Technology Transfer of Chrysanthemum Cultivation in Disaster Area of Mount Merapi To Improve People Revenue

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DOI

(Received: January 12, 2016; Accepted: March 17, 2016)

ABSTRACT

Chrysanthemum flowers are a main income source of surrounding communities of Hargobinangun. Hargobinangun became the main cut flower's supplier to the province of Yogyakarta before the Merapi's eruption. However, due to the shortage of cut flower production, currently those have to be imported from other areas such as Cipanas, Pasuruan and Malang. One of the production's obstacles is the environmental damage. Soil in planting area was covered by thick sand and volcanic ash from Mount Merapi eruption. Research and transfer of technology has been carried out by R & D team of Universities Pembangunan National "Veteran" Yogyakarta (UPNVY). The technologies of amelioran's addition to the soil in the disaster areas was using simple techniques, inexpensive and available materials from surrounding farmers site, such as manure, bamboo leaf litter and garden's compost. The materials were initially fermented by using PGPR that were produced by R & D UPNVY. In addition, application of horticultural mineral oil (HMO) was performed to avoid white rust infection. The results showed the increase of soil fertility and plant growth significantly which is expressed by the increase of plant height, stem diameter, and leaf area after application of ameliorant. The quality of cut flowers also increased in the form of the increase of flower diameter, and numbers of ribbon flower. The application at the farm level showed that farmers have no difficulty in using Enkas, although it is still necessary to conduct further training and mentoring to increase the success rate.

Keywords: Chrysanthemum, Environmental Damage, Ameliorant, HMO, Enkas.

INTRODUCTION

Merapi eruption in 2010 brought tremendous impact on environment, socioeconomic, and agriculture. One of the affected areas was Pakem district. It was exposed by ash and sand directly village on the slopes of Merapi, which has the potential to be developed as center of ornamental plants is Hargobinangun, sub-district Pakem. Since 2005 the region has been considered to be the center of chrysanthemum flower cultivation of Yogyakarta Province, considering the altitude of the area (500-800 m above sea level) are eligible for the growth

of chrysanthemum. Currently, chrysanthemum cultivation has been carried out by more than 100 local farmers, which are grouped in 13 farmer groups to manage an area of 10,000 m² with a production capacity of 15,000 cut flowers per week³.

After the eruption of Merapi on November 5, 2010, Chrysanthemum cultivation in the Hargobinangun village becomes stagnant²⁴. Most farmers did not know what to do because the chrysanthemum planting area conditions were destroyed. This area is very close to the Mount Merapi. The planting area is within the 4-10 km from

the peak of Merapi. Post the eruption, planting area was covered by volcanic ash and sand, and could not be used for cultivating anymore.

Chrysanthemum in the form of cut flowers, produced by farmers in Wonokerso were much degraded, so that consumers were switching on chrysanthemum imported from other regions, such as West Java and East Java. In the field survey showed poor production quality was due to the plant roots were not well developed, brown coloured with short size. Volcanic material with a thickness of 5-15 cm that covers Wonokerso became compressed (compact), hard, waterresistant. Addition of ameliorant was needed to restore the crumb structure.24 reported that the volcanic material that was given ameliorant vermin compost and cow manure was good for dahlias plant growth in the Kinahrejo region. Basically dahlias and chrysanthemums almost the same, so as to improve the planting area of chrysanthemum can be used the same ameliorant.

