Spearmint R2R3-MYB transcription factor MsMYB negatively regulates monoterpane production and suppresses the expression of geranyl diphosphate synthase large subunit (MsGPPS.LSU)

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Summary
Many aromatic plants, such as spearmint, produce valuable essential oils in specialized structures called peltate glandular trichomes (PGTs). Understanding the regulatory mechanisms behind the production of these important secondary metabolites will help design new approaches to engineer them. Here, we identified a PGT-specific R2R3-MYB gene, MsMYB, from comparative RNA-Seq data of spearmint and functionally characterized it. Analysis of MsMYB-RNAi transgenic lines showed increased levels of monoterpenes, and MsMYB-overexpressing lines exhibited decreased levels of monoterpenes. These results suggest that MsMYB is a novel negative regulator of monoterpenes biosynthesis. Ectopic expression of MsMYB, in sweet basil and tobacco, perturbed sesquiterpene- and diterpene-derived metabolite production. In addition, we found that MsMYB binds to cis-elements of MsGPPS.LSU and suppresses its expression. Phylogenetic analysis placed MsMYB in subgroup 7 of R2R3-MYBs whose members govern phenylpropanoid pathway and are regulated by miR858. Analysis of transgenic lines showed that MsMYB is more specific to terpene biosynthesis as it did not affect metabolites derived from phenylpropanoid pathway. Further, our results indicate that MsMYB is probably not regulated by miR858, like other members of subgroup 7.

Introduction

Plants produce an overwhelming variety of specialized metabolites essential for ecological interactions, among which terpenes form the largest and structurally diverse class of natural products. They are involved in mediating plant defence responses and plant pollinator attraction and facilitate plant–environment interactions (Bhargava et al., 2013; Pichersky and Gershenzon, 2002; Singh and Sharma, 2015). Apart from imparting ecological benefits to plants, terpenoids are of great economic importance to humans as well, as they are widely used in fragrances, cosmetics, pharmaceuticals, agricultural industries and chemical industries (Bouvier et al., 2005). Given the commercial and ecological importance of terpenoids, strategies to metabolically engineer them are of considerable interest. Monoterpenes form the C10 class of terpenoids and are generally colourless, lipophilic and volatile. They are responsible for the characteristic aromas and flavours of essential oils, floral scents and resin of aromatic plants (Loza-Tavares, 1999). The essential oil of spearmint (Mentha spicata) mainly consists of two monoterpenes, limonene and carvone, which are extensively exploited for their biological properties (Ringer et al., 2005). In spearmint, the essential oils are produced and stored in specialized structures called peltate glandular trichomes (Figure S1a). These so-called green biofactories are widely found on aerial surfaces of many aromatic plants, and they actively produce and store large quantity of volatile metabolites (Champagne and Boutry, 2013; Lange and Turner, 2013; Turner et al., 2000).

Terpene biosynthesis pathway has been well studied in plants. The universal C5 building blocks for all types of terpenes, isopentenyl diphosphate (IPP) and its isomer, dimethylallyl diphosphate (DMAPP), are synthesized either by the mevalonate (MVA) pathway in the cytosol or by the 2-C-methyl-D-erythritol 4-phosphate (MEP) pathway in plastids (Vranova et al., 2013). The plastidial MEP pathway is largely responsible for producing C5 precursors for monoterpenes and diterpenes production, whereas the cytosolic MVA pathway generates C10 precursors for sesquiterpene and triterpene production (Dubey et al., 2003). However, few studies indicate that, under certain conditions, an exchange of precursor metabolites can occur between the cytosolic MVA and plastidial MEP pathways (Hemmerlin et al., 2012; Vranova et al., 2013). IPP and DMAPP undergo successive condensation reactions, catalysed by a class of enzymes called prenyltransferases, to form intermediates geranyl diphosphate (GPP; C10), farnesyl diphosphate (FPP; C15) and geranylgeranyl diphosphate (GGPP; C20). These terpene diphosphates form the immediate precursors of monoterpenes, sesquiterpenes and diterpenes, respectively (Kellogg and Poulter, 1997; Liang, 2009; Liang et al., 2002; Ogura and Koyama, 1998). Prenyltransferases are key to terpene production as they control the IPP flux into various branches of the terpene family (Liang et al., 2002; Oldfield and Lin, 2012).
Geranyl diphosphate synthase (GPPS) is the prenyltransferase enzyme that is mainly responsible for the production of monoterpenes. GPPS in the plastids catalyses a single condensation of terpene precursors DXP and IPP, to form GPP. Studies in few species of plants revealed that GPPS can function as a homodimer or heterodimer (Nagegowda, 2010). Heteromeric GPPSs have been characterized so far only in angiosperms Mentha piperita, Antirrhinum majus, Clarkia breweri and Humulus lupulus. All these plants produce large amounts of monoterpenes in specialized organs such as PGTs and flower petals (Burke et al., 1998; Tholl et al., 2004; Wang and Dixon, 2009). Heteromeric GPPS in mint consist of a large subunit (LSU) and a small subunit (SSU); both LSU and SSU are catalytically inactive alone (Burke and Croteau, 2002; Burke et al., 1999; Croteau et al., 2005). Interaction between the two subunits results in the formation of an active GPPS. Structural studies show that the LSU acts as the catalytic unit, whereas the SSU serves as the regulatory unit (Chang et al., 2010).

