Gain-of-function of the 1-aminocyclopropane-1-carboxylate synthase gene ACS1G induces female flower development in cucumber gynoecy

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Unisexual flowers provide a useful system for studying plant sex determination. In cucumber (Cucumis sativus L.), three major Mendelian loci control unisexual flower development, Female (F), androecious [α; 1-aminocyclopropane-1-carboxylate (ACC) synthase 11, acs11], and Monoecious (M; ACS2), referred to here as the Female, Androecious, Monoecious (FAM) model, in combination with two genes, gynoecious (g; the WIP family C2H2 zinc finger transcription factor gene WIP1) and the ethylene biosynthetic gene ACC oxidase 2 (ACO2). The F locus, conferring gynoecy and the potential for increasing fruit yield, is defined by a 30.2-kb tandem duplication containing three genes. However, the gene that determines the Female phenotype, and its mechanism, remains unknown. Here, we created a set of mutants and revealed that ACS1G is responsible for gynoecy conferred by the F locus. The duplication resulted in ACS1G acquiring a new promoter and expression pattern; in plants carrying the F locus duplication, ACS1G is expressed early in floral bud development, where it functions with ACO2 to generate an ethylene burst. The resulting ethylene represses WIP1 and activates ACS2 to initiate gynoecy. This early ACS1G expression bypasses the need for ACS11 to produce ethylene, thereby establishing a dominant pathway for female floral development. Based on these findings, we propose a model for how these ethylene biosynthesis genes cooperate to control unisexual flower development in cucumber.
Introduction

Unisexual flower development represents a model experimental system to decipher the regulatory mechanism of sex determination in plants (Tanurdzic and Banks, 2004; Ma and Pannell, 2016). Considerable attention has been devoted to identifying genes that control male and female flower development (Banks, 2008; Aryal and Ming, 2014). Cucumber (Cucumis sativus L.) represents one such system (Malepszy and Niemirowiczczszytt, 1991; Ainsworth and Buchanan-Wollaston, 1997), and the female flower ratio is an important yield trait for cucumber as only female flowers develop economically valuable fruit.

A pioneering genetic study established that different cucumber lines exhibit stable unisexual flower distribution phenotypes (Galun, 1962). These phenotypes included plants that produce only female flowers (termed gynoecious), both male and female flowers (termed monoecious), only male flowers (termed androecious), only bisexual flowers (hermaphrodite), and male and bisexual flowers (termed andromonoecious) (Perl-Treves, 1999). Major genetic loci were identified, named Female (F), androecious (a), and Monoecious (M), which determine the distribution of unisexual flowers (Galun, 1962; Kubicki, 1969a; Kubicki, 1969b, 1969c). For convenience, we term this system the FAM model. Gynoecious cucumbers conferred by the F locus have been used, for almost half a century, specifically in breeding greenhouse varieties with high yield potential.

Early studies also identified ethylene as a key regulator in male and female flower development; for example, application of ethephon promotes female flower development (Wittwer and Hillyer, 1954; Iwahori et al., 1970; Byers et al., 1972). Although other phytohormones were examined for their effects on the ratio of unisexual flowers, ethylene was widely accepted as a key regulator of unisexual flower development in cucumber (Yin and Quinn, 1992).

During this same period, the ethylene biosynthetic pathway was deciphered, and this established that methionine is converted into S-adenosylmethionine (SAM) by methionine adenosyltransferase; SAM is then converted into 1-aminocyclopropane-1-carboxylate (ACC) by ACC synthase (ACS). Finally, ACC is oxidized to ethylene by ACC oxidase (ACO). Here, ACS and ACO function as key enzymes for ethylene production (Adams and Yang, 1979). Almost all major genes controlling flower sex type encode key enzymes involved in ethylene biosynthesis, with WIP1, which encodes a C2H2 zinc finger transcription factor of the WIP family, being the exception. The M gene encodes ACS2, another ACS expressed in the carpel region of the female flower; its inactivation (m) results in formation of a bisexual flower (Boualem et al., 2008; Li et al., 2009). The A gene encodes ACS11 and a dysfunctional ACS11 (a gene) blocks the female flower developmental pathway (Boualem et al., 2015).

The ACO2 was also shown to affect unisexual flower development, through its coordination with ACS11 in floral preferential production of ethylene; defective ACO2 results in the loss of female flowers (Chen et al., 2016). Finally, WIP1 was identified as a C2H2 zinc finger transcription factor required for male flower development; its expression is repressed by ACS11 (Martin et al., 2009; Boualem et al., 2015; Hu et al., 2017). Without WIP1 expression, or when WIP1 is dysfunctional, cucumber plants produce female flowers, but with some bisexual flowers being formed at the lower (earlier) nodes (Hu et al., 2017).

Although many studies have clarified the roles of other loci, the mechanisms by which the F locus affects sex determination in cucumber remain unclear. The discovery that the F locus contains an additional copy of ACS1 (diagrammed in Figure 1A) provided an initial link between ethylene production and female flower development (Trebitsh et al., 1997). However, although the F gene was suggested to encode gynoecious Cucumis sativus ACC synthase (ACS1G) (Mibus and Tatioglu, 2004; Knopf and Trebitsh, 2006), genome-wide mapping of structural variation revealed that the F locus is a 30.2-kb tandem duplicated region. This region contains not only ACS1G but also other genes (Zhang et al., 2015) that, to date, have not been ruled out as participants in the F locus. Additionally, currently, there is no evidence to determine whether the female promotion function of the F locus represents a dosage effect of multiple copies of ACS1, or alternatively, a function conferred by ACS1G by acquiring a novel promoter through recombination.

An earlier study using RNA interference to downregulate ACS1 and ACS1G expressions, in the cucumber FF genotype, reported that this caused a transition from gynoecy to monoecy, which was offered as support for the hypothesis that ACS1G confers a simple dosage effect (Shiber et al., 2008). However, this result could not distinguish between the functions of ACS1 and ACS1G, as the entire untranslated and coding regions of ACS1 and ACS1G are identical, and the cucumber FF genotype contains both ACS1 and ACS1G. Thus, the genetic regulatory mechanism underlying the molecular basis of the F gene remains to be fully elucidated.

Another problem in deciphering the molecular mechanism underlying the genetic control over unisexual flower development is that, in the current FAM model, the criteria to define a unisexual flower are based on the entire flower; i.e., male or female flowers. However, unisexual flowers result from abnormal floral organ development (Dellaporta and Calderon-Urrea, 1993; Ainsworth and Buchanan-Wollaston, 1997). Although it was established that ethylene plays a key role in cucumber female flower development, little is known about whether ethylene is required for carpel development and/or inhibition of stamen development.

