DNA methylation-mediated expression of zinc finger protein 615 affects embryonic development in *Bombyx mori*

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**ABSTRACT**

Cell division and differentiation after egg fertilization are critical steps in the development of embryos from single cells to multicellular individuals and are regulated by DNA methylation via its effects on gene expression. However, the mechanisms by which DNA methylation regulates these processes in insects remain unclear. Here, we studied the impacts of DNA methylation on early embryonic development in *Bombyx mori*. Genome methylation and transcriptome analysis of early embryos showed that DNA methylation events mainly occurred in the 5’ region of protein metabolism-related genes. The transcription factor gene *zinc finger protein 615* (*ZnF615*) was methylated by DNA methyltransferase 1 (*Dnmt1*) to be up-regulated and bind to protein metabolism-related genes. *Dnmt1* RNA interference (RNAi) revealed that DNA methylation mainly regulated the expression of nonmethylated nutrient metabolism-related genes through *ZnF615*. The same sites in the *ZnF615* gene were methylated in ovaries and embryos. Knockout of *ZnF615* using CRISPR/Cas9 gene editing decreased the hatching rate and egg number to levels similar to that of *Dnmt1* knockout. Analysis of the *ZnF615* methylation rate revealed that the DNA methylation pattern in the parent ovary was maintained and doubled in the offspring embryo. Thus, *Dnmt1*-mediated intragenic DNA methylation of the transcription factor *ZnF615* enhances its expression to ensure ovarian and embryonic development.

**Keywords:** DNA methylation; Embryonic development; Transcriptional regulation; Epigenetic

**INTRODUCTION**

The zygote is the starting point of embryonic development. Embryogenesis is a complex process regulated by genetic and epigenetic mechanisms in which a zygote develops into a multicellular individual. DNA methylation via 5-methylcytosine (5mC), i.e., the covalent addition of a methyl (CH₃) group to the fifth carbon of the pyrimidine ring of cytosine residues of DNA molecules (Razin & Riggs, 1980; Yan et al., 2015), is an important epigenetic regulatory mechanism found in all
In this study, we analyzed the methylome and role of Dnmt1 in early embryos of *B. mori*, an economically valuable insect and model lepidopteran. We found that DNA methylation actively participated in embryogenesis by up-regulating the reproduction-specific transcription factor gene zinc finger protein 615 (ZnF615) to enhance the expression of nutrient metabolism-related genes. Furthermore, the regulatory pathway was consistent between parent ovaries and offspring embryos.

**MATERIALS AND METHODS**

**Insects and cell lines**

The *B. mori* strain P50 was obtained from the Research and Development Center of the Sericulture Research Institute of Guangdong Academy of Agricultural Sciences, Guangdong, China. The moths were reared on fresh mulberry leaves at 27 °C under a 12 h light/12 h dark cycle. The *B. mori* Bm12 (DZNU-Bm-12) cell line, originally derived from ovarian tissues (Khourad et al., 2009), was cultured at 28 °C in Grace’s medium (Invitrogen, USA) supplemented with 10% fetal bovine serum (FBS) (HyClone, USA).

**Dot blot assay**

Genomic DNA was extracted from 50–60 silkworm embryos at each stage of embryonic development (i.e., fertilized egg, blastoderm, germ-band, organogenesis, reversal period, and head pigmentation) and digested with RNase A (Promega, USA) to eliminate RNA contamination. Genomic DNA (500 ng) was denatured at 95 °C for 10 min and immediately cooled on ice. The resulting genomic DNA was spotted on a nitrocellulose blot polyvinylidene fluoride (PVDF) membrane (GE Healthcare, China) and dried, followed by UV crosslinking at 27 °C under a 12 h light/12 h dark cycle.

**Whole-genome bisulfite sequencing (WGBS)**

Genomic DNA was fragmented into 100–300 bp fragments by sonication, and a single “A” nucleotide was added to the 3’ end of the blunt fragments. The fragments with the adapter were then bisulfite-converted using a Methylation-Gold Kit (Zymo, USA). Finally, the converted DNA fragments were amplified using polymerase chain reaction (PCR) and sequenced using the Illumina HiSeq™ 2500 platform by Gene Denovo Biotechnology Co. (China).

The obtained clean reads were mapped to the *B. mori* reference genome (Wang et al., 2005; Xia et al., 2004) using BSMAP (v2.90) (Xi & Li, 2009). The methylation level was
calculated based on the methylated cytosine percentages in the whole genome, in each chromosome, or in different regions of the genome for each sequence context (CG, CHG, and CHH). To assess differences in methylation patterns in different genomic regions, the methylation profiles in the flanking 2 kb regions and gene bodies (or transposable elements) were plotted based on the average methylation levels for each window.

