

A Mechanistic insight into chemical cues and interactions involved in herbivory induced jasmonate mediated plant defense mechanism

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Abstract

The first step in plant defense mechanism is to sense the insect attack stimulus. Plant sensitivity of an insect attack is the first step of defense. Molecules generated by the oral secretion of the insect interact with the plant receptors to trigger plant defense mechanisms. We selected some highly cited insect elicitors molecules, volicitin, caeliferin, bruchin which interact with plant defense by interacting with plant elicitors (systemin, inceptin and peps) located on the plant cell surface. This interaction activates plant receptors SYR1, LRR, PEPR and triggers downstream defense signaling. The octadecanoid pathways, involving enzymes allene oxide synthase (AOS) and Hydroxyperoxide lyase (HPL) are activated. These enzymes mediate production of green leafy volatiles and Jasmonic acid by interacting with hydroperoxide molecules. We docked the elicitors with receptors and enzymes with substrates in the pathway of JA production. Phe was found to be an important amino acid that interacts with 13-hydroxyperoxides in the case of AOS to produce JA but not in the case of HPL. JA is converted to JA-Ile which shows strong binding with COI1 and COI1-JA-Ile complex docked with JAZ which showed strong interaction with five hydrogens and one salt bridge bond. AOS and HPL showed less than 40% identity for sequence and structure alignment. AOS and HPL had shown an interaction between each other and showed a common interaction partner of the Lipoxygenase family. HPL shows interaction with ADH2 (Alcohol dehydrogenase) involved in GLVs production. AOS also showed interaction partner AOC, COI1 and OPR1 which are involved in JA-induced plant defense mechanism.

Introduction

The insect plant ratio on our planet is interestingly high. Three hundred thousand plants have one million insect pests to compete with. Both the competitors, plants and insect, have evolved intricate and sophisticated strategies to survive within this aura of competition¹. Some counter defense strategies developed by plants include production of secondary metabolites, repellents, antifeedants and morphological features modification². Right after any damage cue sensed the plants promptly synthesize and discharge organic/inorganic volatile compounds as first step in preparation for defense strategies against attack³. These oral secretions in the form of regurgitates or saliva play key role in plant defense mechanism against herbivory⁴. The insect pest have diversified mode of feeding due to their different types of mouthparts this make them more adaptable than other fauna on earth.

Insect contributes to the largest group of the herbivores on the planet⁵ and this tends to the development of diverse counter defense strategies, against insect attack, by plants⁶. Although the execution and maintenance of this sophisticated mechanism is expensive for plants but this cost is inevitable for their survival in insect pest dominating environment^{7,8}. This defense mechanism is invoked by direct insect pest attack/physical contact or by sensing the chemical cues from neighboring plants attacked by insect^{9,10}. Insect generated signals by physical contact¹¹ ovipositional fluid¹², feces¹³, pheromone¹⁴ and insect feeding vibrations¹⁵ can be sensed by the plants result in initiation of plant defense cascade activation. The production, behavior and effect of insect oral secretion have been studied extensively in

context of plant defense mechanism against herbivory¹⁶. Oral secretions of the insects are divided in two broad categories as per their origin i.e., salivary fluid secreted from salivary gland and those derived from the gut called regurgitate. Both types of the secretions are rich in chemical compounds which can become elicitor or repressor of plant defense mechanism¹⁷.

Plants have developed very sensitive mechanisms to foresee and sense the danger of herbivory. A very intricate and synchronized coevolution has been developed between insects and plants. The key players for the initiation of plant defense are diversified chemical like insect-derived elicitors damage-associated molecules, plant hormone systems activated by insect oral secretion compounds⁴. Volicitin, N-(17-hydroxylinolenoyl)-L-glutamine (volicitin), was first insect oral secretion detected elicitor molecule in beet armyworm larvae, *Spodoptera exigua*¹⁸. Volicitin was also reported to be involved in synthesis of the volatile insect predator attractants after herbivores attacking maize plants¹⁹. Other insect small peptide elicitors like Inceptins have also been reported which are present both in plant and insect regurgitate. Inceptins, firstly reported in *Spodoptera frugiperda* oral secretions, are proteolytic fragments of a chloroplastic ATP synthase c-subunit that are generated in the insect midgut and induce defenses in cowpea and beans²⁰. These disulfide-bonded peptides are synthesized by protein degradation fragments of the chloroplastic ATP synthase (γ -subunit), formed in *Spodoptera frugiperda* gut by the digestion of plant proteins²⁰. The most common inceptin receptors in plant are leucine-rich repeat receptor-like protein (INR)²¹. Caeliferins a disulfoxy fatty acids were identified in the oral secretions of *Shistocerca americana* (American bird grasshopper) oral secretions were reported to contain a disulfoxy fatty acid compound called caeliferin¹⁹. Bruchins is another important elicitor was isolated from the oral secretions of *Callosobruchus maculatus* and *Bruchus pisorum*¹⁸.

Several defense signals are activated in plant right after insect herbivory which results initiation of different plant defense responses²². The first injury-induced peptide signal (elicitor) produced in plant was an eighteen amino acid long peptide named systemin. It was reported to be derived from its inactive form prosystemin right after insect bite in tomato²³. Systemin promotes Jasmonic acid buildup in plant cells and triggers the genes expression for the production of defense related plant proteins²⁴. A leucine-rich repeat receptor kinase (LRR-RK), called SYR1 has been reported as a receptor of systemin in plants²⁵. Later some other injury-induced peptide based plant elicitors had been identified. The application of synthetic twenty three amino acid long maize Peps could mimic the *Spodoptera exigua* attack and found to induce jasmonic acid production²⁶. A short peptide AtPeps acts as signals to activate both jasmonic and salicylic acid signaling pathways in *Arabidopsis*²⁷. The reported receptors of AtPeps—AtPEPR1 and AtPEPR2—are classified under the LRR-RK protein family^{28,29}.

The allene oxide synthase (AOS) and hydroxyl peroxide lipase (HPL) enzymes, responsible for the production of defense signals in the form of jasmonates and aldehydes (GLVs), respectively³⁰, used response of the plants in response to injury or damage is one of the significant components of insect pest control strategies in agriculture for the management of pest population^{31–33}. By increasing the host

plant resistance we can minimize the use of insecticides which in turn. Genetically modified crops have been introduced during last two decades were found to be more economical to cultivate than the use of insecticides extensively.

This study can be proved more helpful in investigations and revelation of plant stress response pathways, interactions structure, and function of proteins involved. In this project we attempt to cover the entire chain of herbivory defense cascade from the recognition of a feeding insect pest, through the production of defense molecules or utilization of physical defenses. Firstly, the primary events of herbivory induced the defense responses are studied and later complex intracellular signaling cascades was predicted, with a special focus on the jasmonate pathway.

Results

The objective of this study is to determine the interactions among the insect/plant elicitors and receptor molecules in herbivory induced plant defense mechanism pathway (Fig. 1).

Three-dimensional protein structure prediction

The first step in a proteomic study is to have the 3D structure of the protein. 3D structures of the selected protein could not be found Protein Data Bank (PDB) so predicted through I-TASSER and MODELLER. The online server I-TASSER send the results via email containing five models, against each submitted query sequence, with different confidence levels (C-score). The models with high scores were selected and evaluated for quality check. The predicted 3D protein models, either from MODELLER or I-TASSER were refined and evaluated and good quality models were selected (Table 2) (Fig. 2–3)

Docking Analysis

Interaction of insect elicitor molecules with plant elicitors

Docking was performed on all selected inhibitors by using AutoDock Vina. The main purpose of docking was to find the potential interactions between insect elicitor molecules and plant receptors to find out how these molecules interact with each other to trigger plant defense pathways (Table 3). The lowest binding energy (-6.9kcal/mol) was recorded for the Volicitin-Systemin complex (Fig. 4a) followed by that of the Volicitin-PEP1 complex (-6.8kcal/mol) (Fig. 4b). The highest number of hydrogen bonds (5) was found in Caeliferin-Systemin (Fig. 4c) and Bruchin-PEP1 complexes (Fig. 4d). The hydrogen bond distance among the docked complexes was ranging from 2.7 to 3.17 Å which reflects good interaction strength (Fig. 4a-f).

