

Induction of Somatic Embryogenesis through Overexpression of *ATRKD4* Genes in *Phalaenopsis* "Sogo Vivien"

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Abstract

Phalaenopsis "Sogo Vivien" is a mini orchid hybrid with beautiful flowers and numerous inflorescences. Mass propagation of this orchid is needed to meet the market demand. Objective of this research was to induce somatic embryogenesis of *P.*"Sogo Vivien" through insertion of *AtRKD4* gene into orchid. T-DNA containing 35S::GAL4::*AtRKD4*::GR was inserted into 16-22 days after sowing orchid protocorms mediated by *Agrobacterium tumefaciens* EHA 105. Activation of the *AtRKD4* gene was induced by glucocorticoid induction system, using 15µM Dexamethasone (Dex). The results showed that 34 out of 2,648 orchid embryos developed into protocorms on hygromycin selection medium, whereas only 4 out of 2,897 non-transformant protocorms developed from embryos. A 500 bp of *HPT* genes was amplified from transformant candidates using specific primers for *HPT* (HygF1 and HygR1) and 380 bp was amplified using specific primers for *AtRKD4* (*AtRKD4* F1 and *AtRKD4* R1), indicated that transgenes have been integrated into orchid genomes. Finally, 17 plantlets were positively carrying *AtRKD4* and *HPT* genes, the efficiency of transformation was 0.63 %. Somatic embryos were also emerged from leaf explants of transformant on hormone-free NP medium and became normal plantlets. It is probably due to the high activity of *AtRKD4* genes in orchids.

Keywords: *AtRKD4*, glucocorticoid, *Phalaenopsis* "Sogo Vivien", orchid hybrid, somatic embryo

Introduction

Phalaenopsis "Sogo Vivien" is an orchid hybrid with beautiful striped purple flowers and dark purple labellum and variegated leaves, define as green leaves having golden yellow ribbon on their edges (McKinley, 2005). With its mini size and many flowers, the orchid is included in the desired criteria for pot plants (Tang and Chen, 2007).

Conventional propagation by using the buds could produce shoots, but it is inefficient because it needs a long time and produces an only small number of plants due

to each bud produces only one plant (Kosir *et al.*, 2004). Propagation using seeds is highly depending on the fruits that resulted from pollination. Therefore, it needs an effective propagation method to produce uniform seedlings in large numbers and in relatively short time.

A breakthrough, which could increase the efficiency of orchid micropropagation is by combining the micropropagation method and the genetic engineering. *Agrobacterium tumefaciens*-mediated genetic transformation method was used to produce a variety of orchid plants (Semiarti *et al.*, 2007; Semiarti *et al.*, 2010; Sreeramanan and Zuraida, 2010).

In plants, some genes involve in somatic embryogenesis. Some of the genes that responsible for the initiation and development of embryogenesis have been

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identified (Radoeva and Weijers, 2014), some of which induce ectopic embryogenesis on the model plant of *Arabidopsis thaliana*, the *LEAFY COTILEDON (LEC)* genes (Braybrook and Harada, 2008), *BABY BOOM (BBM)* gene (Boutilier, 2002), *WUSCHEL (WUS)* homeobox gene (Zuo *et al.*, 2002) and *RWP-RK DOMAIN CONTAINING-4 (RKD4)* (Waki *et al.*, 2011).

RKD4 gene encodes a protein with RWP-RK motif in *Arabidopsis* required for embryonic pattern formation (Chardin *et al.*, 2014). This gene encodes a transcription factor protein that expressed very early in zygotic embryogenesis (Waki *et al.*, 2011). Waki *et al.* (2011) inserted *indRKD4ox* genes into *Arabidopsis* and proved that overexpression of these genes after induction using synthetic steroid hormones such as Dexamethasone (Dex) on the roots of *A. thaliana* transformant, produced somatic embryos.

