

POTENTIAL OF *VlmybA1-2* AS A CANDIDATE MARKER FOR VISUAL IDENTIFICATION OF TRANSGENIC PLANTS: INDUCED ANTHOCYANIN PRODUCTION IN *ARABIDOPSIS* AND TOBACCO

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ABSTRACT

For development of a safe and efficient *in vivo* marker for plant transformation, an *myb*-related gene of anthocyanin biosynthetic pathway, *VlmybA1-2* from grape, was introduced into anthocyanin producing tobacco and *Arabidopsis*, and non- anthocyanin producing spinach under the control of CaMV 35S promoter. Except for the distinguishable purple color, transformed calli and plants of tobacco were not different from controls. RNA gel blot hybridization confirmed the expression of *VlmybA1-2* in purple tobacco seedlings. Completely purple T1 *Arabidopsis* seedlings could not survive as high anthocyanin levels may affect normal growth; Surviving T1 seedlings could produce viable seeds of two distinguishable colors: purple and brown (similar to wild type). Purple seeds germinated on kanamycin medium providing an easy method of transgenic seed identification in *Arabidopsis*. T2 and T3 completely purple seedlings produced purple flowers and seeds. Putative transgenic spinach was not different to control in color, although presence of *VlmybA1-2* was confirmed by DNA gel blot hybridization. *VlmybA* alone, without the aid of an *myc*-related gene partner, could induce complete pigmentation in tobacco and *Arabidopsis* indicating its potential over previously used *myb*- and *myc*-related genes.

Key words: Anthocyanin, *Arabidopsis*, *In vivo* marker, Tobacco, *VlmybA1-2*

INTRODUCTION

Anthocyanins are unique pigments in flowering plants. Structural genes encoding the enzymes of the anthocyanin biosynthetic pathway are conserved among different species (Holton and Cornish 1995). Two classes of regulatory genes, *myb* and *myc* of *C* and *R* families activate the transcription of structural genes in maize. Differences in the regulatory mechanisms controlling the expression of structural genes, and interactions among them would account for the intensity and variety of pigmentation patterns among different plant species. Therefore, the manipulation of regulatory genes in transgenic plants may be versatile over that of structural genes.

Several regulatory genes from *C* and *R* families have been isolated from several plants and one of such gene alone could not induce distinguishable phenotypes in transgenic plants (Lloyd *et al.* 1992). Three types of *myb*-related genes capable of anthocyanin induction in non-colored grape have been isolated from Kyoho grape: *VlmybA1-1*, *VlmybA1-2* and *VlmybA2* (Kobayashi *et al.* 2002). *VlmybA* genes have a unique protein sequence compared with other known plant *myb*-related genes of the anthocyanin biosynthetic pathway (Kobayashi *et al.* 2002). *VlmybA2* could induce anthocyanin pigmentation in *Arabidopsis* and tobacco in a previous attempt (Geekiyanage *et al.* 2007).

Development of alternatives for commonly used marker genes, conferring resistance to antibiotics and herbicides, is useful for safer genetically modified crops to maximize their benefits. If such genes would not affect the normal plant growth and development of the host plant, enhanced anthocyanin production induced by them could be exploited for a simple, non-destructive, efficient identification of transformation at early stages and to investigate the potential of out-crossing of GM crops in the field.

Enhanced anthocyanin production, induced by regulatory genes from maize have been reported in maize, *Arabidopsis*, tobacco and tomato, and their potential as visual markers has been highlighted in those cases (Ludwig *et al.* 1990; Lloyd *et al.* 1992; Goldsbrough *et al.* 1996; Tamaoki *et al.* 2006).

Anthocyanin pigments are useful in human health in a variety of fields and in food industry as a natural food colorant as well (Lila 2004). Attempts to develop callus and plant cell systems from a variety of crops have been made to produce stable anthocyanin pigments *in vitro* to be used for food industry and bio-medicinal purposes. Genetically engineered callus or cell lines must produce high pigment yield and their dependence on irradiance can be low.

In this study, I examined the potential of a grape *myb*-related gene, *VlmybA1-2*, to induce anthocyanin biosynthesis in anthocyanin producing plants, tobacco and *Arabidopsis*, which are previ-

ously tested for some *myb*-related genes and in non-anthocyanin producing spinach, in order to check the functional equivalence of an anthocyanin regulatory gene of a dicot, grape, in them. This was an attempt for the future work of development of a safe *in vivo* marker for plant transformation studies and development of plant cell systems that produce stable anthocyanins for applications as natural food colorants in bio-industry.

