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RESEARCH ARTICLE

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The mitochondrial genome of the ascalaphid owlfly *Libelloides macaronius* and comparative evolutionary mitochondriomics of neuropterid insects

Enrico Negrisolò*, Massimiliano Babbucci and Tomaso Patarnello

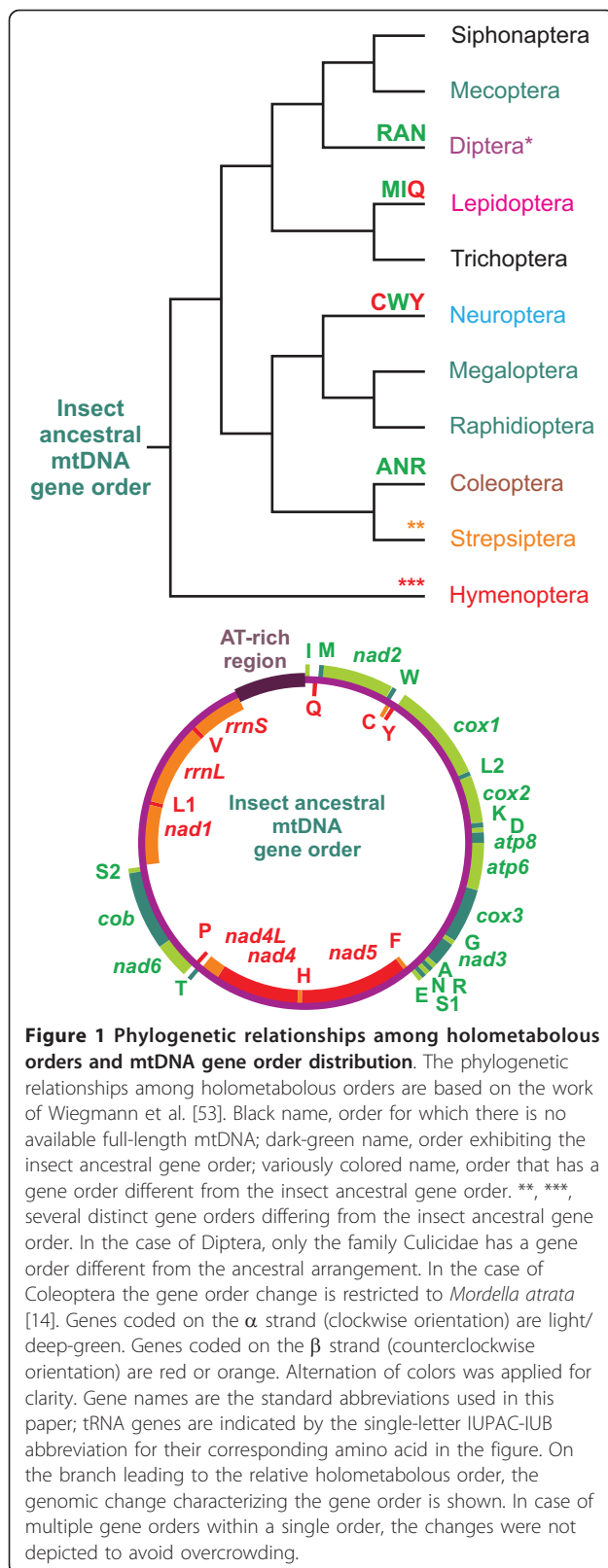
Abstract

Background: The insect order Neuroptera encompasses more than 5,700 described species. To date, only three neuropteran mitochondrial genomes have been fully and one partly sequenced. Current knowledge on neuropteran mitochondrial genomes is limited, and new data are strongly required. In the present work, the mitochondrial genome of the ascalaphid owlfly *Libelloides macaronius* is described and compared with the known neuropterid mitochondrial genomes: Megaloptera, Neuroptera and Raphidioptera. These analyses are further extended to other endopterygotan orders.

