

# MOLECULAR CLONING OF *Elaeis guineensis* PHYTOENE SYNTHASE (*Egpsy*) AND ITS EXPRESSION IN MESOCARP TISSUES

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## ABSTRACT

*Phytoene synthase (psy) gene is responsible for the synthesis of carotenoids in plants. In this study complementary deoxyribonucleic acid (cDNA) and genomic sequences of psy were isolated by rapid amplification of cDNA ends and polymerase chain reaction (PCR) using ribonucleic acid (RNA) and DNA isolated from the mesocarp tissues of oil palm (Elaeis guineensis). The size of the full-length cDNA was 2233 bp. The open reading frame (ORF) was 1299 bp and encoded 433 deduced amino acid (AA) residues. The transcript profile revealed that psy was expressed at a maximum level at 12 weeks after anthesis (WAA). The lowest expression of psy was at 13 WAA, which was speculated as the transition of the leaf-like carotenoid to the formation of storage carotenoids. The AA sequence deduced from the cDNA was 70%-78% identical to PSY from other higher plants. The genomic sequence analysis revealed that the psy gene contains six exons and five introns. Southern blot analysis indicated that oil palm has only one copy of the psy gene in its genome.*

**Keywords:** carotenoids, oil palm, phytoene synthase (*psy*).

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## INTRODUCTION

In 2018, palm oil was a major edible vegetable oil, contributing about 34% of the world's vegetable oils consumption and produced mostly in Malaysia and Indonesia (Kushairi *et al.*, 2019). Crude palm oil is considered as the world's richest natural plant source of carotenoids. Malaysian crude palm oil contains typically 500-700 parts per million (ppm) of carotenoids (Mustapa *et al.*, 2011). Carotenoids such as  $\beta$ -carotene, lycopene and lutein found in vegetable oils, possess anti-oxidant, anti-carcinogenic activities (Uenojo *et al.*, 2007) and provide a primary dietary source of provitamin A in human diets (Tanaka and Ohmiya, 2008). Its deficiency could lead to blindness and premature death (Mayne, 1996). Besides their health-

promoting roles, carotenoids are economically important natural products used mainly in food, animal feed, pharmaceutical, nutraceutical and cosmetic industries. In higher plants, carotenoids are essential for providing photoprotective functions during photosynthesis and as precursors for abscisic acid (ABA) biosynthesis.

Gene cloning and functional research of the carotenoid biosynthesis pathway have been extensively carried out in many photosynthetic and non-photosynthetic organisms. The pathway of carotenoid biosynthesis was elucidated in 1987, starting with phytoene synthase (*psy*) in tomato (Zhang *et al.*, 2016). The enzyme catalyses the first step in the synthesis to produce phytoene from geranylgeranyl pyrophosphate (*ggpp*). Phytoene undergoes a series of four desaturations that result in the formation of lycopene. In higher plants, these steps are catalysed by two enzymes, phytoene desaturase (*pds*) and  $\xi$ -carotene desaturase. Lycopene undergoes cyclisation by lycopene epsilon cyclase (*lcyε*) and lycopene  $\beta$ -cyclase

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(*lcyb*) to produce  $\alpha$ - and  $\beta$ -carotenes. Subsequent hydroxylation reactions produce zeaxanthin and lutein (Nisar *et al.*, 2015).

The molecular basis of oil palm carotenoid synthesis has received little attention compared to other higher plants such as tomato (*Solanum lycopersicum*). Tomato is a well-established model plant for carotenoid biosynthesis in fleshy fruit. It has the wealth of genome resources and genetic transformability. However, it is not considered as a comparative reference to study the transcriptional regulation of carotenoid synthesis pathway especially in oil fruits such as oil palm (Tranbarger *et al.*, 2011). The oil palm (*Elaeis guineensis*) genome has been reported (<http://genomsawit.mpob.gov.my/>). The genome data will be a helpful resource to identify key genes involved in its carotenoid metabolism (Singh *et al.*, 2013). To date, only seven of the oil palm carotenoid genes have been reported, namely, *pds* (Rasid *et al.*, 2005), *lcyb* (Rasid *et al.*, 2003; 2009), 1-deoxy-D-xylulose 5-phosphate synthase (*dxs*) and 1-deoxy-D-xylulose 5-phosphate reductoisomerase (*dxr*) (Khemvong and Suvachittanont, 2005), zeta-carotene desaturase (*zds*) (Rasid *et al.*, 2005), full length of *lcye* (Rasid *et al.*, 2009) and *psy* from *Elaeis oleifera* (Wan Nur Syuhada *et al.*, 2012; Rasid *et al.*, 2008).