METHODOLOGY

Plant materials, growth conditions and culture media

Arabidopsis thaliana ecotype Columbia, *Nicotiana tabacum* cultivar Maryland Mammoth and *Spinacia oleracea* L. cultivar Longstanding Bloomsdale Dark Green were used for the experiment. Growth conditions and culture media for *Arabidopsis* and tobacco were as previously described (Geekiyana *et al.* 2007). Spinach seedlings were grown on MS medium (Murashige and Skoog 1962) under the short-day (SD) condition (8h light and 16h darkness) for cotyledon explants.

Agrobacterium-mediated transformation

Agrobacterium tumefaciens strain LBA4404 harboring the pBI121 vector carrying full-length cDNA of *VlmybA1-2* coding region, cloned by *Xba*I and *Sac*I sites at 5' and 3' ends respectively (Kobayashi *et al.* 2002) in place of the *GUS* gene under the control of the CaMV 35S promoter, and *NPTII* gene under NOS promoter were used for the transformation.

Arabidopsis inflorescences were transformed by the floral dip method (Clough and Bent 1998). The method of Horsch *et al.* (1985) was used for tobacco leaf disc transformation. T1 seeds were selected on kanamycin-containing medium.

Spinach cotyledons were transformed according to the method of Zhang and Zeevaart (1999).

RNA gel blot hybridization

RNA gel blot hybridization was performed as previously described (Geekiyana *et al.* 2007).

Identification and measurement of anthocyanin

Identification of anthocyanin was performed as previously described (Geekiyana *et al.* 2007).

RESULTS

Different phenotypes of 35S: *VlmybA1-2* T1 *Arabidopsis* seedlings

There were five T1 seedlings that showed three phenotypes (Fig 1): two seedlings had purple roots

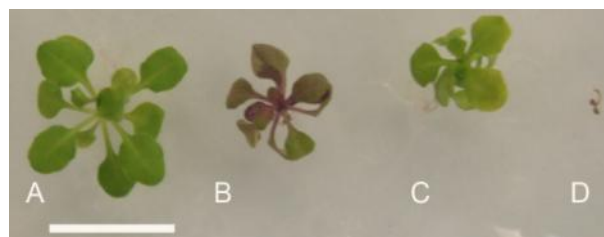


Figure 1 Different phenotypes of T1 *Arabidopsis* plants of 35S:*VlmybA1-2* after 2 weeks of germination. (A) A seedling with purple-green leaves and purple roots (B) A completely purple seedling (C) A seedling with green leaves and purple roots (D) A completely purple seedling which could not grow beyond cotyledon stage. Bar= 2 cm

and, purple-green leaves showing normal growth and flowered and set seeds of two distinguishable colors; purple and brown similar to wild type; three seedlings were completely purple, out of two could not grow beyond the cotyledon stage, while the other was transferred to a pot and died before flowering.

Segregation of *VlmybA1-2* in T2 seeds of *Arabidopsis*

Seeds of two independent T1 lines showed two phenotypes: purple and brown seeds, similar to wild type (Fig 2). The ratio of purple to brown seeds was 3:1 suggesting that homozygous and heterozygous seeds for the transgene are purple in color (data not shown). It was confirmed by the germination test for purple and brown seeds on kanamycin containing medium and kanamycin-free medium (Fig 3).

The purple pigment from *Arabidopsis* could be extracted with water and became dark blue with the addition of 0.1M sodium hydroxide. These results suggest that the pigment is anthocyanin (Rabino and Mancinelli 1986; Uimari and Strommer 1998; Kobayashi *et al.* 2005).

VlmybA1-2 induced anthocyanin production in T2 and T3 *Arabidopsis* plants

Purple T2 seeds of completely purple T1 plant germinated on kanamycin-containing medium. Germination

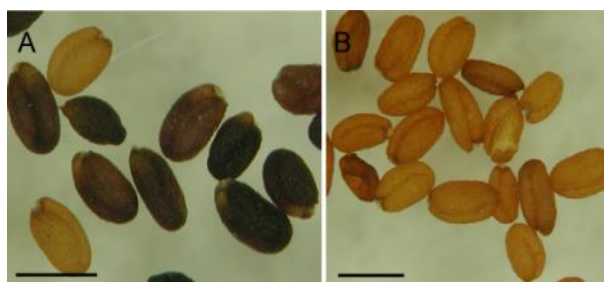


Figure 2 Segregating seed coat color of *Arabidopsis* in T2 generation. (A) Seeds of 35S:*VlmybA1-2* plant showing two colors, purple and brown. Brown seeds were similar to wild type seeds in color (B) Wild type. Bar= 1 mm



