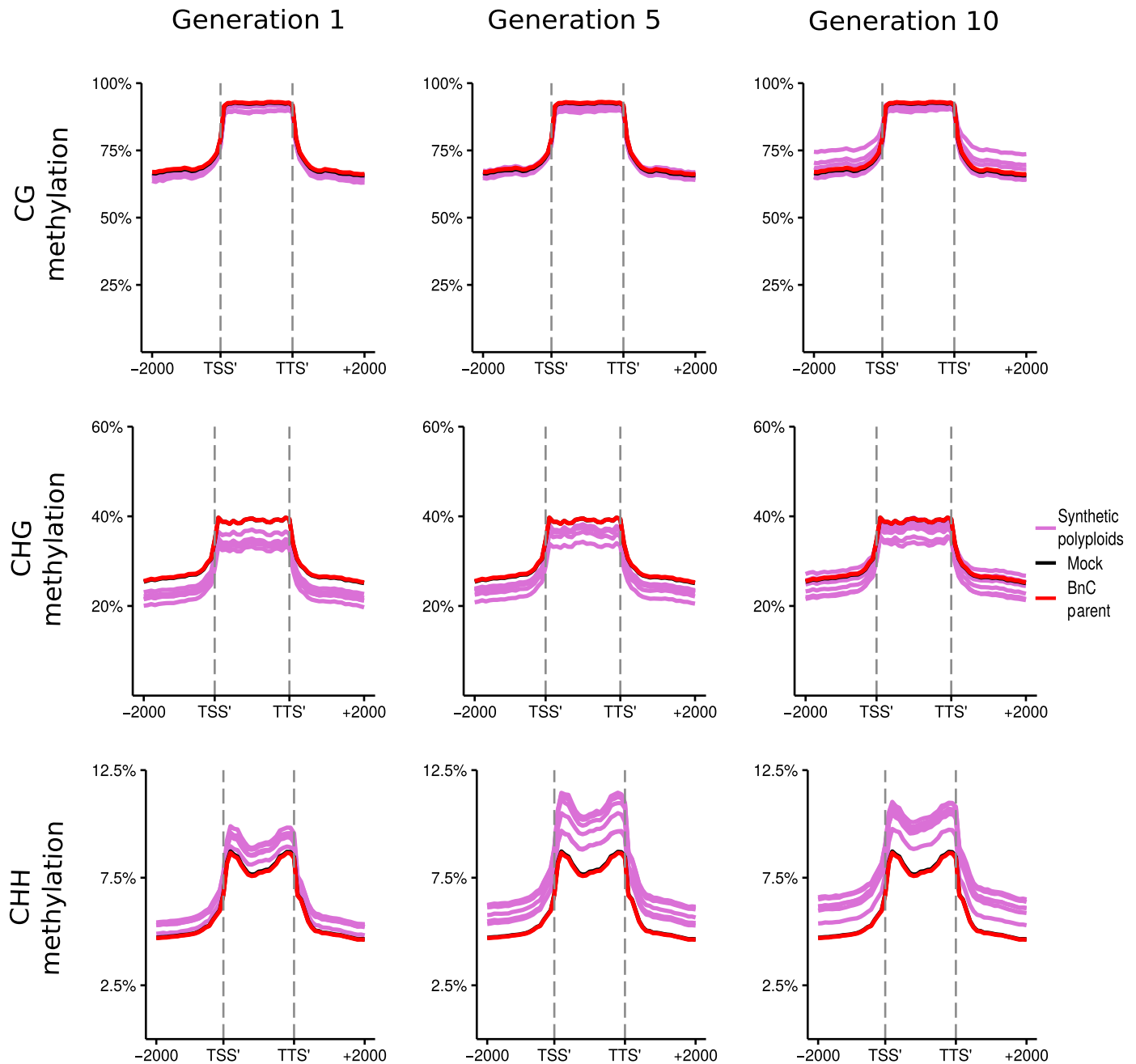


**Fig. 6** DNA methylation of BnA subgenome genes in resynthesised *Brassica napus*. Metaplots of CG, CHG and CHH mean weighted methylation of all gene models on the BnA subgenome for generations 1, 5 and 10 assessed using Bisulfite-seq. The six resynthesised lines are shown in purple, the *Brassica rapa* progenitor is shown in blue and *in silico* mock polyploid in black as a visual control for cross-mapping errors. Methylation levels are shown for the transcription start site (TSS'), gene body, transcription termination site (TTS') and 2 kb upstream and downstream of the TSS and TTS.

progenitor methylation levels in both BnA and BnC subgenomes by the 5<sup>th</sup> and 10<sup>th</sup> generations, although the increase was much larger for the BnC subgenome (Figs 7, 8).

