

Methyl Jasmonate-Elicited Transcriptional Responses and Pentacyclic Triterpene Biosynthesis in Sweet Basil^{1[C][W]}

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Sweet basil (*Ocimum basilicum*) is well known for its diverse pharmacological properties and has been widely used in traditional medicine for the treatment of various ailments. Although a variety of secondary metabolites with potent biological activities are identified, our understanding of the biosynthetic pathways that produce them has remained largely incomplete. We studied transcriptional changes in sweet basil after methyl jasmonate (MeJA) treatment, which is considered an elicitor of secondary metabolites, and identified 388 candidate MeJA-responsive unique transcripts. Transcript analysis suggests that in addition to controlling its own biosynthesis and stress responses, MeJA up-regulates transcripts of the various secondary metabolic pathways, including terpenoids and phenylpropanoids/flavonoids. Furthermore, combined transcript and metabolite analysis revealed MeJA-induced biosynthesis of the medicinally important ursane-type and oleanane-type pentacyclic triterpenes. Two MeJA-responsive oxidosqualene cyclases (*ObAS1* and *ObAS2*) that encode for 761- and 765-amino acid proteins, respectively, were identified and characterized. Functional expressions of *ObAS1* and *ObAS2* in *Saccharomyces cerevisiae* led to the production of β -amyrin and α -amyrin, the direct precursors of oleanane-type and ursane-type pentacyclic triterpenes, respectively. *ObAS1* was identified as a β -amyrin synthase, whereas *ObAS2* was a mixed amyrin synthase that produced both α -amyrin and β -amyrin but had a product preference for α -amyrin. Moreover, transcript and metabolite analysis shed light on the spatiotemporal regulation of pentacyclic triterpene biosynthesis in sweet basil. Taken together, these results will be helpful in elucidating the secondary metabolic pathways of sweet basil and developing metabolic engineering strategies for enhanced production of pentacyclic triterpenes.

Plants are capable of synthesizing an amazing diversity of low molecular weight secondary metabolites that have a variety of applications as pharmaceuticals, flavors, fragrances, and pesticides (Bohlmann and Keeling, 2008; Wink, 2010a). Although some plant-derived secondary metabolites are directly used as drugs, many are leading models for the development of semisynthetic and synthetic drugs (Heilmann, 2010). To date, over 100,000 different structures of secondary metabolites or natural products are known; however, the real number is undoubtedly much higher because only 20% to 30% of the plant species are exploited (Wink, 2010a). Because of the insufficient supply

of raw materials for extraction of many valuable secondary metabolites, increasing the production of these compounds in plants is a well-established target for genetic manipulation through traditional and biotechnological approaches (Goossens et al., 2003; Canter et al., 2005; Alfermann, 2010; Lange and Ahkami, 2013). However, this presents some severe challenges owing to the lack of in-depth knowledge on the secondary metabolite biosynthetic pathways and their regulation (Canter et al., 2005; Lange and Ahkami, 2013). A comprehensive understanding of the metabolic pathway is extremely important to overcome low product yield of the secondary metabolites in plants or plant cell cultures. Once the biosynthetic genes and their regulatory controls have been elucidated, pathway engineering can be employed to increase metabolite accumulation not only in the native plants but also in the organisms that do not even synthesize them (Yun et al., 1992; Chen et al., 2000; Zhang et al., 2004; Lange and Ahkami, 2013; Paddon et al., 2013).

Biologically active phytochemicals are synthesized in plants in tissue-specific, organ-specific, and developmentally specific ways that involve highly complex and sophisticated biosynthetic pathways (Facchini and De Luca, 2008; Murata et al., 2008; Besser et al., 2009; Wink, 2010a). Plants synthesize and accumulate these compounds in specific tissue types perhaps for autotoxicity avoidance, because many of the secondary metabolites are known to be phytotoxic (Reigosa and Malvido-Pazos,

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2007; Ghosh et al., 2013). Moreover, the sites of biosynthesis may not necessarily be the sites of storage, and long-distance transport via the xylem, phloem, or apoplast is also possible (Wink, 2010b). Lipophilic metabolites are usually stored in the cuticle, resin ducts, laticifers, trichomes, or oil cells, whereas hydrophilic metabolites are stored in the vacuole. When plants experience various biotic and abiotic stresses, biosynthesis of the secondary metabolites is triggered that helps plants adapt to the challenging environment (Gershenzon and Dudareva, 2007; Hartmann, 2007; Reichling, 2010). Consequently, transcript and metabolite profiling of stress/elicitor-treated plants or cell cultures represents a powerful approach to determine gene function in the biosynthesis of the secondary metabolites (Aerts et al., 1994; Goossens et al., 2003; Suzuki et al., 2005; Zhao et al., 2005; Naoumkina et al., 2008; De Geyter et al., 2012; Lenka et al., 2012; Yu et al., 2012; Ee et al., 2013; Mishra et al., 2013; Sun et al., 2013).

The plant hormone methyl jasmonate (MeJA) acts as a conserved elicitor of secondary metabolite production across the plant kingdom, from angiosperms to gymnosperms (De Geyter et al., 2012). Several studies have demonstrated that MeJA treatment can trigger the biosynthesis of all three major classes of secondary metabolites (i.e. terpenoids, phenylpropanoids, and alkaloids) through an extensive transcriptional reprogramming of the plant metabolism (Gundlach et al., 1992; Cheong and Choi, 2003; Zhao et al., 2005; Pauwels et al., 2009; De Geyter et al., 2012; Sun et al., 2013). Here, we carried out transcriptome and selected metabolite analyses to identify the MeJA-responsive secondary metabolic pathways of sweet basil (*Ocimum basilicum*), which is a plant of the *Lamiaceae* family with highly valued aromatic and medicinal properties. Sweet basil has been extensively used for centuries in traditional medicine for the treatment of various ailments (Bilal et al., 2012). Several pharmacological studies have demonstrated that sweet basil contains biologically active constituents that are analgesic, anti-inflammatory, antimicrobial, antioxidant, antiulcerogenic, anticarcinogenic, cardiac stimulant, chemomodulatory, central nervous system depressant, hepatoprotective, hypoglycemic, hypolipidemic, immunomodulatory, and larvicidal (Dasgupta et al., 2004; Muralidharan and Dhananjayan, 2004; Lee et al., 2005; Zeggwagh et al., 2007; Meera et al., 2009; Zhang et al., 2009; Arshad Qamar et al., 2010; Choudhury et al., 2010; Dashputre and Naikwade, 2010; Fathiazad et al., 2012).

Sweet basil essential oil contains numerous volatile compounds, including monoterpenes, sesquiterpenes, and phenylpropanoids (Iijima et al., 2004a; Zhang et al., 2009; Verma et al., 2012). In addition, various nonvolatile phytochemicals such as triterpenes, rosmarinic acid, and flavonoids that are present in considerable amounts in sweet basil are shown to have prominent biological properties (Siddiqui et al., 2007; Arshad Qamar et al., 2010; Strazzer et al., 2011; Fathiazad et al., 2012). Although efforts have been made to understand the biochemical and molecular basis of volatile terpene and

phenylpropanoid biosynthesis in *Ocimum* species (Gang et al., 2001, 2002a, 2002b; Iijima et al., 2004a, 2004b; Louie et al., 2007; Rastogi et al., 2013), our knowledge of how nonvolatile compounds are biosynthesized has remained largely incomplete.

Ursane-type and oleanane-type pentacyclic triterpenes (C₃₀ compounds) are known for a variety of pharmaceutical properties, including anti-inflammatory, hepatoprotective, antitumor, antihyperlipidemic, antiulcer, and antimicrobial benefits (Kunkel et al., 2011; Wilkinson et al., 2011; Kurek et al., 2012; da Silva Ferreira et al., 2013; Kong et al., 2013). However, there are limited data on the biosynthesis of these secondary metabolites in plants (Brendolise et al., 2011; Huang et al., 2012; Yu et al., 2013). Moreover, the regulatory controls of these secondary metabolic pathways are completely unknown. *Ocimum* species are known to accumulate these pentacyclic triterpenes, although the biosynthetic genes are yet to be identified and characterized (Silva et al., 2008).

Here we report the identification and characterization of the MeJA-responsive transcripts in sweet basil using a suppression subtractive hybridization (SSH) approach. A total of 388 candidate MeJA-responsive unique transcripts were isolated, which represent various secondary metabolic pathways (e.g. terpenoid, phenylpropanoid/flavonoid), in addition to transcripts associated with plant stress responses. Furthermore, transcript and metabolite analysis revealed MeJA responsiveness of ursane-type and oleanane-type pentacyclic triterpenes (ursolic acid and oleanolic acid), whose biosynthesis is spatiotemporally regulated in sweet basil. Moreover, cloning and functional characterization of two MeJA-responsive oxidosqualene cyclases revealed their involvement in the biosynthesis of the direct precursors (α -amyrin and β -amyrin) of these medicinally important triterpenes.

RESULTS

Cloning and Characterization of the MeJA-Responsive Transcripts

To identify sweet basil transcripts expressed in response to MeJA treatment, the SSH approach was followed because of its high efficiency in enriching low-expressing genes and normalization of the targeted sequences. An SSH library was generated from 2-month-old plants treated with 250 μ M MeJA versus control plants, as described in the materials and methods. A total of 550 individual clones was randomly selected from the SSH library for sequencing using a vector-specific M13 forward primer. This resulted in the isolation of 509 high-quality ESTs with an average sequence length of 481.59 bp, after removing low-quality, vector, and short sequences (Table I). The average guanine-cytosine (GC) content of these high-quality ESTs was 43.92%, which was comparable with that observed for other plant species (Asamizu et al., 2000). EST sequences were deposited in the National Center for Biotechnology Information (NCBI) GenBank database (accession numbers JZ190506

Table 1. An overview of the sweet basil MeJA SSH library

Description	No.
Total clones sequenced	550
High-quality ESTs obtained	509
Average length of ESTs (bp)	481.59
GC content of ESTs (%)	43.92
Unique sequences	388
Average length of unique sequence (bp)	478.82
Total contigs	74
Total singletons	314
Annotated unique sequences ^a	323
Nonannotated unique sequences	65
Novel sweet basil unique ESTs ^b	275

^aBLASTX analysis against NCBI nonredundant protein sequence database. ^bBLASTN analysis against NCBI EST database.

to JZ191014). Furthermore, using the CAP3 program, ESTs were assembled into a total of 388 unique sequences, including 74 contigs and 314 singletons. The contigs consisted of ESTs ranging from 2 to 28, thereby suggesting a high level of MeJA-inducible transcription associated with the corresponding genes, such as mevalonate kinase, oxidosqualene cyclase, lectin, global transcription factor group E, putative ethylene-responsive element-binding protein-type transcription factor, and so forth. The detailed functional annotation of the unique ESTs is presented in Supplemental Table S1. BLASTN analysis of the unique sequences against the sweet basil EST database at the NCBI (<http://www.ncbi.nlm.nih.gov>) revealed that 70.87% of the unique ESTs are novel. To confirm the MeJA-induced expression, some of the transcripts were subjected to semiquantitative reverse transcription PCR (sqRT-PCR) and quantitative reverse transcription PCR (qRT-PCR) analysis (Figs. 1D and 2C).