Geranylgeranyl pyrophosphate (*ggpp*) is the main precursor for carotenoid biosynthesis in plants. However, this precursor is also shared with other vital metabolic pathways including the synthesis of gibberellic acid, chlorophylls, plastoquinones and tocopherols. Therefore, modifying the carotenoid content and/or composition could have potential negative effects on plant growth (Tanaka and Ohmiya, 2008). Since *ggpp* serves as a precursor for other metabolic pathways, PSY plays an important role in controlling the flux (of carbon) in the carotenoid biosynthetic pathway (Lopez-Emparan *et al.*, 2014). The transcription of *psy* genes has been shown to respond to ABA, salt, light, temperature, drought, photoperiod, development cues and post-transcriptional feedback regulation (Cazzonelli and Pogson, 2010). This enzyme has been isolated previously from a range of plant species including tomato (Ray *et al.*, 1992; Fray and Grierson, 1993; Giuliano *et al.*, 1993; Misawa *et al.*, 1994; Fraser *et al.*, 1995), *Arabidopsis thaliana* (Scolnik and Bartley, 1994), tobacco (Busch *et al.*, 2002), maize (Buckner *et al.*, 1996; Wong *et al.*, 2004), melon (Karvouni *et al.*, 1995), citrus (Kim *et al.*, 2001; Kato *et al.*, 2004; Rodrigo *et al.*, 2004), pepper (Huh *et al.*, 2001), sunflower (Salvini *et al.*, 2005) and banana (Mlalazi *et al.* 2012). The modification of this gene could be effective and useful for increasing carotenoid content in industrial crops. In order to study metabolic pathways especially the carotenoid pathway, genes are usually organised in

operons. This operon will be inserted into plastid deoxyribonucleic acid (DNA) and then coexpressed as polycistronic transcripts. These transcripts are often processed further into monocistronic messenger ribonucleic acid (mRNA). For example, Golden rice 2 was developed by introducing maize *psy* in combination with *Erwinia uredovora* carotene desaturase (*crtl*). The amount of  $\beta$ -carotene in the endosperm was increased up to  $37 \mu\text{g g}^{-1}$  of the  $31 \mu\text{g g}^{-1}$  of carotenoids. It is equivalent to the recommended level of vitamin A for a day (Paine *et al.*, 2005).

Manipulation of *psy* in oil palm could potentially increase its carotenoid content as well as alter its composition. However, the molecular basis for the oil palm *psy* is currently unknown especially in *E. guineensis*. Thus, isolation of this gene will enable the investigation of its role in the carotenoid biosynthesis pathway. In this article, we report the isolation and characterisation of *psy* gene from oil palm (*E. guineensis*). The full length of *psy* complementary DNA (cDNA) and its genomic sequence were successfully isolated and characterised. The molecular phylogeny of *psy* and its expression in mesocarp tissues at different developmental stages were determined.

## MATERIALS AND METHODS

### Plant Materials

Oil palm fruits were collected from a field at Malaysian Palm Oil Board (MPOB), Bangi, Selangor, Malaysia. The mesocarp tissues at different fruit developmental stages, namely at 5, 7, 9, 11, 13, 15, 17 and 19 weeks after anthesis (WAA), were harvested for ribonucleic acid (RNA) isolation. Total RNA was extracted from the mesocarp tissues of oil palm according to Zeng and Yang (2002). Total DNA from mesocarp tissue was extracted using the modified cetyl trimethylammonium bromide (CTAB) method (Stewart and Via, 1993). The concentration and quality of the RNA and DNA were determined by spectrophotometer and agarose gel analysis. The DNA and RNA samples were stored at  $-20^{\circ}\text{C}$  and  $-80^{\circ}\text{C}$ , respectively, prior to further analysis.

### Reverse Transcription of mRNA and Reverse Transcription Polymerase Chain Reaction (RT-PCR)

The first-strand cDNA was synthesised from  $1 \mu\text{g}$  of total RNA sample using the SMART<sup>TM</sup> RACE cDNA Amplification Kit (Clontech, USA) according to the manufacturer's instructions. The synthesised cDNA was stored at  $-20^{\circ}\text{C}$  for PCR amplification.

The primers used in the initial reverse transcription PCR (RT-PCR) amplification were designed by using the Consensus- DEgenerate Hybrid Oligonucleotide Primer (CODEHOP) (Rose *et al.*, 2005). The software is available at <http://blocks.fhrcc.org/codehop.html>. Primer sequences were as follows: the sense primer: 5'-GGTGTACAACGTGGTGCTGAARCARGCN GC-3' and the antisense primer: 5'-TGAA GTTGTTGTTAGTCGTTGGCYTCDATYTC-3'. RT-PCR amplifications were performed using AccuPrime™ *Taq* DNA Polymerase System Kit (Invitrogen). The amplifications were carried out in a PTC-100 Thermocycler (MJ Research, USA), in 30-35 sequential cycles programmed as follows: initial denaturing cycle at 94°C for 30 s, 30 s annealing at 55°C-60°C, 2 min elongation at 68°C and a final cycle of 7 min at 68°C.

The 5' RACE primers (PSY15, 5'-TTGGCATA CTCTGCACAGACTTC-3' and PSY19, 5'-TAGCCGAAGGCGTAATGTGAGAAGC-3') were designed and synthesised based on the partial sequence of the initial amplified product. Then, the 5' RACE PCR was performed according to the protocol of the SMART™ RACE cDNA Amplification Kit (Clontech, USA). The first-round PCR was carried out with PSY15 and Universal Primer Mix (UPM) primer set, and the nested-PCR with PSY19 and nested UPM (NUPM) primer set. The nested-PCR product was purified and cloned into PCRII-TOPO vector (TOPO TA Cloning Kit, Invitrogen, USA) and sequenced.

