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Isolation and identification molecular of endophytic fungi isolated from native tropical orchids in Indonesia

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Abstract

*Tricodherma* mycorrhiza has an important role as a biocontrol agent. Its association with *Phalaenopsis amabilis* was molecularly identified through rDNA-ITS sequence analysis. The aims of the study were to identify isolated molecular of orchids mycorrhiza from native tropical orchids in Indonesia, conducted as one of native orchid conservation efforts in Indonesia. One isolate of *Tricodherma* were isolated from the root of orchid plant in Yogyakarta based on morphological and microscopical analysis. Verification analysis molecular of these isolates resulted in 600-750 bp DNA products located on the ITS1-5.8S-ITS4 region. The sequenced products showed insertion and substitution occurances, which may result in strain diversity and possible variation in severity. Reconstruction of phylogenetic trees using *Maximum Parsimony* and *Bootstrap*-1000 approach showed that Indonesian isolates have undergone speciation and have been positioned in the cluster, which are already far apart from the other isolates. Isolate *Tricodherma* from Yogyakarta, Indonesia successfully isolated based on identification of rDNA-ITS sequences. Results of this study were expected to become the basic information in an effort of native orchid cultivation and protection against infectious diseases in Indonesia. The study was the first to report regarding *Tricodherma* isolated from native tropical orchids in Indonesia.

*Keywords:* *Tricodherma*; rDNA-ITS; *Phalaenopsis*; Indonesia

1. Introduction

*Phalaenopsis* is a genus of orchids, which some of its members have important role as parent crosses. The presence in its native habitat (nature) has been reported to have greatly diminished. Thus, Infections by bacteria, fungi, and viruses are still major obstacles in conducting cultivation and development of native orchids in Indonesia (Kumalawati et al., 2011; Mahfut and Daryono, 2014; Mahfut et al., 2016a; Mahfut et al., 2016b; Mahfut et al., 2017). Cultivating and protecting *Phalaenopsis* sp. against diseases in Indonesia could be done through the induction of endophytic microorganisms (Tanawy, 2009), which includes the Orchid Mycorrhizal Fungi (OMF). One type of OMF which have been isolated and identified is *Tricodherma*.

*Tricodherma* could playing a role as biological control agents in crop protection (Otero et al., 2013), in general OMF has a role in stimulating the germination of orchid seeds (Andersen et al., 1996), in supporting the provision of nutrients for growth and development of plantlets. In addition for induce the resistance of *Phalaenopsis* sp. against infections by bacteria, fungi, and viruses (Moreno et al., 2000). In this research, the identification of *Tricodherma* was molecularly conducted through analysis of rDNA-ITS sequence isolated from *Phalaenopsis amabilis* grown in Indonesia. This research was expected to become the basic information on the development of cultivation and protection of nature orchids and where possible, on the prevention of the occurence of the mentioned diseases in Indonesia.

1. Materials and Methods
   1. Plants Materials

The sampling of healthy roots of *Phalaenopsis amabilis* was conducted at four different locations: orchid garden in Condong Catur and Parakan (Yogyakarta), Balikpapan Botanical Garden (East Kalimantan), and Sultan Adam Forest (South Kalimantan).

* 1. Molecular Analysis

Genomic DNA isolation was performed using techniques modified from *cetyltrimethylammonium bromide* (CTAB) method (Doyle and Doyle, 1987) on samples of pure cultures of isolated mycorrhizal endophyte *Tricodherma*. Genomic DNA was PCR amplified according to the manual instructions of *GoTaq® Green PCR mix (Promega)*. Predenaturation reaction and amplification was carried out using methods by Nadarajah et al. (2014), with a pair of universal primers, rDNA-ITS1 (5'-TCCGTAGGTGAACCTGCGG-3') and rDNA-ITS4 (5'-TCCTCCGCTTATTGATATGC-3'). PCR products were visualized using electrophoresis on 2% agarose gel stained with ethidium bromide and 100 bp *Vivantis DNA ladder* was used as marker. The visualized DNA bands indicated the length of the targeted base pairs of rDNA-ITS, which were subjected to sequencing.