**RNA sequencing (RNA-seq) analysis**

Total RNA from *B. mori* embryos was extracted using a TRIzol Reagent Kit (Invitrogen, USA) according to the manufacturer’s protocols. RNA quality was assessed on an Agilent 2100 Bioanalyzer (Agilent Technologies, USA) and checked using electrophoresis on RNase-free agarose gel. RNA-seq was performed on the Illumina NovaSeq 6000 sequencing platform by Gene Denovo Biotechnology Co. (China). The raw reads were filtered to obtain clean reads, which were mapped to the *B. mori* reference genome (Wang et al., 2005; Xia et al., 2004) using HISAT2 v2.4 (Kim et al., 2015) with “RNA-strandness RF” and other parameters set to default. The assembled transcripts were generated after mapping, and their expression levels were normalized using FPKM values (fragments per kilobase of exon per million fragments mapped) by Cufflinks (Trapnell et al., 2012). Analysis of differentially expressed transcripts (DETs) between two groups was performed using DESeq2 software (Love et al., 2014). Transcripts with \( P < 0.05 \) and absolute fold-changes\( \geq 2 \) were considered DETs.

**Western blotting**

For western blot analysis, 40–100 μg of total protein extracted from embryos was separated using 10% sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE), followed by transfer to a nitrocellulose membrane (GE Healthcare, China). The membrane was blocked with 3% (w/v) BSA in TBST for 2 h at room temperature, followed by hybridization overnight at 4 °C in TBST containing 1% BSA and primary antibody (anti-Dnmt1 and anti ZnF 615). After washing in TBST three times, the membrane was then incubated with HRP-conjugated goat anti-rabbit/mouse IgG secondary antibody (1:2 000, Dingguo Biotechnology, China) for 2 h at 37 °C. Tubulin (Cat: TB002-R, Dingguo Biotechnology, China) was used as a reference to verify equal loading of the proteins on the gel.

**Electrophoretic mobility shift assay (EMSA)**

EMSA was conducted using a LightShift Chemiluminescent EMSA Kit (Thermo Scientific, USA). Methy- or biotin-conjugated oligonucleotide probes were heated at 95 °C for 10 min and then slowly cooled to room temperature. Binding assays were performed according to the manufacturer’s protocols. In brief, proteins were incubated for 20 min at room temperature with 20 μL of binding buffer (50 ng of poly(dI-dC), 2.5% glycerol, 0.05% NP-40, 50 mmol/L potassium chloride, 5 mmol/L magnesium chloride, 4 mmol/L EDTA, and 20 fmol of biotinylated end-labeled double-stranded probe). Different concentrations of unlabelled cold probes or unmethylated mutant probes were added to the binding mixture as competitors. After electrophoresis, the proteins were blotted onto positively charged nylon membranes (Hybond N+; Amersham Biosciences, UK), and the bands were visualized using the above EMSA kit according to the manufacturer’s protocols.

**CRISPR/Cas9 knockout of Dnmt1 and ZnF615**

Small guide RNA (sgRNA) sites on the exons of target genes were designed using CRISPRdesign (http://crispr.dbcis.jp/) (Naito et al., 2015) based on the GN19NGG motif. A pair of primers was synthesized to amplify the fragment containing the T7 promoter, target site, and gRNA sequences. The PCR product was purified and used as the template for transcription in vitro. The sgRNA was synthesized in vitro with the MEGAscript T7 Kit (Ambion, USA) following the manufacturer’s instructions and then purified with sodium acetate/ethanol. The quality of the purified product was analyzed by gel electrophoresis.

The fertilized eggs were collected within 2 h after oviposition and immediately placed on a clean glass slide for injection. The sgRNA and Cas9 proteins were mixed to a final concentration of 800 ng/μL and 600 ng/μL, respectively. The mixture was injected into the eggs using a micromanipulator (Eppendorf FemtoJet 4i and 4r, Germany). To prevent contamination, the eggs were disinfected by steaming in 10% formaldehyde for 5 min and then placed in an incubator at 27 °C under a 12 h light/12 h dark cycle. All experimental operations were completed within 2 h of oviposition.

**Identification of mutations at target sites**

To confirm target gene mutation, the epidermis was collected for genomic DNA extraction when the silkworm larvae molted into pupae. Primers including regions upstream and downstream of the target sites were used for PCR to detect knockout of the target sequence. The PCR products were purified and ligated into the pMD-18T vector for sequencing.

