

## Induction of Hairy roots of Pegagan (*Centella Asiatica* (L.) Urban using Several Explant Sources with Several *Agrobacterium Rhizogenes* Strains in Vitro

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**Abstract**— Induction of hairy roots of pegagan using *A. rhizogenes* strains is important to obtain high production of triterpenoid as secondary metabolite. The objectives of this research were to obtain the best *A. rhizogenes* strains and the best explant sources for *C. asiatica* hairy root induction. The research consisted of two stages. Stage I was *A. rhizogenes* arranged in Completely Randomized Design (CRD) with 3 treatments and 6 replications. As a treatment was inoculation with *A. rhizogenes* strains LBA 9457, A4, R1000 and control. Stage II was explant sources trial arranged in CRD with 3 treatments and 6 replications. The treatments were: internodus, apical bud and leaves. The results showed that all three strains of *A. rhizogenes* were able to induce *C. asiatica* plant to produce hairy roots, with R1000 as the best strain. The best explant source was explant leave with 15 day after inoculation and the percentage of *C. asiatica* hairyroot formation was 100 %. The  $\beta$ -glucuronidase (GUS) test result has proven that Ri plasmid T-DNA of *A. rhizogenes* was integrated into the genome of *C. asiatica* plant.

**Keywords**— *Agrobacterium rhizogenes*, *Centella asiatica* (L.) Urban, hairy root.

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### INTRODUCTION

Pegagan (*Centella asiatica* (L.) Urban) belongs to the family of Umbelliferae. This plant produces secondary metabolites that have the potential to be developed into drugs of various diseases [1]. Various types of compounds contained in *C. asiatica* are asiaticoside and madecassic acid which are active ingredient capable of stimulating the production of collagen and protein biosynthesis helpful as a driver for the process of wound healing in humans [2].

This plant contains triterpenoid, secondary metabolites, mainly asiaticoside, asiatic acid and madecassic acid, used to treat bone loss for the olds, increase intelligence in children, youth drug, medicinal skin diseases, for cosmetics, as a potent anticancer drug for ovarian, improve nerve and circulatory disorders, as raw material for drugs useful as anti-dementia and anti-stress [3].

The need for raw materials used as ingredients is increasing along with the increasing need for medication. On the other hand, producing secondary metabolites naturally is still very limited. Secondary metabolites production for commercial sources is effectively done by using plant extracts from field. The problem is often the availability of

secondary metabolites is low. In addition, the content of secondary metabolites is also influenced by the natural conditions such as geographical factors, climate and pests and diseases [4].

One of the technique to induce higher secondary metabolites production is via in vitro culture [5]. In vitro multiplication is an alternative to get the plants in large quantities [6]. Plant tissue culture is a technique to develop the plant, either in the form of organ, tissues and cells under aseptically in vitro conditions. This technique is based on the totipotensi theory cells in aseptically conditions, the use of artificial culture media with a complete nutrient content and plant growth regulators [7].

Hairy root cultures as a form of tissue culture techniques are the result of transformation using *Agrobacterium rhizogenes* genetically. Hairy root cultures provide many benefits, among others are having biosynthesis ability as in normal plant roots with the growth rate is much faster than normal roots or equal to the cell suspension cultures [8], and is able to express the formation of metabolite products stable over the period of culture [9]. In some plants, the production of secondary metabolites was higher in hairy roots compared to normal roots of plants or the ones produced by origin plant [10]

## RESULTS AND DISCUSSION

### A. Test of Some Strains of *A. rhizogenes* on production of hairy roots of *C. asiatica*

The results of testing several strains of *A. rhizogenes* in producing hairy roots of *C. asiatica* plants was performed in Table 1.

TABLE I  
EFFECT OF SEVERAL AGROBACTERIUM RHIZOGENES STRAINS TO HAIRYROOT. *C. ASIATICA* 30 DAYS AFTER INOCULATION (DAI)

No	Treatment	Initiation time (DAI)	Living explants (%)	Hairy roots emergence (%)
1	Control	-	-	-
2	Strain LBA 9457	24 a	33,33 c	33,33 c
3	Strain A <sub>4</sub>	17 b	50 b	50 b
4	Strain R1000	15 b	100 a	100 a

Description: DAI = days after inoculation. The numbers in a column followed by the same lowercase letters are not different according to DNMRT 5%.

