



Transcription factors TgbHLH95 and TgbZIP44 cotarget terpene biosynthesis gene *TgGPPS* in *Torreya grandis* nuts

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Abstract

Terpenes are volatile compounds responsible for aroma and the postharvest quality of commercially important xiangfei (*Torreya grandis*) nuts, and there is interest in understanding the regulation of their biosynthesis. Here, a transcriptomics analysis of xiangfei nuts after harvest identified 156 genes associated with the terpenoid metabolic pathway. A geranyl diphosphate (GPP) synthase (*TgGPPS*) involved in production of the monoterpene precursor GPP was targeted for functional characterization, and its transcript levels positively correlated with terpene levels. Furthermore, transient overexpression of *TgGPPS* in tobacco (*Nicotiana tabacum*) leaves or tomato (*Solanum lycopersicum*) fruit led to monoterpene accumulation. Analysis of differentially expressed transcription factors identified one basic helix-loop-helix protein (TgbHLH95) and one basic leucine zipper protein (TgbZIP44) as potential *TgGPPS* regulators. TgbHLH95 showed significant transactivation of the *TgGPPS* promoter, and its transient overexpression in tobacco leaves led to monoterpene accumulation, whereas TgbZIP44 directly bound to an ACGT-containing element in the *TgGPPS* promoter, as determined by yeast 1-hybrid test and electrophoretic mobility shift assay. Bimolecular fluorescence complementation, firefly luciferase complementation imaging, co-immunoprecipitation, and GST pull-down assays confirmed a direct protein–protein interaction between TgbHLH95 and TgbZIP44 in vivo and in vitro, and in combination these proteins induced the *TgGPPS* promoter up to 4.7-fold in transactivation assays. These results indicate that a TgbHLH95/TgbZIP44 complex activates the *TgGPPS* promoter and upregulates terpene biosynthesis in xiangfei nuts after harvest, thereby contributing to its aroma.

Introduction

Xiangfei (*Torreya grandis*) is an evergreen coniferous tree belonging to the yew family (Taxaceae), subgenus *Torreya*.

Xiangfei nuts have high nutritional value and potential health benefits attributed to a myriad of bioactive compounds, including sciadonic acid, tocopherol, squalene, and

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β -sitosterol (Suo et al. 2019). Roasted xiangfei nuts are valued by consumers for their unique flavor and aroma (Hu et al. 2022), with the latter being an important index for xiangfei nut quality evaluation. The desirable aroma results from a series of coordinated biosynthetic processes that occur under appropriate temperature and humidity conditions during postharvest ripening over a period of 10 to 15 days, resulting in a reduction in astringency, nutrient conversion, and aroma formation (Zhang et al. 2020b; Hu et al. 2022; Song et al. 2022). However, with the rapid development of the xiangfei industry and the associated prioritization of output, postharvest quality has declined, notably a reduction in the unique nut aroma (Li et al. 2005). To address this problem, there is interest in understanding the biosynthetic and regulatory processes that govern aroma formation during the ripening process.

Aroma is a major factor in the sensory quality of fruit and influences food flavor, thereby strongly affecting consumer preferences (El Hadi et al. 2013). Common aroma components in fruits and seeds belong to several chemical groups, including terpenes, esters, alcohols, aldehydes, ketones, and sulfur compounds, the profiles of which vary depending on the variety, developmental stage, postharvest management, and storage conditions (Schwab et al. 2008). The corresponding biosynthetic pathways include terpenoids, fatty acids, amino acids, and carbohydrates (Aragüez et al. 2013). Most studies of the regulation of aroma compounds in fruits have focused on horticulturally important fruits, and there are far fewer equivalent studies of aroma substances in woody nuts. Indeed, those that have been reported for nuts have mainly focused on differences in substance composition and content, and there are only a few reports describing the gene regulation related to nut aroma (Valdés García et al. 2021). In our previous research, 5 major types of aromatic components were identified in xiangfei nuts after harvest: terpenes, aldehydes, alcohols, alkanes, and esters. Among these, terpenes constituted the main aroma compounds during the postharvest ripening stage, accounting for 56% to 87% (Hu et al. 2022).

Volatile terpenoids, which include isoprene (C₅), monoterpenes (C₁₀), and sesquiterpenes (C₁₅), are the most numerous and diverse class of secondary metabolites (Nagegowda 2010). In plants, these C₅-isoprene building units are generated through either the cytoplasmic mevalonate (MVA) pathway or the plastidial 2-C-methyl-D-erythritol 4-phosphate (MEP) pathway. Volatile sesquiterpenes are produced by the MVA pathway, whereas volatile hemiterpenes, monoterpenes, and diterpenes are produced by the MEP pathway (Vranová et al. 2013). Numerous plants have been identified that contain the genes for these 2 pathways' enzymes, e.g. *Arabidopsis* (*Arabidopsis thaliana*) (Hong et al. 2012), tomato (*Solanum lycopersicum*; Zhou and Pichersky 2020), wormwood (*Artemisia annua*; Lv et al. 2016), grape (*Vitis vinifera*; Duchêne et al. 2009), and apple (*Malus × domestica*; Nieuwenhuizen et al. 2013). Interestingly, the identified key terpene biosynthetic genes are different in different species.

For example, the identified key genes for the biosynthesis of grape terpenes encode a 1-deoxy-D-xylulose-5-phosphate synthase (DXS; *VviDXS*) and a terpene synthase (TPS; *VviTPSS2*) (Battilana et al. 2009; Martin et al. 2010), while in apple peel, the identified α -farnesene biosynthesis genes are a 3-hydroxy-3-methylglutaryl coenzyme A reductase (HMGR; *MdHMGR*) and an α -farnesene synthase (AFS; *MdAFS*) (Pechous and Whitaker 2004). Furthermore, biosynthesis of terpenes is affected by many environmental factors, and the biosynthetic genes can respond differently to different environmental factors. For example, UV-B can influence the production of α -farnesene, linalool, and nerolidol in peach (*Prunus persica*) fruit by regulating the terpene biosynthesis genes *PpTPS1/2/3*, respectively (Liu et al. 2017). Notably, the biosynthetic genes involved in terpene biosynthesis in post-harvested xiangfei nuts remain to be identified.

The expression of multiple genes in metabolic pathways is regulated by transcription factors (TFs), which makes them suitable candidates for pathway engineering (Iwase et al. 2009). The synthesis of plant terpenes is known to be regulated by many TFs including members of the APETALA2/ethylene-responsive factor (AP2/ERF), basic-helix-loop-helix (bHLH), v-myb avian myeloblastosis viral oncogene homolog (MYB), NAM, ATAF1/2, CUC1/2 (NAC), WRKY, and basic leucine zipper (bZIP) families (Supplemental Table S1; Xu et al. 2019). For example, CitAP2.10 transcriptionally activates the *CsTPS1* gene and regulates the biosynthesis of the sesquiterpene valencene in sweet orange (*Citrus sinensis*; Shen et al. 2016). The *A. thaliana* bHLH TF MYC2 interacts with DELLA (Asp-Glu-Leu-Leu-Ala motif containing) proteins and regulates the expression of sesquiterpene synthase *TPS21/11* and mediate jasmonic acid-induced sesquiterpene (E)- β -caryophyllene production (Hong et al. 2012). As another example, AaNAC1 can bind to the *AaADS* promoter and upregulate its expression to increase artemisinin content and enhance drought stress tolerance (Lv et al. 2016).

In our previous report, terpenes were identified as the main aroma compounds in xiangfei nuts after harvest, and their content was greatly affected by temperature and humidity (Hu et al. 2022). However, biosynthetic genes and regulatory mechanisms underlying their generation during the postharvest ripening stage remain unclear. This study focused on the biosynthesis of terpenes after harvest with 2 goals: (i) screening and functional validation of key genes for terpene biosynthesis and (ii) analysis of transcriptional regulatory mechanisms of the related biosynthetic genes. We present insights into aroma production to be employed for improving the ripening quality of xiangfei and other woody nuts.

Results

TgGPPS expression positively correlates with terpene contents in harvested xiangfei (*T. grandis*) nuts

Terpenes were found to be the predominant aroma compounds in our previous study that evaluated the aroma components in raw xiangfei nuts throughout the postharvest

ripening stage, and their levels significantly increased at high humidity (Hu et al. 2022). In this current study, the aroma composition of xiangfei nuts changed at various postharvest ripening stages, and 56 volatiles were found using gas chromatography-MS (GC-MS) analysis (Fig. 1A and Supplemental Fig. S1). As expected, terpenes were the main aroma compounds, and the concentrations were low from days 0 to 4, and increased significantly from days 4 to 8, then maintained stable until day 12, reaching a maximum of 96.53 $\mu\text{g/g}$ on day 12 after harvest (Fig. 1B).

As shown in Fig. 2A, putative genes involved in postharvest terpene biosynthesis were identified by combining transcriptome data with reverse transcription quantitative PCR (RT-qPCR) data from days 4 and 8 (Fig. 2A). Transcript analysis revealed the upregulation of biosynthetic genes from the MVA pathway (AACT, HMGS, HMGR, MVK, PMK, and MVD), among which 8 genes were annotated to AACT, 11 genes annotated to HMGS, 27 genes annotated to the HMGR, 5 genes annotated to MVK, 7 genes annotated to PMK, and 3 genes annotated to MVD. DXS, DXR, MCT, CMK, MDS, HDS, HDR, GPPS, and TPS are the key biosynthetic genes in the MEP pathway. Here, we found 13 genes annotated to DXS, 8 genes annotated to DXR, 5 genes annotated to MCT, 3 genes annotated to CMK, 12 genes annotated to MDS, 5 genes annotated to HDS, 5 genes annotated to HDR, 8 genes annotated to GPPS, and 36 genes annotated to TPS (Supplemental Table S2). Highly expressed terpene biosynthetic genes were identified based on the fragments per kilobase of exon per million mapped fragments (FPKM) values, and a heatmap was created (Fig. 2A). A correlation analysis between the FPKM value and the total terpene levels revealed that a geranyl diphosphate synthase (GPPS) unigene (cluster_contig30179), subsequently called *TgGPPS*, was highly and positively correlated with the total terpene content (Fig. 2A). To confirm the transcriptome analysis, RT-qPCR analysis was performed and confirmed that the expression level of *TgGPPS* was significantly higher during days 4 to 8 after harvest (Fig. 2B), which was consistent with the total terpene content ($R = 0.83$, $P < 0.001$; Fig. 2C), consistent with a role for *TgGPPS* in terpene biosynthesis during the postharvest ripening stage.

