Ornacitrus: Development of Genetically Modified Anthocyanin-expressing Citrus with Both Ornamental and Fresh Fruit Potential

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ABSTRACT. Anthocyanins are beneficial bioflavonoids that have numerous roles in human health maintenance, disease prevention, and overall well-being. In addition, anthocyanins are key to the consumer appeal of many ornamental plants. Most citrus (*Citrus* L.) plants do not produce anthocyanins under warm tropical and subtropical conditions. Anthocyanin pigments, responsible for the “blood” color of blood orange [*Citrus sinensis* (L.) Osbeck], are produced after exposure to cold conditions during the fruit's development. The transcription factor *Ruby* is responsible for the production of anthocyanin in blood orange. Functionally, similar genes exist in other fruit crops such as grape [*Vitis vinifera* L. (*VvmybA1* and *VvmybA2*)] and apple [*Malus ×domestica* Borkh (*MdMYB10*)]. Here, *VvmybA1* and *Ruby* genes were constitutively expressed in 'Mexican' lime (*Citrus aurantifolia* Swingle). This cultivar performs optimally under Florida's humid subtropical environment and has a short juvenile phase. Constitutive expression of *VvmybA1* or *Ruby* resulted in anthocyanin pigmentation in the leaves, stems, flowers, and fruit. An increased pigmentation of the outer layer(s) of stem tissue was observed in 'Mexican' lime overexpressing the *VvmybA1*, whereas lower anthocyanin levels were observed in plants overexpressing *Ruby*. Enhanced pigmentation was also observed in the young leaves; however, pigment intensity levels decreased as the leaves matured. Flower color ranged from light pink to fuchsia and the fruit pulp of several 'Mexican' lime lines were maroon; similar to a blood orange. The results demonstrate that expression of anthocyanin-related genes can affect temporal pigmentation patterns in citrus. It also opens up the possibility for the development of modified blood orange and other cultivars adapted to the subtropical environment.

Anthocyanins are plant pigments responsible for the coloration in many fruits, vegetables, and flowers, whereas others are colorized by carotenoids and betalains. These red/violet pigments impart an aesthetic appeal to the cultivars in which they are expressed. Anthocyanin pigments are present in many plant tissues. However, they are more visible in flowers and fruit. They have been consumed by humans and animals since prehistoric times without any side effects (Brouillard, 1982). Anthocyanins belong to the flavonoid group of phytochemicals that provide several therapeutic properties potentially beneficial to human health if consumed in adequate quantities (Bridle and Timberlake, 1997; Saija, 1994). Anthocyanins contribute to free radical scavenging (De Beer et al., 2003; Sánchez-Moreno, 2002) and have potent in vitro antioxidant capacities (Deighton et al., 2000; Lila, 2004; Wang et al., 1997). Numerous pharmacological studies have implicated their intake on the prevention of a number of human health issues such as obesity and diabetes (Tsuda et al., 1996, 2003). In addition to their health benefits, anthocyanins are bacteriostatic (Naz et al., 2007) and have been widely used as a natural source of food colorants (Manach et al., 2004).

Anthocyanins are synthesized by the flavonoid biosynthesis pathway (Holton and Cornish, 1995). These water-soluble pigments are produced through a complex expression of several genes such as MYB transcription factors, basic helix-loop-helix (bHLH) transcription factors, and WD-repeat proteins (Baudry et al., 2004; Irani et al., 2003). Several MYB regulators of anthocyanin production have been cloned in recent years.

When overexpressed in heterologous systems, these transcription factors resulted in an enhanced overexpression and accumulation of anthocyanins (Borevitz et al., 2000; Gonzali et al., 2009; Meng et al., 2014). It has been observed that there is a requirement for the presence of a suitable bHLH partner to achieve full functionality in the heterologous system (Espley et al., 2007; Li et al., 2007; Takos et al., 2006).

The blood orange is rich in anthocyanins (Hillebrand et al., 2004) and arose as a mutation of the common blond orange during the 16th century. Blood orange contains a copia-like retrotransposon inserted in the promoter region of a MYB transcription factor named *Ruby*. This gene is responsible for transcriptional activation of anthocyanin production in blood orange (Butelli et al., 2012). Blood orange is mainly cultivated in Sicily and southern Italy (Bridle and Timberlake, 1997). When grown in the subtropical sweet orange belt, these oranges do not accumulate adequate anthocyanins and, thus, cannot develop the characteristic deep red coloration that is observed in fruit produced in cooler climates (Lee, 2002).