Gene Ontology (GO) annotation was performed to assign putative function to the gene products (Fig. 1, A–C). Altogether, 73.97% of the unique sequences could be functionally categorized based on the GO biological process, cellular component, and molecular function (<http://www.blast2go.com>). The majority of the transcripts were predicted to be associated with the plastid (26.8%), plasma membrane (14.95%), nucleus (10.82%), cytosol (10.31%), mitochondria (10.31%), and vacuole (7.47%), suggesting that these cellular compartments play an important role in mediating MeJA-induced modulation of the cellular metabolic pathways. On the other hand, the majority of the unique sequences under the molecular function category were assigned to binding (20.36%) and catalytic activity (18.56%). Transferase, oxidoreductase, and hydrolase were found to be the major enzyme classes (Supplemental Table S2). Under the biological process, categories such as biosynthetic process (20.1%), response to stress (15.98%), catabolic process (14.95%), and transport (11.08%) were highly represented. This finding highlighted the key role of MeJA in regulating cellular metabolism and stress responses in plants (Cheong and Choi, 2003; Pauwels et al., 2009).

Few transcriptional regulators of APETALA2/Ethylene Response Factor (ERF), WRKY, Plant Homeo Domain

(PHD), and Zinc Finger families were identified in the SSH library (Supplemental Table S3). Functional characterization of these MeJA-responsive transcription factors may lead to a better understanding of the underlying transcriptional regulation of MeJA-induced alteration in cellular metabolism. The SSH library was also enriched with the transcripts that presumably have no direct role in cellular metabolism. Some of them were related to the plant defense responses and are known to be induced by MeJA in other plant species as well (Supplemental Table S4; Cheong and Choi, 2003; Lenka et al., 2012; Ee et al., 2013).

Identification of the Metabolic Pathways

To gain an overview of the sweet basil metabolic pathways that are modulated by MeJA, unique ESTs were analyzed according to the Kyoto Encyclopedia of Genes and Genomes (KEGG; <http://www.genome.jp>), which provides an alternative functional annotation of the transcripts based on the associated biochemical pathways. The analysis revealed a total of 77 KEGG pathways (Supplemental Table S5). Interestingly, six transcripts were found to be associated with the α -linolenic acid metabolism that leads to jasmonic acid biosynthesis (Supplemental Fig. S1; Supplemental Table S5). These transcripts included 13S-lipoxygenase (EC:1.13.11.12), C-acyltransferase (EC:2.3.1.16), and oxidase (EC:1.3.3.6), with three, two, and one unique ESTs, respectively. These results support the existence of a positive feedback regulatory system for the biosynthesis of jasmonic acid (Sasaki et al., 2001).

A list of various secondary metabolic pathways represented by the unique ESTs is provided in Table II. Interestingly, among the secondary metabolic processes, terpenoid and phenylpropanoid/flavonoid biosynthesis pathways were found to be maximally represented (Table II; Supplemental Fig. S2). These secondary metabolite types were previously reported to accumulate in high levels in sweet basil after MeJA treatment; however, the genes involved in the biosynthesis of these compounds were not identified (Kim et al., 2006; Li et al., 2007).

MeJA Responsiveness of the Pentacyclic Triterpene Biosynthetic Pathway

In plants, all diverse types of terpene metabolites are synthesized from two 5C building blocks (dimethylallyl diphosphate and isopentenyl diphosphate) that are formed via two independent pathways: the mevalonic acid (MEV) pathway and the 2C-methyl-D-erythritol-4-phosphate (MEP) pathway (Tholl and Lee, 2011). Isopentenyl diphosphate and dimethylallyl diphosphate derived from the cytosolic MEV pathway serve as the precursors for the biosynthesis of the sesquiterpenes (C15) and triterpenes (C30), whereas the plastidial MEP pathway provides precursors for the biosynthesis of the monoterpenes (C10), diterpenes (C20), and tetraterpenes

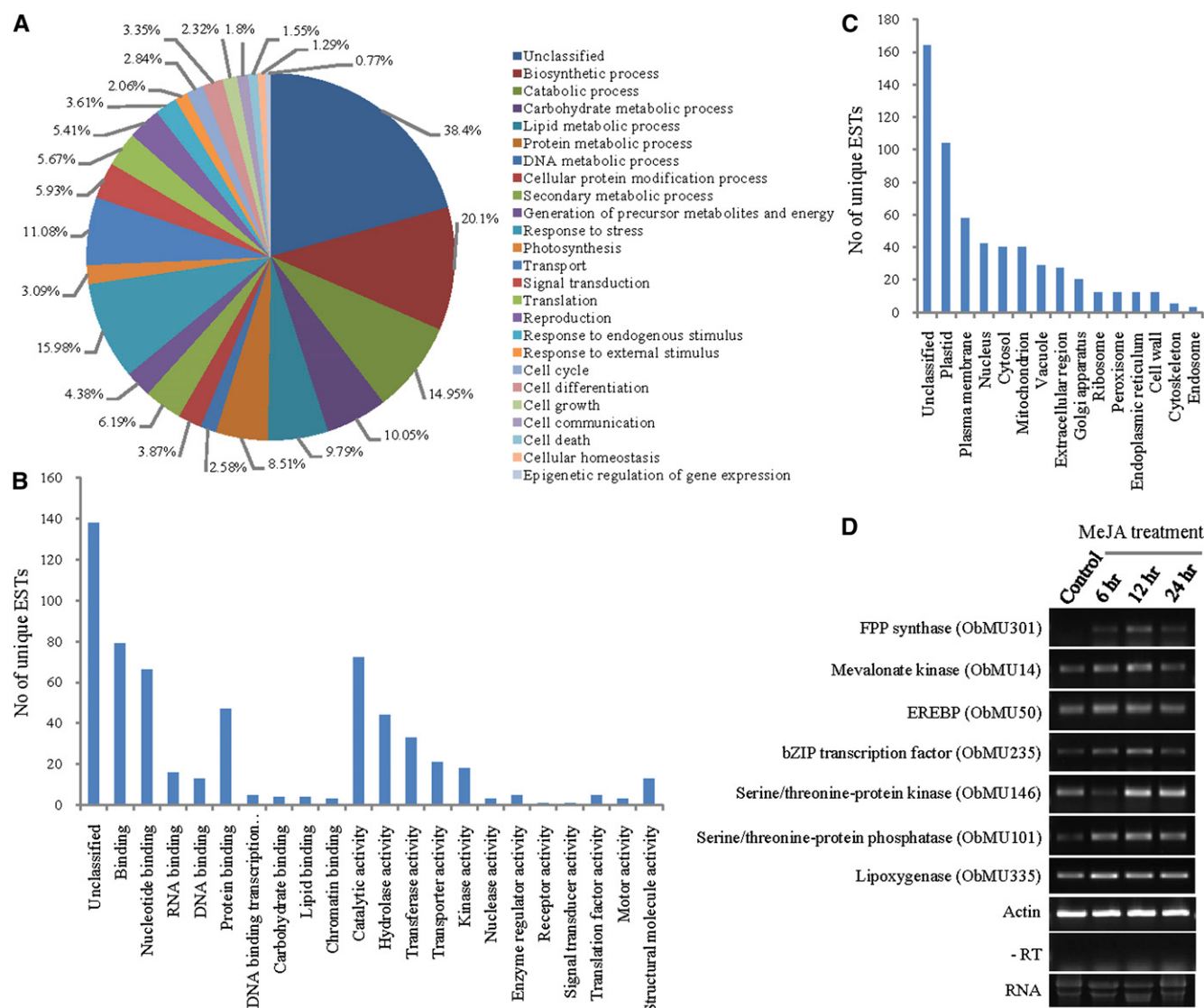
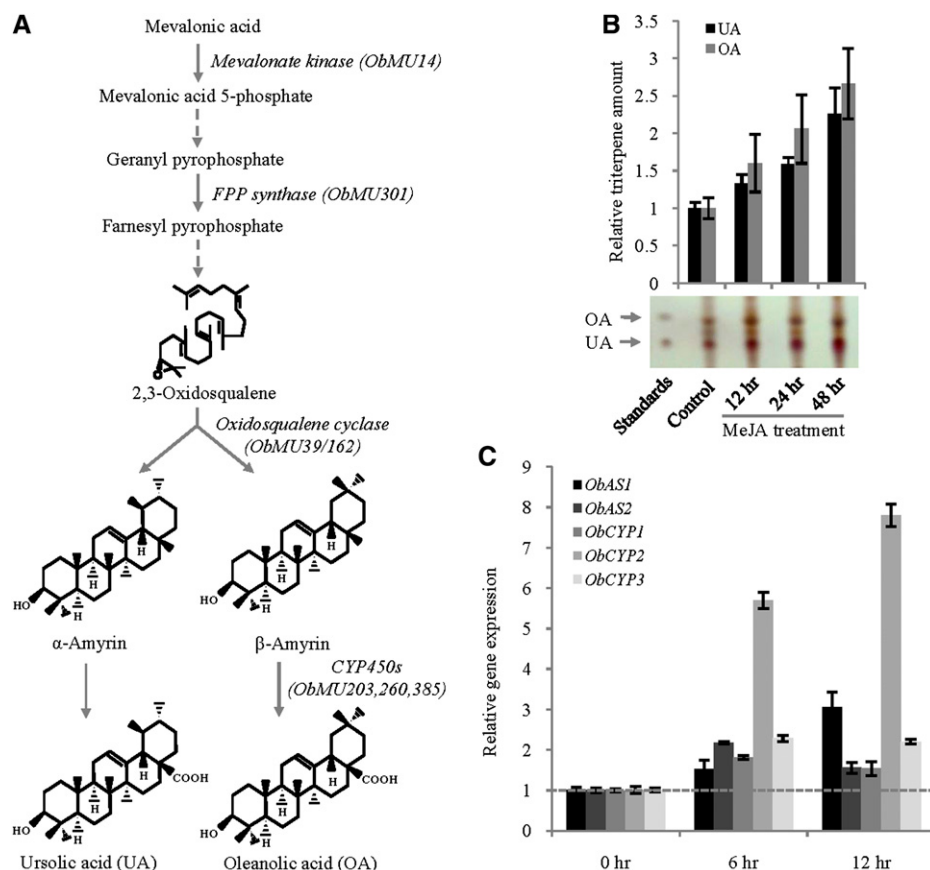


Figure 1. Functional categorization of the MeJA-responsive transcripts. A to C, GO classification of 388 MeJA-responsive transcripts on the basis of the biological process (A), molecular function (B), and cellular component (C). D, MeJA-inducible expression of the transcripts was detected by RT-PCR analysis. Actin amplification was used to normalize the amount of template. The negative control lacked reverse transcriptase in cDNA synthesis reaction (–RT). The experiment was repeated twice independently with similar results. bZIP, basic leucine zipper; EREBP, ethylene-responsive element-binding protein; FPP, farnesyl pyrophosphate. [See online article for color version of this figure.]

(C40) (Sawai and Saito, 2011). The identification of the transcripts similar to mevalonate kinase (three ESTs) and 4-diphosphocytidyl-2-C-methyl-D-erythritol kinase (enzymes of the MEV and MEP pathways, respectively) suggests the up-regulation of these pathways in sweet basil after MeJA treatment. Moreover, MeJA induction of the pentacyclic triterpene biosynthesis was envisaged because the SSH library also represented one EST for putative farnesyl diphosphate synthase and three ESTs for putative oxidosqualene cyclase (OSC) with homology to amyrin synthases of other plant species (Fig. 2A). Furthermore, three cytochrome P450s of the CYP716A subfamily were identified in the SSH library (Fig. 2A).