* 1. Phylogenetic Analysis

Sequencing results were analyzed using *Sequence Scanner* software; the nucleotide sequences were combined using *EditSeq* and *SeqMan* of the *Software Suite for Sequence Analysis DNASTAR Lasergene DM Version 3.0.25*. *BLAST* software was used to determine and to compare the sequence homology with the data contained in the DDBJ database. Comparation between sequences of isolates was carried out using *Algorithm Multiple Alignment Parameters DNA* with *Kimura-2 Parameters*, relationship and phylogenetic analysis using the *Neighbour Joining* of *MEGA 5 Beta* program. Statistical analysis on internal branch was done using *the bootstrap* value with 1000 replication.

1. Materials and Methods
   1. Sample Collection

A total of 12 samples of healthy roots of *Phalaenopsis amabilis* were isolated from 4 different locations, such as orchid garden in Condong Catur and Parakan (Yogyakarta), Balikpapan Botanical Garden (East Kalimantan), and Sultan Adam Forest (South Kalimantan). Positive result of *Tricodherma* isolate was found from orchid garden in Condong Catur.

* 1. Molecular Analysis

ITS rDNA amplification results showed a specific band with a size between 600 and 750 bp similar to that reported by Nadarajah et al. (2014). Internal transcribed spacer (ITS) is an area of ​​the nuclear ribosomal DNA (nrDNA), which has the role of providing important information on the reconstruction of phylogenetic trees at different taxonomic levels (Bayer et al., 1996), as well as similarity at the level of intrageneric (Lee et al., 2010).

Sequencing results were combined and analyzed using *DNASTAR Lasergene DM Version 3.0.25*. Total number of nucleotide of MP isolates which succesfully scanned ​​was 618 with 56.5% GC content. Analysis results were obtained by incorporating sequences to <http://blast.ncbi.nlm.nih.gov/> site and confirmed that isolates were *Tricodherma*. Furthermore, analysis of homologous search sequences carried out using BLAST on NCBI site resulted in 44 sequences indicating a close relationship with high level of similarity of 99%. This result showed that sequences are representative of *Tricodherma* isolates, a non-pathogenic (endophyte) associating with the root of orchid (OMF).

The results of analysis showed another *Tricodherma* isolate from Indonesia, precisely from North Sumatra (KC479808) was isolated from palm oil rhizosphere. The whole isolates originated from various countries in Asia (India, Taiwan, Korea, China), America (USA, Venezuela, Brazil, Costa Rica), and Europe (Austria, Swedia, Polandia). Some isolates of *Tricodherma*, was collected fromsoil (KJ739790; JQ863230; DQ083023; EU330955; KC576709), roots of apple seedling (EU003017), *Populus trichocarpa* (KC007180; KC007211; KC007207), and Picea abies decayed (DQ093713), and from compost of Oyster mushroom (DQ164413). Total bases amplified from each isolate ranged from 583 to 1125 bp with a total amplified base of 1500 bp. Maximum *BLAST* score between isolates ranged from 1033 to 1105 with homology (ident.) 99% and query cover ranged from 84 to 99%.

Analysis on 12 other *Tricodherma* isolates selected based on the distribution area of ​​each different country, showed that isolates from Indonesia were highly different from those of other countries. It was clear that isolates from Indonesia has been undergoing speciation and are different from isolates from other countries. Alignment result of nucleotide sequences showed high occurrence of point mutations in isolates from Indonesian, which were largely insertions and substitutions. Of the 554 total bases aligned, there were 44 bases having substitution consisting of 120 transition, 145 transversion, 11 insertion, and 13 deletion bases.