Effect of overexpressing *TgGPPS* on terpene production in tobacco (*Nicotiana tabacum*) leaves and tomato (*S. lycopersicum*) fruit

A comparison of several GPPS sequences revealed that *TgGPPS* included 2 important aspartate-rich (Asp-rich) motifs of the consensus sequence DD(X)₂₋₄D (where X may be any amino acid) (Supplemental Fig. S2; Wang and Ohnuma 2000). A phylogenetic tree of *TgGPPS* and 21 other plant GPPS proteins from selected gymnosperm and angiosperm species (Fig. 3A) showed that *TgGPPS* clustered with the gymnosperm *Picea abies* PaGPPS, which has been described as a chloroplast-localized regulator of terpene synthesis (Fig. 3A; Schmidt and Gershenzon 2007, 2008). A subcellular

localization assay was performed in *N. benthamiana* leaves, which indicated that *TgGPPS* was also located in the chloroplasts (Fig. 3B), further suggesting a role for *TgGPPS* in terpene biosynthesis.

To confirm *TgGPPS* function, we used a highly efficient tobacco leaf transient expression assay. *Agrobacterium tumefaciens* containing *TgGPPS*-SK or empty SK vector was injected into tobacco (*N. tabacum*) leaves, and volatile substances were detected by GC-MS 1 wk post transfection. Compared with the negative control, *TgGPPS* overexpressing tobacco leaves exhibited 1.6-fold higher monoterpene levels (Fig. 3C and Supplemental Fig. S3). Major monoterpenes showing an increase included α -pinene and myrcene present at trace levels in the control (Fig. 3D). Moreover, α -terpinene and limonene showed 4.8- and 4.4-fold increases in overexpressing leaves, respectively (Fig. 3D).

We also performed a heterologous stable transgenic experiment in tomato (cv. *MicroTom*) plants. Five independent stable transgenic lines were obtained, of which the 3 showing the highest *TgGPPS* expression levels were selected for volatile analysis and RT-qPCR verification. Gene expression analysis confirmed that *TgGPPS* was overexpressed in the fruit (Fig. 3E), and compared with the wild type (WT), the monoterpene levels were 1.3 to 3.3 times higher in 3 independent transgenic plants. The abundance of α -pinene and β -pinene showed particularly significant increases (Fig. 3F and Supplemental Fig. S4). These results confirmed that *TgGPPS* is involved in terpene biosynthesis in *xiangfei* nuts during the postharvest ripening stage (Fig. 3).

TgbHLH95 functions as a positive indirect regulator in terpene biosynthesis

Following the confirmation of *TgGPPS* as a key gene in terpene biosynthesis, we next investigated mechanisms of its regulation. Differentially expressed TFs were identified using the RNA-seq database to further explore transcriptional regulatory mechanisms, and 15 TFs were identified with an expression pattern highly similar to that of *TgGPPS* (correlation index > 0.6 on days 4 to 8 of postharvest ripening) (Fig. 4A). The 15 TF sequences, which collectively were annotated as belonging to the AP2/ERF, MYB, bHLH, bZIP, MADS, and GTE families, were individually inserted into the pGreenII0029 62-SK vector, while the *TgGPPS* promoter sequence was cloned into the pGreen II 0800-LUC vector. The results of a dual-luciferase assay in *N. benthamiana* leaves (threshold was set as 2) suggested that 3 TFs activated the *TgGPPS* promoter: *TgbHLH1*, *TgbHLH2*, and *TgbHLH95*, with *TgbHLH95* displaying the strongest activation effect, of approximately 3.9-fold (Fig. 4B).

TgbHLH95, which showed the strongest activation effect, was selected as a candidate to investigate terpene biosynthesis regulation. First, subcellular localization analysis suggested that *TgbHLH95* was a nucleus-localized protein (Fig. 4C) consistent with the proposed TF function. Besides, a yeast 1-hybrid test was used to analyze the interactions between *TgbHLH95* and the *TgGPPS* promoter. The growth of

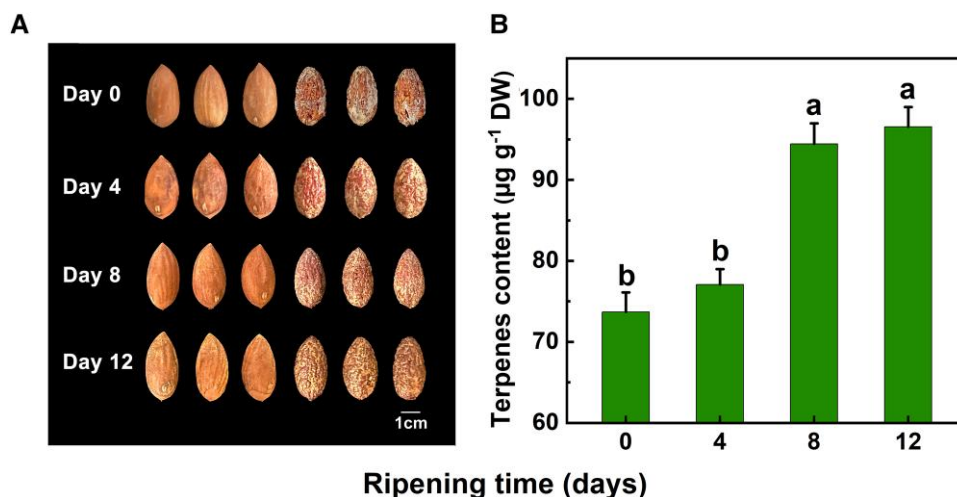


Figure 1. Changes in terpene production in xiangfei (*Torreya grandis*) nuts during the postharvest ripening stage. **A)** Nuts were collected at different postharvest ripening times. Images were digitally extracted for comparison. Scale bars = 1 cm. **B)** Total terpene content of xiangfei nuts during the postharvest ripening stage. Error bars represent \pm SE based on 3 biological replicates. The different letters indicate significant differences ($P < 0.05$) during the postharvest ripening stage by Duncan's multiple range test. DW, dry weight.

yeast containing TgbHLH95 was inhibited as was the empty vector, suggesting that TgbHLH95 binds indirectly to the *TgGPPS* promoter (Fig. 4D). Next, RT-qPCR data indicated that TgbHLH95 expression pattern was similar to that of *TgGPPS*, as both were significantly upregulated on days 4 to 8 after harvest (Fig. 4E). By transiently overexpressing *TgbHLH95* in tobacco leaves, the monoterpene content greatly increased, from 3.35 $\mu\text{g/g}$ toward 3.92 $\mu\text{g/g}$ (Fig. 4F). Collectively, these results revealed that TgbHLH95 functioned as a positive indirect regulator activating *TgGPPS* expression to promote terpene production.

TgbZIP44 acts as a direct regulator in terpene biosynthesis

The gene expression results, the dual-luciferase assay, and the transient overexpression validation in tobacco leaves indicated that TgbHLH95 transactivated the *TgGPPS* promoter, but did not directly bind to it. Therefore, it is essential to discover other TFs that can interact with TgbHLH95, which may serve as a bridge in connecting TgbHLH95 and *TgGPPS*. A yeast 1-hybrid assay was used to analyze the interactions between other differentially expressed TFs and *TgGPPS* promoter. The results revealed that TgERF2, TgRAV3, and TgbZIP44 showed normal growth under 200 ng ml^{-1} aureobasidin A (AbA) compared with the empty vector (Fig. 5A). This indicated that TgERF2, TgRAV3, and TgbZIP44 could directly bind to the *TgGPPS* promoter, suggesting that the activation by TgbHLH95 might occur through a transcriptional complex with TgERF2, TgRAV3, and TgbZIP44. Accordingly, we next investigated whether TgbHLH95 and TgERF2, TgRAV3 and TgbZIP44 are synergistically involved in terpene biosynthesis. In a dual-luciferase assay, TgbHLH95 and TgbZIP44 in combination showed the strongest induction of the *TgGPPS* promoter, with a LUC/REN ratio of 4.68, compared with transfection of TgbHLH95 (1.21-fold activation) or TgbZIP44

(0.64-fold activation) in combination with the empty vector (Fig. 5B). The combination of TgbHLH95 and TgRAV3 showed a relatively lower additive effect (3.64-fold activation), and TgbHLH95 with TgERF2 showed no additive effect (Fig. 5B). Consequently, TgbZIP44 was chosen for further study. The expression pattern of *TgbZIP44* via RT-qPCR analysis was similar to that of *TgGPPS*, which were significantly upregulated on days 4 to 8 (Fig. 5C). In vitro experiments using electrophoretic mobility shift assay (EMSA) revealed that TgbZIP44 could directly recognize and bind to the ACGT-containing elements in the *TgGPPS* promoter (Fig. 5D). When the specific ACGT element was mutated by AAAA sites, the putative TgbZIP44 protein binding was eliminated. Meantime, the binding affinity of the biotinylated probe was substantially reduced as the concentration of the competitor (cold probe) increased (Fig. 5D). In addition, subcellular localization of TgbZIP44 showed a strong fluorescent signal in the nucleus, indicating TgbZIP44 as a nucleus-localized protein (Fig. 5E). Their colocalization implied that TgbHLH95 and TgbZIP44 might form a protein–protein interaction in the nucleus. Collectively, the combination of TgbHLH95 and TgbZIP44 was chosen for further protein–protein interaction assays.

