

GhTCE1–GhTCEE1 dimers regulate transcriptional reprogramming during wound-induced callus formation in cotton

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Abstract

Wounded plant cells can form callus to seal the wound site. Alternatively, wounding can cause adventitious organogenesis or somatic embryogenesis. These distinct developmental pathways require specific cell fate decisions. Here, we identify GhTCE1, a basic helix–loop–helix family transcription factor, and its interacting partners as a central regulatory module of early cell fate transition during in vitro dedifferentiation of cotton (*Gossypium hirsutum*). RNAi- or CRISPR/Cas9-mediated loss of *GhTCE1* function resulted in excessive accumulation of reactive oxygen species (ROS), arrested callus cell elongation, and increased adventitious organogenesis. In contrast, *GhTCE1*-overexpressing tissues underwent callus cell growth, but organogenesis was repressed. Transcriptome analysis revealed that several pathways depend on proper regulation of *GhTCE1* expression, including lipid transfer pathway components, ROS homeostasis, and cell expansion. GhTCE1 bound to the promoters of the target genes *GhLTP2* and *GhLTP3*, activating their expression synergistically, and the heterodimer TCE1–TCEE1 enhances this activity. *GhLTP2*- and *GhLTP3*-deficient tissues accumulated ROS and had arrested callus cell elongation, which was restored by ROS scavengers. These results reveal a unique regulatory network involving ROS and lipid transfer proteins, which act as potential ROS scavengers. This network acts as a switch between unorganized callus growth and organized development during in vitro dedifferentiation of cotton cells.

Introduction

Developmental plasticity in response to environmental conditions is a key distinguishing feature of plants, and this is seen clearly in the response to mechanical wounding. Following wounding, differentiated somatic cells undergo

redifferentiation either to form wound callus that seal the wound site to prevent infection or to undergo de novo organogenesis, such as adventitious rooting or, during tissue culture, somatic embryogenesis (SE) (e.g. [Chen et al., 2014](#);

Yu et al., 2017). A major question is how decisions are made at the molecular level to determine the choice of alternative pathways of either disorganized growth (callus formation) or organized growth, such as de novo meristem formation and SE.

External stimuli such as wounding are critical for callus initiation via cell division, and exogenous hormones are necessary for the propagation of callus in vitro and for the experimental induction of SE (Ikeuchi et al., 2017; Xu et al., 2018). Various environmental stresses such as drought, salt, and reactive oxygen species (ROS) can accelerate adventitious organogenesis or SE by repressing nonembryonic callus proliferation and promoting the transition from nonembryonic callus to callus with morphogenic potential. This can lead to the development of somatic embryos or root or shoot meristems (Jin et al., 2013; Fehér, 2015; Altpeter et al., 2016).

Adventitious organogenesis is under finely tuned regulation, whereby somatic cells re-enter the cell cycle and exhibit pluripotency via chromatin reprogramming (Lee and Seo, 2018). Callus induction is a prerequisite for this, and the callus induction rate, callus cell morphology, and the rate of callus proliferation all influence regeneration efficiency (Li et al., 2013; Zhang et al., 2019). Auxin and cytokinin signaling pathways are essential during organized development from callus (Yang et al., 2012; Liu et al., 2016). Gene expression related to stress responses, control of cell identity, and chromatin remodeling is also required (Ikeuchi et al., 2019). Several genes that function during SE and other types of organ regeneration have been identified via forward and reverse genetic approaches in *Arabidopsis thaliana* (Arabidopsis, Ikeuchi et al., 2017, 2019). More than half of these genes encode transcription factors (TFs), and given that TFs account for less than 6% of total protein-coding genes (Riechmann, 2002), it is likely that TF networks are of central importance as regulators during de novo organogenesis. To date, thousands of genes have been found to be expressed during SE or organogenesis (Yang et al., 2012; Du et al., 2019), but only a small number of them have been verified experimentally for function. Furthermore, callus has typically been studied as a single tissue even though it is heterogeneous in terms of cell morphology, metabolism, and developmental potential (Lindsey and Yeoman, 1983).

In a previous study in cotton (*Gossypium hirsutum*), we found that dedifferentiated callus cells are very different morphologically from highly organized somatic cells, and 466 TFs were identified as differentially expressed genes (DEGs) at different stages of SE (Yang et al., 2012). In the current study, we focused on the cell fate transition at an early stage of callus formation during in vitro dedifferentiation, and found that a basic helix–loop–helix (bHLH) family TF, GhTCE1, specifically promotes callus cell elongation by activating interacting lipid transfer and ROS homeostasis pathway genes and repressing adventitious organogenesis. Our results provide an important new role for lipid transfer

proteins and ROS signaling in regulating early cell fate transitions following wound-induced cell division.

Results

GhTCE1 promotes callus cell elongation during dedifferentiation of cotton cells

GhTCE1 (Gh_D06G2316) was identified during an RNA-seq profiling analysis during cotton SE (Yang et al., 2012), and was chosen for further study because, as a potential TF, it might play a role in early events during callus formation or SE. The 996-bp cDNA encodes a predicted bHLH family TF of 331 amino acid residues. The expression pattern of GhTCE1 was determined in cotton root, stem, leaf, anther, and at different time points during SE from cultured hypocotyl explants, and at earlier time points after explant wounding. GhTCE1 was expressed at low levels in vegetative and reproductive organs, but much more strongly during different stages of SE (Figure 1A). Moreover, it was induced at earlier time points after wounding, with the increased expression being detectable within 60 min, and very strong by 720 min, after wounding (Figure 1B), suggesting a role in the reprogramming process.

Gh_D06G2316 and Gh_A06G0241 are paralogs on the D and A subgenomes of tetraploid cotton, sharing 98.8% sequence identity (Supplemental Figure S1). In order to explore the function of GhTCE1, we created RNAi (Ri) lines, overexpression (OE) lines, and CRISPR-mediated GhTCE1D/A (Gh_D06G2316/Gh_A06G0241) double mutants. Two Ri lines (Ri1 and Ri2), two OE lines (OE1 and OE3), a wild-type line (WT), as well as an Ri-negative line (Null) that segregated from Ri1 in the T₁ generation, were selected for detailed analysis. Immunoblotting analyses revealed that each of these Ri and OE transgenic lines contain a single-copy T-DNA insertion (Figure 1C). The expression levels of GhTCE1 decreased to 24.0% of WT levels in Ri1 and to 21.6% in Ri2, and increased in the OE lines by 833.8% (OE1) and 1036.2% (OE3) compared with WT (Figure 1D). In addition, we created two double mutant lines *ko15* and *ko18* using CRISPR/Cas9-mediated gene editing. These were confirmed by high-throughput sequencing as frame-shift mutations and were selected for further study (Figure 1E). OE and Ri mature plants were morphologically similar to WT (Figure 1F).

Wounded explants from different transgenic lines were cultured on MSB (MS medium plus B5 vitamins) medium according to our previous methods (Yang et al., 2012). At 20 days post wounding, significantly more adventitious roots were produced and a higher ratio of adventitious root regeneration was observed on Ri and mutant explants compared with WT plants. In contrast, the OE lines showed callus growth but no adventitious organogenesis (Figure 2A and Supplemental Figure S2). Both the callus proliferation rate (CPR), which is a measure of the fresh weight of callus tissue produced over time, and callus cell length were measured at 20 days post wounding. The CPR of lines Ri1 and Ri2 was reduced to 45.3% and 56.9%, respectively, compared with WT, and was even lower in *ko15* (24.5%) and *ko18*

