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CRISPR/Cas9-targeted mutagenesis of SlCMT4 causes changes in plant architecture and reproductive organs in tomato

Xuhu Guo1,2,*, Jianguo Zhao3, Zhiwen Chen3, Jun Qiao3, Yongfang Zhang1, Hong Shen1 and Zongli Hu4,*

1School of Life Sciences, Shanxi Datong University, Datong 037009, China
2Protected Agricultural Technology Research and Development Center, Shanxi Datong University, Datong 037009, China
3Institute of Carbon Materials Science, Shanxi Datong University, Datong 037009, China
4Laboratory of Molecular Biology of Tomato, Bioengineering College, Chongqing University, Chongqing 400044, China
*Corresponding author. E-mail: xhguo201010@126.com, huzongli@cqu.edu.cn

Abstract

DNA methylation participates widely in the regulation of gene expression in plants. To date, the regulation and function of DNA methylation is still unknown in tomato plants. Here, we generated SlCMT4 mutants using the CRISPR-Cas9 gene editing system. We observed severe developmental defects in CRISPR-Cas9-mediated SlCMT4 mutants, including small and thick leaves, increased lateral buds, defective stamens and pistils, small fruit size with reduced setting rate, and defective seed development. The alterations at hormonal levels (IAA, tZR, strigol) were consistent with the multibranching phenotype in SlCMT4 mutant plants. CRISPR-Cas9-mediated knockout of SlCMT4 induced the expression of two pollen-specific genes (PMEI and PRALF) that suppressed the development of pollen wall and pollen tube elongation, which is responsible for irregular and defective pollen. The small-sized fruit phenotype is probably associated with upregulated expression of the IMA gene and reduced seeds in the mutant lines. Furthermore, we performed whole-genome bisulfite sequencing (WGBS) of fruits and found that SlCMT4 knockout reduced genome-wide cytosine methylation. A reduction of methylation was also observed in a 2-kb region of the IMA promoter in the SlCMT4 mutant fruits, indicating that the hypermethylation status of the CHH context is critical for the inhibition of IMA and LOXB promoter activity. Our results show that SlCMT4 is required for normal development of tomato vegetative and reproductive organs. This study illuminates the function of SlCMT4 and sheds light on the molecular regulatory mechanism of tomato plant architecture and fruit development and ripening.

Introduction

Epigenetic regulation is an important part of gene mRNA accumulation and has become a hot spot in current research. DNA methylation plays a critical role in gene imprinting, genome stability, development, and response to the environment [1–3]. DNA methylation is mainly catalyzed by DNA methyltransferases (MTases), including methyltransferase 1 (MET1), chromomethylases (CMTs), domains rearranged methyltransferases (DRMs), and DNA methyltransferase 2 (DNMT2) in plants [4]. MET1 encodes the DNA methyltransferase that is associated with maintaining CG methylation. Pleiotropic developmental abnormalities are observed in antisense-MET1 lines of Arabidopsis with decreased methylation level, such as smaller plant size, altered leaf size and shape, reduced apical dominance, altered flowering time, and decreased fertility [5, 6]. Arabidopsis plants with reduced endogenous expression of the MET1 gene display severe phenotypes [7–9], whereas homozygously targeted rice plants shows no apparent phenotypic alterations [10]. Arabidopsis thaliana embryos exhibit reduced viability and altered planes and numbers of cell division due to dysfunction in MET1 and CMT3 [8]. Hypomethylation of DNA in CG sequences is detected in tobacco transgenic lines by expressing an antisense NtMET1 construct [11]. NtMET1 transgenic plants with significantly decreased methylation levels of genomic DNA show severe phenotypic defects, including short internodes, small leaves, and abnormal flower morphology [11]. Distinct developmental defects are observed in slmet1 mutants generated using CRISPR-Cas9 gene editing, including defective inflorescence, small and curly leaves, and parthenocarpy [12].