**Expression, purification, and antibody preparation of ZnF615 protein**

The ZnF615 open reading frame (ORF) (BGIBMGA007439) was cloned using cDNA from *B. mori* embryos with the primers shown in Supplementary Table S9. The cloned ZnF615 cDNA was inserted into the pGEX-6P-1 vector with a GST tag at the N-terminus to generate an expression vector, which was then used to transform *Escherichia coli* (BL21) cells for protein expression. The recombinant protein was purified using a GST Protein Purification Kit according to the manufacturer’s protocols (Beyotime, China) (Supplementary Figure S1). The purified recombinant GST-ZnF615 protein was mixed with Freund’s adjuvant (Dingguo Biotechnology, China) and injected into BALB/c mice obtained from the Guangdong Medical Laboratory Animal Center (China). Antisera were collected after three booster injections, each with 100 μg of the recombinant protein.

**Analysis of DNA affinity purification sequencing (DAP-seq)**

DAP-seq was carried out using recombinant ZnF615 proteins. Genomic DNA extracted from silkworm embryos was sonicated to obtain fragments (~200 bp). The fragments were purified, end repaired, and ligated with Y-shaped adaptors, as
were carried out. To evaluate knockdown efficiency, qRT-PCR was performed using the specific primers.

**Analysis of egg number and hatch rate**

To record egg number, ovaries were dissected from wild-type (WT), Dnmt1<sup>+/−</sup>, and ZnF615<sup>−/−</sup> B. mori moths daily from the first to eighth day of the pupal stage, and the eggs in the dissected ovaries were collected and counted. The larvae hatched from the eggs were recorded on days 6–10.5 after oviposition. The hatch rate was calculated as: hatch rate=number of larvae/number of eggs.

**Statistical analysis**

For all measurements, data are presented as the mean±standard error (SE). P-values for group comparisons were calculated using a paired Student’s t-test (*: P<0.05; **: P<0.01; ***: P<0.001).

**RESULTS**

**Dnmt1 knockout suppresses embryogenesis**

Previous research has shown that Dnmt1 knockdown by RNAi impacts embryonic development in B. mori (Xiang et al., 2013; Xu et al., 2021). To confirm this result, CRISPR/Cas9 gene editing was applied to knockout the Dnmt1 gene. First, sgRNAs targeting Dnmt1 exon 4 were injected into B. mori embryos within 2 h of oviposition. The knockout mutants were detected by aligning the edited target sequences to the WT genomic sequences. One mutant showed a deletion of 52 bases in Dnmt1 exon 4 (Figure 1A), resulting in a defective protein of only 203 amino acids without functional domains, significantly different from the complete Dnmt1 protein of 1 409 amino acids (Figure 1B). The hatching rate in the Dnmt1<sup>−/−</sup> group was only 38.09%, a decrease of 57.66% from WT (Figure 1C). Western blot analysis showed that both Dnmt1<sup>−/−</sup> and Dnmt1<sup>+/−</sup> unedited embryos lacked a complete Dnmt1 protein (Supplementary Figure S2). At day 8 post-oviposition, defects in the unedited Dnmt1<sup>−/−</sup> embryos were found, whereas the WT embryos had developed and started to hatch. The head, thorax, abdomen, gut, and leg segments of the mutant were underdeveloped, suggesting that early development was affected by Dnmt1 knockout (Figure 1D). Ovarian development was also altered in the Dnmt1<sup>−/−</sup> heterozygotes compared to the WT. The egg number generated by Dnmt1<sup>−/−</sup> decreased by approximately 20% compared to the WT (Supplementary Figure S3). These results indicate that DNA methylation catalyzed by Dnmt1 has a considerable influence on ovarian development and early embryogenesis.

