**Article**

**Cytochrome P450 Genes Expressed in Phasmatodea Midguts**

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**Simple Summary:** Cytochrome P450s are a group of detoxification enzymes found in all animals. They are highly diverse, with multiple copies of different families of these genes in individual organisms, especially those that need to digest chemically-defended foods, such as toxic plants. The stick insects are a group of leaf-feeding herbivores whose diets can include toxic species, such as *Eucalyptus*, so this study sought to identify what groups of cytochrome P450s are expressed in the digestive tracts of six species of stick insects and how they evolved. The results show a complement of detoxification genes comparable to or slightly more limited than those of other insects, except for multiple enzymes per species in the subfamily *CYP15A1*, while most insects will only have one enzyme in this group. The functions of these expanded enzyme groups should be investigated further.

**Abstract:** Cytochrome P450s (CYPs) are xenobiotic detoxification genes found in most eukaryotes, and linked in insects to the tolerance of plant secondary chemicals and insecticide resistance. The number and diversity of CYP clans, families, and subfamilies that an organism produces could correlate with its dietary breadth or specialization. This study examined the CYP diversity expressed in the midguts of six species of folivorous stick insects (Phasmatodea), to identify their CYP complement and see if any CYPs correlate with diet toxicity or specialization, and see what factors influenced their evolution in this insect order. CYP genes were mined from six published Phasmatodea transcriptomes and analyzed phylogenetically. The Phasmatodea CYP complement resembles that of other insects, though with relatively low numbers, and with significant expansions in the CYP clades 6J1, 6A13/14, 4C1, and 15A1. The CYP6 group is known to be the dominant CYP family in insects, but most insects have no more than one CYP15 gene, so the function of the multiple CYP15A1 genes in Phasmatodea is unknown, with neofunctionalization following gene duplication hypothesized. No correlation was found between CYPs and diet specialization or toxicity, with some CYP clades expanding within the Phasmatodea and others likely inherited from a common ancestor.

**Keywords:** cytochrome P450; Phasmatodea; stick insects; xenobiotic metabolism

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**1. Introduction**

Many plants are chemically defended against herbivores, producing directly toxic or anti-nutritive chemicals. Different herbivores use different strategies to overcome these. Some have symbiotic microbes that can metabolize and deactivate ingested toxins, while others produce their own endogenous detoxification compounds, and some have both [1]. Xenobiotic detoxification proteins in insects include glutathione-S-transferases (GSTs) [2], carboxy/cholinesterases (CEs) and cytochrome P450 monoxygenases (CYPs). The number of such genes in different insect species varies, and they are often expressed in different tissues performing different functions [3] and/or in different life stages of the host [4]. These detoxification genes are linked to insecticide resistance [5], and so are of significant interest in applied entomology and other industries [6], yet are also informative regarding the evolution of dietary specification. Organisms that specialize in a particularly toxic plant would need to have evolved a powerful detoxification method in order to maintain that niche. Conversely, organisms with broad host and/or geographic ranges may need...
expanded xenobiotic metabolism genes to cover the wider variety of compounds they may be exposed to [7].

The Cytochrome P450s are found in all animals and plants and most microbes. Their high diversity in Metazoa is thought to be a direct result of herbivore–plant warfare [8], and the CYPs are notorious for high levels of gene duplication, transposition, neofunctionalization, and pseudogene development [9,10]. The present nomenclature for CYPs is CYP#X#, with the first numeral for the CYP family, the letter for the subfamily, and the second numeral for the gene [11]. By convention, a CYP family is defined as sharing > 40% sequence similarity along its members’ approximately 500 amino acids [12]. CYPs are found in all insect tissues. Above the family level, insect CYPs are sorted into four clans: the mitochondrial clan, clan 2, clan 3, and clan 4, with clan 3 considered most important for xenobiotic regulation while the others are involved in conserved physiological roles such as hormone synthesis [3,11,13]. The mitochondrial clan is nuclear-encoded and named because of their mitochondria-targeting sequences [14]. The number of CYP genes in insects varies: some parasitoids have less than 30 [5], while mosquitoes have over 200 [13,15]. Those expressed in the midgut, Malpighian tubules, and fat body are integral for plant chemical detoxification and insecticide resistance [16,17], as confirmed by heterologous expression and transcriptomics studies [4,12].