End-to-end PCR was performed to amplify the complete cDNA and genomic sequence of *psy* using the gene-specific primers (PSYA1 and PSYAB4) that were designed based on 5' and 3' regions.

### Bioinformatics Analysis

The AA sequence analyses were performed using open reading frame (ORF) Finder (<http://www.ncbi.nlm.nih.gov/gorf/gorf.html>). Comparison of the sequence was performed by using BLAST of GenBank (<http://www.ncbi.nlm.nih.gov/BLAST/>). Multiple sequence alignment was carried out using Vector NTI software. Prediction of signal peptide was performed with SIGNALP. Full sequences were aligned using ClustalW ver. 2.1, while phylogenetic analysis was done using RAxML (General time reversible nucleotide substitution model with gamma distributed rate variation among sites and Felsenstein's bootstraps) visualisation with ClustalW by ETE3. Protein localisation site was predicted by Wolf PSORT program (Horton *et al.*, 2007). The presence of transmembrane domain was analysed by TMpred algorithm ([http://www.ch.embnet.org/software/TMPRED\\_form.html](http://www.ch.embnet.org/software/TMPRED_form.html)).

### Southern Blot Analysis

A total of 30 µg of genomic DNA were digested with *Bam*HI, *Hind*III, *Eco*RI and *Spe*I, separated on a 0.9% agarose gel and transferred onto Hybond XL membrane (Amersham Bioscience). A specific primed DNA probe was synthesised from the *E. guineensis* phytoene synthase (*Egpsy*) cDNA using the TAKARA random primer (TAKARA) according to the manufacturer's instructions. The membrane was hybridised at 60°C with  $\alpha$ -<sup>32</sup>P-dCTP-labelled probe. After hybridisation, the membrane was washed in 2 × SSC/0.1% SDS at 60°C for 10 min followed by 1 × SSC/0.1% SDS at 60°C for 10 min.

### Reverse Transcription Quantitative Real-time PCR (RT-qPCR)

The expression study of oil palm *psy* was performed using reverse transcription quantitative real-time PCR (RT-qPCR) analysis. Total RNA from mesocarp of 5-19 WAA fruits were used in the analysis. Total RNA (10 µg) was reverse transcribed with the High Capacity cDNA Archive Kit (Applied Biosystem) according to the manufacturer's protocol. The cDNA (100 ng) was subjected to qPCR assays in triplicate on an ABI PRISM 7000 Sequence Detection System (Applied Biosystems, USA) using TaqMan Universal PCR mastermix according to the manufacturer's recommendation (Applied Biosystems, USA). The primers for the *psy* gene were CCGATCTAGTGAGTTGTTCTTCGA (forward) and GGAATTATGCTGTATATACCATATCCGCAAT (reverse) and the probe for the gene was FAM-CTCGGAACAACAAGATC-NFQ. Meanwhile, the primers for the GAPDH were ACTGCTACTCAGAAGACTGTTGATG (forward) and TGCTGCTAGGAATGATGTTAAAGCT (reverse) and the probe for the gene was FAM-ACCCCTCCAGTCCTTG-NFQ. PCR cycling parameters were set at one cycle of 50°C for 2 min, 95°C for 10 min and then 40 cycles of 95°C for 15 s and 60°C for 1 min. Data was acquired and analysed by using the ABI PRISM 7000 Sequence Detection System software (Applied Biosystems, USA).

The expression profile of each cDNA was studied by the relative quantification of the mRNA using the comparative  $\Delta\Delta C_T$  method (Livak and Schmittgen, 2001). First, the threshold cycle ( $C_T$ ) value for each sample and its replicates was obtained from the amplification plots. The value represents the cycle at which the signal is first recorded as statistically significant above the background. Then, the average  $C_T$  values for the target gene in each of the samples were determined from replicates. This was carried out for both the target genes and the internal control glyceraldehyde 3-phosphate dehydrogenase (GAPDH). These values were normalised against

the average  $C_T$  values of the internal control ( $\Delta C_T$ ). The difference in  $C_T$  values ( $\Delta C_T$ ) is equal to the difference in expression after being normalised to the endogenous control.

## RESULTS AND DISCUSSION

### Isolation of *Egpsy*

PSY is located in the chloroplast (Fraser *et al.*, 2000) and it catalyses the first committed step of the carotenoid pathway that converts two molecules of *ggpp* into phytoene. This two-step conversion reaction is considered as a rate-limiting step in the pathway (Cunningham and Gantt, 1998). Consequently, *psy* has become the key target for genetic manipulation in many crop plants to increase the carotenoid content (Fraser *et al.*, 2002; Fray *et al.*, 1995).