Results: The mitochondrial genome of *L. macaronius* is a circular molecule 15,890 bp long. It includes the entire set of 37 genes usually present in animal mitochondrial genomes. The gene order of this newly sequenced genome is unique among Neuroptera and differs from the ancestral type of insects in the translocation of *trnC*. The *L. macaronius* genome shows the lowest A+T content (74.50%) among known neuropterid genomes. Protein-coding genes possess the typical mitochondrial start codons, except for *cox1*, which has an unusual ACG. Comparisons among endopterygotan mitochondrial genomes showed that A+T content and AT/GC-skews exhibit a broad range of variation among 84 analyzed taxa. Comparative analyses showed that neuropterid mitochondrial protein-coding genes experienced complex evolutionary histories, involving features ranging from codon usage to rate of substitution, that make them potential markers for population genetics/phylogenetics studies at different taxonomic ranks. The 22 tRNAs show variable substitution patterns in Neuropterida, with higher sequence conservation in genes located on the α strand. Inferred secondary structures for neuropterid *rns* and *rnl* genes largely agree with those known for other insects. For the first time, a model is provided for domain I of an insect *rnl*. The control region in Neuropterida, as in other insects, is fast-evolving genomic region, characterized by AT-rich motifs.

Conclusions: The new genome shares many features with known neuropteran genomes but differs in its low A+T content. Comparative analysis of neuropterid mitochondrial genes showed that they experienced distinct evolutionary patterns. Both tRNA families and ribosomal RNAs show composite substitution pathways. The neuropterid mitochondrial genome is characterized by a complex evolutionary history.

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Background

Insect mitochondrial genomes (mtDNAs) are usually a double-strand circular molecule containing 13 protein-coding genes (PCGs), 22 transfer RNAs (tRNAs) and 2 ribosomal RNAs, i.e., the small and large subunits [1]. A notable exception is represented by the mtDNA of the human body louse *Pediculus humanus*, which consists of 18 miniature circular chromosomes containing one to three genes [2]. Insect mtDNA size is typically in the range of 14 to 20 kbp, but genomes exceeding these values are known [3]. In these latter cases, the size increase is connected to the expansion of the main non-coding region, named the AT-rich or control region [3]. The gene order is a feature of mtDNA that can provide important evidence to establish evolutionary relationships among taxa at high and/or low taxonomic levels [4,5]. The most widespread gene order in insect mtDNAs is shown in Figure 1. This gene order, initially determined for *Drosophila yakuba* mtDNA, is considered ancestral for the entire class Insecta [5-7]. Several gene orders departing from the ancestral arrangement exist and can be restricted to single species as in the strepsipteran *Mengenilla australiensis* or common to whole groups of higher taxonomic rank, ranging from family (e.g., Culicidae) to order (e.g., Lepidoptera) (Figure 1) [8-12]. In the latter cases, the peculiar gene order becomes an important marker to delimit taxonomic boundaries and constitutes a major signature of mtDNA. To date, full-length mtDNAs in insects have been sequenced in an imbalanced manner, with whole orders still lacking any published information or being poorly represented in data banks. In this respect, the neuropterid orders Megaloptera, Neuroptera (also named Planipennia), and Raphidioptera are underrepresented taxa. Only recently have sequences become available for these taxa [13-15]. Currently, full-length mtDNAs are known for the megalopterans *Corydalus cornutus* and *Protohermes concolorus*, both members of the Corydalidae family, and *Sialis hamata*, of the Sialidae [13-15]. Three mtDNAs are available for the neuropterans *Ascaloptynx appendiculatus* (Ascalaphidae), *Ditaxis latistyla* (Mantispidae), and *Polystoechotes punctatus* (Polystoechotidae) [13,14]. A complete mtDNA sequence exists for the raphidiopteran *Mongoloraphidia harmandi* (Raphidiidae) [14]. Finally, partial mtDNA sequences are available for the neuropteran *Myrmelon immaculatus* (Myrmelontidae) and the raphidiopteran *Agulla* sp. (Raphidiidae) [13].