CHG and CHH (H = A/C/T) methylations are mediated by CMTs and DRMs [2]. A study by Moritoh et al. [13] revealed that targeted disruption of rice OsDRM2 by homologous recombination-mediated gene targeting led to a series of developmental phenotypes in different
growth and development stages, such as defective growth, semi-dwarf stature, reduced tiller number, delayed heading or no heading, abnormal panicle and spikelet morphology, and complete sterility [13], whereas no developmental defects were observed in the drm2 mutants of Arabidopsis. dmt103, loss-of-function lines in maize (Zea mays), shows severe developmental defects in seed morphologies at the reproductive stage, but no morphological alterations at the vegetative developmental stage [14]. In addition, plants with T-DNA insertion of OsTSD2, a putative rice methyltransferase gene, display dwarfishness and serial alterations in the root zones [15]. Bartee et al. [16] reported that Arabidopsis cmt3 mutants in the pai1C251Y background displayed developmental alterations. These alterations include strongly reduced fluorescence at the seedling stage and partially reduced fluorescence at the adult stage, with increased size, decreased bushiness, and increased fertility. Cao and Jacobsen [17] revealed that cmt3 single mutants and drm1, drm2 double mutants all showed no morphological defects, but drm1, drm2, cmt3 triple mutant lines showed pleiotropic alterations in plant development. Recently, Chen et al. [18] showed that suppressed expression of the CMT3 gene through virus-induced silencing (VIGS) in the tomato mutant induced expression of the CNR gene and triggered fruit ripening in the epimutant. Some similar effects on the ripening process in Cnr tomato mutants are also found through VIGS of the SIMET1, SIDRM7, and SICMT2 genes. In addition, Fieldes et al. [19] showed that the level of methylation decreased and flowering time was earlier after treating flax with 5-azaC (5-azacytidine, a methylation inhibitor).

In recent years, the importance of DNA methyltransferases in plant development, transcriptional regulation, and control of metabolic pathways has been increasingly recognized. At present, there are many research reports on genes related to fruit ripening, but there are few research reports that clarify the regulatory mechanism of fruit ripening from the aspect of epigenetics, especially DNA methylation. The epigenetic regulation mechanism of tomato fruit development and ripening remains unclear. Preliminary studies showed that expression of the tomato DNA methyltransferase gene SICMT4 was high in floral organs and immature green fruit, but declined during leaf development and fruit ripening [20]. To investigate DNA methylation in tomato (Solanum lycopersicum 'Ailsa Craig' AC++) , we applied CRISPR-Cas9-mediated gene knockout to generate targeted disruptants of SICMT4, which encode DNA methyltransferases responsible for de novo and non-CG methylation as annotated in Arabidopsis. Furthermore, we analyzed the SICMT4 mutant lines at morphological, physiological, molecular, transcriptome, and methylome levels. This study enhances our understanding about the role of SICMT4 in diverse developmental processes and thus has important theoretical significance.

Results

Efficient mutation of the SICMT4 gene via the CRISPR-cas9 system

SICMT4 (Solyc08g005400.2) was mutated in cv. 'Ailsa Craig' using the CRISPR/Cas9 system to confirm its function. The two targeted sites were located on the first exon of the SICMT4 gene (Fig. 1). In total, six kanamycin-resistant lines were obtained from the T0 transgenic lines. Among them, four lines (CR-01, CR-02, CR-08, and CR-09) were found to have mutations at the first target sites (CR-01, 3-bp insertions; CR-02, 1-bp deletion; CR-08, 1-bp deletion; CR-09, 5-bp deletion) and the second target sites (CR-01, 3-bp deletions; CR-02, 3-bp deletions and 1-bp SNP; CR-08, 1-bp SNP; CR-09, 3-bp deletion and 5-bp SNP) (Fig. 1). We also obtained four types of SICMT4 homozygous mutants from the self-pollinated T0 lines. Through CRISPR-cas9 gene editing and sequencing technology, we obtained four mutant lines, CR-01, CR-02, CR-08, and CR-09 (Fig. 1), among which CR-08 and CR-09 had more obvious phenotypic changes; thus we focused on these two lines as the main studied objects.