To investigate the potential roles and patterns of DNA methylation related to homoeolog expression bias, we focused on homoeologs identified in 2:2 dosage with biased expression in the previous analyses. We compared homoeologs on both subgenomes with different homoeolog expression bias patterns to identify methylation patterns associated with

expression bias. For clarity, each subgenome is coloured in a consistent way as in Fig. 1 (BnA is blue, BnC is red). We observed that homoeolog pairs that had BnC-biased expression showed higher CG, CHG and CHH methylation in the flanking regions of genes (Fig. 9). Furthermore, BnC homoeologs shown to be more highly expressed than their BnA partner had lower CG methylation levels at the TSS, (Fig. 9, top right). While the reciprocal patterns were not observed for dominantly expressed BnA homoeologs at the



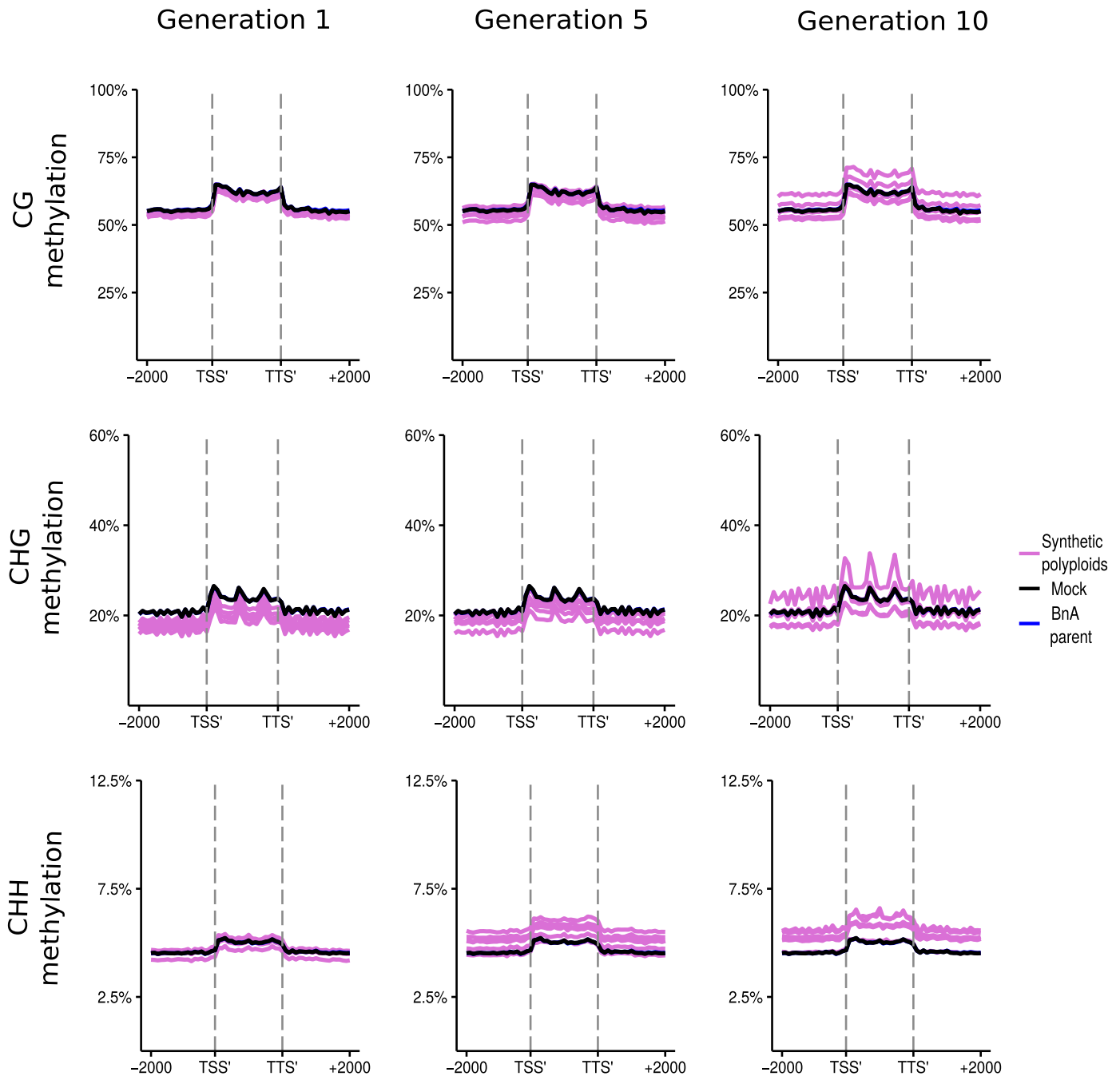
**Fig. 7** DNA methylation of BnC subgenome long-terminal repeat (LTR) TEs in resynthesised *Brassica napus*. Metaplots of CG, CHG and CHH mean weighted methylation of all LTR TEs on the BnC subgenome for generations 1, 5 and 10 assessed using Bisulfite-seq. The six resynthesised lines are shown in purple, the *Brassica oleracea* progenitor is shown in red and *in silico* mock polyloid in black as a visual control for cross-mapping errors. Methylation levels are shown for the transcription start site (TSS'), gene body, transcription termination site (TTS') and 2 kb upstream and downstream of the TSS and TTS.