Chrysanthemum white rust (CWR) can be a serious disease of chrysanthemum crops. According to¹⁰, white rust disease (Puccinia horiana P. Henn.) may decrease freshness of chrysanthemum flowers (vase-life) into only 5 days, significantly shorter than the healthy ones. Its freshness can last up to 12 days at room temperatures (27-29°C). Chrysanthemum yield loss caused by white rust disease is reaches 30% in Indonesia²¹, 80% in Turkey⁹, and 100% in New England⁶. Some insects reported as vector of some virus diseases, i.e. Aphis craccivora, Acyrthosiphon pisum, and Myzus persicae17, Macrosiphoniella sanborni, Rophalosiphum sp. (Aphididae)2 also attack leaves of chrysanthemum. Various pest and disease control measures has been done such as the use of tolerant varieties, culture technique (i.e. cutting infected leaves and setting watering), the use of natural enemies, and the application of synthetic pesticides. However, the intensity of pest and disease still high. Recently, mineral oils were found to be highly effective against citrus pest^{15,15}. Horticultural mineral oil (HMO) is highly refined mineral oils originated from crude petroleum oils. It is paraffinic compound (360% of carbon atoms occur in chains). It has un-sulfonated residue (UR) values 392 % (therefore it contains £8% aromatic molecules). Its molecule weight vary and is reflected in the number of carbon atoms. The lightest oils are oils, and the heaviest oils are generally $n{\rm C}_{25}$ oils. These values reflect the median equivalent n-paraffin carbon numbers and distillation temperatures 1 [4]. Several factors favor the use of horticultural mineral oil including low cost, low mammalian toxicity, and few deleterious environmental effects 7 . Thus the main problem being urgency (virtue) doing in this research is a complete study of various aspects of the chrysanthemum plant cultivation techniques that can improve growing conditions.

MATERIALS AND METHODS

The experiment was conducted in the village Wonokerso, Pakem, Sleman Yogyakarta with the complete randomized block design consists of two factors: material of ameliorant and HMO concentration. The first factor consisted of four levels: vermin compost, fern compost, cow manure and litter of bamboo plants. The second factor consisted of three levels: 0.125, 0.250 and 0.500% v/v of horticultural mineral oil (HMO: nC_{21} Sunspray Ultra Fine®, Amtrade Pty). As control chrysanthemums was grown in media without ameliorant and without HMO application. From these two factors each repeated three times and each block contains 50 plants with five sample plants, so that the overall number was 1800 crop plants plus 150 control plants.

Research was conducted in UV plastic roofed green house, facing east to form a semicircle dome roof. Land management was done as deep as 30 cm and it was mixed with ameliorant material according to treatment then beds were made 10-20 cm height. Chrysanthemum seedlings taken from Horticultural Research Institute at Cipanas, West Java were planted in beds that have been given the nets. Plants were treated for three months, which includes watering, and fertilizing. Watering the plants was conducted twice a day. Fertilization was done at the beginning of the study using 75 g N, 75 g P, and 25 g K fertilizer per plant and leaf fertilizer once a week.

Assessment was conducted fortnightly before oil spray application on five randomly chosen central plants within each plot on plant height, diameter of flower, number of ribbon flower, leaf area, stem diameter, aphid population, and the spread of

white rust disease. Visual observations determined the number of aphid per leaf on five chosen leaves on upper part of tree. The spread of white rust was assessed at same samples as aphid assessment by recording fortnightly the number of leaves infected based on the rating scale of severity (Table 1.)

Severity Rating Description

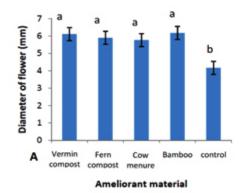
01234 Leaf without any symptom < 25% part of leaf showing symptom25% up to < 50% part of leaf showing symptom50% up to < 75% part of leaf showing symptome" 75% part of leaf showing symptom

Advanced research was conducted in the laboratory to test the ability of plant life to be propagated in vitro using Enkas and Laminair Air Flow cabinet.

Data were subjected to one-way ANOVA (analysis of variance). Duncan's multiple range test (DMRT) was used to determine the differences among treatments when the ANOVA was significant. Significance different was arise at P< 0.05. Analysis was performed using SPSS® version: 10.0.5¹⁹.

RESULTS AND DISCUSSION

Ameliorant material did not affect on diameter of flower (Fig. 1A.), the number of ribbon flowers (Fig. 1B.), plant height (Fig. 2 A.), and the diameter of stem (Fig. 2B.) and there was not any interaction between ameliorant and HMO but significantly higher all these characters than the control. Vermin compost was the best ameliorant material for increasing leaf area (Fig. 3). Diameter of flowers, ribbon flower number, plant height, stem



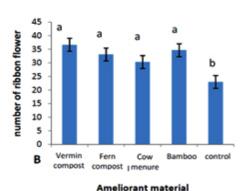
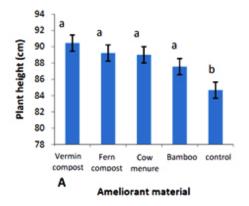