In this work, *psy* cDNA was isolated from *E. guineensis* using degenerate primers (PSY5 and PSY6). A full-length cDNA encoding *psy* was obtained and designated as *Egpsy*. The gene reported in this article has been submitted to the PalmXPlore database (<http://palmxplore.mpob.gov.my/>) under the accession number p5.00\_sc00017\_p0037. The complete *Egpsy* has an ORF of 1299 bp which is preceded by 346 bases of 5-untranslated region (5-UTR) followed by 588 bases of the 3'-UTR (Figure 1). Interestingly, Wan Nur Syuhada *et al.* (2012) also reported that *psy* from *Elaeis oleifera* and *Egpsy* nucleotide sequences were highly conserved at about 98%. The BlastP search results demonstrated that the *Egpsy* clone showed a high sequence identity to PSY from other monocots, such as *Phoenix dactylifera* (91.16%) and Musa AAB Group (79.35%). Overall, the *Egpsy* sequence has a high degree of similarity with various species of plants (80.00%), alga (60.00%) and cyanobacteria (40.00%) but not with bacteria and fungi (16.90%).

The PCR product was also amplified directly from genomic DNA, sequenced on both strands and assembled using standard techniques. The total length of *Egpsy* genomic clone was 4342 bp, consisting of six exons and five introns. The sizes and organisation of the exons and introns are summarised in Figure 1. Comparatively, the number of exons and introns in *Egpsy* is similar to

the *psy* from wheat and maize (Wang *et al.*, 2009; Li *et al.*, 2008).

### *Egpsy* Sequence Analyses

In plants, the synthesis of carotenoids occurs in plastids. Thus, the enzymes involved in the pathway should contain plastid targeting signal for their transport into the plastids. Therefore, we looked for the plastid targeting signal using SIGNALP software and protein localisation using WoLF PSORT program (Horton *et al.*, 2007). The SIGNALP software predicted a 12-amino acid signal peptide with a molecular mass of 48.83 kDa and an isoelectric point of 9.09. The WoLF PSORT program suggested that *Egpsy* might be localised in chloroplast and the protein feature had a similarity of about 69% with PSY from *Arabidopsis thaliana* (<https://www.uniprot.org/uniprot/P37271>). This result may indicate that *Egpsy* is needed for photosynthesis and photoprotection function.

In order to examine the properties and conserved domains of the enzyme, the protein sequence was subjected to National Centre for Biotechnology Information (NCBI) Conserved Domain Search. We found that the protein has a single large prenyltransferase catalytic domain that belongs to the Trans-Isoprenyl Diphosphate, Head-to-Head (1'-1) condensation reaction (Trans\_IPPS\_HH) family (NCBO CDD No. cd00683) (Marchler-Bauer *et al.*, 2011). Analysis of the sequence also indicated the presence of other features corresponding to the Trans-Isoprenyl Diphosphate Synthases family. This domain is present in all PSY, which catalyse the head to head (1'-1) condensation of two *ggpp* molecules to produce phytoene (C30 and C40) (Figure 2). The analysis also indicated the presence of major aspartate-rich motifs that are thought to be responsible for substrate binding. This result was also confirmed through transmembrane region analysis of the sequence using TMpred algorithm. The result showed two strong transmembrane helices, in-to-outside from 3-24 residues and out-to-inside from 257-259 residues. The active site of the enzyme is hydrophobic and located within the transmembrane helical bundle where carbocation is generated, leaving the diphosphate group attached (Zviling *et al.*, 2007). Meanwhile, Lopez-Emparan *et al.* (2014) reported *Brassica napus* L. PSY also had a conserved trans-IPP region and several motives

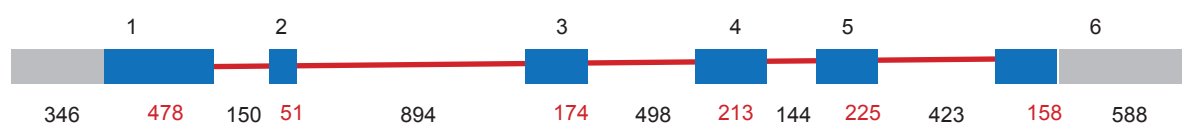


Figure 1. Diagram of exons and introns of *Elaeis guineensis* phytoene synthase (*Egpsy*). Grey and blue bars indicate untranslated and coding region, respectively. Meanwhile, thin red lines indicate introns and the number below them indicate their sizes in basepairs.