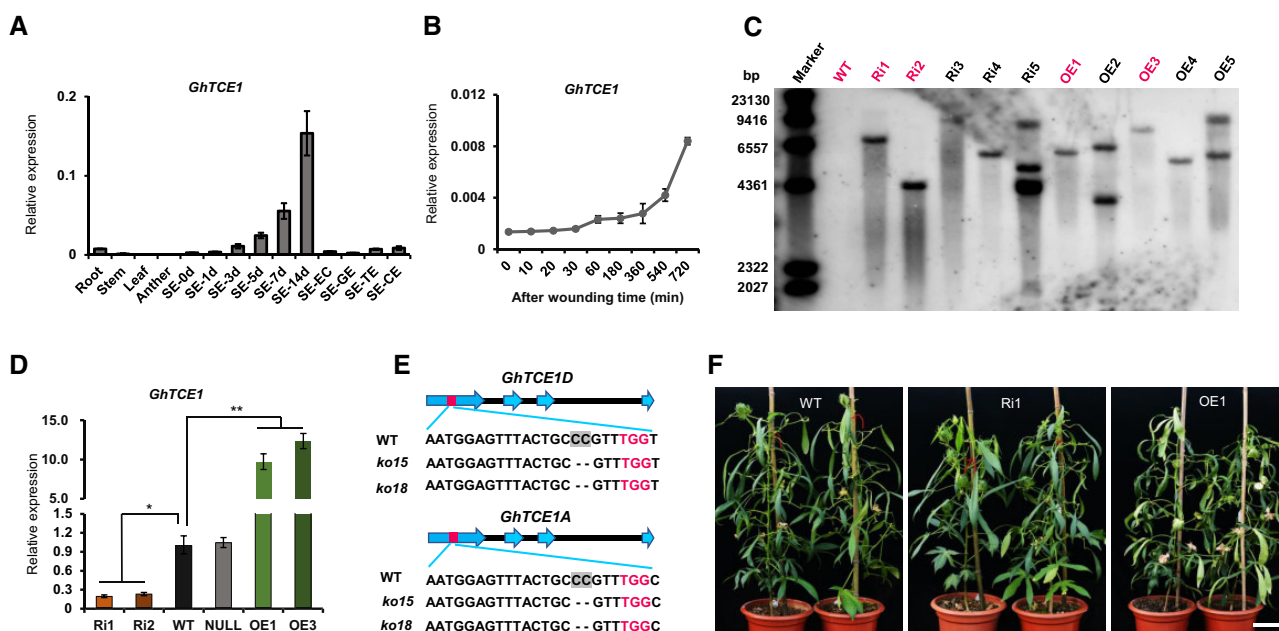


Figure 1 Analysis of *GhTCE1* expression and characterization of transgenic plants. A, Expression pattern of *GhTCE1* in different tissues. SE-0d, -1d, -3d, -5d, -7d, or -14d means the explants of hypocotyls were induced for 1, 3, 5, 7, or 14 days during SE (embryogenic callus [SE-EC], globular stage embryo [SE-GE], torpedo stage embryo [SE-TE], and cotyledon stage embryo [SE-CE]). B, Expression pattern of *GhTCE1* soon after wounding. The expression of *GhUB7* was used as internal control. C, Immunoblot analysis of *GhTCE1* transgenic lines. The first lane represents the λ DNA size marker. The DNA samples were digested with *HindIII*. The magenta front indicates the selected lines for study. D, Relative expression of *GhTCE1* in transgenic lines 7 days post SE induction. WT, wild type; Ri1 and Ri2, Ri lines; OE1 and OE3, OE lines. Null, negative plants segregated from Ri1 in the T_1 generation. The expression of *GhUB7* was used as internal control. E, Sequence analysis of *GhTCE1(A/D)* double mutants *ko15* and *ko18* at the target sites. The PAM motif is marked in magenta and *GhTCE1A* and *GhTCE1D* can be distinguished by the SNP C/T (marked in green) downstream of the PAM motif. The black dotted line represents deletion compared with WT. F, Phenotypes of *GhTCE1* transgenic plant lines grown on soil. Bars = 20 cm. Error bars in (A), (B), and (D) represent \pm standard error of three biological replicates (~ 20 explants were used as one replicate). Significance tests compared each transgenic line with WT plants. * $P < 0.05$, ** $P < 0.01$, Supplemental Data Set S2.

(30.7%) mutants (Figure 2B). However, the CPR increased 173.1% in OE1 and 182.9% in OE3, respectively, compared with WT (Figure 2B). Callus cell length was correspondingly reduced in loss-of-function lines compared with WT to 46.3% (Ri1), 37.9% (Ri2), 42.3% (*ko15*), and 51.8% (*ko18*), and it increased to 144% in OE1 and 135% in OE3 (Figure 2, C and D). These results show that *GhTCE1* promotes callus cell growth and fresh weight (principally a reflection of increased cell length) but represses adventitious organogenesis.

GhTCE1 modulates ROS but not auxin homeostasis during cotton cell dedifferentiation

ROS and auxin play essential roles in cell fate determination and growth (Chapman et al., 2019). To examine the possible role for ROS, we monitored its accumulation by 2,7-dichloro-dihydrofluorescein diacetate (DCFH-DA) staining. DCFH-DA can permeate cell membranes and then be oxidized by ROS into impermeant DCF, which is fluorescent under UV light. The *GhTCE1*-Ri lines and mutant lines showed strong green fluorescence after DCFH-DA staining, but none was seen in WT or OE lines (Figure 2E). H_2O_2 concentrations were assayed and found to be significantly higher in Ri lines than in WT, and were significantly lower in OE lines than WT (Figure 2F). This demonstrates that

GhTCE1 is a regulator of ROS accumulation, and that high levels of *GhTCE1* expression are associated with reduced ROS accumulation, increased callus cell growth, and reduced adventitious organogenesis.

To determine the possible role of auxin in *GhTCE1*-mediated cell reprogramming, we measured IAA concentrations in the different lines at 20 days post wounding. Whereas one Ri line and one OE line had altered IAA levels, the IAA concentrations in all other lines were comparable, and on average similar to levels in WT (Supplemental Figure S3A). We also treated the explants with a synthetic auxin analog indole-3-butyric acid (IBA) to explore whether auxin treatment can mimic the phenotypes of wounded transgenic tissues. Whereas exogenous auxin promoted cell dedifferentiation and callus proliferation at 10 days post wounding, it promoted adventitious root formation in a concentration-dependent manner, and inhibited callus cell proliferation, at 20 days post wounding (Supplemental Figure S3B). Therefore, exogenous auxin treatments phenocopied the OE-like phenotype at 10 days and the Ri-like phenotype at 20 days post wounding (Supplemental Figure S3B). This suggests that auxin may function during organogenesis, but it is not likely to be the cause of differential cell proliferation between *GhTCE1* transgenic lines.

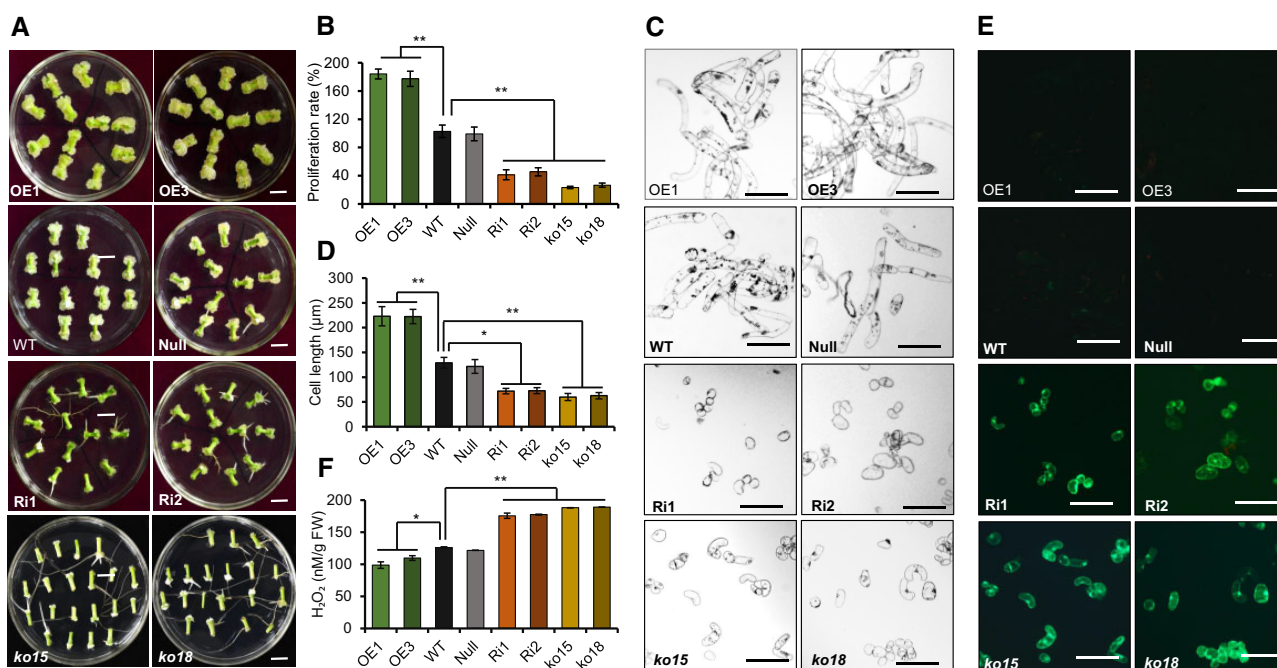


Figure 2 Phenotypic analysis of *GhTCE1* transgenic tissues. **A**, The phenotypes of callus 20 days post wounding from different transgenic lines and mutants. *ko15* and *ko18*, mutation lines. **B**, CPR of different lines 20 days post wounding. Error bars represent \pm standard error of six biological replicates (~ 20 explants each). **C**, Micrographs of callus cells of different lines 20 days post wounding. Bars = 200 μ m. **D**, Statistical analysis of callus cells length of different lines 20 days post wounding. Error bars represent \pm standard error of six biological replicates (~ 50 cells each). **E**, DCFH-DA staining of ROS in *GhTCE1* transgenic plants and mutants. **F**, H₂O₂ concentration in callus from *GhTCE1* transgenic plants 20 days post wounding. Error bars represent \pm standard errors of four biological replicates (~ 20 explants each). Significance tests compared each transgenic line with WT plants. * $P < 0.05$, ** $P < 0.01$, [Supplemental Data Set S2](#).

GhTCE1 targets and activates the expression of cell elongation-related genes during cotton cell dedifferentiation

To explore the underlying mechanism by which GhTCE1 promotes disorganized cell growth, we conducted a transcriptomics analysis using the explants of the different transgenic lines at 7 days post wounding. Differences in cell dedifferentiation among WT, Ri, and OE were first apparent at this stage ([Supplemental Figure S4](#)). DEGs were defined based on a false discovery rate (FDR) of ≤ 0.05 and fold change of $|\log_2| \geq 1$. We identified 331 up-regulated and 168 down-regulated DEGs in OE, and 63 up-regulated and 166 down-regulated DEGs in Ri compared with WT; 62 DEGs overlapped between OE-up and Ri-down DEGs, and included *GhTCE1* ([Figure 3A](#)). GhTCE1 is expected to exhibit transcriptional activation activity, and so the remaining 61 DEGs identified were considered as potential GhTCE1 targets. Their expression levels as fragments per kilobase per million mapped (FPKM) are presented in [Supplemental Data Set S1](#).

Nearly half of the candidates (29 of 61) have annotated functions in sterol synthesis, cell wall modification, cell expansion, ROS homeostasis, and lipid transfer ([Figure 3B](#) and [Supplemental Data Set S1](#)). These functions are all potentially related to cell elongation, and were considered as candidate targets of GhTCE1. The expression levels of the 29 candidate targets were verified by qRT-PCR analysis, and all

were upregulated in both OE1 and OE3, and 28 were down-regulated in both Ri1 and Ri2 ([Figure 3C](#)). The expression levels from RNA-seq highly correlated with qRT-PCR results ([Supplemental Figure S5](#), $R^2 = 0.9258$). The remaining 32 DEGs are involved in various processes such as cellular metabolism, signal transduction, or protein modification ([Supplemental Table S1](#)). It is notable that no differentially expressed auxin metabolism- or signal transduction-related genes were found.