A TgbHLH95/TgbZIP44 complex activates the *TgGPPS* promoter

To test if TgbHLH95 interacts with TgbZIP44, transient bimolecular fluorescence complementation (BiFC) assays were conducted accordingly. *N. benthamiana* leaves cotransformed with PHR2-YC, and SPX4-YN (positive controls) showed a YFP fluorescent signal in the cytoplasm, in contrast to those expressing the combination of a single construct (i.e. TgbHLH95-YFP^N + YFP^C, TgbHLH95-YFP^C + YFP^N, TgbZIP44-YFP^N + YFP^C, and TgbZIP44-YFP^C + YFP^N) and the corresponding bidirectional empty vector (negative controls), which did not produce any YFP signal (Fig. 6A). TgbHLH95-YFP^C cotransformed with

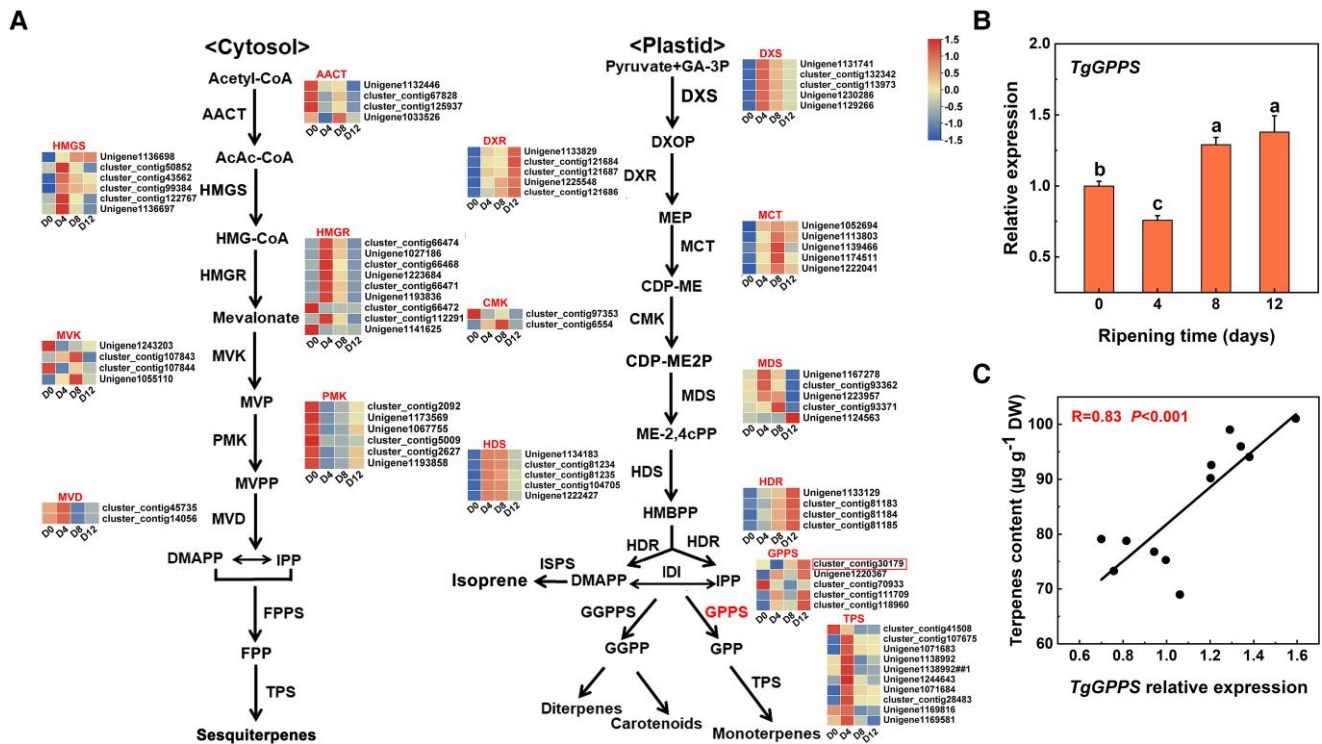


Figure 2. Gene expression and FPKM analysis of terpene biosynthetic genes in xiangfei nuts during the postharvest ripening stage. **A**) Biosynthesis of terpenoid volatile organic compounds and the FPKM values for terpenoid biosynthetic genes. The cytosolic MVA pathway produces sesquiterpenes. The plastidial MEP pathway produces monoterpenes, diterpenes, and volatile carotenoid derivatives. **B**) The *TgGPPS* expression pattern in xiangfei nuts during the postharvest ripening stage. Error bars represent *se* based on 3 biological replicates. The different letters indicate significant differences ($P < 0.05$) during the postharvest ripening stage by Duncan's multiple range test. **C**) Correlation analysis between the total terpene content and *TgGPPS* expression. IPP, isopentenyl pyrophosphate; DMAPP, dimethylallyl pyrophosphate; AACT, acetyl-CoA acetyltransferase; HMGS, hydroxymethylglutaryl-CoA synthase; HMGR, hydroxymethylglutaryl-CoA reductase; MVK, mevalonate kinase; PMK, phosphomevalonate kinase; MVD, mevalonate diphosphate decarboxylase; DXS, 1-deoxy-D-xylulose 5-phosphate synthase; DXR, 1-deoxy-D-xylulose 5-phosphate reductoisomerase; MCT, 2-C-methyl-D-erythritol 4-phosphate cytidyltransferase; CMK, 4-(cytidine 50 -diphospho)-2-C-methyl-D-erythritol kinase; MDS, 2-C-methyl-D-erythritol 2,4- cyclodiphosphate synthase; HDS, (E)-4-hydroxy-3-methylbut-2-enyl diphosphate synthase; HDR, (E)-4-hydroxy-3-methylbut-2-enyl diphosphate reductase; GGPPS, geranylgeranyl pyrophosphate synthase; FPPS, farnesyl pyrophosphate synthase; DW, dry weight.

TgbZIP44-YFP^N or *TgbHLH95-YFP^N* cotransformed with *TgbZIP44-YFP^C* displayed a nuclear YFP fluorescent signal (Fig. 6A), indicating that *TgbHLH95* could interact with *TgbZIP44* in the nucleus. The interaction between *TgbHLH95* and *TgbZIP44* was also examined using a luciferase complementation imaging (LCI) assay to confirm the results obtained by BiFC. When nLUC-*TgbHLH95* and cLUC-*TgbZIP44* or nLUC-*TgbZIP44* and cLUC-*TgbHLH95* were coinfiltrated into *N. benthamiana* leaves, a high LUC luminescence signal showed (Fig. 6B), whereas the negative controls had no detectable LUC activity (the combinations of nLUC + cLUC, nLUC + cLUC-*TgbHLH95*, nLUC-*TgbHLH95* + cLUC, nLUC + cLUC-*TgbZIP44*, nLUC-*TgbZIP44* + cLUC) (Fig. 6B). A co-immunoprecipitation (Co-IP) test was also used to confirm the interaction between *TgbHLH95* and *TgbZIP44*. *N. benthamiana* leaf protein samples were utilized for immunoprecipitation using an anti-GFP antibody. After that, anti-GFP and anti-Flag antibodies were used to immunoblot the eluted protein. The outcome demonstrated that the *TgbZIP44-GFP* fusion protein interacted with *TgbHLH95* in *N. benthamiana*

leaves in vivo, but not the GFP negative control (Fig. 6C). An in vitro GST pull-down test was utilized to further validate the interaction between the *TgbHLH95* and *TgbZIP44* proteins. The recombinant *TgbZIP44-His* protein was incubated in vitro with the recombinant *TgbHLH95-GST* protein and the GST control. Anti-GST and anti-His antibodies were used to immunoblot the protein after it had been eluted using GST-tag beads. The results showed that *TgbHLH95-GST* was pulled down, but GST alone was not. Therefore, *TgbZIP44-His* interacted directly with the *TgbHLH95-GST* protein (Fig. 6D). Collectively, BiFC, LCI, Co-IP, and GST pull-down assays all revealed that *TgbHLH95* and *TgbZIP44* formed a protein–protein complex in vivo and in vitro (Fig. 6).