Results of *A. rhizogenes* inoculation on leaf explants of *C. asiatica* plants after cultures were 30 days old showed that strain R1000, A4 and LBA 9457 were able to induce the formation of hairy roots. Hairy root initiation time, the percentage of living explants and percentage of emergence of hairy root in plants of *C. asiatica*, transformed with R1000 strain, strain LBA 9457 and A4 strain of *A. rhizogenes* bacteria were presented in Table 1. It showed that LBA 9457, A4 and R1000 strains could induce the formation of hairy roots, with different initiation time. The fastest initiation time was shown in strain R1000. Explants inoculated with *A. rhizogenes* R1000 strain of bacteria, hairy roots began to form 15 DAI, while those inoculated with A4 (17 DAI) and LBA 9457 strains, hairy roots began to form 24 DAI. Differences in initiation time of *C. asiatica* hairy roots on each of the *A. rhizogenes* strain due to differences in genes on the Ri plasmid entering into the plant chromosome of *C. asiatica*.

Result of explant inoculation showed that only strain R1000 was able to induce 100% formation of hairy roots of the total replications. The successful formation of hairy roots from inoculated explants was an indication of the transferred of T-DNA from *A. rhizogenes* Ri plasmid into the genome of cells of *C. asiatica*. *C. asiatica* plant is susceptible to infection of *A. rhizogenes*. The three strains used showed the different responses to the infection of *A. rhizogenes*. The highest percentage of hairy roots emergence was obtained on treatment inoculated with *A. rhizogenes*, R1000 strain (100%), in contrast to the control treatment in which hairy roots did not emerge all. Inoculation with A4 strain treatment caused 50% hairy roots emergence and with LBA 9457 was 33.33%. The growth of hairy roots formed from explants of *C. asiatica* was very different both in number and percentage of emergence of hairy roots on each strain tested.

Plant responses to the induction of T-DNA from *A. rhizogenes* were different. Some formed callus first, some forms the hairy root and others remained alive without forming hairy roots, and some even had no response at all (dead). According to [16], in several species of plants, hairy

Several factors are known to affect the success and efficiency of transformation via *Agrobacterium* including tissue culture conditions and co-cultivation conditions, the suitability of the bacterial strains and selection of plant tissue as starting materials (explants source) [11]. From a research conducted on two types of quinine (*Cinchona ledgeriana* and *C. succirubra*), out of 10 strains tested, only *A. rhizogenes* strain LBA 9457 were able to induce hairy roots of both types of quinine. Furthermore, from the two types of quinine it is known that the explants derived from apical buds and leaves in vitro is the best source of explants to form hairy roots [12]. Explant source selection is very influential for the success of the hairy root culture. Research conducted by [13] on plants of Patchouli (*Pogostemon cablin* Benth) showed the best explant to induce hairy root is from the leaves. The results of a study done by [14], on *Ophiorrhiza pumila* plant, indicated that the best explants to induce hairy roots derived from the buds of shoots.

A research was conducted entitled, "Induction of Hairy Root of *Centella asiatica* (L.) Urban Using Several Explant Source With *Agrobacterium rhizogenes* In Vitro". The objectives of the research were to obtain the best strains of *A. rhizogenes* and the best explant source to induce hairy roots of *C. asiatica*.

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### B. Working Procedure

*A. rhizogenes* strain R1000, LBA 9457 and A4, were maintained on agar slant YMB (Yeast Mannitol Broth) before inoculation. Inoculation was carried out following the methods as listed in Noli [12]. Explant of *C. asiatica* used to induce hairy roots derived from one week old cultured explants in vitro.