The Phasmatodea, or leaf and stick insects, are an order of obligatorily herbivorous insects in the Polynoptera clade. They are not known to have microbial symbionts and process their food with endogenous digestive and detoxification proteins [18,19]. Several are of economic significance [20]. Graeffea crouanii (Le Guillou 1841) is a pest of coconut plantations in the south Pacific [21]. Outbreaks of Diapheromera femorata (Say 1924) (Diapheromeridae) in the USA can cause severe forest defoliation [22]. In Australia, plague-like outbreaks of Didymuria violescens Leach 1815 (Phasmatidae), Anchiale austrotesellata Brock & Hasenpusch 2007 (Phasmatidae), and Podacanthus wilkinsoni Macleay 1882 (Phasmatidae) can cause severe defoliation to Eucalyptus L’Her. trees [23]. The latter is particularly remarkable given how toxic Eucalyptus is [24]. Understanding how these species evolved the ability to metabolize, eliminate, or otherwise endure Eucalyptus secondary compounds has basic and applied science significance.

This study compared the CYPs expressed in the digestive tracts of six different Phasmatodea species, representing four families and five subfamilies and fed three different species of the host plant, to understand how CYPs evolved within Phasmatodea and attempt to identify CYP families that may play important roles in plant chemical detoxification.

2. Materials and Methods

This study uses published Phasmatodea transcriptome data (GenBank Accession numbers PRJNA238833 and PRJNA221630) from the physiologically distinct anterior and posterior midguts [19] of Arctaon asperrimus (Redtenbacher 1906) (Heteropterygidae: Obriminae), Extatosoma tiaratum (Macleay 1826) (Phasmatidae: Extatosomatinae), Medauroidea extradentata Brunner von Wattenwyl 1907 (Phasmatidae: Cliummineae), Peruphasma schultei Conle & Hennemann 2005 (Pseudophasmatidae), Rhamulus artemis (Westwood 1859) (Phasmatidae: Cliummineae), and the anterior midgut only of Sipyloidea sipylus (Westwood 1859) (Lonchodidae) [25]. The Extatosoma were reared on Eucalyptus sp., the Peruphasma were reared on privet (Ligustrum sp. L.), and the others fed on rose leaves (Rosa sp. L.).

In June 2021, representative sequences for insect cytochrome P450s were downloaded from NCBI, limiting the search to those in the UniProtKB database [26,27]. The resulting 111 sequences were used as a query to mine the above transcriptomes using tblastn [28] with an expected value threshold of e−10. These were manually annotated by removing truncated sequences, using the ExpASy online translation tool [29] to obtain the complete amino acid sequences, removing duplicates using the sRNA toolbox webserver [30], and confirming that the sequences were cytochrome P450s by identifying them using blastp against the NCBI database. This search also enabled the identification of the sequences to the CYP family: according to standard CYP nomenclature, sequences with more than 40% identity
are in the same family [31]. The resulting sequences were combined with the representatives from NCBI, aligned using the Clustal W program [32] built into the software MEGA version X [33]. Any sequences missing the heme-binding domain FXGXXXCGA/A [34], which is a signature motif for CYPs [6], were deleted. The sequences were also checked for the presence and absence of other four signature motifs from insect CYPs: helix C (WxxxR), helix I (GxE/DTT/S), helix K (ExLR), and PERF (PxxFxPE/DRE/F) [35]. Fasta files for the amino acid sequences of these putative CYPs from the six transcriptomes were uploaded to Zenodo (DOI: 10.5281/zenodo.7049585).

All validated Phasmatodea species’ CYPs and select insect CYPs from the NCBI/UniProt databases were aligned using Clustal Omega [36], and phylogenetic analysis was done using the neighbor-joining method [37] with 10,000 bootstraps in MEGA version X [33], to confirm and if necessary correct the identities assigned to each sequence via blastp. This tree would also show if Phasmatodea genes that have multiple copies in one species also have multiple copies in another, which could help time any gene duplication or expansion events across Phasmatodea evolutionary history. Phylogenetic tree inference using maximum likelihood with rapid bootstrapping was made using the IQ-TREE web server [38] with 1000 ultrafast bootstrapping replicates [39] and the best-fitting model LG+I+G4 as identified using ModelFinder [40]. The tree was visualized using FigTree v1.4.4.

