Study of micro-trichome (mict) reveals novel connections between transcriptional regulation of multicellular trichome development and specific metabolism in cucumber

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Abstract
Trichomes that cover the epidermis of aerial plant organs play multiple roles in plant protection. Compared with a unicellular trichome in model plants, the development mechanism of the multicellular trichome is largely unclear. Notably, variations in trichome development are often accompanied by defects in the biosynthesis of cuticle and secondary metabolites; however, major questions about the interactions between developmental differences in trichomes and defects in metabolic pathways remain unanswered. Here, we characterized the glabrous mutant mict / csgl1/cstbh via combined metabolomic and transcriptomic analyses to extend our limited knowledge regarding multicellular trichome development and metabolism in cucumber. Mict was found to be explicitly expressed within trichome cells. Transcriptomic analysis indicated that genes involved in flavonoid and cuticle metabolism are significantly downregulated in mict mutants. Further metabolomic analysis confirmed that flavonoids, lipids, and cuticle compositions are dramatically altered in mict mutants. Additional studies revealed that Mict regulates flavonoid, lipid, and cuticle biosynthesis by likely directly binding to downstream functional genes, such as CsTT4, CsFLS1, CsCER26, and CsMYB36. These findings suggest that specific metabolic pathways (e.g., flavonoids and cuticle components) are co-regulated by Mict and provide insights into transcriptional regulation mechanisms of multicellular trichome development and its specific metabolism in cucumber.

Introduction
Plant trichomes are highly specialized structures located on the surfaces of aerial organs. They play essential roles in diverse biotic and abiotic adaptation, for instance, deterring insects, herbivores, and microbes; secreting ions and contaminated metals1; reducing mechanical wear; regulating surface temperature; collecting and dispersing pollen; and absorbing water and nutrients2. Morphologically, trichomes can be glandular trichomes or non-glandular trichomes, unicellular or multicellular, and branched or unbranched. In addition, they have become an excellent model system for the analysis of cell differentiation as they are easily accessible in model plants such as Arabidopsis thaliana3. Trichome development was reported to have a close relationship with increased metabolite levels, such as flavonoids, anthocyanins, and phenylpropanoids4.

In the model plant Arabidopsis with non-glandular and unicellular trichomes, expression maps of leaf trichomes revealed high anthocyanin and flavonoid pathway activities, indicating the roles of trichomes in secondary metabolism and defense4. Similarly, in Brassica rapa with
non-glandular trichomes, overexpression of BraLTP2 leads to increased trichome density and affects the accumulation of secondary metabolites. In Solanum lycopersicum L., jasmonic acid–insensitive1 (jai1) is defective in JA signaling, and there was a significant reduction in the density and type of trichomes and monoterpene levels in the jai-1 mutant. In addition, Trichomes in the hairless (hl) mutant show abnormal morphology and density, reduced sesquiterpene and polyphenolic compounds, and increased susceptibility to insects and pathogens. In addition, ScCycB2, which encodes a B-type cyclin gene, is involved in the initiation of multicellular trichomes and terpenoid biosynthesis. Nevertheless, little is known about the developmental process and underlying regulatory molecular mechanisms of multicellular non-glandular trichomes.

Cuticles are composed of polymer matrix cutin that is covered and embedded with waxes, and both are synthesized by underlying epidermal cells. Cutin is a biopolyester composed of C16 and C18 hydroxy and epoxy fatty acids, while wax is a complex mixture of very long chain fatty acids (VLCFAs), predominantly of chain lengths from 26 to 34 carbons. Nevertheless, wax also includes secondary metabolites and triterpenoids, such as flavonoids and sterols. Wax aliphatic compounds are produced via the alcohol-forming and alkane-forming pathways, leading to VLC alkanes and their derivatives. eceferum (cer) mutants present abnormal wax compositions or constituents in Arabidopsis thaliana. According to mutant phenotype expression patterns and heterologous expression in yeast, CER6 participates in producing fatty acids with 26 and 28 carbons. Notably, it has been proven that CER26 has a major role in C30 elongation, with high tissue and substrate specificity.