In the case of spearmint, the monoterpenes biosynthetic pathway is well characterized (Champagne and Boutry, 2013; Croteau et al., 1991; Lange et al., 2011; Munoz-Bertomeu et al., 2008). Strategies to increase the yield of peppermint oil by manipulating genes that code for structural pathway enzymes have been reported (Diemer et al., 2001; Mahmoud et al., 2004). However, the developmental regulation of this secondary metabolite pathway still remains elusive. Transcription factors (TFs) can activate or repress multiple genes in a metabolic pathway; hence, they are ideal targets for pathway engineering (Grotewold, 2008; Iwase et al., 2009). Only one TF MsYABBY5 has been reported from spearmint that regulates monoterpenes production (Wang et al., 2016) but several TFs have been identified from other plants like Artemisia, cotton, Taxus, rubber and rice that control terpene biosynthesis (Chen et al., 2012; Li et al., 2013; Miyamoto et al., 2014; Shen et al., 2016; Xu et al., 2004; Zhang et al., 2012).

The R2R3-MYBs represent one of the largest families of plant TFs (Kranz et al., 1998; Stracke et al., 2001) and can function as activators or repressors of genes (Adato et al., 2009; Legay et al., 2007). Many R2R3-MYBs have been characterized as transcriptional activators of flavonoid, phenylpropanoid and glucosinolate pathway genes in plants (Dubos et al., 2010; Zhao and Dixon, 2011; Zhong et al., 2010). R2R3-MYBs belonging to subgroup 4 are known to negatively regulate monolignol and flavonoid biosynthetic pathways (Fornalé et al., 2010; Jin et al., 2000; Legay et al., 2007; Preston et al., 2004; Tamagnone et al., 1998). Further, a single R2R3-MYB gene was also found to act as both repressor and activator of genes (Bhargava et al., 2010). Redundancy, dual function and opposite action as activators and repressors on common target genes by R2R3-MYB help fine tune the regulation of plant secondary metabolite pathways. Despite several studies on R2R3 MYB-mediated regulation of plant phenylpropanoid pathway, knowledge of its role in terpene secondary metabolism is lacking (Matus, 2016). Besides TFs, microRNAs also are known regulators of various processes in plant growth and development. However, knowledge of their role in controlling secondary metabolic pathways is limited. microRNAs mainly target TFs to regulate various plant developmental processes (Sunkar et al., 2012; Wu et al., 2013; Zhang, 2015; Zhang and Wang, 2015). In plants, MYB TFs are potential targets for miR858 (Addo-Quaye et al., 2008; Guan et al., 2014; Jeong et al., 2013; Jia et al., 2015; Xia et al., 2012). Further, many studies have shown that miR858 can target R2R3-MYBs in apple, grapes, salvia and Arabidopsis (Li and Lu, 2014; Rock, 2013; Sharma et al., 2016; Xia et al., 2012). Presently, miR858 is considered as a potential regulator of plant phenylpropanoid metabolite pathway through its action on R2R3-MYBs (Fahlén et al., 2007).

In this study, we identified a R2R3-MYB transcript which is preferentially expressed in spearmint PGTs from the comparative transcriptome data (Jin et al., 2014) of spearmint leaf, leaf stripped of PGT (L-PGT) and PGT. Phylogenetic analysis assigned MsMYB to subgroup 2 of R2R3-MYBs of Arabidopsis. Members of subgroup 2 are closely related R2R3-MYBs which control flavonoid accumulation in different parts of the Arabidopsis seedling, and are regulated by miR858 (Fahlén et al., 2007; Sharma et al., 2016; Yang et al., 2012). Apart from Arabidopsis, several other R2R3-MYBs from different plants, similar to the members of subgroup 7, are also involved in flavonoid pathway (Adato et al., 2009). We functionally characterized the role of this R2R3-MYB and R2R3-MYB and found it to be a negative regulator of monoterpenes production in spearmint. Suppression of MsMYB expression, by RNAi approach, led to increase in monoterpenes levels and its overexpression led to decrease in monoterpenes levels in transgenic lines. This is the first report of regulation of monoterpenes production by a R2R3-MYB. Ecopic expression of MsMYB in sweet basil (Ocimum basilicum), an aromatic plant similar to spearmint, and in tobacco (Nicotiana sylvestris), affected the production of terpenes but not flavonoids. We further found that MsMYB suppresses the expression of geranyl diphosphate synthase large subunit (MsGPPS.LSU) gene in spearmint. To investigate whether MsMYB is regulated by miR858, similar to other R2R3-MYBs of subgroup 7, we performed small RNA-Seq of spearmint PGT and L-PGT. We found low amounts of miR858 in spearmint PGT but high amounts in L-PGT. Although in silico analysis predicted MsMYB as a target for miR858, we were unable to detect its cleaved products. This suggests that the differential expression of miR858 and MsMYB in separate tissues prevents MsMYB regulation by miR858. Collectively, the above results provide insight into the production of essential oils in a valuable crop plant, such as spearmint, and expand our knowledge of R2R3-MYB-mediated regulation of plant secondary metabolism.