3. Results

Table 1 lists the number of isogroups (comp#_c#) and isotigs (comp#_c#_seq#, or individual transcript sequences) of each identifiable CYP family in each of the Phasmatodea, limiting the results to non-truncated sequences. All valid sequences containing the heme-binding domain had a blastp sequence similarity > 40% to a known CYP subfamily; however, some sequences were equally similar to known 6A13 and 6A14 genes, or to the 4C1 and 4C3 genes. Most CYP families had a single gene per species, while a few were highly expanded: 6J1, 6A13/14, 4C1, and 15A1. Medauroidea had the largest number of CYP transcripts, followed by the other two Phasmatidae species, Ramulus and Extatosoma, largely due to isogroup and isotig expansions in the 6J1 family. Peruphasma also had many 6J1 isotigs, but few of the others. Aretaon had comparatively many 15A1 transcripts but the fewest 6J1. Sipyloidea had the fewest isotigs and tied with Peruphasma for the fewest isogroups. The Sipyloidea transcriptome had zero representatives from the mitochondrial clan, though Peruphasma and Extatosoma each only had two.

| CYP450 Clan | Family | Aa | Et | Me | Ps | Ra | Ss * |
|-------------|--------|----|----|----|----|----|-----|
| Mito. Clan  | 44     | 1  | 0  | 1  | 1  | 1  | 0   |
|             | 49A1   | 1  | 1  | 0  | 0  | 1  | 0   |
|             | 302A1  | 1  | 0  | 1  | 0  | 0  | 0   |
|             | 314A1  | 1  | 1  | 1  | 1  | 1  | 0   |
|             | 315A1  | 1  | 0  | 0  | 0  | 1  | 0   |
| Clan 2      | 304A1  | 1  | 1  | 1  | 0  | 1  | 1   |
|             | 307A1  | 1  | 0  | 0  | 0  | 1  | 1   |
|             | 15A1   | 3  | 5  | 4  | 3  | 3  | 1   |
| Clan 3      | 6J1    | 5  | 8  | 7  | 4  | 10 | 6   |
|             | 6A13/14| 5  | 4  | 4  | 3  | 7  | 4   |
|             | 6K1    | 1  | 1  | 1  | 1  | 1  | 1   |
|             | 9E2    | 1  | 1  | 1  | 1  | 0  | 1   |
| Clan 4      | 4AA1   | 0  | 1  | 2  | 0  | 0  | 0   |
|             | 4CI    | 6  | 6  | 6  | 4  | 5  | 4   |
|             | 4GI02  | 0  | 1  | 0  | 0  | 0  | 0   |

Isogroup names are based on the original transcriptome data. Values in parentheses are the number of isotigs per isogroup if >1. * Transcriptome based on the anterior midgut only, while other species’ transcriptomes are the complete anterior and posterior midgut. Aa = Aretaon asperrimus. Et = Extatosoma tiaratum. Me = Medauroidea extradentata. Ps = Peruphasma schultei. Ra = Ramulus artemis. Ss = Sipyloidea sipylus.

Table 1. Cytochrome P450s of the Phasmatodea.
Figure 1 shows the maximum likelihood tree of the representative CYP isogroups from Phasmatodea and other insects. The CYP clans form well-supported monophyletic groups, and the tree clusters according to the CYP family rather than insect taxonomy. Most isogroups had one representative from each Phasmatodea, with Peruphasma and Sipyloidea the most frequent missing species. Any isogroups expanded in one species were typically also expanded in another, with isotigs clustering according to species.

4G102 0 0 1 (1) 0 0 0

Isogroup names are based on the original transcriptome data. Values in parentheses are the number of isotigs per isogroup if >1. * Transcriptome based on the anterior midgut only, while other species’ transcriptomes are the complete anterior and posterior midgut.