Interactions of plant elicitors with plant receptors

Protein-Protein docking was done by the PatchDock server. The input of this server is two molecules of any type: proteins, DNA, peptides, drugs. The output is a list of potential complexes sorted by shape complementarity criteria. The amino acid interaction between docked proteins was observed through DIMPLOT (Table 4). Overall a strong interaction was observed as the hydrogen bond distance for all the

interactions were less than 3 Å (Fig. 5a-c). Systemin-SYR1 formed the highest numbers (8) of hydrogen bonds (Fig. 5a). Inceptin, although released from insect regurgitate, acts as a plant elicitor and interacts with plant receptor LRR by forming a strong hydrogen bond of 2.59Å bond distance (Fig. 5c).

Jasmonate biosynthesis

Interaction of AOS and HPL with their ligand molecules

Hydrogen bonding in the protein-ligand complex was found for both proteins (Table 5). The lowest binding energy (-6.7kcal/mol) has resulted in the 13(S) Hydroperoxide-HPL complex (Fig. 6a). Four hydrogen bonds, with bond distances ranging between 3.20 and 2.55 Å, were found in 13(S) Hydroperoxide-AOS complex (Fig. 6b). The hydrogen bond interactions of 9(S) Hydroperoxide with HPL and AOS are shown in Figs. 6c-d.

Jasmonate signaling and regulation

Interaction of COI1 with JA-Ile

Coronatine Insensitive Protein 1 (COI1) is a nucleus receptor that binds with the JA-Ile (Jasmonate Isoleucine) to degrade JAZ (Jasmonate ZIM domain) protein. The complex was docked successfully with - 5.8kcal of binding energy. Three good strength hydrogen bonds (2.81–3.01 Å) were also observed (Fig. 7a-b).

Interaction of COI1 receptor with JAZ protein

JAZ proteins inhibit the activity of MYC2 by binding to their respective transcription factors. The production and deposition of Jasmonic acid in damaged cells is the first response herbivory or mechanical damage to the plants. The jasmonic acid further converted to (+)-7-iso-JA-L-Ile. This binds to coronatin-insensitive 1 (COI1) protein. This event leads to the binding of COI1 with Jasmonate ZIM-domain (JAZ) protein. This complex is recognized by 26S proteasome and JAZ is degraded by ubiquitination process⁴.

Keeping in view evidence from literature, that JA-Ile- COI1 binds to JAZ which leads to its ubiquitination of JAZ by 26S proteasome⁴, we performed protein-protein docking of JA-Ile- COI1 complex with JAZ by assigning JA-Ile- COI1 as ligand and JAZ as a receptor. The resulting docked complex showed a lower binding affinity of -8.20 kcal/mol. Seven hydrogen bonds were formed with bond distance ranging from 2.61 to 3.21 Å and one salt bridge bond between Arg211 (JAZ) and Glu203 (COI1) (Fig. 8a-b).

Protein-protein interaction)

STRING database was used to find out the interaction partner of Hydroperoxide lyase and Allene oxide synthase.

Allene Oxide Synthase (AOS)

Eleven nodes and thirty-six edges with an average clustering coefficient of 0.829 were observed in the case of AOS (Fig. 9). The interaction with ten proteins in the network was based on co-expression and

text mining. AOC showed the highest homology score (0.979) through co-expression followed by that of HPL (0.907) through text mining and co-expression and loxD (0.863) through co-expression.

Hydroperoxide lyase (HPL)

Eleven nodes and twenty-nine edges with an average clustering coefficient of 0.471 were observed in the case of HPL (Fig. 10). The interaction with ten proteins in the network was based on co-expression and text mining. ADH2 showed the highest homology score (0.971) through text mining followed by that of loxC (0.693) through text mining and co-expression and Solyc09g055900.2.1 (0.958) through co-expression and text mining.

Protein sequence and structure alignment

The sequences show only a percent similarity of 60.7 while percent identity 35.9 was observed (Fig. 11). In the case of 3D structure superposition, the percent identity was 37.3 was observed (Fig. 12a-b).

Discussion

The plants and insects have an intricate evolution history. Both coevolved sometimes as mutualistic relations and mostly as competitors. Insect plant co evolution is the most speedily occurring event in nature. As the insects acquire advanced capabilities to damage plants in turn the plants response back to the damage by modifying it defense machinery. This is the interplay of insect-plant chemicals to become winner. The first and foremost event in initiation of plant defense mechanics is the recognition of the stimulus by plant receptor cells. Insect oral secretions induces several internal signals from the wounded tissues, including jasmonate signaling^{39,40}. Although many oral secretions might not serve a specific function to the insect, salivary proteins play a more active role in modulating the interactions between plant and insect herbivore⁴¹⁻⁴³. The plants release volatile compounds such as repellents for insect pest and attractants for parasitoides/predators act as an alarm for herbivore attack⁴⁴. The signals in the form insect oral secretion elicitor compounds are perceived by plant elicitors and induce defense responses afterwards. Jasmonic acid is the key player of insect herbivory induced plant defense mechanisms in plants.

In the current study, we selected literature that reported insect-derived elicitors and predicted their interaction with plant elicitors and receptors in the Jasmonic acid pathway. The 2D structures were downloaded from PubChem in .sdf format and later converted into .pdb format through Chimera. The sequence of plant-derived elicitors was retrieved through UniProt in FASTA format. The 3D structure of plant receptors was not available in Protein Data Bank (PDB) those were modeled through homology-based modeling (Modeller) and *Ab-initio* method (I-TASSER). I-TASSER server uses multiple templates to predict protein structure⁴⁵. It generates five models with different confidence scores. Good quality models with high C scores^{46,47} were selected for further analysis.

During insect herbivory there are lots of elicitor molecules that are released from the oral secretions of insects, however, for this study we have selected highly cited elicitor molecules i.e., volicitin, caeliferin,

bruchin and inceptin. Volicitin (N-(17-hydroxylinolenoyl)-l-Gln) a small fatty acid derived molecule was first reported in the corn seedling wounds caused by beet armyworm feeding¹⁹ In the current study, findings show a strong interaction between plant elicitor peptides and their receptors with hydrogen bond distance less than 3Å which reflects a strong bonding⁴⁸ between insect oral secretion chemical compounds and plant elicitors, Systemin and PEP1. This confirmed the findings that insect oral secretions contain the compounds which can induce defense responses of plants and cause higher levels of JA production as compared only to mechanical damage^{40,49}. Our results confirmed previous findings as reported by Chen & Mao⁴ who reported that elicitors derived from insects can activate plant defense by interacting with plant elicitors Systemin and PEP1. The receptors of these elicitors present on the epithelial cells are SYR1, LRR PEPR1 upon receiving signals they initiate a cascade of plant defense signaling network. After this several changes occur in a plant cell all these changes consider as early plant defense responses by the production of Jasmonic acid⁵⁰. These changes activate jasmonate acid (JA) production in the plant. Afterward, certain changes occur in the peroxisome of a plant cell. All these changes in plant cells activate the OPDA pathway. Jasmonic acid and its primary precursor fatty acid derived 12-oxophytodienoic acid (OPDA) enzymes are the most studied oxylipins⁵¹. These OPDA enzymes, include Hydroperoxide lyase and Allene oxide synthase, play the role of key mediator in the production pathway of green leafy volatiles and jasmonic acid respectively. Molecular docking of AOS and HPL, with their ligands 9(S) and 13(S) hydroperoxides⁵², was done through AutoDock Vina and analyzed the results by using LIGPLOT. Both AOS and HPL showed good interaction with the ligands. Allene oxide synthase belongs to the cytochrome P450 family of enzymes, interacts with 9- or 13-hydroperoxides and act as catalytic agent in jasmonate production in plant cell chloroplast³⁷ hence proved. Hydroperoxide lyase another important enzyme catalyzes the production of GLVs by using the same substrates⁵³. Our finding confirmed that not only AOS but HPL interact with 9(S) and 13(S) hydroperoxides to mediate the Jasmonic acid synthesis pathway. JA and JA-Ile induced by systemin act as signals and are transported to adjacent sites for defensive responses⁵⁴. The Jasmonic acid shifts towards cytoplasm then converted to JA-Ile (Jasmonate Isoleucine) and interact with nuclear receptor COI1 (Coronatine Insensitive Protein 1). This interaction causes the degradation of JAZ protein that holds transcription factor MYC2. this induces downstream gene expression for defense protein formation⁵⁵. Our results show a strong interaction between JA-Ile and COI1 with three hydrogen bonds at a bond distance of less than 4Å. The MYC transcription factors by bind to JAZ protein and activates JA-Ile-COI1 complex and degrade of JAZ with the help of 26S proteasome³⁹. Three dimensional structure COI1 and JAZ complex is evident of its high binding affinity with JA-Ile compound⁵⁶. We docked the COI1-JA-Ile complex with JAZ and predicted strong interaction showing five hydrogen bonds each with less than 4Å distance. Seven hydrogen bonds were formed and a salt bridge bond was also found. The salt bridge most often arises from the anionic carboxylate (RCOO⁻) of either aspartic acid or glutamic acid and the cationic ammonium (RNH₃⁺) from lysine or the guanidinium (RNHC(NH₂)₂⁺) of arginine⁵⁷. The bond distance required for good interaction must be ≤ 4 Å (400 pm)⁵⁸.