Discussion

TgGPPS is a key terpene biosynthetic gene in xiangfei (*T. grandis*) nuts after harvest

GPPS (EC 2.5.1.1) is the main enzyme responsible for the production of the monoterpene precursor GPP

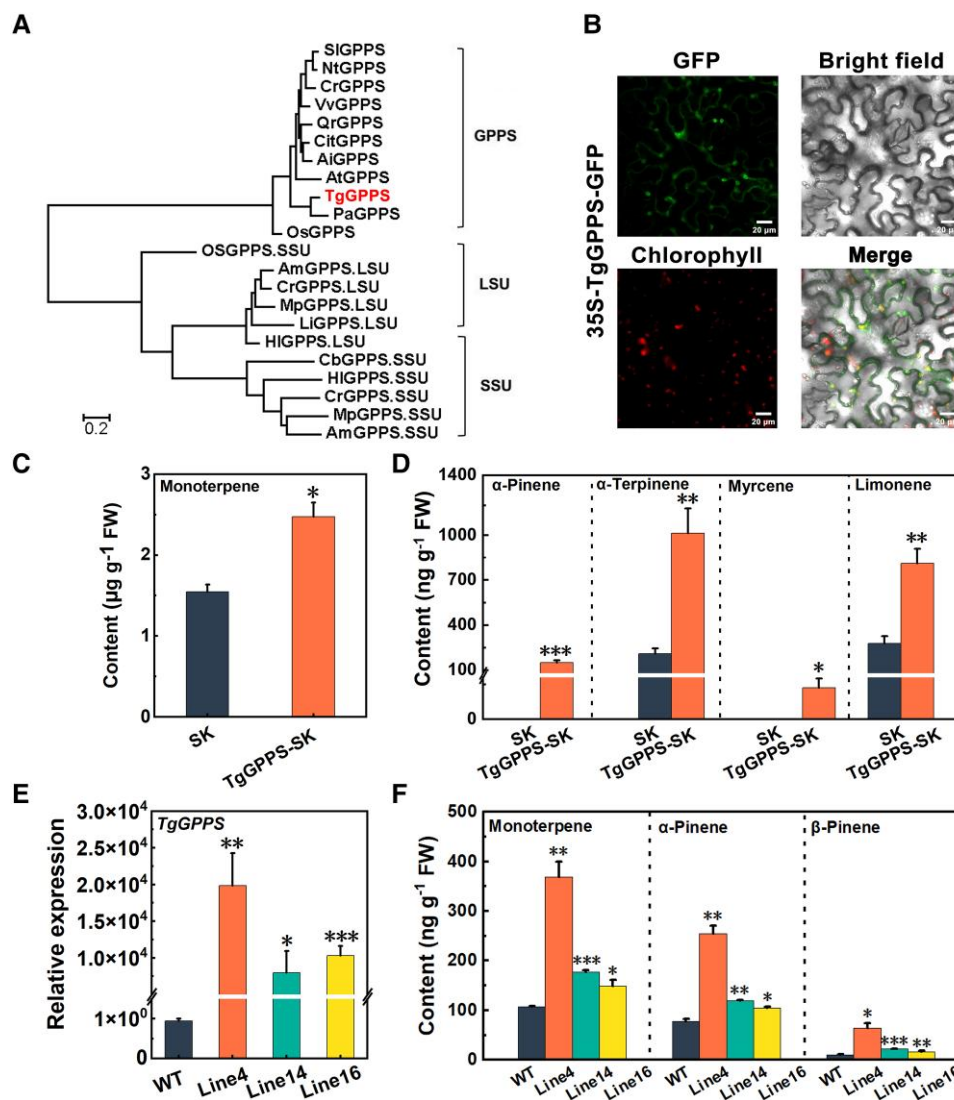


Figure 3. Functional characterization of *TgGPPS*. **A)** Phylogenetic analysis of *TgGPPS* and GPPS proteins from other species. All the sequences were obtained from the NCBI database. Accession numbers for these GPPSs are given in Materials and methods section. **B)** Subcellular localization analysis of *TgGPPS*. The GFP fluorescence indicates *TgGPPS* localization and the red fluorescence indicates chloroplast fluorescent protein (CFP). Scale bars = 20 μm . **C)** Transient overexpression of *TgGPPS* increases total monoterpene content in tobacco leaves. GC-MS identified 10 monoterpene components in tobacco leaves: α -pinene, β -pinene, myrcene, limonene, γ -terpinene, α -terpinene, terpinolene, linalool, geraniol, and menthol. **D)** Levels of α -pinene, α -terpinene, myrcene, and limonene in tobacco leaves transiently overexpressing *TgGPPS* or in leaves transformed with the empty vector (SK) as a control. The relative content was calculated according to the peak area of the internal standard (2-octanol). **E)** Relative expression of *TgGPPS* in transgenic tomato fruit. **F)** Overexpressing *TgGPPS* induces monoterpene synthesis in tomato. GC-MS detected 4 monoterpene components in tomato fruit: α -pinene, β -pinene, limonene, and linalool. SEs are all obtained from 3 biological replicates, and the significant differences are indicated with asterisks using Student's test (* $P < 0.05$, ** $P < 0.01$ and *** $P < 0.001$). FW, fresh weight.

(Dudareva et al. 2006), and studies in several plant species have shown that GPPS can form homodimers or heterodimers (Rai et al. 2013). Heterodimeric GPPS has been identified in the leaves of the angiosperms peppermint (*Mentha piperita*), snapdragon (*Antirrhinum majus*), and hops (*Humulus lupulus*) (Bouvier et al. 2000; Schmidt and Gershenzon 2008; Wang and Dixon 2009) and consist of large subunits (LSUs) and small subunits (SSU). For example, LSU and SSU in heterogenous GPPS in peppermint have no catalytic activity alone, while the interaction

between the 2 subunits leads to the formation of active GPPS. Structural studies also showed that LSU acts as a catalytic unit, while SSU acts as a regulatory unit (Chang et al. 2010). In gymnosperms, only homodimer GPPS complexes have been identified to be involved in terpene biosynthesis, such as PaGPPS in *Picea abies* (Schmidt and Gershenzon 2007). This current study showed a phylogenetic clustering of *TgGPPS* and PaGPPS, consistent with its involvement in terpene biosynthesis (Fig. 3A). GPPS has also been located in the nongreen plastids of the active

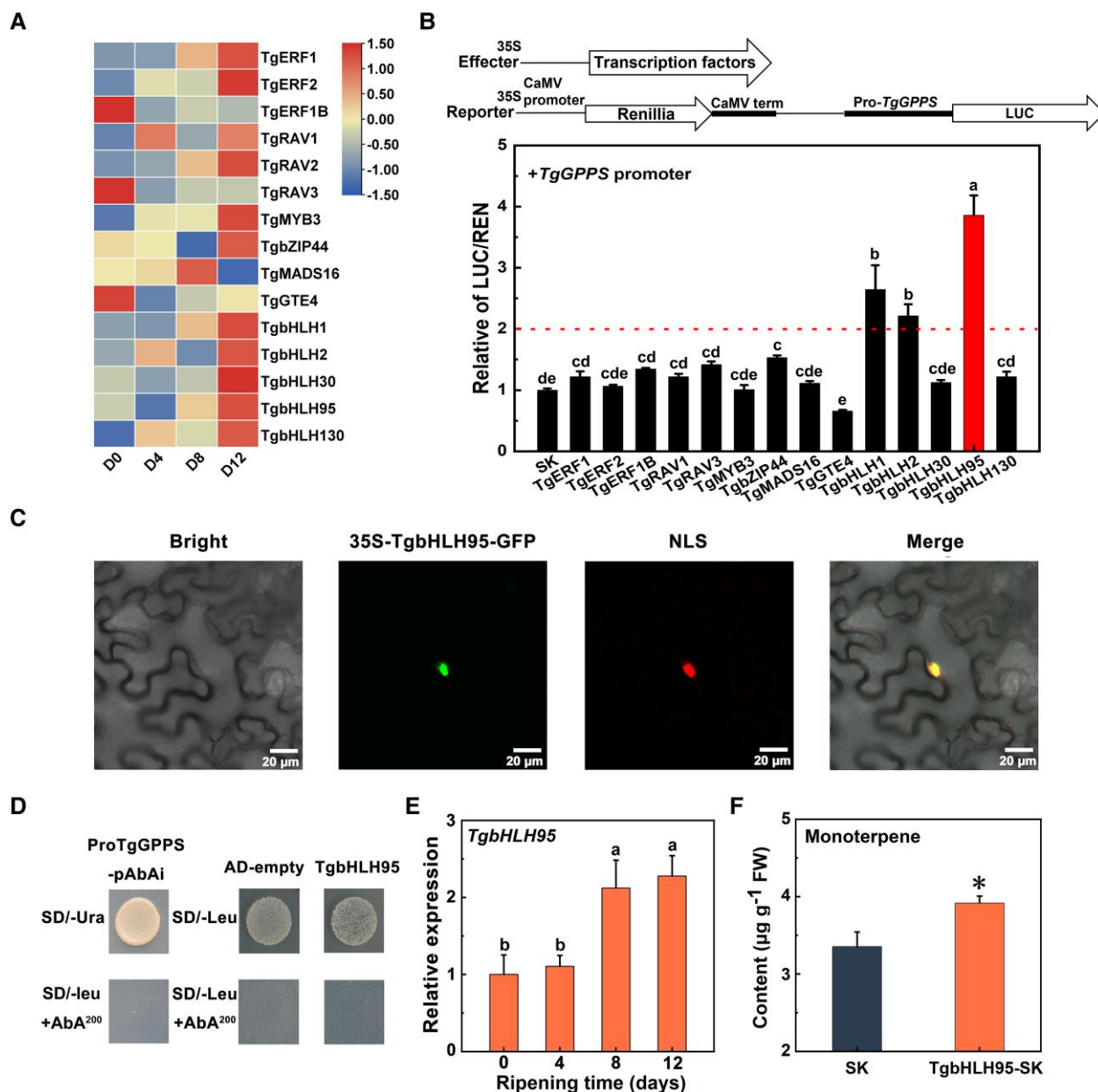


Figure 4. Functional characterization of *TgbHLH95*. **A**) Heatmap analysis of differentially expressed TFs. **B**) The regulatory effect of TFs on the *TgGPPS* promoter, the LUC/REN value of the empty SK vector on the *TgGPPS* promoter was set to 1.0, and \pm SE values were calculated using 3 biological replicates. Different letters indicate significant differences ($P < 0.05$) during the postharvest ripening stage by Duncan's multiple range test. **C**) Subcellular localization analysis of *TgbHLH95*. The fluorescent signals were detected in tobacco cells cotransfected with *TgbHLH95*-GFP and the nuclear RFP-NLS marker. Scale bar = 20 μ m. **D**) Yeast 1-hybrid analysis of *TgbHLH95* binding capacity to the *TgGPPS* promoter, with the empty vector pGADT7 (AD) used as a negative control. **E**) The *TgbHLH95* expression pattern in xiangfei nuts during the postharvest ripening stage. Error bars represent \pm SE based on 3 biological replicates. The different letters indicate significant differences ($P < 0.05$) during the postharvest ripening stage by Duncan's multiple range test. **F**) Transient overexpression of *TgbHLH95* increases total monoterpene content in tobacco leaves. Error bars represent \pm SE based on 3 biological replicates. Significant differences are indicated with asterisks using Student's *t* test (* $P < 0.05$).