Aa = Aretaon asperrimus.
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Figure 1. Maximum-likelihood phylogenetic tree of the Phasmatodea CYP450s. The evolutionary history was inferred using the Maximum likelihood method with the model LG+I+G4, identified as the best via Bayesian information criterion scores. The consensus tree with the log-likelihood \(-129,334.593\) is shown, with the sum of branch lengths 72.0129. The percentage of replicate trees in which the associated taxa clustered together in the bootstrap test (1000 replicates) are depicted as ten possible sizes of circles at the nodes, with the largest nodes indicating 100% [41]. The tree is drawn to scale, with branch lengths in the same units as those of the evolutionary distances used to infer the phylogenetic tree. The tree was edited using FigTree v 1.4.4 and Adobe Photoshop. Phasmatodea CYP
isotigs are labeled with a two-letter code for the species (Aa = Aretaon asperrimus. Et = Extatosoma tiaratum. Me = Medauroidea extradentata. Ps = Peruphasma schultei. Ra = Ramulus artemis. Ss = Sipyloidea sipylus) followed by the Trinity [42] output comp#_c#_seq#, then a “_” and the putative CYP family, subfamily, and gene in #X# format. Sequences from GenBank are labeled sp_[Genbank Accession number] | CP#x#_[UniProt organism code]. The CYP clans are labeled: Mitochondrial clan in blue, Clan 2 in red, Clan 3 in black, and Clan 4 in green. Credit: M. Shelomi.

Modifications of motifs clustered by CYP subfamily, not by species. CYP307 is missing helix C completely, and one monophyletic clade of Phasmatodea CYP13/14 has the motif as WxxxxK instead of WxxxxR. Helix I (GxE/DTT/S) is completely absent in CYP315 and CYP304, including the non-Phasmatodea sequence. It is mutated as GxDTX in CYP49, GxEST or GxE/DTx in the Phasmatodea CYP15A’s, GxxTT in CYP44, GxxxxT in CYP314, GxE/Dxx in CYP4AA and CYP392, and Gxxxx in CYP307. Helix K mutations were widespread, especially in the CYP15A and CYP6K subfamilies. The PERF motif was mostly conserved with scattered exceptions.

4. Discussion

It is not always possible to tell whether individual transcripts/isotigs represent truly different genes, different alleles from different individuals, splice variations, or errors in assembling the original transcriptomes. However, the original study compared the Peruphasma schultei transcriptome data to genome data from the same individuals and concluded that the isotigs represent true isoforms within each isogroup, and thus are closer to the actual gene number [25]. The numbers identified from this analysis are still likely an underestimate of the total number of CYP genes for each species, as only midguts were sampled, and all from insects that had not been given a toxin, pathogen, or stress challenge that could induce expression of different CYPs. As the original study was unable to produce a transcriptome from the posterior midgut of Sipyloidea sipylus, it is likely that some CYPs differentially expressed in that subregion of the midgut are missing from that species’ dataset: probable examples based on Table 1 are the 44, 314A1, and some 4C1 and 15A1 CYPs. Moreover, as 5’ and 3’ truncated transcripts were deleted, some misassembled genes would not be in this dataset, so the isotig numbers at least are likely underestimates. These caveats do not affect the general conclusions of this study or the ability of the data to test the qualitative hypotheses of this study, and only mean the quantitative numbers in Table 1 are meant to be preliminary rather than definitive.

Peruphasma and Sipyloidea each had 19 different CYP isogroups, while the others had 27–33. Some isogroups had multiple paralogous isotigs, and such isogroups in one species would also have paralogues in another, suggesting the gene duplications occurred in a common ancestor of these six species. The low isogroup number in Sipyloidea likely reflects the lack of posterior midgut tissue that includes the Phasmatodea-specific excretory organs, the “appendices of the midgut,” which are known to have differentially expressed CYPs and play a role in xenobiotic metabolism [43]. Why the mitochondrial clan CYPs are expressed in the posterior midgut but not the anterior midgut is worth examining further. If one assumes that a Sipyloidea transcriptome that included the posterior midgut would have higher transcript numbers, then the relatively low isogroup number in Peruphasma is likely due to it being in a separate lineage from the others: a recent Phasmatodea phylogenomic work separated the New World Occidophasmata, which includes the Peruvian Peruphasma, and the Old World Oriophasmata that includes the other five species [44]. The expansions within the different CYP families do not all match this phylogeny. Some of the 6A13/14 and 15A1 CYP branches show higher diversity among the Oriophasmata, or have the P. schultei gene as a sister to the others. The 6J1s show many duplications that occurred after the individual species evolved, not before. As most recent phylogenomic analyses of the Phasmatodea disagree on the relative placements of the non-Phasmatidae families [44–46], one cannot currently be certain of where the expansions of these CYP families occurred. For example, recent studies placed P. schultei as more closely related to Medauroidea, Ramulus, and Extatosoma, with Aretaon and Sipyloidea as more basal groups [45,47]. From Figure 1,
this type of branching is seen in the CYP group 304A1, but not in the clan 4 CYPs, for example, though the low levels of Sipyloidea CYP transcripts affect the data. Ultimately one cannot currently approach the different Phasmatodea phylogenetic hypotheses using CYP expansion data.