Fig. 1: Affect of various ameliorant materials on diameter of flower (A) and number of ribbon flower (B) of chrysanthemum. Bars with the same letter are not significantly different (DMRT, α 5%)



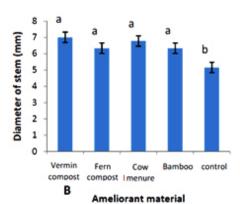
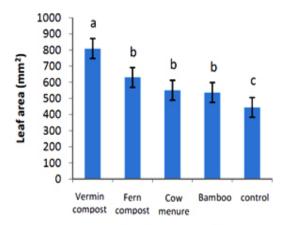


Fig. 2: Affect of various ameliorant materials on plant height (A) and diameter of stem (B) of chrysanthemum. Bars with the same letter are not significantly different (DMRT, α 5%)

diameter and leaf area on treatment of ameliorant were ranging from 5.76 to 6.18 cm, 30.33 to 36.67, 87.55 to 90.44 cm, 6, 33 to 7.00 mm and 550.60 to 808.90 mm², respectively. While in control, the flower diameter, number of ribbon flower, plant height and stem diameter were 4.17 mm, 23.00, 84.67 cm, and 5.15 mm respectively.

Leaf tissue anatomy in monocotyledon is composed of a set of cells that have almost the same shape. It is composed of upper and lower epidermis tissue, and mesophyll tissue, which is composed of palisade tissue and spongy tissue. The epidermis covers the upper and lower surfaces of leaves up to epidermal stem [16]. Mesophyll layer is the most important leaf part for photosynthesis.



Ameliorant material

Fig. 3: Affect of various ameliorant materials on leaf area of chrysanthemum. Bars with the same letter are not significantly different (DMRT, α 5%)

Table 1: Disease severity rating scale used to access the spread of white rust

Severity rating	Description
0	Leaf without any symptom
1	< 25% part of leaf showing
	symptom
2	25% up to < 50% part of leaf
	showing symptom
3	50% up to < 75% part of leaf
	showing symptom
4	≥75% part of leaf showing
	symptom

Most of palisade layer is composted by chloroplasts, and affects the products of photosynthesis. Damage by high temperatures that occurs in mesophyll, specifically in palisade, will give the highest impact on the photosynthesis activities. The common histological changes in leaf damage by high temperature are plasmolysis, granulation or disorganization of cells constituent, destruction or disintegration of cells and tissue pigmentation²².

¹³mentioned that pollutants are able to induce physiological damage in the plant long before of the occurrence of physical damage. Other experts said that it is a hidden damage. Hidden damage may be a decrease in the ability of plants to absorb water, slow cell growth or imperfect stomata opening. According to17, mechanism of stomata opening dependents on the changes of cover cells turgid. Cover cell containing a high concentration of starch will be closed, especially at night. When the sun is able to generate photosynthesis in chlorophyll, CO₂ level will decline, and it is reduced into CH₂O. This process is followed by the increase of pH and will increase the activity of posporilase enzymes to convert starch into glucose. The glucose formation will increase the osmosis level of cover cell, causing water influx from neighbouring cells. This condition causes stomata turgid and stomata will open.

There was no significant difference between oil concentration treatments (Table 2.). It seemed that oil film could not provide a barrier by masking the feeding and oviposition stimulants, hence preventing the aphid from locating, accepting

Table 2: Effect of oil concentration on aphid population and diseases severity of white rust on chrysanthemum leaf

Oil concentra	Disease	
(% v/v)	Population	severity (%)
0	24.11 ± 2.86 q	88.89 ± 2.47 a
0.125	23.89 ± 2.20 q	86.67 ± 2.36 a
0.250	21.67 ± 2.66 q	$69.44 \pm 2.27 b$
0.500	$18.00 \pm 2.25 \mathrm{q}$	$65.00 \pm 2.89 b$
Probability (P)	0.298	< 0.001