**Identification of DNA-methylated genes required for embryogenesis**

To study how DNA methylation affects embryonic development, 5mC levels in embryos at different developmental stages were examined by qRT-PCR and dot blot analysis. Results showed that 5mC was detectable throughout embryonic development but was higher during the early embryonic stages (i.e., fertilized egg, blastoderm, and germ-band) than during the late embryonic stages (i.e., organogenesis, reversal period, and head pigmentation).
Illumina WGBS of genomic DNA from blastoderm-stage embryos, when the highest Dnmt1 mRNA level was detected (Figure 2A), showed a total of 439,751 5mCs in the whole genome, representing 0.67% of all cytosine sites in the B. mori genome (Figure 2B; Supplementary Table S1). CG methylation (70.92%) was higher than CHG (4.99%) and CHH (24.09%) methylation (H represents A, C, or T). CG, CHG, and CHH methylation occurred at 1.34%, 0.44%, and 0.50% of the genome-wide CG, CHG, and CHH sites, respectively. In the CG context, DNA methylation occurred predominantly in gene bodies, particularly in the exons (Figure 2B). Gene annotation showed that the methylated genes were mainly enriched in pathways related to protein metabolism, including ribosome, ribosome biogenesis in eukaryotes, proteasome, aminoacyl-tRNA biosynthesis, protein processing in endoplasmic reticulum (ER), ubiquitin-mediated proteolysis, and protein export (Supplementary Figure S4). In addition, four groups of genes were identified according to mean expression levels. The relationship between the expression levels of these genes and DNA methylation rates was analyzed (Figure 2C). Most methylation changes in the genes occurred around the TSSs. The gene expression levels were positively correlated with the methylation rates (Figure 2C), suggesting that the methylated protein metabolism-related genes were expressed at high levels, whereas unmethylated genes were expressed at low levels.

To further investigate the transcription products of the target genes of DNA methylation, RNA-seq was implemented after Dnmt1 RNAi. The fertilized zygotes were injected with dsDnmt1 and collected after 48 h to evaluate RNAi efficiency and conduct RNA-seq. The Dnmt1 mRNA level was significantly decreased by RNAi (Supplementary Figure S5). Transcriptome data showed that 24 up-regulated genes and 105 down-regulated genes were identified between the Dnmt1 RNAi treatment and control (Figure 2D; Supplementary Table S2). Differentially expressed genes (DEGs) were mainly enriched in nutrient metabolic pathways, including lipid, amino acid, and carbohydrate metabolism (Figure 2E). Among the 129 DEGs, 23 had a methylation rate of more than 0.5% (Figure 2D), which may be positively correlated with the mRNA level. Among these 23 DEGs, the mCpG sites of 20 genes were enriched around their TSSs (Supplementary Figure S6 and Table S3). To investigate whether Dnmt1 RNAi affects embryogenesis through these 23 genes, their expression patterns were compared with that of Dnmt1 from the fertilized egg to head pigmentation stages of embryonic development via the transcriptome (Xu et al., 2022a). Four genes (BGIBMGA007439, BGIBMGA005446, BGIBMGA012542, and BGIBMGA006364) were found to have identical expression patterns to Dnmt1 (Figure 2F; Supplementary Table S4), implying that they may be positively regulated by DNA methylation. Therefore, these four genes were selected as target genes of Dnmt1 for further study.
To determine the roles of these candidate genes in embryogenesis, dsRNA targeting BGIBMGA007439, BGIBMGA005446, BGIBMGA012542, and BGIBMGA006364 was injected into eggs on the first day of oviposition (Supplementary Figure S7). Knockdown of the BGIBMGA007439 gene, corresponding to the zinc finger protein ZnF615 containing six zinc finger motifs and a zinc finger associated domain (Figure 3A), significantly reduced the hatching rate (Figure 3B), whereas RNAi of the other three candidate genes did not have an obvious effect on hatching (Supplementary Figure S8). Furthermore, the mRNA (Figure 3C) and methylation levels (Figure 3D) of the promoter and gene body of ZnF615 were significantly decreased after Dnmt1 RNAi, indicating that ZnF615 is downstream of Dnmt1 and involved in embryogenesis.

DNA methylation in ZnF615 gene body enhances its expression

To investigate how 5mCG is involved in ZnF615 expression, 100 bp regions that showed the largest differences in mCG levels in the promoter (regions 1–4) or gene body (regions 5–8) of ZnF615 were selected and analyzed to determine their regulatory activities using a luciferase assay (Figure 4A; Supplementary Figure S9). The transcriptional activities of regions 5 and 7 in the gene body were significantly inhibited.
by Dnmt1 RNAi, whereas that of region 6 was enhanced. The transcriptional activities of regions 1 and 4 in the promoter were also regulated by Dnmt1 RNAi, but activity was extremely low (Figure 4B). Mutations in regions 5 and 7, in which CG at the mCpG sites was replaced by AT (Supplementary Figure S9), did not cause any change in transcriptional activity after Dnmt1 RNAi (Figure 4C). Thus, we assume that 5mC in the gene body rather than in the promoter of the ZnF615 gene is primarily responsible for regulation of gene expression. These results indicate that methylation in regions 5 and 7 in the gene body enhances ZnF615 gene expression, which offsets the inhibitory effect of region 6 during embryogenesis in silkworms.