Utilization of glucocorticoid regulation system is a general tool for regulating gene expression in plant cells and whole plants. Expression of the glucocorticoid receptor protein from a stage- or tissue-specific promoter in plants could conceivably provide a way to target glucocorticoid-inducible gene expression to specific times in development or to specific plant tissues. Experiment with transgenic plants expressing the glucocorticoid receptor was conducted by Schena *et al.* (1991) that determined the extent to which glucocorticoid induction can be achieved in *A. thaliana*. The mammalian steroid receptors can be used function as hormone-dependent inducers of gene expression in heterologous systems, and the feasibility of using mammalian steroid hormones as selective inducers of plant gene expression was investigated. Schena *et al.* (1991) found that in glucocorticoid induction system, the treatment with glucocorticoid hormone can activate the glucocorticoid receptor expression in plant cells. This system can be used to activate a test gene linked to glucocorticoid response element. These findings indicate that glucocorticoid

induction of steroid responsive promoters provide a general method for regulating gene expression in plant cells, and imply that such a system might ultimately function in whole plants such as *A. thaliana* and others. Overexpression of the transgene in these systems could only be induced in transgenic plants when there is glucocorticoid response element in the plant genome and the existence of glucocorticoid hormone in the medium (Kang *et al.*, 1999). Dexamethasone as glucocorticoid hormone was successfully used to induce the expression of *Myc-XA21* gene in transgenic rice plants (Park *et al.*, 2012) and to activate *Luc* genes in *A. thaliana* transgenic (Aoyama and Chua, 1997). However, the technique of embryo somatic induction using glucocorticoid system in orchids, mainly *P. "Sogo Vivien"*, has not been reported.

In this paper we report the use of glucocorticoid selective inducible system to overexpress *AtRKD4* transgene in orchid for inducing somatic embryogenesis of *P. "Sogo Vivien"* orchid hybrid.

Materials and Methods

Plant material and culture conditions

Plant material used in this study was *P. "Sogo Vivien"* orchid hybrid obtained from CV. Amabilis nursery, Kiaradong, Bandung. Four-year-old flowering plants were planted in the pots with sphagnum moss medium, maintained in a greenhouse under natural lighting at temperatures between (25 and 30)° C. Mature pods from self-pollinated flowers of variegated *P. "Sogo Vivien"* plants were used as plant materials for all of the experiments.

Seed plantation for protocorms preparation

A three-month-old orchid pod from self-pollinated variegated *P. "Sogo Vivien"* was used as a seed source. The pod was subsequently sterilized by dipping it in alcohol and then flame it on a Bunsen lamp. Next, the seeds were sowed on NP medium

(Islam *et al.*, 1998). Intensive observation was conducted every day to observe the development stage of the seeds that grew into protocorms, which was carefully examined using a stereo microscope (Eschenbach Binoculars DM150DB, Germany), and were photographed using digital camera (Canon A2300 HD, Japan), and Optilab (Miconos, Indonesia) with LED light system.

Plasmid vector and bacterial strain

The plasmid used was pTA7002 with the T-DNA construction of 35S::GAL4::AtRKD4::GR (obtained from Dr. Jose Gutierrez-Marcos, University of Warwick, UK; Fig.1), which was contained in *Agrobacterium tumefaciens* strain EHA105. Isolation of plasmid was conducted using the alkali lysis method (Green and Sambrook, 2012). The T-DNA consists of multiple strong promoters 35S, GVG, GAL 4 and VP16 with Glucocorticoid response element (GRE) for selective inducible system with glucocorticoid hormone (Dexamethasone). For antibiotic selective system, *Hygromycin phosphotransferase* (HPT) was driven by *Nopaline Synthase* (NOS) promoter and the target gene *AtRKD4* from *A. thaliana*.

Transformation and regeneration of transformants

Overnight cultures of *Agrobacterium* in LB medium were diluted 1 : 4 (v/v) using NP liquid medium supplemented with 2% (w/v) sucrose, a phenolic natural product acetosyringone 50 mg.L⁻¹ (Bio Basic, Canada). Three-weeks-old protocorms were immersed in the diluted culture of *Agrobacterium* for an hour, then transferred its onto sterile nylon cloth to dry in laminar air flows and then transferred onto modified NP solid medium

containing 2 mg.L⁻¹ 2,4-D (Sigma), 100 g.L⁻¹ tomato extract, and 25 mg.L⁻¹ acetosyringone, for cocultivation. At day-3, protocorms were washed 3 times using sterile distilled water, then soaked it in a half-strength of NP liquid medium without antibiotics, protocorms then shaken using a shaker with the speed of 100 rpm for 2 nights, the medium was renewal every 24 hours. Next, protocorms were rinsed 3 times using sterile distilled water, drained again for 30 min. Once dried, protocorms were transferred onto solid NP medium containing 25 mg.L⁻¹ meropenem (Sanbe) to inhibit the growth of *Agrobacterium*, and were cultured for a week. After that, protocorms were moved onto NP medium (selection medium) containing 25 mg.L⁻¹ meropenem and 10 mg.L⁻¹ hygromycin (Wako, Japan), for selection of the transformants. The developing protocorms were later transferred onto a new medium for every 2 weeks for further selection of the transformants. Developing shoots as the candidate of transformants were screened by PCR using *AtRKD4* and *HPT*-specific primers to confirm that they were the successful transformants. Efficiency of transformation was measured from the number of positive transformants for both *HPT* and *AtRKD4* transgene in the genome per total protocorms examined, multiply by 100%.