Figure 3 Co-segregation of *VlmybA1-2* and kanamycin resistance in T2 *Arabidopsis* seeds on MS medium after one week of seed germination. (A) Purple seeds could germinate on 50 mg l⁻¹ kanamycin containing MS medium (B) Brown seeds could not germinate on kanamycin containing MS medium.

nating seedlings were with green leaves and purple roots. Some of them produced all purple T3 seeds while some produced both purple and brown seeds. Germinating all purple seeds and some purple seeds of both colors, produced completely purple plants, out of which could produce purple flowers (Fig 4; A and D), plants with purple-green leaves and well grown purple roots, which could produce white flowers (Fig 4; B and E) and plants with translucent green leaves and purple roots (Fig 4; C).

Plant regeneration from transgenic tobacco callus

Purple protrusions originated from leaf explants after 2 weeks of transformation by the pBI121 vector carrying *VlmybA1-2* cDNA. Purple callus proliferated from such purple protrusions (Fig 5) and purple shoots regenerated from them. Purple roots were subsequently regenerated and were distinguishable from plants regenerated from wild type or control vector transformants by color (Fig 6). The transgenic plants grew normally and produced flowers (Fig 7). The corolla of the transgenic flow-

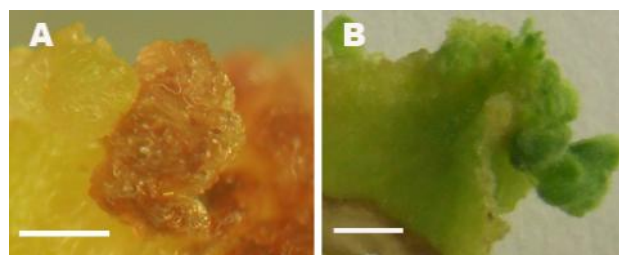


Figure 5 Expression of *VlmybA1-2* in transformed tobacco leaf derived callus. (A) 35S:*VlmybA1-2*. (B) Wild type. Bar = 1 mm.

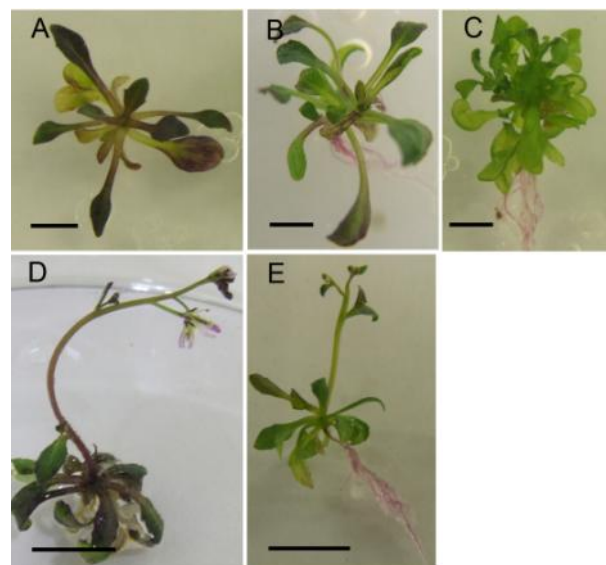


Figure 4 Different phenotypes of T3 *Arabidopsis* plants of 35S:*VlmybA1-2* after 3 weeks of seed germination. (A) A completely purple plant (B) A plant with purple-green leaves and purple roots (C) A plant with abnormal green leaves and purple roots (D) A purple plant with purple flowers at flowering (E) A purple-green plant at flowering. Bar = 1 cm

ers was dark purple and distinguishable from the light pinkish corolla of wild type flowers.

VlmybA1-2 expression in transgenic tobacco plants

Five independent transgenic regenerants were checked for transgene expression by RNA gel blot hybridization. All transgenic regenerants expressed high levels of the transgene (Fig 8).