For bacterial decontamination, subcultures were performed on the same medium by adding Cefotaxin 100 mg/L. Subcultures were repeated 3 times until they were free of bacteria. Furthermore, explants were planted again on solid medium without cefotaxin.

A trial of  $\beta$ -glucuronidase (GUS) was performed to confirm the integration of T-DNA of Ri plasmid into the plant genome of *C. asiatica*. Working procedures followed methods of Jefferson [15].

roots emerged directly at the site of inoculation, but in certain species initially appear on the inoculant callus which eventually form hairy root and hairy roots in normal circumstances would appear in the range of 1 to 4 weeks after inoculation.

Leaf explants of *C. asiatica* plants inoculated with some strains of *A. rhizogenes* also showed the formation of hairy root response (Figure 1) and the initial response of the explants was hairy root formation and explants that did not form the hairy roots (Figure 1). The growth of the hairy roots of *C. asiatica* transformed with each strain of *A. rhizogenes* was very different, especially in the number of hairy roots emerged (Figure 1).

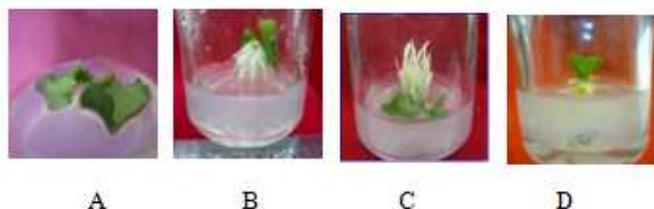


Figure 1. Response of *C. asiatica* leaf explants inoculated with some strains of *A. rhizogenes*

A. Control; B. strain A4; C. strain R1000; D. strain LBA 9457

#### B. Test of different sources of explants of *C. asiatica* plants transformed with *A. rhizogenes* bacteria, R1000 strain

The results of tests on several sources of explants producing the hairy roots of *C. asiatica* transformed with *A. rhizogenes* were presented in Table 2.

TABLE II  
EFFECT SEVERAL EKSPANS SOURCES WITH *A. RHIZOGENES* R. 1000 TO HAIRY ROOT

Explant source	Average time of initiation (DAI)	Average growth response		
		Living explants (%)	Hairy roots (%)	Number of Roots
1. Leaves	15 b	100	100 a	27 a
2. Stolon	24 a	100	77,7 b	16,7 b
3. Apical shoots	19,6 a	100	66,7 b	14 b

Remarks: The figures are marked with the same lowercase letters are not significantly different according to DNMR5 5%. DAI = days after

The difference in initiation time due to differences in the response of the plant taken as a source of explants. Explants derived from the leaves need a shorter time for initiation compared to explants derived from the stolon and apical buds. According to [17], the success factors of hairy root culture was on selection of plant tissue used as sources of explant.

The percentage of explants living in all parts of the plantlets used either leaves, apical buds and internodus showed insignificant results. Source of explants showed significant difference in the percentage of living explants. All explant sources can be used to form hairy roots, because after explants with *A. rhizogenes* transformation was

initiated, all explants sources showed growth responses, good survival and the ability to form hairy roots. Percentage of living explants for all explants sources was 100%.

Growth response of *C. asiatica* transformed with several strains of *A. rhizogenes* was shown by the emergence of hairy root. The highest response (100%) was on leaf explants which was significantly different from the apical bud and stolon explants. The difference in the emergence of the hairy root might be possible because not all parts of the plantlets were suitable with bacterial isolates, so that not all bacterial T-DNA were integrated into the genome of the plants to form hairy roots. [18] stated that in higher plants there is a fairly large DNA sequences that do not encode active proteins. If the position of the T-DNA inserted in the genome was not random and only integrated plant in the DNA that are not active, then it can not be expressed by the plant.

The difference on the percentage of emergence of the hairy roots was also possible because of differences in age and physiological condition of explants used. Selection of the tissues as the initial material used was one of the most important factors affecting the success and efficiency of transformation. It depends on the age and physiological condition of the tissue. Differences in the ability of explant sources to form hairy roots after transformation with *A. rhizogenes* also affected the number of hairy roots formed. The average number of hairy roots was 16.7 from stolon explants which was insignificantly different from apical bud explants, 14 but significantly different from the leaf explants, 27. Results of some studies showed differences in the number of hairy roots produced on explants transformed with *A. rhizogenes* were influenced by the type of bacterial strains inoculating explants.