CYP716A subfamily cytochrome P450 enzymes were recently found to be involved in triterpene biosynthesis in *Panax notoginseng*, *Medicago truncatula*, and *Catharanthus roseus* (Carelli et al., 2011; Han et al., 2011; Huang et al., 2012). To confirm MeJA induction of the pentacyclic triterpene biosynthesis in sweet basil, the levels of ursolic acid and its isomer oleanolic acid were determined by thin-layer chromatography (TLC) analysis (Supplemental Fig. S3; Wójciak-Kosior, 2007). The analysis revealed 2.2- and 2.6-fold higher level accumulation of ursolic acid and oleanolic acid, respectively, in sweet basil leaves at 48 h after MeJA treatment (Fig. 2B). The MeJA-inducible expression of the candidate pentacyclic triterpene

Figure 2. MeJA responsiveness of the pentacyclic triterpene biosynthetic pathway. **A**, Ursolic acid and oleanolic acid biosynthetic pathway. Transcripts identified in the MeJA SSH library and the corresponding biosynthetic steps are shown. Solid and dashed arrows denote single and multiple enzymatic steps, respectively. **B**, The relative levels of ursolic acid and oleanolic acid in sweet basil leaves were determined by TLC analysis after MeJA treatment for the indicated time periods. Data are the mean $\pm$ SE from three to four independent experiments. The photograph of one representative TLC plate is presented. **C**, qRT-PCR expression analysis of the OSCs (*ObAS1*, *ObAS2*) and cytochrome P450s (*ObCYP1*, *ObCYP2*, *ObCYP3*) to show their MeJA-inducible expression. Data are mean $\pm$ SE from two independent experiments ($n = 5$ to 6). FPP, farnesyl pyrophosphate; OA, oleanolic acid; UA, ursolic acid. [See online article for color version of this figure.]



biosynthesis genes (amyrin synthases and CYP716A subfamily cytochrome P450 enzymes) of sweet basil was validated by qRT-PCR analysis (Fig. 2C). Altogether, these data suggest MeJA responsiveness of the ursane-type and oleanane-type pentacyclic triterpene biosynthetic pathways in sweet basil.

Cloning and Sequence Analysis of the OSCs

2,3-Oxidosqualene, the precursor of ursolic acid and oleanolic acid, serves as the endogenous substrate for the enzyme OSC that produces α -amyrin and/or β -amyrin, which then act as the substrates for the cytochrome P450 enzymes for the biosynthesis of the ursolic acid and oleanolic acid, respectively (Fig. 2A). To shed light on pentacyclic triterpene biosynthesis in sweet basil, two MeJA-responsive unique ESTs encoding putative OSCs (ObMU39 and ObMU162) were targeted for the functional analysis. The full-length complementary DNAs (cDNAs) specific to the OSCs were obtained from leaf total RNAs by a combination of reverse transcription PCR (RT-PCR) amplification of the unique sequences and rapid amplification of the cDNA ends (5' and 3' RACE). Analysis of the full-length sequences revealed that both of the unique sequences, ObMU39 and ObMU162, are the common gene products with a match to the 5' and 3' regions, respectively, of the complete cDNA sequence.

To identify additional OSCs of sweet basil, we conducted RT-PCR using degenerate oligonucleotides followed by 5' and 3' RACE. Sequence analysis of the amplified cDNAs revealed one more OSC in addition to the previous one. Both of the complete cDNA sequences were submitted to the GenBank database under the accession numbers KF636411 (*ObAS1*) and JQ809437 (*ObAS2*). Sequence analysis revealed that *ObAS1* and *ObAS2* encode for 60.32% identical polypeptides of 761 and 765 amino acids, respectively, with calculated molecular masses of 87.73 kD and 87.49 kD. *ObAS1* and *ObAS2* were subjected to phylogenetic analysis with the OSCs of other plants (Fig. 3). At the amino acid sequence level, *ObAS1* and *ObAS2* shared 83.05% and 86.61% identities, respectively, with the *Panax ginseng* β -amyrin synthase and the *Olea europaea* mixed-function OSC that catalyzes the production of α -amyrin and β -amyrin as well as small amounts of ψ -taraxasterol and butyrospermol (Kushiro et al., 1998; Saimaru et al., 2007). On the contrary, *ObAS1* and *ObAS2* shared 61.37% and 61.07% identities with the *O. europaea* mixed-function OSC and the *P. ginseng* β -amyrin synthase, respectively. *ObAS1* and *ObAS2* have all of the conserved amino acid motifs characteristic of an OSC (Fig. 4). These include the DCTAE motif that is considered a part of the active site that binds the substrate 2,3-oxidosqualene, the MWCYCR motif, and few β -strand turn QW motifs that are known to be shared by the OSC superfamily (Poralla

Table II. *MeJA-responsive unique ESTs of sweet basil related to secondary metabolite biosynthesis*

Sl No	Contig ID	Length <i>bp</i>	Functional Annotation to GenBank Database	E- Value
Terpenoid				
1	ObMU14	893	Mevalonate kinase (AEZ55674.1)	7.00E-120
2	ObMU39	771	Mixed amyrin synthase (BAF63702.1)	3.00E-160
3	ObMU118	513	Nerolidol synthase (AER36088.1)	3.00E-59
4	ObMU156	480	Acetoacetyl-coenzyme A thiolase (ABV08820.1)	9.33E-109
5	ObMU162	673	Mixed amyrin synthase (AFJ19235.1)	2.00E-138
6	ObMU203	442	Cytochrome P450 CYP716A52v2 (AFO63032.1)	9.00E-31
7	ObMU260	130	Cytochrome P450 CYP716A17 (NP_001268076.1)	4.00E-15
8	ObMU261	169	Linalool synthase (Q5SBP3.1)	7.00E-23
9	ObMU262	726	4-Diphosphocytidyl-2-C-methyl-D-erythritol kinase (ABP96842.1)	2.00E-18
10	ObMU301	589	Farnesyl diphosphate synthase (AAK63847.1)	5.00E-128
11	ObMU385	306	Cytochrome P450 CYP716A52v2 (AFO63032.1)	1.00E-36
Phenylpropanoid/ flavonoid				
12	ObMU78	526	Cytochrome P450 82A3 (XP_002281995.2)	4.00E-51
13	ObMU89	356	Isoflavone reductase homolog P3 (XP_002282110.1)	3.00E-54
14	ObMU96	375	Dihydroflavonol reductase (XP_002276827.1)	2.00E-37
15	ObMU135	80	Flavonol synthase (ABB53382.1)	6.00E-06
16	ObMU350	180	Flavonoid O-methyltransferase 3 (AFU50297.1)	2.00E-27
17	ObMU355	312	Cinnamoyl-CoA reductase (XP_002520066.1)	3.00E-19
18	ObMU374	187	Eugenol synthase (Q15GI4.1)	2.00E-25
Alkaloid				
19	ObMU1	339	Polyphenol oxidase (AAD53122.1)	8.00E-05
20	ObMU187	522	Strictosidine synthase (XP_002273764.1)	7.00E-80
21	ObMU190	221	Polyamine oxidase 2-like (XP_004251556.1)	1.14E-33
22	ObMU253	144	Primary amine oxidase-like (XP_004298688)	5.00E-25
Tocopherol				
23	ObMU131	271	Homogentisate phytyltransferase 2 (XP_002276728.1)	9.00E-25
Carotenoid				
24	ObMU314	331	Cytochrome P450 CYP97C10 (ABD97103.1)	3.00E-61
Others				
25	ObMU92	877	3-Ketoacyl-CoA thiolase 1 (AGC59769.1)	7.19E-163
26	ObMU209	628	Lipoxygenase (XP_002516771.1)	2.00E-73
27	ObMU272	879	Lipoxygenase 6 (XP_003531186.1)	1.00E-177
28	ObMU306	408	Acetoacetyl-CoA thiolase (ABB45810.1)	8.75E-52
29	ObMU335	408	Lipoxygenase (NP_001233812.1)	2.00E-53

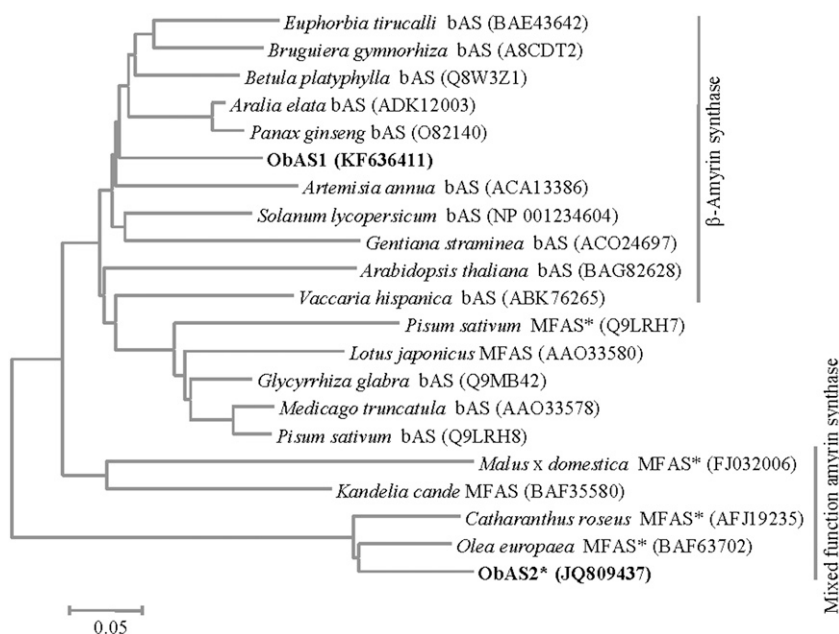
et al., 1994; Abe and Prestwich, 1995). These results suggest that ObAS1 and ObAS2 are the candidate OSCs that possibly provide the direct precursors for ursolic acid and oleanolic acid biosynthesis in sweet basil. Moreover, *ObAS1* and *ObAS2* exhibited MeJA-induced expression, suggesting their potential role in MeJA-induced biosynthesis of these pentacyclic triterpenes in sweet basil (Fig. 2, B and C).

Functional Expression of *ObAS1* and *ObAS2* in Yeast

To determine the product specificities of ObAS1 and ObAS2, the full-length cDNAs (2286 and 2298 nucleotides, respectively) were expressed in yeast (*Saccharomyces cerevisiae* strain BY4741) under the control of the Gal-inducible GAL1 promoter. In yeast, endogenous 2,3-oxidosqualene can serve as an *in vivo* substrate for ObAS1 and ObAS2. The expression of *ObAS1* and *ObAS2* in transgenic yeasts was confirmed by RT-PCR analysis and by purifying poly(His) (6×His)-tagged

proteins by using a nickel-nitrilotriacetic acid agarose column (Fig. 5, A and B). ObAS1 and ObAS2 were resolved as 97-kD recombinant proteins on 10% (w/v) SDS-PAGE (Fig. 5B). This was in agreement with the calculated molecular masses of 6×His-tagged recombinant proteins, indicative of the complete translation of the full-length cDNA sequences in yeast. To determine the enzyme function of ObAS1 and ObAS2, Gal-induced transgenic yeast cells were analyzed for the accumulation of pentacyclic triterpene metabolites. Yeast cells were extracted with hexane after alkaline treatment and metabolites were identified through gas chromatography-mass spectrometry (GC-MS) analysis. Compared with the vector control strain (*S. cerevisiae* transformed with pYES2/NTB empty vector), the yeast cells expressing *ObAS1* and *ObAS2* produced one and two new products, respectively (Fig. 5C). By comparing the retention time and mass spectra of those new products with those of the authentic standards (α -amyrin and β -amyrin), we found that *ObAS1*-expressing transgenic yeast accumulated β -amyrin, which was not

Figure 3. Phylogenetic tree of plant OSCs and their relationships to sweet basil *ObAS1* and *ObAS2*. The complete amino acid sequences of OSCs obtained from the GenBank database, together with that of *ObAS1* and *ObAS2* were analyzed by MEGA5. The phylogenetic tree was built using the Neighbor Joining Method. MFASs with product preference for α -amyrin are marked with an asterisk. bAS, β -amyrin synthase; MFAS, mixed-function amyirin synthase.



present in the control yeast cells transformed with empty vector (Fig. 5, C and E). By contrast, *ObAS2*-expressing transgenic yeast accumulated both α -amyrin and β -amyrin as new products in a 5:3 ratio (Fig. 5, C–E). The yeast expression experiments were repeated three times, always with very similar results. These results clearly indicated that *ObAS1* encodes for a β -amyrin synthase, whereas *ObAS2* encodes a mixed-function amyirin synthase with a product preference for α -amyrin.