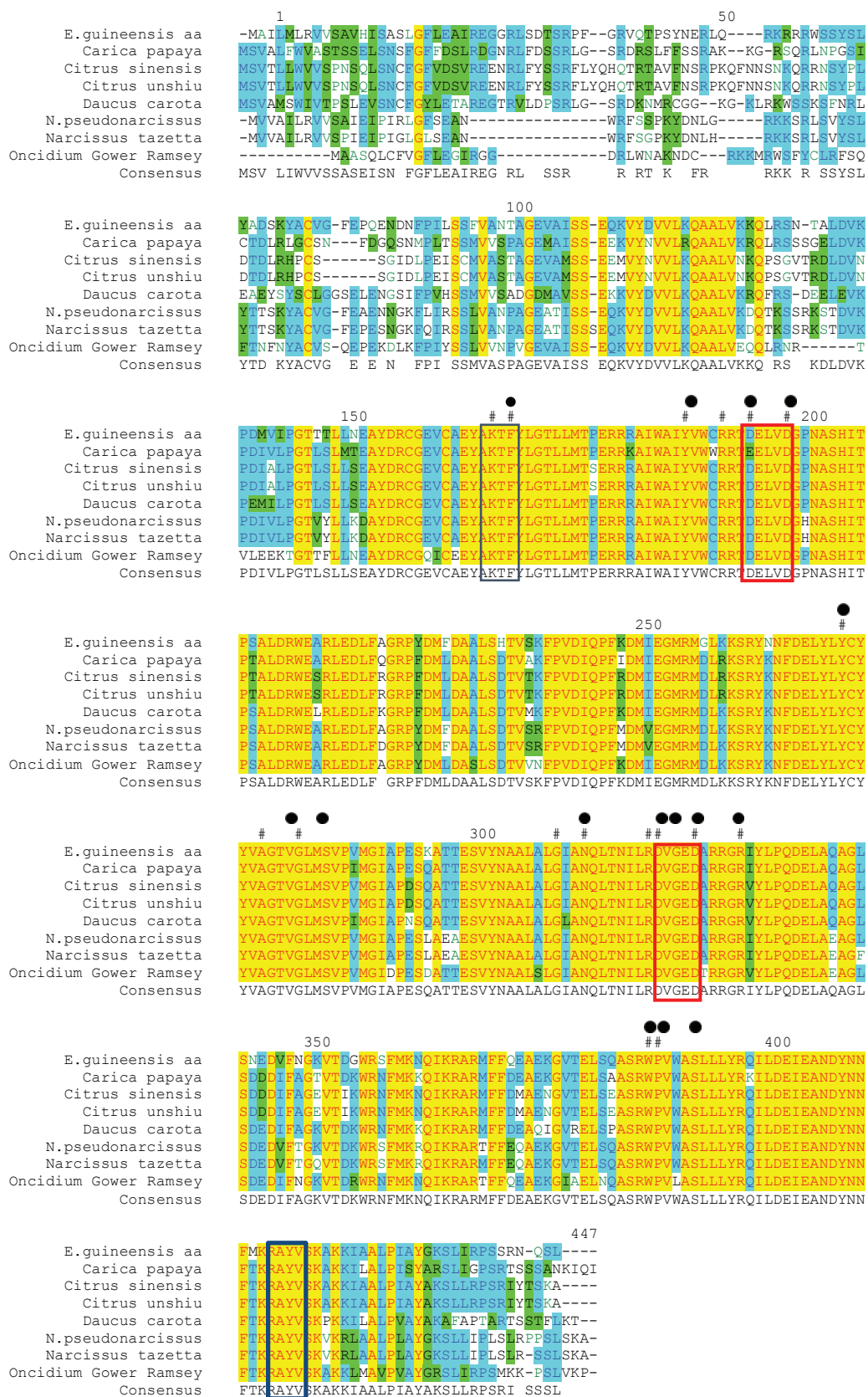


Figure 2. Amino acid sequence alignment of phytoene synthase (PSY) from oil palm and PSY from several other plants. Yellow colour shows highly conserved amino acid residues. Numbers above the alignment indicate the amino acid positions along the *Egpsy* protein. The two aspartate binding regions (DXXXD) and substrate-Mg<sup>2+</sup> binding site are indicated in red box. The active site lid residues are indicated in blue box. The substrate binding pocket are indicated (#). The catalytic residues are indicated (●). The sequences used in this analysis were as follows: *Carica papaya* (GenBank accession No. NW\_019019210), *Citrus sinensis* (GenBank accession No. NM\_001288886), *Daucus carota* subsp. *Sativus* (GenBank accession No. NM\_001329167), *Narcissus pseudonarcissus* (GenBank accession No. X78814), *Narcissus tazetta* (GenBank accession No. DQ984674), *Oncidium Gower Ramsey* (GenBank accession No. FJ859988).

such as plastid transit peptide and a putative *psy* active site (DXXXD) with conserved aspartate residues which was similar to *Egpsy*.

The deduced AA sequence for *Egpsy* was also subjected to phylogenetic analysis using ClustalW by ETE3. The sequence was compared with its respective orthologues (Figure 3). The phylogeny is rooted on the branch joining the outgroup to other sequences. PSY sequences were shown to fall into three defined clusters. One comprises the enzymes from cyanobacteria. The other clusters comprise of algae and higher plants. *Egpsy* has high identity with PSY sequences from the higher plants. The phylogenetic tree indicates that PSY sequences from higher plants, algae and cyanobacteria have clear divergence from each other. In higher plants, *Egpsy* is categorised into the monocot group together with *Phoenix dactylifera*, commonly known as date palm. This result showed that the PSY sequences from these two palm plants were closely related to each other. Bourgis *et al.* (2011) and Singh *et al.* (2013) also reported that oil palm and date palm

nucleotide sequences were highly conserved which was about 92.0%-94.4% identity. Interestingly, these two palms showed a completely different carbon-partitioning pattern with oil palm mesocarp containing mostly lipid while date mesocarp accumulates mostly sugars.