At the end Jasmonic acid activate defense responsive genes and the plant shows the JA mediated response. the F-box COI1 protein, which functions as part of the JA receptor, interacts with the Skp1/Cullin counterparts to form the Skp1–Cullin–F-box (SCF) COI1 ubiquitin E3 ligase complex and interacts physically with JAZ repressor proteins in the presence of JA conjugated with the amino acid isoleucine and convert to jasmonoyl-L-isoleucine (JA-Ile). This is an active form of JA⁵⁹ which switch on the defense related plant genes^{60,61}.

The study of Arabidopsis AOS protein-protein interaction mechanism and structure elucidation has been reported before by Lee et al. ^{62,63}. The Protein-Protein interaction of AOS and HPL was done by using the STRING database to further confirm the interaction partners of these proteins in the JAS production pathway. The biosynthesis of JA starts with -linolenic acid which is converted to its 13(S) - hydroperoxylinolenic acid by 13-LOX; the LOX product is subsequently converted to jasmonic acid by sequential action of allene oxide synthase (AOS), allene oxide cyclase (AOC) and 12-oxo-phytodienoic acid reductase (OPR) while HPL and ADH2 lead to the biosynthesis of GLV ⁶⁴⁻⁶⁷. The results confirm the above findings. Both AOS and HPL had shown an interaction between each other and showed common interaction partners of the Lipoxygenase family i.e., loxC, Solyc09g0559002.1, 101263024 and loxc. AOS also showed interaction partner AOC, COI1 and OPR1. Lipoxygenases (lox) are a ubiquitous family of non-heme iron enzymes widely distributed in plants, initiate hydroperoxidation of polyunsaturated fatty acids and produce phytooxylipins (jasmonic acid and GLVs) responsible for physiological processes like seed germination, fruit ripening, senescence and defense (against biotic and abiotic stress) ⁶⁸.

HPL shows interaction with ADH2 (Alcohol dehydrogenase). Alcohol dehydrogenases (ADHs) belong to the dehydrogenase superfamily ⁶⁹. GLVs originate in the hydroperoxide lyase (HPL) branch of the oxylipin pathway, with the help of ADH2 ⁷⁰⁻⁷². Which is a crucial plant defense signal of in result of herbivory^{73, 74}.

The pairwise alignment of AOS and HPL showed 39.5% identity and 60.7% similarity. Structural superposition showed a similar trend with 37. 5% identity these results confirm that although AOS (CYP74A) and HPL (CYP74B) are closely related members of the CYP74 family and interact with the same set of substrates (13-h) yet result in diverse products. Our findings confirmed the information given by Tyagi et al.⁷⁵. They found the F amino acid plays important role in binding with 13- hydroxy peroxide for the production of jasmonic acid. They also observed that the F in AOS sequence (involved in binding with substrate) is replaced by L in HPL of Japanese rice. In our study we found to F amino acids in AOS were found to interact with 13-hydroxy peroxide but no F was involved in HPL-13 hydroxy peroxide interaction. The HPL also showed interaction with ADH2 which leads to producing GLVs but commonly interacting with lox.

Our findings suggest that these two branches (AOS and HPL) of the oxylipin pathway exhibit crosstalk with regards to biosynthesis and signaling and cooperate to function in multiple abiotic stress responses⁷⁶

Materials And Methods

Selection and retrieval of Insect/plant elicitors and receptors

Insects release various chemical compounds, from their oral secretions (regurgitate and saliva) during herbivory which activate defense mechanisms in plants. We have selected highly cited elicitor molecules involved in plant defense pathway activation. The structures of selected molecules were downloaded in sdf (structure-data file) format through PubChem and converted to .pdb format through PyMol (Table 1). Inceptin is a proteolytic fragment produced by insect larvae that previously ingested the plant protein and mediates plant perception of herbivory. This is categorized both as insect/plant elicitor. The amino acid sequences of some plant elicitors (and inceptin) and receptors were retrieved from UniProt (Table 2).

Protein 3D Structure Prediction

Three-dimensional (3D) structures of the proteins not available in PDB were predicted through homology and *Ab-initio* methods. Protein BLAST of the protein sequences was done through NCBI BLASTp (selecting PDB dataset) to get good templates for use in homology modeling. The 3D structures of proteins, having more than 35% identical templates, were modeled by homology modeling. Homology modeling was done by using an offline tool Modeller 9.1. The protein which could not result in an identical template (< 35%) were predicted through *Ab-initio* based structure modeling method. The protein FASTA sequences were submitted to an online available server

I-TASSER for *Ab-initio* based structure modeling. ModRefiner a web-based server was used to by using *Ab-Initio* modeling³⁴. The quality of the predicted models have to be evaluated by different evaluation tools to ensure its quality. The refinement of low quality structures was done Galxyweb server. The refined models were submitted to the SAVES server, which uses different sets of structure evaluation platforms and qualified models were selected.

Docking Analysis

Docking analysis of protein-ligand interaction was done by using Auto Dock Vina while protein-protein docking was done by using the online available server Patch Dock. The strongest binders were identified based on lower binding energy and hydrogen bonding.

Protein-ligand with Auto Dock Vina

Auto Dock Vina is an offline sever it has GUI-based MGL tools, ligand and protein preparations and a Command line-based interface for the execution of molecular docking. First of all, the pdb files of ligand and protein files were subject to charges addition and converted to .pdbqt format. Grid was set for blind docking then a configuration file was prepared to run in the command line for docking. Five models were generated and the model with the lowest binding energy and RMSD value was selected. All the models were deleted from the output.pdbqt file except the selected one. This output.pdbqt file was used for interactive visualization.

Protein-Protein docking with Patch Dock

Protein-Protein docking was done by an online available server Patch Dock. It works on a molecular docking algorithm based on shape complementarity principles³⁵. Three dimensional models of proteins (.pdb) to be docked were uploaded to server. The results were received via email in the form of the compressed folder of the top 20 selected complexes. The docked complexes were subjected for refinement through the online available server Fast Interaction REfinement in molecular DOCKing (FireDock). The protein-ligand and receptor pdb files and a Patch Dock generated, Tarsansformations.txt file was uploaded to the server for refinement. The resulting best-scored docked complex was selected based on lower binding energy.

Docking result visualization

Complex. pdb files were uploaded to LIGPLOT + to visualize interactions in the docked complex. Protein-ligand was visualized by selecting the option LIGPLOT while Protein-Protein interaction was visualized by selecting option DIMPLOT. The 2-D structure of the complex was generated showing hydrogen bonding and hydrophobic interactions. BIOVA DISCOVERY STUDIO®³⁶ was used for 3D visualization of a docked protein complex.

Jasmonate biosynthesis pathway

The interactions of insect elicitors with plant receptors activates different plant defense through different pathways. We selected the Octadecanoid pathway, which produces jasmonic acid (JA), for our study. JA is the main regulator of the plant defense mechanism.

AOS (Allene oxide synthase) and HPL (Hydroxyperoxide lyase) are the two main enzymes of jasmonic acid (JA) and Green leafy volatiles (GLV) production pathway. AOS and HPL were selected to study how they are involved in Jasmonic acid production. The amino acid sequences of both enzymes were obtained from UniProt under Q9LLB0 (AOS) and K4CF70 (HPL). Homology-based modeling was done using Modeler 9.1.

Interaction of AOS and HPL with their ligand molecules

It was reported in the literature that Allene oxide synthase (AOS) catalyzes the first step of jasmonate biosynthesis in the chloroplast by interacting with 9 and 13- hydroxyperoxides³⁷. The molecules of 9 and 13- hydroxyperoxides were selected as ligands for AOS and HPL. Protein-ligand docking was performed by using AutoDock Vina and protein-ligand interaction was visualized from LIGPLOT.