The highest number of hairy roots formed was on R1000 strain of *A. rhizogenes* treatment with leaf explants was 27 hairy roots and followed by stolon and shoot apical (Figure 2). Based on the average number of hairy roots, hairy root emergence, percentage of living explants and the average of initiation time, it could be stated that inoculation using R1000 strains showed the best results and the best source was leaf explants.



Figure 2. Response of some explants of *C. asiatica* inoculated with R1000 strain ( A. Stolon, B. apical buds, leaf explants of *C. asiatica*)

Explant competence is determined by several factors, such as species or genotype explant origin, type of organ used as explants, the level of organ development, even history of each explant used.

### C. Integration of Plasmid Ri T-DNA into the genome of plants of *C. asiatica* (L.) Urban

GUS test was performed to determine the integration of Ri T-DNA plasmid R1000 strain of *A. rhizogenes* bacteria into the genome of *C. asiatica* plants. In the hairy roots of *C. asiatica* which had been soaked in a solution of X-gluc for 24 h showed the formation of blue spots (Figure 3) in the root cells. This proved that the T-DNA of plasmid was integrated into the plant cell. [13] suggested that the gene *w* Gus gene was gene reporter that encoded the synthesis of  $\beta$ -glucuronidase, an enzyme that could hydrolyze a colorless substrate 5-bromo-3 indolyl 4 chloro glucuronide (X-gluc) became blue. [19] stated that the expression of  $\beta$ -glucuronidase gene activity (GUS) in transformed plant cells of the T-DNA could be used as an indicator of T-DNA integration into the plant genome. Expression of the cell strains of *Agrobacterium* infection is shown as blue spots by giving 5-bromo 4-3-indolyl cloro glucuronic acid (X-gluc). [20] explained that the expression of transgenic cells would be clear when analyzed using GUS parameters and could also describe the expression and function of the tissues of the transgene.



Fig. 3 Cross-section of hairy roots transformed with *A. rhizogenes* with blue color (designated by arrows).

Description: The blue color is the site where the DNA is integrated (as indicated by arrow). From the test results of GUS in the hairy roots it was showed that there was a blue color that covered the whole root or root portion only, or even not visible at all. This was possible because the transformation occurred as a whole (cellular) or partially (chimera). [13] research also found the presence of hairy roots after being tested with GUS but blue spots was not formed. The GUS test was called positive if blue spots were formed on the tissues. The thicker and bluer, the higher the GUS gene expression. Other cause of invisible blue spots was possibly because the hairy roots was taken randomly, so it was possible that the hairy roots cut in GUS test was not part of the tissues of integrated with T-DNA.

### CONCLUSIONS

The best strains of *A. rhizogenes* to induce hairy roots of *C. asiatica* was R1000 strain. Leaf explants was the best source of explants in inducing hairy roots.

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4	Strain R1000	15 b	100 a	100 a

Description: DAI = days after inoculation. The numbers in a column followed by the same lowercase letters are not different according to DNMRT 5%.

Results of *A. rhizogenes* inoculation on leaf explants of *C. asiatica* plants after cultures were 30 days old showed that strain R1000, A4 and LBA 9457 were able to induce the formation of hairy roots. Hairy root initiation time, the percentage of living explants and percentage of emergence of hairy root in plants of *C. asiatica*, transformed with R1000 strain, strain LBA 9457 and A4 strain of *A. rhizogenes* bacteria were presented in Table 1. It showed that LBA 9457, A4 and R1000 strains could induce the formation of hairy roots, with different initiation time. The fastest initiation time was shown in strain R1000. Explants inoculated with *A. rhizogenes* R1000 strain of bacteria, hairy roots began to form 15 DAI, while those inoculated with A4 (17 DAI) and LBA 9457 strains, hairy roots began to form 24 DAI. Differences in initiation time of *C. asiatica* hairy roots on each of the *A. rhizogenes* strain due to differences in genes on the Ri plasmid entering into the plant chromosome of *C. asiatica*.