Each of these mutations had an influence on changes in amino acids formed. Of the total 184 amino acids belonging to isolates from Indonesia, some of which had very different percentage from those of other isolates. Some amino acids of isolates from Indonesia had increased their number dramatically compared to those of other isolates, i.e. Cys, Asp, Glu, Gly, His, Ile, Lys, Met, Gln, Arg,, Ser, and Tyr by 0.17%, 0.59%, 2.2%, 2.7%, 0.6%, 0.57%, 0.67%, 1.12%, 1.62%, 0.59%, 2.2%, 1.23%, and 1.10%, respectively, of total average. Several other amino acids had also decreased in number, i.e. Leu, Asn, Pro, Thr, Val, by 3.65%, 4.22%, 4.15%, 1.50%, and 0.44%, respectively, of total average.

It appeared that the resulted amino acid changes had enormous influence in the process of adaptation to the environment in Indonesia. Results of the relationship analysis between isolates through the reconstruction of phylogenetic tree showed that isolates from Indonesia were on separated branches and far apart from 13 other isolates (Fig. 1).



DQ164413

Fig. 1. Reconstruction of phylogenetic trees of *Tricodherma* isolates based on nucleotide sequences with Maximum Parsimony method with Bootstrap-1000 approach

1. Discussion

*Maximum Parsimony* method with *Bootstrap-1000* approach classify the data based on specific characters (discrete character states) on which nucleotide sequences are used to obtain information about evolution through evolutionary data changes. The results of analysis were classified based on distribution on the geographical areas. A lengthy branch, which well separating between the two groups, indicated that isolates from Indonesia had considerably evolved and it led to speciation events. Such changes in sequence through mutation was supported by the ability of the mycorrhizal itself to adapt to environmental changes as well as to its host range. The adaptability of mycorrhiza is strongly influenced by the presence of amino acid changes due to mutations that alter the function of genes which they arrange.

The reconstruction of phylogenetic tree, in addition to explaining relationship based on geographic regions, also determine the origin and history of distribution of the isolates. Trading activities of plants, fruits, and seedlings from several countries which are reported to have been infected by *Tricodherma* is most likely the cause of its spreading. Indonesia is known as an importer country of orchid plant seedlings from Asian countries such as Thailand, Singapore and Taiwan(Mahfut et al., 2016a), and possibly have imported seedling and fruit of apple and strawberry plants from other countries such as India and Italy that had been reported infected by *Tricodherma* isolates. Proximity with Singapore in terms of geographical location is also very possible to allow distribution activities of orchids between the two countries.

This study results proved that nucleotide sequences and amino acid of *Tricodherma* isolates from Indonesia were different with other isolates from another country (Asia, Europe, and America) due to adaptation to various environmental conditions. Furthermore, the mutations are capable of causing the *Tricodherma* to lose its ability to infect and to associate with host plants. Another possible result of mutations is that *Tricodherma* could have the ability to infect plants while showing pathogenic characters or otherwise become mycorrhizal endophytic.

One way to control pathogens of orchids is by using biological control, such as the mechanisms of mycorrhizal resistance induction. *Tricodherma* as OMF has a role as a biological agent in disease control(Otero et al., 2013) including infection of ORSV. This study results clarified the MP sequence as OMF *Tricodherma* isolates from Indonesia based on identification of its rDNA-ITS sequences. Furthermore, this study was expected to be the basic information beneficial for the improvement of cultivation effort as well as for the development of biocontrol agents through natural orchid protection against pathogens and diseases.

1. Conclusion

Molecular analysis based on rDNA-ITS resulted in isolate from the root of orchid plant in Yogyakarta being *Tricodherma*. These isolates showed 600-750 bp in length DNA products located on the ITS1-5.8S-ITS4 region. The sequence products showed insertion and substitution occurances, which may result in strain diversity and possible variation in severity. Reconstruction of phylogenetic tree using *Maximum Parsimony* and *Bootstrap*-1000 approach showed that Indonesian isolates have undergone speciation and have been positioned in the cluster, which are already far apart from the other isolates.

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