To confirm whether Dnmt1 directly binds to regions 5 and 7 in the gene body, thereby catalyzing the formation of mCG, EMSA was conducted with the biotin-labeled region 5 or 7 probes and nuclear proteins isolated from silkworm embryos (Figure 5). Nuclear proteins bound to labeled region 5 or 7 probes could be competed off with 50× or 100× nonlabeled probes (cold probe) but not with mutated nonlabeled probes (Figure 5A, B). To confirm whether the binding protein is Dnmt1, a supershift assay was performed using an anti-Dnmt1 antibody (Xu et al., 2021). A supershifted band was detected when the anti-Dnmt1 antibody was present (Figure 5C), suggesting that the protein bound to the of region 5 or 7 probes was Dnmt1. Thus, the transcriptional activity assay and EMSA results demonstrated that DNA methylation catalyzed by Dnmt1 in gene body regions promoted ZnF615 expression.

Knockout of ZnF615 suppresses embryogenesis

To confirm the function of ZnF615 in embryogenesis, CRISPR/Cas9 gene editing was applied to knockout the ZnF615 gene. The qRT-PCR results showed that Dnmt1 and ZnF615 had similar expression patterns, with higher mRNA levels in embryos and gonads at the pupal stages in the WT silkworms (Figure 6A). A ZnF615 mutant, in which two bases in exon 4 were deleted (Figure 6B), formed a defective protein of 205 amino acids, lacking five of the zinc finger motifs in the complete protein of 565 amino acids (Figure 6C), and the full-length ZnF615 protein could barely be detected in ZnF615−/− (Figure 6D top). The hatching rate in the homozygous ZnF615−/− mutant was similar to that in the Dnmt1−/− mutant (Figure 6D bottom), and the embryos were unable to develop normally, with only 53.54% of the eggs hatching (Figure 6D bottom). Similar to Dnmt1−/− B. mori, both ZnF615−/− and ZnF615−/− unhatched embryos lacked the complete ZnF615 protein (Supplementary Figure S10).

Paraffin sections of ZnF615−/− mutant embryos at day 8 post-oviposition revealed that the unhatched ZnF615−/− embryos did not undergo complete embryogenesis and the organs were undifferentiated and unformed at the early stage (Figure 6E). Some of the hatched ZnF615−/− silkworms also died during the larval period, although about 30% survived to adulthood (Figure 6E). However, the ZnF615−/− embryos hatched and developed normally (Supplementary Figure S11). This phenotype was similar to that of the Dnmt1−/− mutant. In addition to the low hatching rate, egg number in the ZnF615−/− mutant decreased by approximately 20% compared to the WT
Identification of genes required for embryogenesis regulated by DNA methylation-mediated ZnF615

To determine which downstream target genes are regulated by ZnF615, DAP-seq analysis was carried out. Immunofluorescence assays showed that the ZnF615 protein was localized to the nucleus of Bm12 cells (Supplementary Figure S13), implying that it is a nuclear factor acting on its target genes. Therefore, we used DAP-seq to identify the target genes acted upon by ZnF615 in the silkworm genome (Figure 7A). By mapping the DAP-seq reads to the B. mori genome, ZnF615 binding sites and motifs were identified (Figure 7A). ZnF615 had binding sites on each scaffold, especially high peak number regions on Ps01–06 (Supplementary Figure S14). The ZnF615 binding sites were enriched in the regions near the TSS and TTS (Figure 7B). The most frequently appearing sequence motif was TTTTTTATTTGGTTT (Figure 7C), suggesting that this may be the specific motif for ZnF615 binding. A total of 1,209 ZnF615 candidate binding genes were annotated (Supplementary Table S6). To further investigate the genes regulated by DNA methylation-mediated ZnF615, we combined the ZnF615 peak-related genes from DAP-seq and DEGs identified by RNA-seq after Dnmt1 RNAi (Figure 7D). Nine DEGs after Dnmt1 RNAi were enriched in ZnF615 DAP-seq (Supplementary Table S7). The mRNA levels of these nine genes were determined by qRT-PCR after ZnF615 RNAi, with six showing significant reductions (Figure 7D), i.e., adhesive plaque matrix protein, synaptic vesicle glycoprotein 2B, uncharacterized protein LOC101738308, cuticular protein glycine-rich 19, obscurin, and AGAP005312-PA-like protein (Supplementary Figure S15). These findings suggest that the genes directly bound by ZnF615 and down-regulated by Dnmt1 RNAi are not nutrient metabolism-related genes. Therefore, in the ZnF615-knockdown embryos, we detected changes in the expression of nutrient metabolism-related DEGs enriched after Dnmt1 RNAi, including amino acid, carbohydrate, and lipid metabolism pathways (Figure 7E; Supplementary Table S8). The qRT-PCR results showed that nine of the 13 nutrient metabolism-related genes enriched by...
Dnmt1 RNAi were significantly decreased after ZnF615 RNAi (Figure 7E), including lipase, glucose dehydrogenase, and glucosidase metabolism-related genes. These results indicate that DNA methylation may enhance the expression of these genes via its effects on ZnF615.