Polymerase chain reaction analysis of transformants

Genomic DNA from the putative 35S::GAL4::AtRKD4::GR transformants was analyzed by direct PCR using KAPA3G Plant PCR Kit (KapaBiosystem), using specific primers for the *AtRKD4* gene: *AtRKD4F*, 5- GTTCATTTTCATTTGGAGAGGACG-3 ; *AtRKD4R*, 5- CTTCCATATCTAGGAGAG AAT CAAG-3 and specific primers for the

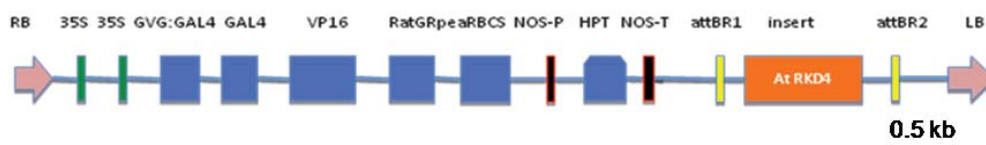


Figure1. The structure of T-DNA carrying 35S::GAL4::AtRKD4::GR in plasmid pTA7002/EHA105

HPT gene: *HygF*, 5-TCGGACGATTGCGTCCATC-3; *HygR*, 5-AGGCTATGGATCGATCGCTG-3. PCR for both genes was performed with 30 cycles of 95 °C for 30 s, 58°C for 30 s, and 72°C for 30 s using a Thermal Cycler (Biorad T100™). Amplified DNA fragments were separated on a 1,4% agarose gel (Sigma), stained with SYBR Safe DNA Gel Stain (Invitrogen), and visualized under Ultraviolet Viewer (ExtraGene).

Induction of embryo somatic and Analysis of *AtRKD4* gene expression

The leaves originated from plantlets, which grown on selection medium were subsequently taken and cut into pieces, and then planted on NP medium containing 15 µM Dexamethasone (Dex, Merck) for a week in order to induce somatic embryo formation. After that, the leaf pieces were transferred onto hormone-free NP medium. Observations were performed on the development of leaf explants to examine and count the number of somatic embryos using a stereo microscope, Optilab and photographed using a digital camera.

Results and Discussion

The growth of protocorms as the target of transformation

The seeds were obtained from a healthy mother plant of *P. "Sogo Vivien"*, that produced good quality of seeds in three months old mature fruit/pod (Fig. 2).

Development of zygotic embryos in the seeds of *P. "Sogo Vivien"* into protocorms on NP medium was observed intensively for one month (Fig. 3). At day 0, the embryo was white yellowish in color surrounded by testa (Fig. 3a). It gradually grew into swollen embryo at day 5 with the volume become twice bigger than before (Fig. 3b). Later, the color of embryo were gradually changed into yellow-green (Fig. 3d-e) and then turned to green (Fig 3f). At 29 days after sowing (DAS), absorbing hairs were gradually emerged from the basal part of the embryos and the growth of embryo significantly increased up to 6 times (Fig. 3f). In this study, seeds were planted on solid NP medium containing 150 mL.L⁻¹ coconut milk and covered by nylon cloth to facilitate convenient technique on protocorm harvesting prior to genetic transformation. Based on morphological change (size, color, shape, and the emergence of absorbing hairs) of embryo into protocorm, it can be classified into five phases as follow: Phase 1 (5 DAS)-seeds had swollen embryo due to water uptake from the medium, white yellowish in color, and most of testa (seed coat) had broken (Fig.3b); Phase 2 (11 DAS)- enlarged embryos, they were white yellowish in color and globular in shape (Fig. 3c); Phase 3 (16 DAS)- the embryo enlarged and changed the color from white to yellow, the polarity of the embryo appeared bipolar, where the anterior part was yellow-green in color, and the posterior part was yellowish (Fig. 3d). Phase 4 (21 DAS)-the embryo enlarged