**Results**

MsMYB shows preferential expression in spearmint PGTs

From the transcriptome data of three tissues of spearmint, namely leaves, PGTs and leaves devoid of PGTs (Jin et al., 2014), we annotated 45 R2R3-MYBs (Figure 1; Supporting Data S1). Of these, only two R2R3-MYBs, MsMYB and MsMYB112, showed high expression in PGTs. MsMYB was chosen for further characterization. RNA-Seq expression of MsMYB along various tissues was further validated by quantitative RT-PCR (qRT-PCR) (Figure 2a). Full-length open reading frame (ORF) of MsMYB was amplified from PGT cDNA and consisted of 813 base pairs (bp) encoding a polypeptide of 271 amino acids (Figure 3). It contained a R2, R3 repeat and five tryptophan residues within the edge of R2R3-MYB-mediated regulation of plant secondary metabolism.
under subgroup 7 along with AtMYB111, AtMYB11 and AtMYB12 (Dubos et al., 2010). All these three Arabidopsis MYBs function as activators of flavonoid pathway (Stracke et al., 2007).

To examine subcellular localization of MsMYB, the ORF was fused with yellow fluorescent protein (YFP) and expressed under the control of CaMV 35S promoter in Nicotiana benthamiana leaves by agroinfiltration. As shown in Figure 2b, the recombinant protein specifically localized to the nucleus.

MsMYB promoter shows trichome specific expression

To identify the cis-acting regulatory elements in the promoter of MsMYB, we cloned a 615-bp genomic DNA fragment upstream of the translation start site by genome walking. Apart from the common TATA and CAAT box, several other cis-acting regulatory elements were identified (Figure 4a) using PlantCARE tool (http://bioinformatics.psb.ugent.be/webtools/plantcare/html/). Many of these elements were for hormones like ABRE-motif which is known for abscisic acid responsiveness (Basu et al., 2014), AuxRE-motif which is an auxin-responsive element (Ulmasov et al., 1995), CGTCA-motif and TGACG-motif which are known for MeJA responsiveness (Zhou et al., 2012). Interestingly an AC-I element was also found. In general, AC-rich regions are known to be bound by R2R3-MYBs (Koschmann et al., 2012; Prouse and Campbell, 2012). This suggests that MsMYB might be regulated by other R2R3-MYBs. To check for the expression pattern, the 615-bp promoter fragment was fused with a β-glucuronidase (GUS) reporter gene and transformed into N. benthamiana plants. Trichome-specific staining was observed in leaves and stems of tobacco transgenic plants (Figure 4b,c).

Manipulation of MsMYB expression levels affects monoterpene production in spearmint

To characterize the function of MsMYB, we generated transgenic spearmint lines harbouring RNAi construct and overexpression construct where MsMYB was under the control of CaMV 35S promoter. Four independent lines each confirmed by Southern blot were selected for further characterization (Figure S1b,c). qRT-PCR analysis showed significant reduction in levels of MsMYB transcripts in RNAi lines (Figure 5a) and higher levels of MsMYB transcripts in overexpression lines when compared to wild type (WT) (Figure 5b). Both overexpression and RNAi transgenic plants appeared phenotypically similar to WT plants. Scanning electron microscopy was performed on these plants to take a closer look at leaf cells and PGTs. No phenotypical changes were observed.

Further, gas chromatography–mass spectrometry (GC-MS) analysis was performed to evaluate any changes in volatiles production. Major secondary metabolites of spearmint are monoterpenes; limonene and carvone (Figure S3a). GC-MS analysis revealed significant changes in the amount of these secondary metabolites in transgenic lines. Compared to WT plants, total monoterpene production in MsMYB-RNAi lines was higher; about 2.3- to 4.5-fold increase was observed (Figure 5c, d), whereas in MsMYB-overexpressing lines a decrease of 0.5- to 0.7-fold was seen (Figure 5e, f). The above results indicated that MsMYB might be a negative regulator of secondary metabolism in spearmint.

MsMYB negatively regulates MsGPPS.LSU

To understand how MsMYB affects terpene metabolism, it is essential to know its downstream target genes. Towards this, we checked the expression levels of several transcripts encoding enzymes in various terpene precursor pathways, transporters and flavonoid pathway (Table S2), but none of them were significantly changed except for MsGPPS.LSU. It showed

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increased levels in MsMYB-RNAi lines and decreased levels in MsMYB-overexpressing lines (Figure 6a,b). GPPSs are known to localize to both nongreen plastids and chloroplasts of photosynthetic cells where MEP pathway is active (Bouvier et al., 2000). We checked the localization of MsGPPS.LSU and found it to be localized in the chloroplast (Figure 9b). To check whether the promoter of MsGPPS.LSU has putative MYB binding sites, we cloned a 549-bp genomic DNA fragment upstream of the translation start site by genome walking. Using PlantCARE tool we identified two MYB binding sites (MBS) along with several elements for hormones and light responsiveness (Figure 7a).