Jasmonate signaling and regulation

Jasmonic acid (JA) is generated in the peroxisome and transferred into cytoplasm. In cytoplasm, the active signal in defense signaling appears to be the amide-linked isoleucine conjugate JA-Ile rather than JA itself. This JA-Ile promotes the interaction of Coronatine Insensitive Protein 1 (COI1) and Jasmonate ZIM domain (JAZ) protein. Afterwards, 26S proteasome perform the ubiquitination of JAZ proteins and expression of downstream genes are activated by specific transcription factors.

Interaction of COI1 with JA-Ile

Coronatine Insensitive Protein 1 (COI1) is a nucleus receptor that binds with the JA-Ile (Jasmonate Isoleucine) to degrade JAZ protein. The 2D structure of JA-Ile was obtained from PubChem while the 3D structure of COI1 was downloaded from RCSB PDB (PDB ID: 3OGM). 3OGM is the Structure of COI1-ASK1 in complex with coronatine and the JAZ1 degnon. Chain B of the complex is COI1 and it was cut by using Chimera 1.8. The ligand JA-Ile was docked with COI1 by Auto Dock Vina.

Interaction of COI1 receptor with JAZ protein

The 3D structure of the JAZ binding domain was downloaded from RCSB PDB (PDB ID: 4RS9) which is a structure of MYC3 N-terminal JAZ-binding domain complex with Jas motif of JAZ9. The chain B, JAZ degnon, was cut and retrieved by Chimera 1.8.

The docked COI1-JA-Ile complex was docked with JAZ by submitting to the Patch Dock server and complexes were refined through FireDock. The best-docked complex was downloaded and 2D interactions were visualized through DIMPLOT while for 3D complex visualization BIOVA DISCOVERY STUDIO®

Protein-Protein interactions

Protein-Protein interaction networks³⁸ were used to understand biological processes in organisms. Protein-Protein interaction was explored, from protein-protein interaction database STRING, to find out the interactions of the target protein to other proteins which might be selected for alternate target site in future.

Protein sequence and structure alignment

Hydroperoxide lyase (HPL) and Allene oxide synthase (AOS) are involved in a same pathway for almost same functions that's why they are thought to be closely related. The pairwise sequence alignment was done by the EMBL_EBI Pairwise sequence alignment tool. Three-dimensional structure superposition was by Matchmaker of Chimera 1.8.

Declarations

Conflict of interest

The authors declared no conflict of interest

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Tables

Due to technical limitation the tables are available as a download in the Supplementary Files.