TSS (Fig. 9, top left), differences in methylation patterns were observed in the flanking regions of these genes which were not observed for dominantly expressed BnC homoeologs. Homoeologs on the BnC subgenome more highly expressed than their BnA partner also showed lower CHH and CHG methylation levels in coding sequences. These patterns were not observed for the more highly expressed homoeologs on the BnA subgenome. No clear DNA methylation pattern differences were observed to explain higher expression for homoeologs on the BnA subgenome than their BnC partners.

## Discussion

Subgenome expression bias is consistent and mirrors expression differences of the diploid progenitor species

We observed a significant expression bias towards the maternal *Brassica oleracea* (BnC) subgenome in all six resynthesised lines. We saw *c.* 70% of biased homoeolog pairs showing the same bias relationship in all six lines and in the parents. This further bolsters the claim that subgenome expression dominance may be



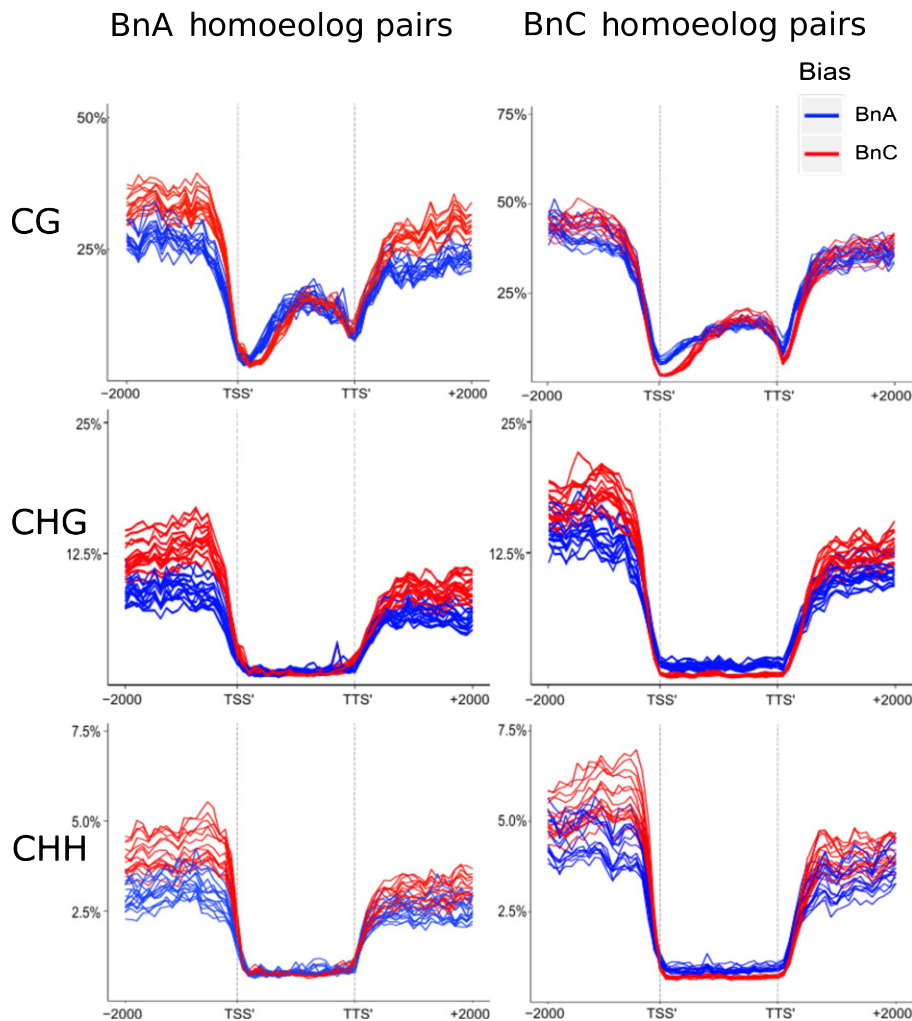
**Fig. 8** DNA methylation of BnA subgenome long-terminal repeat (LTR) transposable elements in resynthesised *Brassica napus*. Metaplots of CG, CHG and CHH mean weighted methylation of all LTR transposable elements on the BnA subgenome for generations 1, 5 and 10 assessed using Bisulfite-seq. The six resynthesised lines are shown in purple, the *Brassica rapa* progenitor is shown in blue and *in silico* mock polyploid in black as a visual control for cross-mapping errors. The data here show unique and unexpected oscillations for CHG methylation, which may be tracking nucleosomes, but are unlikely to be an artefact as CG and CHH methylation is unaffected. Methylation levels are shown for the transcription start site (TSS') and gene body, transcription termination site (TTS') and 2 kb upstream and downstream of the TSS and TTS.