Spatial and Temporal Regulation of Pentacyclic Triterpene Biosynthesis

To determine the involvement of *ObAS1* and *ObAS2* in the biosynthesis of ursane-type and oleanane-type pentacyclic triterpenes in sweet basil, the spatiotemporal expression patterns of these genes were correlated with in planta metabolite accumulation patterns. The relative transcript levels of *ObAS1* and *ObAS2* in different plant organs, tissue types, and leaf developmental ages were determined by qRT-PCR analysis and compared with the ursolic acid and oleanolic acid accumulation patterns (Figs. 6, 7, and 8). The level of these metabolites in sweet basil was determined by TLC analysis after iodine derivatization (Supplemental Fig. S3; Wójciak-Kosior, 2007).

In some plant species, pentacyclic triterpenes accumulate in a substantial amount in surface cuticular wax, possibly because of epidermal cell specialization of the biosynthetic pathways (Murata et al., 2008; Yeats and Rose, 2013). The cuticular wax component of sweet basil was selectively extracted from stems and leaves by brief dipping in chloroform and analysis was conducted with TLC (Fig. 6A). The presence of ursolic acid and oleanolic acid in surface extracts suggests epidermal cell specialization of their biosynthesis in sweet basil. To

substantiate these results, the expression levels of *ObAS1* and *ObAS2* were determined in leaf epidermal cells using the small subunit of Rubisco, a marker for the mesophyll cells, as the negative control. The carborundum abrasion method (Levac et al., 2008) was followed to harvest leaf epidermis-enriched RNAs and the relative gene expression was determined by qRT-PCR analysis (Fig. 6, B and C). Compared with the whole leaf, *ObAS1* and *ObAS2* exhibited 4.1- and 3.4-fold higher expression levels in the leaf epidermis, respectively (Fig. 6B). By contrast, the expression level of Rubisco small subunit was very low in the leaf epidermis compared with the whole leaf (Fig. 6C). Together, these results suggest epidermal cell localization and possible involvement of *ObAS1* and *ObAS2* in the biosynthesis of ursolic acid and oleanolic acid.

The expression levels of *ObAS1* and *ObAS2* in different plant organs and leaves of different developmental ages were determined and correlated with the ursolic acid and oleanolic acid levels (Figs. 7 and 8). *ObAS1* and *ObAS2* showed the highest expression in inflorescences and leaves, respectively, and the lowest expression in roots (Fig. 7B). On the other hand, both genes exhibited higher expression levels in older leaves compared with the youngest leaves (Fig. 8C). These differential expression patterns of *ObAS1* were quite well correlated with the accumulation profiles of oleanolic acid in plant organs and leaves of different developmental ages (Figs. 7A and 8B). Oleanolic acid accumulated 3.6 times more in inflorescences compared with the roots and 1.8 to 3.5 times more in older leaves compared with the youngest leaves. Therefore, enzyme function (Fig. 5) and gene expression patterns of *ObAS1* (Figs. 6–8) strongly suggest that this β -amyrin synthase is involved in the biosynthesis of oleanolic acid in sweet basil. However, the level of ursolic acid was found to be 2.7 times higher in inflorescences compared with the leaves and the

		QXXXGXW		QXXXGXW		MMCYCR		QXXXGXW						
Malus	107	ALQSPHGHWPA	157	NHQNEDGGWGL	236	NMPPEFW	254	SKMFCYRLT	361	NSRYITIGVEKPL	410	QSFGSQSW	443	HEFLKISQVR
Kandelia	107	ALQASDGHWCA	157	CHQNEDEGGWGL	236	NMPPEFW	254	AKMLCYRLT	361	NSRYITIGVEKPL	410	QSFGSQAW	443	HNFIKDSQVT
Solanum	107	ALQATDGHWPA	157	CHQNEDEGGWGL	236	NMPPEFW	254	AKMWCYRMV	361	NSRYITIGVEKVL	410	QSFGSQEW	443	HDFIKSQSVT
Gentiana	107	ALQSSDGHWPA	157	CHQNEDEGGWGL	236	NMPPEFW	254	GKMGYCYRMV	361	NSRYITVGSVEKVL	410	QSFGSQLW	443	HDFIKNSQVK
Aralia	109	ALQASDGHWPA	159	CHQNEDEGGWGF	238	NMPPEFW	256	AKMWCYRMV	363	NSRYITIGVEKVL	412	QSFGSQEW	445	HDFIKKSQVK
Panax	109	ALQASDGHWPA	159	CHQNEDEGGWGL	238	NMPPEFW	256	AKMWCYRMV	363	NSRYITIGVEKVL	412	QSFGSQEW	445	HDFIKKSQVK
Euphorbia	107	ALQASDGHWPA	157	CHQNEDEGGWGL	236	NMPPEFW	254	AKMWCYRMV	362	NSRYITIGVEKVL	411	QSFGSQEW	444	HDFIKKSQVK
Bruguiera	107	ALQASDGHWPA	157	YHQNEDEGGWGL	236	NMPPEFW	254	AKMWCYRMV	362	SSRYITIGVEKVL	411	QSFGSQEW	444	HDFIKKSQVR
Betula	107	ALQASDGHWPA	157	YHQNEDEGGWGL	236	NMPPEFW	254	AKMWCYRMV	362	NSRYITIGVEKVL	411	QSFGSQEW	444	HDFIKKSQVK
ObAS1	107	ALQASDGHWPA	157	CHQNEDEGGWGF	236	NMPPEFW	254	AKMWCYRMV	361	NSRYITIGVEKVL	410	QSFGSQEW	443	HEFIKNSQVK
Artemisia	107	ALQADDGHWPA	157	CHQNEDEGGWGF	236	NMPPEFW	254	AKMWCYRLV	361	NSRYITIGSVEKAL	410	QSFGSQEW	443	HEFIKASQVK
Vaccaria	107	AIQAEDGHWPA	157	YHQNEDEGGWGL	236	NMPPEFW	254	AKMWCYRMV	362	NSRYITIGVEKVL	411	QSFGSQEW	445	HYFIKESQVK
Glycyrrhiza	107	ALQTSDDGHWPA	157	YHQNEDEGGWGL	236	NMPPEFW	254	AKMWCYRLV	362	TSRYITIGVEKVL	411	QSFGSQEW	444	HDFIKKSQVR
Medicago	107	ALQTSDDGHWPA	157	CHQNEDEGGWGL	236	NMPPEFW	254	AKMWCYRLV	362	NSRYLTIGVEKVL	411	QSFGSQEW	444	HDFIKKSQVT
Lotus	107	GLQTSDDGHWPA	157	VHQNEDEGGWGL	236	NMPPEFW	254	AKMWCYRLV	362	NSRYITIGVEKVL	411	QSFGSQEW	444	HDFIKKSQVK
Pisum	107	TLQSDNDGHWPA	157	CHQNEDEGGWGL	236	NMPPEFW	254	AKMLCYRLV	362	NSRYLTIGVEKVL	411	HSFGSQEW	444	HEFIKKSQVR
Arabidopsis	108	ALQASDGHWPA	158	CHQKEDGGWGL	237	NMPPEFW	255	AKMWSYRMV	363	NSRYITIGVEKVL	412	QSFGSQLW	445	HEFIKNSQVG
Olea	109	AVQASDGHWPA	159	NHQNDGGWGF	237	NPLPEFW	255	AKMWCYRRT	362	ESRYITIGVEKSL	411	QSFGSQEW	444	HFYIKESQIK
ObAS2	112	AVQASDGHWPA	162	NHQNEDEGGWGF	240	NPLPEFW	258	AKMWCYRRT	365	ESRYITIGVEKIL	414	QSFGSQLW	447	HFYIKESQVK
Catharanthus	109	AIQATDGHWPA	159	NHQNEDEGGWGF	237	NPLPEFW	255	AKMWCYRRT	362	ESRYITIGVEKSL	411	QSFGSQLW	444	HFYIKESQVK
		*: .*** *		**::*****:		***:***:*		..* .*** .		***:***:*		:***** *		*::* **:

		DCTAE		QXXXGXW		QXXXGXW				
Malus	464	RHISKGAWTFSDRDHGWQ	482	VSDCTAEAL	599	DIQYPDGSWYGNWG	648	KTQRADGGWGE	712	DDGDFPQQE
Kandelia	464	RHISKGSWTFSDKDHWGQ	482	VSDCTAESL	599	EIQKPDGSWYGNWG	648	RTQREDGGWGE	712	ELGDFPQQE
Solanum	464	RHISKGSWTFSDQDHWGQ	482	VSDCTAEAL	599	DVQMPDGSWYGNWG	648	RTQREDGGWGE	712	EDGDFPQQE
Gentiana	464	RHMSKGSWTFSDQDHWGQ	482	VSDCTAEBL	599	DVQMPDGSWYGNWG	648	TTQKQDGGWGE	712	ENGDFPQQE
Aralia	466	RHISKGSWTFSDQDHWGQ	484	VSDCTAEBL	601	DIQMPDGSWYGNWG	650	KSQMDGGWGE	714	EDGDFPQQE
Panax	466	RHISKGSWTFSDQDHWGQ	484	VSDCTAEBL	601	DTQMPDGSWYGNWG	650	KSQMDGGWGE	714	EDGDFPQQE
Euphorbia	465	RHISKGSWTFSDQDHWGQ	483	VSDCTAEBL	600	DVQTADGGWYGNWG	649	RTQKQDGGWGE	713	EDGDFPQQE
Bruguiera	465	RHISKGSWTFSDQDHWGQ	483	VSDCTAEBL	600	SIQTSDDGGWYGNWG	649	RIQRDNGWGE	713	EDGDFPQQE
Betula	465	RHISKGSWTFSDQDHWGQ	483	VSDCTAEBL	600	VIQMPDGSWYGNWG	649	RAQRDNGWGE	713	EDGDFPQQE
ObAS1	464	RHISKGSWTFSDQDHWGQ	482	VSDCTAEBL	599	DVQRPDGSWYGNWG	648	KSQREDGGWGE	712	EDGDFPQQE
Artemisia	464	RHISKGSWTFSDQDHWGQ	482	VSDCTAEAL	599	KIQMEDGSWYGNWG	648	ETQLEDGGWGE	712	ETGDFPQQE
Vaccaria	466	RHISKGSWTFSDQDHWGQ	484	VSDCTAEBL	601	DTQYPNGSWYGNWG	650	KSQKEDGGWGE	714	ESGDFPQQE
Glycyrrhiza	465	RHISKGSWTFSDQDHWGQ	483	VSDCTAEBL	600	DTQTADGSWYGNWG	649	TTQREDGGWGE	713	EEGDWPPQE
Medicago	465	RHISKGSWTFSDQDHWGQ	483	VSDCTAEBL	600	DIQTADGSWYGNWG	649	TTQREDGGWGE	713	EEGDWPPQE
Lotus	465	RHISKGAWTFSDQDHWGQ	483	VSDCTAEBF	600	DIQTADGSWYGNAG	649	TTQSKDGGWGE	713	EDGDWPPQE
Pisum	465	RHISKGSWTFSDKDHWGQ	483	VSDCTAESL	600	YKQTSDDGSWYGNWG	649	TTQREDGGWGE	713	EEGDWPPQE
Arabidopsis	466	RHISKGAWTFSDRDHWGQ	484	VSDCTAHL	601	NMQTRDGSWYGNWG	650	AAQKNDGGWGE	714	ESGDFPQQE
Olea	466	RHFTKGAWTFSDQDQGWV	484	VSDCTAEAL	601	GRQWPDGSWYGYWG	650	STQNEEGWGE	714	DDGDFPQQE
ObAS2	469	RHFTKGSWTFSDQDQGWV	487	VSDCTAEAL	604	GKQWPDGSWYGYWG	653	STQNEEGWGE	717	DDGDFPQEE
Catharanthus	466	RHFTKGSWTFSDQDQGWV	484	VSDCTAEAL	601	GRQWPDGSWYGYWG	650	STQNEEGWGE	714	DDGDFPQQE
		::*:*****:***		*****:..:		*::*.*** *		*::*.*** *		:***:***:

Figure 4. Amino acid sequence comparisons of the sweet basil ObAS1 and ObAS2 with plant OSCs. Multiple sequence alignment was carried out by ClustalW2 and manually edited to present the regions having high homology (complete alignment is presented in Supplemental Figure S4). Identical amino acid residues are depicted with an asterisk, whereas conserved and semiconserved substitutions are marked with a colon and period, respectively. [See online article for color version of this figure.]