Result of explant inoculation showed that only strain R1000 was able to induce 100% formation of hairy roots of the total replications. The successful formation of hairy roots from inoculated explants was an indication of the transferred of T-DNA from *A. rhizogenes* Ri plasmid into the genome of cells of *C. asiatica*. *C. asiatica* plant is susceptible to infection of *A. rhizogenes*. The three strains used showed the different responses to the infection of *A. rhizogenes*. The highest percentage of hairy roots emergence was obtained on treatment inoculated with *A. rhizogenes*, R1000 strain (100%), in contrast to the control treatment in which hairy roots did not emerge all. Inoculation with A4 strain treatment caused 50% hairy roots emergence and with LBA 9457 was 33.33%. The growth of hairy roots formed from explants of *C. asiatica* was very different both in number and percentage of emergence of hairy roots on each strain tested.

Plant responses to the induction of T-DNA from *A. rhizogenes* were different. Some formed callus first, some forms the hairy root and others remained alive without forming hairy roots, and some even had no response at all (dead). According to [16], in several species of plants, hairy

roots emerged directly at the site of inoculation, but in certain species initially appear on the inoculant callus which eventually form hairy root and hairy roots in normal circumstances would appear in the range of 1 to 4 weeks after inoculation.

4 Leaf explants of *C. asiatica* plants inoculated with some strains of *A. rhizogenes* also showed the formation of hairy root response (Figure 1) and the initial response of the explants was hairy root formation. 46 explants that did not form the hairy roots (Figure 1). The growth of the hairy roots of *C. asiatica* transformed with each strain of *A. rhizogenes* was very different, especially in the number of hairy roots emerged (Figure 1).

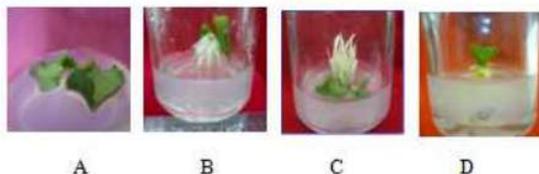


Figure 1. Response of *C. asiatica* leaf explants inoculated with some strains of *A. rhizogenes*  
A. Control; B. strain A4; C. strain R1000; D. strain LBA 9457

#### B. Test of different sources of explants of *C. asiatica* plants transformed with *A. rhizogenes* bacteria, R1000 strain

The results of tests on several sources of explants producing the hairy roots of *C. asiatica* transformed with *A. rhizogenes* were presented in Table 2.

TABLE II  
EFFECT SEVERAL EKSPANS SOURCES WITH *A. RHIZOGENES* R. 1000 TO HAIRY ROOT

Explant source	Average time of initiation (DAI)	Average growth response		
		Living explants (%)	Hairy roots (%)	Number of Roots
1. Leaves	15 b	100	100 a	27 a
2. Stolon	24 a	100	77,7 b	16,7 b
3. Apical shoots	19,6 a	100 5	66,7 b	14 b

Remarks: The figures are marked with the same lowercase letters are not significantly different according to DNMR 5%. DAI = days after

The difference in initiation time due to differences in the response of the plant taken as a source of explants. Explants derived from the leaves need a shorter time for initiation compared to explants derived from the stolon and apical buds. According to [17], the success factors of hairy root culture was on selection of plant tissue used as sources of explant.

The percentage of explants living in all parts of the plantlets used either leaves, apical buds and internodus showed insignificant results. Source of explants showed significant difference in the percentage of living explants. All explant sources can be used to form hairy roots, because after explants with *A. rhizogenes* transformation was

initiated, all explants sources showed growth responses, good survival and the ability to form hairy roots. Percentage of living explants for all explants sources was 100%.