MEP pathway and in chloroplasts of photosynthetic cells (Bouvier et al. 2000). Subcellular localization analysis indicated that *TgGPPS* was localized in chloroplasts (Fig. 3B), further revealing its role in terpene biosynthesis. In addition, the *TgGPPS* transcript levels were positively correlated with the total terpene content after harvest (Figs. 1

and 2). Furthermore, transient heterologous overexpression of *TgGPPS* in tobacco (*N. tabacum*) leaves or tomato (*S. lycopersicum*) fruit resulted in a significant increase in monoterpene levels (Fig. 3, C to F), indicating that *TgGPPS* contributes to terpene biosynthesis in nuts after harvest.

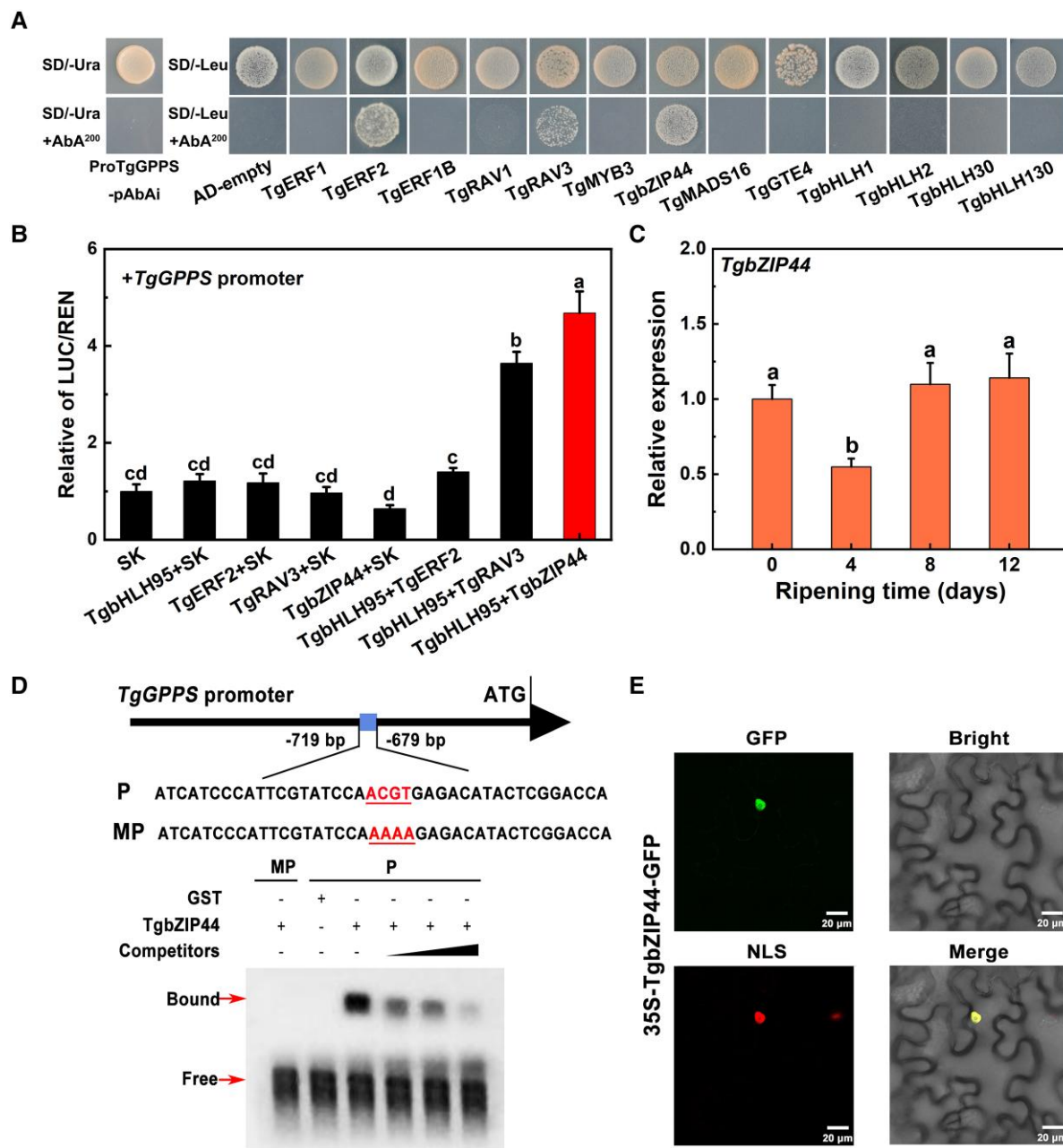


Figure 5. Functional characterization of *TgbZIP44*. **A**) Yeast 1-hybrid analysis of other TFs binding capacity to the *TgGPPS* promoter, with the empty vector pGADT7 (AD) used as a negative control. **B**) Synergistic transactivation effect of different TFs on the *TgGPPS* promoter. The LUC/REN ratio of the empty vector (SK) plus the promoter was used as a calibrator (value set to 1.0). Error bars represent \pm SE based on 3 biological replicates. The different letters indicate significant differences ($P < 0.05$) by Duncan's multiple range test. **C**) The *TgbZIP44* expression pattern in xiangfei nuts during the postharvest ripening stage. Error bars represent \pm SE based on 3 biological replicates. The different letters indicate significant differences ($P < 0.05$) during the postharvest ripening stage by Duncan's multiple range test. **D**) In vitro binding ability of *TgbZIP44* to the *TgGPPS* promoter performed by EMSA. The presence (+) or absence (–) of specific probes is marked. The black arrows become larger from left to right indicating an increase in the concentration of competitive probes. GST protein was used as a negative control. **E**) Subcellular localization analysis of *TgbZIP44*. The fluorescent signals were detected in tobacco cells cotransfected with *TgbZIP44*-GFP and the nuclear RFP-NLS marker. Scale bar = 20 μ m.

TgbHLH1, *TgbHLH2*, and *TgbHLH95* are positive regulators of terpene biosynthesis in xiangfei nuts after harvest

There are many reports describing the transcriptional regulation of terpene biosynthesis, but the regulatory mechanisms

acting on GPPS are less well studied. One such study, performed in spearmint (*M. spicata*), showed that MsMYB inhibits *GPPS.LSU* expression to reduce the terpene content (Reddy et al. 2017); however, the role of other TFs on GPPS expression remains to be elucidated. In the current study, 3