predominately inherited from progenitors rather than an outcome of interspecific hybridisation and whole genome duplication (Buggs *et al.*, 2014). However, given that 7 of 16 individuals had a proportion of BnC-biased homoeologs that significantly deviated from parental expectations, and that 30% or more of BnC-biased homoeolog pairs were not shared by all resynthesised lines and/or the parental genomes, there appears to be some

variability in the extent to which homoeologous pairs are biased towards or away from the BnC subgenome.

The observed excess of BnC-biased pairs is also reflective of the pattern identified in natural *Brassica napus* cultivar Darmorbzh (Chalhoub *et al.*, 2014), but runs counter to observations of more BnA-biased pairs genome-wide and in genes related to cyto-nuclear interactions in other resynthesised *Brassica napus*





**Fig. 9** DNA Methylation of biased homoeologs in resynthesised *Brassica napus*. Average weighted methylation for homoeolog pairs in 2 : 2 dosage for all 16 lines assessed using Bisulfite-seq. BnC-biased expressed homoeologs in red and BnA-biased expressed homoeologs in blue. Results are shown for methylation in CG, CHG and CHH contexts. Methylation levels are shown for the transcription start site (TSS'), gene body, transcription termination site (TTS') and 2 kb upstream and downstream of the TSS and TTS.

lines (Wu *et al.*, 2018; Ferreira de Carvalho *et al.*, 2019). The differences between the results presented here are likely, in part, to be due to *B. rapa* (BnA) being used as the maternal parent in the other two studies, while *B. oleracea* (BnC) is the maternal parent in this study. Furthermore, differences in methodology may have affected the results obtained in these other studies. For example, we controlled for homoeologous exchanges that altered gene dosage by excluding genomic regions that were not in a balanced 2 : 2 ratio between both parents. Gene dosage is known to produce dosage-dependent expression changes (Conant *et al.*, 2014; Lloyd *et al.*, 2018). Furthermore, the differences between our study and these two previous studies (Wu *et al.*, 2018; Ferreira de Carvalho *et al.*, 2019) could, in part, be due to genetic differences in the accessions used to resynthesise *Brassica napus*. For example, previous work in autopolyploid *Arabidopsis thaliana* showed that parental genotype affected transcriptomic changes upon duplication (Yu *et al.*, 2010). The current model to explain subgenome expression dominance patterns is related to methylated TE density differences near genes in parental genomes (Bird *et al.*, 2018), and that gene and TE content are known to be highly variable within a species (Golicz *et al.*, 2016; Anderson *et al.*, 2019). Due to this intraspecific

variation, the progenitor genomes may differ in such a way as to yield variable, and even opposite subgenome dominance patterns in different resynthesised allopolyploid lines.