In this regard, it was demonstrated that primordial anther-specific DNA damage is closely correlated with stamen arrest in the female cucumber flower (Hao et al., 2003). Subsequently, it was shown that this anther-specific DNA damage was caused by the stamen preferential downregulation of the ethylene receptor (ETR) gene, Cs-ETR1 (Wang et al., 2010), and such organ preferential downregulation of Cs-ETR1 is caused by a B-class APETALA (AP)-type MADS (MCM1, AG, DEF, SRF)-box gene, Cs-AP3 (Sun et al., 2016).
Data are presented as mean YB-Mo and 09-Mo are monoecious; YB-Gy and 09-Gy are gynoecious. YB-Mo and YB-Gy, and 09-Mo and 09-Gy, are isogenic lines, respectively.

Finally, ethylene-induced expression of a calcium-dependent DNase gene, Cs-CaN, is correlated with the anther-specific DNA damage (Gu et al., 2011). Taken together, these findings support the hypothesis that ethylene plays a direct role in the inhibition of anther primordia development, in female cucumber flowers, through a stamen-specific ethylene perception and downstream response. However, a remaining question is why are ethylene synthesis genes playing key roles in promoting female flower development?

Based on current information, it would appear that all genes known to be involved in unisexual flower development, such as M (ACS2), A (ACS11), WIP1, and ACO2, are expressed preferentially in the carpel primordia (Yamasaki et al., 2001; Boualem et al., 2008; Li et al., 2009; Martin et al., 2009; Boualem et al., 2015; Chen et al., 2016). Although there is no direct evidence for cucumber as to whether ethylene is required for carpel development, in tobacco (Nicotiana tabacum), it has been shown that ethylene is required for ovule development (De Martinis and Mariani, 1999).

Analysis of the pre-microRNA distribution in male, female, and hermaphrodite cucumber flowers suggested that ethylene is required for carpel development, by rescuing stress inhibition of carpel primordia (Sun et al., 2010). This ethylene-mediated inhibition of anther primordia development, in the female flower, might have been a side-effect of carpel rescue, and such an inhibition could well have been selected through advantages conferred by outcross pollination. In our previous study, we proposed that the F gene might have been co-opted, after the M and A genes, to a regulatory mechanism to enhance carpel development (Sun et al., 2010). From this perspective, clarification of the role played by the F gene is central to deciphering the molecular mechanism of unisexual flower development in cucumber; this would likely provide the last piece to complete the puzzle of the genetic FAM model.

In the present study, we sought to clarify the molecular basis of the F locus to provide a complete scenario for the FAM model of cucumber unisexual flower regulation, at a molecular level. To this end, we used a combination of CRISPR/Cas9 gene editing and ethyl methanesulfonate (EMS) mutagenesis to assess the functions of the genes within the F locus in determining female flower development. In addition, we transformed monoecious cucumber with the entire ACS1G genomic region and these lines became gynoecious. Our findings revealed that ACS1G, not ACS1 or other genes in the F locus, is the gene responsible for development of female flowers. We further demonstrate that ACS1G is strongly expressed in every floral bud, at early stage 2, bypassing the need for ACS11, thereby establishing a dominant pathway for female floral development. This ACS1G expression pattern is conferred by the promoter of ACS1G. Based on these findings, we propose a model to explain how the various ethylene biosynthesis genes function to ensure successful carpel development in cucumber.

**Results**

**F locus genomic structure**

The F locus contains two annotated genes, CsaV3_6G044400 (ACS1) and CsaV3_6G044410, and a part of CsaV3_6G044420 (Branched Chain Amino Acid Transaminase, BCAT). CsaV3_6G044410 encodes a cucumber MYB family transcription factor, hereafter identified as MYB1. In gynoecious...
plants carrying the F locus, the entire MYB1 region is duplicated. In addition, within this region, ACS1G gained a part of the BCAT gene, likely functioning as a distal promoter, but it retained the same proximal promoter, untranslated, and coding regions as ACS1. We also ascertained that the gynoecious lines (Ff genotype) retained a complete copy of the ACS1 and BCAT genes, as in the monococious lines (Figure 1A).

Considering the complicated structure of the F locus, for clarity, we developed nomenclature to distinguish the genes in this region. ACS1G stands for the gynoecious-specific ACS1 copy with a novel promoter; the ACS1 copies on the F and f haplotypes are designated as ACS1F and ACS1f, respectively, and have identical genomic and flanking sequences. However, ACS1F is always genetically linked to ACS1G. The MYB1 copy in the f haplotype was named MYB1f and the two MYB1 copies in the F haplotype were termed MYB1a (S’ to ACS1G) and MYB1b (S’ to ACS1G); these three MYB1 copies have identical genomic and flanking sequences (Figure 1A).

To explore the F gene identity, we first assessed the expression level, within floral buds, of the three candidate genes from near-isogenic gynoecious and monococious lines. ACS1G has a different promoter as both ACS1F and ACS1G share identical untranslated and coding sequences; therefore, we were unable to distinguish between their expression levels. Through RT-qPCR (Real-time Polymerase Chain Reaction) assays, we determined that ACS1F/ACS1G and MYB1 had elevated expression levels in floral buds (pre-stage 4) from gynoecious compared to near-isogenic monococious lines (Figure 1, B–D). Based on these expression profiles, we concluded that BCAT is unlikely to be the F gene (since its expression was unchanged between lines), but we could not discount the possibility that MYB1 functions as the F gene.

**Gynoecy requires a functional ACS1G**

To assess the impact of ACS1F, ACS1G, MYB1a, and MYB1b on the gynoecious phenotype, we sought to generate loss-of-function mutations in each of these genes. The FF genotypes are recalcitrant to current transformation protocols (Hu et al., 2017); therefore, we employed a CRISPR/Cas9 gene-editing (CR) approach, using the transformable cucumber ff genotype, to generate loss-of-function MYB1CR mutants. These transformed T0 seedlings were then used as the male parent and crossed with FF near-isogenic lines (Supplemental Figure 1, A and B). Pollen containing the Cas9 transgenic fragment was next employed to generate F1 plants in which the target genes, located on the F haplotype, were further edited (Figure 2A). It is important to stress that, in the F1 plants, generated from pollen containing the transformed T0 seedlings but not containing Cas9, only the target gene located on the f haplotype was modified, as gene-editing occurred in the T0 generation of the ff genotype plants (Figure 2A; Supplemental Figure 1B).