As Dnmt1-mediated ZnF615 methylation affects the development of embryos and ovaries, we further investigated whether ZnF615 is methylated at the same sites and at the same level in the ovary and embryo. Analysis showed that the methylation regions in the ZnF615 gene were similar between the ovary and embryo, but the methylation level of the ZnF615 gene in the embryos was twice that in the ovaries (Figure 7F) (Xu et al., 2021). These observations suggest that DNA methylation of the ZnF615 gene is involved in the regulation of early embryogenesis in B. mori and that modification of ZnF615 methylation in the embryos likely originates in the ovaries and is strengthened by Dnmt1 activity during embryogenesis.

**DISCUSSION**

Properly regulated gene expression is essential for embryonic development. In B. mori, the fertilized zygote progresses through the blastoderm, germ-band, organogenesis, reversal period, and head pigmentation stages to develop into a mature embryo. Many genes related to DNA replication, transcription, protein synthesis, and epigenetic modifications are up-regulated before embryonic organogenesis, while genes related to hormone synthesis and signaling, neuromeres, and cuticle proteins are up-regulated at or after the organogenesis stage (Xu et al., 2022a). Thus, DNA methylation modifications may regulate early embryonic development by influencing the expression of genes necessary for cell division and differentiation. DNA methylation is associated with embryonic development in hemimetabolous and holometabolous insects (Bewick et al., 2019; Li et al., 2020; Ventós-Alfonso et al., 2020; Xiang et al., 2013). However, how DNA methylation affects insect embryonic development remains unclear. In this study, we confirmed the critical role of DNA methylation in B. mori embryogenesis by Dnmt1 knockout (Figure 1). We found that DNA methylation level was higher in early embryogenesis than in later embryogenesis (Figures 1, 2A). Based on genome-wide methylation analysis in early embryos, we also found that 5mC methylation mainly occurred in the gene bodies of highly expressed protein metabolism-related genes (Figure 2C; Supplementary Figure S4) required for early embryonic development. We determined that the nuclear protein ZnF615 gene was downstream of Dnmt1 (Figure 3). Finally, we revealed that the expression of nutrient metabolism-related genes was inhibited by Dnmt1 and ZnF615 RNAi (Figures 2E, 7D, E). Based on these findings, we hypothesize that downstream genes regulated by Dnmt1-induced ZnF615 methylation are vital for early B. mori embryonic development.

Research has reported that 5mC affects early embryonic development in insects. In Tribolium castaneum, DNA methylation occurs in the early stage of embryogenesis, followed by global demethylation as the embryo develops (Feliciello et al., 2013). Dnmt1 RNAi in B. germanica, which has both Dnmt1 and Dnmt3, results in defective embryos (Ventós-Alfonso et al., 2020). In this study, we demonstrated that knockout of Dnmt1, the only DNA methyltransferase in B. mori, stopped early embryonic development and even induced death (Figure 1), similar to that of Dnmt1 RNAi (Lyu et al., 2021; Xiang et al., 2013; Xu et al., 2021). However, nearly 40% of the homozygous knockout mutants hatched and grew...
normally. We speculate that the defective Dnmt1 protein in the knockout mutant was still partially active after mutation, or the organism was able to activate other mechanisms to compensate for the damage caused by the lack of DNA methylation. These results highlight the necessity of Dnmt1 expression and DNA methylation in early embryogenesis to ensure proper embryonic development in insects (Figure 1).