Development of genetically modified citrus with the ability to overexpress anthocyanins can have two potential benefits. The first benefit is that it would allow the cultivation of these healthy fruit in the major subtropical citrus belt and/or the production of ornamental citrus plants, depending on the cultivar. The second benefit is that novel fruit, leaf, and flower colors could be produced by regulating anthocyanin biosynthesis (Xiaonan and Qixiang, 2002). Here, we demonstrate the stable expression of two MYB transcription factors and subsequent anthocyanin production in 'Mexican' lime (*VvmybA1* cloned from the red grape 'Ruby Seedless' and *Ruby* cloned from the 'Moro' blood orange) and results are presented on their expression.

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Materials and Methods

VECTOR CONSTRUCTION. Fruits of the red grape ‘Ruby Seedless’ were purchased from a local supermarket, whereas fruits from the blood orange ‘Moro’ were obtained from R. Krueger (USDA-ARS, Riverside, CA). RNA was isolated from both ‘Ruby Seedless’ skin and ‘Moro’ blood orange pulp using an RNeasy plant mini kit (Qiagen, Valencia, CA). Reverse transcription polymerase chain reaction (RT-PCR) amplicons were generated using a OneStep RT-PCR kit (Qiagen), 500 ng of the extracted total RNA, and gene-specific primers designed from *VvmybA1* (AB097923) and *Ruby* (JN402334) sequences [obtained from the National Center for Biotechnology Information database (Table 1)]. These primers were designed to introduce a *Bam*HI site immediately upstream of the translation start site and a *Not*I site immediately following the stop codon. Amplicons were visualized under ultraviolet light on a 1% agarose gel containing GelRed™ (Biotium, Hayward, CA). Amplicons were isolated, purified, and cloned into pGEM®-T plasmid (Promega, Madison, WI). The sequences were verified by Sanger sequencing. Fragments were ligated into a unique *Bam*HI/*Not*I cloning site that lay between a double enhanced CaMV 35S promoter (D35S) and a CaMV 35S terminator (35S-3′). The promoter-gene cassette was inserted into a plant transformation vector containing the *nptII* selectable marker gene. The resulting binary plasmid was introduced into the *Agrobacterium tumefaciens* (Smith and Towns) Conn strain EHA105 (Hood et al., 1993).

PLANT MATERIALS. Epicotyl segments, originating from juvenile nucellar seedlings of ‘Mexican’ lime, were excised and cut obliquely into 2.5-cm-long segments to expose the cambial ring. The *A. tumefaciens* cells were grown overnight and epicotyl segments were transformed as described by Dutt and Grosser (2009). Plants were selected based on the level of anthocyanin production in the regenerating shoots (either on the leaves or the stem). Elongated transgenic shoots were micrografted in vitro onto Carrizo citrange [*Citrus sinensis* × *Poncirus trifoliata* (L.) Raf.] as outlined by Dutt and Grosser (2010) to expedite the recovery of transgenic plants. The

grafted shoots were potted into a peat-based commercial potting medium and acclimated to greenhouse conditions.

HORTICULTURAL MANIPULATIONS TO INDUCE RAPID FLOWERING. An ex vitro micrografting technique (Dutt and Grosser, 2010) was used to clonally propagate the transgenic anthocyanin over-expressing shoots. Tender shoots obtained from greenhouse acclimated mother plants were micrografted onto actively growing 6-month-old Carrizo citrange rootstocks and maintained in a greenhouse that was shady (60% to 80%) and humid (80% to 90%). Plants with active growth shoots were potted in 18.9-L pots and maintained in a greenhouse with 40% shade. Trees were sprayed with micronutrients and fertilized with a 20N–4.4P–16.6K fertilizer (Peat Lite; JR Peters, Allentown, PA) as per manufacturer’s recommendations. Trees were allowed to grow under the above conditions for 2 years before girdling. Girdling was performed on the rootstock ≈15 cm above the potting medium surface in the fall. A sharp cut (≈1 mm wide) was made to completely encircle the stem, and the bark was not removed during girdling. The cut was made deep enough to disrupt the phloem tissues only.