Cuticles are also special structures on the epidermis of aerial plant organs. However, different from trichomes, which are generally considered “biofactories” for producing valuable natural products, the cuticle is the interface between the plant and its surrounding environment. In Artemisia annua, a novel HD-ZIP IV/MIXTA complex plays a significant role in regulating epidermal development, including cuticle formation and glandular trichome initiation.

In cucumber, trichomes are multicellular, unbranched cells that are regularly distributed on most aerial surfaces. tril/csgfl3 and csgfl/mict are important cucumber mutants that enable the dissection of trichome development into distinct, genetically controlled steps, as follows: Trichomeless (Tril) regulates the initiation and density of the trichome and Micro-trichome (Mict) allows trichomes to differentiate correctly. Both Tril and Mict are HD-ZIP transcription factors, different from the well-known MYB-WD40-bHLH complex in Arabidopsis. Unlike the unicellular trichomes in Arabidopsis, trichomes in cucumber allow cells to divide and to form into specialized trichome cells, such as fruit spines and fruit tumors. The regulatory network for the formation of trichomes and its relevance to cuticles in cucumber remains largely unknown.

In a previous study, we found that Mict regulates trichome initiation. In this study, we investigated the underlying metabolomic and transcriptomic changes in mict mutants and uncovered Mict regulatory pathways in the cuticle and secondary metabolism. The results provide valuable molecular information for further studies on cucumber fruit external quality traits related to market value.

Results

Mict loss of function mutants exhibit profoundly altered leaf and fruit epidermis properties

Mict has previously been reported to be involved in the development of trichomes in cucumber fruits and leaves. However, many other profound phenotypic changes were found in mict mutants; for example, mutant leaves looked brighter and glossier (Fig. 1a, b) and mutant fruit surfaces were smooth (Fig. 1c). However, the mict mutant sarcocarp did not show visible phenotypic differences in color, gloss, or cell morphology compared with wild-type (Fig. 1d). Gloss levels (gloss unit value) of the leaves and fruits were examined by a gloss meter, which showed that mict leaf but not fruit surfaces were glossier than those of the WT (Fig. 1e). The gloss levels of the mict leaves were significantly higher than those of WT leaves even when trichomes were removed artificially, indicating that the gloss unit value is not affected by the presence or absence of trichomes. This result implied that mutation of Mict affects not only epidermal properties, but also epidermal cell metabolism in cucumber.

Mict expression is highly associated with trichome development

To understand the in-depth function of Mict in the development of trichomes, comparative phenotypic observations of early developmental stages of trichomes were carried out (Fig. 2a).

The most apparent differences in trichomes between the wild-type and mict mutant occurred from Stages III to V, where the mutant trichomes maintained a bulge shape without cell division or elongation that occurred in the wild-type (Fig. 2a). Therefore, Mict regulates trichome development at early stages via its function in cell division or elongation.

We further examined the spatiotemporal expression pattern of Mict and revealed that Mict mRNA levels were dramatically elevated during the early trichome developmental stages, initiating from Stage I, sharply increasing at Stage II, and peaking at Stage IV while decreasing slightly.
in Stage V (Fig. 2b). Mict was highly expressed in the fruit spine and barely expressed in tissues without the spine (Fig. 2b). Noticeably, in situ hybridization showed that Mict transcripts were specifically detected in the first cell of the multicellular trichome (Fig. 2c–f), which indicated that Mict is involved in the early stage of trichome development, particularly in initiating the transformation of trichomes from single-cell bulges to cone-shaped multicellular trichomes.