Here we determined the complete mtDNA sequence of the ascalaphid owlfly *Libelloides macaronius* (Scopoli, 1763) (Neuroptera, Ascalaphidae). The newly determined

genome was compared with available neuropterid mtDNAs (complete and partial) as well as with genomes obtained from other holometabolous insects (Table 1) [6,9-52]. The analyses were performed following a comparative and evolutionary perspective.

Results and discussion

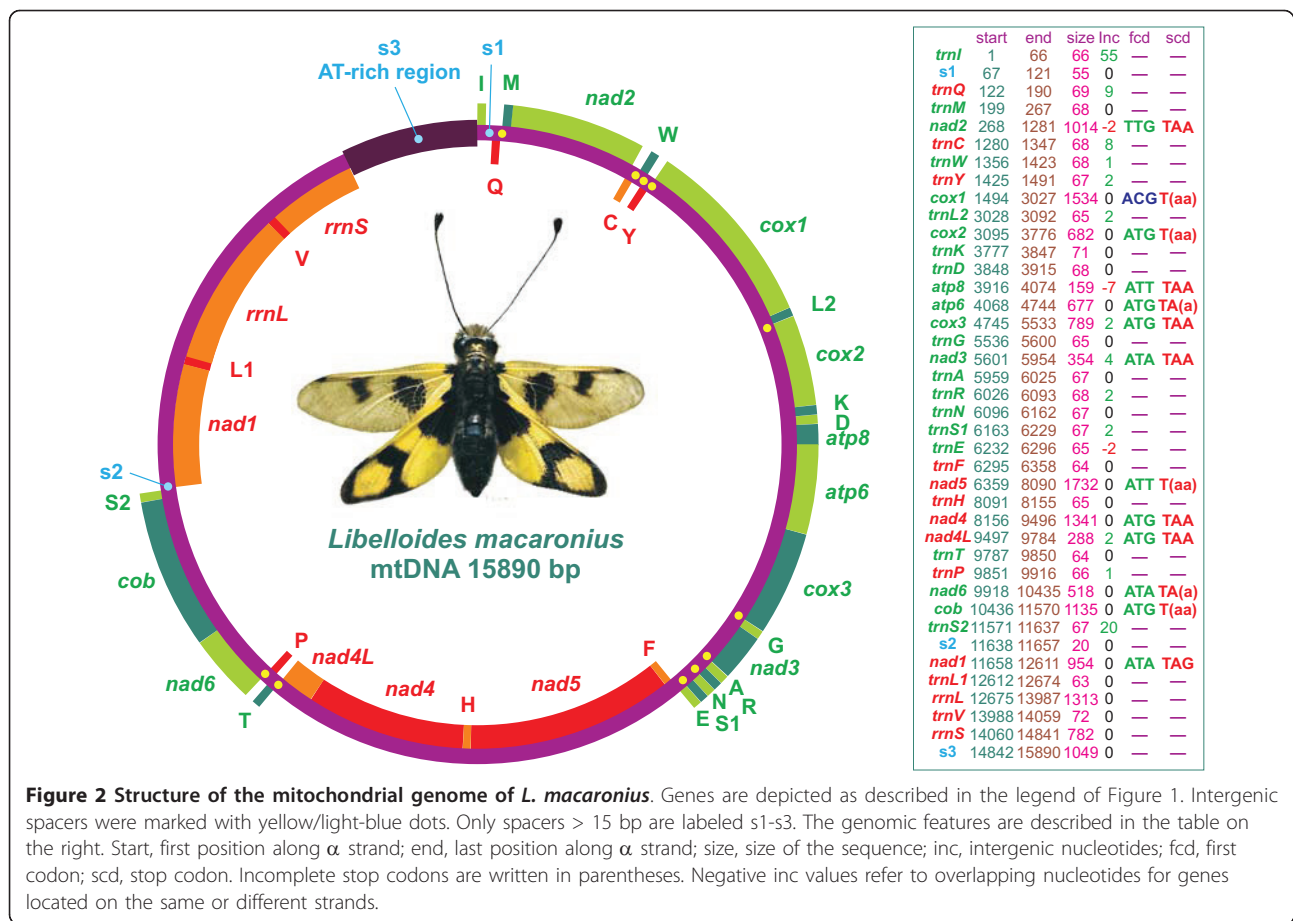
Genome structure

The mtDNA genome of *L. macaronius* is a circular molecule 15,890 bp long (Figure 2). It contains the entire set of 37 genes usually present in animal mtDNAs