MOLECULAR ANALYSIS OF TRANSFORMANTS. DNA was isolated from 100 mg of leaf tissue using a GenElute™ Plant Genomic DNA Miniprep Kit (Sigma-Aldrich, St. Louis, MO). PCR was carried out to confirm presence of the transcription factor transgene using a GoTaq® Green Master PCR Mix (Promega), 25 nm of the isolated DNA, and the gene-specific primers described above. PCRs were run on a thermal cycler (C1000 Touch; Bio-Rad Laboratories, Hercules, CA). To quantify the production of *VvmybA1* or *Ruby* in the different plant parts, RNA was isolated from 100 mg of leaf tissue as described earlier and reverse transcription real-time PCR (RT-qPCR) primers and probes were designed to detect *VvmybA1* or *Ruby* amplicons using PrimerQuestSM online software (Integrated DNA Technologies, Coralville, IA). The probes were labeled with a 6-carboxy-fluorescein reporter dye at the 5′-end and with Black Hole Quencher (BHQ)-1 at the 3′-end. The housekeeping gene *18S rRNA* was used as an endogenous control (Table 2). RT-qPCR reactions were performed with a final 20-μL reaction volume using a TaqMan® RNA-to-Ct™ OneStep kit (Applied

Table 1. Forward and reverse primer sequences used in cloning of the two MYB genes. RNA was used as a template and a one-step reverse transcription polymerase chain reaction was used to amplify the selected cDNA.

Gene	Forward primer	Reverse primer
<i>VvmybA1</i>	GGATCCATGGAGAGCTTA GGAGTTA	GCGGCCGCTCAGATCAAGTGATTTACT
<i>Ruby</i>	GGATCCATGGCGGATTCCTTAGGAGT	GCGGCCGCTTACTTTGCATTGAGAAGA

Table 2. Primers and probe used in the one-step real-time, reverse transcription polymerase chain reaction assay for the detection of the transcription factor transcripts.

Target gene	Amplicon length (bp)	Primer/probe sequence 5′ → 3′ ^{1/2}
<i>VvmybA1</i>	105	TGGCATAGTCACCACTTCAAA FAM-5′-AGGTTTCAGTTCCAGGAAGAAGGGAGA-3′ CTTGTGAGGGTGAGGCTTTAT
<i>Ruby</i>	104	CCCGAATATCAAACGAGGAGAA FAM-5′-TCCATAAGCTGCTGGGCAACAGAT-3′ GAAGCCTGCCCAACATCA
<i>18S rRNA</i>	112	TCGGGTGTTTTACGTCTCA HEX-5′-TGGAACCTCTGGATTTTGCC-3′ CGCCGTAGGTGAGGTAGC

^{1/2}Primer/probe sequences listed in the column include the forward primer in the first line, followed by the probe sequence in the second, and the reverse primer sequence in the third line.

Fig. 1. (A) An anthocyanin-expressing bud emerging in a selection medium after *Agrobacterium tumefaciens*-mediated transformation of 'Mexican' lime explants. (B) An anthocyanin overexpressing shoot is ready for micrografting to Carrizo citrange rootstock. (C) An anthocyanin overexpressing transgenic line is exhibiting intense pigmentation, curled leaves, and a lack of vigor. (D and E) Normal anthocyanin overexpressing in a transgenic line vigorously growing in the greenhouse.

Biosystems, Foster City, CA) and reactions were run on a thermocycler (ABI PRISM 7700, Applied Biosystems) according to manufacturer's instructions. The one-step kit parameters consisted of 20 min incubation at 48 °C followed by 10 min incubation at 95 °C and 40 cycles at 95 °C for 15 s and 60 °C for 1 min. RNA from a nontransgenic greenhouse 'Mexican' lime plant was used as a negative control, and the data were normalized using the house-keeping gene as described above. Experiments were conducted in triplicate and were repeated at least twice. Relative quantitation was measured and analyzed by the comparative Cq method ($2^{-\Delta\Delta C_t}$). The fold change in the relative expression was then determined by calculating $2^{-\Delta\Delta C_t}$ as described by Livak and Schmittgen (2001).

LIGHT MICROSCOPY. Stems were cut into pieces (1-cm pieces for longitudinal sections and 2.5-mm pieces for cross-sections) and processed for microscopy. Stem pieces were arranged and mounted in Tissue Tek Cryo-OCT Compound (Thermo-Fisher Scientific, Waltham, MA) on a brass stub. The stubs were frozen and 40- μ m sections of tissue were cut using a WRC cryostat microtome (Harris Manufacturing, North Billerica, MA). The sections were mounted in Optic Mount I (Mercedes Medical, Sarasota, FL) on glass slides and a cover slip was placed over the top of the sections. The sections were observed using a fluorescence microscope (BX61; Olympus America, Melville, NY). Light micrographs were captured using a digital camera (14MP; OMAX, Irvine, CA).