4 Growth response of *C. asiatica* transformed with several strains of *A. rhizogenes* was shown by the emergence of hairy root. The highest response (100%) was on leaf explants which was significantly different from the apical bud and stolon explants. The difference in the emergence of the hairy root might be possible because not all parts of the plantlets were suitable with bacterial isolates, so that not all bacterial T-DNA were integrated into the genome of the plants to form hairy roots. [18] stated that in higher plants there is a fairly large DNA sequences that do not encode active proteins. If the position of the T-DNA inserted in the genome was not random and only integrated plant in the DNA that are not active, then it can not be expressed by the plant.

The difference on the percentage of emergence of the hairy roots was also possible because of differences in age and physiological condition of explants used. Selection of the tissues as the initial material used was one of the most important factors affecting the success and efficiency of transformation. It depends on the age and physiological condition of the tissue. Differences in the ability of explant sources to form hairy roots after transformation with *A. rhizogenes* also affected the number of hairy roots formed. The average number of hairy roots was 16.7 from stolon explants which was insignificantly different from apical bud explants, 14 but significantly different from the leaf explants, 27. Results of some studies showed differences in the number of hairy roots produced on explants transformed with *A. rhizogenes* were influenced by the type of bacterial strains inoculating explants.

The highest number of hairy roots formed was on R1000 strain of *A. rhizogenes* treatment with leaf explants was 27 hairy roots and followed by stolon and shoot apical (Figure 2). Based on the average number of hairy roots, hairy root emergence, percentage of living explants and the average of initiation time, it could be stated that inoculation using R1000 strains showed the best results and the best source was leaf explants.



Figure 2. Response of some explants of *C. asiatica* inoculated with R1000 strain (A. Stolon, B. apical buds, leaf explants of *C. asiatica*)

Explant competence is determined by several factors, such as species or genotype explant origin, type of organ used as explants, the level of organ development, even history of each explant used.

**C. Integration of Plasmid Ri T-DNA into the genome of plants of *C. asiatica* (L.) Urban**

GUS test was performed to determine the integration of Ri T-DNA plasmid R1000 strain of *A. rhizogenes* bacteria into the genome of *C. asiatica* plants. In the hairy roots of *C. asiatica* which had been soaked in a solution of X-gluc for 24 h showed the formation of blue spots (Figure 3) in the root cells. This proved that the T-DNA of plasmid was integrated into the plant cell. [13] suggested that the gene w Gus gene was gene reporter that encoded the synthesis of  $\beta$ -glucuronidase, an enzyme that could hydrolyze a colorless substrate 5-bromo-3 indolyl 4 chloro glucuronide (X-gluc) became blue. [19] stated that the expression of  $\beta$ -glucuronidase gene activity (GUS) in transformed plant cells of the T-DNA could be used as an indicator of T-DNA integration into the plant genome. Expression of the cell strains of *Agrobacterium* infection is shown as blue spots by giving 5-bromo 4-3-indolyl cloro glucuronic acid (X-gluc). [20] explained that the expression of transgenic cells would be clear when analyzed using GUS parameters and could also describe the expression and function of the tissues of the transgene.



Fig. 3 Cross-section of hairy roots transformed with *A. rhizogenes* with blue color (designated by arrows).

Description: The blue color is the site where the DNA is integrated (as indicated by arrow). From the test results of GUS in the hairy roots it was showed that there was a blue color that covered the whole root or root portion only, or even not visible at all. This was possible because the transformation occurred as a whole (cellular) or partially (chimera). [13] research also found the presence of hairy roots after being tested with GUS but blue spots was not formed. The GUS test was called positive if blue spots were formed on the tissues. The thicker and bluer, the higher the GUS gene expression. Other cause of invisible blue spots was possibly because the hairy roots was taken randomly, so it was possible that the hairy roots cut in GUS test was not part of the tissues of integrated with T-DNA.

**CONCLUSIONS**

The best strains of *A. rhizogenes* to induce hairy roots of *C. asiatica* was R1000 strain. Leaf explants was the best source of explants in inducing hairy roots.

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