GhTCE1 is annotated as a bHLH TF and subcellular localization assays show that it is localized to the nucleus ([Supplemental Figure S6A](#)). bHLH family TFs can bind the E-box (5'-CANNTG-3') or G-box (5'-CACGTG-3') of target gene promoters, depending on sequences in the basic region ([Toledo-Ortiz et al., 2003](#)), and sequence analysis suggests that GhTCE1 belongs to the E-box binding sub-family. To confirm the transcriptional activity of GhTCE1, activation assays were carried out in yeast (*Saccharomyces cerevisiae*). The full length GhTCE1 coding sequence was fused to a GAL4 DNA binding domain (BD), designated Full:BD. The yeast strain Y2HGold harboring Full:BD can activate the GAL4 reporter gene ([Supplemental Figure S6B](#)), demonstrating that GhTCE1 is a transcriptional activator. To profile the region responsible for its transcriptional activation, we constructed a series of truncated GhTCE1:BD sequences in yeast Y2HGold and mated yeast strains containing them with yeast strain Y187 harboring the empty vector pGADT7

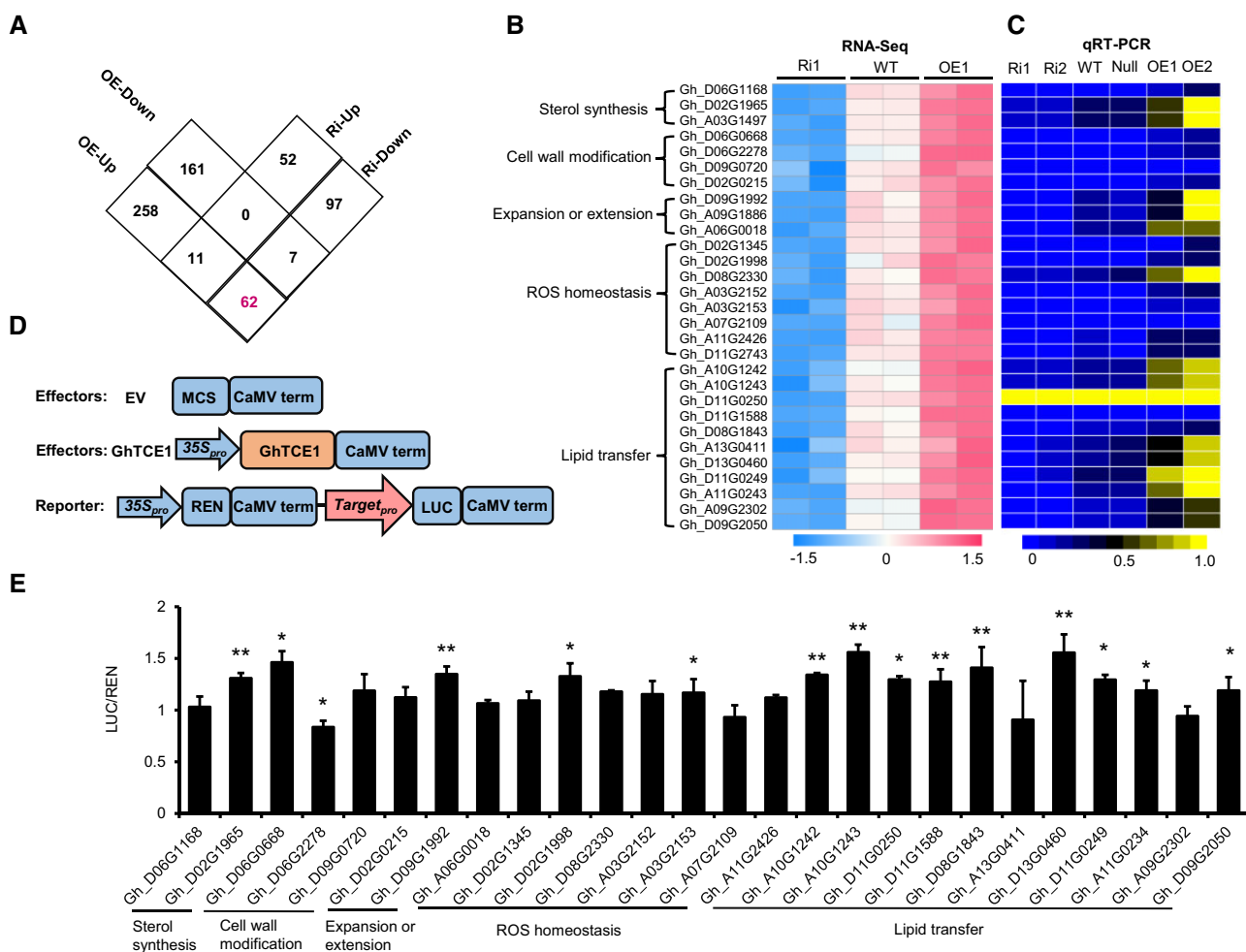


Figure 3 Analysis of downstream targets of *GhTCE1*. **A**, Differential gene (DEG) analysis in *GhTCE1* Ri and OE lines compared with WT. Screening for up or down regulated genes used a stringency of $P \leq 0.05$ and a \log_2 fold change of ≥ 1 or ≤ -1 compared with WT. Hypocotyls at 7 days post wounding were used for RNA-seq analyses, with two biological replicates (~ 50 explants each) for each line. The magenta front indicates 62 DEGs overlapped between OE-up and Ri-down. **B**, Heat-map analysis of 29 DEGs from enriched pathways that were up-regulated or down-regulated in OE and Ri lines, respectively. **C**, Heat-map analysis of the 29 DEGs from (B) using qRT-PCR, with three biological replicates for each line. **D**, Schematic diagram of effector and reporter constructs. Orange rectangle, the effector; pink arrow, the reporter; blue rectangles, other vector components; blue arrows, 35S promoters. The empty vector (EV) was used as a negative control. **E**, Transactivation analysis between *GhTCE1* and its candidate targets in cotton callus protoplasts using the LUC reporter system. Promoters from the candidate targets were fused to upstream of LUC in pGreenII 0800-LUC. 35S_{pro}:REN in pGreenII 0800-LUC was used as internal control. *GhTCE1* was fused downstream of the CaMV 35S promoter (35S_{pro}) in pGreenII 62-SK. pGreenII 62-SK empty vector was used as a negative control and designated EV. Relative activation level represents the ratio of fluorescence intensity of LUC/REN with *GhTCE1* to LUC/REN without *GhTCE1*. Error bars in (E) represent \pm standard errors of three biological replicates (10 samples each). Significance tests compared the expression of Promoter:LUC + 35S_{pro}:REN + 35S_{pro}:*GhTCE1* with Promoter:LUC + 35S_{pro}:REN + EV, $P < 0.05$ (*) or $P < 0.01$ (**), [Supplemental Data Set S2](#).

([Supplemental Figure S6C](#)). In diploid yeast, the N-terminal region (1–160 aa) can activate the reporter gene while the C-terminal region (161–331 aa) cannot ([Supplemental Figure S6D](#)). Sequences comprising either 80–240 aa or 80–331 aa can also activate the reporter gene, suggesting that the region from 80 to 160 aa is critical for *GhTCE1* transcriptional activation activity ([Supplemental Figure S6D](#)).

We downloaded 2-kb promoter region sequences (i.e. upstream of the translation start site) of 29 candidate target genes with potential roles in cell elongation from the cotton genome database (<https://cottonfgd.org/>) and then determined the presence and number of E-boxes. The number of

E-boxes ranged from 2 to 13 ([Supplemental Table S2](#)). We cloned 26 promoters from the 29 candidates for testing in transactivation studies, which used a luciferase (LUC) reporter activation system in tobacco (*Nicotiana benthamiana*) leaves ([Figure 3D](#)). Of the 26 candidate promoters, 14 were activated by *GhTCE1* to relatively low levels, less than two-fold above basal levels ([Figure 3E](#)). One possible explanation for this relatively low activation is that the full transcriptional activation of downstream targets may require protein partners which are absent in leaf.

As *GhLTPs* are the most abundant members among the candidate *GhTCE1* targets, and both *GhLTP2* and *GhLTP3*

are highly expressed during SE, we selected the promoters of *GhLTP2* and *GhLTP3* for transcriptional activation assays with both full length GhTCE1 and the N-terminal truncated GhTCE1 (Δ GhTCE1). The results show that GhTCE1 can interact with the promoters of both *GhLTP2* (*LTP2_{pro}*) and *GhLTP3* (*LTP3_{pro}*), promoting LUC expression in tobacco leaves. In contrast, the N-terminal truncated version of GhTCE1 did not activate *LTP2_{pro}* and *LTP3_{pro}* (Figure 4, A and B). Meanwhile, we analyzed the activated expression of *LTP2_{pro}* and *LTP3_{pro}* by GhTCE1 and Δ GhTCE1 in cotton callus protoplasts using the LUC expression system, and our results show that GhTCE1 can activate LUC expression, but Δ GhTCE1 was unable to activate LUC expression (Figure 4C). To confirm that GhTCE1 can interact with the promoters of *LTP2* and *LTP3*, we carried out yeast-one-hybrid (Y1H) experiments. There was no detectable interaction between GhTCE1 and the empty pGAD vector on SD/-Trp-Leu-His medium. However, in the presence of 3-amino-1,2,4-triazole (3-AT), GhTCE1 interacted with *LTP2_{pro}* and *LTP3_{pro}* (Figure 4D), confirming that GhTCE1 can bind to the promoters of *LTP2* and *LTP3*.

GhTCE1 belongs to the E-box (5'-CANNTG-3') binding bHLH TF family, and in order to identify directly the transactivation region, we used a LUC reporter activation system in tobacco leaves and cotton callus protoplasts using the full-length and truncated *GhLTP2* promoter (*LTP2_{pro}* Δ 1 to *LTP2_{pro}* Δ 4) regions and the E-boxes denoted as V–I, respectively (Figure 4E). The results showed that, in addition to the full-length promoter, *LTP2_{pro}* Δ 1, *LTP2_{pro}* Δ 2, and *LTP2_{pro}* Δ 3 can each be activated for target LUC fusion expression in tobacco leaves (Figure 4F). Similarly, we examined *LTP3_{pro}* and its two truncated forms (*LTP3_{pro}* Δ 1 to *LTP3_{pro}* Δ 2) and the E-boxes II and I (Figure 4G). The results show that both *LTP3_{pro}* Δ 1 and *LTP3_{pro}* Δ 2 can be activated for LUC expression (Figure 4H). To further verify the specificity of binding between GhTCE1 and the CANNTG motif, we generated a series of 26 bp probes for different E-boxes in the *LTP2_{pro}* and *LTP3_{pro}* to conduct electrophoretic mobility shift assays (EMSAs) (Figure 4I). Probes V–II, but not I, of *LTP2_{pro}*, and both I and II of *LTP3_{pro}* were bound by the GhTCE1 protein (Figure 4, J and K). In addition, chromatin-immunoprecipitation quantitative PCR (ChIP-qPCR) analysis shows that the promoter region of E-box II–IV of *LTP2_{pro}* and E-box I–II of *LTP3_{pro}* was enriched after immunoprecipitation of GhTCE1 (Figure 4, L and M).