### Southern and Expression Analyses

The copy number of *Egpsy* gene in the *E. guineensis* genome was examined by performing Southern blot hybridisation. Using the 1.3-kb coding region of *Egpsy* as a probe, Southern blot analysis revealed the hybridisation pattern of 1-3 bands per lane (Figure 4). However, a single major hybridisation band was observed for two of the digestions, namely *Bam*HI (~7 kb) and *Eco*RI (~5 kb). Digestion using *Hind*III and *Spe*I produced two (~3 kb and ~6 kb) and three (~1 kb, ~3 kb and ~4 kb) major hybridisation bands, respectively. The three sites of *Spe*I were located in the intron whereas *Hind*III was in exon 5 (Figure 1). Overall, the result

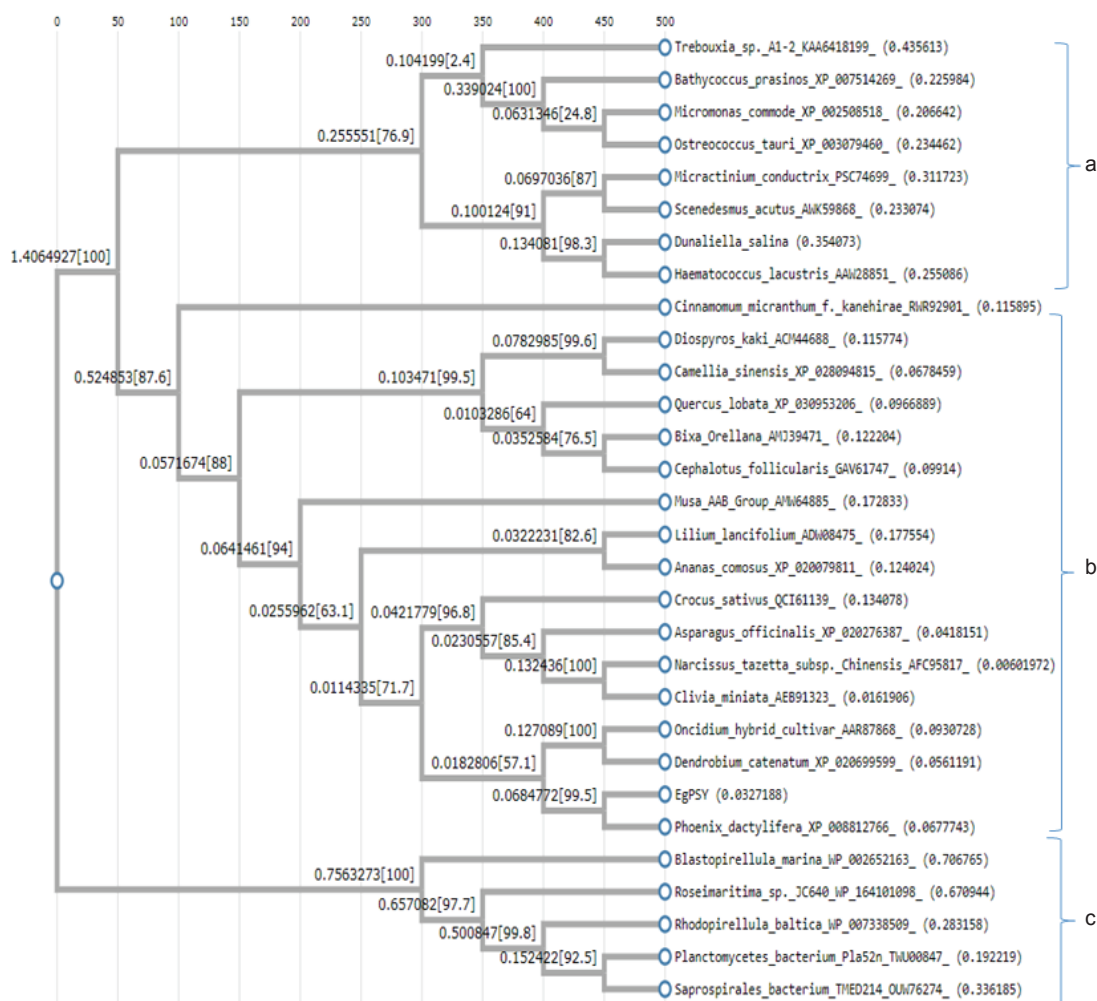


Figure 3. Phylogeny of phytoene synthase (PSY) amino acid sequences. The value at the nodes and branches represent the length of each branch. The values in bracket indicate the bootstrap support of each branch. The main clades labelled algae (a), plant (b) and cyanobacteria (c). The GenBank accession numbers are indicated at the end of the species names.