Segregation of *VlmybA1-2* in T1 seedlings of tobacco

Primary transgenic plants produced fertile T1 seeds. Kanamycin resistance and *VlmybA1-2*-induced purple color were co-expressed. Surviving seedlings were completely purple and greenish purple. Completely green seedlings could not grow beyond the cotyledon stage and died on kanamycin-

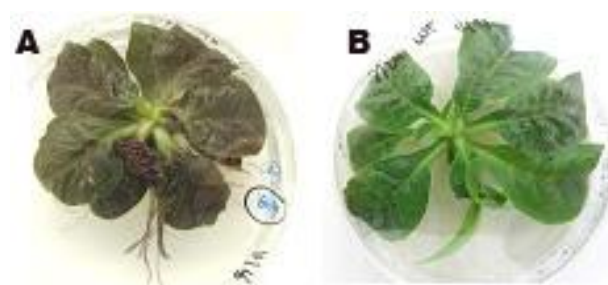


Figure 6 A 35S:*VlmybA1-2* tobacco plant regenerated from leaf callus on root formation medium. (A) A 35S:*VlmybA1-2* transgenic plant with purple leaves and purple roots was distinguishable from a control vector transformant (B).

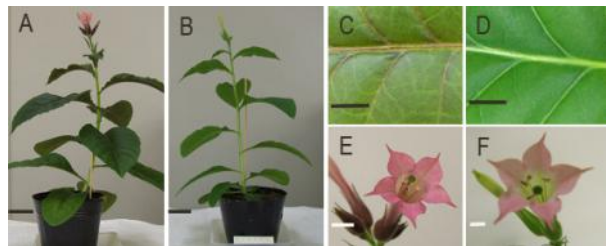


Figure 7 A regenerated 35S:VlmybA1-2 tobacco plant grown under the SD condition in a pot (A). Flowering occurred around 2 months after pot transfer. Bar = 4 cm. (C) A magnified view of a part of a 35S:VlmybA1-2 leaf. Bar = 5 mm. (E) A 35S:VlmybA1-2 flower. Bar = 1 cm. (B)(D) and (F) Wild type.

containing medium (Fig 9). Ratio between surviving completely purple and greenish purple seedlings to dead green seedlings was 53:17 (approximately 3:1) in an observed progeny of a line (line one), indicating that the Mendelian inheritance of a single gene locus.

***VlmybA* induced anthocyanin production in tobacco**

The purple pigments from tobacco leaf callus, regenerated plants and T1 seedlings could be extracted with water and became dark blue with the addition of 0.1M sodium hydroxide. These results suggest that the pigment is anthocyanin. The absorbance values at 525nm of the acetic acid extracts per g fresh weight tissue were interpreted as the anthocyanin content. The absorbance values were 235.51 ± 12.31 (standard error, $n=3$) and 250.31 ± 20.45 (SE, $n=3$) in purple line 1 plants and line 2 plants, respectively, while they were 153.14 ± 1.47 (SE, $n=3$) and 136.67 ± 7.18 (SE, $n=3$) in greenish purple line 1 plants and line 2 plants, respectively. An average absorbance of 1.42 ± 0.19

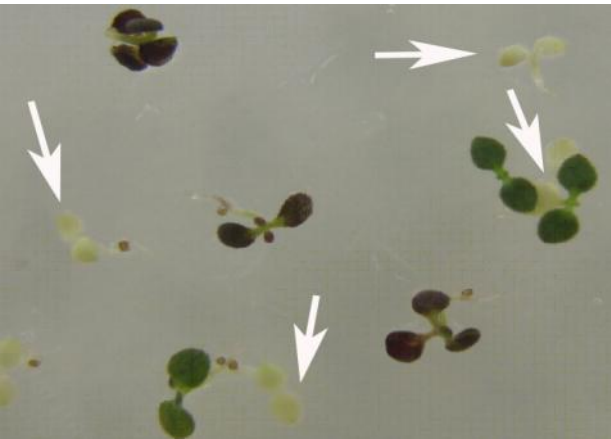


Figure 9 Segregation of 35S:VlmybA1-2 tobacco seedlings of T1 generation on kanamycin-containing medium. Complete purple seedlings and seedlings with partially purple-green leaves could survive on kanamycin containing medium. White arrows indicate dead seedlings. These plants are derived from the primary transformant corresponding to line 2 in Figure 4.

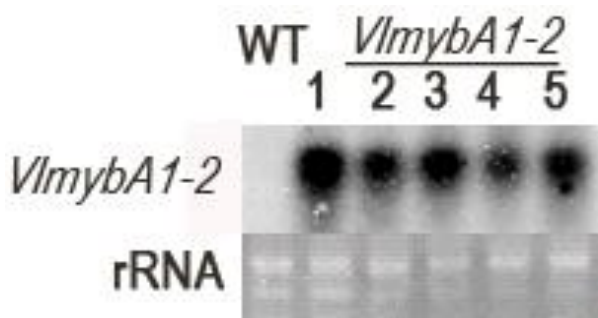


Figure 8 RNA gel blot hybridization of tobacco primary transformants. WT: Wild type. Numbers 1 to 5 are blots for five regenerated independent 35S:VlmybA1-2 plant lines.