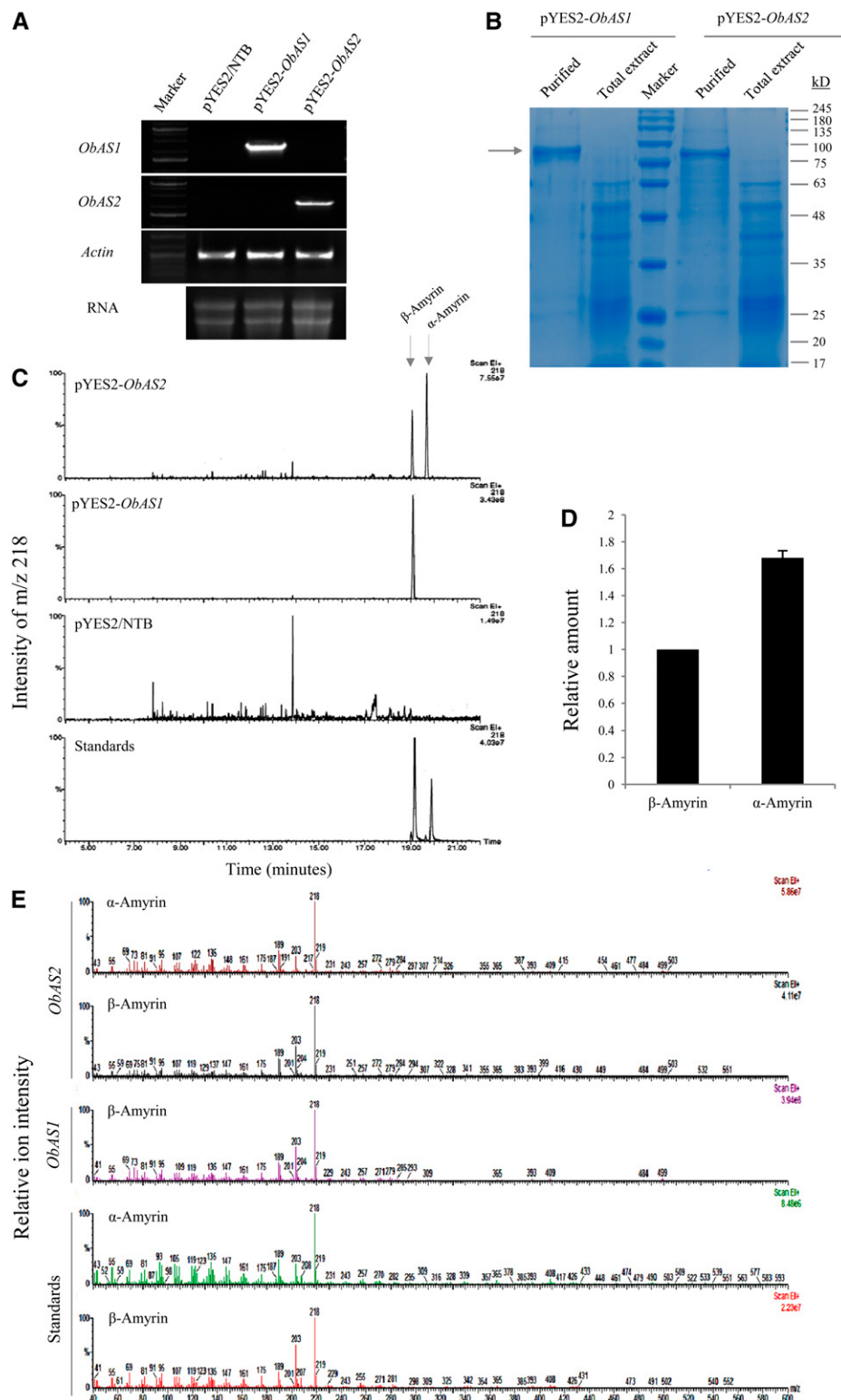
highest level was detected in the youngest leaves compared with the older leaves. The role of *ObAS2* for in planta biosynthesis of the ursolic acid was envisaged based on the epidermal cell enrichment of the *ObAS2* mRNA and accumulation of the ursolic acid in the plant surface (Fig. 6, A and B). However, it is also possible that the accumulation of ursolic acid in different plant organs and during leaf development is also influenced by its catabolism/metabolism and transport after biosynthesis. Moreover, the presence of the additional α -amyrin synthase that is involved in high-level biosynthesis of the ursolic acid in inflorescences and youngest leaves cannot be excluded, because very limited transcript information is available and the genome of sweet basil is yet to be sequenced.

DISCUSSION

Although a number of structurally and functionally diverse classes of secondary metabolites with important

pharmacological traits are identified from sweet basil (Iijima et al., 2004a; Meera et al., 2009; Zhang et al., 2009; Arshad Qamar et al., 2010; Choudhury et al., 2010; Dashputre and Naikwade, 2010; Verma et al., 2012), their biosynthetic pathways remained largely unexplored. The plant endogenous signaling molecule MeJA, in addition to regulating the normal plant developmental processes, induces defense responses during herbivore and pathogen attacks by accumulating diverse secondary metabolites (Cheong and Choi, 2003). Because MeJA has been shown to be a powerful inducer of secondary metabolite production in diverse plants (Aerts et al., 1994; Suzuki et al., 2005; Kim et al., 2006; De Geyter et al., 2012; Lenka et al., 2012), we studied MeJA-elicited transcriptional changes with the goal to identify the candidate genes involved in the biosynthesis of secondary metabolites in sweet basil. SSH is a powerful genomics tool for identification of differentially expressed genes for nonmodel organisms such as sweet basil, in which genomic information is currently scarce (Ohlrogge and Benning, 2000). The SSH approach

Figure 5. Functional expression of *ObAS1* and *ObAS2* in yeast and identification of the amyrin products. A and B, The expression of the full-length cDNAs in *S. cerevisiae* was confirmed by RT-PCR analysis (A) and purifying 6×His-tagged recombinant proteins (B). C, GC-MS analysis of hexane extracts of yeast cells expressing *ObAS1* and *ObAS2*. Yeast transformed with empty vector pYES2/NTB was used as the control. Products were monitored on the basis of intensity of the base peak (mass-to-charge ratio of 218) and retention time using a mixture of α -amyrin and β -amyrin as standards. Total ion chromatograms are presented in Supplemental Fig. S5. D, The relative comparison of α -amyrin and β -amyrin accumulated in the *ObAS2*-expressing yeast strain based on GC peak area. Data are the mean $\pm$ SE from three biological replicates. E, Mass spectra of the peaks correspond to α -amyrin and β -amyrin in standards and *ObAS1* and *ObAS2* expressing yeast extracts are compared. [See online article for color version of this figure.]



led to the isolation of 388 candidate MeJA-responsive transcripts that included several (70.87%) new transcripts of sweet basil. Many of these transcripts comprise the secondary metabolic pathways, including terpenoids and phenylpropanoids/flavonoids, in addition to the

transcripts that are associated with MeJA biosynthesis and plant stress responses (Table II; Supplemental Figs. S1 and S2; Supplemental Tables S1 and S4).

Some of the MeJA up-regulated transcripts are potential candidates for regulation of jasmonic acid biosynthesis

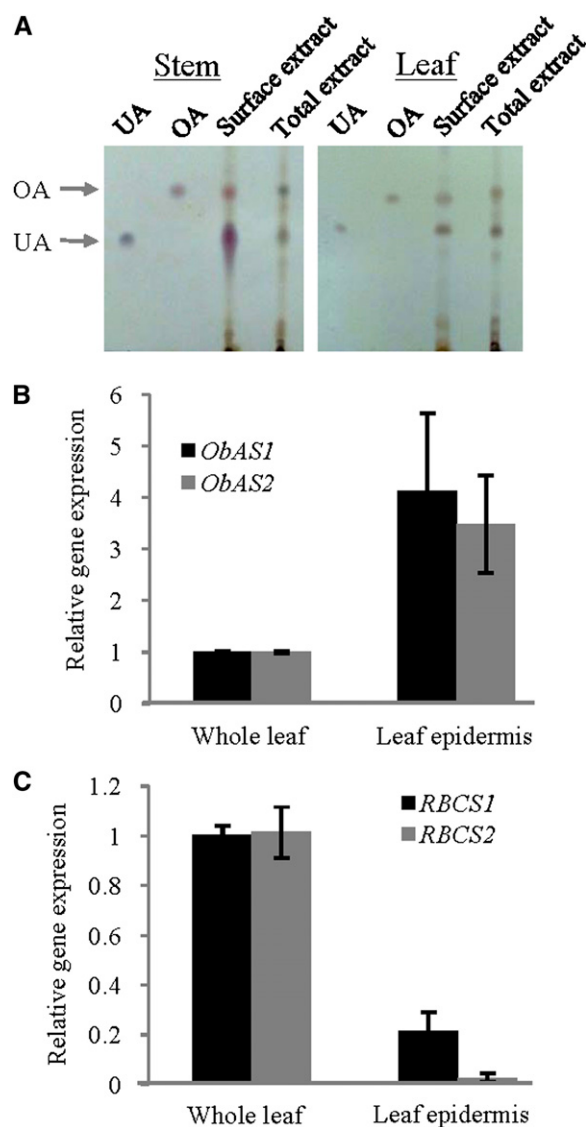


Figure 6. Epidermal cell localization of the pentacyclic triterpene biosynthesis. A, Stem and leaf surface extracts were obtained by dipping stem and leaves, respectively, into chloroform as described in the materials and methods. Total extracts were obtained after complete homogenization of stem and leaf tissues followed by chloroform extraction. Each extract was analyzed by TLC using 200 ng of ursolic acid (UA) and oleanolic acid (OA) as the standards. The analysis was repeated thrice independently with similar results. B, qRT-PCR expression analysis of *ObAS1* and *ObAS2* using whole leaf and epidermal-enriched RNAs. Data are the mean $\pm$ SE from seven biological replicates. C, The expression levels of two unique transcripts of sweet basil Rubisco small subunit (RBCS) were determined using whole leaf and epidermal-enriched RNAs. Data are mean $\pm$ SE ($n = 4$, from two biological replicates). GenBank accession numbers for RBCS1 and RBCS2 are DY342732 and DY336269, respectively. [See online article for color version of this figure.]

(Supplemental Fig. S1). Among these, lipoxygenases are known to catalyze the oxygenation of polyunsaturated fatty acids, such as linolenic acids, to produce fatty acid hydroperoxides that serve as intermediates in the formation of jasmonic acid (Sasaki et al., 2001).