As in the majority of insects [3,12,15,48], the Phasmatodea CYPs are mostly in the CYP6 family. This clan 3 family unique to insects is accepted as their primary xenobiotic metabolism CYP. CYP6J and CYP6A seem to be the conserved, dominant CYP subfamilies in the Phasmatodea, inherited from a common ancestor but with frequent gene duplication events that could reflect neofunctionalizations. Differentiating between CYP6J1, 6A13, and 6A14 is difficult, and these three genes and other CYP6’s are closely related as evidenced in Figure 1 and the literature [49,50]. All Phasmatodea also expressed a single CYP6K inherited from a common ancestor. CYP6J1 and CYP6K1 were originally described as microsomal CYPs in the cockroach Blatella germanica (L.), where they are expressed in all life stages but at higher levels in the adult abdomen, but their function was not speculated [49]. CYP6D1 affects pyrethroid resistance in the housefly Musca domestica L. 1758, while CYP6B’s in Papilio L. 1758 butterflies detoxify plant furanocoumarins [4]. CYP6G1 expression in the Drosophila Malpighian tubules directly correlates with DDT resistance [17]. Knockdown of four different CYP6 genes in Locusta migratoria L. 1758 increased nymph mortality following insecticide exposure [51]. A CRISPR-Cas9 knockout of CYP6AE in the extremely polyphagous Helicoverpa armigera (Hübner 1808) reduced its resistance to both host plant chemicals and insecticides [52], with similar results obtained from RNAi knockdown of CYP6AB60 in the cutworm Spodoptera litura (Fabricius 1775) [53].

CYP4’s are in CYP clan 4, and produce isozymes linked to xenobiotic metabolism as well as odorant metabolism [3]. CYP4Cl is one of the earliest known CYPs from insects [12], and is induced by endogenous hypetrehalosemic hormone [4]. Most research on this gene focuses on its relationship to hormones, not xenobiotics [54]. In addition to CYP4C, some but not all Phasmatodea also expressed CYP4AA and CYP4G genes. CYP15A is in CYP clan 2, which is primarily involved in basic physiological functions. CYP15A1 in particular is involved in juvenile hormone metabolism, specifically the epoxidation of methyl farnesoate to juvenile hormone III [55,56].

The other families, with the exception of 4AA1 in Medauroidae, had no more than one representative isogroup per species and most only had one isotig each. Even assuming that these numbers are underestimates, it is still unlikely that less conservative criteria for including and excluding sequences in the final analysis would have produced more hits for these families: these Phasmatodea likely inherited one copy of each gene from their common ancestor. These families include several from the mitochondrial clan of CYPs, including 44, 49A, 302A, 314A, and 315A, which are highly conserved and essential for physiological functions such as molting and development [3,11,57]. CYP304 and CYP307 are in clan 2 with CYP15, and also hold non-defensive functions, such as ecdysteroid synthesis [3]. CYP9 is in clan 3 along with CYP6 and is also involved in xenobiotic metabolism [3].