Loss-of-function of Mict strongly affects global changes in genes involved in flavonoid pathways and cuticle metabolism

To reveal the global effects of Mict on leaf trichomes and fruit spine development, transcriptome analysis was performed to identify significantly changed metabolic pathways and putative downstream targets of Mict. For the RNA-seq of mict and wild-type plants, we collected expanded leaves at the four leaves seedling stage and fruits (the blooming female flower’s ovary). FPKM values were used to calculate gene expression levels and differential expression was defined by statistical parameters \((P < 0.05\) and fold change \(>2\) or \(<−2\)). A total of 2548 (leaf) and 711 (fruit) genes exhibited differential expression, including 1367 (leaf) and 481 (fruit) upregulated genes and 1181 (leaf) and 230 (fruit) downregulated genes (Table S1).

An analysis of Kyoto Encyclopedia of Genes and Genomes (KEGG) pathways showed that differentially expressed genes (DEGs) were significantly enriched in flavonoids, fatty acids, and cuticle biosynthesis metabolic pathways in both the leaves and fruits (red asterisks in Fig. 3a, and Table S1). Venn analysis identified a common set of 359 DEGs between the leaves and fruits, and heatmap analysis showed that they showed consistent expression patterns between these two tissues (Fig. 3b, c). STRING analysis\(^{18}\) was used to predict the functional association networks of the homologs to Arabidopsis among these 359 DEGs, which uncovered two arrays of functionally related genes that were closely associated with epicuticular cuticle formation, including waxes, cutin, and flavonoids (Fig. 3d, e). This was in agreement with previous reports of the components of cuticles in plants\(^{8}\).

Compositions of wax and cutin are largely changed in mict leaves and fruits

Gas chromatography-mass spectrometry (GC-MS) analysis was carried out to characterize the chemical composition of the leaf and the enzymatically isolated cuticle from mict fruit. Overall, compared with wild-type, the abundance of cutin monomer was increased approximately 2-folds in both the mict leaf and fruit. The significant increase in the amounts of fatty acids and midchain hydroxylated fatty acids in the leaf, and fatty acids and terminal hydroxylated fatty acids in the fruit mainly contributed to the accumulation of cutin monomers (Tables S2 and S3).

The total waxes in mict were reduced to 23.66% (leaf) and 53.33% (fruit) of the values in wild-type. The significant reductions in the amounts of alkanes contributed to reduced wax compounds in the mict leaf (Tables S4 and S5). The concentration of fatty acids, alcohols, and alkanes changed little in the mict fruit, and the reduction in total wax was mainly caused by the significant reduction in the concentrations of unidentified monomers. The cuticle
component differences in the leaf were more significant than in the fruit, which was consistent with the gloss measurements (Fig. 1e). These metabolic analyses clearly indicated that the loss-of-function of Mict also affects the leaf and fruit cuticle composition biosynthesis.

The profiles of flavonoids and lipids are altered in mict leaves and fruits

To further reveal the details of the metabolic changes, we performed a comparative metabolomic analysis in leaves and fruits of mict and wild-type plants using UHPLC-Q-TOF-MS. A total of 129 and 162 compounds with known structures were identified from the leaves and fruits, respectively. Among those, levels of 71 and 80 compounds were significantly different between the wild-type and mict mutant in leaves and fruits ($P < 0.05$), respectively (Table S6).

The most notable metabolomic changes were observed in flavonoids and lipids (Fig. 4). In mict leaves, 70.37% and 14.81% of detected flavonoids were decreased and increased, respectively. In mict fruits, 80.4% and 15.22% of detected flavonoids were decreased and increased, respectively. For lipids, in mict leaves, 69.23% and 30.77% of detected lipids were decreased and increased, respectively. In mict fruits, 36.54% and 63.46% of detected lipids were decreased and increased, respectively.

This metabolomic results suggest that Mict significantly affects lipid and flavonoid metabolic pathways.