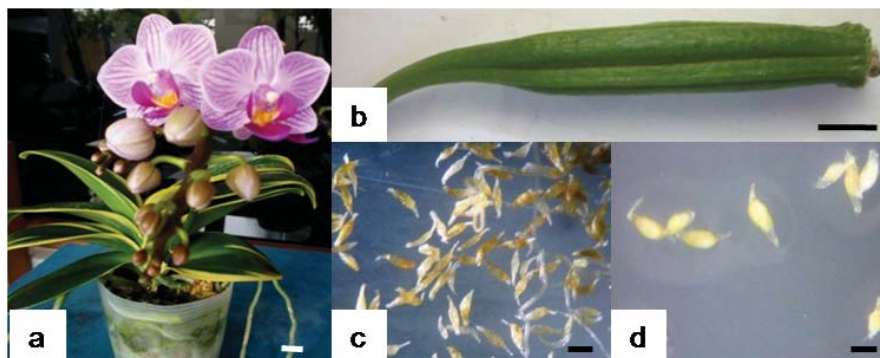


Figure 2. Phenotype of *Phalaenopsis "Sogo Vivien"* orchid hybrid. (a) Mother plant; (b) self-pollinated fruit; (c) seeds before planting and (d) seeds at 2- days after sowing (DAS) on NP medium. Bars: 5 mm.

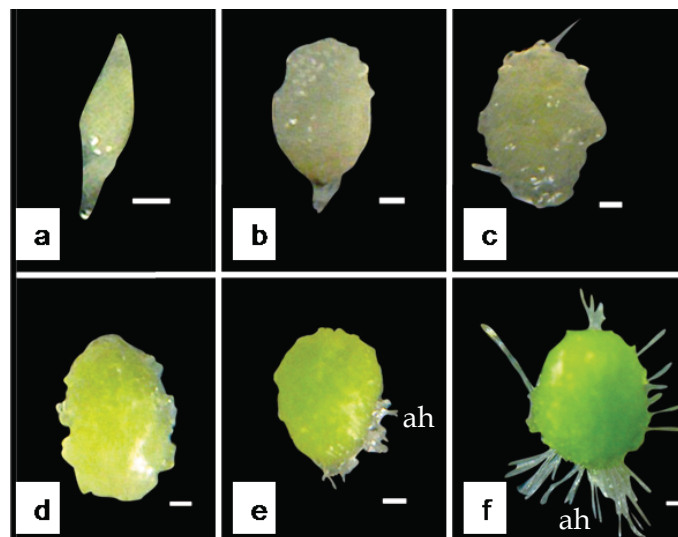


Figure 3. Development of *Phalaenopsis* "Sogo Vivien" orchid embryos on NP medium. (a). 0 DAS; (b). Swollen embryo with broken testa at 5 DAS; (c). yellowish embryo at 11 DAS; (d). Yellow-green embryo at 16 DAS; (e). Greenish embryo with absorbing hair at 21 DAS; (f). Bright green embryo with many numbers of absorbing hairs (ah) at 29 DAS. Bars 0.1 mm.

and changed the yellow color to green, and started to exhibit absorbing hairs (Fig. 3e). Phase 5 (29 DAS)- the embryo enlarged with green color, the anterior part showed leaf primordia, and had much number and more dense of absorbing hairs (Fig. 3f). Based on the morphological changes during embryo development of *P. "Sogo Vivien"*, therefore protocorms in phase 4 (21 DAS) (Fig. 3a) were selected in this study as the target for genetic transformation using *Agrobacterium*.

This phenomenon was very similar to that of embryo development in the wild orchid *Phalaenopsis amabilis* as reported by Semiarti *et al.* (2007), in which 21 DAS protocorms were used as the target for *Agrobacterium*-mediated transformation. Bunnag and Pilahome (2012) have also used protocorms as the transformation target. Although many researchers used protocorm-like bodies (plb) as the target for the genetic transformation in orchids (Chan *et al.*, 2005; Shrestha *et al.*, 2007; Subramaniam *et al.*, 2009; Zhang *et al.*, 2010).