Importantly, in this study, we identified and characterized the nuclear protein gene ZnF615, which is modified by 5mC and acts downstream of Dnmt1. Here, ZnF615 knockout resulted in embryo and ovary phenotypes similar to those of Dnmt1 (Figure 6); moreover, 70% of the nutrient metabolism-related genes down-regulated by Dnmt1 RNAi were regulated by ZnF615 (Figure 7E; Supplementary Table S8), including lipid, amino acid, carbohydrate, and glycan metabolism-related pathways. Early embryonic development requires glycosidases to hydrolyze carbohydrates to produce glucose and provide energy; amino acids and fatty acids are also important for insect embryonic development (Levin et al., 2017). Knockdown of pancreatic lipase-related protein 2 in Nilaparvata lugens results in a decrease in the hatching rate (Xu et al., 2017). Thus, Dnmt1 RNAi may impact insect development by affecting these few genes. Dnmt1 RNAi inhibited ZnF615 expression, leading to the down-regulation of genes required for embryogenesis. Bombyx mori ZnF615 is a typical C2H2 zinc finger protein; C2H2 zinc fingers comprise the most common and diverse family of transcription factors (Vaquerizas et al., 2009; Weirauch & Hughes, 2011; Wu et al., 2019). However, as the defective ZnF615 protein in knockout mutants contained a zinc finger motif and zinc finger associated domain and may be partially active, more than 50% of mutant embryos hatched normally.

In insects, 5mC occurs in genes with conserved housekeeping functions (Bonasio et al., 2012; Lyko et al., 2010; Sarda et al., 2012; Simola et al., 2013). Our WGBS and RNA-seq data revealed a positive correlation between high methylation rates and high expression levels of numerous housekeeping genes (Figure 2C; Supplementary Figure S4). However, Dnmt1 RNAi did not down-regulate these housekeeping genes, as reported in other insects. For example, suppression of DNA methylation by Dnmt1 RNAi in Oncopeltus fasciatus (Bewick et al., 2019) or by 5mC inhibitor

Figure 6  Loss-of-function analysis of ZnF615 in B. mori embryos
A: Expression patterns of Dnmt1 and ZnF615 in different tissues, including epidermis, silk gland, testis, ovary, gut, wing disc, and fat body (top), and at different developmental stages, including embryo, 1<sup>st</sup>-5<sup>th</sup> instar larval, wandering larval, pupal, and adult stages (bottom). B: Schematic of nucleic acid base deletion site in ZnF615 (top). C: Functional domain analysis of ZnF615 protein in WT and ZnF615<sup>−/−</sup> mutant. D: Western blot analysis of ZnF615 protein in WT and ZnF615<sup>−/−</sup> mutant embryos (top). Hatching rates in WT and ZnF615<sup>−/−</sup> mutant (bottom). Each point represents an embryo hatching rate of female WT or ZnF615<sup>−/−</sup> silkworms. E: Morphology and structure of WT and ZnF615<sup>−/−</sup> mutant embryos. Significant differences were determined by t-test (*: P<0.05; **: P<0.01; ***: P<0.001).
Figure 7 DAP-seq analysis of ZnF615 in B. mori embryos
A: Overview of DAP-seq experimental process. cDNA of ZnF615 ORF fused to Halo affinity tag was expressed in vitro and recombinant protein was bound to ligand-coupled beads. Genomic DNA at blastoderm stage of B. mori embryo was ultra-sonicated to 200 bp fragments, which were ligated with Illumina-based sequencing adaptors. HaloTag-ZnF615 protein was then incubated with adapter-ligated genomic DNA library. After unbound DNA fragments were washed away, the ZnF615-bound fragments were released. Released DNA fragments were then purified and sequenced. B: Peak frequency in different gene body regions between 2 kb upstream and downstream. C: Top-ranked motif in ZnF615 DAP-seq data was TTTTTATTGT. TSS: transcription start site, TTS: transcription termination site. Motifs were determined by MEME analysis using top-ranked peaks. D: Venn diagram comparing DEGs identified by RNA-seq after Dnmt1 RNAi and ZnF615 binding genes by DAP-Seq at blastoderm stage of embryo (top). Change in mRNA levels of genes both peak related and differentially expressed after Dnmt1 RNAi in ZnF615 RNAi B. mori embryos (bottom). E: Changes in mRNA levels of DEGs related to nutrient metabolism pathways enriched by Dnmt1 RNAi in ZnF615 RNAi B. mori embryos. F: Analysis of mCG levels and regions of ZnF615 in B. mori ovary and blastoderm stage of embryo. Significance of results was determined by t-test (\( P<0.05; \) \( ** P<0.01 \)).
treatment in *B. mori* (Xu et al., 2021) only down-regulates a small number of genes but has a significant impact on insect development. We previously speculated that highly expressed housekeeping genes may be regulated by multiple pathways or mechanisms, including DNA methylation (Xu et al., 2021). Therefore, the transcriptional inhibition caused by the loss of gene body methylation may be easily compensated by other regulatory pathways. Second, the transient loss of methylation caused by Dnmt1 RNAi did not change the chromosomal structure of these genes, resulting in no change in gene expression.