Loss-of-function of Mict downregulates the expression of an array of genes involved in flavonoid and cuticle metabolism

We performed a metabolite-gene correlation analysis to further identify the association of flavonoids and cuticle
metabolism pathways with the putative Mict target genes. Among the DEGs, 15 putative cucumber orthologues of reported genes involved in cuticle formation in Arabidopsis showed significant downregulation in mict mutants (Fig. 5a). We further performed a qRT-PCR assay to test the expression levels of these genes in trichomes removed from leaves and fruit peels, which were in agreement with the cuticle composition analysis. We found that expression of MAH1-like and LACS3 were not significantly changed, the other cuticle-related genes were downregulated in trichomes removed from mict leaves (Fig. S1). In fruit peel, except for CER6 and HTH, genes related to the cuticle were also downregulated in mict, which was nearly consistent with the transcriptome data (Fig. S1).

According to the metabolomic and transcriptomic results, flavonoid metabolism was largely hampered in the mict mutant, particularly at the step catalyzing p-Cinnamoyl-CoA to kaempferol derivates (Fig. 6a). Moreover, the contents of naringenin chalcone, naringenin, and kaempferol derivates were substantially lower than that of the wild-type (Fig. 6c). Accordingly, the expression of the four key enzymes (i.e., TT4, CHI, F3H, and FLS1) that catalyze the above processes was repressed in the mict mutant. Other important genes, such as ANS, IFR, and HCT, were also downregulated to varying degrees in the mict mutant (Fig. 6b), implying that the flavonoid and flavone biosynthesis pathways are defective if Mict is dysfunctional.

Mict positively regulates the expression of CsTT4, CsFLS1, CsCER26, and CsMYB36 by binding to their promoters

Based on the multi-omic results that loss-of-function of Mict had a major impact on cuticle component metabolism (Fig. 4) and our finding that the downstream genes may contribute to the abnormal metabolic phenotype
(Figs. 5 and 6), we examined whether Mict could interact with the promoter of the revealed downstream target genes. We used a yeast one-hybrid assay to test the interaction between Mict and the promoters of 47 putative downstream target genes that are associated with the flavonoid and cuticle pathways (out of the 359 DEGs). The results showed that Mict could physically interact in vitro with four of these promoters (Fig. 7a), namely, CsTT4, CsFLS1, CsCER26 (Csa2G147920), and CsMYB36 (Csa2G352940). Meanwhile, co-infiltration of 35S:Mict
and promoter-LUC constructs resulted in significant transactivation of the promoters of the four tested putative Mict targets (Fig. 7b, c). To further explore the Mict binding region in the promoters, we segmented the promoters into four 500 bp long fragments (Fig. S3a). The results of yeast one-hybrid assay suggest that the CsFLS1, CsCER26, and CsMYB36 binding sites are located in the −2000 to −1500 bp regions of the promoter. For CsTT4,
the binding site is located in the −500 to 0 bp region in the promoter (Fig. S3b). In addition, these four genes were also significantly downregulated in the other two trichome mutants (trichome-less (tril) and glabrous 3 (gl3)), which exhibit recessive epistasis toward mict and show a similar phenotype to the mict mutant with bright leaves and reduced wax upload (Fig. 7d).

To further confirm the biological functions of CsTT4, CsFLS1, and CsCER26, we created CsTT4, CsFLS1, and CsCER26 overexpression Arabidopsis transgenic lines. The naringenin chalcone and kaempferol-3-O-glucoside contents were significantly increased in the oxCsTT4 and oxCsFLS1 transgenic plants, respectively (Fig. S2a, b), which suggests that CsTT4 and CsFLS1 are functionally conserved in the flavonoid biosynthesis pathway. The CsCER26-overexpressing plants showed a glossy phenotype compared with the wild-type plant stem (Fig. S2c), which was consistent with the AtCER26-overexpressing Arabidopsis plants13. Taken together, these results suggest that Mict likely affects flavonoid and cuticle metabolism as a direct action on the gene expression of its target genes and is associated with epidermal cell development and patterning.

