

## Eggplant (*Solanum melongena* L.) polyphenol oxidase multi-gene family: a phylogenetic evaluation

Aravind Kumar Jukanti · Ramakrishna Bhatt

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**Abstract** Polyphenol oxidases (PPOs) in different *Solanum* species including eggplant have been studied. PPOs have been implicated in undesirable enzymatic browning of eggplant fruit and also in plant defense. The main objective of this study was to identify and accelerate the further functional characterization of additional eggplant PPOs that are involved in food biochemistry and defense-related functions. Eggplant PPOs identified earlier were used in “Basic local alignment search tool (BLAST)” search against expressed sequence tag and nucleotide databases. We have identified seven additional sequences which were almost complete in length. The sequences of the PPOs were aligned and their phylogenetic and evolutionary relationships established. The sequences are quite diverse, broadly falling into two major clusters; three PPOs form a separate branch/minor cluster. The thirteen sequences had conserved copper A binding sites but copper B binding sites differed considerably in two new PPO sequences (AFJ79642 and ACR61398). A third conserved ‘Histidine-rich’ region has been identified at the ‘C’ terminus of the eggplant PPOs. In addition, all the seven new PPOs exhibited at least one glycosylated sequon in the mature PPO sequence. Identification of additional PPO genes will further help in functional and biological characterization of these PPOs.

**Keywords** Eggplant · Polyphenol oxidase · Phylogenetic analysis · Multi-gene family · N-glycosylation

### Introduction

Polyphenol oxidases (PPOs) can oxidize specific phenolic substrates in the presence of oxygen in contrast to peroxidases which oxidize phenols in presence of H<sub>2</sub>O<sub>2</sub>. PPOs are ubiquitously distributed in plants (Mayer and Harel 1979); they play a role in food quality and in plant defense against pest and pathogens (Thipyapong et al. 1995; Thipyapong and Steffens 1997; Wang and Constabel 2004). PPOs are also involved in: time-dependent darkening and discoloration of cereal-based products (Baik et al. 1994), biosynthesis of flavonoids (Ono et al. 2006) and in oxidation of flavonoids (Pourcel et al. 2005). PPOs come into contact with phenolic substrates that are released due to tissue damage. The phenols are oxidized to highly reactive *o*-quinones which either self-polymerize or further react with nucleophiles to produce dark colored pigments that are usually undesirable in fresh or processed foods (Anderson and Morris 2001).

PPOs contain two copper (Cu) binding sites (Cu A and Cu B) and the Cu ion is bound by conserved histidine residues. PPOs interact with molecular oxygen and phenolic substrates at Cu-A and Cu-B sites (Van Gelder et al. 1997). PPOs are nuclear encoded enzymes, synthesized as precursor proteins in cytosol, processed to mature proteins and imported into plastidial thylakoid membranes (Koussevitzky et al. 1998; Sommer et al. 1994). The typical N-terminal transit peptide of PPOs is about 80–100 amino acids in length and during the chloroplast import the molecular weight of the enzyme is reduced from ~65–70 to <60 kDa (Dry and Robinson 1994; Van Gelder et al. 1997). PPOs are basically three types based on the substrates they catalyze: cresolases (monophenol oxidases), *o*-diphenol oxidases (catecholases) and laccase-like multi-copper oxidases (Shetty et al. 2011). Among the

A. K. Jukanti (✉) · R. Bhatt  
Central Arid Zone Research Institute, Jodhpur 342003,  
Rajasthan, India  
e-mail: jukanti5@yahoo.com; aravindjukanti@gmail.com

three classes of PPOs, *o*-diphenol oxidases have been extensively characterized in plants. Catecholases mostly occur as multi-gene families, introns are absent among the dicotyledonous PPOs genes reported so far (Shetty et al. 2011). But introns have been reported in monocot species like pineapple, banana and wheat (Massa et al. 2007). Up-regulation of PPO genes upon mechanical wounding and damage due to pests has been reported in several crop species (Thipyapong et al. 1995, 1997; Wang and Constabel 2004).

The role of PPOs in plant defense and enzymatic discolouration/browning affecting quality of crop/plant products has led to extensive identification and characterization of PPO genes in several plant species. Identification and characterization of all/most PPOs in a plant species will aid in better understanding the structural and functional differences among the multi-gene family. Solanaceae crops like potato, tomato and eggplant/brinjal form an important part of the daily diet in many parts of the world. Specifically, eggplant is an important constituent of the Indian cuisine. The role of PPOs in enzymatic browning of eggplant fruit has been vastly studied and also chlorogenic acid is shown to be the most predominant phenolic in the flesh of its fruit (Whitaker and Stommel 2003; Singh et al. 2009). Studies on eggplant PPOs describing their biochemistry, enzymatic action and genes have been published (Pérez-Gilabert and García Carmona 2000; Shetty et al. 2011). But due to their role in plant defense and food quality the identification of any additional PPO genes could be critical. Therefore, the main objective of this manuscript was to identify any additional eggplant PPOs utilizing bioinformatic tools and publicly available databases.