Numbers in columns followed by the same letter are not significantly different (DMRT, α 5%)

or using the host plant. It was not consistent with the report on the adult females of two spotted mite (*Tetranychus urticae* Koch [Acari: Tetranychidae])¹¹,

Asiatic citrus psylla (*Diaphorina citri* Kuwayama)¹⁴, whiteflies (*Bemesia argentifolii* Bellows and Perring [Hemiptera: Aleyrodidae])²⁰, greenhouse thrips

Table 3: The number of days for shoots growing process, the percentage growing process, shoot height, number of shoots and number of roots on Laminair Air Flow and Enkas at 30 days after sowing

Treatments						
M1		Shoots Growing	of the growing	Height		
(½MS + 1 ppm NAA + 0 ppm Kinetin) M2 9 a 60 c 4,6 b 3 b 6 b (½MS + 1 ppm NAA + 0,1 ppm Kinetin) M3 12 b 65 b 5,1 b 4 ab 6 b (½MS + 1ppm NAA + 0,2 ppm Kinetin) M4 12 b 85 a 6,7 a 5 a 11 a (½MS + 1 ppm NAA + 0,3 ppm Kinetin) M5 13 b 70 b 5,2 b 2 b 4 b (½MS + 1 ppm NAA + 0,4 ppm Kinetin) Enkas M1 15 c 50 d 2,9 b 2 b 3 b (½MS + 1 ppm NAA + 0,4 ppm Kinetin) M2 15 c 60 c 3,3 ab 2 b 3 b (½MS + 1 ppm NAA + 0,1 ppm Kinetin) M3 14 b 70 b 3,7 ab 3 a 9 a (½MS + 1 ppm NAA + 0,2 ppm Kinetin) M3 14 b 70 b 3,7 ab 3 a 9 a (½MS + 1 ppm NAA + 0,2 ppm Kinetin) M4 9 a 80 a 4,5 a 4 a 11 a (½MS + 1 ppm NAA + 0,3 ppm Kinetin) M4 9 a 80 a 4,5 a 4 a 11 a (½MS + 1 ppm NAA + 0,3 ppm Kinetin) M5 16 c 45 d 2,9 b 3 a 4 b	Laminair Air Flow					
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M2 9 a 60 c 4,6 b 3 b 6 b (½MS + 1 ppm NAA + 0,1 ppm Kinetin) M3 12 b 65 b 5,1 b 4 ab 6 b (½MS + 1 ppm NAA + 0,2 ppm Kinetin) M4 12 b 85 a 6,7 a 5 a 11 a (½MS + 1 ppm NAA + 0,3 ppm Kinetin) M5 13 b 70 b 5,2 b 2 b 4 b (½MS + 1 ppm NAA + 0,4 ppm Kinetin) Enkas M1 15 c 50 d 2,9 b 2 b 3 b (½MS + 1 ppm NAA + 0,1 ppm Kinetin) M2 15 c 60 c 3,3 ab 2 b 3 b (½MS + 1 ppm NAA + 0,1 ppm Kinetin) M3 14 b 70 b 3,7 ab 3 a 9 a (½MS + 1 ppm NAA + 0,2 ppm Kinetin) M4 9 a 80 a 4,5 a 4 a 11 a (½MS + 1 ppm NAA + 0,3 ppm Kinetin) M4 9 a 80 a 4,5 a 4 a 11 a (½MS + 1 ppm NAA + 0,3 ppm Kinetin) M5 16 c 45 d 2,9 b 3 a 4 b	(½MS + 1 ppm NAA + 0					
(½MS + 1 ppm NAA + 0,1 ppm Kinetin) M3	ppm Kinetin)					
ppm Kinetin) M3	M2	9 a	60 c	4,6 b	3 b	6 b
M3	(1/2MS + 1 ppm NAA + 0,1	I				