**Discussion**

Based on previous studies, the determination and morphogenesis of the multicellular trichomes of cucumber are regulated by HD-ZIP transcription factors15−17,19,20, which show differences from unicellular trichomes. The known genes related to the cucumber trichome (spine) development are tril, mict, and Tu (tuberculate fruit). Mict mediates the morphogenesis of trichomes, and disruption of Mict makes most trichomes remain in the single-cell bulge state. In this study, evidence from our multi-omic analysis shows that Mict acts as a regulator in the very beginning phase to turn on an array of biological processes related to trichome development, directly or indirectly. Although the functions of Mict target genes need to be tested in further research, our results indicate that Mict can activate genes involved in flavonoid and cuticular lipid biosynthesis. The main purpose of this study was to identify the connection between transcriptional regulation and metabolic pathways in trichome development via multi-omics analysis and provide evidence to support that Mict regulates metabolic pathways (e.g., flavonoids and cuticle) both directly and indirectly.

**Mict turns on the regulatory network at the beginning of trichome development**

Based on our spatial-temporal expression analysis results, Mict is highly expressed at Stage III and Stage IV and is reduced at Stage V, implying that Mict functions at a particular stage during trichome development in leaf. Moreover, transcription of Mict can hardly be detected in any other organs except fruit spines, which implies that Mict acts as a regulator in the very beginning of trichome initiation, modulating a large number of downstream target genes for a transient period. Then, the regulated genes continually promote processes, such as cell division and secondary metabolite biosynthesis, to promote proper trichome development. Other evidence that Mict acts as a turn-on switch is that Mict is accurately detected in the top cell of the multicellular trichome, inducing successive trichome cells to develop spontaneously after Mict activates the regulatory network in the top cell. The expression of transcription factors (e.g., CsWIN1, CsMYB6, and CsMYB36) is absent in mict mutants, which implies complex interactions between transcription factors are involved in the transcript regulatory network. As a result, the mict mutant shows a defective phenotype in wax biosynthesis. In this study, we narrowed the Mict binding sites into a 500 bp region in the promoter of the four downstream genes. Common cis-element analysis of these four promoters suggested that only light response elements were identified in all four promoters (Fig. S3c). Taken together, Mict might bind novel elements in the promoters that specifically regulate trichome development, which merits further investigations.

**Mict regulates cuticle metabolism in epidermal cells**

The plant cuticle is an extracellular lipid structure deposited on the aerial surfaces of organs to protect the plant against environmental stresses21. Cuticles are complex mixtures of VLCFAs, with chain lengths of more than 26 carbons. In mutant leaf, the wax composition with more than 26 carbons changed significantly compared to that of WT. However, in mutant fruit, similar significant changes in wax components were lacking. Therefore, the change in VLCFA contents might contribute to observed phenotypes between the two organs. Transcriptomic study revealed significant DEGs involved in fatty acid and cuticle biosynthesis pathways, and led us to examine the cuticle’s wax composition. In combination with metabolomics data, we constructed a schematic diagram to explain wax changes, such as VLCFAs, alkanes, and secondary alcohols, in mict mutants (Fig. 5b). Down-regulation of CER26 and CER26-like indicated the elongation of 30 carbons or more was blocked in the mict mutant13. Downregulated CER8 in the mict mutant caused the lacing of alkanes and redundancy of free fatty acids22.

Moreover, low expression of CER1 and CER3, which act together to catalyze the formation of alkanes from VLCFA-CoA, also explains the lack of alkanes in the mict mutant23. MAH1, a midchain alkane hydroxylase that catalyzes secondary alcohol formation, was down-regulated and related to the absence of 2-C18:OL in the mict mutant24. In addition, downregulated ABCG13 and
ABCG32, functional ABC transporters involved in wax component transport, may also influence the transport of wax from the plasma membrane (PM) to the exterior of the cell. In summary, the downregulated expression of the genes mentioned above could partly explain some of the compound changes in cuticle biosynthesis pathways, which means that Mict positively regulates the downstream genes directly or indirectly to control these pathways in cucumber leaves and fruits. These results are consistent with the mutation of genes related to trichomes, such as GL1, MYB16, MYB106, and CER6, which can also affect cuticle formation in Arabidopsis.