Table 1 List of endopterygotan taxa analyzed in present paper

ORD	TAXON	ACN	REF	ORD	TAXON	ACN	REF
COL	<i>Acmaeodera</i> sp.	FJ613420	[16]	DIP	<i>Drosophila simulans</i>	AF200833	[31]
COL	<i>Adelium</i> sp	FJ613422	[16]	DIP	<i>Drosophila yakuba</i>	X03240	[6]
COL	<i>Anoplophora glabripennis</i>	DQ768215	UNP	DIP	<i>Haematobia irritans</i>	DQ029097	[33]
COL	<i>Apatides fortis</i>	FJ613421	[16]	DIP	<i>Lucilia sericata</i>	AJ422212	[34]
COL	<i>Chaetosoma scaritides</i>	EU877951	[17]	DIP	<i>Mayetiola destructor</i>	GQ387648	[35]
COL	<i>Chauliognathus opacus</i>	FJ613418	[16]	DIP	<i>Rhopalomyia pomum</i>	GQ387649	[35]
COL	<i>Chrysochroa fulgidissima</i>	EU826485	[18]	DIP	<i>Simosyrphus grandicornis</i>	DQ866050	[30]
COL	<i>Crioceris duodecimpunctata</i>	AF467886	[19]	DIP	<i>Trichophthalma punctata</i>	DQ866051	[30]
COL	<i>Cyphon</i> sp.	EU877949	[17]	HYM	<i>Abispa ephippium</i>	EU302588	[36]
COL	<i>Hydroscapha granulum</i>	AM493667	UNP	HYM	<i>Apis mellifera ligustica</i>	L06178	[37]
COL	<i>Lucanus mazama</i>	FJ613419	[16]	HYM	<i>Bombus ignitus</i>	DQ870926	[38]
COL	<i>Macrogyrus oblongus</i>	FJ859901	[14]	HYM	<i>Cephus cinctus</i>	FJ478173	[39]
COL	<i>Mordella atrata</i>	FJ859904	[14]	HYM	<i>Diadegma semiclausum</i>	EU871947	[40]
COL	<i>Priasilpha obscura</i>	EU877952	[17]	HYM	<i>Evania appendigaster</i>	FJ593187	[41]
COL	<i>Psacotheta hilaris</i>	FJ424074	[20]	HYM	<i>Melipona bicolor</i>	AF466146	[42]
COL	<i>Pyrocoelia rufa</i>	AF452048	[21]	HYM	<i>Orussus occidentalis</i>	FJ478174	[39]
COL	<i>Pyrophorus divergens</i>	EF398270	[22]	HYM	<i>Vanhornia eucnemidarum</i>	DQ302100	[39]
COL	<i>Rhagophthalmus lufengensis</i>	DQ888607	[23]	LEP	<i>Acraea issoria</i>	GQ376195	[43]
COL	<i>Rhagophthalmus ohbai</i>	AB267275	[23]	LEP	<i>Adoxophyes honmai</i>	DQ073916	[44]
COL	<i>Rhopaea magnicornis</i>	FJ859903	[14]	LEP	<i>Antheraea pernyi</i>	AY242996	[45]
COL	<i>Sphaerius</i> sp.	EU877950	[17]	LEP	<i>Antheraea yamamai</i>	EU726630	[46]
COL	<i>Tetraphalerus bruchi</i>	EU877953	[17]	LEP	<i>Artogeia melete</i>	EU597124	[47]
COL	<i>Trachypachus holmbergi</i>	EU877954	[17]	LEP	<i>Bombyx mandarina</i>	AB070263	[48]
COL	<i>Tribolium castaneum</i>	AJ312413	[24]	LEP	<i>Bombyx mori</i>	AF149768	[48]