The activation of other candidate targets of GhTCE1 in leaf or other tissues which undergo cell expansion or dedifferentiation following wounding was also investigated. The expression of *GhTCE1* was barely detectable in young leaves of WT and Ri lines, whereas high expression levels (comparable to those seen in SE-7d tissues) were detected in the OE lines (Supplemental Figure S7A). Among the 29 candidate target genes, only 6 were expressed at higher levels in OE lines than in WT (Supplemental Figure S7A). However, the expression levels of all the activated genes in young leaf were much lower than in callus at 7 days post wounding

(SE-), and the average relative expression ratio of the activated candidates between young leaf and callus 7 days post wounding (Leaf/SE) ranged from 0.0008 to 0.0764 (Supplemental Figure S7A). This suggests that GhTCE1 cannot fully activate its candidate targets in young leaves, an idea consistent with the lack of morphological differences between *GhTCE1* OE, Ri lines, and WT (Figure 1F). To investigate *GhTCE1* expression during cell dedifferentiation, callus formation was induced by stem grafting (Supplemental Figure S7B). Expression of *GhTCE1* changed a little during grafting-induced callus formation (Supplemental Figure S7C). Among the 29 candidate target genes of *GhTCE1*, 7 were induced during callus formation (Supplemental Figure S7C), suggesting possible roles in cell dedifferentiation and callus formation, but in a GhTCE1-independent manner. Given that OE of *GhTCE1* inhibits organogenesis from wounded explants, we conclude it must be down-regulated after the early callus induction phase and prior to organogenesis.

Reduced expression of *GhLTP2* and *GhLTP3* results in increased ROS accumulation and reduced callus cell growth during dedifferentiation

The above analysis shows that GhTCE1 activates the expression of both *GhLTP2* and *GhLTP3*. To determine whether *GhLTP2* and *GhLTP3* (*GhLTP2/3*) also function in the control of callus formation or cell differentiation in cotton, *GhLTP2*-Ri13/Ri16 and *GhLTP3*-Ri8/Ri10 transgenic lines were obtained by Ri, and their expression was highly suppressed (Figure 5, A and B). The wounded explants from different transgenic lines showed significantly decreased CPR compared with WT, having smaller cell size during callus formation (Figure 5, C–F). DCFH-DA staining showed that callus from both *GhLTP2* and *GhLTP3* Ri lines accumulate more ROS than WT, and H₂O₂ concentrations increased significantly in the Ri lines (Figure 5, G and H).

To further investigate the possible role for GhTCE1-*GhLTP2/3* in the regulation of ROS homeostasis and a link with cell elongation, tissues were grown in the presence of the ROS scavenger 1,3-dimethylthiourea (DMTU) to determine any effects on callus formation or growth. All *GhLTP2* and *GhLTP3* (*GhLTP2/3*) Ri lines, but particularly *GhLTP2*-Ri13 and *GhLTP2*-Ri16, had more callus growth in the presence of 10 mM DMTU than did mock controls at 20 days post induction, but they had no additional lateral roots (Figure 5, D and I). The cell length of the Ri lines was also significantly greater in the presence of DMTU (Figure 5, F and J). *GhLTP2/3* Ri lines both exhibited less ROS fluorescence in the presence of DMTU, with H₂O₂ concentrations also lower than those seen with mock control treatments (Figure 5, H and K). In addition, we also analyzed the ratio of adventitious root regeneration of *GhLTP2/3* transgenic lines and WT plants, and the results were consistent with those for knockout or Ri *GhTCE1* tissues. Compared with WT plants, *LTP2/3* Ri lines showed a higher ratio of adventitious root regeneration, and this was significantly reduced in the presence of 10 mM DMTU (Supplemental Figure S8).

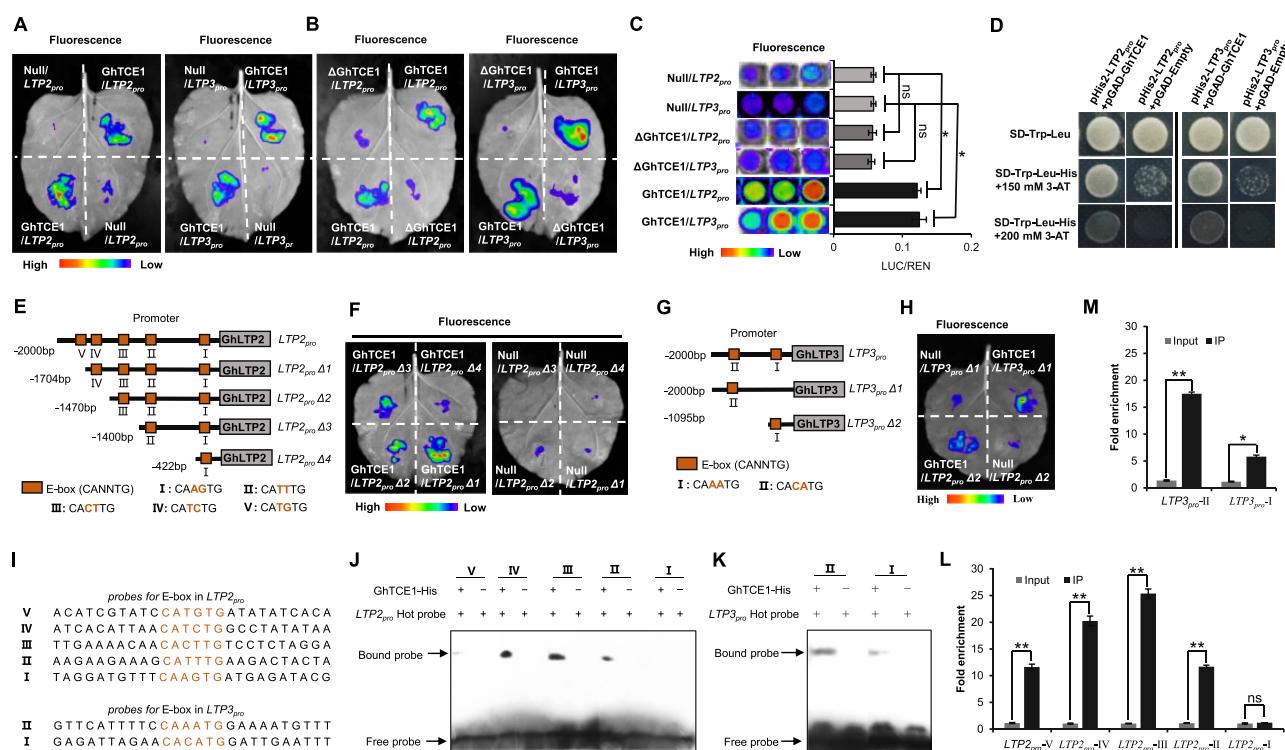


Figure 4 GhTCE1 binds to the promoters of *GhLTP2* and *GhLTP3* to activate their expression. A and B, LUC signals detected in tobacco leaves. Promoters of *GhLTP2* and *GhLTP3* (*LTP2_{pro}* and *LTP3_{pro}*) were separately fused upstream of LUC in the vector pGreenII 0800-LUC and designated *LTP2_{pro}:LUC* and *LTP3_{pro}:LUC*. As an internal control, *35S_{pro}:REN* in vector pGreenII 0800-LUC was used. *GhTCE1* and Δ *GhTCE1* were fused downstream of the 35S promoter in pGreenII 62-SK. The pGreenII 62-SK empty vector was used as negative control and named Null. For each tobacco leaf, half was used to express pGreenII 0800-LUC:*LTP_{pro}* + pGreenII 62-SK:*GhTCE1* + *35S_{pro}:REN* (named *GhTCE1/LTP_{pro}*) and half was used to transiently express pGreenII 0800-LUC:*LTP_{pro}* + pGreenII 62-SK empty vector:*REN* (named Null/*LTP_{pro}*) following *Agrobacterium*-mediated transfection, Δ *GhTCE1* in the same way. C, Activated expression of *LTP2_{pro}* and *LTP3_{pro}* by *GhTCE1* in cotton callus protoplasts using the LUC expression system. Relative activation level represents the ratio of fluorescence intensity of LUC/REN with *GhTCE1* to LUC/REN without *GhTCE1*. D, Y1H assays of *GhTCE1* binding to *LTP2_{pro}* and *LTP3_{pro}*. The full-length coding sequence of *GhTCE1* was fused to the pGADT7 vector, and *LTP2_{pro}* and *LTP3_{pro}* were cloned into the pHis2 vector. Interaction was determined on selective medium lacking Trp + Leu + His in the presence of 3-AT. E, A series of truncations (*LTP2_{pro}* Δ 1 to *LTP2_{pro}* Δ 4) with E-box deletions were used in the LUC activation system vector. F, *GhTCE1* activates the expression of truncated *LTP2_{pro}* by detected LUC signals on tobacco leaves. The truncated *LTP2_{pro}* constructs were fused to the vector pGreenII 0800-LUC, and combined with pGreenII 62-SK:*GhTCE1* + *35S_{pro}:REN* as a positive control. The pGreenII 62-SK empty vector was used as a negative control. They are labeled the same way as in (A) and (B). G, *GhTCE1* activates the expression of truncated *LTP3_{pro}* (*LTP3_{pro}* Δ 1 and *LTP3_{pro}* Δ 2) by LUC signals detected on tobacco leaves. H, *GhTCE1* activates the expression of truncated *LTP3_{pro}* (*LTP3_{pro}* Δ 1 and *LTP3_{pro}* Δ 2) by LUC signals detected on tobacco leaves. The process same to (F). I, The probe sequences of *LTP2_{pro}* and *LTP3_{pro}* are noted and the E-boxes are shown in red. J and K, EMSA showing the specificity of in vitro binding between *GhTCE1* and the E-box elements of *LTP2_{pro}* (J) and *LTP3_{pro}* (K). The WT probes were subjected to biotin-labeling. No *GhTCE1*-His tag protein lines as control. The arrows indicate the up-shifted binds. L and M, Validation of the binding of *GhTCE1* to the promoters E-box elements of *LTP2_{pro}* (L) and *LTP3_{pro}* (M) by ChIP-qPCR. Error bars in (C), (L), and (M) represent \pm standard errors of three biological replicates (10 samples each in [C], three samples each in [L] and [M]). Statistical significance is indicated by **P* < 0.05, ***P* < 0.01, Supplemental Data Set S2.