Final plates were constructed in Photoshop CC (Adobe Systems, San Jose, CA).

Results and Discussion

PRODUCTION OF TRANSGENIC PLANTS. A population of transgenic lines expressing either the *VvmybA1* or the *Ruby* transcription factor under the control of the d35S promoter was obtained. These transcriptional activator (MYB) genes cloned from blood orange and grapevine appear to enhance the expression of all genes in the anthocyanin biosynthesis pathway (Azuma et al., 2007; Butelli et al., 2012; Jeong et al., 2006). 'Mexican' lime is considered to be recalcitrant to *A. tumefaciens*-mediated transformation (Dutt et al., 2009); however, it was relatively easy to select the few transgenic shoots from the majority of nontransformed ones based on color in

Fig. 2. (A) Flower clusters on a *VvmybA1* overexpressing 'Mexican' lime transgenic line. (B and C) Close-up of flowers on two independent *VvmybA1* overexpressing lines. (D) Close-up of flowers on a *Ruby* overexpressing line. (E) A fruit cluster on a *VvmybA1* overexpressing 'Mexican' lime line. (F) A cross-section of a fruit from a *VvmybA1* overexpressing 'Mexican' lime line demonstrating the production of anthocyanin in the pulp.

Fig. 3. Cross-sections of a transgenic *Ruby* overexpressing 'Mexican' lime fruit with a control nontransgenic fruit for comparison.

Fig. 4. Cross-sections of a transgenic *VvmybA1* overexpressing 'Mexican' lime leaf (A) and a nontransformed control (B). The arrows indicate anthocyanin overexpression in leaf epidermal cells.

Fig. 5. (A) Cross-section of a transgenic *VvmybA1* overexpressing 'Mexican' lime stems demonstrating anthocyanin expression localized to the epidermis, phloem, and xylem of transgenic plants compared to the control (B). Longitudinal sections of stem confirm that only transgenic stems (C) exhibit anthocyanin expression in the epidermis, phloem, and xylem (not shown). Anthocyanins were absent in longitudinal sections of the nontransformed control (D).

either leaves or stem or both (Fig. 1A and B). It is possible that some lines that could potentially express the transgene(s) in other plant organs or later on developmentally were not selected by this process.

VISUAL CHARACTERIZATION OF THE TRANSGENIC PLANTS. Transgenic plants were observed to be phenotypically normal and grew well after being micrografted onto standard Carrizo

rootstocks. Plants could be greenhouse acclimated in 6–8 weeks post-transformation. Fourteen transgenic lines expressing the *VvmybA1* transcription factor and six lines expressing the *Ruby* transcription factor were successfully acclimated. *VvmybA1* overexpression potentially had a detrimental effect on two of our regenerated transgenic lines. These plants exhibited intense pigmentation, curled leaves, and a lack of vigor (Fig. 1C). These lines remained stunted and did not grow beyond 15 cm, even after 2 years in the greenhouse. Accordingly, they were excluded from this study. Similar results have been observed in previous studies following overexpression of *Lc* (Goldsbrough et al., 1996) and *VvmybA1* (Stover et al., 2013) transcription factors. Anthocyanin production is metabolically expensive. It is probable that there is an anthocyanin production threshold where the overexpression of the MYB genes results in toxic levels of anthocyanin that inhibit plant growth and development (Bradley et al., 1998). It is also possible that certain T-DNA insertions could have resulted in the disruption of specific gene(s) in the citrus genome that are needed for normal growth and development (Vlachonasios et al., 2003; Zheng et al., 2005).