CRISPR-Cas9-mediated knockout of SICMT4 promotes branching

SICMT4 mutant lines CR-08 and CR-09 exhibited increased lateral buds at the seedling stage (Fig. 2a). Statistical analysis showed that the number of lateral buds in mutant CR-08 and CR-09 lines were significantly

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Figure 1. SICMT4 gene knockout mediated by the CRISPR-Cas9 system. Schematic map of the sgRNA targeted sites on the genomic regions of SICMT4, and sequencing results of the SICMT4 homozygous mutant lines from the T1 generation. The edited exon is shown as lines; the adjacent intron is shown as a blue box. Other exons and introns are shown as dashed lines. PAMs (protospacer adjacent motifs; NGG) are shown in red boxes, black arrows represent insertions, and dashes indicate deletions.
higher than in wild type (WT) at the mature stage (3 months old) (Fig. 2b). This change in the number of lateral buds was also observed in SlCMT4 RNAi lines (Supplementary Fig. S1). By comparing lateral buds grown for 5 days (Fig. 2c), we found that the length and diameter of the lateral buds in the SlCMT4 mutant line CR-08 were obviously larger than those of the WT (Fig. 2d and e). Anatomical data revealed that the parenchyma cells in pith of the lateral buds from the SlCMT4 gene mutant line became larger in transverse and longitudinal sections (Fig. 3), which was similar to the size alteration of cortical cells in the mutant lines (Supplementary Fig. S2). Given the increased lateral branches phenotype of SlCMT4 mutant plants, we further studied the effects of SlCMT4 knockout on lateral branches at hormone levels. The results revealed the content of Indole-3-acetic acid (IAA) and trans-Zeatin-riboside (tZR) in the lateral buds of the SlCMT4 mutant lines increased (Fig. 4a and b); tZR content was increased ~4-fold in lateral branches of the mutant line CR-08. Consistent with the reduced content of strigol in SlCMT4 RNAi lines (Supplementary Fig. S3), the content of strigol decreased in the CR-08 line (Fig. 4c).

**Knockout of SlCMT4 changes leaf morphogenesis**

The SlCMT4 mutant lines displayed compact architecture (Fig. 5a) and small internodes (Fig. 5b). The mutant lines also exhibited small leaves (Fig. 5c), which was manifested in the significantly smaller leaf length, width, perimeter, and area (Fig. 5d-g). RNAi suppression of SlCMT4 also produced a similar leaf phenotype (Supplementary Fig. S4). Longitudinal section analysis of WT and mutant line leaves (Fig. 5h) further revealed that the thickness, palisade, and sponge tissue thickness of the mutant lines increased significantly, and the gap between the palisade and sponge tissue was larger (Fig. 5k-m). These observations illustrated that leaf structure differed at the microscopic level between WT and mutant lines.

**CRISPR-Cas9-mediated knockout of SlCMT4 affects floral organ morphology and pollen development**

The floral organs of CR-08 lines were obviously defective and everted at anthesis (Fig. 6a). Scanning electron microscopy revealed that the stamen cells (Fig. 6c) and pollen grains (Fig. 7) were irregular and defective in the mutant lines. We also studied the expression of tomato pollen-specific genes in CR-08 by quantitative real-time PCR (qRT-PCR) analysis, and found that PMEI and PRALF were markedly upregulated in the pollen of the mutant line CR-08 (Fig. 8). In addition, the mutant line CR-08 showed short and thick pistils compared with WT (Supplementary Fig. S5).

**CRISPR-Cas9-mediated knockout of SlCMT4 leads to reduced fruit size**

The SlCMT4 mutant lines displayed small fruit compared with WT (Fig. 9a). The data showed that the horizontal
Figure 3. Anatomical analysis of lateral buds from WT (a, d) and CR-08 (b, e) lines. a and b Parenchyma cells in pith from transverse section of lateral buds. c Parenchyma cell area was estimated in transverse sections of lateral buds. d and e Parenchyma cells in pith from longitudinal sections of lateral buds. f Parenchyma cell area was estimated in longitudinal sections of lateral buds. Scale bars = 100 μm. The samples were prepared from 5-day-old lateral buds.