DIP	<i>Aedes aegypti</i>	EU352212	UNP	LEP	<i>Coreana raphaelis</i>	DQ102703	[49]
DIP	<i>Aedes albopictus</i>	AY072044	UNP	LEP	<i>Diatraea saccharalis</i>	FJ240227	UNP
DIP	<i>Anopheles gambiae</i>	L20934	[9]	LEP	<i>Eriogyna pyretorum</i>	FJ685653	[50]
DIP	<i>Anopheles quadrimaculatus A</i>	L04272	[10]	LEP	<i>Lymantria dispar</i>	FJ617240	UNP
DIP	<i>Bactrocera carambolae</i>	EF014414	UNP	LEP	<i>Manduca sexta</i>	EU286785	[11]
DIP	<i>Bactrocera dorsalis</i>	DQ845759	[25]	LEP	<i>Ochrogaster lunifer</i>	AM946601	[12]
DIP	<i>Bactrocera oleae</i>	AY210702	[26]	LEP	<i>Phthonandria atrilineata</i>	EU569764	[51]
DIP	<i>Bactrocera papayae</i>	DQ917578	UNP	LEP	<i>Saturnia boisduvalii</i>	EF622227	[52]
DIP	<i>Bactrocera philippinensis</i>	DQ995281	UNP	NEU	<i>Ascaloptynx appendiculatus</i>	FJ171324	[13]
DIP	<i>Ceratitis capitata</i>	AJ242872	[27]	NEU	<i>Ditaxis biseriata</i>	FJ859906	[14]
DIP	<i>Chrysomya putoria</i>	AF352790	[28]	NEU	<i>Libelloides macaronius</i>	FR669150	[**]
DIP	<i>Cochliomyia hominivorax</i>	AF260826	[29]	NEU	<i>Myrmelon immaculatus</i>	FJ207458-9	[13]
DIP	<i>Culicoides arakawae</i>	AB361004	UNP	NEU	<i>Polystoechotes punctatus</i>	FJ171325	[13]
DIP	<i>Cydistomyia duplonotata</i>	DQ866052	[30]	MEG	<i>Corydalus cornutus</i>	FJ171323	[13]
DIP	<i>Dermatobia hominis</i>	AY463155	UNP	MEG	<i>Protohermes concolorus</i>	EU526394	[15]
DIP	<i>Drosophila littoralis</i>	FJ447340	UNP	MEG	<i>Sialis hamata</i>	FJ859905	[14]
DIP	<i>Drosophila mauritiana</i>	AF200830	[31]	RPH	<i>Mongoloraphidia harmandi</i>	FJ859902	[14]
DIP	<i>Drosophila melanogaster</i>	U37541	[32]	RPH	<i>Agulla</i> sp.	FJ207460-1	[13]
DIP	<i>Drosophila sechellia</i>	AF200832	[31]	MCP	<i>Neopanorpa pulchra</i>	FJ169955	UNP

ORD, order; N, identification number; ACN, accession number; REF, reference. COL, Coleoptera; DIP, Diptera; HYM, Hymenoptera; LEP, Lepidoptera, NEU, Neuroptera; MEG, Megaloptera; RPH, Raphidioptera; MCP, Mecoptera. UNP, unpublished mtDNA genome; **, present paper. Within each order taxa are listed alphabetically.



and 14 non-coding portions. Three of these spanning at least 15 bp are labeled as intergenic spacers (s1-s3) in Figure 2.