Results presented here are for four randomly selected, phenotypically normal, transgenic lines generated with each transcription factor construct. Regenerating *in vitro* shoots overexpressing the *VvmybA1* transcription factor were deep purple in color. Visually, the anthocyanin levels remained steady in the micrografted plants and the pigmentation levels decreased as the leaves matured. In contrast, shoots overexpressing the *Ruby* transcription factor had little or no initial *in vitro* coloration. Anthocyanin levels increased in the micrografted plants with expression confined to the younger leaves. Anthocyanin levels

subsequently decreased as the leaves matured. Stems were also highly pigmented in *VvmybA1* transcription factor overexpressing juvenile plants, and decreased on maturity. This expression pattern has also been observed in other plants overexpressing anthocyanin (Bradley et al., 1998; Qiu et al., 2014). Anthocyanins are produced by the flavonoid biosynthetic pathway via the phenylpropanoid pathway and a number of genes are

Fig. 6. The epidermis of a transgenic *VvmybA1* overexpressing 'Mexican' lime flower petal (A and B) and control petal (C). Anthocyanin production was localized in the vacuole (see arrows).

Fig. 7. Polymerase chain reaction amplicons of the *VvmybA1* and *Ruby* genes from genomic DNA of transgenic 'Mexican' lime plants. Amplification was carried out using gene-specific primers which yielded 0.7-kb *VvmybA1* fragments (lanes 1–4) and 0.8-kb *Ruby* fragments (lanes 5–8) that matched the expected band size determined by a 1-kb DNA ladder (lane M).

involved (Jaakola, 2013). It could be that one or more of these genes are developmentally regulated in maturing citrus. In *Hedera helix* L., dihydroflavonol reductase activity ceased as the plants progressed from the juvenile to the mature phase, resulting in a lack of anthocyanin accumulation in the mature phase (Murray and Hackett, 1991). Similarly, downregulation of an anthocyanidin glucosyltransferase resulted in decreased accumulation of anthocyanins during fruit ripening in strawberry [*Fragaria xananassa* Duch. (Griesser et al., 2008)].

Anthocyanin levels varied among the individual lines and were observed to be temperature dependent. In grape, high temperature decreased anthocyanin accumulation in the berries. This was attributed to a combinatory effect of anthocyanin degradation and inhibition of mRNA transcription of the anthocyanin biosynthetic genes (Mori et al., 2007). MYB transcript levels can also be differentially regulated by cold stress (Agarwal et al., 2006; Camm et al., 1993) and light (Talos et al., 2006). Similar to that observed in other plant species (Lin-Wang et al., 2011; Reyes et al., 2004;

Stiles et al., 2007), anthocyanin levels in both sets of plants were enhanced during winter months and diminished during the summer (results not shown). 'Mexican' lime does not naturally produce anthocyanins, and control plants remained green for the duration of this study. Transgene copy number, insertion location, and position effects can result in differential gene expression (Peach and Velten, 1991). Our transgenic lines produced anthocyanin in tissue-specific areas (Figs. 1

and 2). Rapid flowering following the girdling of 'Mexican' lime trees has been demonstrated earlier by Dutt et al. (2011), and we observed similar results with the transgenic lines in this study. Girdling disrupts movement of photosynthates and metabolites through the phloem (Goren et al., 2004) and increases the carbohydrate levels in leaves (Li et al., 2003). Flowers from different lines had variable anthocyanin production. The fruit pulp was dark purple in the *VvmybA1* and fushia in the *Ruby* overexpressing lines (Figs. 2F and 3). Even though our constructs had MYB genes driven by a constitutive 35S promoter, anthocyanin expression was not detected in the fruit peel or in the mesocarp in any of the *VvmybA1* and *Ruby* overexpressing transgenic lines. This corroborates previous data where tissue-specific expression of transcription factors governed the distribution of anthocyanins in specific organs (Steyn et al., 2002). This also suggests that there are certain regulatory elements, likely including bHLH *R* genes, requiring coactivation before transcriptional activation occurs. These regulatory elements could be missing in the 'Mexican' lime fruit peel, preventing anthocyanin production.

HISTOLOGICAL ANALYSIS. Anthocyanin expression in the leaf, stems, and petals of transgenic plants was examined by histological analyses. Cross-sections of leaves showed that anthocyanin production was localized to the epidermis in transgenic plants [Fig. 4A (see arrows)], but there was no production of anthocyanins in control leaves (Fig. 4B). Likewise, cross-sections of transgenic plant stems showed that the pigment was expressed in the epidermis, phloem, and xylem (Fig. 5A). These results were also confirmed in longitudinal sections (Fig. 5B). There was no pigmentation observed in control sections (Fig. 5C and D). Anthocyanin expression was localized to the vacuole in the flower petals [Fig. 6A and B (see arrows)]; however, anthocyanins expression was absent in control petals (Fig. 6C). This agrees with previous data where anthocyanins have been observed to be sequestered in the vacuole in *Arabidopsis thaliana* (L.) Heynh. and many other plants (Markham et al., 2000; Pourcel et al., 2010; Zhang et al., 2006). Anthocyanins are generally transported to the vacuole by the secretory pathway in cells resulting in a high accumulation of anthocyanin in the vacuole (Poustka et al., 2007).