## Materials and methods

The PPO gene sequences published by Shetty et al. (2011) were utilized to analyze public databases for additional eggplant PPOs using basic local alignment search tool (BLAST; Altschul et al. 1990) against National Center for Biotechnology Information (NCBI; <http://www.ncbi.nlm.nih.gov/blast>). The *S. melangena* hits were exported to San Diego Super Computer Center biology workbench (<http://seqtool.sdsc.edu>) for sequence analysis. Sequences were aligned using CLUSTALW tool (Thompson et al. 1994) after translation to protein sequences. Phylogenetic relationships among the identified sequences were calculated using PHYLIP (Felsenstein 1989). Sequence identity matrix of all the thirteen PPOs was computed using LALIGN program (<http://workbench.sdsc.edu>). The molecular weight (statistical analysis of protein sequence (SAPS), Brendel et al. 1992) and isoelectric point (<http://seqtool.sdsc.edu/CGI/BW.cgi>) of different PPO sequences

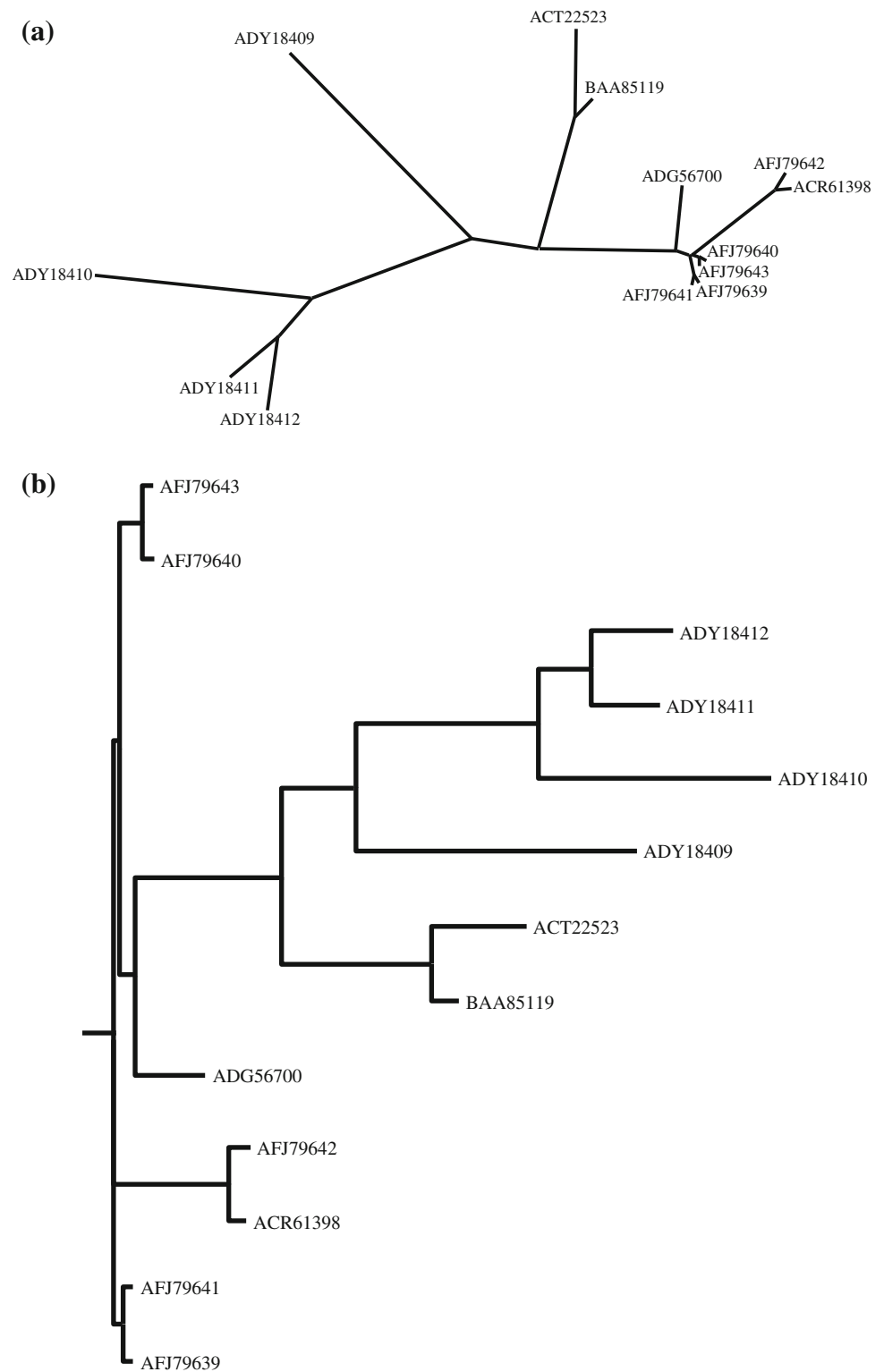
were also determined. The N-glycosylation sites and their position in the mature novel PPO sequences were analyzed using NetNGlyc software ([www.cbs.dtu.dk/services/NetNGlyc/](http://www.cbs.dtu.dk/services/NetNGlyc/)). All the additional sequences obtained (except one) were almost complete with minimal gaps, therefore, were further not sequenced. All thirteen eggplant PPOs are aligned and analyzed as described above.