Figure 4. Effect of SlCMT4 knockout on lateral buds at hormone levels. a–c Contents of IAA, tZR and strigol, respectively, in lateral buds of WT and mutant lines. Each value represents the mean ± standard deviation of three replicates. Asterisks indicate statistically significant differences relative to WT, determined using t-tests. ∗P < .05, ∗∗P < .01.

diameter, vertical diameter, and weight of the fruit at different maturity periods (immature green fruit (IMG), mature green fruit (MG), breaker (B), 3 days after breaker (B3)) was decreased in the SlCMT4 mutant lines (Fig. 9b–d). A similar result was observed in the SlCMT4-RNAi plants (Supplementary Fig. S6). Transverse anatomical analysis of fruit at IMG stage showed reduced seeds in the CR-08 lines compared with WT (Fig. 10a). The expression level of the fruit size-related gene IMA was upregulated in fruits at IMG stage of the CR-08 line based on RNA-seq (Fig. 10b). qRT–PCR analysis validated that the fruit size-related gene IMA was upregulated by ~2.3-fold in fruit at the IMG developmental stage of the mutant lines (Fig. 10c). Besides, seeds from IMG fruit in the CR-08 line were smaller compared with WT (Fig. 10d and e).

Differentially expressed genes and enrichment analysis
We collected pericarp tissue samples at breaker stage from WT and CR-08 lines to perform a comparative transcriptome analysis. The RNA-seq data were reliable on the basis of Pearson correlation coefficients (> .90) of the three biological replicates (Supplementary Fig. S7A). Two different levels of gene expression were recorded in WT and CR-08 from principal component analysis of six pericarp samples (Supplementary Fig. S7B). We specifically paid close attention to genes that were differentially expressed (DEGs) in the pericarp samples in the CR-08 line. Differential expression of genes was defined using a 1.5-fold change and a P-value of < .05 as the cutoffs (Supplementary Fig. S7C). We identified 4102 DEGs, among which 2524 were upregulated and
Figure 5. Leaf morphologies of WT and mutant lines. a) Gross morphologies of WT, CR-08, and CR-09 lines, from left to right. b) Internode length of 3-month-old WT and mutant lines. c) Leaf morphologies of WT and mutant lines. d–g) Length, width, perimeter, and area of leaves, respectively, from WT and mutant lines. h–j) Longitudinal sections of leaves from WT and mutant lines (CR-08, CR-09). Scale bars = 50 μm. k–m) Thickness of leaves, palisade tissue, and sponge tissue, respectively, of WT and mutant lines. Each value is the mean ± standard deviation of three replicates. The asterisk indicates a significant difference (P < 0.05) between WT and mutant lines.

1578 were downregulated (Supplementary Fig. S7D). Supplementary Fig. S8 lists the results of Gene Ontology (GO) category enrichment analysis using GOseq [21] for DEGs. GO terms associated with critical cellular components were enriched in pericarp of the CR-08 line, including organelle part, membrane-enclosed lumen, macromolecular complex, cell junction, extracellular region, and symplast. Molecular functions that were enriched included structural molecular activity, nucleic acid-binding transcription factor activity, transporter activity, molecular transducer activity, and nutrient reservoir activity. Supplementary Fig. S9 shows that most DEGs were categorized as belonging to functional pathways responsible for the following: (1) metabolism, including carbon metabolism, biosynthesis of amino acids, phenylpropanoid biosynthesis, amino sugar and nucleotide sugar metabolism, and starch and sucrose metabolism; (2) genetic information processing, such as protein processing in the endoplasmic reticulum; and (3) environmental information processing, such as plant hormone signal transduction.

Transcriptional analysis of fruit ripening-related genes in SlCMT4 mutants

Using WT and CR-08 line fruits (B stage) as materials, transcriptome sequencing technology was used to study the transcriptional levels of tomato fruit ripening-related genes (such as PE1, PG2, LOXB, and ACO1). The results revealed that all ripening-related genes were upregulated in the CR-08 line based on RNA-seq (Fig. 11a–d). To validate the results of RNA-seq, we performed qRT-PCR analysis. The results showed that PE1 and PG2 genes, related to fruit wall metabolism, were respectively upregulated by ~5- and 6-fold in the fruits of mutant line CR-08. In addition, the lycopene synthesis-related gene LOXB was significantly upregulated in B-stage fruits of the mutant
line. The ACO1 gene, related to ethylene synthesis, was also upregulated in fruits of the mutant line during B stage (Fig. 11e–h).