In the 31 tested F1 plants not carrying Cas9, only MYB1f was mutated (MYB1fCR, Figure 2A; Supplemental Figures 1, A and 2); in the 32 tested F1 plants carrying Cas9, each of the three MYB1 copies was mutated (MYB1AllCR, Figure 2A; Supplemental Figures 1B and 3). Sequencing and sex phenotyping derived from 15 randomly selected seedlings are shown here (Figure 2C; Supplemental Figures 2 and 3). Given that both mutant types were gynoecious (Figure 2C), we could exclude MYB1 as the F gene.

Similarly, we generated Ff plants with only mutated ACS1f (ACS1fCR, Figure 2B; Supplemental Figures 1B and 4) or all copies of ACS1 mutated (ACS1AllCR, Figure 2B; Supplemental Figures 1B and 5). Here, ACS1fCR remained gynoecious, whereas the ACS1AllCR plants were identified as monococious (Figure 2C). These findings indicated that either ACS1F or ACS1G, or both ACS1F and ACS1G together, in the F locus, conferred the gynoecy phenotype.

Since ACS1F and ACS1G have identical untranslated and coding sequences, and the distance on chromosome 6 between ACS1F and ACS1G is only 30.2 kb, it is hard to distinguish their function simply by using CRISPR/Cas9 (Clustered Regularly Interspersed Short Palindromic Repeats/Crispr associated protein 9) or RNAi methods. However, whether gynoecy is conferred by a simple dosage effect, caused by two copies of ACS1, or by a novel ACS1G promoter conferring a special expression pattern, is an important question.

To further distinguish the functions of ACS1F and ACS1G, we constructed an EMS mutation library based on the ff genotype sub-gynoecious line. It is extremely difficult to acquire seeds of EMS-treated gynoecious or sub-gynoecious lines containing the F locus because they are much more sensitive to the effects of EMS than the monococious lines. These lines rarely bear male flowers, and it is not possible to use the Ag+ treatment to induce male flowers because this typically exacerbates the toxicity of the EMS and hinders seedling growth and flowering. In any event, we performed EMS mutagenesis on three different Ff-background cucumbers several times, screened 1,500 M1 plants, and obtained two nonsynonymous mutations, ACS1fE353K and ACS1GHI64Y, from the YB-Ff and 09-Ff backgrounds, respectively (Figure 3, A and B; Supplemental Figure 6, A and B).

These two mutations, located in nonconserved domains (Figure 3C), did not compromise either the sub-gynoecious phenotype (Figure 3D; Supplemental Figure 6C), or the ACS1 enzyme activity of these mutant proteins in terms of ethylene production (Figure 3E; Supplemental Figure 6D). However, these two single-nucleotide polymorphisms (SNPs) provided us with an opportunity to distinguish transcripts from ACS1f, ACS1F, or ACS1G, at the key developmental stages when the majority of transcripts from the F gene would be expected, since it causes gynoecy.

To use these SNPs to look at transcript levels of these ACS1 genes, RNA was extracted from early-stage female-desined floral buds (pre-stage 4) located at nodes 10 and above on ACS1fE353K (FF) and ACS1GHI64Y (Ff) mutant plants (Figure 3D). The mixed transcripts from ACS1f, ACS1F, and ACS1G were amplified, using primers common to their coding regions, and the PCR products were then cloned into a T-vector. At least 8 randomly selected positive clones were...
sequenced for each plant, and at least 20 independent plants were analyzed for each mutant type. We next used the number of clones corresponding to the different SNPs to infer a ratio for the ACS1f, ACS1fCR, and ACS1G transcripts. In the ACS1fE353K(Ff) line, only wild-type transcripts were detected (Figure 3F), indicating that ACS1f was not expressed at the key floral developmental stage in the Ff background. This finding indicated that ACS1f is not involved in promoting female floral development, consistent with the observation that ACS1fCR(Ff) plants are gynoecious (Figure 2C).

As ACS1f was not expressed at the key developmental stage, in the Ff background, the ACS1GH164Y(Ff) line was used to distinguish transcripts from ACS1F (wild-type) and ACS1G(CR) transcripts.
Figure 3 ACS1G transcripts are dominant in flowers at the key stage of female flower determination in gynoecious cucumber lines. (A and B) Schematic representation of the ACS1G<sup>H164Y</sup> (A) and ACS1f<sup>E353K</sup> (B) mutant genotypes with the Ff background. (C) Alignment of ACS1 homologous proteins from representative species. (D) The ACS1G<sup>H164Y</sup> and ACS1f<sup>E353K</sup> mutations did not affect the gynoecious flower sex types in the Ff background. Orange arrow indicates the boundary node between lower male nodes and upper female nodes, usually located at about node 10. (E) Enzymatic activity indicated by the ethylene production of wild-type ACS1/1G (green bars; WT), ACS1G<sup>H164Y</sup> (orange bars) and ACS1f<sup>E353K</sup> (purple bars) protein isoforms at various concentrations of SAM. Data are presented as mean ± SE, n = 3. *p < 0.05 and **p < 0.01 in Student’s t-test compared to the wild-type ACS1/1G protein isoform (Statistical data are provided in Supplemental Data Set 1). (F) Number of T-clones containing WT and ACS1f<sup>E353K</sup>-type transcripts in the ACS1f<sup>E353K</sup> mutant line with the Ff background. A total of 20 individuals was analyzed, and for every individual at least 8 independent T-clones were sequenced to identify the SNP. (G) Number of T-clones containing WT ACS1F and ACS1G<sup>H164Y</sup> in the ACS1G<sup>H164Y</sup> mutant line with the Ff background. A total of 24 individuals was analyzed, and for every individual at least 12 independent T-clones were sequenced to identify the SNP.
ACS1G (H164Y SNP) plants. Our sequencing results revealed that during key stages determining the female flower, the major transcripts were derived from the ACS1G allele (Figure 3G); ACS1F transcripts were detected in only 3 of 24 plants, equating to 3 of 411 clones. In the independent 6 ACS1F<sup>WT</sup>ACS1G<sup>H164Y</sup>ACS1F<sup>E353K</sup>(ff) plants acquired from the cross between ACS1f<sup>E353K</sup>(ff) and ACS1G<sup>H164</sup>(ff) mutant plants (Supplemental Figure 7, A and B), the ACS1F clone was detected only once in a total of 65 clones (Supplemental Figure 7C), whereas all the other clones were ACS1G<sup>H164Y</sup>. These findings provided strong support for the hypothesis that ACS1G is the F gene.

To further test our hypothesis that ACS1G is the F gene, we generated transgenic monoecious cucumber expressing the entire ACS1G genomic region, including a 4.5-kb region upstream from the start codon. Compared to the untransformed cucumber ff genotype (Figure 4B), expression of this ACS1G genomic region resulted in a gynoecious phenotype (Figure 4, A, C, and D). These findings established that ACS1G alone is necessary and sufficient to confer gynoecy in cucumber.