This proposed explanation may seem to contradict the metaphor of 'replaying the tape of life' or the general determinism of subgenome dominance, however our model of subgenome dominance is based around aspects of genotypes rather than assumptions about fixed interspecies genomic differences. If subgenome dominance was due to 'genomic shock' of interspecific hybridisation and genome doubling, we might expect individual genotypes to exhibit variable subgenome dominance outcomes (i.e. chance of either subgenome being dominant). However, if subgenome dominance is due to genotype-specific genomic features, we can reliably predict the subgenome dominance based on the parental genomes. The results presented here strongly suggested that an intermediate position holds true. Subgenome expression dominance in these resynthesised allopolyploids is strongly dependent upon progenitor genomic features, and crosses from the same genotypes seem to result in predictable and repeatable subgenome dominance patterns for a majority of genes in a given tissue type and environment. However, there is a sizeable proportion of genes that are variable

between lines, suggesting that dynamic processes are also occurring and affecting homoeolog expression.

GO and pathway enrichment analyses of the *Arabidopsis thaliana* protein–protein interaction network revealed that *Arabidopsis thaliana* homologs of BnC-biased homoeologs are highly enriched with primary metabolic and organellar functions, while homologs of BnA-biased homoeologs were not enriched with any known biological functions. Additionally, the protein–protein interaction networks constructed from BnC-biased homoeologs were more highly interconnected and clustered compared with BnA-biased homoeologs. However, there was still observed enrichment for interactions in the PPI network for BnA-biased gene pairs in generations 1 ( $P=9.75e^{-07}$ ) and 5 ( $P=0.00681$ ), but not generation 10 ( $P=0.16$ ). These results suggested that the hundreds of BnC-biased homoeologs are, to a greater extent, regulating vital cellular functions. The enrichment of organellar functions towards the BnC subgenome, which is the maternally contributed subgenome in our resynthesised lines, may implicate cyto-nuclear interactions as a driving force of subgenome dominance (Sharbrough *et al.*, 2017). One potential explanation is that genes on the maternal genome need to be expressed to interact with the plastid and mitochondria and paternal gene copies need to be silenced, and this is carried out by genomic imprinting. However, this is not definitive until similar analyses are repeated with reciprocal crosses. To further investigate the potential role of needing to maintain proper cyto-nuclear interactions, we compared our results with those of Ferreira de Carvalho *et al.* (2019) who analysed 110 nuclear-encoded components of plastid protein complexes and failed to find maternal progenitor homoeolog expression bias in their resynthesised *Brassica napus*. Of these 110 genes, we found 41 genes biased toward the BnC subgenome in all six lines in generation 1, 24 in generation 5 and 14 in generation 10. None of the 110 genes were biased towards the BnA subgenome in any individual across the six lines. However, because our analyses are genome-wide rather than just focused on the 110 gene subset, we were able to identify a BnC subgenome bias towards the organelles based on the thousands of known nuclear-encoded organellar genes (Savage *et al.*, 2013) as indicated by the 263 BnC-biased genes with GO cellular component annotation for the chloroplast in the 1<sup>st</sup> generation.

Homoeolog DNA methylation patterns are consistent, are correlated with observed expression biases and partially reflect differences in progenitors

We also identified consistent DNA methylation pattern differences between subgenomes. These differences appear to be a combination of inherited progenitor methylation patterns and methylation changes following interspecific hybridisation and polyploidisation. Globally, CG, CHG and CHH methylation of genes and LTR retrotransposons are higher in the dominant BnC subgenome than in the BnA subgenome, similar to results from natural *B. napus* (Chaloub *et al.*, 2014). These methylation patterns were also present when comparing diploid progenitor epigenomes. The observed methylation responses are possibly

due to ‘genomic shock’ following interspecific hybridisation, however without F1 hybrids we cannot rule out that it is a response to genome doubling rather than hybridisation. When we investigated the methylation level differences of biased expressed homoeologs, we saw that BnC homoeologs that were more highly expressed than their BnA partner showed markedly lower CG methylation at the TSS and lower CHG and CHH methylation across the coding sequence. These DNA methylation patterns were each associated with higher gene expression (Niederhuth *et al.*, 2016). However, it remains unclear whether these methylation pattern differences are the causal mechanism or are possibly just the result of expression differences of biased homoeologs.