## Results

Six PPO sequences were reported in eggplant by Shetty et al. (2011). However, large scale work on eggplant PPO ESTs/gene sequences has been reported in past few years. Therefore, a detailed analysis of PPOs in eggplant was performed; it resulted in identifying a multi-gene family of thirteen PPOs, grouped into two major clusters and a third minor group (Fig. 1). The published sequences (Shetty et al. 2011) were used as a search tool for NCBI BLAST search, seven additional eggplant PPO sequences with a sequence identity >62 % in overlaps of at least 266 amino acids were identified in this study. A phylogeny tree and evolutionary relationships of all the PPOs was constructed using PHYLIP (Fig. 1a, b). It was observed that six novel sequences identified are paired up with ADG56700 as a separate cluster and the seventh sequence (BAA85119) was closely related to ACT22523. Four previously characterized sequences (ADY184109, 18410, 18411 and 18411) form a second major cluster with ADY18409 forming a separate branch. Based on sequence comparison with the other eggplant PPOs the new sequences represent almost full length sequences except for one (BAA85119). The eggplant PPOs were compared with other plant PPOs (data not shown) including potato (*Solanum tuberosum* L.), tomato (*Lycopersicon esculentum* L.), tobacco (*Nicotiana tabacum* L.) and grape (*Vitis vinifera* L.). The tentative start site of mature PPOs is indicated based on the sequence alignment with higher plant and previously characterized eggplant PPOs. Tentative proteolytic processing sites for both stromal and thylakoid peptidases are also indicated. The length and molecular weight of mature PPOs ranged 584–601 amino acids and 65.9–67.7 kDa, respectively (Table 1). The molecular weight of mature PPOs was about ~56–58 kDa (after proteolytic processing). The isoelectric point of different PPOs was determined and it ranged from 6.062 to 7.956 (Table 1)

The percentage identity among the different PPOs was also calculated using LALIGN software. Eggplant PPO with GenBank accession number ACR61398 was identified to be an additional eggplant PPO. This sequence demonstrates ~61–87 % identity with the sequences used as search tool, the new sequence shows highest identity to an eggplant PPO with GenBank accession number,

**Fig. 1 a** Phylogenetic analysis (*unrooted tree*) of eggplant PPOs, **b** phylogenetic analysis (*rooted tree*) of different eggplant PPOs



ADG56700 (Table 2). ACR61398 sequence has relatively higher sequence identity (71–99 %) to the additional five PPO sequences (AFJ79639, AFJ79640, AFJ79641, AFJ79642, and AFJ79643) identified in this study, being almost identical to AFJ79642 (Table 2). Another novel

sequence with an accession number, AFJ79639 was similar to two other novel sequences, AFJ79640 and AFJ79643 differing only at six different amino acid (Aa) sites over a length of 595 Aa (data not shown). AFJ79639 also shows a similar trend to ACR61398 in sequence identity. AFJ79641

**Table 1** Characteristics of eggplant PPOs

PPO sequence	Amino acid number	Molecular weight (kDa)	Isoelectric point
ACR 61398	593	66.9	6.488
AFJ 79639	594	66.5	6.062
AFJ 79640	595	66.7	6.147
AFJ 79641	594	66.6	6.147
AFJ 79642	590	66.7	6.488
AFJ 79643	595	66.7	6.236
ADG 56700	601	67.6	6.549
ADY 18409	590	67.1	7.465
ADY 18410	584	66.3	6.294
ADY 18411	586	66.0	6.162
ADY 18412	584	65.9	6.832
ACT 22523	600	67.7	7.233
BAA 85119	266	30.4	7.956

is ~98 % identical to AFJ79639, AFJ79640 and AFJ79643. The sequence AFJ79643 follows same trend as the other new sequences. But the amino acid identity of

AFJ79642 was comparatively lower with the other novel sequences identified. Overall, the newly reported sequences vary considerably in their similarity to the search tool sequences ranging ~61–95 % (Table 2).