The novel ACS1G promoter confers a new expression pattern and female-promoting function

Next, we asked whether the female promotion function of ACS1G is conferred by a simple dosage effect of multiple copies of ACS1, or alternatively, by a new expression pattern due to its promoter. To explore these possibilities, we first performed in situ hybridization assays to monitor the expression pattern of ACS1G (Figure 5, A–C; Supplemental Figure 8, A–E). Given that, transcripts in the developing Ff floral buds, at pre-stage 4, were derived predominantly from ACS1G (Figure 3G), we reasoned that in situ hybridization, based on FF genotype plants, would reflect the expression pattern of ACS1G, rather than that of ACS1F. Here, our assays revealed that ACS1G was expressed mainly during the early stages 1 and 2 of floral development (Figure 5, A and B; Supplemental Figure 8, A and B). At stage 2, ACS1G was expressed just beneath the central zone of the floral meristem, the region that can later develop into the carpels (Figure 5B). At stage 4, weak ACS1G signals were detected in the developing petal, stamen, and carpel primordia (Figure 5C; Supplemental Figure 8C). In monoecious cucumber (ff genotype), we detected no specific ACS1f expression pattern in floral bud stage 1 or 2 (Figure 5, D and E; Supplemental Figure 8, K and L). However, ACS1f was weakly expressed later, at stage 4 (Figure 5F; Supplemental Figure 8M).

To address whether these two expression profiles were conferred by the different promoters of ACS1G and ACS1f, we next generated ProACS1G:GUS and ProACS1f:GUS transgenic plants. GUS (β-glucuronidase) staining assays established that, from stages 1 to 3, the ACS1G promoter was expressed strongly (Figure 6, A–D); however, in the ProACS1f:GUS transgenic floral buds, at stages 1–3, no GUS signal was detected (Figure 6, E–H). These patterns are similar to those obtained by in situ hybridization.

In later floral development, at stages 4–8, the ProACS1G:GUS remained active within the lower region of the floral bud (Supplemental Figure 9, A–D), whereas ProACS1f:GUS generated only a weak GUS signal (Supplemental Figure 9, E–H). This ProACS1G:GUS pattern differed somewhat from that observed by in situ hybridization. This might reflect more distant cis-elements regulating the ACS1G expression, at these later developmental stages, which may be absent from our ProACS1G:GUS construct. In any event, collectively, these findings support the conclusion.

Figure 4 Cucumber transgenic lines expressing the ACS1G genomic region exhibit the transition from monoecy to gynoecy. (A) Flower sex type of transgenic CU2 (ff) carrying the ACS1G genomic region. Scale bar = 10 cm. (B) Flower sex type of CU2 (ff). Scale bar = 10 cm. (C) Genomic verification of the transgenic fragments using primers for the right border. (1) transgenic CU2 plant expressing the ACS1G genomic region; (2) CU2; (3) plasmid as the positive control. (D) Genomic verification of the transgenic fragments using primers for bar, an antibiotic selection gene. (1) transgenic CU2 plant expressing the ACS1G genomic region; (2) CU2; (3) plasmid as the positive control.
that a recombination in the cucumber genome gave ACS1G a new promoter and function.

Genetic relationship among the genes involved in determining the ratio of unisexual flowers

In monoecious cucumber, Androecious (A) is encoded by ACS11, which is expressed specifically in the carpel regions of floral buds destined to be female, where it cooperates with ACO2 to release the ethylene essential for initiating the female developmental pathway (Boualem et al., 2015; Chen et al., 2016). In contrast, in the presence of the female-promoting function of ACS1G, ACS2 expression is up-regulated dramatically in the gynoecious lines (Figure 7C). Based on these findings, we concluded that the spatial and temporal expression pattern of ACS1G, rather than a simple dosage effect of ACS1 due to copy number variation, confers the female promoting function of ACS1G.

Loss-of-function mutations in WIP1 also promote female flowers in cucurbits (Martin et al., 2009; Hu et al., 2017; Zhang et al., 2020). Furthermore, WIP1 expression is repressed by ACS11 expression in cucumber and melon (Boualem et al., 2015). Therefore, we next investigated whether ACS1G expression and ethylene production repress WIP1 expression. Compared with the isogenic monoecious lines, WIP1 expression was reduced significantly in the gynoecious line, aco2/aco2 (acs1g/acs1g), all nine ACS1G/aco2/aco2 genotype plants were androecious (Table 1). This result is consistent with our earlier findings that the application of ACC does not rescue the androecious phenotype of the aco2 homozygous mutants (Chen et al., 2016) and demonstrated that F gene function is dependent on ACO2 and, further, that ACS1G must cooperate with ACO2 to produce the required ethylene dosage.

We next investigated whether the ACS1G/aco2/aco2 genotype plants (where ACS1G/aco2/aco2 includes ACS1G/ACS1G and ACS1G/acs1g) could produce female flowers. ACS1G is linked to ACO2 on chromosome 6, with a physical distance of 2 Mb. Of 187 F2 plants obtained from a cross between YB-Gy (ACS1G/ACS1G ACO2/ACO2) and 406a (acs1g/acs1g aco2/aco2), all nine ACS1G aco2/aco2 genotype plants were androecious (Table 1). This result is consistent with our earlier findings that the application of ACC does not rescue the androecious phenotype of the aco2 homozygous mutants (Chen et al., 2016) and demonstrated that F gene function is dependent on ACO2 and, further, that ACS1G must cooperate with ACO2 to produce the required ethylene dosage. Based on these findings, we concluded that the spatial and temporal expression pattern of ACS1G, rather than a simple dosage effect of ACS1 due to copy number variation, confers the female promoting function of ACS1G.
Discussion

New technology provided the opportunity to complete the puzzle of the FAM model

The ACS1G gene was the first ethylene biosynthesis gene reported to be linked with the gynoecy phenotype in the FAM model (Trebitsh et al., 1997). However, due to the complexity of the F locus genome structure (Zhang et al., 2015), definitive evidence demonstrating that ACS1G is responsible for female flower determination, as opposed to other genes within the F locus, was lacking. Furthermore, a full molecular dissection of the F locus is essential for understanding unisexual flower development in cucumber.

In this study, we demonstrate that only ACS1G, not BCAT, MYB1a/b or ACS1, functions as the F gene (Figures 1–4). In addition, we discovered that the ACS1G expression pattern was created by a combination of the BCAT intron and the ACS1-coding sequence. Expression driven by this promoter resulted in ACS1G acquiring dominance over other ethylene synthesis genes, in regulating female flower development. Thus, we have now obtained the final molecular information to complete the puzzle of the FAM model.