In insects, 5mC usually occurs in the gene bodies of highly transcribed genes (Glastad et al., 2014; Wang et al., 2013; Xiang et al., 2010; Zemach et al., 2010) and does not inhibit transcription but leads to gene activation (Hellman & Chess, 2007; Meng et al., 2015; Wolf et al., 1984). This may occur because DNA methylation shapes chromatin and gene expression status by affecting the structure of nucleosomes and/or regulating other factors (Chodavarapu et al., 2010). In addition, methyl-CpG-binding proteins (MeCPs and MBDS) specifically recognize DNA methylation and change the local chromatin structure by recruiting histone-modifying enzymes or chromatin-remodeling complexes (Meng et al., 2015). Gene body methylation can also facilitate transcription elongation by inhibiting the activation of alternative promoters and controlling alternative splicing (Meng et al., 2015). In the *B. mori* ovaries, 5mC in the 5′ regions of the gene bodies recruits acetyltransferase through MB2D3 to enhance gene expression (Xu et al., 2021). Thus, consistent mechanisms of methylation targeting gene bodies (i.e., intragenic methylation) of protein metabolism-related genes may facilitate and promote embryogenesis. We demonstrated that Dnmt1 directly binds to the 5′ region of the ZnF615 gene body, thereby promoting its transcription efficiency by catalyzing 5mC formation (Figures 4, 5) and subsequent transcription of the ZnF615 gene.

We found similar intragenic methylation enrichment in genes associated with protein metabolism-related pathways in the *B. mori* ovary (Xu et al., 2021) and embryo (Supplementary Figure S4). Furthermore, 5mC-modified ZnF615 affected both ovarian and embryonic development by directly binding to protein metabolism-related genes and by indirectly regulating nutrient metabolism-related genes (Figures 2E, 7E; Supplementary Figure S4). The effect of 5mC-mediated ZnF615 in ovarian and embryonic development was achieved through DNA methylation at the same ZnF615 gene sites (Figure 7G). We revealed that, in insects, intragenic DNA methylation in the ovary can be inherited by the fertilized egg. In addition, the higher methylation levels of ZnF615 in the embryos than in the ovaries (Figure 7G) may explain why Dnmt1 expression is up-regulated at the beginning of embryogenesis (Lyu et al., 2021). The up-regulation of DNA methylation in ZnF615 enhanced its own expression and that of its downstream genes. Thus, the similar methylation patterns and levels in the ovary and embryo ensured the expression of embryogenesis-related genes.

In summary, this study demonstrated that intragenic DNA methylation enhances the expression of protein metabolism-related and nutrient metabolism-related genes through the DNA methyltransferase Dnmt1 and transcription factor ZnF615 to ensure ovarian and embryonic development, which may facilitate high fecundity in insects.

**DATA AVAILABILITY**

The raw Illumina sequencing data from WGBS of the *B. mori* early embryos were deposited in the NCBI Sequence Read Archive (SRA) under accession No. SRP343081 and GSA database under accession No. CRA006748. The raw Illumina sequencing data from RNA-seq after Dnmt1 RNAi in *B. mori* embryos were deposited in the NCBI SRA under accession No. SRP342894 and GSA database under accession No. CRA006752. The raw Illumina sequencing data from DAP-seq were deposited in the NCBI SRA under accession No. SRP342920 and GSA database under accession No. CRA006730.

**SUPPLEMENTARY DATA**

Supplementary data to this article can be found online.

**COMPETING INTERESTS**

The authors declare that they have no competing interests.

**AUTHORS’ CONTRIBUTIONS**

G.F.X. conducted most of the experiments, participated in data analyses, and drafted the manuscript. C.C.G., Y.L.T., T.Y.F., and Y.L.P. conducted Dnmt1 and ZnF615 knockout, egg number and hatch rate analysis, RNAi, gene cloning, and protein expression analysis. H.L., Y.G.L., and C.M.T. assisted with the experiments and insect rearing. Q.L.F. and Q.S.S. provided technical and material support, participated in discussions, and helped draft and revise the manuscript. S.C.Z. conceived the study design, supervised the study, and drafted and finalized the manuscript. All authors read and approved the final version of the manuscript.

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