The use of protocorms as transformation target has several advantages, i.e. it can easily

and quickly produced in large quantities as it comes from the growth of seeds, it has a high level of regeneration, and it allows transfer of genes into protocorm (Mishiba *et al.*, 2005).

Regeneration and selection of hygromycin-resistant protocorms

Genetic transformation using protocorms was conducted in several steps, namely the pre-culture of protocorms, co-cultivation of protocorm with bacteria carrying the recombinant DNA, elimination of bacteria, selection of transformants candidate, and regeneration of transformant candidates. The medium used for pre-culture contained 2,4-D, tomato extract and acetosyringone, which aimed to keep the meristematic level of protocorm, protected from oxidative damage, and to improve the response level of protocorm when gene transfer was performed (Mishiba *et al.*, 2005 ; Semiarti *et al.*, 2010; Dwiyani *et al.*, 2010). In this research, co-cultivation was carried out in 3 days to optimize the integration of T-DNA into the genome of protocorms. For the elimination of bacteria,

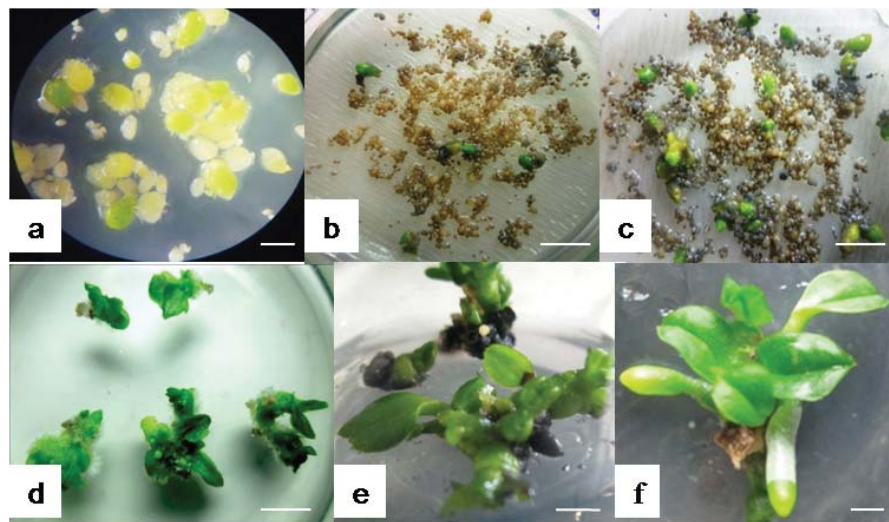


Figure 4. Regeneration of *Phalaenopsis* "Sogo Vivien" transgenic plants from protocorms. (a) 3 weeks after sowing (WAS)-protocorms used as transformation target; (b-c) 9-WAS of Non-transformant protocorms grown on hygromycin selection medium (b); and Transformant line #1 (T#1) protocorms which are able to form buds on selection medium (c); (d) 16-WAS T#1 protocorms, which are able to regenerate into plantlets; (e-f) Regenerated T#1 plants at 18-WAS (e) and at 22-WAS (f). Bars 1mm in a, 5 mm for b-f.

antibiotics meropenem was used as it could effectively disturbed the growth of *Agrobacterium* (Mishiba *et al.*, 2005). The selection of transformant candidates was done in 6 weeks with intensive subcultured for every 2 weeks on selection medium containing hygromycin 10 mg.L^{-1} . The use of the antibiotics was in accordance with the existence of *HPT* gene in T-DNA that caused it resistance to hygromycin (Fig.4).

Total number of protocorm that able to grow on hygromycin selection medium was 46 protocorms, generated from 3-times-experiments with 2,648 transformed protocorms examined (Table 1, Fig.4). A total 34 protocorms formed buds on selection medium (Fig.4c) and of these only 20 protocorms were able to regenerate into plantlets (Fig.4d). Almost all of non-transformant protocorms were unable to grow on selection medium containing hygromycin, only 6 out of 2,897 protocorms could grow, the others embryos were turned the color into dark brown, due to browning process and necrosis (Fig.4b). Overall, the

morphology of the transformant plant candidates were relatively similar to their wildtype plant (Fig. 4e, 4f). The average of transformation frequency obtained from 3-times-experiment was relatively low, about 0.63%, as shown by that very view number of protocorms undergone genetic transformation.