(SE, $n=3$) at 525nm was recorded from the wild type.

***VlmybA1-2* could not induce anthocyanin production in spinach**

Both infected and wild type spinach cotyledon bases were purple after 3 days of culture, giving no indication for transformed tissue identification. A few regenerated putative transgenic spinach plants, similar to wild type in color and appearance, could survive on kanamycin- containing medium (data not shown) and the presence of *VlmybA1-2* was confirmed by DNA gel blot hybridization in one of them, which could survive on 50mg l^{-1} kanamycin-containing medium for more than 4 months (Fig 10): number of regenerated plants after transformation was limited due to low regeneration frequency of spinach.

DISCUSSION

The over-expression of *VlmybA1-2* alone could induce anthocyanin pigmentation in whole plant in *Arabidopsis* and tobacco transformants. As *VlmybA* are transcription factor genes in the anthocyanin biosynthetic pathway in a dicot, it may bind to the promoters of the structural genes of the anthocyanin biosynthetic pathway in them, which may be conserved among anthocyanin-producing dicots.

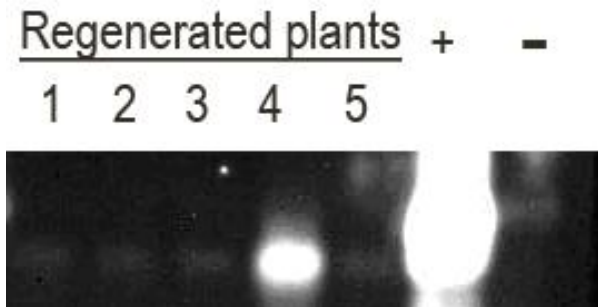


Figure 10 DNA gel blot hybridization of spinach primary transformants: Numbers 1 to 5 are blots for five regenerated independent putative 35S:VlmybA1-2 plants. (+) Positive control. (-) Wild type.

Together with the previous result of *VlmybA2* (Geekiyanage *et al.* 2007), and results of present study for *Arabidopsis* and tobacco, it is postulated that *VlmybA* genes can function in a variety of anthocyanin-producing dicots.

The copy number of the transgene or the localization of transgene insertion in the genome may determine the level of transgene expression resulting in phenotypic differences in T1. Furthermore, all T3 *Arabidopsis* purple seeds and some of T2 purple seeds derived from both color T1 plants, produced completely purple plants including flowers. Normal plant growth was disturbed in some T1 transformants of *Arabidopsis*: Two completely purple seedlings did not survive beyond the cotyledon stage, while the surviving partially purple seedlings with purple roots produced seeds that segregated according to the expected Mendelian ratio of 3:1 for the inheritance of a single gene locus. Furthermore, the completely purple T1 seedlings showed retarded root growth or had no roots at all. A similar phenotype was lethal in tomato transformed by *Lc* (Goldsbrough *et al.* 1996). Flavonoids act as negative regulators of auxin transport in *Arabidopsis* (Brown *et al.* 2001). The effect on root formation could be due to the negative regulation of auxin transport by the over-expressed flavonoids in *Arabidopsis*.

VlmybA genes induce remarkably the expression of the *UFGT* gene, while the expression of the other anthocyanin structural genes is slightly increased in grape (Kobayashi *et al.* 2002). In *Arabidopsis*, the color of testa is mainly due to the oxidation of proanthocyanidins (Chapple *et al.* 1994). In the present study, the pigmentation in *Arabidopsis* seeds, making it a distinguishable phenotype, could be due to the enhanced production of anthocyanin or proanthocyanidins. This distinguishable phenotype could be observed in 35S: *VlmybA2* as well (Geekiyanage *et al.* 2007) and suggests that in *Arabidopsis*, the *VlmybA2* and *VlmybA1-2* gene product/s may activate not only *UFGT* but also other structural genes that are upstream or at the branched end towards proanthocyanidin in seed coat. Due to this phenotype, *VlmybA* genes provided an efficient and simple method for the identification of transformed seeds in *Arabidopsis*.