To verify MsMYB interaction with GPPS.LSU promoter, yeast one-hybrid assays were performed. The whole GPPS.LSU promoter along with two truncated regions, viz. site 1 and site 2, which had the conserved MYB binding domains and mutated versions of site 1 and site 2 was used as bait to generate five Y1HGold [Bait/AbAi] strains, that is pAbAi-GPPS-full, pAbAi-site1, pAbAi-site2, pAbAi-mut-site1 and pAbAi-mut-site2. After cotransformation with functional MsMYB gene in pGADT7 vector (pGADT7-MYB), three Y1HGold strains, that is pAbAi-GPPS-full and pAbAi-site1 and pAbAi-site2, could grow on SD minus leucine with aureobasidin (SD/-Leu/AbA, 800 ng of AbA) auxotrophic medium but not the two mutant strains pAbAi-mut-site1 and pAbAi-mut-site2, confirming interaction of MsMYB with GPPS.LSU promoter in vivo (Figure 8). TFs can act as activators or repressors of transcription. Transgenic studies indicated that MsMYB is a negative regulator. To further determine the transcriptional repression activity of MsMYB in planta, transient expression assay in N. benthamiana was performed. We used 35Spro:GFP or 35Spro:MsMYB as effectors and MsGPPS.LSUpro:GUS as reporter. Promoter activity of MsGPPS.LSUpro:GUS was significantly reduced in 35Spro:MsMYB expressing leaves when compared to 35Spro:GFP expressing leaves (Figure 9a). These results show that MsMYB binds to the promoter of MsGPPS.LSU and suppresses its activity.
Terpene biosynthesis is known to be regulated by light in many plants (Cordoba et al., 2009). As we found light-responsive elements in MsGPPS.LSU promoter, we checked whether MsGPPS.LSU is also regulated by light. WT spearmint plants were grown in dark and light conditions. Leaf samples were collected at 0-, 24-, 48- and 72-h interval to quantify MsGPPS.LSU transcript levels. Expression of MsGPPS.LSU was highly reduced in the plants grown under dark conditions suggesting that MsGPPS.LSU is regulated by light (Figure 7b). As GPPS is a key enzyme in the monoterpene pathway, multiple layers of gene regulation to fine tune the pathway is plausible.

Ectopic expression of MsMYB affects secondary terpene metabolism in sweet basil and Nicotiana sylvestris

Similar to spearmint, sweet basil too produces its essential oil in PGTs. The constituents of sweet basil essential oil are derived from terpene and phenylpropanoid pathways (Figure S3b). It is a mixture of monoterpenes, sesquiterpenes and the main phenylpropanoid derivative is eugenol. Ectopic expression of MsMYB in

Figure 4 MsMYB promoter analysis and expression pattern in tobacco. (a) Regulatory elements present in the promoter region (–615 bp) of MsMYB. (b), (c) N. benthamiana plants transformed with MsMYB:GFP showing trichome specific expression in leaves and stems. Black arrows point GFP stained trichome heads.

Figure 5 Transcript levels and GC analysis of transgenic plants. Reduced (a) and increased (b) levels of MsMYB in transgenic spearmint MsMYB-RNAi and MsMYB-overexpressing lines when compared to WT. (c), (d) Increased levels of limonene and carvone in MsMYB-RNAi lines when compared to WT. (e), (f) Reduced levels of limonene and carvone in MsMYB-overexpressing lines when compared to WT. GFP: GFP-overexpressing line. Data are indicated as mean ± SE. *P < 0.05; **P < 0.01; ***P < 0.001.
sweet basil was pursued to evaluate whether MsMYB can affect the sesquiterpene pathway and phenylpropanoid pathway as suggested by phylogenetic analysis. Three independent lines (line 5, line 7 and line 10) confirmed by Southern blot were further selected for characterization (Figure S1d). Ectopic expression of MsMYB in these lines was confirmed by qRT-PCR (Figure 10a). Analysis of CBT-diols production in transgenic plants showed a 42%–50% reduction (Figure 10b). Similar to sweet basil plants, the leaves of tobacco transgenic lines were smaller in size when compared to WT (Figure S4c). The above results indicate that MsMYB can affect the flux in various branches of terpene metabolism.

**MsMYB is probably not a target of microRNA858**

microRNA858 is known to regulate several R2R3-MYBs, specifically the subgroup 7 R2R3-MYBs, AtMYB11, AtMYB111 and AtMYB112 (Fahlgren et al., 2007; Sharma et al., 2016). As MsMYB fell under subgroup 7, we were interested to know whether it is regulated by microRNA858. We generated small RNA sequencing data from PGTs and L-PGT and identified ms-miR858. The sequence similarity of ms-miR858 is shown in Figure 11a. Expression analysis from the sequencing data showed that ms-miR858 had higher expression in L-PGT tissue when compared with PGT. This was validated by TaqMan® qPCR assay using miR396 as internal control due to its stable expression in L-PGT and PGT tissues (Figure 11b). The Plant Small RNA Target Analysis Server (Dai and Zhao, 2011) predicted MsMYB as a probable target of ms-miR858 (Figure 11c). To test whether low levels of miR858 expression can target MsMYB, we looked for cleavage products through RNA ligase-mediated 5’ amplification of cDNA ends (RLM-RACE) and poly(A) polymerase-mediated (PMP)-RACE. However, no cleavage products of MsMYB were found in our conditions, thus reducing the possibility of it being regulated by miR858. MsMYB and miR858 appears to be differentially regulated at transcriptional level making their interaction less feasible.