Atp8 and *atp6* are the only genes, located on the same strand, that overlap. The *atp8-atp6* partial superimposition is a common feature of neuropterid mtDNAs and, in general, of animal mtDNAs [1,13,15]. Other genes located on the same strand (both α and β) are contiguous (e.g., *nad4L* and *nad4*) or separated by a few nucleotides (e.g., *nad3* and *trnA*). Genes on opposite strands overlap (e.g., *nad2* and *trnC*), are contiguous (e.g., *trnQ* and *trnM*), or are separated by some nucleotides (e.g., *trnT* and *trnP*) or intergenic spacers (e.g., *trnS2* and *nad1*) (Figure 2). Similar patterns can be observed in other neuropterid mtDNAs. It must be noted that *nad4L* and *nad4* are contiguous in *A. appendiculatus* and *L. macaronius* (both members of the family Ascalaphidae). Conversely, in all other sequenced neuropterid mtDNAs, *nad4L* and *nad4* overlap by 7 bp [13,15].

The gene order of *L. macaronius* mtDNA differs from the insect ancestral gene order in the translocation of *trnC*, which is located upstream of *trnW* in contrast to its

traditional location downstream of *trnW* (Figures 1, 2). This translocation is shared by and exclusive to all neuropteran mtDNAs that are fully or partly sequenced to date. Conversely, Megaloptera and Raphidioptera exhibit the typical insect ancestral gene order as shown in Figure 1[13,15]. This figure presents the distribution of various gene orders in holometabolous insects, mapped on the tree obtained by Wiegmann et al. [53], but alternative phylogenetic hypotheses exist ([e.g., [54]]).

According to Aspöck et al., Neuroptera includes three major groups, Hemerobiiformia, Myrmelontiformia and Nevrothiformia [55]. The first two taxa are sister groups and encompass most neuropteran species, while Nevrothiformia includes the single small family Nevrothidae, which represents an early branching-off taxon within Neuroptera [55,56]. This view has been challenged by Winterton et al., who favor a paraphyletic Hemerobiiformia with respect to Myrmelontiformia and identifies the Coniopterygidae (Hemerobiiformia) family as the sister taxon of all other Neuroptera [57]. Irrespective to the arrangement of internal taxa, all studies strongly support a monophyletic Neuroptera order [55-57].

Available neuropteran mtDNAs, fully or partly sequenced, have been obtained from species belonging to distinct phyletic lines within Neuroptera. Indeed *D. latistyla* and *P. punctatus* belong to the divergent hemerobiiform families Mantispidae and Polystoechotidae, while *M. immaculatus* (Myrmelontidae), and *A. appendiculatus* + *L. macaronius* (Ascalaphidae) are members of two myrmelontiform families [13,14], present paper]. The known distribution of *trnC* translocations was mapped (data not shown) based on the recent phylogeny of Neuropterida produced by Winterton et al., which also contains estimations of the divergence times of major phyletic lineages [57]. It appears that such genomic re-arrangement occurred at minimum 238 million years ago. The estimation could be further extended back to 294 million years if this genomic change is confirmed for the whole order Neuroptera.

Further work is needed to establish whether the *trnC* translocation represents a strong molecular signature for the entire Neuroptera clade.

Within Endopterygota, the translocation of *trnM* characterizes all lepidopteran species sequenced to date and seems to be a strong molecular mitochondrial signature for the whole order (Figure 1) (e.g., [11,12]). At the family level, the mosquitoes (family Culicidae) are characterized by the *trnR-trnA* arrangement instead of the traditional *trnA-trnR* (Figure 1) [9,10]. Several other re-arrangements are known for endopterygote mtDNAs but are restricted to single species (e.g., *Xenos vesparum*) or not consistent within broad taxonomic groups (e.g., Hymenoptera) [37,39,58-63]. The insect ancestral gene order remains the most widespread arrangement among holometabolous (Figure 1) and heterometabolous insects, suggesting that the gene order cannot play a pivotal role in resolving the relationships at the interordinal level among pterygote insects [7]. Conversely, mtDNA re-arrangements represent strong potential signatures to define taxa at the order and lower taxonomic levels (i.e., family, genus and lower) [9-14,39], present paper].