Effects of SlCMT4 knockout on seed and fruit set rate
The SlCMT4 mutant lines exhibit a low fruit set rate (Fig. 12). The seed number per fruit (Fig. 13c), weight per 50 seeds (Fig. 13d), and germination rate (Fig. 13e) in the mutant lines decreased; in particular, the seed number per fruit of CR-08 line decreased by ~70%, and some seeds failed to develop normally (Fig. 13a and b). Additionally, the middle of the seeds was sunken (Fig. 13i), and the number of epidermal hair of seeds from the CR-08 line was increased, as confirmed by scanning electron microscopy (Fig. 13j and k).

SlCMT4 dysfunction reduces genome-wide cytosine methylation
The single-base resolution methylome of SlCMT4 mutant fruit was investigated by whole-genome bisulfite sequencing (WGBS). The genome-wide CHH methylation level was markedly reduced to 5.82% in SlCMT4 mutant lines.
compared with WT plants (Fig. 14a). CHH hypomethylation was observed across each of the 12 chromosomes in SlCMT4 mutants (Supplementary Fig. S10). The strong reduction in global CHH methylation level is consistent with the role of CMT4 being responsible for CHG and CHH methylation in plants [2]. In addition to genome-wide CHH hypomethylation, SlCMT4 mutants also exhibited substantially decreased DNA methylation levels in the CHG context (Fig. 14a, Supplementary Fig. S10). On the whole, SlCMT4 dysfunction displayed a stronger impact on genome-wide DNA methylation compared with WT; gene regions and transposable element (TE) regions both showed CHH hypomethylation on a whole-genome scale in the SlCMT4 mutant (Supplementary Fig. S11). Thus, disruption of SlCMT4 function extensively changed the tomato DNA methylome due to integrated regulation of DNA methylation in different cytosine contexts.

Since SlCMT4 loss of function differentially influences DNA methylation in CG, CHG, and CHH cytosine contexts, we investigated differentially methylated regions (DMRs) of CG, CHG, and CHH separately throughout the genome using DMRcaller [22]. In the CHH context, SlCMT4
mutants show many more hypo DMRs than hyper DMRs (Fig. 14b). These results are consistent with the assessment of global DNA methylation levels shown in Fig. 14a and Supplementary Figs S10 and S11. That CHH DMRs in SlCMT4 mutants are overwhelmingly dominated by hypomethylation supports the role of SlCMT4 as a CHH
methylnase. We further categorized DMRs into eight types, i.e. CGI, gene, DOWN3K, intron, intergenic, UP3K, repeat, and exon regions. Among the SlCMT4 mutant CHH hyper DMRs, UP3K and DOWN3K region accounted for 43.46 and 28.13%, respectively (Fig. 14c). Meanwhile, in the SlCMT4 mutant CHH hypo DMRs, repeat and CGI regions accounted for 74.18 and 7.82%, respectively (Fig. 14d).

CRISPR-Cas9-mediated knockout of SlCMT4 reduces cytosine methylation in the promoter region of candidate genes

We further examined the DNA methylation levels of candidate gene (IMA and LOXB) target promoters (2 kb) in tomato WT and SlCMT4 mutant lines. A marked reduction of methylation was observed in a 2-kb region of the

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**Figure 13.** Characteristics of seeds in WT and mutant lines. **a** Transverse sections of fruits at different stages from WT and mutant lines. **b** Digital photographs of seeds of single fruits. **c** Seed number per fruit. **d** Seed weight per 50 seeds at B3 stage. **e** Seed germination rate. **f** and **i** Close-up photographs of seeds; **g**, **h**, **j**, and **k** are scanning electron microscope images (B stage). Data represent the mean ± standard deviation of three replicates. Asterisks indicate statistically significant differences relative to WT, determined using t-tests. *P < .05, **P < .01.
IMA and LOXB promoters in the SlCMT4-mutant fruits (Fig. 15).