Moreover, additional transcripts of the jasmonic acid biosynthesis pathway, such as C-acyltransferases and oxidases, were also identified (Supplemental Fig. S1; Supplemental Table S5). These results suggest that MeJA controls its own biosynthesis by a positive feedback

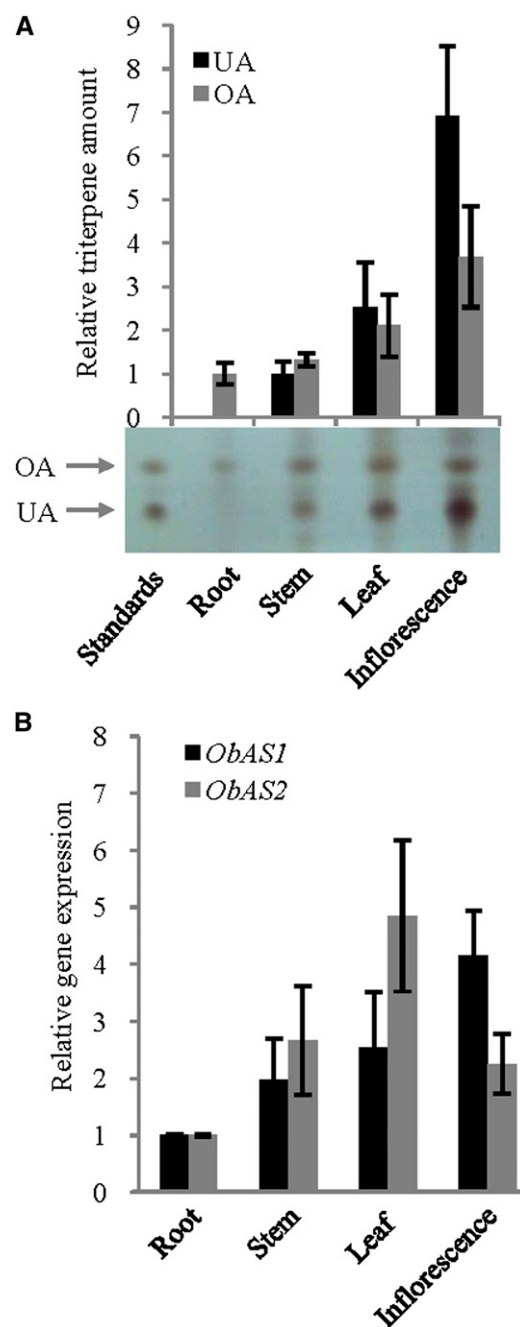


Figure 7. Tissue-specific expression profiles of *ObAS1* and *ObAS2* and differential accumulation of the pentacyclic triterpenes. A, The relative levels of ursolic acid (UA) and oleanolic acid (OA) in different tissues of sweet basil were determined by TLC analysis. Data are the mean $\pm$ SE from six independent experiments and the photograph of one representative TLC plate is presented. B, The relative transcript level was determined by qRT-PCR analysis. Data are the mean $\pm$ SE from four biological replicates. [See online article for color version of this figure.]

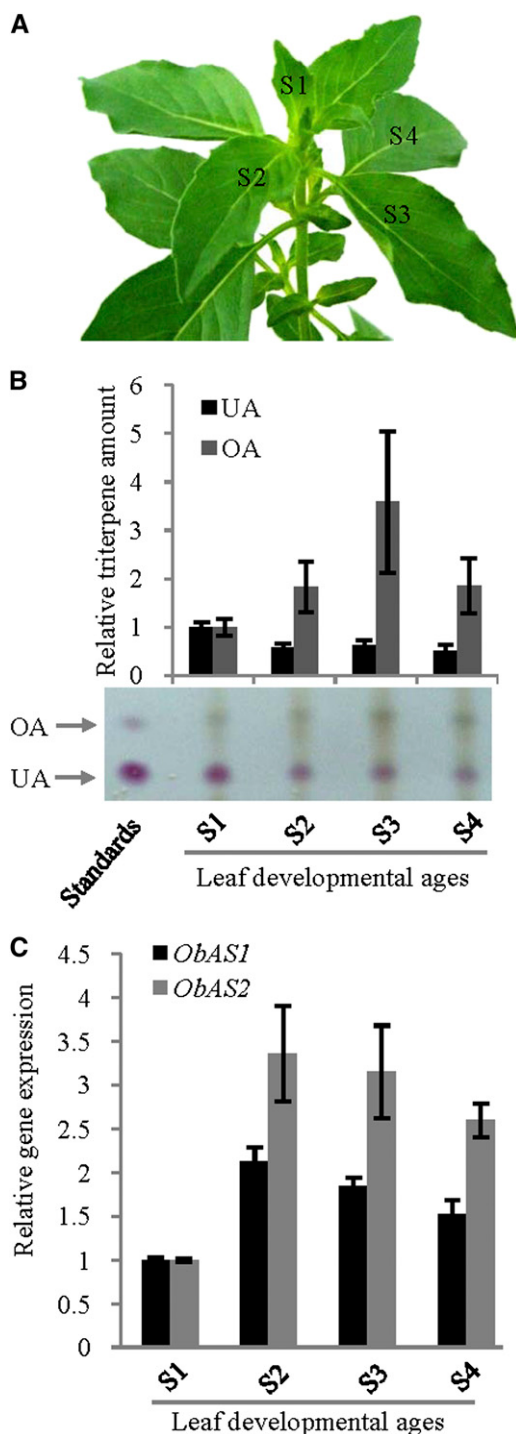


Figure 8. Differential expression of *ObAS1* and *ObAS2* and differential accumulation of the pentacyclic triterpenes in leaves of different developmental ages. A, Four stages (S1 to S4) of leaf pairs used for the gene expression and metabolite analysis. B, The relative levels of ursolic acid (UA) and oleanolic acid (OA) in different leaf ages were determined by TLC analysis. Data are the mean $\pm$ SE from five independent experiments and the photograph of one representative TLC plate is presented. C, The relative transcript levels of *ObAS1* and *ObAS2* were determined by qRT-PCR analysis. Data are the mean $\pm$ SE from four biological replicates. [See online article for color version of this figure.]

mechanism (Sasaki et al., 2001). Few MeJA up-regulated transcripts are the putative transcriptional regulators of MeJA responses in sweet basil (Supplemental Table S3). The APETALA2/ERF and WRKY transcription factors were previously shown as regulators of secondary metabolite biosynthesis in plants in response to MeJA (Yu et al., 2012; Mishra et al., 2013), suggesting their possible involvement in the regulation of secondary metabolite biosynthesis in sweet basil. Lectins, callose synthase, and thaumatin-like proteins that presumably do not have any role in secondary metabolite accumulation were also identified among the MeJA up-regulated transcripts (Supplemental Table S4). These genes function as plant defense components and were also found to be induced by MeJA in other plant species (Cheong and Choi, 2003; Lenka et al., 2012; Ee et al., 2013). The data also suggest MeJA responsiveness of the MEP and MEV pathways that provide 5C building blocks for the biosynthesis of the diverse terpene metabolites (Table II). The MeJA-induced expression of the linalool synthase, eugenol synthase, and polyphenol oxidase substantiates the earlier reports of MeJA-induced biosynthesis of the corresponding metabolites in sweet basil (Table II; Kim et al., 2006; Li et al., 2007).

Our transcript and metabolite analysis revealed MeJA induction of the ursane-type and oleanane-type pentacyclic triterpene biosynthesis pathways of sweet basil (Fig. 2, A to C). OSCs (*ObAS1* and *ObAS2*) and CYP716A subfamily cytochrome P450s (*ObCYP1*, *ObCYP2*, and *ObCYP3*) that showed induced expression after MeJA treatment could be involved in MeJA-induced biosynthesis of ursane-type and oleanane-type pentacyclic triterpene skeletons in sweet basil (Fig. 2C). The OSC-mediated cyclization of 2,3-oxidosqualene is the initial diversifying step for the formation of a large group of structurally diverse triterpene compounds that are widespread throughout the plant kingdom (Abe and Prestwich 1995; Phillips et al., 2006). Hydroxylation by cytochrome P450 enzymes, alkylation by methyltransferases, and glycosylation by glycosyltransferases are among the further enzymatic modifications of the triterpene skeletons. In contrast with animals and fungi that generally have a single OSC (lanosterol synthase) for biosynthesis of the membrane sterol, the higher plants contain several OSCs for biosynthesis not only of the sterols (cycloartenol synthase and lanosterol synthase) but also of diverse triterpenes (Phillips et al., 2006; Sawai and Saito, 2011). The functional diversity of the OSCs and evolution of the mixed-function OSCs that produce more than one specific compound enable over 100 different carbon skeletons of naturally occurring triterpenes in plants (Sawai and Saito, 2011). The product specificities of *ObAS1* and *ObAS2* were determined by heterologous expression in yeast (Fig. 5). In yeast, *ObAS1* formed β -amyrin as its sole product, whereas *ObAS2* catalyzed the formation of both α -amyrin and β -amyrin, with α -amyrin as the major product. Moreover, phylogenetic analysis also indicated that *ObAS1* is closely related to the β -amyrin synthases, whereas *ObAS2* is more similar to the mixed-function

OSCs (Fig. 3). Although OSCs that produce β -amyrin specifically are known from other plant species, no enzyme producing α -amyrin as a sole product has been identified to date. All of the identified mixed-function OSCs that catalyze the formation of α -amyrin also produce β -amyrin and few other minor triterpenes (Morita et al., 2000; Saimaru et al., 2007; Brendolise et al., 2011; Yu et al., 2013). Further studies on ObAS1 and ObAS2 can be directed to elucidate the molecular mechanisms and evolutionary perspectives of the amyirin product specificities for OSCs.

The conversion of 2,3-oxidosqualene into α -amyrin and β -amyrin involves an initial protonation step followed by a polyene addition cascade that leads to the formation of tetracyclic C-20 dammarenyl carbocation to pentacyclic C-20 lupenyl, C-19 oleanyl, C-20 taraxasteryl, and C-13 ursanyl carbocation intermediates. Finally, a series of 1,2-shifts of hydride and/or methyl groups, followed by a deprotonation step, give a stable pentacyclic triterpene skeleton (Fig. 9; Phillips et al., 2006; Abe, 2007). The cyclization mechanism of the α -amyrin synthase and β -amyrin synthase is identical up to the C-19 oleanyl carbocation stage. The hydride shifts and deprotonation of the C-19 oleanyl carbocation result in the formation of β -amyrin. However, the migration of a methyl group from C-20 of oleanyl carbocation gives a C-20 taraxasteryl carbocation intermediate that is further converted to α -amyrin following 1,2-hydride shifts and deprotonation (Morita et al., 2000). The reaction mechanisms of α -amyrin synthase and β -amyrin synthase suggest that the conserved amino acid residues in the enzyme active sites take part in a common reaction

up to the C-19 oleanyl carbocation stage. However, some specific amino acid residues of these enzymes determine which of two alternative paths to follow after the formation of the C-19 oleanyl carbocation intermediate (Fig. 9).

ObAS2 produced more α -amyrin than β -amyrin; therefore, it must have characteristic amino acid residues that favor the formation of the ursane-type over the oleanane-type pentacyclic triterpene by directing the reaction toward C-13 ursanyl carbocation following a C-20 methyl shift, instead of completely stabilizing the C-19 oleanyl carbocation (Fig. 9). At the amino acid sequence level, ObAS1 and ObAS2 shared 60.32% identity and all of the conserved amino acids within the DCTAE, MWCYCR, and QG motifs were identical (Fig. 4). Therefore, the observed difference in their enzymatic functions could be due to the variation in amino acid residues that are outside these conserved motifs. On the basis of the amino acid sequence comparison of apple (*Malus domestica*) MdOSC1 with other OSCs, it was proposed earlier that substitution of Phe to Trp within the MWCYCR motif may possibly be responsible for the increased product specificity to α -amyrin (Brendolise et al., 2011), because site-directed mutagenesis of Trp residue led to loss of the product specificities for some OSCs (Kushiro et al., 2000). However, the presence of Trp in the MWCYCR motifs of both ObAS1 and ObAS2 suggests that α -amyrin product specificity is a result of the changes at other amino acid positions.