The methylation patterns observed here appear to diverge from expectations based on the conceptual model outlined in Bird *et al.* (2018) and with observations in other polyploid systems that exhibit subgenome expression dominance. This model expects spill-over from methylation of TEs to affect neighbouring gene expression via RNA-directed DNA methylation pathways, predicting that subgenomes with lower TE density and lower methylation of TEs and/or upstream of genes will be dominantly expressed (Bird *et al.*, 2018). For example, in *Mimulus peregrinus* resynthesised and natural polyploids, the dominant subgenome had lower TE density and tended to have equal or lower expression than the nondominant subgenome (Edger *et al.*, 2017). Edger *et al.* (2017) highlighted that TEs on the dominant subgenome do not return to parental levels of CHH methylation, while they do on the nondominant subgenome and argued that this lower-than-parental methylation contributed to subgenome dominance. In maize, the dominant subgenome has consistently been observed to have lower methylation upstream of genes, in particular lower CHH methylation around 500 bp upstream of the TSS. (Renny-Byfield *et al.*, 2017; Zhao *et al.*, 2017). By contrast here, we found higher CHH methylation upstream for both the BnC subgenome on average and for BnC-biased homoeologs (Figs 5, 7, 9). There are several possible explanations for this. Regions of high CHH methylation, called ‘CHH islands’, have been found to be associated with highly expressed genes in maize (Gent *et al.*, 2013; Li *et al.*, 2015) and other species (Niederhuth *et al.*, 2016). It was proposed that this could serve as a boundary to reinforce TE silencing (Li *et al.*, 2015). Alternatively, it was recently shown in *Arabidopsis thaliana* that highly methylated CHH regions near genes recruited a protein complex that enhanced transcription and counteracted the deleterious effects of neighbouring TEs on genes (Harris *et al.*, 2018). It has yet to be demonstrated that this mechanism operates in other species, but it could explain paradoxical associations of high proximal CHH methylation with higher gene expression. Both of these cases provide alternatives to the current subgenome expression dominance model and potentially explain why higher expression and CHH methylation levels were observed on the dominant subgenome of our resynthesised lines.

Additionally, there are some patterns in common between our results and previous investigations of subgenome expression

dominance. In *Mimulus peregrinus*, there is markedly lower CG methylation at the TSS on the dominant subgenome, similar to what we observed with BnC homoeologs with higher expression than their BnA partner. Additionally, when Renny-Byfield *et al.* (2017) compared higher and lower expressed paralogues, regardless of subgenome of origin, they observed lower CG, CHG and CHH methylation at the TSS and across the coding sequence for the higher expressed paralogues. However, this lower expression was also seen upstream of the TSS, unlike our observations in resynthesised *B. napus*. These results may be due to particular genomic features of the progenitors.

For example, in soybean where there is no observable subgenome expression dominance, the paralogues from the two subgenome groups showed no differences in the distance to the nearest TE or in methylation levels (Zhao *et al.*, 2017). However, few genes in the soybean genome have a TE nearby (<1 kb), but more highly expressed soybean genes tended to be further away from TEs (Zhao *et al.*, 2017). These results suggested that multiple processes contributed to subgenome expression dominance, especially in genomes in which genes do not have TEs closely flanking upstream regions. This may also help to explain the observed spectrum of subgenome dominance in different species. For example, maize, *Mimulus peregrinus*, *Brassica rapa*, garden strawberry (*Fragaria x ananassa*), and others exhibited strong transcriptome-wide subgenome expression dominance, where genes on the dominant subgenome are, on average, significantly more highly expressed than the nondominant subgenome, and significantly more gene pairs that are biased toward the dominant subgenome. In addition, soybean exhibited neither transcriptome-wide bias, nor differences in how many gene pairs are biased to each subgenome. In our resynthesised *Brassica napus*, as well as natural *B. napus*, there is little if any transcriptome-wide bias, but significant bias in the number of homoeolog pairs that are biased toward the BnC subgenome. This may be a promising new avenue to further dissect the role of the various DNA methylation contexts as the underlying mechanism(s) of subgenome expression dominance.