Discussion
DNA methylation plays an important function in manipulating many developmental pathways in plants. In this study, we constructed a CRISPR-Cas9 vector of the SlCMT4 gene, transformed WT tomato, and screened mutants of the SlCMT4 gene. Based on the phenotypes of mutant lines, some possible functions of the SlCMT4 gene were characterized at the morphological, physiological, biochemical, anatomical, and molecular levels. The SlCMT4 mutants showed severe developmental defects with a deletion mutation, including small and thick leaves, increased lateral buds, defective floral organs, small fruit size with reduced setting rate, and defective seed development, suggesting that hypomethylation of plant DNA can cause abnormal development and that DNA methylation may regulate many developmental processes in plants. Similar experiments were performed in Arabidopsis, in which antisense-MET1 lines with decreased methylation exhibited a series of phenotypic and developmental alterations, including smaller plant size, reduced apical dominance, altered leaf size and shape, altered flowering time, and decreased fertility [5, 6]. In addition, Arabidopsis ddm mutants with decreased DNA methylation [23] developed morphological abnormalities after several generations of self-fertilization [24].

CRISPR-Cas9-mediated knockout of SlCMT4 altered tomato plant morphogenesis and promoted lateral branch growth, which was confirmed through the increased number of lateral buds and faster growth rate. Anatomical analysis demonstrated that cortical cells and parenchyma cells in the pith of the lateral buds became larger in transverse and longitudinal sections, and the thickness of the cortical cell layers increased in SlCMT4 gene mutant lines. In our study, the contents of IAA and tZR in the lateral buds of SlCMT4 mutant lines increased, while the content of strigol decreased. The above results at hormonal levels were consistent with the multibranching phenotype in the SlCMT4 mutant plants. In addition, this study found that the mutant lines showed compressed architecture, short internodes, and small and thick leaves, which was confirmed through leaf morphological measurements and anatomical analysis. These results indicate that SlCMT4 acts as a DNA methyltransferase and plays a critical function in tomato plant morphogenesis.
CRISPR-Cas9-mediated knockout of \( \text{SlCMT4} \) seriously affected the structure and function of tomato flower organs, including defective and everted stamens, and short and thick pistils. Scanning electron microscopy revealed that the stamen cells and pollen grains of the mutant lines were irregular and defective. Pectin methylesterase (PME) is an important enzyme for the modification of side chains of pectin, and is regulated by the pectin methylesterase inhibitor \( \text{SlPMEI} \) gene in tomato [25]. The modification processes of pectin as the main component of the intine will directly influence the development of the intine. The development of the pollen wall is significantly crucial for maintenance of pollen morphology and function. The gene responsible for rapid alkalinization, \( \text{SlPRALF} \), encodes a preproprotein that negatively regulates pollen tube elongation in tomato [26]. In this study, we further assessed the transcriptional levels of two pollen-specific expression genes (\( \text{SlPMEI} \) and \( \text{SlPRALF} \)) in tomato. The results showed that \( \text{PMEI} \) and \( \text{PRALF} \) genes were respectively upregulated by \( \sim 4 \) - and \( \sim 2.5 \)-fold in pollen of mutant line CR-08. Our results indicate that CRISPR-Cas9-mediated knockout of \( \text{SlCMT4} \) induces the expression of these two pollen-specific genes, which are responsible for irregular and defective pollen. A possible reason is that the upregulated expression of \( \text{PMEI} \) and \( \text{PRALF} \) genes suppressed the development of the pollen wall and pollen tube elongation, resulting in the developmental abnormalities with reduced pollen fertility seen in the CR-08 line.