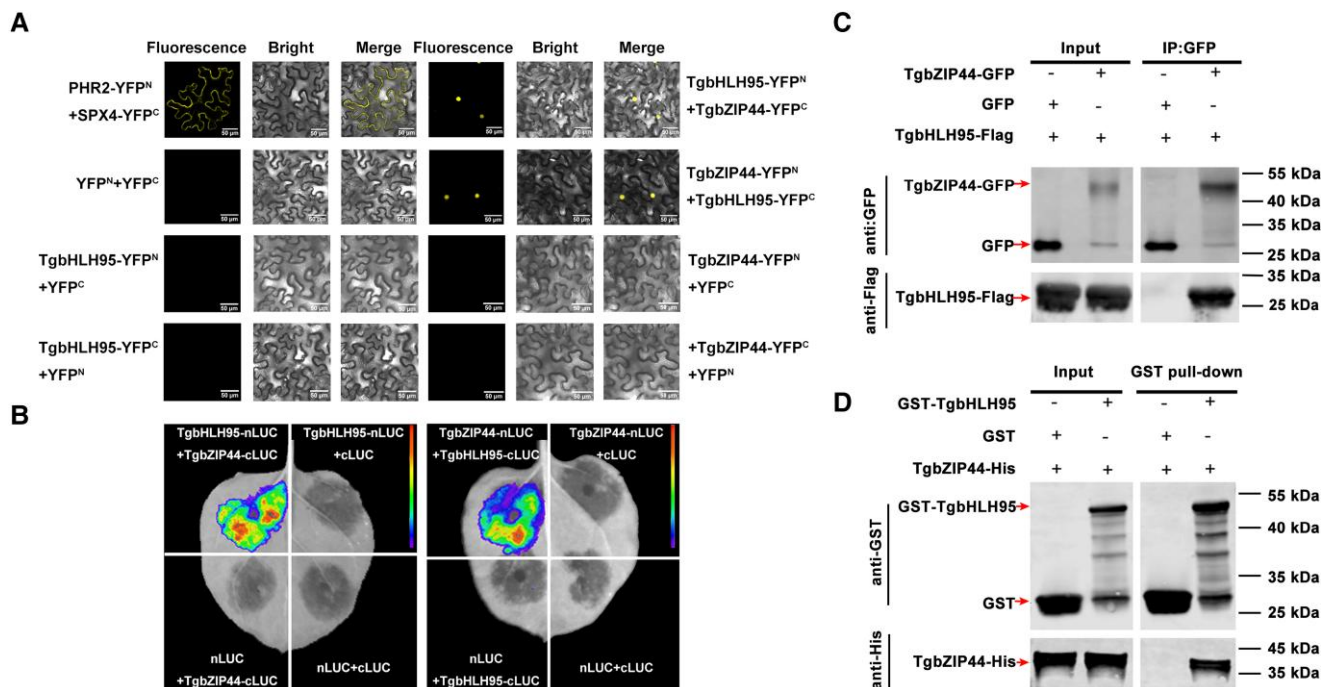


Figure 6. TgbHLH95 protein interacts with TgbZIP44. **A**) BiFC analysis of the interaction between TgbHLH95 and TgbZIP44. The pairs of fusion proteins tested were TgbHLH95-YFP^N + TgbZIP44-YFP^C and TgbZIP44-YFP^N + TgbHLH95-YFP^C. PHR2-YFP^N and SPX4-YFP^C were used as the positive control; YFP^N and YFP^C were used as negative controls. Nuclear location signal (NLS) indicates the position of the nucleus. Scale bars = 50 μ m. **B**) LCI analysis of the interaction between TgbHLH95 and TgbZIP44. Tobacco leaves were infiltrated with *Agrobacterium* strain GV3101 harboring different constructs. The luminescence signals of the infiltrated areas were measured 48 h after infiltration. The experiment was repeated at least 3 times. **C**) Co-IP assay of TgbHLH95 and TgbZIP44. TgbHLH95-Flag plus TgbZIP44-GFP proteins were immunoprecipitated with an anti-GFP antibody and immunoblotted with anti-GFP and anti-Flag antibodies. **D**) Pull-down assay detection of the interaction of TgbHLH95 and TgbZIP44. The recombinant TgbHLH95-GST or GST was incubated with TgbZIP44-His. Blots were probed with anti-GST and anti-His antibody.

bHLH TFs were identified that exerted significant activation effects on the *TgGPPS* promoter. TgbHLH95 showed the highest activation effect, gene expression, subcellular localization analysis, and transient overexpression assays in tobacco indicated that it was indeed a positive regulator of terpene biosynthesis, although it did not bind directly to the *TgGPPS* promoter in Y1H assay (Fig. 4). The expression analysis also revealed that TgbHLH1 and TgbHLH2 expression increased significantly on days 4 to 8 after harvest (Supplemental Fig. S5). Furthermore, tobacco overexpression assays showed that overexpression of *TgbHLH1* significantly increased terpene content, although overexpression of *TgbHLH2* did not (Supplemental Fig. S6), indicating that TgbHLH1 also indirectly regulates *TgGPPS*, while TgbHLH2 is not involved in terpene biosynthesis. This study suggested that TgbHLH95 and TgbHLH1 activate *TgGPPS* expression to promote terpene production.

The proteins in the plant bHLH TF family, the second-largest TF family, have a conserved bHLH domain that has a length of about 60 amino acids (Feller et al. 2011). Members of the bHLH family have been identified and characterized in many species, such as *A. thaliana* and apple (Pires and Dolan 2010; Mao et al. 2017). These bHLH TFs play important roles in various metabolic pathways,

including hormone synthesis (Qi et al. 2014), stress response (Fan et al. 2014), and metabolite (flavonoid, alkaloid, terpenoid) biosynthesis (Xie et al. 2017). In many plant models, the contribution of bHLH members to terpene biosynthesis has been demonstrated (Hong et al. 2012; Xu et al. 2018). In tomato, the bHLH TF SIMYC1 inhibits sesquiterpene production in stem trichomes while acting as a positive regulator of monoterpene biosynthesis in leaf and stem trichomes (Xu et al. 2018). In peach fruit, PpbHLH1 directly bound to the *PpTPS3* promoter to regulate production of the key aroma compound linalool (Wei et al. 2021). In xiangfei nuts, TgbHLH1 and TgbHLH95 were shown to be involved in post-harvest terpene biosynthesis through indirect regulation of the *TgGPPS* promoter. These studies suggested that bHLH family members play an important role in the regulation of terpene synthesis.

A TgbHLH95/TgbZIP44 transcription complex promotes terpene production via activation of the *TgGPPS* promoter

Terpene production has been linked to a number of TFs, but most of these TFs act alone on individual biosynthetic genes. For example, MdERF3 and MdMYC2 regulate α -farnesene

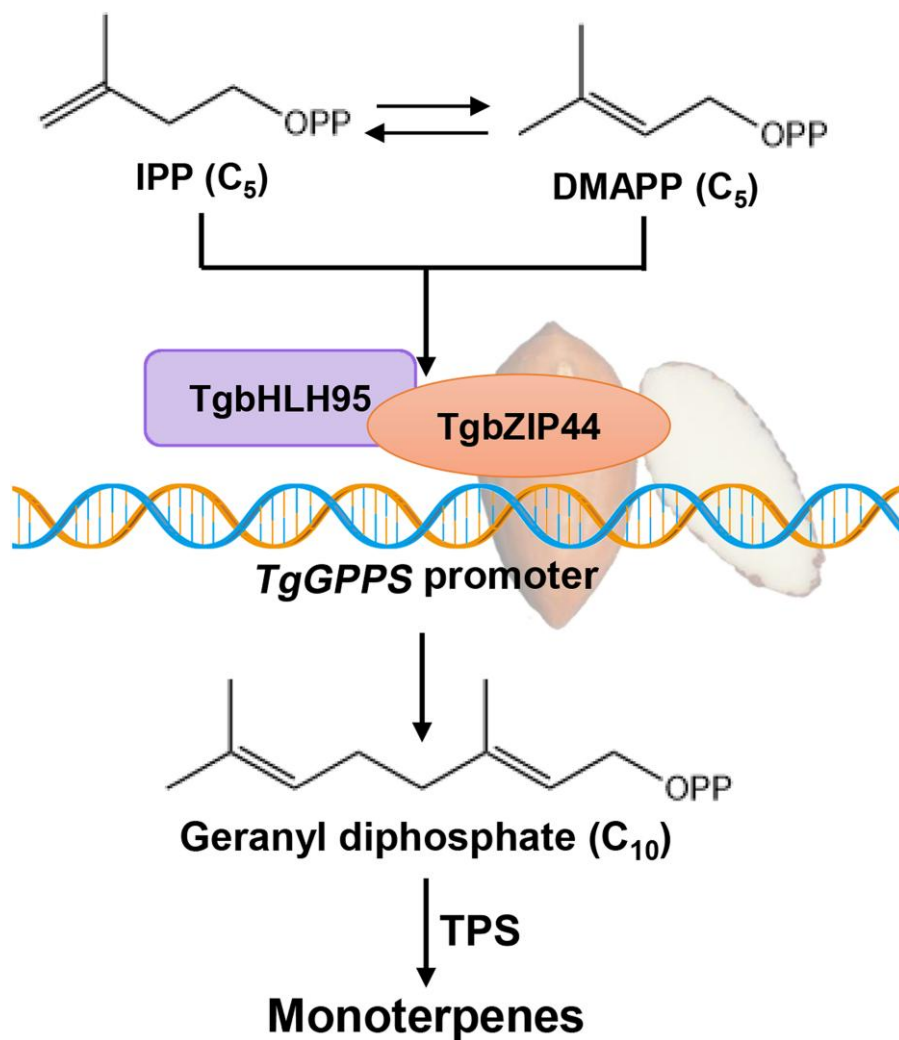


Figure 7. A proposed regulatory model of the TgbHLH95-TgbZIP44 protein complex promoting terpene accumulation via activation of *TgGPPS* expression in xiangfei nuts during the postharvest ripening stage.

synthesis through the activation of expression of the terminal enzyme *MdAFS*. Among these, *MdERF3* can directly bind to the DRE element of the *MdAFS* promoter (Wang et al. 2020). Xu et al. (2017) reported that *MdMYC2* can activate *MdERF3* transcription by binding to the *MdERF3* promoter, i.e. *MdMYC2* can act as an upstream regulator of *MdERF3* and coregulate it to induce α -farnesene synthesis in apple, further expanding this regulatory network. Less is known about the transcriptional complexes that regulate terpene production, although a MYB-bHLH protein complex has been identified (Yang et al. 2020). In flowers of *Freesia hybrida*, *FhMYB21* substantially activated *FhTPS1* expression, while *FhMYC2* showed an inverse expression pattern compared with *FhTPS1*. Upon *FhMYC2* coexpression with *FhMYB21*, *FhMYC2* reduced the activation effect of *FhMYB21* through interactions between the 2 TFs, demonstrating that the MYB-bHLH complex cooperated in the regulation of linalool synthesis. The mechanism of MYB-bHLH complexes in regulating terpene aromatics is conserved in plants such as *A. thaliana* (Yang et al. 2020). This

report shows a bHLH-bZIP transcription complex regulating terpene synthase genes (Fig. 7). TgbHLH95 indirectly activates the *TgGPPS* promoter, while TgbZIP44 binds directly to it, with formation of a TgbHLH95/TgbZIP44 protein complex participating in terpene biosynthesis (Figs. 4 to 6). These findings add to our understanding of the regulatory mechanisms affecting terpene biosynthesis in nuts and provide a foundation for further research into terpene biosynthesis in other woody nuts.

Conclusions

Accumulation of terpenes in xiangfei nuts after harvest is important for nut quality and consumption. Here, based on both in vitro and in vivo assays, a *TgGPPS* gene was cloned and characterized encoding a protein capable of promoting terpene production. TgbHLH95 activated *TgGPPS* transcription indirectly, and TgbHLH95 overexpression promoted the terpene content in tobacco leaves. Y1H and EMSA analysis showed that TgbZIP44 directly bound to the *TgGPPS*

promoter, and protein binding assays further confirmed an interaction between TgbHLH95 and TgbZIP44, with a synergistic effect in combination on the *TgGPPS* promoter to regulate the synthesis of terpenes. These hypotheses, identified in xiangfei nuts, have yet to be examined in other tree nut species. Knowledge of the regulation of volatiles' biosynthesis pathways can facilitate future efforts to enhance nut flavor quality and screen for in breeds or genetically modified plants showing the best aroma composition targeting certain terpenes.