(½MS +1ppm NAA + 0,2 ppm Kinetin) M4 12 b 85 a 6,7 a 5 a 11 a (½MS +1 ppm NAA + 0,3 ppm Kinetin) M5 13 b 70 b 5,2 b 2 b 4 b (½MS +1 ppm NAA + 0,4 ppm Kinetin) Enkas 4 b 2 b 3 b M1 15 c 50 d 2,9 b 2 b 3 b (½MS + 1 ppm NAA + 0 ppm Kinetin) M2 15 c 60 c 3,3 ab 2 b 3 b (½MS + 1 ppm NAA + 0,1 ppm Kinetin) M3 14 b 70 b 3,7 ab 3 a 9 a (½MS +1 ppm NAA + 0,2 ppm Kinetin) W4 9 a 80 a 4,5 a 4 a 11 a (½MS +1 ppm NAA + 0,3 ppm Kinetin) M5 16 c 45 d 2,9 b 3 a 4 b (½MS +1 ppm NAA + 0,4 4 b 4 b 4 b 4 b 4 b	ppm Kinetin)					
ppm Kinetin) M4	M3	12 b	65 b	5,1 b	4 ab	6 b
M4 12 b 85 a 6,7 a 5 a 11 a (½MS +1 ppm NAA + 0,3 ppm Kinetin) M5 13 b 70 b 5,2 b 2 b 4 b (½MS +1 ppm NAA + 0,4 ppm Kinetin) Enkas M1 15 c 50 d 2,9 b 2 b 3 b (½MS + 1 ppm NAA + 0,1 ppm Kinetin) M2 15 c 60 c 3,3 ab 2 b 3 b (½MS + 1 ppm NAA + 0,1 ppm Kinetin) M3 14 b 70 b 3,7 ab 3 a 9 a (½MS +1 ppm NAA + 0,2 ppm Kinetin) M4 9 a 80 a 4,5 a 4 a 11 a (½MS +1 ppm NAA + 0,3 ppm Kinetin) M5 16 c 45 d 2,9 b 3 a 4 b (½MS +1 ppm NAA + 0,4						
(½MS +1 ppm NAA + 0,3 ppm Kinetin) M5 13 b 70 b 5,2 b 2 b 4 b (½MS +1 ppm NAA + 0,4 ppm Kinetin) Enkas M1 15 c 50 d 2,9 b 2 b 3 b (½MS + 1 ppm NAA + 0 ppm Kinetin) 60 c 3,3 ab 2 b 3 b (½MS + 1 ppm NAA + 0,1 ppm Kinetin) 70 b 3,7 ab 3 a 9 a (½MS + 1 ppm NAA + 0,2 ppm Kinetin) 9 a 80 a 4,5 a 4 a 11 a (½MS +1 ppm NAA + 0,3 ppm Kinetin) M5 16 c 45 d 2,9 b 3 a 4 b (½MS +1 ppm NAA + 0,4 16 c 45 d 2,9 b 3 a 4 b	ppm Kinetin)					
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(½MS +1 ppm NAA + 0,4 ppm Kinetin) Enkas M1	• •					
ppm Kinetin) Enkas M1			70 b	5,2 b	2 b	4 b
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ppm Kinetin) M2		15 C	50 a	2,9 b	2 D	3 D
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(½MS + 1 ppm NAA + 0,1 ppm Kinetin) M3		15.0	60.0	3 3 ah	2 h	3 h
ppm Kinetin) M3			00 0	5,5 ab	20	3.0
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M5 16 c 45 d 2,9 b 3 a 4 b (½MS +1 ppm NAA + 0,4						
(½MS +1 ppm NAA + 0,4	• • • • • • • • • • • • • • • • • • • •	16 c	45 d	2,9 b	3 a	4 b
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The figures followed by the same letter in the columns and rows are not significantly different based on DMRT Test (α = 5%). The (-) sign indicates no significance between LAF and Enkas.