**Discussion**

Extensive exploitation of aromatic plants for their commercially valued secondary metabolites calls for novel strategies for metabolic engineering to increase yield. A complete understanding of the secondary metabolism pathway by unravelling the enzymes and TFs controlling the pathway is indispensable for the same. PGTs are found on the aerial parts of many aromatic plants but very few studies have focused on identifying regulators of secondary metabolite production from these organs (Wang, 2014). In this study, we identified and characterized a R2R3-MYB gene, MsMYB from spearmint which is highly expressed in PGTs and acts as a negative regulator of monoterpene synthesis. GPPS enzyme is responsible for the formation of monoterpene precursor C15-GPP in plastids and regulates flux into this pathway. Details about their function and regulation in plants are very scarce. The heteromeric GPPS in spearmint consists of a small and large subunit and interaction between them results in an active transgenic plants (Figure 6c), but levels of ObFPPS were unaltered. Additionally, we measured the total flavonoid content in the leaves of sweet basil transgenic plants and found the total flavonoid content to be unaltered (Figure 10f). This suggests that MsMYB is a R2R3-MYB specific to terpene metabolism.

The sweet basil data showed that MsMYB can affect both monoterpene and sesquiterpene synthesis. To investigate whether it has a role in diterpene synthesis, we ectopically expressed MsMYB in *N. sylvestris*. Glandular trichomes of *N. sylvestris* mainly produce diterpenes; cembranoids (CBT-diol) are generally derived from the same MEP pathway as the monoterpenes. Three independent lines were chosen for further characterization and ectopic expression of MsMYB was confirmed by qRT-PCR (Figure S5a). Analysis of CBT-diols production in transgenic plants showed a 42%–50% reduction (Figure S5b).
GPPS. Structural and mutagenic studies of these two subunits have shown that the large subunit is catalytically active and presumably through protein–protein interactions, and the SSU shapes the active site cavity of LSU towards producing C10-GPP and restricts the production of long-chain C15 or C20 prenylphosphates (Chang et al., 2010). In spearmint, both the subunits of GPPS are preferentially expressed in PGTs (Turner and Croteau, 2004). MsMYB was found to interact with MsGPPS.LSU promoter and repress its activity. The increased production of monoterpene in MsMYB-RNAi lines can be attributed to enhanced production of GPP through greater production of MsGPPS.LSU. Misexpression of either the regulatory or catalytic subunit are known to alter enzyme activity in plants and animals (Gergs et al., 2004; He et al., 2004; Hu et al., 2016; Tiessen et al., 2002). In snapdragon and hop where heteromeric GPPS are found, only GPPS.SSU is closely correlated with monoterpene production, whereas the GPPS.LSU is more ubiquitous. But spearmint GPPS.LSU expression is more PGT-specific (Jin et al., 2014). Unlike the purified mint GPPS.LSU, which is inactive by itself, the potential GPPS.LSU from snapdragon and hops are active alone and functions as GGPPS (Tholl et al., 2004; Wang and Dixon, 2009). This suggests function of GPPS subunits from different plants might vary and also the mechanisms that regulate them.

Ectopic expression of MsMYB in sweet basil and tobacco revealed that it could also perturb metabolites derived from sesquiterpene and diterpene pathways. Sesquiterpenes and diterpenes are derived from C15-FPP and C20-GGPP which are produced in cytoplasm and plastids through MVA and MEP. Figure 7

**Figure 7**  Promoter analysis of MsGPPS.LSU. (a) Regulatory elements present in the promoter region (~549 bp) of MsGPPS.LSU. (b) Decreased levels of MsGPPS.LSU in plants grown under dark when compared to plants grown under light. Data are indicated as mean ± SE.
pathway, respectively. Studies have demonstrated that exchange of precursor metabolites can occur between the cytosolic MVA and plastidial MEP pathway. Metabolic intermediates like IPP, GPP and GGPP can be transported across plastidial membranes through transporters (Hemmerlin et al., 2012; Vranova et al., 2013). FPPS catalyses the sequential condensation of two molecules of IPP with DMAPP to form C15-FPP, the immediate precursor for all sesquiterpenes (Cornish, 1993; Lange et al., 2000). But FPPS is also known to accept C10-GPP as an allylic substrate to generate FPP (Guo et al., 2015; Hemmerlin et al., 2003). Further sesquiterpenes synthesized in the cytoplasm can be derived from plastidial GPP too (Adam et al., 1999). Hence, an alteration in GPP produced in plastids can have an effect on sesquiterpene production. Ectopic expression of MsMYB in sweet basil resulted in reduced expression of ObGPPS.LSU. The decrease in sesquiterpene production observed in sweet basil indicates that GPP produced might be used towards synthesizing sesquiterpene especially as no change was observed on sweet basil FPPS. The monoterpene and diterpene pathway are both largely localized to plastids. GGPPS is responsible for producing GGPP for diterpenes (C20) and tetraterpenes (C40) synthesis. Apart from IPP and DMAPP, in vitro data show that GPP or FPP can also be used by GGPPS as allylic substrates to generate GGPP (Hefner et al., 1998; Okada et al., 2000; Takaya et al., 2003). The decrease in diterpene produced in MsMYB-overexpressing tobacco plants can be due to a reduction in endogenous pool of GPP. Tobacco generally produces low levels of monoterpene and there is no information currently about endogenous GGPP from it. Although introduction of monoterpene synthases from other plants does result in the emission of volatile monoterpenes in tobacco, suggesting some amount of GPP availability (Lückner et al., 2004).
overexpression of MsMYB in tobacco and sweet basil generated smaller leaves. Several major plant hormones are produced through terpene pathway such as gibberellin, cytokinin, abscisic acid and strigolactone which control various aspects of plant development (Gomez-Roldan et al., 2008; Umehara et al., 2008). Further experiments need to be carried out to identify perturbation in which hormone/hormones leads to smaller leaf size. This phenotype was not observed in spearmint. TFs can have additional targets when ectopically expressed. For example, maize ZmMYB31 cannot bind to the promoter of maize 4CL and C3H genes; however, it can repress the expression of Arabidopsis 4CL and C3H when overexpressed in Arabidopsis (Fornalé et al., 2010). There is a possibility that additional target genes related to primary terpene metabolism are regulated when MsMYB is overexpressed heterologously in these plants. Ectopic expression of MsMYB did not lead to any observable defect in chlorophyll formation which is a primary metabolite derived from diterpenes. An earlier study showed that GGPP used in chlorophyll biosynthesis is formed directly from DMAPP and IPP rather than from GPP and IPP. Thus, any alterations in the amount of GPP formed do not affect chlorophyll biosynthesis (Van Schie et al., 2007).