VlmybA1-2 expression in *Arabidopsis* seems to be promising for visual identification of transformants as all T1 transformants and their progenies showed distinguishable purple color to the wild type. Plants with green leaves and normal growth were even distinguishable by purple roots, and ability to induce purple color in roots is a unique feature in *VlmybA* genes compared to previously used other *myb* or *myc* genes. Furthermore, although anthocyanin pigmentation can be induced by increased irradiation or stress in other parts of the

plant (Rabino and Mancinelli 1986), there are no reports on such pigmentation in roots. Expression of *VlmybA1-2* in seeds provided an efficient and unique method of transgenic seed identification in *Arabidopsis*. However, level of expression may affect the normal growth in *Arabidopsis*.

Numerous environmental, developmental and tissue- and cell-specific factors are involved in anthocyanin biosynthesis. In the present work, the distribution and intensity of purple color of the entire regenerated tobacco shoots were affected as purple color was prominent in the veins of leaf upper surface and lower leaf blade of pot-grown plants (Fig 7). The differences in anthocyanin distribution in the leaves of 35S: *VlmybA1-2* plants could have resulted from the availability of endogenous factors, which may be dependant on developmental stage, and/or on environmental conditions.

Anthocyanin content in 35S: *VlmybA1-2* tobacco calli/ plants were several times higher than the control, indicating its potential for development of plant cell systems that produce stable anthocyanins for applications as natural food colorants in bio-industry.

Spinach belongs to Caryophyllales, which produce betacyanins instead of anthocyanins. Anthocyanin and betacyanins are produced in two mutually exclusive pathways in plants (Stafford, 1994). The presence of the first committed structural gene in anthocyanin biosynthetic pathway, dihydroflavonol 4-reductase (*DFR*), in betalain-producing plants suggests that the lack of transcription factor genes could be a reason for inability to produce anthocyanin in them (Shimada *et al.* 2004). *VlmybA* genes (Kobayashi *et al.* 2002) and previously cloned regulatory genes such as *C1* (Cone *et al.* 1986; Paz-Ares *et al.* 1987), *AN2* (Quattrocchio *et al.* 1998), *DELILA* (Goodrich *et al.* 1992), *Lc* (Ludwig *et al.* 1990), *AN11* (Quattrocchio *et al.* 1993) and *TTG1* (Shirley *et al.* 1995) would provide a means to test this hypothesis in transgenic betalain-producing plants. Lack of distinguishable pigmentation from control in the *VlmybA1-2* spinach plant and regenerated shoots on kanamycin medium, may be due to the inability of *VlmybA1-2* to induce structural genes of anthocyanin biosynthesis pathway in non anthocyanin producing spinach (Shimada *et al.* 2007).

VlmybA1-2 expression in tobacco seems to be promising for visual identification of transformants as all tobacco primary transformants and their progenies are apparently not different from the wild type except for the purple color. Flavonoids act as negative regulators of auxin transport in *Arabidopsis* (Brown *et al.* 2001). Negative effect on normal plant growth by over-expressed flavonoids in *Arabidopsis* was not observed in transgenic tobacco plants.

Most commonly used markers are the bacterial genes conferring resistance to antibiotics kanamycin and hygromycin or to herbicide glufosinate ammonium. There is a growing social concern over use of these genes in transgenic plants in eco systems and human consumption. In addition, there are some evident drawbacks of use of these genes because selection chemicals, like antibiotics may either interfere with plant regeneration, resulting in a fewer transgenic plants or may be less effective as plant tissues may exhibit a high level of intrinsic antibiotic resistance, which is misleading during selection.

A versatile gene for anthocyanin production could be a good candidate for a simple and non-destructive visual marker for plant transformation and may replace controversial antibiotic marker genes. *VlmybA* genes show higher potential than the other anthocyanin regulatory genes previously tested (Ludwig *et al.* 1990; Lloyd *et al.* 1992; Mooney *et al.* 1995; Goldsbrough *et al.* 1996; Deluc *et al.* 2006). Its potential should be exploited in the genetically modified plant production process, starting from the efficient recovery of regenerants during transformation up to the monitoring of transgenic plants at field level for risk assessment, and development of plant cell systems that produce stable anthocyanins for applications as natural food colorants in bio-industry.

CONCLUSIONS

- 35S:*VlmybA1-2* can induce anthocyanin pigmentation in whole plants of tobacco and *Arabidopsis*.
- High levels of anthocyanin production are associated with lethality and abnormality in *Arabidopsis* seedlings.
- Color of 35S:*VlmybA1-2 Arabidopsis* seeds can be used for transgenic seed identification.
- *VlmybA1-2* shows a potential to be used as a visual marker for anthocyanin producing plants

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