These results further indicate that *GhLTP2/3* expression is associated with low ROS accumulation and increased disorganized cell growth.

GhTCE1 enhances the activation ability of GhTCE1 by forming heterodimers

We have shown here that *GhTCE1* activates cell elongation-related genes during somatic cell dedifferentiation and callus formation, whereas the activation of these genes by *GhTCE1* is much weaker in cotton and tobacco leaves (Figure 3E). This suggests that *GhTCE1* alone is not sufficient for full

transcriptional activation and that other partners may be also involved during cell reprogramming. To identify potential partner proteins, the activation domain-truncated version of *GhTCE1* (C-terminal from 161 to 331 aa), which contains the bHLH domain, was used as bait for yeast two-hybrid screening with the prey library constructed from callus cDNA isolated early after wounding. A total of 10 *GhTCE1* candidate interacting proteins were identified (Supplemental Table S3). It has been reported that bHLHs usually regulate downstream targets by dimerization (Toledo-Ortiz et al., 2003), and so we focused on members of this family of TFs. We found that three bHLH proteins

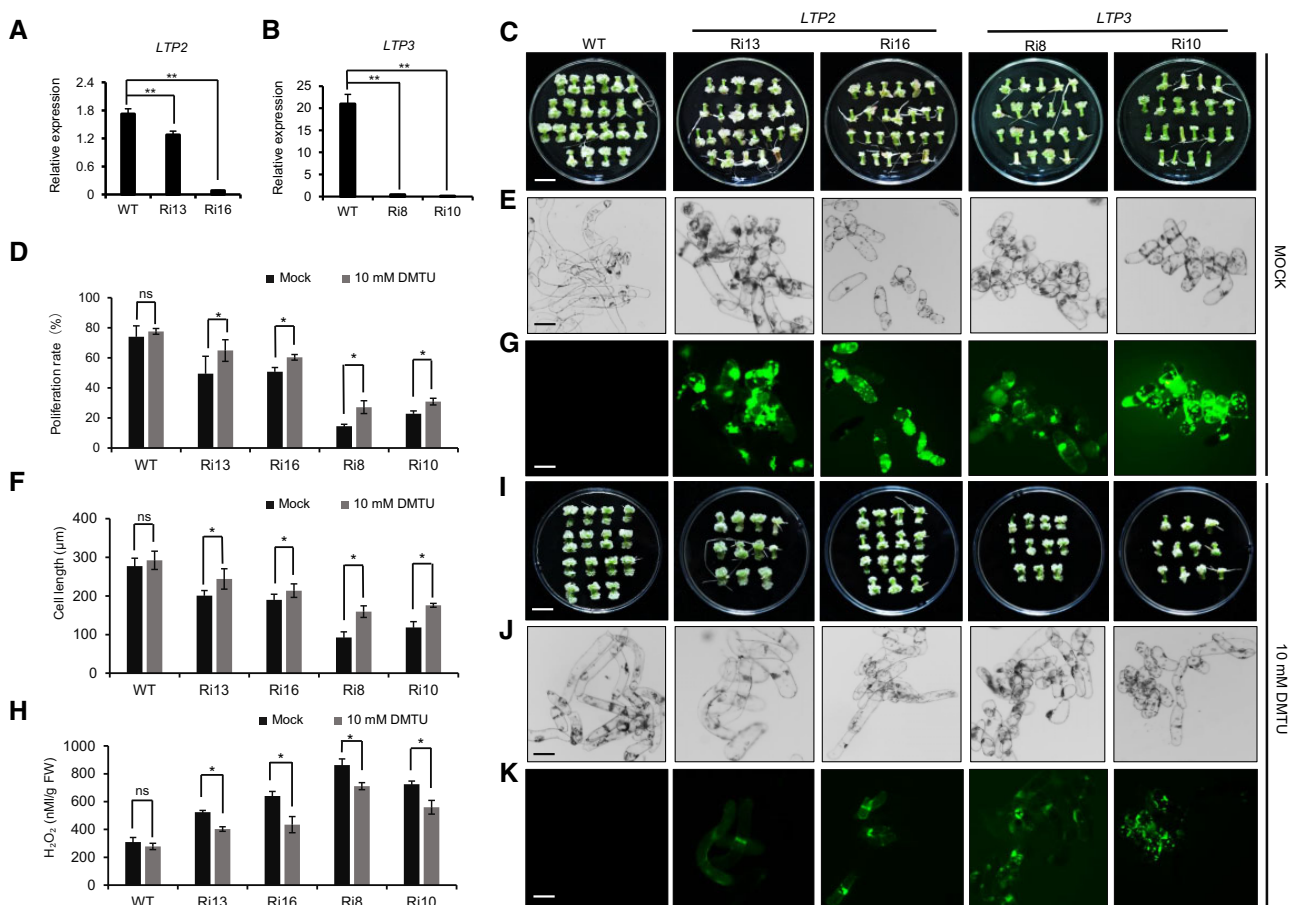


Figure 5 Phenotypic analysis of *GhLTP2* and *GhLTP3* Ri transgenic tissues. A and B, Relative expression of *GhLTP2* and *GhLTP3* in WT and respective Ri transgenic lines. Error bars represent \pm standard error of three biological replicates as described in Figure 1A. C, Phenotype of callus 20 days post wounding for different transgenic lines following mock treatment. Bar = 1 cm. D, CPR of different lines 20 days post wounding following mock treatment or treatment with 10 mM DMTU. Error bars represent \pm standard error of six biological replicates as described in Figure 2B. E, Micrographs of callus cells, under mock treatment. Bar = 100 μ m. F, Statistic analysis of callus cell lengths of different lines 20 days post wounding under mock treatment and 10 mM DMTU. Error bars represent \pm standard error of six biological replicates as described in Figure 2D. G, DCFH-DA staining of ROS in WT, *GhLTP2*, and *GhLTP3* transgenic callus cells. Bar = 100 μ m. H, H₂O₂ concentration in callus from transgenic plants 20 days post SE induction. Error bars represent \pm standard errors of three biological replicates as described in Figure 2F. I–K, Phenotype, micrograph, and DCFH-DA staining of ROS in callus cells, 20 days post wounding from different transgenic lines under 10 mM DMTU treatment. Bar = 1 cm in I and 100 μ m in J and K. Significance tests compared each transgenic line with WT plants. * $P < 0.05$, ** $P < 0.01$. Results of statistical analyses are presented in Supplemental Data Set S2.

(Gh_D11G3523, Gh_D11G0415, and Gh_D11G0650) interacted with GhTCE1 (Figure 6A and Supplemental Figure S9). The interaction between GhTCE1 and GhTCEE1 (Gh_D11G3523) was further verified by bimolecular fluorescence complementation (BiFC) (Figure 6B). Consistent with interaction between the two proteins, the expression of *GhTCEE1* was induced at earlier time-points after wounding, was detectable within 540 min after wounding, and increased strongly beyond this to 720 min after wounding, showing a co-expression pattern with *GhTCE1* (Supplemental Figure S10).

The LUC reporter activation assay was also used to verify the enhanced activation ability of GhTCE1 by GhTCEE1. The results showed that GhTCEE1 can activate *GhLTP2_{pro}* and *GhLTP3_{pro}* for LUC expression in a similar way as does GhTCE1 (Figure 6C). Furthermore, co-expression of GhTCE1

and GhTCEE1 led to significantly higher levels of *GhLTP2_{pro}* and *GhLTP3_{pro}* for LUC expression than either GhTCE1 or GhTCEE1 alone (Figure 6D), showing that GhTCE1 and its interacting protein GhTCEE1 function cooperatively to activate the expression of both *GhLTP2* and *GhLTP3*.

Discussion

GhTCE1 regulates callus and adventitious root formation in an antagonistic manner

Callus and adventitious roots are typically induced following wounding or exogenous hormone application. The mutants *alf4-1* and *clf-50 swm-1* are defective in callus formation and also failed to produce adventitious roots, suggesting that callus and adventitious roots share some common genetic pathways (Liu et al., 2014). It has also been suggested that callus, formed from multiple organs, shares a gene