CRISPR-Cas9-mediated knockout of \( \text{SlCMT4} \) influenced fruit growth and ripening. In our study, the fruit size and weight of the \( \text{SlCMT4} \) mutant lines were reduced compared with WT, which was confirmed through measuring the horizontal diameter, vertical diameter, and weight of fruits at different maturity stages. Besides, transverse anatomical analysis of fruits at the IMG stage displayed reduced seeds in the CR-08 line. Based on the fruit phenotypic alterations mentioned above, we performed a relative transcriptional analysis of the fruit size-related gene \( \text{IMA} \), which has been reported to play a critical role in tomato fruit growth [27]. The \text{INHIBITOR OF MERISTEM ACTIVITY (IMA)} gene from tomato is involved in multiple regulatory pathways related to cell division, differentiation, and hormone regulation of tomato fruits. In plants overexpressing \( \text{IMA} \), the number of carpel cells was reduced, the carpels became smaller, and smaller flowers and fruits were produced, indicating that the expression of \( \text{IMA} \) inhibited cell division [27]. Our results showed that the fruit size-related gene \( \text{IMA} \) was upregulated in the fruits of the mutant lines at the IMG stage. Therefore, we deduced that the small-sized fruit phenotype in the mutant lines can be partially explained by reduced seeds and upregulated expression of the \( \text{IMA} \) gene in the mutant lines.

The function of methylation in determining the onset of ripening was first revealed by Zhong et al. [28]. They found that treatment of immature tomato fruits with the methyltransferase inhibitor 5-azacytidine...
induced premature ripening and demonstrated that DNA methylation promotes the regulation of fruit ripening. The binding sites of tomato transcription factor MADS-RIN, related to fruit ripening, are usually located in the demethylation regions of the promoters of many ripening-related genes. The binding of MADS-RIN to these promoters coincides with the demethylation of these sites [28], indicating that DNA methylation is also an important way to regulate fruit ripening. Furthermore, the mature mutant Cnr shows a spontaneous epigenetic change caused by methylation of the promoter of the SBP-CNR gene [29, 30]. Considering that fruit ripening is related to cell wall alterations [31] and synthesis of carotenoid and ethylene, we further studied the relative transcriptional levels of four ripening-related genes, PE1 (pectinesterase), PG2 (polygalacturonase) [33], LOXB [34, 35], and ACO1 [36, 37], in the WT and mutant tomato fruits at B stage. In this study, the PE1 and PG2 genes related to fruit wall metabolism were significantly upregulated during the B stage of fruit development in mutant lines. In addition, the lycopene synthesis-related gene LOXB was also significantly upregulated at the B stage of fruits of the mutant line. The ACO1 gene, related to ethylene synthesis, was upregulated during the B stage of fruit development in mutant lines. These results suggest that CRISPR-Cas9-mediated knockout of the DNA methyltransferase gene SlCMT4 may promote fruit ripening in tomato, which was partially demonstrated by the finding that ripening time of SlCMT4 RNai fruits was accelerated by 3–5 days compared with WT fruits in our study (Supplementary Table S1).

In this study, the number and weight of seeds in the mutant lines decreased; in particular, the number of seeds per fruit in the CR-08 line decreased by up to 70% and some seeds failed to develop and mature normally. The middle of the seeds in the CR-08 line was sunken, and the epidermal hairs of the seeds in CR-08 line were seen by scanning electron microscopy to be increased. We performed pollen observation by scanning electron microscopy of the WT and mutant lines. The results revealed that most pollen grains in the CR-08 line were aberrant and defective, suggesting that knockout of SlCMT4 significantly affected pollen grain development and led to the formation of defective pollen grains. Additionally, SlCMT4-knockout stamens and pistils were twisted and shorter, respectively. These changes in stamens, pollen grains, and pistils reduced the success of pollination. Therefore, the mutant line CR-08 had a lower fruit set rate. In angiosperms, branching patterns greatly determine plant architecture and affect nutrient allocation, height, light-harvesting efficiency, and visibility for pollinators [38]. Thus, vigorous vegetative growth of the SlCMT4 mutants probably accounts for the low fruit set percentage.

Measurement of organ architecture parameters
To characterize the differences in organ morphology between WT and mutant lines, we counted and measured the number, length, and diameter of lateral branches as well as the length, width, perimeter, and area of compound leaves and internode length from WT and mutant lines. The length, width, perimeter, and area of compound leaves from 3-month old plants were determined with a leaf area meter (Yaxin-1241, Beijing). The length and diameter of lateral branches were measured after 5 days of growth. Fruit size and weight were measured using a Vernier caliper (0–150 mm, Shanghai) and electronic balance (0–100 g, Shanghai). The fruit set of WT and mutant lines was also determined.