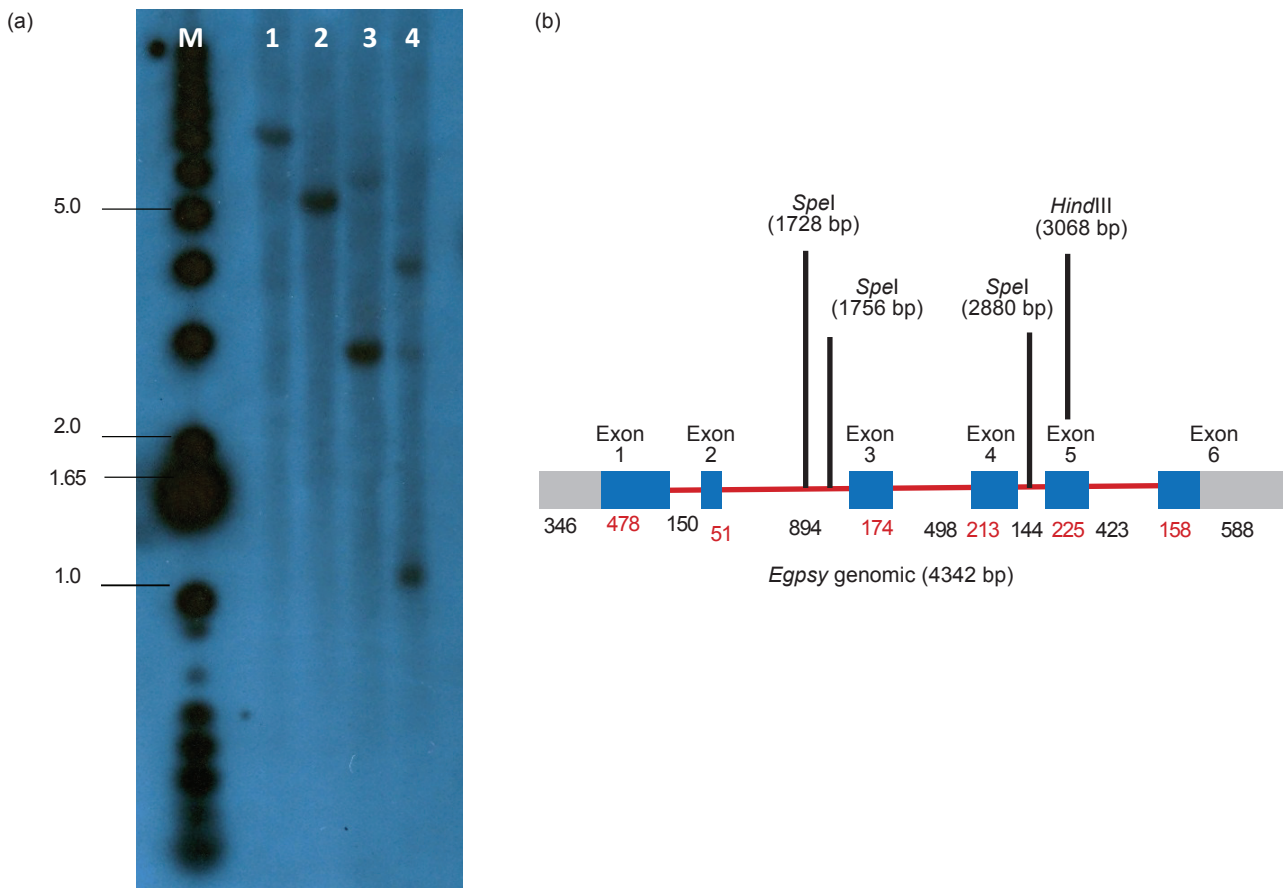


Figure 4. Southern blot analysis of *E. guineensis* phytoene synthase (*Egpsy*). (a) Thirty micrograms of total DNA of *E. guineensis* were digested using four different restriction enzymes namely BamHI, EcoRI, HindIII, and SpeI. Lane 1 is *E. guineensis* DNA digested with BamHI; Lane 2, *E. guineensis* DNA digested with EcoRI; Lane 3, *E. guineensis* DNA digested with HindIII; Lane 4, *E. guineensis* DNA digested with SpeI. (b) SpeI and HindIII restriction sites in the *Egpsy* genomic sequence.

suggested that there is only one copy of *psy* gene in the genome of *E. guineensis*. The copy number of *psy* gene in *E. guineensis* genome was also examined using the available oil palm genome data (<http://genomsawit.mpob.gov.my>). The search result indicated that there is a single copy of *psy* gene in *E. guineensis*. The gene was predicted to be located on chromosome 16.