TFs are known to regulate multiple genes. MsGPPS.LSU is probably one of the many downstream targets regulated by MsMYB. Hence, we cannot negate the fact that the effects seen on terpene metabolism in spearmint, sweet basil and tobacco can be due to other genes as well. Transcriptome analysis of MsMYB-RNAi lines can provide us with more candidate genes that are potentially regulated by MsMYB. We found that MgGPPP.LSU is further regulated by light. Several genes of MEP pathway are known to be regulated by light (Cordoba et al., 2009). Secondary metabolite production is mainly for plants biotic and abiotic stress responses, so stringent multiple regulatory mechanisms involving activators and repressors must exist to determine the timing, amount and patterning of these compounds.

Majority of R2R3-MYBs characterized in plants are involved in the regulation of different classes of phenylpropanoid-derived compounds (Liu et al., 2015). Phylogenetic analysis showed that MsMYB is very similar to subgroup 7 of Arabidopsis R2R3-MYBs which include MYB11, MYB12 and MYB111. These MYBs have been characterized as activators of biosynthetic enzymes of flavonoid pathway which is derived from the general phenylpropanoid pathway. Analysis of major

Figure 10 Ectopic expression of MsMYB in sweet basil alters the amount of secondary metabolites. (a) MsMYB expression in basil transgenic plants. (b), (c) Reduced levels of monoterpenes and sesquiterpenes. (d) Decreased levels of total terpenes. (e) Unaltered levels of phenylpropenes (eugenol). (f) Unaltered levels of total flavonoids. Data are indicated as mean ± SE. *P < 0.05; **P < 0.01; ***P < 0.001.

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flavonoid pathway genes from spearmint showed no change in MsMYB transgenic lines. Further no change was seen in either eugenol production which is derived from phenylpropanoid pathway or in total flavonoid content of leaves from sweet basil MsMYB-overexpressing lines. These results suggest MsMYB as a regulator of terpene metabolism and not affecting flavonoid production. This indicates towards diversification of function within the same subgroup. Although the N-terminus of R2R3-MYBs is conserved, C-terminus amino acids are very diverse and unique providing opportunity for divergence. Comparison of MYB proteins from different species will furnish new insights into the evolution of this important family of TF. In the past few years numerous studies have demonstrated miR858 can target R2R3-MYBs from various plants including the members of subgroup 7 of Arabidopsis R2R3-MYBs (Addo-Quaye et al., 2008; Rock, 2013; Sharma et al., 2016; Xia et al., 2012). Small RNA-Seq data of spearmint PGTs and L-PGTs showed an inverse pattern of expression for miR858 and MsMYB. High expression was seen in L-PGTs and low in PGTs for miR858 which is opposite to MsMYB expression, indicating a differential transcriptional regulation. MsMYB promoter is able to drive trichome specific expression as seen in tobacco. As genome of spearmint is not available, we cannot verify the promoter activity of miR858. Although in silico analysis predicts MsMYB to be a potential target for miR858, no cleavage products could be observed. This suggests that the PGT specific expression of MsMYB is governed mainly by the cis-elements present in its promoter and may be by other trans-factors but not by miR858. 

In conclusion, MsMYB is the first R2R3-MYB gene identified from spearmint that is related to terpene secondary metabolism. MsMYB represses MsGPPS.LSU and is able to modify terpene production. This opens up new avenues and targets to metabolically engineered plants for altering pathways.

**Experimental procedures**

**Plant material and transformation**

Secondary metabolites of commercial spearmint variety (M. spicata) and sweet basil (O. basilicum) were analysed by GC-MS and grown in green house under natural light conditions. Agrobacterium mediated transformation of spearmint was performed according to previously published protocol (Niu et al., 1998, 2000) with several modifications. The complete details of transformation are provided in the supplementary data. Agrobacterium mediated transformation of sweet basil was performed as previously described by Wang et al. (2016). Tobacco transformation was performed as described previously (Gallois and Marinho, 1995).