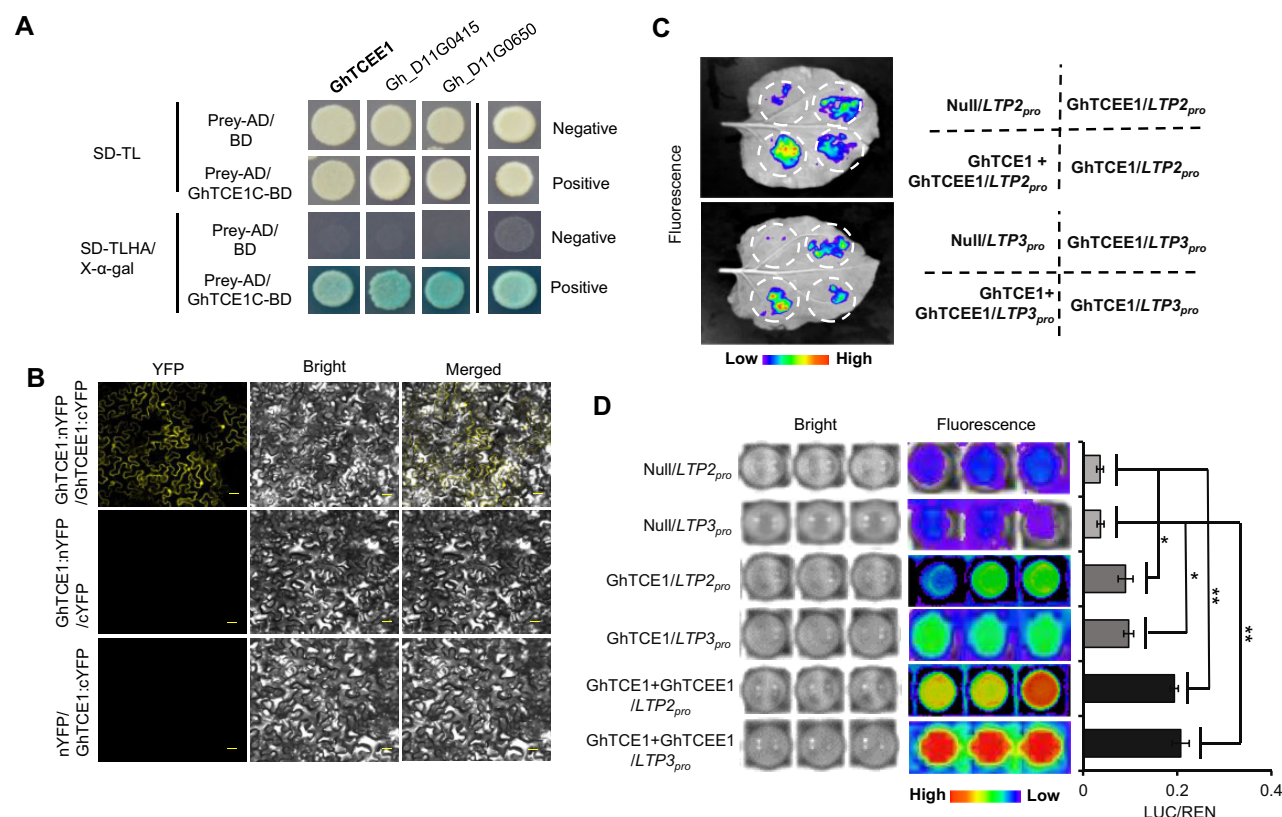


Figure 6 GhTCEE1 enhances the activation activity of GhTCE1 by forming heterodimers. **A**, Sequences encoding the GhTCE1 C-terminus from aa residues 160–331 without transactivation activity were fused to sequences encoding a GAL4 BD domain and designated GhTCEE1-BD. The prey AD was linked to callus cDNAs. Negative represents no growth, positive represents growth. SD-TL/TLHA represents synthetically defined medium dropouts Trp, Leu or Trp, Leu, His, and Ade, respectively. **B**, BiFC assay between GhTCE1–nYFP and GhTCEE1–cYFP in tobacco epidermal cells. Bars = 30 μ m. **C**, LUC assays in tobacco leaves and cotton callus protoplasts. Promoters of *GhLTP2* and *GhLTP3* were, respectively, fused upstream of LUC. *35S_{pro}*:REN was used as an internal control. GhTCE1 and GhTCEE1 were fused downstream of *35S_{pro}* in pGreenII 62-SK. The pGreenII 62-SK empty vector was used as control named Null. For each tobacco leaf, four constructs were expressed: (i) pGreenII 0800-LUC:*GhLTP2_{pro}* + *35S_{pro}*:REN + pGreenII 62-SK empty (Null/*LTP2_{pro}*); (ii) pGreenII 0800-LUC:*GhLTP2_{pro}* + *35S_{pro}*:REN + pGreenII 62-SK:GhTCE1 (GhTCE1/*LTP2_{pro}*); (iii) pGreenII 0800-LUC:*GhLTP2_{pro}* + *35S_{pro}*:REN + pGreenII 62-SK:GhTCEE1 (GhTCEE1/*LTP2_{pro}*); and (iv) pGreenII 0800-LUC:*GhLTP2_{pro}* + *35S_{pro}*:REN + pGreenII 62-SK:GhTCE1 + GhTCEE1 (GhTCE1 + GhTCEE1/*LTP2_{pro}*) following *Agrobacterium*-mediated transfection. **D**, Activation of LUC expression by GhTCE1/GhTCEE1 in cotton callus protoplasts. Relative activation level represents the ratio of fluorescence intensity of LUC/REN with GhTCE1 to LUC/REN without GhTCE1. Error bars represent \pm standard errors of three biological replicates as described in Figure 3E. Statistical significance is indicated by * $P < 0.05$, ** $P < 0.01$, Supplemental Data Set S2.

expression pattern with some similarities to the root meristem (Sugimoto et al., 2010). LATERAL ORGAN BOUNDARIES DOMAIN (LBD) proteins are key transcriptional regulators for both lateral root development and callus induction, and their expression is directly regulated by WUSCHEL RELATED HOMEODOMAIN11 (WOX11) and WOX12 (Fan et al., 2012; Liu et al., 2014, 2018). In addition, WOX13 also plays a key role in callus formation and organ reconnection following grafting by regulating cellular reprogramming and cell wall modification (Ikeuchi et al., 2021). LBDs can interact with specific bHLH proteins to co-regulate downstream targets (Husbands et al., 2007). GhTCE1 acts as a bHLH protein and was found to interact with multiple TFs in yeast two-hybrid screens (Supplemental Figure S7). Future studies could determine whether GhTCE1 also interacts with LBDs during callus or adventitious organ development.

Although callus and adventitious roots are regulated by some common factors, there exists an antagonism between extensive callus formation, in which differentiated cells reprogramme and undergo elongation as they lose their original differentiated state, and adventitious root formation, in which cells form de novo meristems which, in contrast to callus cells, are small and highly organized. High concentrations of the auxin IBA promote callus production while lower concentrations promote adventitious roots (Supplemental Figure S2B), consistent with previous reports for indole acetic acid (IAA) treatment (Liu et al., 2014). Previously we found that GhL1L1 promotes adventitious root development rather than callus proliferation (Xu et al., 2019). In the current study, we show that explants expressing reduced levels of *GhTCE1* showed reduced callus proliferation but enhanced adventitious root formation (Figure 2A),

suggesting that *GhTCE1* has an opposite or competing function to *GhL1L1*, further illustrating the antagonistic relationship between callus formation and adventitious root development.

***GhTCE1* regulates callus cell proliferation and elongation by regulating multiple pathways**

The transcriptomic analysis presented shows that multiple pathways linked to cell proliferation and elongation are regulated by *GhTCE1*. Among the 61 DEGs activated by *GhTCE1*, genes related to sterol synthesis, cell wall modification, cell extension/expansion, ROS homeostasis, and lipid transfer were highly enriched. Sterols such as brassinosteroids (BRs) are required for cell elongation and plant growth (Clouse, 2002). BR treatment of cultured cotton ovules results in a significant increase in fiber length, while the BR synthesis inhibitor BRZ strongly inhibits fiber elongation (Sun et al., 2005). BR-defective mutants show extremely dwarfed phenotypes with reduced cell length (Azpiroz, 1998). The BR-inducible gene *EXORDIUM* (*EXO*) was also found among the DEGs. *EXO* was originally identified in a promoter trap screen for early embryo genes in *Arabidopsis* (Farrar et al., 2003), and it encodes an extracellular protein required for cell expansion in *Arabidopsis* (Schroder et al., 2009).

Cell wall modification and cell expansion pathways regulate cell elongation. The identified DEGs contain genes predicted to be involved in cell wall metabolism, including pectin methylesterase, putative pectin acyl transferase, β -1,3-glucanase, and BBE-like enzyme for monolignol oxidation, expansion, and extension. Pectin de-methylesterification is important for hypocotyl and internode cell elongation in *Arabidopsis* (Pelletier et al., 2010; Peaucelle et al., 2011). Also, pectin acetylation is involved in hypocotyl elongation, and mutation of pectin acyltransferase results in short hypocotyls under dark growth conditions (Sinclair et al., 2017). Callose (β -1,3-glucan) can accumulate under stress conditions to impregnate the cell wall and restrict cell growth (Piršelová and Matušíková, 2012). Degradation of callose by β -1,3-glucanase can increase cell wall flexibility and promote pollen tube germination and elongation (Parre and Geitmann, 2005). BBE-like was reported to participate in the oxidation and mobilization of monolignols (Daniel et al., 2015), and a direct target of bZIP59-LBD16 in *Arabidopsis* is required for callus initiation during early SE (Xu et al., 2018), suggesting that cell wall metabolism is important for cell dedifferentiation and callus growth. Extensins and expansins are involved in cell wall assembly and are required for elongation of root hairs and hypocotyl cells (Cannon et al., 2008; Zdanio et al., 2020). We found that two homologous genes, encoding an extensin and an expansin, were activated by *GhTCE1* during dedifferentiation of cotton somatic cells (Figure 3E). Ri of the extensin gene (Gh_D09G1992) resulted in decreased callus proliferation and cell elongation (Supplemental Figure S11). These data show that *GhTCE1*

likely promotes callus cell growth, at least in part, by regulating the cell wall modification machinery.

***LTPs* participate in cell fate switching via ROS homeostasis**

A key feature of this work is the discovery that *GhTCE1* regulates the transcription of the lipid transfer protein genes *LTP2* and *LTP3* through promoter binding (Figure 4), and data strongly suggest a link with ROS homeostasis in cell fate determination. ROS, including hydrogen peroxide (H_2O_2), singlet oxygen ($^1\text{O}_2$), the superoxide anion (O_2^-), and the hydroxyl radical (OH^\bullet), can either promote or repress cell growth according to the particular species and distribution (Passardi et al., 2004). H_2O_2 promotes cell differentiation and strengthens the cell wall by promoting the crosslinking of cell wall components (Raggi et al., 2015). In contrast, OH^\bullet promotes cell elongation by cleaving polysaccharides or activating calcium channels (Foreman et al., 2003; Muller et al., 2009). In the current study, two NADPH oxidase and six peroxidase genes were identified as candidate targets of *GhTCE1*. NADPH oxidase participates in the production of O_2^- , which can be converted into H_2O_2 spontaneously or by superoxide dismutase (SOD). Peroxidases are involved in H_2O_2 scavenging by substrate oxidation or in the production of OH^\bullet via the Haber–Weiss reaction (Shigeto and Tsutsumi, 2016). We provide evidence that *GhTCE1* is also involved in ROS homeostasis, and potentially promotes cell elongation by activating the expression of peroxidases and NADPH oxidase.