Figures

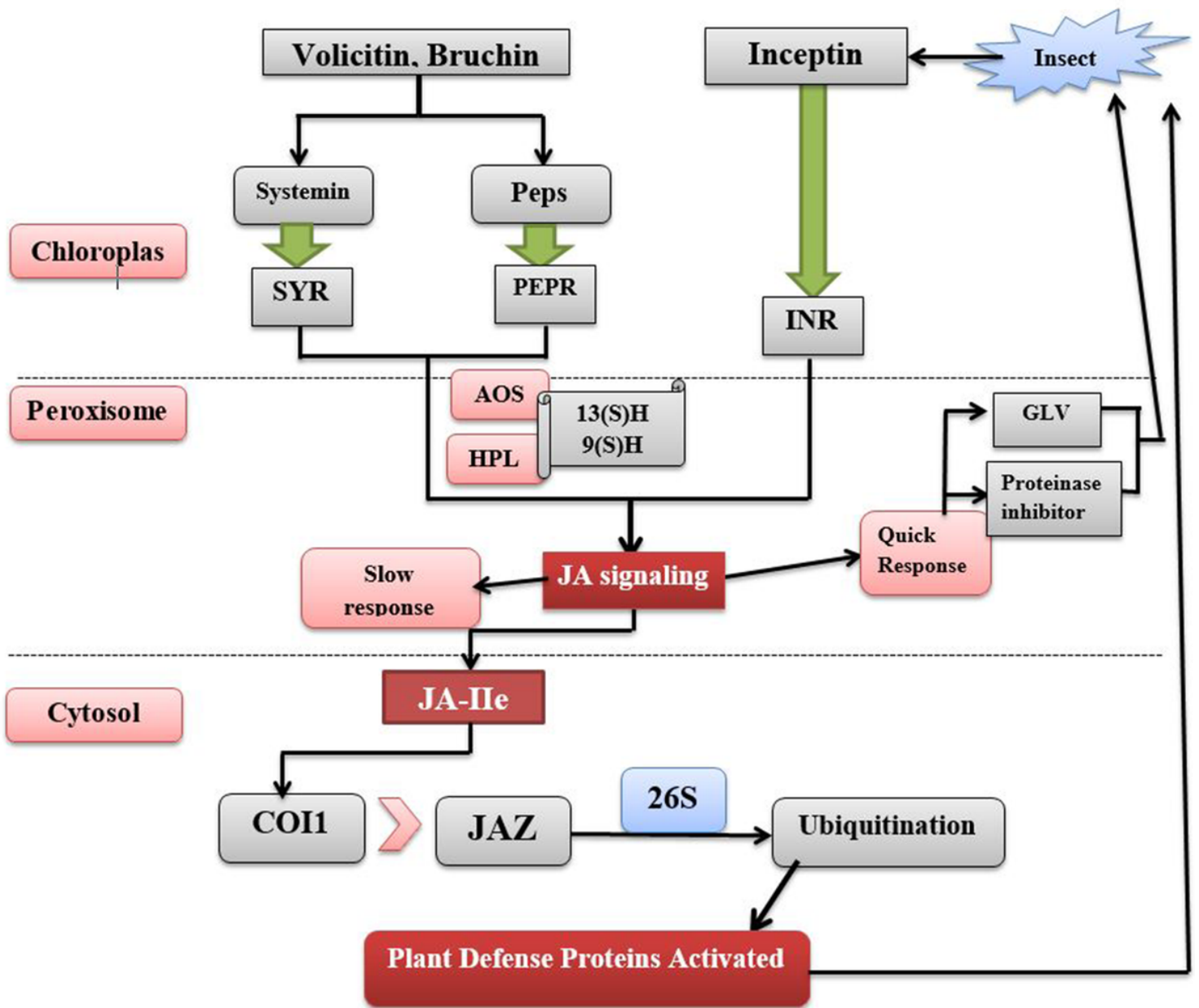


Figure 1

Herbivory induced plant defense pathway

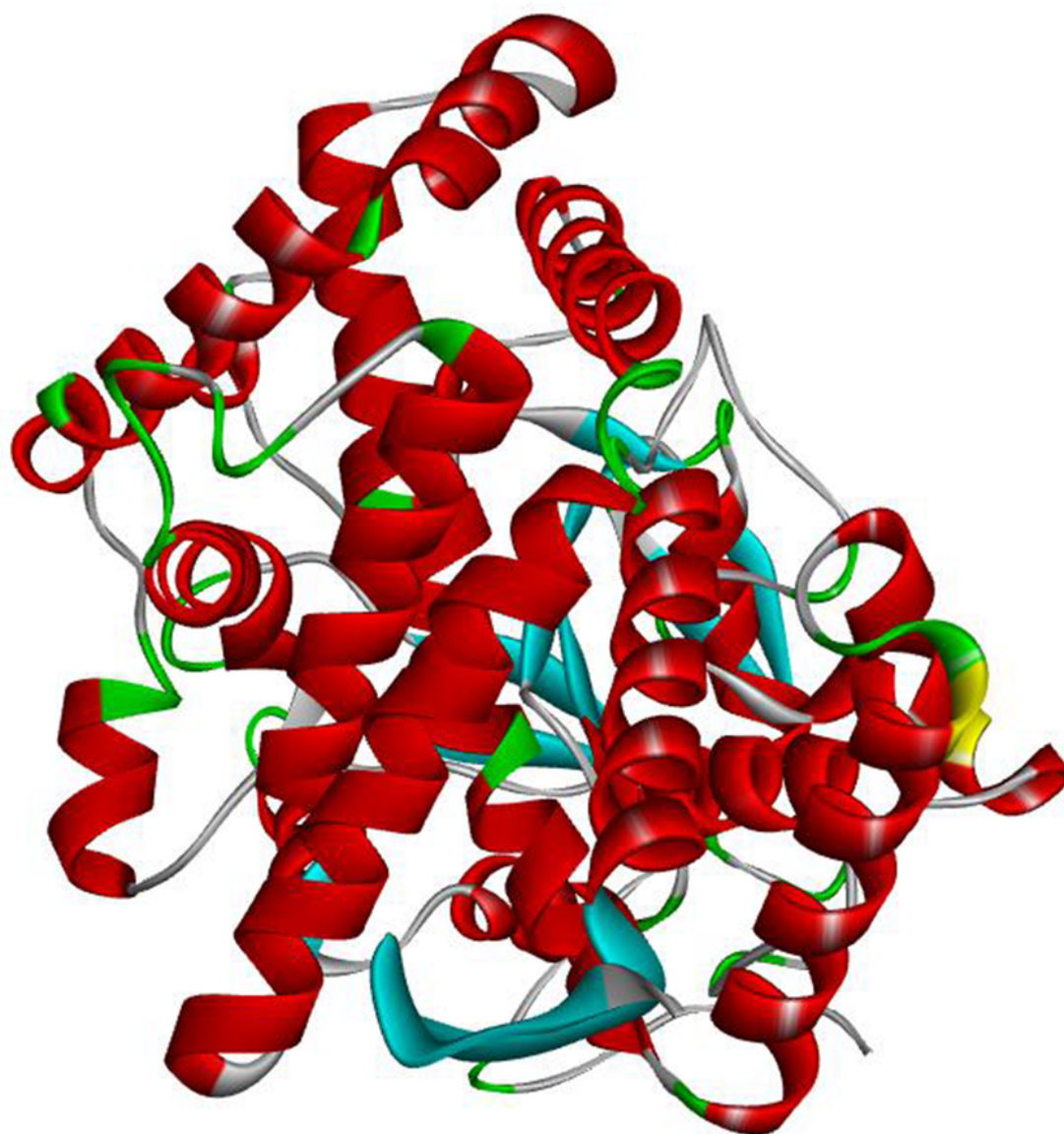


Figure 2

Predicted 3D structure of allene oxide synthase (AOS) of *Solanum lycopersicum*

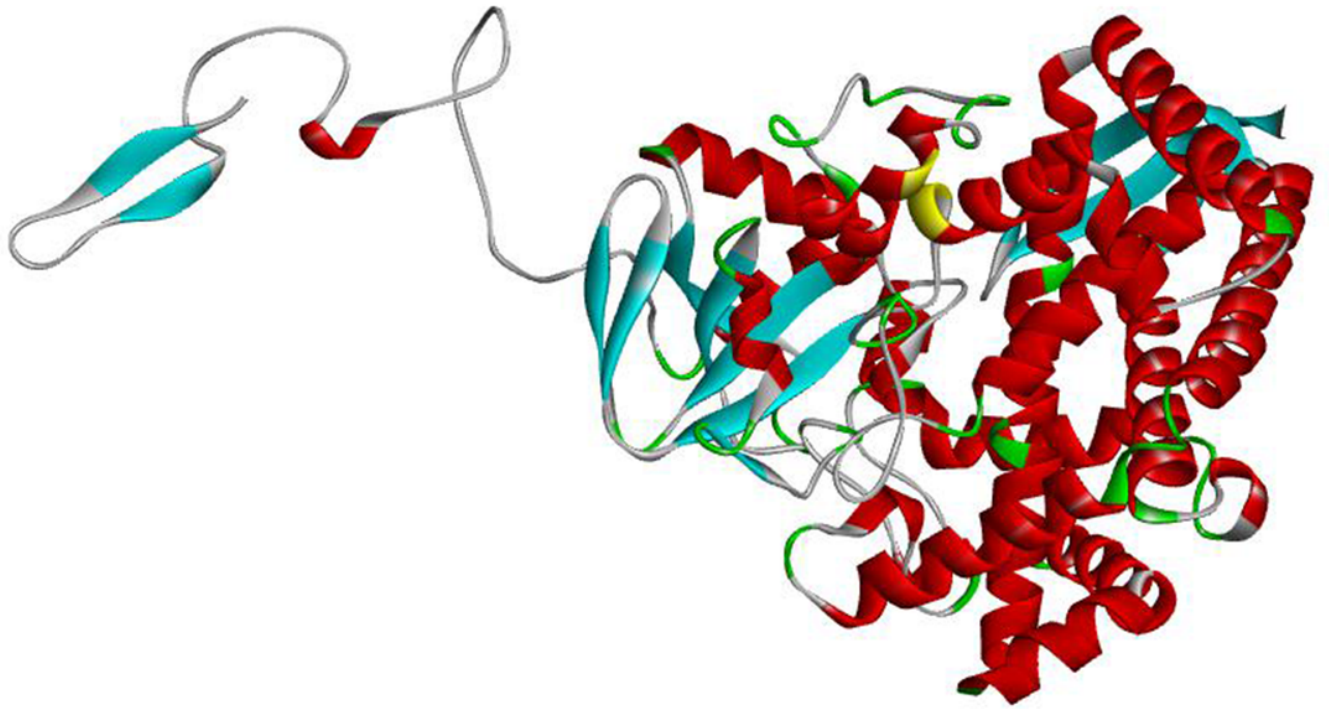


Figure 3

Predicted 3D structure of hydroxperoxide lyase (HPL) of Solanum lycopersicum

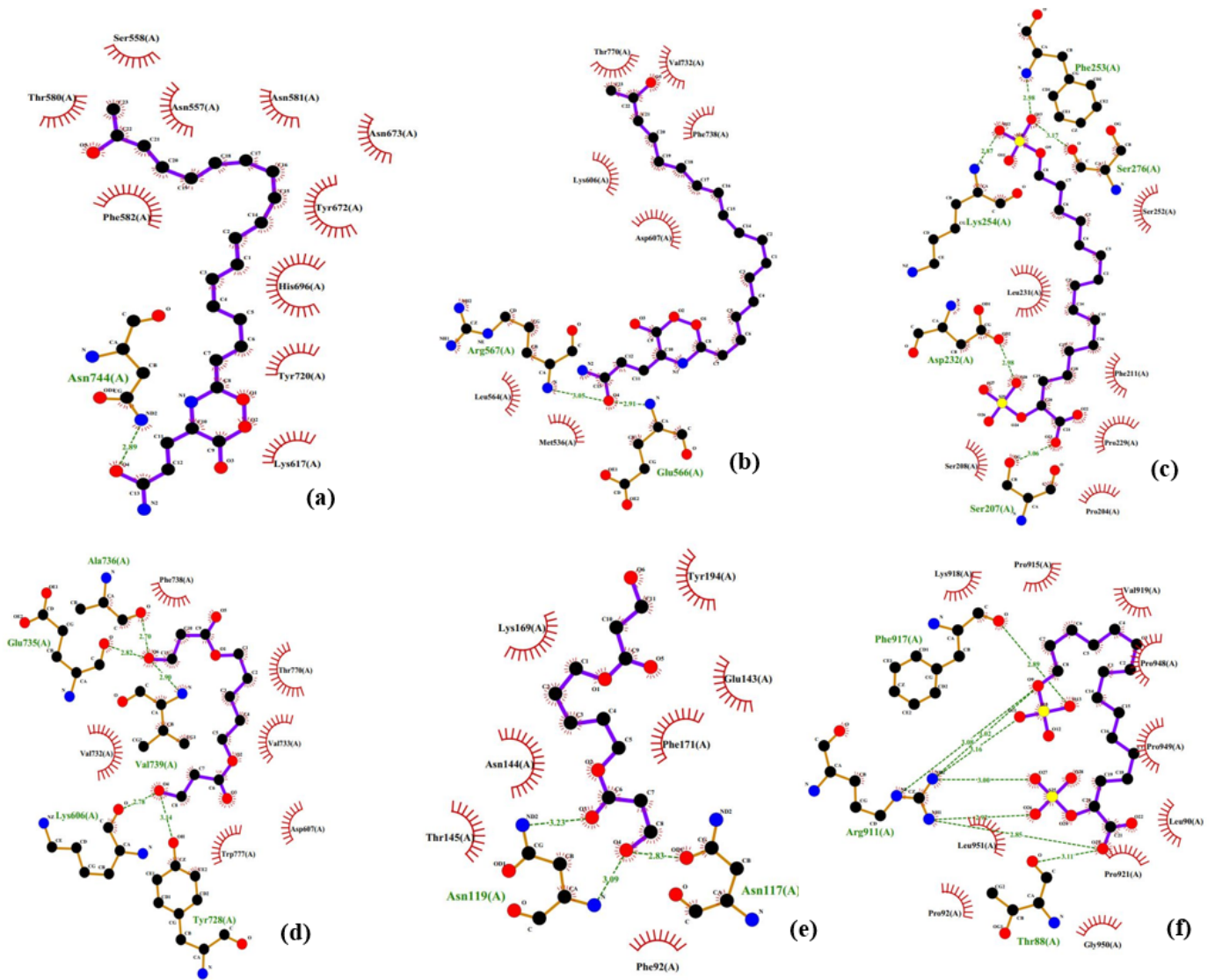


Figure 4

Protein-ligand 2D interaction map a. Systemin-voliticin b. PEP-voliticin c. Systemin-caeliferin d. PEP-bruchin e. Systemin-bruchin f. PEP-caeliferin