It is intriguing that it took two decades to establish that ACS1G is the functional gene of the F locus. Naturally, in the absence of an annotated cucumber genome, the complex structure of the F locus represented a significant challenge. As an example, the presence of MYB1, in the F locus, was not revealed until 2015 (Zhang et al., 2015). Next, downregulation of ACS1/ACS1G, based on RNAi, could transform gynoecious into monoecious lines (Shiber et al., 2008). However, as these two genes share identical untranslated and coding regions, a functional analysis of each gene was impossible to achieve through RNAi. Finally, the other genes in the F locus, such as ACS1, MYB1a, and MYB1b, as well as BCAT, are tightly linked with ACS1G, and hence, their segregation from ACS1G was not possible. With the advent of CRISPR/Cas9 gene editing technology, we were able to use this method, in combination with an EMS mutant library, to identify mutations that allowed us to test, directly, whether ACS1 or ACS1G functions as the F gene.

Identification of ACS1G as the F gene revealed an interesting situation in that all players in the FAM model are different members of the ACS gene family. Interestingly, previous efforts to discover the identity of the FAM genes uncovered two players in the regulation of female flowers, namely ACO2 and WIP1 (Martin et al., 2009; Chen et al., 2016; Hu et al., 2017). Here, ACO2 is a key enzyme in the ethylene biosynthesis pathway, while WIP1 is repressed by ACS11 (A gene), which relieves its repression of ACS2 (M gene; Boualem et al., 2008, 2015). This raises the
rhetorical question of why might the ethylene synthesis genes have been selected to play a pivotal role in female flower development? Furthermore, it is intriguing to ponder what series of events led to the involvement of multiple ACS genes, and what might control the interaction between these genes? Answers to these questions can be sought through interrogation of the extensive cucurbit genomics databases, which includes genetic diversity from undomesticated and landrace accessions to commercial cultivars.

Differences between the Female (F) and gynoecy (g) genes
Although both the F and g genes promote female flower formation, there are important differences. The F gene (ACS1G) is dominant, and it has been discovered only in cucumber, but not in other cucurbit species. However, the g gene (dysfunctional WIP1) is recessive, and its function is conserved in melon, cucumber, and watermelon (Martin et al., 2009; Hu et al., 2017; Zhang et al., 2020). In addition, the wip1 mutants bear some bisexual flowers, at the lower nodes, but cucumbers carrying the F locus produce no bisexual flowers.

To further an understanding, the genetic relationship between F and WIP1, we obtained F_ wip1 seeds (where F_ can be FF or Ff) from an F2 population derived from a cross between CU2-FF and our wip1 mutants (CRISPR/Cas9 edited). As with cucumbers containing the F locus, these F_ wip1 plants produced female, but no bisexual flowers (Supplemental Figure 10). This established that the ethylene, produced by ACS1G and ACO2, could activate ACS2 expression, not only by repressing WIP1 expression, but also through another WIP1-independent pathway (Supplemental Figure 13D). Of note here, in Cucurbita pepo, females can be produced when WIP1 expression is not downregulated (Garcia et al., 2020), which is consistent with the notion that other pathways must exist to overcome the repression of CpACS27 (ortholog of ACS2 and ACS7) by WIP1. Identification of the factors that participate in such pathway(s) will be the focus of future studies.

As we mentioned above, the F gene is epistatic to the a gene, which means that F acs11 plants are gynoecious (Kubicki, 1969a, 1969c). In the present study, we demonstrated that F aco2/ac02 is androecious. To further our understanding of the differences between the two female promoting genes, ACS1G and wip1, we obtained the wip1 acs11 (from an F2 population between wip1 and Erez) and wip1 aco2 (from an F2 population between wip1 and 406a) double mutants. Similar to the findings in melon, these wip1

Figure 7 Expression levels of WIP1 and ACS2 in the apices of cucumber isogenic lines. (A) WIP1 expression in gynoecious (Gy) and monoecious (Mo) isogenic lines. Mo, monoecious; Gy, gynoecious. "YB" and "09" are two different genetic backgrounds. Data are presented as mean ± ss, n = 3. *p < 0.05 and **p < 0.01 in Student’s t-test compared to the isogenic monoecious lines (Supplemental Data Set 1). (B) WIP1 expression in shoot apices of monoecious lines. Con, control, treated with water; ETH, treated with ethephon. Data are presented as mean ± ss, n = 3. *p < 0.05 and **p < 0.01 in Student’s t-test compared to the control (Supplemental Data Set 1). (C) ACS2 expression in shoot apices of monoecious CU2 (WIP1WIP1 genotype) and gynoecious CU2 (wip1wip1 genotype) lines. Data are presented as mean ± ss, n = 4. *p < 0.05 and **p < 0.01 in Student’s t-test compared to CU2 (WIP1WIP1) genotype (Supplemental Data Set 1). (D) Model for regulating gynoecy and monoecy in cucumber.
acs11 double mutants produced both bisexual and female flowers, like the wip1 single mutant (Supplemental Figure 11). These findings indicate that, in cucumber, WIP1 also functions downstream of ACS11. However, the wip1 aco2 double mutants produced bisexual, but not female, flowers (Supplemental Figure 12, A–D). Furthermore, RT-qPCR assays showed that ACS2 was not upregulated significantly in these wip1 aco2 double mutant plants (Supplemental Figure 12E). Hence, the downregulation of WIP1, or dysfunctional WIP1, is not sufficient to activate ACS2 expression, and functional ACO2 is necessary for activating ACS2 expression.

The need for ACO2 to activate ACS2 expression and ensure female flower formation in the wip1 aco2 double mutant is difficult to reconcile with the current model. If ACO2 cooperates with ACS11 in repressing WIP1 and upregulating ACS2, then the wip1 aco2 and wip1 acs11 plants should have similar phenotypes. Thus, another ACS could cooperate with ACO2 to release the ethylene in these wip1 acs11 double mutants, but this ACS gene would be repressed by WIP1 in monoeocious cucumber. Whether ACS2 is a candidate, or if another ACS could perform this function remains to be determined.