Interestingly, a recent report found that the specific cuticular wax composition varies between different epidermal cell types, i.e., trichome and pavement cells. However, in this context, it is remarkable that Mict has much stronger expression in trichome cells than in other types of cells, leading us to speculate that Mict regulates the specific cuticular wax components. Notably, CER26 has much stronger expression in trichome cells than in other epidermal cells in Arabidopsis, making it reasonable that Mict positively regulates CsCER26 to synthesize the trichome-specific cuticular wax composition. It is technologically challenging for us to isolate multicellular trichomes from cucumber leaves, limiting an accurate examination of the trichome-specific cuticular wax composition. In the future, we believe that identification of the trichome-specific cuticular wax composition could reveal the function of Mict in cooperative regulation of trichome development and material metabolism.

**Mict regulates flavonoid metabolism in trichome development**

The trichome is regarded as a bio-factory of secondary metabolites in plants. The advantages of trichome-specific secondary metabolite biosynthesis could be maximizing the protective effects while minimizing the harm of their overaccumulation. Flavonoids are frequently used as pigments and involved in response to the environment and developmental processes. Transcriptional control of the structural genes in the flavonoid biosynthetic pathway has been most extensively studied in plants. In our research, Mict was shown to regulate catalytic enzymes, such as CsTT4 and CsFLS1, which participate in flavonoid biosynthetic pathways. The expression of catalytic enzymes usually determines the content of metabolites as they are always located in the down-stream regulation area. FLS is a key enzyme in flavonoid biosynthesis, and TRANSPARENT TESTA (TT) family members were found to work as transcription factors or enzymes in flavonoid biosynthesis pathways in Arabidopsis. Among those, CHALCONE SYNTHASE (CHS), which is encoded by the TT4 gene, catalyzes the first step of flavonoid biosynthesis for producing naringenin chalcone, a common precursor of various flavonoids. In addition, IF7MAT (isoflavone 7-O-glucoside-6-O-malonyltransferase), ANS (anthocyanidin synthase), HCT (O-hydroxycinnamoyltransferase), and IFR (isoflavone reductase) also play key roles in flavone, flavonol, and flavonoid biosynthesis pathways. The aforementioned genes were greatly downregulated in mict mutants, suggesting they may participate in the trichome-specific secondary metabolite biosynthesis pathway via Mict-mediated regulation. Accordingly, together with the genes–metabolites association analysis in this study, Mict may act upstream in a way that activates regulatory networks in trichome morphogenesis and specific metabolic pathways, which ensures that trichomes develop correctly.

**Mict activates CsMYB36 to form fruit wax-powders**

The cucumber fruit wax-powder is a thin layer of white powder that surrounds the fruit, and it is mainly composed of VLCFAs (and its derivatives), such as alkanes, aldehydes, and primary and secondary alcohols. Several cucumber mutants show no wax-powder on the fruits, such as csmyb36, tril/csgl3, and mict. It is clear that tril exhibits recessive epistasis with mict, as mict does to csmyb36, which suggests that CsMYB36 is the main regulator of wax-powder formation. In this study, we found that Mict could bind to the CsMYB36 promoter and stimulate its transcription. This result explains the absence of wax-powder on mict and tril fruits. Accordingly, the interaction between Mict and CsMYB36 directly bridges trichome development and wax-powder formation, and it provides us with a better understanding of the significance of trichome cell differentiation.