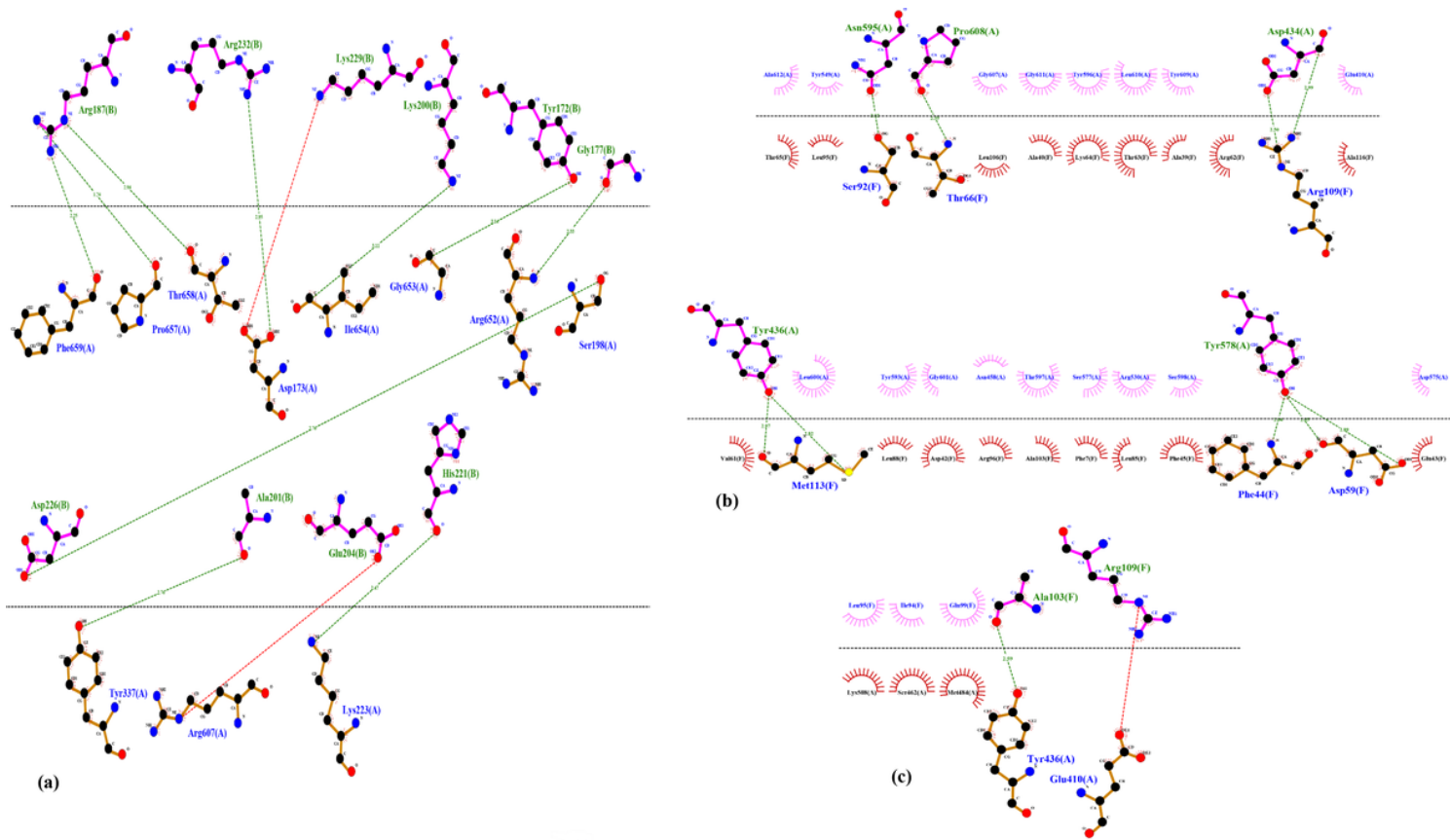


Figure 5

Protein-protein 2D interaction map a. Systemin-SYR b. PEP-PEPR c. Inceptin-LRR

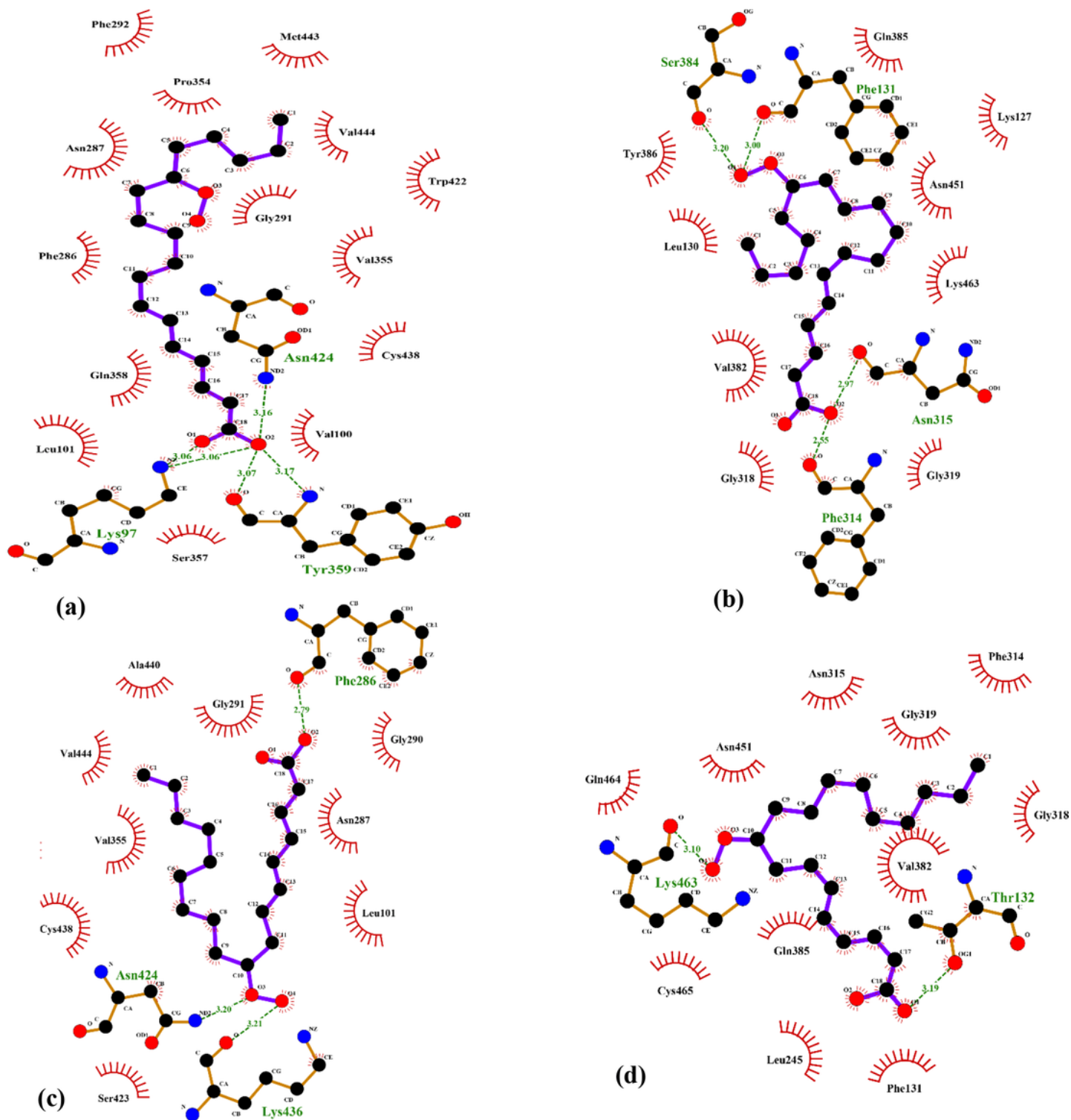


Figure 6

2D interaction map for Protein-ligand in jasmonate biosynthesis pathway a.13(S) Hydroperoxide-HPL complex b. 13(S) Hydroperoxide-AOS complex c. 9(S) Hydroperoxide-HPL complex d. 9(S) Hydroperoxide-AOS complex