New model of unisexual flower development in cucumber
The question as to which specific ethylene synthesis genes interact in flower development is addressed by our current spatiotemporal examination of the expression patterns of ACS1G, ACS1, and ACO2 (Figure 5), when assessed together with known ACS2, ACS11, as well as WIP1, expression patterns (Saito et al., 2007; Boualem et al., 2008, 2015; Li et al., 2009; Martin et al., 2009). Previous evidence indicated that, during carpel development, ACS2 functions as a basal player, regardless of whether a floral bud is male or female (Chen et al., 2016). In addition, for lines lacking the F locus, after floral buds have developed to stage 4, when ACO2 is expressed along the receptacle (Supplemental Figure 13A), the ACS11 (A gene) is expressed in some floral buds, which then represses WIP1 expression, thereby activating ACS2 (M gene) expression (Martin et al., 2009; Boualem et al., 2015). In this situation, these floral buds are developmentally programmed to become female. However, in those floral buds in which ACS11 is not being expressed, the male developmental program is active. In this way, the presence or absence of ACS11 activity leads to the monoeocious phenotype, in that both female and male flowers are produced on the same plant (Figure 7D).

An elimination of ACS11 function, as in the aa genotype (Boualem et al., 2015), or of ACO2 function (Chen et al., 2016), will cause WIP1 repression to be released, and as a consequence, ACS2 (M gene) expression is repressed, thereby giving an insufficient ethylene level to activate carpel development; the result is male floral bud development (Supplemental Figure 13B). In contrast to the regulatory roles of ACS11 (A gene) and WIP1, the ACS2 (M gene) appears to be the last player in the FAM regulatory system. In any event, with functional ACS2 expression, regardless of the reason, there will be sufficient ethylene for carpel development, and so the floral bud will develop as female. In the situation in which ACS2 is lacking, carpel development occurs only when both ACO2 and ACS11 provide marginal, but sufficient ethylene, to meet the requirement for carpel development: this level of ethylene does not arrest stamen development, and thus, the flowers are bisexual (Supplemental Figure 13C). In this regard, WIP1 and ACS11 may be coopted, through a regulatory complex, to regulate or stabilize ACS2 expression, a basic requirement for carpel development.

The F gene (ACS1G) functions uniquely in that it is expressed only in lines carrying the F locus, and it is expressed as early as ACO2, at bud stage 2 (Figures 5 and 7D; Supplemental Figure 13D). In the homozygous background, ACS1G expression in the floral buds gives rise to female flowers, regardless of the existence of ACS11 (A gene). These findings indicate that ACS1G can bypass the need for ACS11 and work directly with ACO2 to upregulate ACS2 and thereby ensure carpel development (Figure 7D); in this way, ACS1G exhibits dominance over ACS11. Here it should be mentioned that, at times, the genetic data offer only a potential regulatory framework, and thus, biochemical experiments may be needed to provide the detailed relations between the involved components, or elucidate other possibilities for the underlying gene interactions.

Identification of additional factors controlling expression patterns of ACS genes is needed for an understanding of unisexual flower development
The dominance of ACS1G over other ACS genes implies that these different members might be co-opted, in a stepwise manner, dependent upon the requirement for carpel development; here, the robustness of the rescuing mechanism might well contribute to its selection. This observed ACS1G dominance may reflect its duplication origin, and is consistent with the hypothesis that it is the newest integration of an ACS gene family member into the regulatory complex for cucumber carpel development (Sun et al., 2010).

Currently, we know that, in monoeocious cucumber, selective expression of ACS11 triggers female flower development, whereas in gynoecious cucumber, ACS1G dominates in female flower development. Irrespective of whether the cucumber line is monoeocious or gynoecious, upregulation of ACS2 is essential for female flower development. Clearly, gaining an understanding of the regulatory system that controls ACS11, ACS1G, and ACS2 expression would advance our understanding of unisexual flower development. Except for expression in the carpel region, ACS11 is also expressed in the underlying phloem of the female floral bud. This raises the question as to whether there might be factors, delivered via the phloem, which could serve to integrate both physiological and environmental inputs to impact the local
decision on floral bud development, as to whether to produce male or female flowers.

ACS1G is expressed at a very early stage in the carpel of every floral bud. The mechanism by which this new gene, ACS1G, acquired this novel function, within an already existing transcriptional regulatory network, needs to be elucidated in future studies. Finally, although WIP1 is a repressor of ACS2, it seems there are still other pathways and factors that can regulate ACS2 expression. Identification of such additional factors that can regulate ACS expression patterns, thereby leading to unisexual flowers development, should advance our understanding of the mechanisms regulating unisexual flower development in the cucurbits.

Materials and methods

Plant materials and growth conditions
Seeds of the three pairs of isogenic cucumber (Cucumis sativus L) inbred lines, CU2-Gy (Gynoecious, FFMM) and CU2-Mo (Monoecious, ffMM); YB-Gy (Gynoecious, FFMM) and YB-Mo (Monoecious, ffMM); 09-Gy (Gynoecious, FFMM) and 09-Mo (Monoecious, ffMM) were obtained from Hunan Xingshu Seed Industry Co., Ltd. (Chang Sha, Hunan, China). EMS-induced mutants were based on the YB and 09 backgrounds. Requests for these EMS-induced mutants require a material transfer agreement with Hunan Xingshu Seed Industry. If the proprietary inbred line is requested, then Hunan Xingshu Seed Industry will provide a hybrid, derived from the requested inbred, at its discretion. Seeds of the inbred line CU2-Mo were used for transformations. The CU2-Gy lines were crossed with the transgenic lines with a backbred line CU2-Mo were used for transformations. The CU2-Mo (Monoecious, ffMM) and YB-Mo (Monoecious, ffMM); YB-Gy (Gynoecious, FFMM) and 09-Gy (Gynoecious, FFMM) and 09-Mo (Monoecious, ffMM) were obtained from Hunan Xingshu Seed Industry Co., Ltd. (Chang Sha, Hunan, China). EMS-induced mutants were based on the YB and 09 backgrounds. Requests for these EMS-induced mutants require a material transfer agreement with Hunan Xingshu Seed Industry. If the proprietary inbred line is requested, then Hunan Xingshu Seed Industry will provide a hybrid, derived from the requested inbred, at its discretion. Seeds of the inbred line CU2 were peeled and sterilized, then germinated on Murashige and Skoog medium containing 2 mg/L 6-benzylaminopurine (BA) and 1 mg/L abscisic acid (ABA) for 2 days in the dark at 28°C. Explants were prepared by removing the germ and transverse cutting of the cotyledon. The verified vector was introduced into the A. tumefaciens strain EHA105 for cucumber transformation. Primer sequences are listed in Supplemental Table 1.