While many of the methylation differences between subgenomes appear to be inherited from the diploid progenitor genomes, we also identified methylation patterns that appeared to have been repeatedly modified following interspecific hybridisation and/or polyploidy. CHH methylation of both BnA and BnC genes and LTR retrotransposons showed transgressive hypermethylation patterns, first matching parental methylation levels in the 1<sup>st</sup> generation, and then surpassing parental methylation levels in 5<sup>th</sup> and 10<sup>th</sup> generations. This increase in LTR methylation over parental and mid-parent value in later generations matched previous observations in resynthesised *B. napus* that LTR-derived 21 and 24 nucleotide (nt) siRNAs are transgressively expressed in later generations (Martinez Palacios *et al.*, 2019). Additionally, for the BnC subgenome, CG methylation also appeared to be transgressively hypermethylated in the later generations. The functional effect, if any, of hypermethylation at CG sites on driving subgenome expression dominance remains poorly understood and should also be the focus of future studies.

## Conclusions

By analysing genomic, transcriptomic and epigenomic datasets of six isogenic resynthesised polyploids, we were able to observe repeated patterns of subgenome expression dominance across multiple plays of the 'evolutionary tape'. Collectively, results from this study suggested that subgenome expression dominance in this *B. napus* system was largely, but not exclusively, based on pre-existing parental gene expression differences and related to nuclear genome interactions with organelles. Similarly, we observed subgenome methylation patterns consistent with pre-existing differences between parental epigenomes, although many cases of transgressive hypermethylation suggested that the epigenome was not as constrained by parental legacy. The observed DNA methylation differences between subgenomes were distinct from expectations of conceptual models and some previous findings. However, these patterns were consistent with patterns and mechanisms that could explain the observed expression bias, some not previously reported in cases of subgenome expression dominance to our knowledge. The causal role of DNA methylation patterns, particularly the pre-existing differences among the diploid progenitor genomes, in affecting subgenome expression differences remains poorly understood. Future progress in these areas will bring us closer to answering looming questions on the underlying mechanisms of subgenome dominance and the connections between subgenome expression and DNA methylation differences in polyploid genomes.








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## Author contributions

KAB, ZX, RV and PPE planned and designed the research. KAB, CH, SO, MG, JCP, ZX, RV and PPE performed data analysis, collection and/or interpretation. KAB, RV and PPE wrote the manuscript. KAB, CH, SO, MG, JCP, ZX, RV and PPE reviewed and edited the manuscript.

## ORCID

Kevin A. Bird  <https://orcid.org/0000-0002-3174-3646>  
 Patrick P. Edger  <https://orcid.org/0000-0001-6836-3041>  
 Malia Gehan  <https://orcid.org/0000-0002-3238-2627>  
 Chad E. Niederhuth  <https://orcid.org/0000-0003-3268-9079>  
 Shujun Ou  <https://orcid.org/0000-0001-5938-7180>  
 J. Chris Pires  <https://orcid.org/0000-0001-9682-2639>  
 Robert VanBuren  <https://orcid.org/0000-0003-2133-2760>



## Data availability

Raw data from this project are available on the NCBI Sequence Read Archive (SRA) Project PRJNA577908. Intermediate files and code to reproduce plots can be found at <https://doi.org/10.5061/dryad.h18931zjr>.

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## Supporting Information

Additional Supporting Information may be found online in the Supporting Information section at the end of the article.

**Figs S1–S5** Homoeolog bias in the parents and across generations of the five lines.

**Fig. S6** Common shared biased homoeolog pairs.

**Fig. S7** DNA methylation of BnC syntenic genes.

**Fig. S8** DNA methylation of BnA syntenic genes.



**Fig. S9** DNA methylation of BnC nonsyntenic genes.

**Fig. S10** DNA methylation of BnA nonsyntenic genes.

**Table S1** homoeolog expression bias chi-squared table.

**Table S2** homoeolog expression bias vs parent chi-squared table.

**Table S3** Enrichment analyses of BnC-biased gene pairs in the 1<sup>st</sup> generation.

**Table S4** Enrichment analyses of BnC-biased gene pairs in the 5<sup>th</sup> generation.

**Table S5** Enrichment analyses of BnC-biased gene pairs in the 10<sup>th</sup> generation.

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