MOLECULAR CHARACTERIZATION OF THE TRANSGENIC PLANTS. Screening of putative regenerated plants by PCR analysis confirmed the presence of the *VvmybA1* and the *Ruby* transcription factor transgene in our transgenic lines (Fig. 7). In addition, gene expression of these transcription factors was evaluated by RT-qPCR. Transcription factor expression was always higher in the younger leaves compared with mature leaves (Fig. 8). In the present study, a maximum of an 8-fold

Fig. 8. Quantification of *VvmybA1* and *Ruby* activity using reverse transcription real-time polymerase chain reaction. Four selected transgenic lines from each group were evaluated. Three independent clones were tested from each transgenic line. Total RNA from a nontransgenic control plant was also included to verify the accuracy of the amplification process. Primers used to amplify the *VvmybA1* and *Ruby* gene(s) are described in Table 2.

increase in expression of *VvmybA1* was observed, whereas only a 2-fold increase in expression of *Ruby* was observed in transgenic ‘Mexican’ lime lines (Fig. 8). VVMyb-4 had the highest relative expression among the transgenic lines evaluated. This line exhibited an 8-fold increase in *VvmybA1* gene expression compared with the control in young leaves. Gene expression in the other *VvmybA1* overexpressing transgenic lines ranged from a 2-fold to a 6½-fold increase in juvenile leaves compared with the control. In several plant species, juvenile reddening decreases as the leaves mature (Chalker-Scott, 1999) and this plays a role in the tender leaf’s defense from insect herbivory and excessive light (Karageorgou and Manetas, 2006; Steyn et al., 2002).

In certain purple-leaved ornamentals, color is maintained in the mature tissues. This phenomenon helps the plant to reduce the frequency and severity of photoinhibition (Gould et al., 2002; Neill and Gould, 2003) as well as protection from ultraviolet radiation or free radical scavenging (Burger and Edwards, 1996; Wang et al., 1997). This was not observed in our transgenic lines, where the anthocyanin levels dramatically decreased in mature leaves (Figs. 1E, 2A, and 8). We observed a 4-fold decrease in VVMyb-4 expression in mature leaves. The VVMyb-1 and VVMyb-2 lines also showed a 3-fold decrease in expression in the mature leaves, whereas expression of VVMyb-3 was reduced by half.

Anthocyanin production in *Ruby* overexpressing lines was much lower than that observed in the *VvmybA1* overexpressing transgenic lines. The *Ruby* lines had a 2-fold increase in the expression levels in young leaves. The transgenic line Ruby-4 had a slight reddish hue in the mature leaves, whereas leaves from the other lines were visibly indistinguishable from control leaves. In red grape berries, 29 anthocyanins have been identified of which malvidin derivatives were the most abundant anthocyanins (Liang et al., 2008). However, in blood oranges, the major anthocyanins were characterized as cyanidin 3-glucoside and cyanidin 3-(6"-malonylglucoside) (Hillebrand et al., 2004). This could indicate that the *Ruby* transcription

factor may act in a different manner on the structural genes of the anthocyanin biosynthetic pathway in citrus when compared with the activity of the *VvmybA1* transcription factor. Validation of these observations with RNA sequencing studies or by analyzing the anthocyanin composition differences between the different lines is required for a conclusive interpretation.

Conclusions

Our report outlines the successful production of transgenic ‘Mexican’ lime plants overexpressing the *VvmybA1* or the *Ruby* MYB transcription factor, resulting in the production of anthocyanins in a citrus cultivar that does not produce anthocyanin naturally. These transgenic plants exhibited unique leaf pigmentation, flower coloration, and pulp enhanced by anthocyanin overproduction. These transgenic lines can be directly used as novel ornamentals or released for fresh fruit consumption, as anthocyanins have been shown to be beneficial to human health. In addition, our research opens up the possibility for the development of modified sweet orange or grapefruit cultivars that are anthocyanin rich and adapted to the subtropical environment. Additional information on these plants can be searched at <http://www.crec.ifas.ufl.edu/>.

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