The Phasmatodea CYP complement can be compared to those of other insects. A study of the Plutella xylostella (L.) genome similarly found many copies of CYP6, albeit from different subfamilies, but also found many genes of CYP340, which is a clan 4 CYP associated with xenobiotic detoxification, multiple CYP9 genes, and only one CYP15 gene [3]. The Drosophila genome contains 86 CYP genes in 25 families, more than half of which were CYP4 and CYP6 [58]. An RNA-Seq study of the polyphagous stink bug Halyomorpha halys Stål 1855 found a minimum 163 CYP genes, of which 105 were in clan 3, 46 in clan 4, and six each in clan 2 and the mitochondrial clan. The H. halys CYP6s were primarily in the B subfamily, and only one CYP15 gene was found [13]. Genome analysis of the mosquito Aedes aegypti found 44 CYP6 and 37 CYP9 genes, and only one CYP15 [15]. That only one CYP9 gene was ever found in the Phasmatodea midgut transcriptomes despite this family being expanded in other insects could suggest the gene family is more highly expressed in other tissues not sampled in the original transcriptome study, such as the ‘fat body’. It is not clear why Phasmatodea have so many more CYP15 genes than other
insects: there seems to be no precedent for an insect with multiple CYP15A1 genes, nor is this juvenile hormone epoxidase gene commonly associated with the midgut [56]. Recall that CYP pseudogenes are relatively common, so the possibility exists that some of these transcripts are for nonfunctional proteins [9]. The functions of the Phasmatodea CYP15A1s are for now an open mystery.

*Extatosoma tiaratum* was particularly interesting for this study due to its ability to eat *Eucalyptus*. CYPs play roles in detoxifying *Eucalyptus* in marsupial mammals, such as CYP2C and CYP4A in the koala, *Phascolarctos cinereus* (Goldfuss 1817) [59,60]. No CYP clade was particularly expanded in *Extatosoma* but not in other phasmds, so one cannot presently speculate which of the *Extatosoma* CYPs is most important for *Eucalyptus* detoxification. That question could be answered with an RNA-Seq experiment comparing transcriptomes of *Extatosoma* reared on *Eucalyptus* and those reared on another suitable diet, such as *Rosa*. One should also note that *E. tiaratum* is not a *Eucalyptus* specialist, but rather a polyphagous species that happens to also eat *Eucalyptus* in the wild. Comparing *Extatosoma* with other *Eucalyptus*-feeding insects, particularly specialists and including but not limited to other Phasmatodea, would be informative. Presently we see no evidence from the Phasmatodea data that either toxic diets or polyphagy correlate with CYP diversity.

The other Phasmatodea with a highly chemically defended diet, *Peruphasma*, similarly did not show any expansion in any CYP families relative to other organisms. Their natural food plant is *Schinus moll*, the Peruvian pepper or *pirul*, whose leaf essential oils are known insecticides [61], but it remains unknown how exactly *P. schultei* survives such a toxic diet. Their food plant in captivity, *Ligustrum*, produces an iridoid glycoside, oleuropein, which decreases lysine levels and makes the plant proteins non-nutritious. Metabolizing oleuropein with a CYP is not the only means of nullifying it, nor is it a common strategy. Lepidoptera and Hymenoptera that specialize in *Ligustrum* secrete large amounts of glycine or, less often, GABA or β-alanine, into their gut that counteract the lysine-eliminating effect [62]. *Peruphasma* may use a similar strategy, though this hypothesis has yet to be tested.

5. Conclusions

In conclusion, the Phasmatodea have conserved, orthologous CYPs likely inherited from a common ancestor, with the likely primary xenobiotic-detoxification proteins in the CYP6f and CYP6A families. Within those families, typically one or two isogroups had an unusually large number of isotigs, suggesting gene duplication events, splice variants, or other modifications that in turn suggest possible neofunctionalization as observed with Phasmatodea digestive enzymes [63]. These would be the primary genes to study in any future knockdown experiments. The curious expansion in CYP15A1 needs to be investigated further, with hypothesized explanations including neofunctionalization of xenobiotic metabolism ability or pseudogenes: the sequences with missing or mutated CYP conserved motifs may or may not be examples thereof. The Phasmatodea gut CYP arsenal seems relatively limited. While many species are oligo- or polyphagous, individual insects often prefer to feed on the same species they first fed on [64], so it is possible that the Phasmatodea genomes may contain more CYP genes than need to be expressed by any individual over its lifetime. However, as other authors have noted, it is difficult to draw conclusions about diets from comparative CYP expressions in different species, even within the same genus or insect family [15]. CYPs in an organism likely reflect its evolutionary origin more than its diet, although broader studies are needed to confirm this.

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