Base composition and AT- and GC-skews in the mitochondrial genomes of endopterygote insects

The nucleotide-compositional behavior of mitochondrial genomes is usually investigated through the calculation of the three parameters: AT-skew, GC-skew, and A+T content (AT%), expressed in percentages (e.g., [8,12]). Here these parameters were studied for 84 complete endopterygotan mtDNAs (Table 2; Additional file 1, Table S1) and combined, for the first time, in a single three-dimensional scatter-plot analysis (Figure 3). Arbitrary partitioning schemes, further detailed below, were applied to the estimated values to facilitate the presentation/discussion of the results. Each empirical

clustering scheme was defined in order to maximize the appreciation of the uneven distribution of various mtDNAs within the continuum of variation of the considered parameter.

A+T content

The range of variation of AT% spanned from 66.99 to 87.41, with the average value equal to 77.77. Different mtDNAs exhibited an uneven distribution within this range. Five principal clusters (A1-A5) were obtained that divided the whole range of AT% into intervals encompassing 5% of the variation.

The majority (61/84) of endopterygote mtDNAs exhibited an AT% in the range of 74-82 (A2 partially; A3; A4 partially), with a strong concentration in the interval 75-80 (43/84) (A3). Most of neuropterids were included in cluster A3, while *L. macaronius* and *C. cornutus* belonged to cluster A2. The mtDNA of *M. harmandi* was comprised in cluster A4. Very high AT% (>84) contents (A4 partially; A5) were characteristic of and limited to a few hymenopterans and dipterans. Conversely, low AT% (< 70) were restricted to coleopterans (A1).

AT-skews

The average AT-skew was 0.039, while the whole range of variation extended from -0.047 to 0.248. The endopterygote mtDNAs were grouped into six clusters (B1-B6) through the partitioning of the whole range of AT-skews in intervals spanning 0.050.

The majority of mtDNAs exhibited a positive AT-skew (71/84) with two peaks, in the intervals 0.000-0.050 (44/84; B2) and 0.050-0.100 (16/84; B3). In cluster B2 were placed the mtDNAs of *C. cornutus*, *S. hamata*, *D. biseriata* and *M. harmandi*. The cluster B3 contained also the mtDNAs of *A. appendiculatus* and *L. macaronius*. Only some coleopteran mtDNAs exceed values of 0.100 (B4-B6), and the skews were higher than 0.200 solely in *C. fulgidissima* and *T. bruchi*; (B6). Negative AT-skews (13/84) were modest (>-0.005; B1), and lepidopteran taxa predominated in this group. The mtDNAs of *P. punctatus* and *P. concolorus* exhibited negative AT-skews.

GC-skews

The average GC-skew value was -0.213, and all 84 mtDNAs displayed negative skews. The whole set of genomes was grouped into six clusters (C1-C6), each spanning 0.050 of the range of GC-skew variation.

The majority of the GC-skews were located within the range -0.30/-0.15 (68/84; C3-C5), with two peaks represented by clusters C4 (30 genomes) and C5 (24 mtDNA). The neuropterids *C. cornutus* and *P. concolorus* were included in C3, *M. harmandi* and *A. appendiculatus* in C4, while *D. biseriata*, *L. macaronius*, *S. hamata*, and *P. punctatus* were inserted in cluster C5.