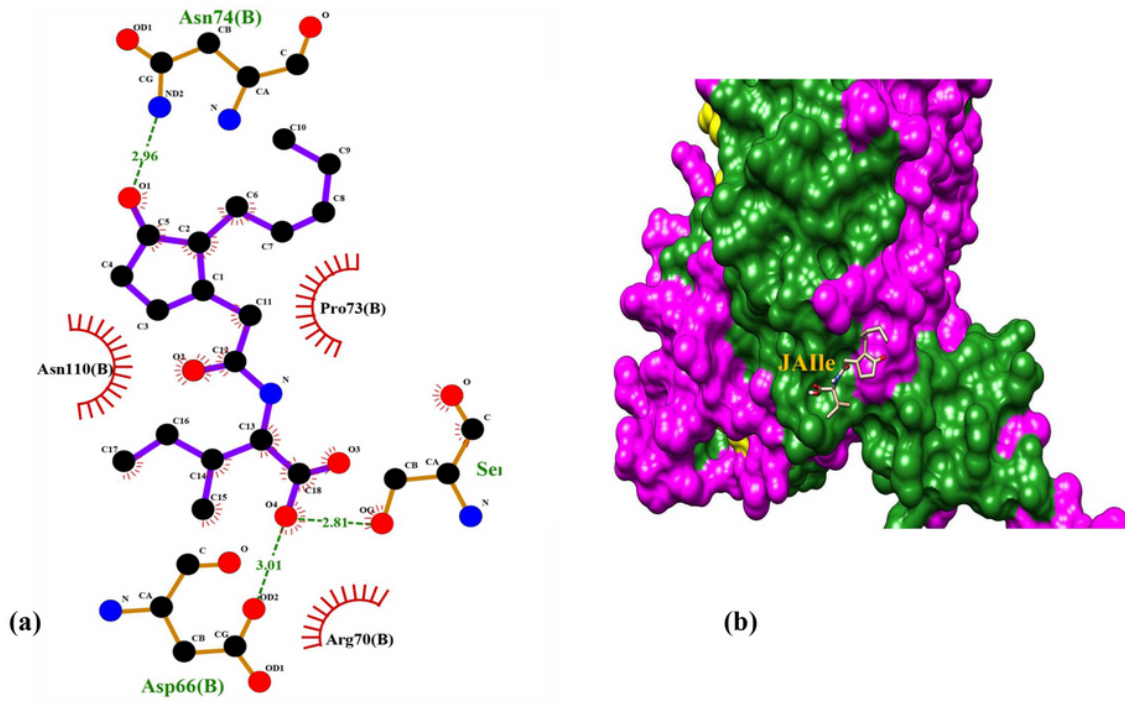


Figure 7

Interaction of COI1 with JA-Ile in Jasmonic acid induce plant defense mechanism a. 2 D interaction plot b. 3D complex

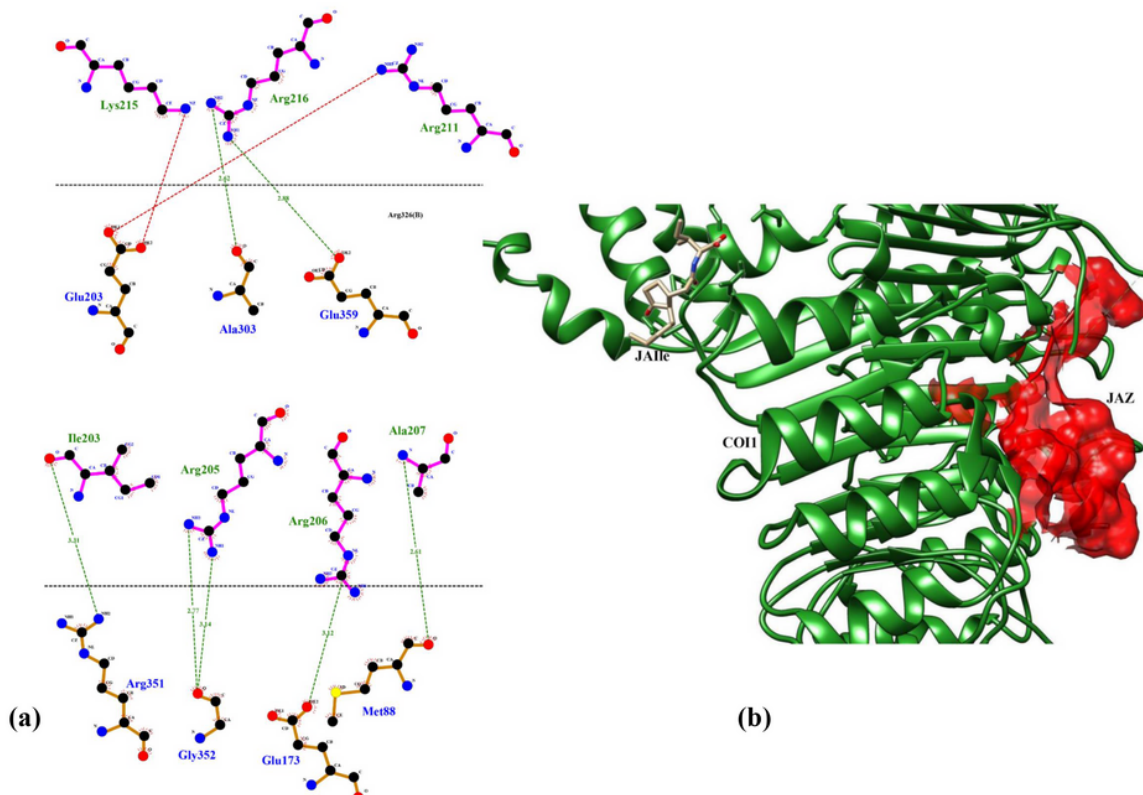


Figure 8

Interaction of COI1- JA-Ile complex with JAZ protein degron in Jasmonic acid induce plant defense mechanism a. 2 D interaction plot b. 3D complex