(*Heliothrips haemorrhoidalis* Bouche [Thysanoptera: Thripidae])¹². They do not lay their eggs on plant treated with oils. Density of damage spots caused by citrus red mite (*Panonychus citri* McGregor [Acari: Tetranychidae]) feeding activity was reduced significantly on plant treated with these oils⁵. Greenhouse thrips preferred untreated fruit to HMO-treated fruit as feeding site ¹¹.

Oil application in concentrations of 0.250 and 0.500% were able to decrease the spread of white rust with the severity of 69.44 and 65.00 respectively (Table 2.). Oil might be caused deformation of aprezoria and affected on uredospores germination to infect chrysanthemum plants. Similar result has been reported by 18 on wheat rust (*Puccinia recondite* f. sp. *Tritici*.). Oils could also provide a mechanical barrier to prevent the invasion of uredospores germ tube²². The success of oil in suppressing plant disease was also achieved on powdery mildew⁷.

Significant suppression of aphid would likely require either higher concentration of oil or more frequent application of oil then were used in this study to form appropriate oil layer density on leaves surfaces. The level of suppression related to the number of applications [7]. It should be thick enough to prevent the emitting of leaves volatile

Wonokerso area is used as a center for the cultivation of chrysanthemum. This will be a "pilot project" or as a model for other areas to cultivate chrysanthemum. Observations in the field indicated that the closure of volcanic material with a thickness of 5-15 cm in the ground can cause delays in the entry of air into the soil. This can lead to land

degradation as a habitat for flora and fauna that can support the growth of cultivated plants as a source of farmer's revenue.

Advanced research was conducted in the laboratory to test the ability of plant life to be propagated in vitro using Enkas and Laminair Air Flow cabinet. Observations during the incubation period since the explants are planted until the end of the study showed various changes in the explant. These changes include the shoots growing process, the percentage of the growing process, shoots height, total number of shoots, and total number of roots.

The initial response that occurred in the majority of explants after implantation is the tissue swelling of explants. According⁸, changes in the explant are linked to the level of osmolarity of the medium used.²⁴, stated that the tissue swelling is due to the effect of the addition of nutrients and growth regulators.

The results showed that the sowing explants implemented on Enkas and Laminair Air Flow (LAF) was not significantly different for all important parameters of callus growth observed, as presented in table 3. This means that the Enkas equipment, although it is pretty simple, easy to use, and cheap, it can create products that are equally good compared to Laminair Air Flow (LAF) that looks more complicated and expensive.

Table 3 shows that during the shoots growing process, the treatments between Enkas and LAF had no significant difference. Meanwhile,



Fig. 4: Plantlets produced using Enkas (left) and LAF (right); at 60 days after sowing.

between the four kinds of media used, shoots from M1 medium emerged faster than the other media. The initial response that occurred in the majority of explants planted is the tissue swelling of explants. According to²¹, changes in the explant are linked to the level of osmolarity of the medium used. Media culture is one of the critical success factors of plant propagation in vitro. In tissue culture media, the addition of rapid growth regulator substances is required to support the growth of explants. This is in line with the opinion of8, which stated that the higher the concentration of cytokines, the longer the shoots will grow. It also affects the percentage of the shoots growing process, in which the influence of kinetin to the cell is its role in stimulating the process of cell division and cell differentiation. The main influence is in DNA replication that will affect the growth of shoots. The planting medium used also determines the growth. M3 medium (MS + 2 ppm NAA + 0.3 ppm Kinetin) was able to spur the growth of shoots to make them taller and in great quantities, as well as the total number of roots. Plantlet rooting is important for further growth due to the growing number of roots, which are good for the media to absorb nutrients. This is because the more number of roots produced, the vaster the field of nutrients absorption from the roots media becomes. The number of root crops will be useful for acclimatization in a glass house. Therefore the increasing number of roots will also improve the field of nutrients absorption²³.

The application at the farm level showed that farmers have no difficulty in using enkas, although

it is still necessary to conduct further training and mentoring to increase the success rate. The use of Enkas and LAF was able to provide results that are equally good for the in vitro development of the potato, although the need of advanced skills is still necessary to make it more flexible in its use.

CONCLUSION

The addition of ameliorant increased soil fertility, plant growth, and cut flower production. Plant growth in the form of plant height, stem diameter, and leaf area was higher than the control. Cut flower production in the form of flower diameter and number of ribbon flower was also better than the control. Oil application on concentration of 0.250 and 0.500% was able to suppress the severity of white rust on chrysanthemum. Higher concentration or frequency was required to achieve significant control on aphid population. The application at the farm level showed that farmers have no difficulty in using enkas, although it is still necessary to conduct further training and mentoring to increase the success rate.

ACKNOWLEDGEMENT

Our gratitude goes to LPDP-RISPRO who has funded this research through grants implementation.

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