In this study, we identified the metabolic changes related to leaf trichome and fruit spine development in cucumber, and linked flavonoid and cuticle metabolism with transcriptional regulation. However, there are still many metabolites that we could not associate them with corresponding genes due to the limited cucumber molecular pathway databases. Fortunately, in recent years, with the rapid advances in trichome-related gene identification in cucumber, we believe our results will provide important clues to discover biochemical mechanisms in cucumber and reveal the molecular regulatory network involved in trichome development.

**Materials and methods**

**Plant materials**

The wild-type is cucumber (Cucumis sativus) North China type inbred line 06-1. The spontaneous mict mutant line 06-2 from 06-1 was grown under standard water, fertilizer management, and pest control with appropriate management in the glasshouse of Shanghai Jiao Tong University in Shanghai. Tobacco (Nicotiana benthamiana) and Arabidopsis were grown at 22 °C in 24 h continuous light indoors.
Gloss levels measurement

Fully expanded leaves and mature fruits of cucumber were samples for gloss levels measurement. The equipment we used is a Novo Curve Gloss Meter (RHOPPOINT, England). Each measurement was repeated four times.

Nucleic acid extraction and qRT-PCR

Genomic DNA was extracted from the expanded leaves using the CTAB reagent. Fully expanded leaves and ovaries of female flowers were sampled on the day before flowering for RNA and were frozen in liquid nitrogen immediately and stored at −80°C. The RNA extraction, cDNA reverse transcription, and qRT-PCR assays were performed as reported previously35. The gene-specific primers used for qRT-PCR were designed using Primer 3 software. Cucumber Actin3 and UBIQUITIN were used as internal controls to normalize and verify the expression data36,37. Three biological replicates were used per gene. Each qRT-PCR experiment was performed with three technical replicates. The gene-specific primers are listed in Table S7.

Materials for RNA-seq

Total RNA was extracted from fully expanded leaves and ovaries of female flowers on the day before flowering. RNASeq for comparative transcriptomic analyses of the two phenotypes was performed with three biological replicates. The library was constructed and sequenced using a BGISEQ-500 by the Beijing Genomic Institution (BGI, China). The genomic DNA was removed with two digestions with amplification grade DNase I (Epigenetics, USA). Then, the material was treated as previously reported36,37. The clean data have been uploaded to the China National GeneBank (CNGB) (Project ID: CNS0230824, CNS0230826, CNS0230828, CNS0230830, CNS0230832, CNS0230834, CNS0230836, CNS0230838, CNS0230840, CNS0230842, CNS0230844, and CNS0230846).

In situ hybridization

Young cucumber fruits at different stages of growth (1 cm, 1.5 cm, and 2 cm) were fixed with FAA. In situ sense and antisense probes were amplified with gene-specific primers with SP6 and T7 RNA polymerase-binding sites, respectively. Sample fixation, embedding and pretreatment of sections, hybridization, and immunological detection were performed as described previously38. Primers of antisense and sense probes are given in Table S7.

Bioinformatic analysis

We conducted transcriptome profiling experiments using the DGE approach39. Samples were collected from the ovaries of female flowers on the day of flowering and fully expanded leaves from the wild-type and mict mutant. Venn tools (Oliveros, 2007–2015) was used for Venn diagram analysis. To identify the homolog in Arabidopsis, we used the cucumber protein IDs40 to batch query the Arabidopsis proteins (TAIR10) using BLASTP on the Cucurbit Genomics Database (http://cucurbitgenomics.org) with an e-value cut-off of 1e−1. Based on the cucumber homolog in Arabidopsis, we predicted the protein interaction network using STRING (https://string-db.org).