Sequence comparison of ObAS1 with ObAS2 highlighted some amino acid residues that are also conserved in two other mixed-function OSCs from *C. roseus* and *O. europaea*, which have α -amyrin product preference

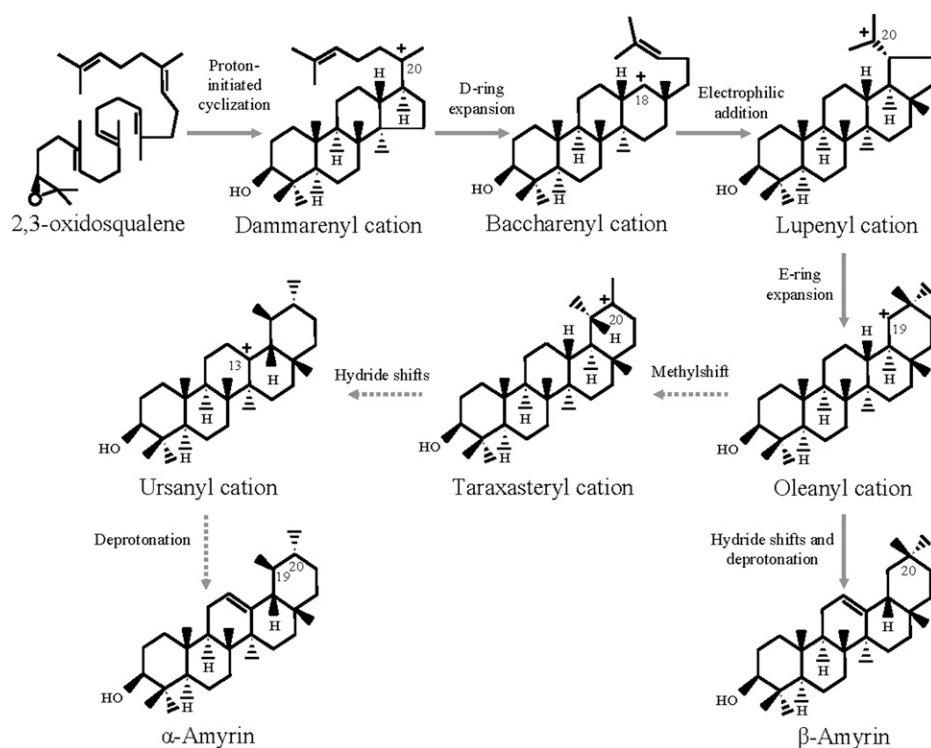


Figure 9. The mechanism of ObAS1 and ObAS2-mediated cyclization of 2,3-oxidosqualene into pentacyclic triterpenes. The protonation of 2,3-oxidosqualene triggers a cyclization cascade that leads to the generation of different carbocation intermediates, which further undergo hydride and methyl shift rearrangement steps before being deprotonated. The differential stability of the C-19 oleanyl and C-13 ursanyl carbocations bound to the enzyme is likely to affect the product specificities of ObAS1 and ObAS2. The reaction steps are common for ObAS1 and ObAS2 up to C-19 oleanyl carbocation intermediate. After that, the reaction path favored by ObAS2 is depicted by dashed arrows.

such as ObAS2 (Saimaru et al., 2007; Huang et al., 2012; Yu et al., 2013). Some of these residues include Tyr-173, Tyr-227, Leu-242, Tyr-255, ThrThr-266, TyrHis-279, Tyr-358, Glu-392, Tyr-439, PheTyr-448, PheThr-470, Val-485, Arg-516, Tyr-526, Phe-535, ProTyr-545, His-565, Trp-607, Tyr-615, LeuGlu-665, and MetTyr-694 (Supplemental Fig. S4). Aromatic amino acid substitutions are particularly interesting because they are thought to play an important role in stabilizing the high energy carbocation intermediates of the terpene cyclase reactions (Christianson, 2006; Phillips et al., 2006). To determine the functional significance of these amino acid substitutions in ObAS2, further studies using domain swapping/site-directed mutagenesis approaches are needed.

The tissue-specific expression profiles of *ObAS1* and *ObAS2* suggest that these enzymes are localized exclusively in the epidermal cells (Fig. 6B). Moreover, high levels of ursolic acid and oleanolic acid were also extracted as cuticular wax components coating the epidermal cells (Fig. 6A). This correlation in transcript and metabolite abundance implies that the biological role of *ObAS1* and *ObAS2* is to synthesize the triterpenes destined for the cuticular wax component of the plant surface. Taking together differential gene expression patterns and metabolite levels, it is very likely that *ObAS1* and *ObAS2* are the major enzymes responsible for the plant organ and leaf age-specific biosynthesis of oleanolic acid (Figs. 7 and 8). By contrast, plant organ and leaf age-specific accumulation of ursolic acid was somewhat different from *ObAS2* expression patterns. Although the data suggest the involvement of *ObAS2* in the biosynthesis of the ursolic acid in stems and leaves, the role of other OSCs in high-level biosynthesis in inflorescences cannot be excluded. Moreover, it is also possible that metabolite transport function determines ursolic acid accumulation in high levels to the actively growing parts and reproductive organs of the plants, as shown for other secondary metabolites (Wink, 2010b). This is quite likely because cuticular triterpenes are thought to play a role in plant defense, which signifies their accumulation in these organs (González-Coloma et al., 2011). Although the biological role of the pentacyclic triterpenes has not been clearly established, the potential insecticidal and antimicrobial activities of the pentacyclic triterpenes suggest a protective role against pests and pathogens (González-Coloma et al., 2011; Kurek et al., 2012).

Pentacyclic triterpenes are synthesized inside the epidermal cells but are accumulated in the cuticle (Fig. 6, A and B; Murata et al., 2008), suggesting the involvement of the transporter for secretion into the cuticle. The transport is possibly facilitated by the proteins, because it is unlikely that these lipophilic metabolites travel from the epidermal cell to the cuticle through diffusion. Although ATP binding cassette transporters that are involved in secretion of antifungal diterpene and cuticular lipid are known in plants, no triterpene transporter has been identified to date (Jasiński et al., 2001; Pighin et al., 2004). Moreover, long-distance transport of the secondary

metabolites from the site of synthesis to the site of storage by means of the xylem and phloem or via the apoplast has been shown for some instances (Wink, 2010a, 2010b). It is also possible that sterol or lipid transporters that have broad substrate spectra are involved in the export of pentacyclic triterpenes (Buschhaus and Jetter, 2012).

In conclusion, this work led to the identification of several MeJA-responsive transcripts in sweet basil and functional characterization of two OSCs that are involved in the biosynthesis of the direct precursors of two medicinally important pentacyclic triterpenes, ursolic acid and oleanolic acid. This study will be useful in expanding our understanding of the molecular basis of the secondary metabolite accumulation in *Ocimum* species and in metabolic engineering of pentacyclic triterpene biosynthesis. Further work on the enzyme structure-function relationship using *ObAS1* and *ObAS2* could reveal the molecular mechanism of α -amyrin and β -amyrin product specificity of the OSCs.

MATERIALS AND METHODS

Plant Materials and MeJA Treatment

Sweet basil (*Ocimum basilicum* 'CIM-Saumya') seeds were obtained from the National Gene Bank for Medicinal and Aromatic Plants at the Central Institute of Medicinal and Aromatic Plants. Seeds were germinated in presterilized soil. Seedlings at the second true leaf stage were transplanted into earthen pots (15 cm height and internal diameter) containing a mixture of soil and farmyard manure in a 2:1 ratio. The plants were grown under natural light in a glass house at 24°C to 26°C and watered daily with application of Hoagland solution once a week. Roots, stems, matured green leaves, and inflorescences were collected from 90-d-old plants (average height, 63.71 $\pm$ 3.77 cm; n = 7), washed with reverse osmosis water, frozen immediately in liquid nitrogen, and stored at -80°C. MeJA (Sigma-Aldrich) treatment was given to 60 d-old plants (average height, 26.66 $\pm$ 3.7 cm, n = 9; number of matured green leaves, 48.66 $\pm$ 6.76, n = 9) with each group consisting of at least three plants. Plants were thoroughly sprayed until run-off with 250 μ M of MeJA solution prepared in MilliQ water containing 0.2% (v/v) dimethyl sulfoxide and 0.05% (v/v) TritonX-100 and covered with polyethylene bags. After 2 h of treatment, the polyethylene bags were removed and the plants were left in open air for 2 h to completely remove the remaining MeJA and then were later transferred to the glass house. The control plants were sprayed with same solution without MeJA, bagged, and kept in isolation from the MeJA-treated plants, however, under the same treatment/growth conditions as mentioned for the MeJA-treated plants. After the specified time of treatment, leaves (stage S3; Fig. 8A) from the plants were harvested, frozen immediately in liquid nitrogen, and stored at -80°C for further metabolite and RNA isolation.

Construction of the Subtraction cDNA Library

The SSH library was generated according to the PCR-Select cDNA Subtraction kit instruction manual (Takara) as previously described (Ghosh et al., 2013). In brief, poly(A⁺) RNA was isolated from the total RNAs using a mRNA purification kit (Qiagen) and 2 μ g of purified poly(A⁺) RNA was reverse transcribed using avian myeloblastosis virus reverse transcriptase. To produce a cDNA library representative of MeJA-induced transcripts, sweet basil plants treated with 250 μ M MeJA were used as the tester samples, whereas mock-treated plants were used as the driver samples. Double-stranded tester and driver cDNAs were prepared from poly(A⁺) RNA of leaf tissues (stage S3; Fig. 8A) harvested from the plants after treatment for 6, 12, and 24 h. Total RNAs were isolated individually from each time point and pooled together in equal ratios to prepare tester and driver RNA samples for poly(A⁺) RNA isolation. Both tester and driver cDNAs were digested with *Rsa*I for producing blunt end cDNAs. Two populations of adapter-ligated tester cDNA samples were prepared and two rounds of hybridization between the driver and adaptor-linked tester were performed to remove common or noninduced transcripts.

Tester-specific cDNAs were then PCR amplified (Advantage 2 PCR kit; Takara). PCR products were ligated into pTZ57R/T vector (Thermo Scientific) and transformed into *Escherichia coli* (DH5 α) for propagation. Recombinant clones were selected based on blue-white selection on a Luria-Bertani agar plate containing 50 $\mu\text{g mL}^{-1}$ ampicillin and grown in Luria-Bertani media at 37°C for plasmid isolation and glycerol stock preparation.

Sequencing and Analysis of ESTs

To obtain the sequences of ESTs, plasmid DNAs were isolated from the individual recombinant clones. Purified plasmid DNAs were analyzed on 0.8% (w/v) agarose gel before sequencing. Plasmid DNAs were sequenced using the BigDye Terminator kit (Applied Biosystems) with the M13 forward primer in an ABI Prism 3700 DNA analyzer (Applied Biosystems). A total of 550 independent recombinant clones were randomly picked; after sequencing, 509 high-quality sequences were obtained that were submitted to GenBank (accession numbers JZ190506 to JZ191014). These sequences were assembled into contigs after cleaning of vector sequences (Masoudi-Nejad et al., 2006). Functional annotation of the unigenes (contigs and singletons) was carried out by a homology search against the nonredundant protein databases of NCBI (<http://www.blast2go.com>) and UniProtKB/The Arabidopsis Information Resource (<http://www.agbase.msstate.edu>) and by following the GO scheme and KEGG pathways (<http://www.blast2go.com>).