Anatomical and cytological analyses of lateral branches and leaves
The 5-day-old lateral branches and mature leaves collected from WT and mutant lines were immediately fixed with 2.5% glutaraldehyde. The experimental steps for tissue sectioning and staining were as follows: (i) dewaxing and dehydration; (ii) Safranin O staining; (iii) decolorization; (iv) fixed green staining; (v) placing sections into three cylinders of xylene for 5 minutes; and (vi) finally mounting the tissue sections with neutral balsam. The sections were observed under a microscope (Eclipse E100, Nikon, Japan) fitted with an image acquisition system (DS-U3, Nikon, Japan). The cell morphologies of lateral branches and leaves were measured and visualized using CaseViewer 2.3 software and photographed. The number
Scanning electron microscopy

Flower organs (stamens, pistils, and pollen grains) were collected from the inflorescences of WT and mutant lines at anthesis. Samples were fixed in 2.5% glutaraldehyde for 2 hours and subsequently dehydrated in an ethanol–water series. After dissection, the samples were dried in a critical point dryer and sputtered with gold for 30 seconds. The samples were observed and images were taken using a scanning electron microscope (SU8100, Hitachi, Japan).

Quantitative real-time PCR analysis

Total RNA was extracted from WT and mutant lines using an RNeasy Plant Mini Kit (Qiagen,74904), and RNA concentration was detected using a NanoQuant infinite (Amersham). RT–PCR was carried out according to a previous method [39]. The primers SlCMT4-F and SlCMT4-R were used to determine the expression levels of SlCMT4 in WT and mutant lines. All qPCR primers designed by Primer Express Software v2.0 are shown in Supplementary Table S2.

Extraction and quantification of plant hormones

Lateral branches were collected from 3-month-old plants (1.0 g per plant) and homogenized in liquid nitrogen. Extraction and quantification of plant hormones followed the methods reported by Kojima et al. [42] and Pan et al. [43].

Whole-genome bisulfite sequencing and data analysis

Genomic DNA was extracted from the pericarp tissues of WT and CRISPR line #08 fruits, and bisulfite treatment was implemented using the Zymo EZ DNA Methylation-Gold Kit. The raw sequencing data were filtered using SOAPnuke software [44]. Clean data were aligned to the tomato reference genome (NCBI version GCF_000188115.4_SL3.0) using Bismark software [45], and statistical information such as the comparison rate and bisulfite conversion rate of each sample was calculated. Bismark with default parameters was also used to remove duplicated sequence reads. Differentially methylated regions (DMRs) refer to certain DNA fragments in different samples that exhibit different methylation patterns in the genome. The DNA methylation level is the ratio of the number of reads that support methylation to the number of reads covering the site [46] and was calculated as follows:

\[ Rm_{\text{average}} = \frac{Nm_{\text{all}}}{(Nm_{\text{all}} + Nm_{\text{all}})} \times 100\% \]

DMRcaller [22] R was used to calculate and analyze DMRs. We used BEDTools [47] to calculate DMR-associated genes or other genome elements based on position. By default, a 1-bp overlap between DMR and genes or other elements is the DMR-associated gene or DMR-associated element. Furthermore, we performed GO enrichment and Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway enrichment analysis on these related genes.

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Author contributions

X.G. and Z.H. conceived and designed the research; J.Z. provided the experimental conditions; X.G., J.Q., Z.C., Y.Z., and H.S. performed the research; X.G. wrote and corrected the article. All authors have read and approved the final version of the manuscript.

Data availability

Raw transcriptome sequencing data reported in this paper have been deposited (PRJCA007796) in the Genome Sequence Archive in the BIG Data Center, Chinese Academy of Sciences, under accession code CRA005782 at https://ngdc.cnbc.ac.cn/.

Conflict of interest

The authors declare that they have no conflict of interest.

Supplementary data

Supplementary data is available at Horticulture Research online.

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