Opabode (2006) stated that the main factors causing low efficiency of transformation is the hypersensitive plant defense reaction against *Agrobacterium* infection inducing excessive production of reactive oxygen species (ROS). This compound is highly reactive towards biomolecules, which cause damage of plant tissue followed by browning and necrosis (Kuta and Tripathi, 2005). Several studies of genetic transformation using *Agrobacterium* with transformation targets in the form of protocorm-like bodies (plb) and orchid protocorm obtained transformation efficiency of 4.3% in *Cattleya* (Zhang *et al.*, 2010); 1.91% in *Phalaenopsis* hybrid (Mishiba *et al.*, 2005); 1.7% in *Phalaenopsis amabilis* (Semiarti *et al.*, 2007); and 1.3 to 2.7% in

Table 1. Efficiency of transformation of *Phalaenopsis* "Sogo Vivien"

	Experiment	No. of protocorms examined	Positive selective medium	No. of protocorm producing shoot	No. of regenerated plants	PCR Detection		Efficiency of transformation (%)*
						HPT+	AtRKD4+	
Non-transformant (NT)	1	762	6	3	2	0/2	0/2	0
	2	869	6	3	2	0/2	0/2	0
	3	1266	8	4	3	0/5	0/5	0
	Total	2897	20	10	7	0/9	0/9	0
35S::GAL4::AtRKD4::GR	1	1044	19	14	9	8/9	8/9	0.77
	2	842	15	11	7	5/7	5/7	0.59
	3	762	12	9	4	4/4	4/4	0.52
	Total	2648	46	34	20	17/20	17/20	0.63 (Average)

* The calculation of the efficiency of transformation see materials and methods

Oncidium and *Odontoglossum* (Raffener *et al.*, 2009). To improve the transformation efficiency, there are many studies using a variety of treatments, such as the provision of antioxidants, desiccation of transformation target before and after infection with *Agrobacterium*, length of co-cultivation time, density of *Agrobacterium tumefaciens* used in the transformation, temperature of co-cultivation process, and the addition of surfactant at the time of *Agrobacterium tumefaciens* infection to the explants (Opabode, 2006). The low transformation efficiency in *P. "Sogo Vivien"* might caused by the low capability of protocorms to face the oxidation processes as a response to infection by *Agrobacterium*.

Molecular analyses of transformants

Molecular analysis to confirm the existence of *AtRKD4* and the *HPT* genes in the genome of transformant plants was performed on plantlets of transformant candidates that grown on medium containing 10 mg.L⁻¹ hygromycin. Seventeen plantlets out of 20 transformant plant candidates showed positive results for both the existence of *AtRKD4* and *HPT* genes in the genome, indicated by the detection of amplified DNA fragments in the size of 500 bp of *HPT* genes in electrophoresis gel (Fig. 5 lane 4-6) and

380 bp of *AtRKD4* (Fig.5 lane 11-13) and the summarize data in Table 1. The bands did not appear on the samples of non-transformant plantlets (Fig.5 lane 3 and lane 10). As a positive control, the PCR reaction and electrophoresis was performed using plasmid pTA7002 that contained T-DNA carrying 35S::GAL4::AtRKD4::GR construct, and resulted a 380 bp DNA band for *AtRKD4* gene (Fig.5 lane 2) and 500bp DNA band for *HPT* gene (Fig. 5 lane 9). These data showed that the T-DNA constructs were successfully integrated into the genome of the *P. "Sogo Vivien"* transformant candidates.

Phenotypes of orchid transformants expressing *AtRKD4* genes

To induce overexpression of *AtRKD4* gene in the genome of *P. "Sogo Vivien"* transformant candidates, glucocorticoid-inducible system using Dexamethasone (Dex) were generated. This is important to do because the T-DNA construct in pTA7002 plasmid contained glucocorticoid response element (GRE) that can be activated after the treatment of glucocorticoid hormone to the plants. For inducing *AtRKD4* gene expression, leaves were taken from 3.5 months old positive transformants plantlet #1 on selection medium (T#1). The T#1 leaves were cut into pieces and planted on