The amino acid sequences of the newly identified PPO sequences were analyzed for determining the most conserved features of all PPOs: two copper-binding sites and transit peptide sequence (Koussevitzky et al. 1998). The chloroplast transit peptide sequence harboring both thylakoid and stromal targeting domains extending ~80–90 amino acids is also found in the novel sequences. The transit peptide sequences of all the eggplant PPOs demonstrated conserved cleavage sites for both stromal peptidases (V, S, C, K/N) and thylakoid peptidases (L, A/T, A, S/N, A; Fig. 2). Both the copper-binding (A and B) regions of all the thirteen PPOs show considerable conservation of amino acid sequence. But among the two, 'A' site is relatively more conserved than 'B'. Two new sequences identified (AFJ79642 and ACR61398) in this study differed drastically from other PPOs especially in the copper 'B' region (Fig. 2). In these two sequences, the most conserved

**Table 2** Identity matrix of eggplant PPO sequences as percentages

	ACR 61398	AFJ 79639	AFJ 79640	AFJ 79641	AFJ 79642	AFJ 79643	ADG 56700	ADY 18409	ADY 18410	ADY 18411	ADY 18412	ACT 22523	BAA 85119
ACR 61398	100	92	92	93	99	92	87	67	61	66	66	74	71
AFJ 79639	92	100	99	98	92	99	94	71	64	70	70	75	83
AFJ 79640	92	99	100	98	92	98	93	71	64	69	69	75	82
AFJ 79641	93	98	98	100	93	98	94	70	64	70	70	75	82
AFJ 79642	99	92	92	93	100	92	89	67	61	67	67	76	71
AFJ 79643	92	99	98	98	92	100	95	70	64	69	69	75	82
ADG 56700	87	94	93	94	89	95	100	69	63	68	68	77	78
ADY 18409	67	71	71	70	67	70	69	100	65	69	68	70	75
ADY 18410	61	64	64	64	61	64	63	65	100	81	80	61	62
ADY 18411	66	70	69	70	67	69	68	69	81	100	92	66	70
ADY 18412	66	70	69	70	67	69	68	68	80	92	100	65	66
ACT 22523	74	75	75	75	76	75	77	70	61	66	65	100	93
BAA 85119	71	83	82	82	71	82	78	75	62	70	66	93	100



**Fig. 2** Sequence alignment of predicted amino acid sequence of thirteen eggplant PPOs. Transit peptide is boxed and proteolytic processing sites are indicated by (stromal peptidase, green) and (thylakoid peptidase, yellow). Conserved copper binding regions (A & B) are in bold font with underline, conserved histidine and cysteine

(thio-ether linkage) are highlighted in light grey. Third ‘His-rich’ region is highlighted in dark grey at the ‘C’ terminus. N-glycosylated (Asn-Xaa-Thr/Ser) sequons are highlighted in red. (\*) - single, fully conserved residue; (:) - conservation of strong groups; (.) - conservation of weak groups

amino acid (histidine, ‘H’) was absent. In addition to the two known copper-binding domains of all PPOs, it was noted that the eggplant PPOs consists of a third conserved ‘Histidine (His)-rich’ region at the ‘C’ terminus (Fig. 2). NetNGlyc, N-glycosylation software identified four glycosylated sites in the different mature PPO sequences. The ‘NLT’ sequon was the most widely conserved among the different eggplant PPOs (except ADG56700). The other glycosylated sites identified are: NGT/NTS—ACT22523 and BAA85119; NGT/NAS—ADY18409.

**Discussion**

During the past few years higher plant PPOs have been extensively studied due to their potential role in food biochemistry (Feillet et al. 2000) and plant defense (Constabel et al. 1995). Specifically, eggplant PPOs have been investigated due to their perceived role in browning of eggplant

fruit which is rich in phenols (Shetty et al. 2011). Chlorogenic acid is the major phenolic compound present in the flesh of eggplant fruit, accounting for ~70–95 % of the total phenolics (Whitaker and Stommel 2003; Singh et al. 2009). The biological importance of PPOs demands a comprehensive study of the number and role of different PPOs present in eggplant. Analysis of publicly available data has identified seven additional eggplant PPO sequences, constituting the eggplant PPO multi-gene family of thirteen genes. The phylogenetic and identity matrix analysis of the thirteen eggplant PPOs indicate the presence of two major clusters. In addition, it was also observed that three sequences (ACT22523 and BAA85119; ADY18409) probably represent different branch/clusters, these could include a few more yet to be identified PPO sequences.

The N-terminal region of the eggplant PPOs contained chloroplast transit peptide, these regions consists of both stromal and thylakoid targeting domains. Stromal and thylakoid targeting domains help in importing the mature



primarily involved in browning of eggplant fruit. In addition, the data presented in this manuscript could help in better understanding the implicated role of eggplant PPOs in defense against pests and pathogens.

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