LTPs represent other potential participants in cell elongation, but their functions have not been well characterized. A number of possible mechanisms by which they influence cell elongation should be considered. LTPs are small compact proteins with lipid binding and transport activity (Edqvist et al., 2018). They are usually localized to extracellular spaces, with targeting mediated by N-terminal signal peptides (Boutrot et al., 2008). LTPs are involved in the deposition of wax and lipid-based polymers in the defense response, as well as in cell wall loosening during cell elongation (Liu et al., 2015). LTPs can interact with, and promote the activity of the pectin-degrading enzyme polygalacturonase (PG), thereby loosening the cell wall (Tomassen et al., 2007). The lipid binding hydrophobic cavity of LTP may also interact with hydrophobic cell wall compounds, thereby disrupting their noncovalent bonding and promoting cell wall extension (Nieuwland et al., 2005). In the current study, reduced expression of *LTP2* and *LTP3* by Ri resulted in ROS accumulation and restricted callus cell growth, demonstrating a role for LTPs in mediating dedifferentiation control via ROS. Of particular significance here is the previously reported potential role for LTPs as ROS scavengers (e.g. Wang et al., 2014; Gangadhar et al., 2016; Xu et al., 2018). These studies describe how LTPs are upregulated under a range of abiotic stress conditions and can confer stress tolerance by scavenging ROS. Our data are consistent with a role for LTPs in scavenging ROS, regulating the transition from

differentiated cell type (in the explant) to disorganized callus growth following wounding.

In vitro dedifferentiation of somatic cells post wounding is orchestrated by TFs

Callus induction is a clear example of plant developmental plasticity in which wounding leads to cell dedifferentiation, requiring a reprogramming of gene expression by induced TFs. Auxin response factor (ARF)- and cytokinin response factor (CRF)-mediated auxin and cytokinin responses are essential for callus formation under appropriate auxin/cytokinin ratios. The *arf7 arf9* double mutant displays defective callus initiation, but ectopic expression of their downstream targets *LBD16/17/18/29* is sufficient to induce autonomous callus production in the absence of exogenous phytohormones (Fan et al., 2012; Xu et al., 2018). Ectopic expression of *CRF3* also results in autonomous callus formation (Xu et al., 2012). Furthermore, *ARF10* represses the expression of *ARABIDOPSIS RESPONSE REGULATOR 15* (*ARR15*) to promote callus initiation, highlighting the interplay between auxin and cytokinin (Liu et al., 2016). To date, more than 50 TFs have been reported to function during regeneration from callus in Arabidopsis, and nearly half of them participate in the regulation of callus growth itself (Ikeuchi et al., 2019). In the current study, we show that *GhTCE1*, with its partner protein *GhTCEE1*, is a part of a critical regulatory module involving LTPs and ROS. This module promotes cell dedifferentiation and callus growth in an auxin-independent manner, and it acts as a molecular switch to determine cell fate control by promoting callus growth and repressing more organized meristematic growth (Figure 7).

Materials and methods

Plant materials, vector construction, and genetic transformation

The cotton used in this study is *G. hirsutum* cv YZ1. Transgenic lines were planted in the greenhouse at Huazhong Agricultural University, Wuhan, China. The greenhouse was kept at 26°C–32°C, 14-h light/10-h dark photoperiod. The explants from transgenic lines and tobaccos that were used for LUC reporter assays were grown at 25°C in a light cultivation room (16-h light/8-h dark photoperiod, 3,000 lex). Ri vectors for *GhTCE1* and *GhLTP2/3* were constructed in the pHellsgate 4 vector (Wesley et al., 2001). The OE construct for *GhTCE1* was in the pK2GW7 (Karimi et al., 2002) vector. The CRISPR-Cas9-mediated gene editing vector was constructed as described previously (Wang et al., 2018). All the primers used for vector construction are listed in Supplemental File S1. Transgenic plants were created by Agrobacterium-mediated transformation as described previously (Jin et al., 2006). The expression levels of *GhTCE1* and *GhLTP2/3* in transgenic plants were determined by qRT-PCR, using explants at 7 days post wounding, using *GhUB7* as an internal control. T-DNA insertion copy numbers were determined by immunoblotting as described previously (Wang et al., 2018). The mutation analysis of *GhTCE1*

CRISPR-Cas9 transgenic plants was achieved by the Hi-TOM method as described previously (Liu et al., 2019). Mutants of *GhTCE1(A/D)* carrying frameshift mutations were chosen for phenotypic analysis.

Callus induction, CPR, cell length analysis, and adventitious root formation

Hypocotyls of etiolated seedlings were cut into 5–7 mm sections as explants for culture on MSB (MS medium plus B5 vitamins) medium and growth conditions of explants as described previously (Yang et al., 2012). The CPR represents the gained weight of per unit weight explants at 20 days post wounding induction. Callus cells were separated in 1 × PBS (10 mM sodium phosphate, 137 mM NaCl, 2.7 mM KCl, and pH 7.5) and photographed under a light microscope (ZEISS, AXIO Scope A1). Cell length was measured from the micrograph using Image J software (Rha et al., 2015). Adventitious root formation was measured as a ratio of the frequency of roots formed per explant cultured.

DCFH-DA treatment, ROS staining, and H₂O₂ measurement

Hypocotyls of etiolated seedling were cut into 5–7 mm sections as explants, 10 mM DMTU was added to MSB medium as the treatment group, and MSB medium lacking DMTU was the negative control.

For ROS staining, fresh callus cells were collected 20 days post wounding and stained in the DCFH-DA solution described below for 5 min. Stained cells were rinsed three times in ddH₂O and examined under a fluorescence microscope (Nikon, SMZ25) with 480 nm excitation, and 100 ms exposure time was used for each image. The DCFH-DA storage solution (pH 7.5) was 10 mmol/L prepared in dimethylsulfoxide, and it was diluted 1,000-fold with 1 × PBS for ROS staining. The concentration of H₂O₂ from fresh callus was determined using a H₂O₂ quantification kit (#C500069, Sangon Biotech, Shanghai, China).

Endogenous IAA concentration assays and IBA treatment

Explants were collected 20 days post wounding and the IAA levels were determined by HPLC-MS as reported previously (Xu et al., 2019). In standard MSB medium, the IBA concentration (0.5 mg/L) was defined as 1 ×. The final concentrations of IBA used were 0, 0.5, 1, 2.5, 5, and 10 mg/L in MSB medium and defined as 0 ×, 1 ×, 2 ×, 5 ×, 10 ×, and 20 ×, respectively, for IBA treatments. Explants were photographed 10 and 20 days post wounding.

RNA-seq

Hypocotyls of WT and *GhTCE1* Ri and OE lines were used for RNA extraction 7 days post wounding. RNA-seq was performed by the Beijing Genomics Institute. DEGs were determined using the threshold of $P \leq 0.05$ and the absolute value of the log₂ fold change ≥ 1 . The annotations of DEGs were based on the functions of their homologues in Arabidopsis unless known in cotton.

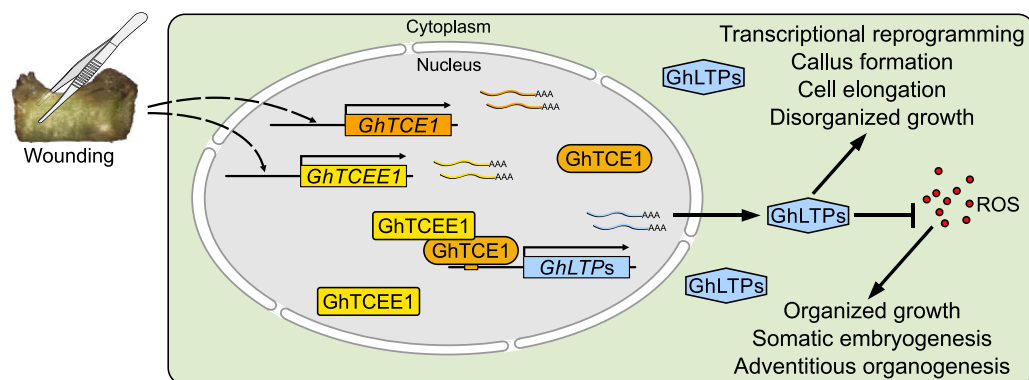


Figure 7 Model outlining the proposed relationship between mechanical wounding, GhTCE1 and GhTCEE1 (bHLH dimers), LTPs, ROS, and cell fate determination. We propose that following wounding and callus initiation, GhTCE1 and GhTCEE1 are simultaneously at the transcriptional level, and the proteins interact to activate the transcription of LTP genes *GhLTP2* and *GhLTP3*. The LTP proteins repress ROS activity, possibly by acting as ROS scavengers, as suggested by previous LTP studies in relation to roles in abiotic stress tolerance. Repression of ROS leads to the maintenance of callus cell identity and growth, whereas an increased level of ROS activity is associated with reduced cell growth and cell differentiation. We propose that the TCE1/TCEE1 transcriptional module regulates cell fate determination through these effects on ROS-mediated cell growth and differentiation control.

Subcellular localization

Full length *GhTCE1* was cloned into the vector pMDC43 with GFP fused to the N-terminus (Cai and Goodman, 2015). The $35S_{pro}$:GFP-*GhTCE1* construct was transiently expressed in tobacco epidermal cells following *Agrobacterium*-mediated transfection. The pMDC43 empty vector harboring $35S_{pro}$:GFP was used as control. GFP fluorescence was detected under a confocal microscope (Olympus FV1200) at 2 days following *Agrobacterium* transfection.

BiFC assay

For the BiFC assay, the full-length sequences of *GhTCE1* and *GhTCEE1* were cloned into pDONR vector (Invitrogen), then into the destination vector pBiFCt-2in1-NC by Gateway Technology (Grefen and Blatt, 2012). The recombinant vectors were transformed into *Agrobacterium tumefaciens* strain GV3101, and transient expression was performed in tobacco epidermal cells. The YFP fluorescence in BiFC assays was examined by confocal microscopy (Olympus FV1200).