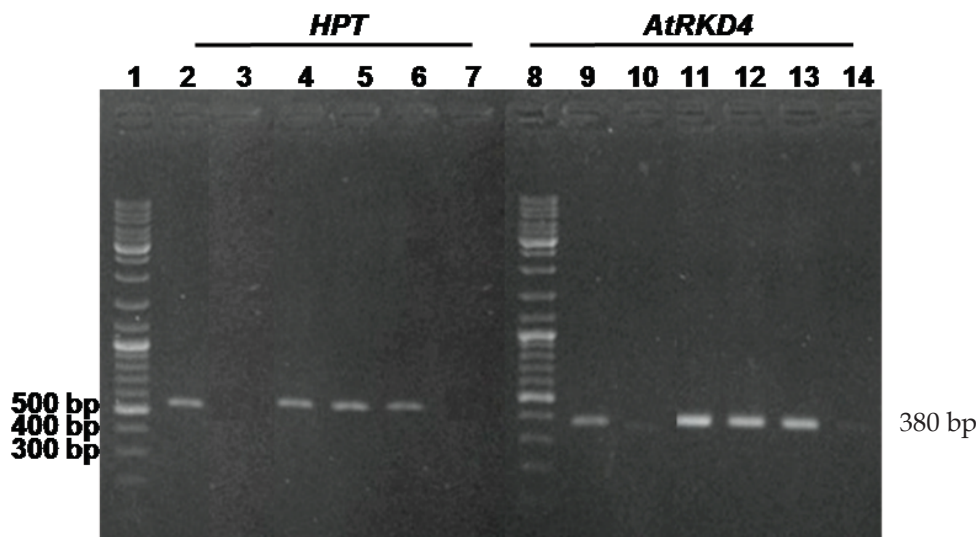


Figure 5. Detection of *HPT* and *AtRKD4* genes integration in the genome of *P. "Sogo Vivien"* transformants. Lane 1 and 8: 1000 bp DNA ladder as size maker; lane 2: amplified 500 bp DNA fragment of *HPT* gene from pTA7002 plasmid; lane 4-6 transformant candidates T#1, T#6, T#11, respectively; lane 7: T#13 transformant candidate that produced no band. Lane 9-14 indicated 380 bp *AtRKD4* DNA fragment: Lane 9 from pTA7001; lane 10: Non transformant plants; lane 11-13 Transformants candidates T#1, T#6, and T#11, respectively; Lane 14: T#13 transformant candidate that produced no band.

NP solid medium containing 15 μ M Dex for 1 week. Furthermore, the leaf pieces were subcultured on Dex-free NP medium.

The results showed that somatic embryos were emerged from T#1 leaf pieces that grown on hormone-free NP medium (Fig.6). However, there was not observed in all pieces of leaves from NT plants, in which they quickly became browning and finally death. In T#1 leaf pieces, the somatic embryo formation began with the swelling leaves on day-8, followed by the formation of pre-globular somatic embryos on day-18, then it developed to globular embryo at day-33, the shape was very clear after reach day-59. The observation showed that the formation of somatic embryos was occurred on the entire surface of the leaf on the average 72 somatic embryos produced per 0.5 cm² leaf pieces at day-80. Regenerated plantlets from somatic embryos were obtained at day-140, that showed normal phenotype similar to the phenotype of original non transformant plant (Fig.6). This data showed that overexpression of *AtRKD4* gene had been induced by

Dexamethasone and was able to function in orchid to form somatic embryos from leaf pieces of T#1 transformant plant, that finally produced 413 plantlets (Table 2).

Glucocorticoid-inducible expression system is a system that uses regulatory mechanisms of steroid hormone receptors to induce the expression of a gene. This system contains components GVG gene that produces the protein as a transcription factor that regulates glucocorticoids, and components of the gene to be inserted into a plant (Aoyama and Chua, 1997). Transgenic plants which carry plasmids with glucocorticoid-inducible system will express the inserted genes when the glucocorticoid compound was present (Kang *et al.*, 1999). Waki *et al.* (2011) reported that in *Arabidopsis thaliana* plants, the steroid compound in 10 μ M Dexamethasone can induce *AtRKD4* gene expression to the formation of somatic embryos from cotyledons and roots of *A. thaliana* transformants. One hour treatment of Glucocorticoid hormone (in Dexamethasone) in this research could induced cell