Cucumber transformation
Cucumber transformation was performed as previously described (Hu et al., 2017). Briefly, the seeds of the inbred line CU2 were peeled and sterilized, then germinated on Murashige and Skoog medium containing 2 mg/L 6-benzylaminopurine (BA) and 1 mg/L abscisic acid (ABA) for 2 days in the dark at 28°C. Explants were prepared by removing the germ and transverse cutting of the cotyledon. The verified vector was introduced into the A. tumefaciens strain EHA105, and transformed clones were incubated, overnight, in Luria-Bertani medium containing 50 mg/L kanamycin and 25 mg/L rifampicin, at 28°C, then incubated in 30 mL LB medium until the culture OD_{600} (the optical density of a sample measured at a wavelength of 600 nm) reached 0.4–0.8. The cell culture was then collected by centrifugation, washed with sterilized water and transferred to the shoot regeneration medium (2 mg/L BA, 1 mg/L ABA, 200 μM acetosyringone, and 10 mM MES [pH 5.2]) to an OD_{600} of 0.2. The explants were then immersed in 10 mL inoculation medium, contained within a 20-mL syringe, and vacuum infiltration was performed by pulling up on the plunger. After the infection, explants were co-cultured with the Agrobacterium on three layers of damp filter papers for ~72 h, in the dark, then washed with sterilized water and transferred to the shoot regeneration medium (2 mg/L BA, 1 mg/L ABA, 100 mg/L kanamycin, and 200 mg/L timentin). Shoots appeared in 3–4 weeks, and the transformed buds were selected by screening for GFP (Green Fluorescent Protein) fluorescence. These transgenic buds were excised from the explants and planted onto elongation MS medium (1.0 mg/L gibberellic acid, 0.1 mg/L BA, 0.01 mg/L 1-naphthaleneacetic acid, 2 mg/L AgNO₃, and 100 mg/L timentin), and cultured at 25°C until they reached ~4 cm, at which time shoots were transferred onto rooting medium (6.5 mg/
L hemin and 100 mg/L timentin). Successfully rooted seedlings were cultured for 5 days under low light and then transplanted to soil. Transgenic constructs were verified using PCR and by monitoring GFP fluorescence at later growth stages.

EMS mutagenesis and targeting induced local lesions in genomes (TILLING)

EMS mutagenesis and a targeting induced local lesions in genomes (TILLING) screen were performed, as previously described (Tadmor et al., 2007), with some modifications to meet the needs of isolating mutations in the region of the F locus. Since the gynoecious lines with the FF background rarely form male flowers, and were particularly susceptible to the effects of EMS, seeds with the 09-Ff and YB-Ff backgrounds, from the crosses between 09-Gy and 09-Mo, and YB-Gy and YB-Mo, were used for the EMS mutagenesis, thereby allowing the use for self-pollination of male flowers that developed at the lower nodes. This circumvented the need for the AgNO3 treatment to induce male flowers, generated from the self-pollination of these M1 plants. For each M1 plant containing a mutation in the ACS1G locus, the offsprings (M2 plants) were genotyped to select ACS1G H164Y (which is known to severely aggravate the growth and inhibition of fruit development) (Tadmor et al., 2007), with some modifications to the protocol. Instead, a TILLING screen was applied to the fertile M1 plants. For each M1 plant containing a mutation in the ACS1G locus, its genomic DNA was extracted. Primers covering the genomic duplicated region, the ratio of point mutations was sequenced to verify the mutation type; each M2 individual was then analyzed to determine its homo- or heterozygosity for this mutation.

The leaf closest to the first fruit in each M1 plant was collected and its genomic DNA was extracted. Primers covering the entire coding region of ACS1/ACS1G (Supplemental Table 2) were used for PCR amplification. After denaturation and annealing, the PCR products were digested with CIL I (Colbert et al., 2001) and analyzed using polyacrylamide gel electrophoresis. The M2 population, generated from the M1 plants containing mutations in ACS1/ACS1G, was used to assess whether the mutations were linked to the F locus. The specific primers used to identify the F locus are listed in Supplemental Table 2. Mutations not linked to the F locus were considered to be located in ACS1f. The possibility that the mutations in ACS1f or ACS1G were linked to the F locus was investigated by sequencing the amplification products with primers specific to the distal promoters of ACS1F and ACS1G (Supplemental Table 2). The homozygous and heterozygous alleles were estimated from the sequencing chromatograms, and then verified using the derived cleaved amplified polymorphic sequence (dCAPS) method (Neff et al., 1998). Primers for dCAPS are listed in Supplemental Table 2.

The M2 plants containing ACS1G H164Y and ACS1f E353K mutations were backcrossed, individually, with 09-Mo (ff) and YB-Gy (FF). The offsprings (M3 plants) were genotyped to select ACS1G H164Y (ff) and ACS1f E353K (ff) lines for further analysis; the segregation ratio was determined using this generation.

Expression and purification of recombinant ACS1 isoforms

Coding sequences for the wild-type ACS1, ACS1G H164Y, and ACS1f E353K isoforms were introduced into the pET30a (Merck Millipore, Burlington, MA, USA) plasmid. The constructs were verified by sequencing and then individually transformed into the BL21 (DE3) Escherichia coli strain. Transformants were placed in LB medium supplemented with kanamycin (50 mg/L) and incubated overnight at 37°C, with shaking at 200 r.p.m. The pre-culture cells were incubated in 500 mL LB medium containing 50 mg/L kanamycin and were shaken at 37°C until the culture had an OD600 of 0.6. Isopropyl β-D-thiogalactopyranoside was then added to a final concentration of 0.05 mM. After shaking the culture at 180 r.p.m. for ~16 h, at 16°C, cells were harvested by centrifugation and resuspended in 50 mL lysis buffer (0.3 M NaCl, 50 mM KH2PO4, and 10 mM imidazole [pH 8.0]) supplemented with 1 mg/mL lysozyme (Amresco, Solon, OH, USA; 0663) and were incubated on ice for 4 h before being disrupted by sonication on ice. The supernatant was collected after centrifugation at 12,000 rpm at 4°C for 15 min, then 2 mL of a 50% slurry of Ni-NTA Agarose (Qiagen, Hilden, Germany; 30210), equilibrated with lysis buffer, was added to each sample. The mixture was incubated on ice for 30 min with gentle shaking and then placed on an Econo-Pac Chromatography Column (Bio-Rad Laboratories, Hercules, CA, USA; 7321010). Unbound protein was removed by washing the Ni-NTA agarose, three times, with a column volume of wash buffer (0.3 M NaCl, 50 mM KH2PO4, and 20 mM imidazole [pH 8.0]), then 2-mL elution buffer (0.3 M NaCl, 50 mM KH2PO4, and 250 mM imidazole [pH 8.0]) was applied to elute the His-fusion protein. Next, centrifugal filter devices (Merck Millipore; UFC201024) were used to replace the elution buffer with 1 × Phosphate buffer saline (PBS) buffer (0.14 M NaCl, 2.7 mM KCl, 10 mM Na2HPO4, and 1.8 mM KH2PO4). Purified protein was quantified using a Pierce BCA Protein Assay Kit (Thermo Fisher Scientific; 23227), then aliquoted, stored at −80°C, and used for the enzyme activity assay. Primers used for vector construction are listed in Supplemental Table 3.