**RNA isolation and quantitative real-time PCR (qRT-PCR)**

For PGT RNA isolation, initially, PGTs were isolated from 2- to 3-cm leaves as described previously (Jin et al., 2014). Later, total RNA was extracted from PGT using the Spectrum Plant total RNA kit from Sigma (UK). Total RNA from other tissues was extracted using an RNaseasy Plus Mini kit from Qiagen (Germany). 500 ng of RNA was reverse-transcribed to cDNA using Scrypt cDNA Synthesis kit form Bio-Rad (CA, USA). Expression levels of genes along various tissues were analysed using qRT-PCR that was carried out as described previously (Jin et al., 2014). In current study, elongation factor 1 (ef1) was used as internal control for both spearmint and sweet basil, due to its stable expression in plant (Nicot et al., 2005). For tobacco, elongation factor TuB (EF-TuB) was used as internal control. Error bars represent mean ± SE.

**TaqMan qPCR assay**

7 ng of total RNA was reverse-transcribed using the TaqMan microRNA reverse transcription kit (Applied Biosystems, USA) according to the manufacturer’s protocol. Custom-made primers were used specifically for miR858 and miR396 cDNA synthesis. Custom TaqMan® small RNA assays (Applied Biosystems, USA) were carried out for microRNA qPCR validation using custom-made TaqMan® probe for miR858 and miR396. Expression values were calculated relative to the internal control miR396.

**Promoter cloning and analysis**

Genomic DNA was isolated from young leaves of spearmint using CTAB method. The flanking sequences of MsMYB and MsGPPS.LSU genes were amplified using a GenomeWalker Universal kit (Clontech, USA) and later ligated to pGEM®-T vector. The resulting product was transformed into Escherichia coli (E. coli) XL1-Blue cells and sequenced. The promoter was
amplified with Phusion® High-Fidelity DNA Polymerase (NEB, USA) and subcloned into a gateway donor vector pENTR™/D-TOPO™ (Invitrogen, Germany). Further, the recombinant plasmid was introduced into destination vector pBGWFS7 by LR recombination. The destination plasmid was further transformed into Agrobacterium EHA105 by heat shock. The recombinant Agrobacterium EHA105 strain was used to generate transgenic tobacco lines.

Gene amplification and plasmid construction

Full-length ORF encoding MsMYB was obtained by performing 3’ and 5’ RACE using the SMARTer™ RACE cDNA amplification kit from Clontech (CA, USA). For sequencing of ORF, purified fragments were ligated with pGEM®-T vector and transformed into E. coli XL1-Blue cells. To overexpress MsMYB, firstly, the sequences were amplified with Phusion® High-Fidelity DNA Polymerase (NEB, USA). For MsMYB-RNAi, four primers with restriction enzymes located at flanking region were used to amplify the fragment showing low similarity to other MYB genes. The purified PCR products were then cloned into the donor vector and subsequently introduced into the destination vector pK7WG2D via LR recombination. The MsMYB gene was driven by 35S promoter in both overexpression and RNAi plants. All destination plasmids harbouring target gene were transformed into Agrobacterium EHA105 by heat shock. The recombinant Agrobacterium EHA105 strains were used for plant transformation. Sequences of primers used in this study are listed in Table S1.

Subcellular localization of TFs

The full-length cDNA of MsMYB without the stop codon was cloned into the Gateway donor vector pENTR/D-TOPO (Invitrogen, Germany) and then subsequently transferred into the destination vector pSG5 (Zhang et al., 2005) which contains the CaMV 35S promoter and C-terminal in frame with YFP, to generate MsMYB-YFP. The construct was then introduced into Agrobacterium strain EHA105 by heat shock. 4’,6-Diamidino-2-phenylindole (DAPI) was used as marker to stain nucleus. N. benthamiana seeds were germinated on MS plate and transferred into soil. Three weeks after growing in the glasshouse, the seedlings were used for agroinfiltration. Subcellular localization was performed as described by Wang et al. (2016).

Selection of transgenic lines

Initially, visual screening using GFP filter was pursued to isolate GFP-positive plants. Later, DNA was isolated from GFP-positive plants and insertion of gene of interest was checked using PCR. DNA-positive lines were then subjected to Southern blot using DIG wash and block buffer set from Roche (IN, USA). DNA probe followed by antibody binding and chemiluminescent reaction development by CDP-Star. Finally, the membrane was exposed in a film and the number of T-DNA insertions were analysed. Spearmint plants were clonally propagated thrice before analysis. Basil plants were analysed in T2 generation and a minimum of six plants were sampled for each line. Tobacco plants were analysed in T1 generation. For each line, ~32 plants were used for phenotypical screening and six plants for GC-MS analysis.