Materials and methods

Plant materials and experimental treatments

Xiangfei (*T. grandis* cv. "Merrillii") nuts were harvested 525 days after full bloom from an orchard in Xinchang, Zhejiang, China. On the day of harvest, the nuts were brought to the laboratory. The arils were manually removed, thoroughly cleaned with clear water, and then allowed to air dry until there was no more moisture present outside the shell. Nuts with uniform size and free of visible wounding were selected and divided into 4 groups. They were kept at controlled temperature conditions ($20\text{ }^{\circ}\text{C} \pm 2\text{ }^{\circ}\text{C}$) and relative humidity (RH) levels ($90\% \pm 2\% \text{ RH}$), and 30 nuts were sampled on days 0, 4, 8, and 12 in 3 biological replicates. After removing the shell, the kernels were frozen in liquid nitrogen and stored at $-80\text{ }^{\circ}\text{C}$ for further studies.

Tobacco (*N. tabacum*) plants for transient overexpression, *N. benthamiana* plants for dual-luciferase assay, and tomato (*S. lycopersicum*, cv. *MicroTom*) plants for stable overexpression were grown in a greenhouse with a light/dark cycle of 16:8 h at $25\text{ }^{\circ}\text{C}$.

Analysis of volatiles by GC-MS

Volatiles were analyzed according to Zhang et al. (2010). Frozen xiangfei kernel tissue (1 g) and whole tomato fruit tissue (5 g) were ground into a powder in liquid nitrogen, which was then transferred to vials containing 5 mL of saturated NaCl solution. Frozen and ground *N. tabacum* leaves (1 g) were transferred to a vial containing 1.5 mL 200 mM EDTA and 1.5 mL 20% (w/v) CaCl_2 solution. Subsequently, 2-octanol (0.8 mg/mL, 10 μL) was added as the internal standard, and then the vials were sealed. Agilent 7890A gas chromatograph and 5975C mass spectrometer linked to a solid-phase microextraction autosampler (Combi PAL, CTC Analytics, Agilent Technologies) homogenized and analyzed the mixture. A fiber coated with 65 μm of polydimethylsiloxane and divinylbenzene from the Supeclo Co. in Bellefonte, Pennsylvania, was used to capture volatiles. On a DB-WAX column (30 m \times 0.25 mm, 0.25 μm phase film thickness, Agilent, USA), volatiles were separated. The flow rate of the helium carrier gas was 1 mL/min. The temperature program began at $40\text{ }^{\circ}\text{C}$ and increased to $100\text{ }^{\circ}\text{C}$ and $245\text{ }^{\circ}\text{C}$ at rates of $3\text{ }^{\circ}\text{C}/\text{min}$ and $5\text{ }^{\circ}\text{C}/\text{min}$, respectively. The column effluent was ionized at an energy of 70 eV with a source temperature of $230\text{ }^{\circ}\text{C}$ and a transfer temperature of $250\text{ }^{\circ}\text{C}$.

Volatiles were discovered by comparing electron ionization mass spectra and retention time data with the NIST Mass Spectral Library (NIST-08 and Flavor). The peak area of the internal standard was used as a base to calculate the amount of volatiles using the total ion chromatogram.

RNA-seq analysis

Total RNA was isolated from xiangfei kernel using the Hexadecyl trimethyl ammonium Bromide (CTAB) method (Chang et al. 1993). Then using agarose gel electrophoresis and a NanoDrop ND-1000 spectrophotometer (NanoDrop Technologies Inc., Wilmington, USA), the quantity and quality of the total RNA were evaluated. The cDNA libraries were created using a total of 2 g of RNA samples in accordance with the recommended procedure. The Illumina HiSeq 4,000 platform was used for the paired-end sequencing (Illumina Inc., California, USA). Using the Trinity software, the high-quality sequence reads were further combined to obtain unigenes after the low-quality reads, and adaptor sequences were filtered out (version 6.0). Six public databases, including the gene ontology, NCBI nonredundant protein sequences, NCBI nonredundant nucleotide sequences, SwissProt protein sequence database, and Clusters of Orthologous Groups of Proteins, were used to annotate putative terpene associated biosynthetic genes in xiangfei transcriptome database (Supplemental Table S2). Differentially expressed genes were those with a $|\log_2\text{FoldChange}| > 1$ and an adjusted *P*-value ≤ 0.05 .

RNA isolation and RT-qPCR

Genome DNA in the total RNA was eliminated with the gDNA eraser. The PrimeScript RT reagent Kit (Takara, Dalian, China) was used to create first-strand cDNA from 1 g of RNA after which it was diluted with water (1: 10). Utilizing the CFX96 equipment and the Ssofast Eva Green Supermix Kit, RT-qPCR was carried out (Bio-Rad, USA). Melting curves and product sequencing were used to confirm the specificity of primers before use. NCBI/Primer-BLAST was used out for primer design. The reaction mixture, which had a total volume of 20 μL , consisted of the following ingredients: 10 μL of SYBR PCR supermix (Bio-Rad, California, USA), 6 μL of diethylpyrocarbonate-treated water, 2 μL of cDNA template diluted 5 times, and 1 μL of each primer (10 μM) (Zhang et al. 2020c). Data were processed, and relative expression levels were computed using the $2^{(-\Delta\Delta\text{Ct})}$ method to look at the differential expression of genes (Zhang et al. 2020b). The relative expression of control *N. tabacum* leaves expressing an empty vector (SK) was set as 1 for the transient overexpression assay. The relative expression in WT fruit was set as 1.0 for overexpression in tomato (cv. *MicroTom*). Supplemental Table S3 provides a list of the RT-qPCR primers.

Gene isolation, promoter cloning, and analysis

The coding sequence of the differentially expressed TFs and the promoter were cloned based on the RNA-seq data. The

1,426 bp-long *TgGPPS* promoter was amplified by PCR, sequenced, and subjected to an in-silico study starting from the start codon. The online tool <http://bioinformatics.psb.ugent.be/webtools/plantcare/html/> was used to analyze the *TgGPPS* promoter *cis*-element. [Supplemental Table S4](#) contains a list of the primers.

Phylogenetic tree construction

The MEGA 7.0 program ([Kumar et al. 2016](#)) was used to align the full-length GPPS amino acid sequences and create a neighbor-joining tree. NCBI accession numbers are as follows: pedunculate oak QrGPPS (*Quercus robur*; XP_050287822.1), Norway spruce PaGPPS (*Picea abies*; ACA21459.1), grape VvGPPS (*V. vinifera*; XP_002268229.2), tomato SIGPPS (*S. lycopersicum*; NP_001234089.1), *Clarkia brewer* CbGPPS.SSU (AY534745), tobacco NtGPPS (*N. tabacum*; NP_001312253.1), satsuma mandarin CitGPPS (*Citrus unshiu*; AAN86061.1), neem AiGPPS (*Azadirachta indica*; AIG15448.1), peppermint MpGPPS.SSU (*M. piperita*; AAF08792.1), MpGPPS.LSU (AAF08793.1), hops HIGPPS.SSU (*H. lupulus*; ACQ90681), HIGPPS.LSU (ACQ90682), rice OsGPPS (*Oryza sativa*; XP_015644451.1), OsGPPS.SSU (EAY87007), Arabidopsis AtGPPS (*A. thaliana*; CAC16849.1), snapdragon AmGPPS.SSU (*A. majus*; AAS82859), AmGPPS.LSU (AAS82860.1), madagascar periwinkle CrGPPS (*Catharanthus roseus*; AGL91647.1), CrGPPS.SSU (AGL91646.1), CrGPPS.LSU (AGL91645.1), and lavender LiGPPS.LSU (*Lavandula × intermedia*; QHN60320.1).

Subcellular localization analysis

The full-length *TgGPPS*, *TgERF2*, *TgRAV3*, *TgbZIP44*, and *TgbHLH95* coding sequences without stop codons were independently ligated into the pCAMBIA1302 vectors. [Supplemental Table S4](#) provides a list of the primers used in vector construction. Following electroporation into *A. tumefaciens* GV3101, all of the recombinant vectors were transiently expressed in transgenic *N. benthamiana* leaves ([Huang et al. 2013](#)). The *A. tumefaciens* culture was brought to an OD₆₀₀ of 0.75 using the infiltration buffer, which contains 10 mM MES, 10 mM MgCl₂, and 150 mM acetosyringone at a pH of 5.6. Using a confocal laser scanning microscope (LSM510, Karl Zeiss, Germany), GFP fluorescence was measured and observed 2 days after infiltration ([Zhang et al. 2020b](#)). The excitation wavelength for GFP fluorescence was 488 nm, and fluorescence was detected at 490 to 520 nm. The excitation wavelength for NLS fluorescence was 594 nm, and fluorescence was detected at 612 to 635 nm.

Transient overexpression in *N. tabacum* leaves

Full-length cDNA of *TgGPPS* and *TgbHLH95* sequences was separately cloned into the pGreenII0029 62-SK (SK) vector and transformed into Agrobacterium strain GV3101. Cultures were grown to OD₆₀₀ = 0.75 with the same infiltration buffer as in subcellular localization analysis, and 1 mL was infiltrated into *N. tabacum* leaves until the whole leaf was infiltrated. Seven days after infiltration, volatile contents

were evaluated. Negative control was empty vector infiltration (SK).