Yeast one-hybrid assay

For the yeast one-hybrid assay, the Mict open reading frames (ORFs) were amplified from 06-1 genomic DNA and then cloned into the pB42AD vector (digestion with the EcoRI and XhoI restriction enzymes) using ClonExpress II (Vazyme, Nanjing, China). The 2 kb promoter of CsTT4, CsFLS1, CsCER26, and CsMYB36 from 06-1 were inserted into the vector (digestion with the EcoRI and XhoI restriction enzymes). pB42AD-Mict and placZ-CsTT4Pro, placZ-CsFLS1Pro, placZ-CsCER26Pro, or placZ-CsMYB36Pro were co-transformed into the yeast strain EGY48a. The empty vectors were used as negative controls. These transformants were cultivated to confirm whether Mict can bind the promoter of the CsTT4, CsFLS1, CsCER26, and CsMYB3641. All of the primers used in this study are listed in Table S7.

Dual-luciferase assay

The assay was carried out as described previously20. Briefly, Mict ORFs were cloned into pHB driven by the 2’35S promoter to obtain overexpression vectors. Then, the vectors were separately transformed into Agrobacterium tumefaciens GV3101 as effectors. The promoters of CsMYB36, CsFLS1, CsTT4, and CsCER26 from 06-1 were cloned into pGREEN 0800 to drive the firefly luciferase reporter gene. The CsMYB36, CsFLS1, CsTT4, and CsCER26 promoter vectors were then co-transformed into the yeast strain EGY48a. The empty vectors were used as negative controls. These transformants were cultivated to confirm whether Mict can bind the promoter of the CsTT4, CsFLS1, CsCER26, and CsMYB3641. All of the primers used in this study are listed in Table S7.

Ectopic expression of CsTT4, CsFLS1, and CsCER26 in Arabidopsis

To generate lines overexpressing CsTT4, CsFLS1, and CsCER26, the full-length coding regions of these three genes were amplified using specific primers containing the BamHI (5’ end) and SacI (3’ end) endonuclease sites, and inserted in the reverse orientation into the pHB vector. The recombinant plasmids were transformed into Col (WT) using the floral dip method42.

Metabolite extraction and profiling

The fully expanded leaves and ovaries of female flowers on the flowering day from the wild-type and mict mutant were ground into a fine powder. Twenty milligram of fine
powder was used for metabolite extraction prior to UHPLC-Q-TOF-MS analysis. As the compound content was calculated by weight, the trichomes and fruit spines were not removed from the samples, which was similar to the transcriptome analysis. The metabolite extraction was analyzed as previously described. All measurements were performed with three replicates.

Analysis of cuticular wax and cutin chemical analysis

As the compound content is calculated from the surface area, we removed the trichomes and fruit spines from the samples to reduce the area errors caused by the trichomes. For the fruit peels, we collected the peels from fruits at the 10th day after flowering so that the fruits were elongated enough that we could isolate the peel from the trichomes. Cucumber cuticular wax was extracted from fully expanded leaves and the fruit pericarp on the 10th day after flowering. Three independent wild-type (WT) and mict mutant lines were analyzed. For cutin analysis, fully expanded leaves and fruits were delipidated in a 10 ml mixture of methanol–chloroform (1:1, v:v) for 2 weeks (solvent changed daily), after which the material was dried, weighed, and used for analysis.

Statistical analyses

All data are expressed as mean value ± standard deviation (SD) of biological replicates. Statistical significance of the treatment differences was assessed using the Student’s t-test when only two components were compared. In addition, P values are from one-way ANOVA (Tukey’s multiple comparisons test, α = 0.05). GraphPad Prism version 8.2.1 for Windows (GraphPad Software, La Jolla, CA, USA) was used for the statistical analyses.

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JS.P, G.W., J.P., and L.Z. conceived the project and designed the research. J.P. and L.Z. performed most of the experiments with help from G.C., H.W., Y.C., H.D., and J.Z.; J.P. and L.Z. analyzed the data and wrote the article with assistance from HH., H.L., H.C., J.S., R.C., G.W., and JS.P.; J.P., J.Y., and L.Z. supervised the project and completed the writing.

Conflict of interest
The authors declare that they have no conflict of interest.
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