GC-MS and total flavonoid content analysis

For GC-MS studies, spearmint plants were clonally propagated three times before collecting about 4–6 leaves of 2–3 cm which were ground to a fine powder using liquid nitrogen and homogenized using 500 µl ethyl acetate. Camphor was added as an internal control. For sweet basil, the analysis was performed in T-2 plants. From bottom, leaves of 3–4 cm at fourth node were used. For tobacco, 4- to 5-cm leaves were used from T1 plants. Diethyl sebacate was added as an internal standard in sweet basil and tobacco samples. Samples were incubated for 10 min at room temperature with vigorous shaking followed by a centrifugation for 10 min at 15871g. The top organic layer was transferred to a new tube and dehydrated using anhydrous Na2SO4. The samples were analysed using GC-MS (7890A with 5975C inert MSD with triple axis detector, Agilent Technologies, USA). 2 µl of samples was injected, and separation was achieved with a temperature programme of 50 °C for 1 min and increased at a rate of 8 °C/min to 300 °C and held for 5 min, on a 30 m HP-5 MS column (Agilent Technologies, USA). Estimation of total flavonoid content was performed using the aluminum chloride (AlCl3) colorimetric method as described previously (Vaidya et al., 2014).

Yeast one-hybrid assay

For ascertaining the interaction between MsMYB and MsGPPS.LSU promoter, a Matchmaker Gold yeast one-hybrid library screening system (Clontech, USA) was used. As MsGPPS.LSU promoter encompasses two different MYB binding sites, two promoter regions approximately 40–45 bp in size harbouring the two sites, namely site 1 and site 2, were cloned into the MCS of pAbAi vector independently in addition to two mutant versions of these two sites. Furthermore, complete promoter was also cloned in pAbAi vector and used as bait for further confirmation. The MsMYB gene was cloned into the pGADT7 vector to be used as prey in the form of functional fusion protein along with GAL4 transcription activation domain (GAL4 AD). As a positive control, p53 bait and prey provided in the kit was used. Mutated site 1 and mutated site 2 were used as negative control. Yeast transformation and validation of positive interactions were implemented as described in the Matchmaker Gold Yeast one-hybrid system user manual (www.clontech.com/xclt_1bcGetAttachment.jsp?cltemId=17599). The primers used are listed in Table S3.

Transactivation activity assay

The 549-bp promoter region of MsGPPS.LSU was amplified and inserted into pENTR™/D-TOPO™. The resulting plasmid was transformed into pBGWFS7 by LR recombination and further introduced into Agrobacterium EHA105. Leaves of N. benthamiana were agroinfiltrated with effector and reporter at a ratio of 1:1. Two days after infiltration, leaves were harvested to isolate crude protein. GUS quantitative assay was performed in triplicate as described in Li et al. (2014).
RLM-RACE and PPM-RACE

For RLM-RACE, 5’ RACE adapter was ligated to 1 μg of total RNA, and for PPM-RACE, poly(A) tailing of 1 μg of total RNA was performed using poly(A) polymerase (Invitrogen, Germany). RNA was then reverse-transcribed and amplified for 5’ and 3’ ends using FirstChoice RLM-RACE kit (Thermo Fisher, USA) according to the manufacturer’s protocol.

Small RNA library construction and high-throughput sequencing

The RNA libraries were prepared using the TruSeq RNA library prep Kit v2, set A (Illumina Inc., USA) according to manufacturer’s instructions. The quality and size of DNA libraries for sequencing were checked using the Agilent 2200 TapeStation system. Three libraries were run on single lanes on HiSeq™ 2000 (Illumina Inc., USA), individually.

Bioinformatics analysis of sequencing data

After removing adapters by Cutadapt tool (Martin, 2011), modified sequences from 18 nt to 30 nt were used for further analysis. To begin with, sequences of rRNAs, tRNAs, snRNAs and snoRNAs available in Rfam11.0 database (ftp://ftp.ebi.ac.uk/pub/databases/Rfam/11.0/) were removed from the small RNA sequence reads. Deposited miR858 sequences from miRBase server. deposited miR858 sequences from miRBase database (http://www.mirbase.org/search.shtml) were mapped to the remaining small RNA sequence reads by bowtie tool (Langmead and Salzberg, 2012) to identify ms-miRNA858. A maximum of one mismatch per sequence was set as a parameter.

Phylogenetic analysis

Phylogenetic tree was constructed using MEGA6 software by the neighbour-joining method with bootstrap values of 1000 replicates and edited in FigTree.

Statistical analysis

Data are indicated as ‘mean ± SE’ of three to six biological replicates each performed in triplicate. Statistical significance between transgenic plants and WT was analysed using a two-tailed Student’s t-test and indicated by asterisks. * indicates P < 0.05; ** indicates P < 0.01; *** indicates P < 0.001.

Accession numbers

Sequence data of A. thaliana R2R3-MYB sequences were obtained from TAIR website.

Competing interests

The authors declare that they have no competing interests.

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MsMYB negatively regulates terpene synthesis

Figure S1. Scanning electron micrograph (SEM) of leaf surface and Southern blot analysis of transgenic plants.
Figure S2. Phylogenetic tree showing the similarity of MsMYB to known Arabidopsis thaliana R2R3-MYBs.
Figure S3. GC profiles of wild-type plants.
Figure S4. Transgenic plants overexpressing MsMYB show smaller leaf size.
Figure S5. Ectopic expression of MsMYB in tobacco.

Table S1. Primers used in this study.
Table S2. List of genes analysed in spearmint transgenic plants.
Table S3. Oligonucleotides used for the generation of bait sequences.
Data S1. Sequence data and expression values of spearmint R2R3-MYBs.
Data S2. Accession numbers of A. thaliana R2R3-MYBs.