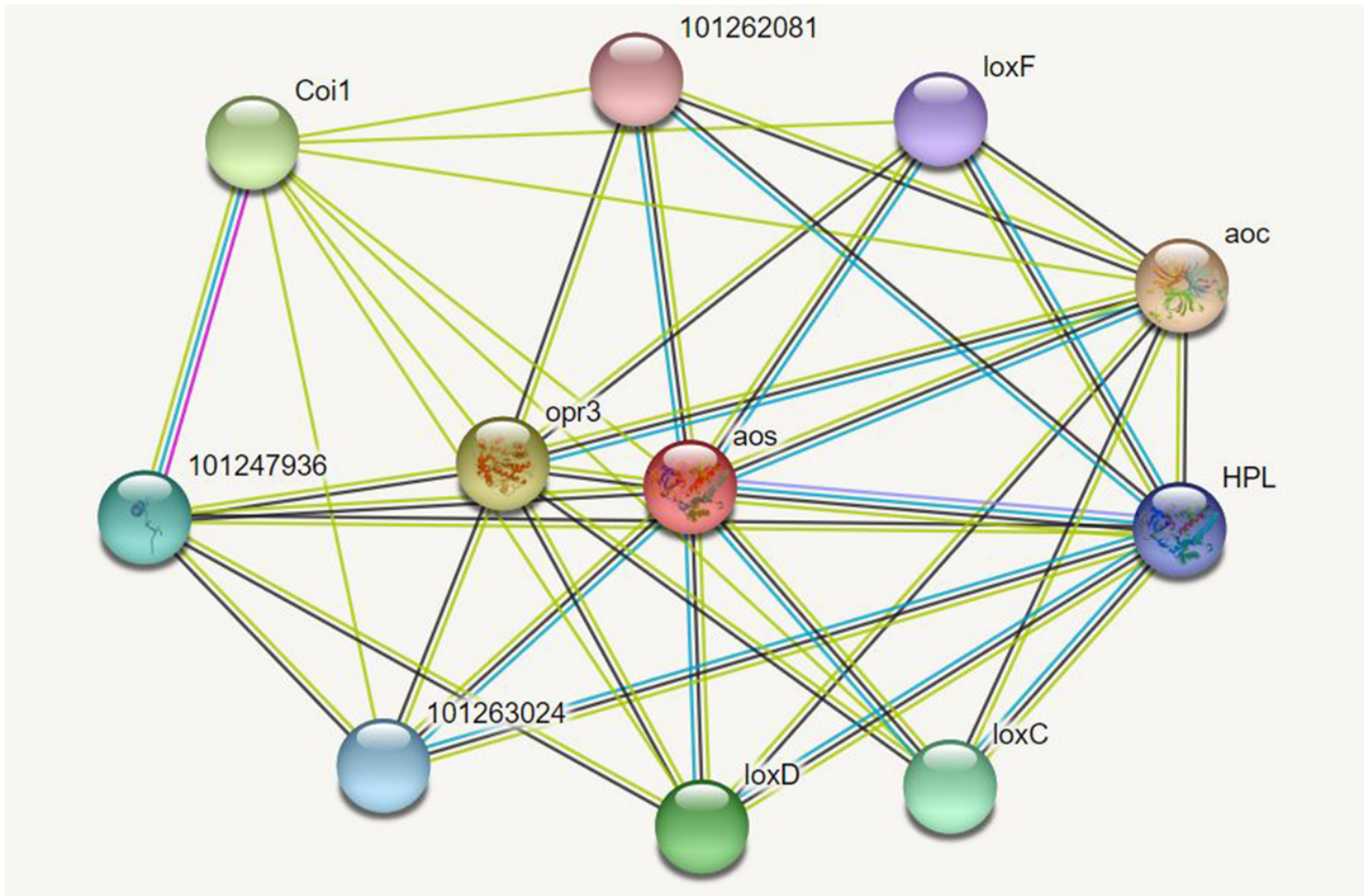


Figure 9

Protein interaction partners of allene oxide synthase (AOS) of *Solanum lycopersicum*

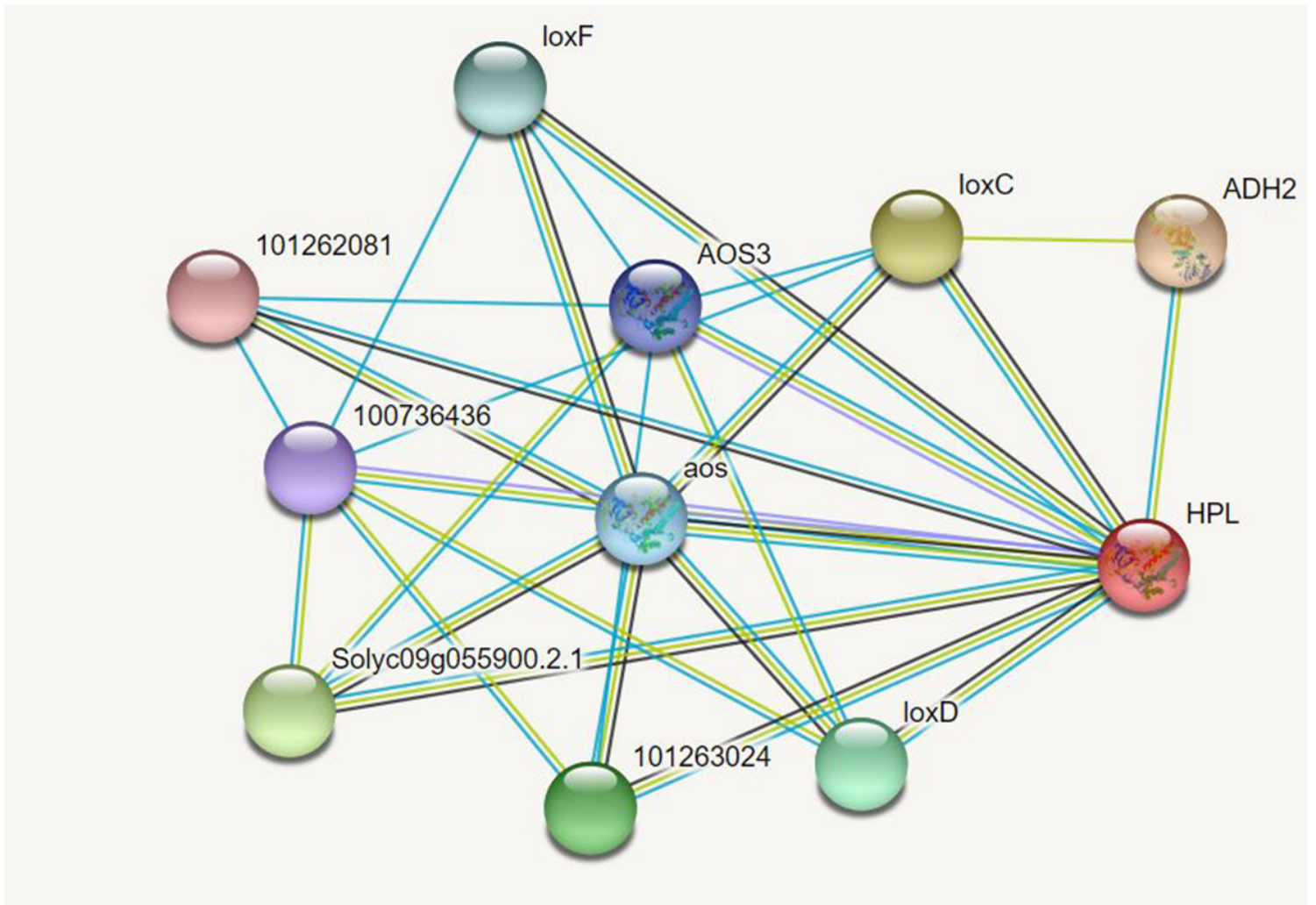


Figure 10

Protein interaction partners of hydroxyperoxide lyase (HPL) of *Solanum lycopersicum*

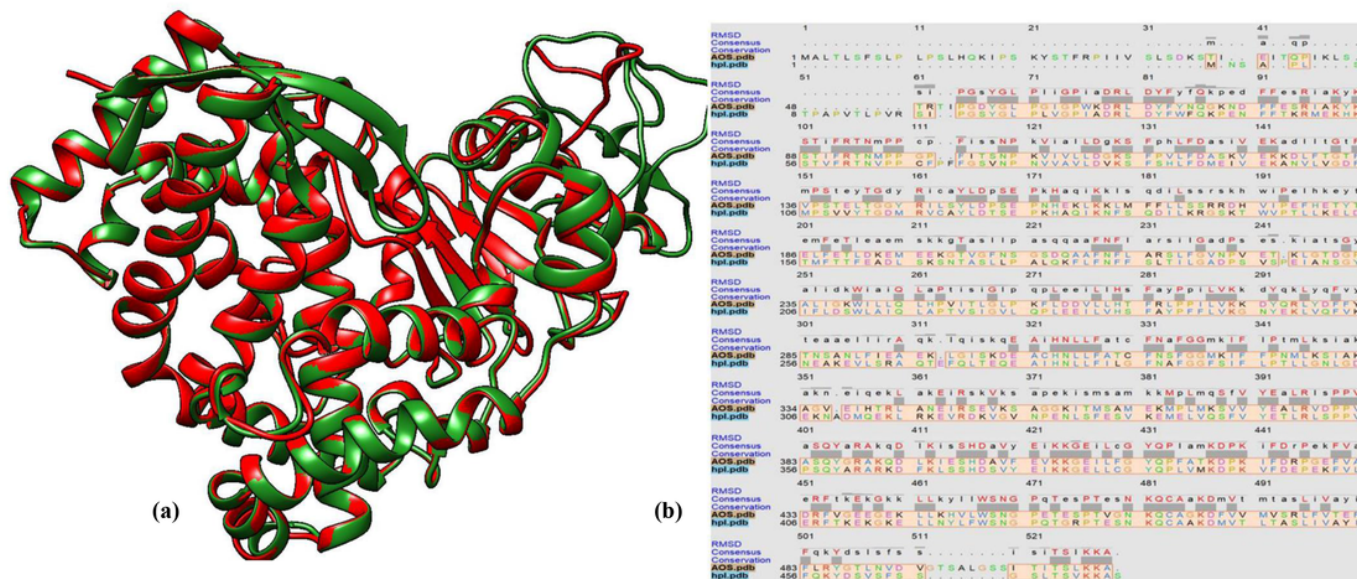


Figure 12

Three dimensional structure superposition of allene oxide synthase (AOS) and hydroxyperoxide lyase (HPL) of *Solanum lycopersicum* a. 3D superposed complex b. pairwise sequence alignment during structure superposition.

Supplementary Files

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