ACS enzyme activity assay

The ACS activity assays were performed, as previously described. This method measures ethylene production, thereby indicating the ACS activity within the tested tissues. The reaction solution, containing 200 mM tricine (pH 8.0), 3.5 μM pyrodoxal-L-phosphate, 10 mM dithiothreitol, and 1.2 mM S-(5'-adenosyl)-L-methionine chloride, was freshly prepared. The reaction was initiated by adding 3 μg of purified recombinant enzyme to 1.6 mL reaction solution and incubated for 2 h at 25°C with gentle shaking. The reaction was terminated with 200 μL 100 mM HgCl2, then 950 μL of the reacted mixture and 850 μL distilled water were combined in a 10-mL glass vial, which was sealed with a cap before...
being injected with 0.2 mL NaOH–NaOCl mixture (freshly prepared by mixing two units of 5% NaOCl and one unit of 6 M NaOH on ice). The sample was vortexed for 5 s and placed on ice for 4 min to react, before being vortexed for 5 s again to release the ethylene into the headspace of the glass vial. A 250-μL aliquot of gas was withdrawn from the headspace, using a syringe, and its ethylene content quantified using an Agilent 6890N gas chromatograph (Agilent Technologies, Santa Clara, CA, USA) fitted with a flame-ionization detector. Results shown are the mean ± SEM (n = 3 replicates).

In situ hybridization assay
Sense and antisense RNA probes were transcribed, in vitro, using a digoxigenin RNA labeling kit (Roche, Basel, Switzerland; #11750259101), with the ACS1-specific region as a template. Primers used to amplify the template are listed in Supplemental Table 1. Tissue sections were prepared, as previously described (Schneitz et al., 1998), with minor modifications. Briefly, shoot tips were excised and fixed, over-night; four treatments of TBST with gentle shaking, 15 min; RNase A in 1× NTE at 37°C, 5 min; RNase A in 1× NTE at 37°C, 30 min; two treatments of 1× NTE at 37°C, 5 min; 2× SSC at 45°C, 20 min; 1× Tris-buffered saline, 5 min; 1% blocking reagent, 45 min; 1% bovine serum albumin in Tris-buffered saline with Tween-20 (TBST), 45 min; 1:1,000 anti-DIG-antibody at 4°C, overnight; four treatments of TBST with gentle shaking, 15 min; detection buffer (100 mM Tris–HCl, 100 mM NaCl [pH 9.5]), 5 min; Nitroblue tetrazolium chloride/5-bromo-4-chloro-3-indolyl-phosphate (NBT/BCIP) solution, for several hours. When the signal was strong enough, the reaction was terminated using 1× TE (Tris–EDTA buffer). The sections were then transferred onto slides, and images were taken using a digital camera mounted on a Leica DM5500B microscope.

Construction of ProACS1G:GUS and ProACS1f:GUS fusion genes
For the ProACS1G:GUS (β-glucuronidase) reporter construct, DNA fragments covering the upstream 2 kb of the start codon of ACS1G were amplified from the genomic DNA of the gynoecous cucumber, CU2-Gy. For the ProACS1f:GUS reporter construct, DNA fragments covering the upstream 2 kb of the ACS1f start codon were amplified from the genomic DNA of the monoecious cucumber, CU2-Mo. The Sal I and Xba I sites were inserted into the end of the two forward primers and the end of the same reverse primer, respectively. Then, the promoters of ACS1G and ACS1f were cloned into a binary vector, pCAMBIA1305.4, before the GUS coding region. Each fusion construct was confirmed by DNA sequencing. Sequences for the primers are listed in Supplemental Table 4.

Histochemical GUS staining
Histochemical GUS staining of transgenic cucumber floral buds was performed, as previously described (Wang et al., 2014). Shoot tips containing the floral buds at were cut, using a scalpel, placed in the GUS staining solution, and then vacuum-infiltrated. After incubation at 37°C for 24 h, the GUS staining was terminated. Chlorophyll was removed by using ethanol and then methanol. Before imaging, tissues were cleared using chloral hydrate. The GUS-positive plant tissues were examined under a bright field microscope (Leica DM5500B, Cambridge, England).

Accession numbers
Sequences from this article can be found in the Cucurbit Genomics Database (www.icugi.org) under accession numbers CsaV3_6G044400 (ACS1), CsaV3_6G044410 (MYB1), CsaV3_6G044420 (BCAT), CsaV3_1G040170 (ACS2), CsaV3_2G025850 (ACS11), CsaV3_4G024150 (WIP1), CsaV3_6G048630 (AC02), CsaV3_6G041900 (ACTIN2).

Supplemental data
Supplemental Figure 1. Genotype categories of the CRISPR/Cas9-edited ff cucumber plants with dysfunctional MYB1 or ACS1/1G.

Supplemental Figure 2. Sanger sequence analysis of MYB1 in MYB1ffCR plants.

Supplemental Figure 3. Sanger sequence analysis of MYB1 in MYB1_ALLCR plants.

Supplemental Figure 4. Sanger sequence analysis of ACS1 in ACS1fCR plants.

Supplemental Figure 5. Sanger sequence analysis of ACS1/1G in ACS1_ALLCR plants.

Supplemental Figure 6. Genomic verification of the ACS1G<sup>HIS64T</sup> (ff) and ACS1f<sup>L353K</sup> (ff) mutants.
Supplemental Figure 7. In gynoecious lines, ACS1G transcripts are dominant in floral buds at the key stage of sex determination.

Supplemental Figure 8. In situ hybridization with sense probes of ACS1/G and ACO2.

Supplemental Figure 9. GUS staining pattern in flower buds of ProACS1G::GUS and ProACS1fGUS transgenic cucumbers.

Supplemental Figure 10. Flower sex type of F_ wip1 double mutants.

Supplemental Figure 11. Flower sex type of wip1 acs11 double mutants.

Supplemental Figure 12. Flower sex type of wip1 aco2 double mutants.

Supplemental Figure 13. Model for the regulation of unisexual flower development in cucumber.

Supplemental Table 1. RNA primers used in this study.

Supplemental Table 2. Genome editing, TILLING, and genotyping primers.

Supplemental Table 3. Protein expression primers.

Supplemental Table 4. Transgenic vector and genomic verification primers.

Supplemental Data Set 1. Data for all statistical analyses performed in this study.

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