Stable overexpression in tomato fruit

The full-length *TgGPPS* cDNA was ligated into the pART-CAM vector ([Xu et al. 2014](#)). [Supplemental Table S4](#) provides a list of the primers used in vector construction. Seeds were first placed in sterile distilled water for 30 min; sterilized with 75% (v/v) ethanol for 30 s, sodium hypochlorite solution (available chlorine is 2%) for 15 min, rinsed 4 times with sterile water, and inoculated on 1/2 mS solid medium. According to Wang et al., tomato (cv. *MicroTom*) transformation was accomplished using *A. tumefaciens* ([Wang et al. 2005](#)). The WT and transgenic T₁ generations of tomato plants were grown in a greenhouse at 25 °C with 16 h of light and 8 h of darkness. Tomato fruits at the stage (breaker + 7 days) were frozen in liquid nitrogen and stored at –80 °C for later use. Three plants were selected from each tomato line to symbolize 3 biological replicates, each of which contained 10 fruits.

Dual-luciferase assay

The dual-luciferase assay was used to examine the transactivation capabilities of different TFs on target promoters. Full-length sequences of 14 TFs were amplified and inserted into the pGreen II 0029 62-SK (SK) vector, while the *TgGPPS* promoter sequence was recombined to the pGreen II 0800-LUC (LUC) vector ([Zhang et al. 2020b](#)). [Supplemental Table S4](#) provides a list of the primers used in vector construction. GV3101 of the *A. tumefaciens* was electroporated with each of the SK and LUC vectors and then suspended in the same infiltration buffer as in the subcellular localization analysis, with an optimal density OD₆₀₀ of 0.75. *N. benthamiana* leaves were infiltrated with *A. tumefaciens* combinations of promoters and TFs at a volume of 1:10 using needleless syringes. The LUC/REN fluorescence was assessed with the Dual Luciferase Reporter Assay System (Promega, Madison, USA) after 3 days of infiltration. Each TF–promoter interaction experiment consisted of 3 biological replicates.

Yeast 1-hybrid assay

The interaction of TFs with the *TgGPPS* promoter was tested using the Matchmaker Gold Yeast One-Hybrid Library Screening System (Clontech, Mountain View, California). The pGADT7 vector was used to insert the full-length TF sequences, while the pAbAi vector was used to amplify and ligate the *TgGPPS* promoter sequence. [Supplemental Table S4](#) provides a list of the primers used in vector construction. To test the promoter autoactivation, the Y1HGold yeast strain was transformed using the recombinant *TgGPPS* promoter-pAbAi vector after linearization in accordance with the system's instructions. The TF-pGADT7 plasmids and the empty vector (pGADT7) were individually transfected into the Y1HGold strain carrying the *TgGPPS* promoter.

Electrophoretic mobility shift assay

The full-length *TgbZIP44* was inserted into pET32a vector resulting in His-tagged fusion proteins. The recombinant construct was purified and transformed into *Escherichia coli* BL21 (DE3). Recombinant *TgbZIP44*-His protein was expressed and purified as described by Zhang et al. (2020b). The Lightshift Chemiluminescent EMSA kit (ThermoFisher Scientific, Waltham, MA, USA) was used following the manufacturer's instructions. Sequences (40 bp) containing ACGT motif derived from the *TgGPPS* promoter (Supplemental Table S4) were labeled with 5'biotin by HuaGene. The identical DNA fragments without labels were used as competitors (cold probes). As mutant probes, DNA fragments with the ACGT motif were replaced with a polyA sequence.

Bimolecular fluorescence complementation

Using the methods described by Zhang et al. (2020a), the protein–protein interaction between *TgbHLH95* and *TgbZIP44* can be examined via the BiFC assay. Full-length *TgbHLH95* and *TgbZIP44* without termination codons were cloned into vectors containing the N-terminal and C-terminal sequences of yellow fluorescent protein (YFP), respectively. Supplemental Table S4 provides a list of the primers used in vector construction. All of the recombinant vectors were transiently expressed in *N. benthamiana* leaves by *Agrobacterium* transfection (GV3101). A confocal laser scanning microscope (LSM510, Karl Zeiss, Germany) was used to image the YFP fluorescence of infiltrated leaves after 2 days of infiltration. The fluorescence was measured at a wavelength of 520 to 540 nm. The excitation wavelength for YFP fluorescence was 488 nm, and fluorescence was detected at 490 to 520 nm. The excitation wavelength for NLS fluorescence was 594 nm, and fluorescence was detected at 612 to 635 nm.

Firefly LCI

Validation of the BiFC data was confirmed using the LCI assay. According to the methods described by Xiang et al. (2019), full-length *TgbHLH95* and *TgbZIP44* sequences were ligated into the pCAMBIA-nLUC vector and pCAMBIA-cLUC vectors, respectively. Supplemental Table S4 provides a list of the primers used in vector construction. By using *Agrobacterium*-mediated transformation, each recombinant vector was simultaneously introduced into *N. benthamiana* leaves and resuspended in infiltration buffer at a final density of $OD_{600} = 1$. Prior to infiltration, equal quantities of each *Agrobacterium* strain (i.e. nLUC + cLUC, *TgbHLH95*-nLUC + cLUC, *TgbHLH95*-cLUC + nLUC, *TgbZIP44*-nLUC + cLUC, *TgbZIP44*-cLUC + nLUC, *TgbHLH95*-nLUC + *TgbZIP44*-cLUC, *TgbHLH95*-cLUC + *TgbZIP44*-nLUC) were mixed and incubated at 25 °C for 3 h. The transformed tobacco leaves was used for LCI assays after 2 days of infiltration. After spraying luciferin (0.2 mM) onto the leaves' infiltrated areas, they were left in the dark for 30 min. The luciferase signal was detected using a microplate luminometer (LSM510, Karl Zeiss).

Co-IP assay

The full-length of *TgbHLH95* without stop codon was fused with 3× flag tags and cloned into pCAMBIA1302 vector, and *TgbZIP44* was fused with GFP tag and cloned into pBinGFP2 vector. Following electroporation into *A. tumefaciens* GV3101, all of the recombinant vectors were transiently expressed in *N. benthamiana* leaves. Liquid nitrogen was used to grind the tobacco leaves, and an extraction buffer (25 mM Tris-HCl pH 7.4, 150 mM NaCl, 1 mM EDTA, 1% NP-40, 5% glycerol, 1 mM PMSF, and 0.5 × proteinase inhibitor) was used to extract the proteins. After 15 min of centrifugation at 23,000 g, the extracts were incubated with anti-GFP antibody (Sigma-Aldrich, St. Louis, MO, USA) at 4 °C for 3 h. The immunoprecipitated protein was boiled in 30 μL of protein loading solution for 5 min after being washed 4 times with dilution buffer (10 mM Tris-HCl pH 7.5, 150 mM NaCl, and 0.5 mM EDTA). This protein was then examined by immunoblotting using an anti-GFP or anti-Flag antibody (Sigma-Aldrich, St. Louis, MO, USA).

GST pull-down assay

The full-length of *TgbHLH95* was ligated into pGEX-6P-1 vector to produce GST-tagged fusion protein, while *TgbZIP44* was ligated into pET32a vector to produce His-tagged fusion protein, respectively. According to the manufacturer's instructions, the pull-down assay was carried out using a Pierce Pull-Down PolyHis Protein: Protein Interaction Kit (ThermoFisher Scientific, Waltham, MA, USA). Western blotting was used to identify the eluted samples using anti-GST and anti-His antibodies (Sigma-Aldrich, St. Louis, MO, USA).

Statistical analysis

The Origin 8.0 was used for drawing the figures (Microcal Software). When comparing significant differences, SPSS Statistics 20.0 was used (SPSS Inc., Chicago, IL, USA). Additionally, a Student's *t*-test was applied to determine whether there were any significant differences between the 2 groups (**P* < 0.05; ***P* < 0.01; ****P* < 0.001).

Accession numbers

Sequence data from this article can be found in the Supplemental Table S5.

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Author contributions

L.S. and J.W. conceived the research plans and supervised the experiments; Z.Z. and L.T. carried out most of the experiments and analyzed the data; L.G., Y.G., J.S., W.Y., Y.H., and C.W. provided technical assistance to Z.Z.; Z.Z. and L.T. wrote

and revised the article. M.A.F. took part in the discussion and revision of the manuscript; all authors read and approved the final manuscript.

Supplemental data

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

Supplemental Figure S1. Gas chromatography-mass spectrometry (GC-MS) chromatogram of volatiles in xiangfei (*Torreya grandis*) nuts determined on different days during the postharvest ripening stage.

Supplemental Figure S2. Alignment of amino acid sequences of TgGPPS and GPPS proteins from other species.

Supplemental Figure S3. GC-MS chromatogram of volatiles determined 7 days after infiltration of tobacco (*Nicotiana tabacum*) leaves.

Supplemental Figure S4. GC-MS chromatogram of volatiles determined 7 days after the breaker stage of tomato (*Solanum lycopersicum*, cv. *MicroTom*) fruit.

Supplemental Figure S5. Relative expression of *TgbHLH1* and *TgbHLH2* on days 0 to 12 during the postharvest ripening stage of xiangfei nuts.

Supplemental Figure S6. Transient overexpression of *TgbHLHs* in tobacco (*Nicotiana tabacum*) leaves.

Supplemental Table S1. Isolated and identified transcription factors involved in regulating terpenes biosynthesis in different plant species.

Supplemental Table S2. The fragments per kilobase of exon per million mapped fragments (FPKM) value of all terpenoid synthesis genes from the 2-C-methyl-D-erythritol 4-phosphate (MEP) and mevalonate (MVA) pathways through annotation in Xiangfei transcriptome database.

Supplemental Table S3. Primers used for RT-qPCR.

Supplemental Table S4. Primers used for vector construction.

Supplemental Table S5. The full-length sequences of *Tfs* and *TgGPPS*.

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Conflict of interest statement. The authors declare no conflict of interest.

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