Cloning and Sequence Analysis of *ObAS1* and *ObAS2*

Gene-specific oligonucleotide primers were designed based on the sequences of two MeJA-responsive unique ESTs encoding for putative OSCs (ObMU39, ObMU162) and 5' and 3' end sequences were amplified as previously described (Ghosh et al., 2011). In addition, gene-specific degenerate oligonucleotides were also designed based on the conserved motifs of OSCs identified from the multiple sequence alignment. Total RNAs were isolated from the sweet basil leaves and reverse transcribed to generate cDNAs using a poly(A) tail-specific oligonucleotide (3' RACE adapter primer; Invitrogen). The left primer corresponding to either peptide HQNEDGGWG (5'-CAYCAAAAYGARGAYGGIGITGG-3') or NMPPEFW (5'-AAYYCIATGCCCHCCIGARTTYTGG-3') and the right primer corresponding to peptide DQDHGWQV (5'-ACITGCCAICCRTRGRT-CYTGRTC-3') were used to amplify partial sequences of OSCs that were later cloned into pTZ57R/T vector (Thermo Scientific) and sequenced. The remaining 5' and 3' regions were amplified using the RACE kit (Invitrogen/Takara). Related protein sequences from other species were taken and phylogenetic analysis (Tamura et al., 2011) and sequence alignments (ClustalW2; <http://www.ebi.ac.uk>) were performed.

RNA Isolation, sqRT-PCR, and qRT-PCR

RNA was isolated according to the LiCl precipitation method (Meli et al., 2010), treated with DNase I (Thermo Scientific), and purified using the RNeasy Mini Kit (Qiagen). At least two independent isolations consisting of three plants for each group were performed. RNA quality was checked by running 1.2% (w/v) agarose gel and by determining the A260/280 ratio. Four micrograms of total RNA, quantified using a nanodrop (Thermo Scientific), was reverse transcribed using superscript II (Invitrogen) following the manufacturer's instructions. cDNAs were diluted three times before use in the sqRT-PCR and qRT-PCR reactions. sqRT-PCR was performed in a 25- μL reaction volume with 200 nm of each primers using one unit of recombinant Taq DNA polymerase (Invitrogen) for 28 cycles of denaturation, annealing, and extension steps. qRT-PCR reactions (10 μL reaction volume) consisted of 5 μL 2X Power SYBR Green PCR Master Mix (Applied Biosystems), 0.5 μL of diluted cDNA, and 500 nm of each forward and reverse gene-specific primer. qRT-PCR was performed using 7900 HT Fast Real-Time PCR (Applied Biosystems). A list of the oligonucleotide primers used is shown in Supplemental Table S6. Melting curves were analyzed at the dissociation step to examine the specificity of amplification. Relative gene expression was analyzed following the $2^{-\Delta\Delta C_t}$ (cycle threshold) method, using actin (DY329457) as the endogenous control.

Yeast Expression, Protein Purification, and GC-MS Analysis of Yeast Extracts

The *ObAS1* and *ObAS2* were expressed in yeast (*Saccharomyces cerevisiae*) as N-terminal poly(His)-tagged proteins under the control of Gal-inducible

GAL1 promoter. The full-length coding sequences of the *ObAS1* and *ObAS2* were PCR amplified using Phusion high-fidelity DNA polymerase (Finnzymes) with gene-specific forward and reverse primers that contained *NotI* and *XbaI* restriction sites, respectively. The PCR products were digested with *NotI/XbaI* and ligated into the *NotI/XbaI* sites of the yeast expression vector pYES2/NTB to create pYES2-*ObAS1* and pYES2-*ObAS2*. After verifying the integrity of the gene through sequencing, expression plasmids were transformed into *S. cerevisiae* (strain BY4741) following the standard lithium acetate method. Transformants were selected on synthetic dextrose medium without uracil after 3 d of incubation at 30°C and were confirmed by colony PCR.

To induce protein expression, yeast transformants were initially grown at 30°C in synthetic dextrose medium without uracil for 16 h and cells were collected by centrifugation. Cells were then washed twice with MilliQ water, resuspended in synthetic complete medium lacking uracil, containing 2% (w/v) Gal and 1% (w/v) raffinose, and incubated at 30°C for protein induction. After 12 h of growth, cells were collected by centrifugation, crushed with liquid nitrogen using a pestle and mortar, and resuspended in lysis buffer (50 mM Na₂PO₄, 300 mM sodium chloride, 20 mM imidazole, pH 8.0 with 0.1 mM phenylmethylsulfonyl fluoride and protease inhibitor cocktail; Sigma-Aldrich). After dialysis for 12 h in lysis buffer, poly(His)-tagged proteins were purified using nickel-nitrilotriacetic acid agarose beads according to the manufacturer's protocol (Qiagen). The purity of the proteins was determined by resolving on 10% (w/v) SDS/PAGE gel.

For analysis of yeast products, cells were harvested by centrifugation after 48 h of Gal induction and saponified with 20% (w/v) KOH in 50% (v/v) ethanol at 80°C for 45 min with intermittent vortexing. After extraction with hexane twice, the hexane phase was dried and the residue was derivatized using bis-*N,O*-(trimethylsilyl) trifluoroacetamide (Sigma-Aldrich) at 80°C for 45 min prior to GC-MS analysis. The PerkinElmer Clarus 680 GC system equipped with an Elite-5MS column (30 m length, 0.25 mm i.d., 0.25- μm film thickness) and coupled with a PerkinElmer Clarus SQ 8C Mass Spectrometer was used for GC-MS analysis. The sample was injected into a split injection mode (1:25) and the carrier gas was helium with a flow rate of 1.2 mL/min. The injection temperature was 250°C. The GC oven temperature was programmed from 80°C (held for 2 min) to 290°C at 20°C/min and held for 20 min. For metabolite identification, full mass spectra were generated by scanning within the mass-to-charge ratio range of 40 to 600. Triterpene products in yeast were monitored on the basis of the intensity of the base peak (mass-to-charge ratio of 218), with pYES2/NTB empty vector as a negative control and a mixture of authentic standards, α -amyrin and β -amyrin (Sigma-Aldrich), as positive control. Triterpenes were confirmed by comparing both the retention time and mass spectra with the authentic standards.

Extraction and Analysis of Triterpenes

Fresh tissues from different plant parts were harvested separately, frozen in liquid nitrogen, and grinded into fine powder using a pestle and mortar. The ground tissue (500 mg) was extracted twice with 10 mL of chloroform. The obtained extract was evaporated to dryness and the residue was dissolved in methanol to a fixed volume (200 μL). To obtain surface extracts, tissues (500 mg) were dipped in 10 mL chloroform, vortexed for 2 min, and incubated for 5 min at room temperature. The surface extracts were dried and finally dissolved in 200 μL of methanol.

Triterpene analysis was carried out as described previously (Wójcik-Kosior, 2007) with few modifications. Ursolic acid and oleanolic acid stock solutions (0.2 mg mL⁻¹) were prepared in methanol. For analysis of the plant extracts, TLC plates (silica gel 60 F254, 20×20 cm; Merck) were washed with methanol and dried before use. Standards and samples were derivatized with 1% (w/v) iodine solution in chloroform for 30 min at dark (30°C) and spotted onto a TLC plate. Excess iodine from the TLC plate was removed by drying. Further, the plates were developed with a mixture of petroleum ether:ethyl acetate:acetone (8:2:1, v/v/v) as a mobile phase. After drying, the plates were sprayed with 2% (w/v) ceric sulfate solution prepared in 5% (v/v) H₂SO₄, incubated at 37°C for 10 min, and heated to 110°C for 5 min. The visualized spots were documented with a digital camera in daylight. The quantification was carried out by densitometric scanning (Camag TLC Scanner 3) in absorbance/transmittance mode at $\lambda = 530$ nm.

Isolation of Leaf Epidermis-Enriched RNAs

Leaf epidermis-enriched RNAs were isolated as described previously (Levac et al., 2008), with few modifications. Sweet basil leaves (1 g fresh weight) of developmental stage S2 (Fig. 8A) were freshly harvested and mixed

with 7.5 mL of Trizol reagent (Invitrogen) and 1g of carborundum powder (300 grit; Fisher Scientific) in a 50-mL conical bottom centrifuge tube (Tarsons). The tube was vortexed at maximum intensity for 1 min using a cyclomixer (REMI CM 101) to ensure that majority of the leaves in the tube were well abraded. The tube was then kept at room temperature for 5 min and abraded leaves were removed with forceps. The whole process was repeated once with an additional 1 g of fresh leaves. The extract was centrifuged at 3,000g for 5 min and the supernatant was transferred to a fresh centrifuge tube. To the supernatant 0.2 volume of 5 M sodium chloride was added and vortexed for 30 s. Further, a 0.6 volume of chloroform was added, vortexed for 30 s, and centrifuged at 3,000g for 30 min. The aqueous solution was transferred to 1.5-mL centrifuge tubes together with 0.9 volume of isopropanol and 10 μ g of linear polyacrylamide and incubated at -20°C for 12 h. Samples were then centrifuged at 20,800g for 30 min to collect the RNA precipitate. The RNA pellet was washed with 70% (v/v) ethanol, air-dried, and dissolved with diethyl pyrocarbonate-treated MilliQ water. For RNA isolation from whole leaf tissues, 0.5 g of freshly harvested stage S2 leaves were ground into powder using liquid nitrogen and 7.5 mL of Trizol was added. The rest of the isolation steps were the same as the epidermal cell RNA isolation. RNA quality and quantity were determined using a nanodrop (Thermo Scientific) and resolving on 1.2% (w/v) agarose gel. Through carborundum extraction, approximately 2 μ g of total RNAs were obtained from 1 g of fresh leaves.

Sequence data from this article can be found in the GenBank/EMBL data libraries under accession numbers JZ190506–JZ191014 (ESTs), KF636411 (*ObAS1*), JQ809437 (*ObAS2*), FJ032006 (*Malus*), BAF35580 (*Kandelia*), NP 001234604 (*Solanum*), ACO24697 (*Gentiana*), ADK12003 (*Aralia*), O82140 (*Panax*), BAE43642 (*Euphorbia*), A8CDT2 (*Bruguiera*), Q8W3Z1 (*Betula*), ACA13386 (*Artemisia*), ABK76265 (*Vaccaria*), Q9MB42 (*Glycyrrhiza*), AAO33578 (*Medicago*), AAO33580 (*Lotus*), Q9LRH7 (*Pisum*), BAG82628 (*Arabidopsis*), BAF63702 (*Olea*), and AF19235 (*Catharanthus*).

Supplemental Data

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

Supplemental Figure S1. Alpha-linolenic acid metabolism and jasmonic acid biosynthetic pathway obtained from the KEGG database (<http://www.genome.jp/kegg/kegg2.html>).

Supplemental Figure S2. A simplified representation of the phenylpropanoid/flavonoid pathway.

Supplemental Figure S3. TLC separation of the isomeric pentacyclic triterpene compounds ursolic acid (UA) and oleanolic acid (OA) after iodine derivatization.

Supplemental Figure S4. Amino acid sequence comparisons of sweet basil *ObAS1* and *ObAS2* with mixed-function OSCs of *C. roseus* and *O. europaea*.

Supplemental Figure S5. Total ion chromatograms obtained from GC-MS analysis of hexane extracts of yeast cells expressing *ObAS1* and *ObAS2*.

Supplemental Table S1. Functional annotation of the MeJA-responsive unique ESTs of sweet basil.

Supplemental Table S2. List of the enzyme classes identified in the MeJA SSH library.

Supplemental Table S3. List of the sweet basil transcription factors identified in the MeJA SSH library.

Supplemental Table S4. Some of the plant defense-related transcripts identified in the MeJA SSH library.

Supplemental Table S5. KEGG pathway analysis of the MeJA-responsive unique ESTs.

Supplemental Table S6. List of the primers used in this study.

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