Autoactivation analysis of GhTCE1

The full-length sequence of *GhTCE1* was inserted into the pGBKT7 vector to produce the Full:BD construct. Full:BD was transferred into the Y2HGold yeast strain and cultured on SD-T/X- α -gal plates for 3 days to detect its autoactivation (based on colony color). To further investigate the autoactivation in detail, four different truncated *GhTCE1* versions were constructed in pGBKT7 and designated 1–160, 161–331, 80–240, and 80–331 according to the amino acid fragments contained in each. The different constructions of *GhTCE1* were transferred into the Y2HGold yeast strain and mated with Y187 yeast harboring the pGADT7 empty vector. The positive and negative controls used were according to the manufacturer's instructions (Clontech, PT4084-1). The diploid yeasts were cultured in SD-TL and SD-TLHA/X- α -gal plates for 3 days to detect autoactivation. SD means

synthetically defined medium, SD-T/TL/TLHA represent SD dropouts Trp, Trp/Leu, and Trp/Leu/His/Ade, respectively.

Transactivation assays using the LUC reporter system

Transactivation assays were conducted as reported previously (Hu et al., 2018). In brief, the promoters of *GhTCE1* candidate targets were cloned and inserted upstream of sequences encoding the LUC reporter in the pGreenII 0800-LUC vector. $35S_{pro}$:REN in the same vector was used as an internal control. *GhTCE1* and *GhTCEE1* were cloned downstream of the cauliflower mosaic virus 35S promoter ($35S_{pro}$) in pGreenII 62-SK and the same vector without *GhTCE1* was used as control. Activation of E-box motifs assays was conducted using the 2,000-bp full-length *GhLTP2* promoter with five E-box motifs (I, II, III, IV, and V) and the *GhLTP3* promoter with two E-box motifs (I and II). The truncation promoters were created by successively deleting one E-box in turn from the 5'-end and were named *LTP2_{pro}* Δ 1 to *LTP2_{pro}* Δ 4 for deletions of *GhLTP2_{pro}* and named *LTP3_{pro}* Δ 1 to *LTP3_{pro}* Δ 2 for *GhLTP3_{pro}*. The truncated promoters were cloned into the pGreenII 0800-LUC vector. All plasmids were transformed individually into *A. tumefaciens* strain GV3101 for transient expression in tobacco (*N. benthamiana*). The concentration of *A. tumefaciens* harboring the pGreenII 0800-LUC or pGreenII 62-SK vectors was adjusted to the same concentration (e.g. pGreenII 62-SK:*GhTCE1*, OD = 0.2, pGreenII 0800:*LTP2_{pro}*, OD = 0.2) and mixed in volumes of 1:1 (pGreenII 0800-LUC: pGreenII 62-SK), and all of the pairs of the suspensions carrying the appropriate constructs were adjusted to the same OD₆₀₀ = 0.6 prior to coinfiltration into tobacco leaves with same infiltrated areas to ensure comparability. The infiltrated plants were grown at 25°C in a light cultivation room (16-h light/8-h dark photoperiod, irradiance of 135 $\mu\text{mol m}^{-2}\text{s}^{-1}$). After 60–72 h, the LUC luminescence of leaves was detected using a

cryogenically cooled CCD camera (Lumazome PyLoN 2048B). For each tobacco leaf, one half was transfected with experimental vectors and the other half with control vectors by agroinfection. However, for four truncations, 62-SK:GhTCE1+ pGreenII 0800:*LTP2_{pro}*Δ constructs were evenly distributed on one leaf. Three biological replicates were carried out for each combination.

The LUC assays in cotton protoplasts were performed as described (Hellens et al., 2005). In brief, vector each of effector (1:1) and reporter constructs (total effectors:reporter = 1:1) were cotransformed into protoplasts using 40% polyethylene glycol 4000 (v/v) (Sigma), then cultured at 25°C in darkness for 16 h. LUC and REN activity were detected (the relative ratio of the LUC/REN ratio with GhTCE1 and GhTCEE1 versus the LUC/REN ratio without GhTCE1s and GhTCEE1) for transactivation assays using GhTCE1 and its candidate targets by using the dual-LUC assay reagents (Promega, Madison, WI) with the Multimode Plate Reader (Perkin-Elmer).

Yeast two-hybrid screening

The activation domain of truncated GhTCE1 (C-terminal from 160 to 331 aa) was cloned into pGBKT7 (designated pGBKT7-ΔGhTCE1) and transformed into yeast strain Y2HGold. The cDNA library from early time points after wounding were constructed in pGADT7 and transformed into yeast strain Y187. Yeast two-hybrid screening was performed according to the manufacturer's instructions (Clontech, PT1172-1, PT4084-1). The vectors from positive clones were recovered and transferred into *Escherichia coli* for sequencing. After removing the unmatched or frameshift preys, the remaining protein coding preys (designated prey-AD) were transferred into the Y187 strain separately and mated with the Y2H strain containing either the pGBKT7 empty vector (BD) or pGBKT7-ΔGhTCE1 (ΔGhTCE1-BD). Both prey-AD/BD and prey-AD/ΔGhTCE1-BD combinations showing activated reporter genes were considered as false interactions. Only prey-AD + ΔGhTCE1-BD activated reporter genes while the prey-AD/BD did not were considered as true interactions.

Electrophoretic mobility shift assays

Biotin-labeled single-stranded DNA was synthesized and annealed to produce double strands. For EMSA, a sequence encoding full-length GhTCE1 was cloned into the pET28a (Novagen) vector, and His-tagged proteins were induced and expression in *E. coli* and proteins were purified by chromatography. Binding reactions were carried out using the LightShift Chemiluminescent EMSA Kit (Thermo scientific, 20148). Signals were captured by X-ray film and the film was developed according to the manufacturer's instructions. The primers used for vector construction are listed in Supplemental File S1.

Chromatin-immunoprecipitation quantitative PCR

For ChIP experiments, *GhTCE1*-GFP transgenic lines were used for chromatin extraction and immunoprecipitation as

described previously by Huang et al. (2018). Briefly, the samples were fixed with 1% (v/v) formaldehyde (Sigma, Cat. # F8775) for 10 min at room temperature, and neutralized with 0.125 M glycine for 5 min. The samples were collected by centrifuging (Beckman Coulter, Avanti JXN-26) at 1,000 g at 4°C for 5 min. A Bioruptor (KQ5200DE) was used at high power with 30-s on/30-s off cycles for 15 times until the average chromatin size was approximately 300 bp. Anti-GFP (ABclonal, AE012) antibodies (5 μg antibodies were added per 25 μg chromatin DNA) were used to perform immunoprecipitations. The immunoprecipitated chromatin fragments were isolated and preabsorbed with sheared salmon sperm DNA/protein A-agarose (Sigma-Aldrich) to remove nonspecific binding (IP). After DNA purification, the GhTCE1-binding sites on *LTP2_{pro}* and *LTP3_{pro}* were evaluated using qPCR and normalized by total chromatin (Input). The fold enrichment calculated as the ratio between the samples of input and IP samples. *GhUBQ7_{pro}* was used as reference promoter and the primers are listed in Supplemental File S1.

Y1H assay

Y1H screening was performed using the Matchmaker Gold Yeast One-Hybrid Library Screening System User Manual (Clontech, PT4087-1). The full-length sequence of GhTCE1 was cloned into the pGADT7 vector and transformed into yeast strain Y187, the promoter sequences of *LTP2_{pro}* and *LTP3_{pro}* were cloned into the pHis2 vector and transformed into yeast strain AH109, and the pGAD-empty vector was used as negative control vector. Ten microliters of each for combined yeast strains Y187 and AH109 were cultured in 200 μL YPDA liquid medium (1 L YPDA contains 10 g yeast extract, 20 g tryptone, 20 g glucose, 100 mg adenine, fill ddH₂O to 1 L, pH = 6.5) at 30°C for 20–24 h. Then, 6 μL yeast that mated successfully was spotted onto SD-Trp-Leu-His containing 200 mM 3-AT solid medium at 30°C for 3–5 days, then analyzed the interaction between GhTCE1 and *LTP2_{pro}* and *LTP3_{pro}*. The primers used for vector construction are listed in Supplemental File S1.

Accession numbers

Sequence information for the cotton genes in this study can be found in the Cotton Genome Database (<https://cot tonfgd.org/>, *G. hirsutum* AD1, upland cotton, NAU assembly) according to the accession numbers as shown in Supplemental Data Set S1 and Supplemental Tables S1–S3.

Supplemental data

The following materials are available in the online version of this article.

Supplemental Figure S1. Protein sequence alignment of Gh_D06G2316 (GhTCE1D) and GhA_06G0241 (GhTCE1A).

Supplemental Figure S2. GhTCE1 is involved in adventitious root regeneration during wound-induced callus formation.

Supplemental Figure S3. Auxin detection in *GhTCE1* transgenic plants.

Supplemental Figure S4. Callus formation in GhTCE1 transgenic lines.

Supplemental Figure S5. Correlation analysis of 29 DEGs comparing RNA-seq versus qRT-PCR.

Supplemental Figure S6. Subcellular localization and autoactivation analysis of GhTCE1.

Supplemental Figure S7. Expression analysis of *GhTCE1* and its candidate targets in different tissues.

Supplemental Figure S8. *GhLTP2* and *GhLTP3* are involved in adventitious root regeneration during wound-induced callus formation.

Supplemental Figure S9. Yeast two-hybrid screening for proteins interacting with GhTCE1.

Supplemental Figure S10. Expression pattern of GhTCEE1 at early time points following wounding.

Supplemental Figure S11. EXT1A and EXT1D are involved in cell elongation during SE.

Supplemental Data Set S1. FPKM values of 62 DEGs identified by RNA-seq analysis.

Supplemental Data Set S2. Statistical analysis tables.

Supplemental Table S1. Function annotation of 32 left candidate targets of GhTCE1.

Supplemental Table S2. Characterization of 29 GhTCE1 candidate targets.

Supplemental Table S3. Annotation of proteins interacting with GhTCE1 from yeast two-hybrid screening.

Supplemental File S1. List of primers used in this study.

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Conflict of interest statement. The authors declare that there is no conflicts of interest.

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