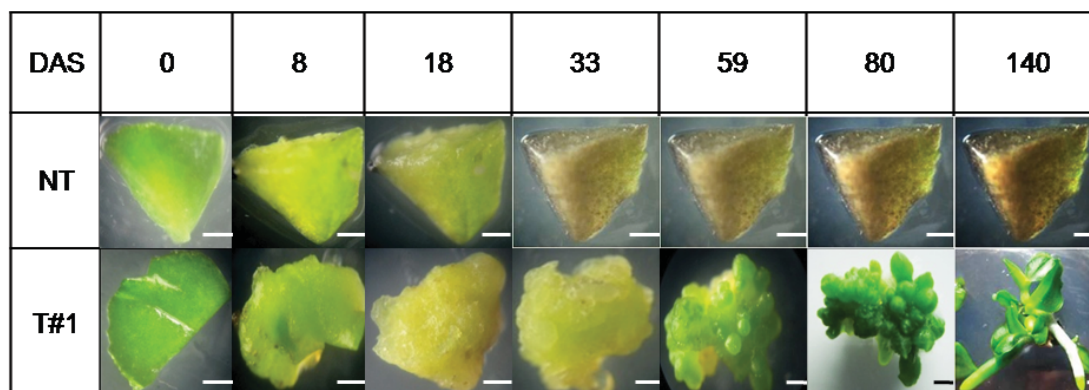


Figure 6. Formation of somatic embryos from leaf explants of *P. "Sogo Vivien"* T#1 transformant after induction by dexamethasone (Dex). Somatic embryos were induced at 8 DAS by 15 μ M Dex that showed enlarged size of leaf explants, followed by the formation of embryo at 18 DAS. Pre-globular embryos were observed at 33 DAS, followed by the formation of propagules at 59 DAS, leaf primordia were emerged from the propagule produced shoots at 80 DAS. Finally, each shoot continued their growth into plantlet at 140 DAS. On the other hand, the growth of somatic embryo was not observed from non-transformant (NT) leaf explants. Bars: 1 mm.

Table 2. The growth of somatic embryo from leaf explants of T#1 transformants after dexamethasone (Dex) induction

Experiment	No. of explants examined	No. of explants responded	the occurrence of somatic embryo (days)	No. of somatic embryo
1	5	3	18	187
2	5	2	17	81
3	5	3	20	145
Total	15	8		413

proliferation leaf pieces, that could grow into whole plants after subcultured on Dex-free medium, indicates that undergone each cell proliferation will form a structure resembling embryo and will further develop into plantlet. This is in accordance to the result of Schena *et al.* (1991) that nanomolar concentrations of glucocorticoids are sufficient to induce gene expression more than 150-fold, without causing detectable alterations in the physiology of the cultured plant cells.

The advantages of glucocorticoid-inducible expression system are non-toxic, have no physiological negative effect, allowing the induction of the target genes without pleiotropic effect, induce the target gene rapidly (glucocorticoid can easily permeate plant cell), no visible damage to leaves if glucocorticoid is accumulated, very

flexible because it possible to develop another steroid induction system in combination with the GVG system, widely applicable to many genes and in different species of transgenic plants (Aoyama and Chua, 1997). The use of the induction system with glucocorticoid is highly efficient, when compared to the other methods. In transgenic technology of plants, generally induction of foreign gene expression was conducted by providing strong promoter(s) such as the Cauliflower Mosaic Virus-35S promoter (Semiarti *et al.*, 2007), regulatory protein Gal4 from *Saccharomyces cerevisiae* (Liu and Lehman, 2008), or the herpes simplex virus VP16 as a regulatory switch (Wysocka and Herr, 2003) that positioned upstream of the transcription unit of the foreign gene to be regulated, so as to force the foreign gene can be expressed. But, it could not in this case,

because of the constitutive promoters always active continuously, therefore we could not set when the gene to be expressed, whereas using glucocorticoid induction system it could be determined when a foreign gene to be expressed. This will give great benefit to researchers who want to know the expression and function of the foreign gene in a specific region and period during transgenic plants development.

Conclusion

Somatic embryos of *Phalaenopsis* "Sogo Vivien" can be induced by overexpression of *AtRKD4* gene of *Arabidopsis thaliana* through *Agrobacterium*-mediated genetic transformation method. Insertion of T-DNA carrying *35S::GAL::AtRKD4::GR* construct into 21 day old protocorms through *A.tumefaciens* strain EHA 105 produced 17 transformant plants, the efficiency of transformation was 0.63%. A total 413 plants were regenerated from leaf pieces of Orchid Transformant line #1 after 1 hour induced by 15 μ M Dex, then maintained on Dex-free NP medium. All somatic embryos were able to regenerate into normal phenotype of plantlets, which were morphologically similar to the mother plants.

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