Mesocarp tissue is the storage site of carotenoid accumulation in oil palm. Since PSY is the important control point of flux in carotenoid biosynthesis, the expression of its gene should be highly regulated in this tissue. The expression level of *psy* was examined during oil palm fruit development at weekly-intervals starting from 5 WAA until 19 WAA (Figure 5). The expression of *psy* in 5 WAA mesocarp tissue was about 1.72 folds higher than the calibrator (13 WAA). The expression increased to about 2.38 folds at 6 WAA, before decreasing to about 1.7 folds at 7 WAA. The *psy* expression remained at about the same level before decreasing at 10 WAA to about 1.3 folds, rebounded at 11 WAA and reached the peak at about 2.4 folds at 12 WAA. Surprisingly, mRNA level of *psy* decreased to the lowest level at

13 WAA. At 14 WAA, the *psy* expression increased and remained at about the same level thereafter. Overall, *Egpsy* was highly expressed in 6, 11, 12, 17 and 19 WAA as compared to the calibrator. This pattern suggested that *Egpsy* is involved in early and ripening stages of oil palm carotenoid biosynthesis, which is consistent with the findings from *E. oleifera* reported by Rasid *et al.* (2008). Oil synthesis and carotene storage in mesocarp commence at 15 to 16 WAA, reaching a peak at 20WAA (Kaur and Sambanthamurthi, 2008; Singh *et al.*, 2013). In young and mature tissues, the major product of carotogenesis was leaf-like xanthophylls which is mainly lutein and neoxanthin (Tay and Gwendoline, 2006). However, during fruit ripening, the lutein will decrease drastically and shift to carotene, mainly  $\alpha$ - and  $\beta$ -carotene (Kaur and Sambanthamurthi, 2008). The low expression at 13 WAA could possibly be due to the transition from the synthesis of leaf-like carotenoids to the formation of carotenes. The expression pattern observed for *psy* is also quite similar with the expression of a gene encoding phytochrome-interacting factor (*pif*). This protein was shown to repress *psy* and down-regulate the

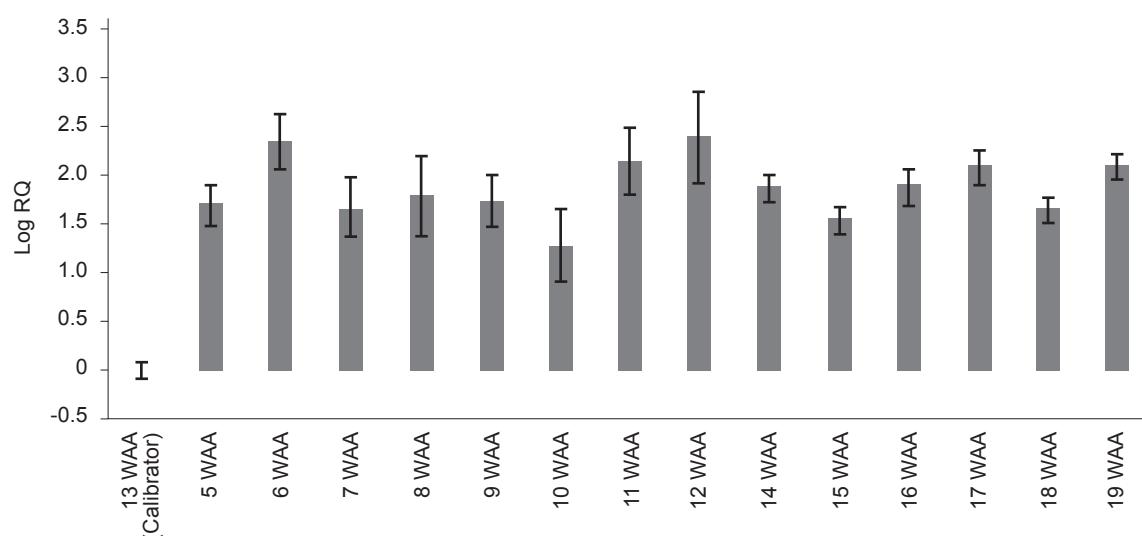


Figure 5. Relative quantities of phytoene synthase (PSY) mRNA in mesocarp tissue of oil palm fruit at different developmental periods. Each week after anthesis (WAA) sample was individually assayed in triplicates. Values shown represent the log Real-Time Quantitative (RQ) and bars indicate the standard error. The expression in the 13 WAA tissue was used as the calibrator.

accumulation of carotenoids (Toledo-Ortiz *et al.*, 2010). Interestingly, the *pif* gene was concomitant with the gene encoding phytochromes (*phy-A*, -B, and -C) in oil palm which was reported to be only expressed at 100 DAP (14 WAA) (Tranbager *et al.*, 2011). Therefore, the expression data of *psy* could suggest its important role, possibly along with other regulators, in the accumulation of carotenoids in oil palm fruit.

## CONCLUSION

We have isolated a cDNA clone and genomic DNA encoding *E. guineensis psy*. The sequence of this clone was shown to have high identity to *psy* sequences from other plants. We also examined its expression patterns during fruit development. The expression of *psy* was shown to be differentially expressed along the fruit developmental stages. This result suggests that *Egpsy* may regulate the synthesis and accumulation of carotenoids in the *E. guineensis* fruit. In essence, the sequence and expression analyses of the *psy* gene provided useful information for further understanding of its roles in the molecular regulation of carotenoid production and biosynthesis in *E. guineensis*.

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