Overall comparison

When the three variables analyzed above were considered together, some patterns emerged. Even if the

Table 2 AT%, AT-skew and GC-skew in endopterygotan complete mtDNAs

ORD	N	TAXON	AT%	AT-skew	GC-skew	ORD	N	TAXON	AT%	AT-skew	GC-skew
COL	1	<i>Acmaeodera</i> sp.	A1 68.41	B4 0.1142	C4 -0.2492	DIP	43	<i>Drosophila sechellia</i>	A3 77.57	B2 0.0092	C6 -0.1351
COL	2	<i>Adelium</i> sp	A2 72.71	B4 0.1367	C4 -0.2222	DIP	44	<i>Drosophila simulans</i>	A3 77.93	B2 0.0084	C6 -0.1352
COL	3	<i>Anoplophora glabripennis</i>	A3 78.31	B2 0.0116	C4 -0.2066	DIP	45	<i>Drosophila yakuba</i>	A3 78.59	B2 0.0050	C6 -0.1364
COL	4	<i>Apatides fortis</i>	A1 67.19	B5 0.1704	C3 -0.2946	DIP	46	<i>Haematobia irritans</i>	A3 79.07	B2 0.0044	C6 -0.1245
COL	5	<i>Chaetosoma scaritides</i>	A3 79.04	B2 0.0427	C4 -0.2008	DIP	47	<i>Lucilia sericata</i>	A3 77.61	B2 0.0159	C5 -0.1666
COL	6	<i>Chauliognathus opacus</i>	A3 76.86	B4 0.1120	C5 -0.1532	DIP	48	<i>Mayetiola destructor</i>	A4 84.12	B3 0.0614	C6 -0.1100
COL	7	<i>Chrysochroa fulgidissima</i>	A1 69.92	B6 0.2026	C4 -0.2373	DIP	49	<i>Rhopalomyia pomum</i>	A5 85.15	B2 0.0468	C6 -0.1230
COL	8	<i>Crioceris duodecimpunctata</i>	A3 76.89	B2 0.0442	C5 -0.1634	DIP	50	<i>Simosyrphus grandicornis</i>	A4 80.84	B1 -0.0038	C6 -0.1328
COL	9	<i>Cyphon</i> sp.	A3 75.17	B3 0.0721	C4 -0.2282	DIP	51	<i>Trichophthalma punctata</i>	A2 73.96	B3 0.0912	C4 -0.2447
COL	10	<i>Hydroscapha granulum</i>	A3 77.29	B2 0.0297	C5 -0.1519	HYM	52	<i>Abispa ephippium</i>	A4 80.61	B1 -0.0186	C1 -0.3795
COL	11	<i>Lucanus mazama</i>	A1 67.11	B3 0.0743	C3 -0.2723	HYM	53	<i>Apis mellifera ligustica</i>	A4 84.86	B2 0.0183	C3 -0.2686
COL	12	<i>Macrogyrus oblongus</i>	A3 78.00	B2 0.0421	C4 -0.2080	HYM	54	<i>Bombus ignitus</i>	A5 86.78	B2 0.0028	C3 -0.2719
COL	13	<i>Mordella atrata</i>	A2 71.94	B3 0.0691	C3 -0.2556	HYM	55	<i>Cephus cinctus</i>	A4 81.95	B2 0.0343	C3 -0.2855
COL	14	<i>Priasilpha obscura</i>	A3 76.5	B3 0.0523	C3 -0.2646	HYM	56	<i>Diadegma semiclausum</i>	A5 87.41	B2 0.0086	C5 -0.1984
COL	15	<i>Psacothaea hilaris</i>	A3 76.63	B2 0.0114	C4 -0.2129	HYM	57	<i>Evania appendigaster</i>	A3 77.77	B2 0.0267	C2 -0.3491
COL	16	<i>Pyrocoelia rufa</i>	A3 77.41	B4 0.1058	C5 -0.1584	HYM	58	<i>Melipona bicolor</i>	A5 86.72	B2 0.0155	C3 -0.2511
COL	17	<i>Pyrophorus divergens</i>	A1 69.44	B5 0.1650	C3 -0.2915	HYM	59	<i>Orussus occidentalis</i>	A3 76.21	B2 0.0169	C2 -0.3076
COL	18	<i>Rhagophthalmus lufengensis</i>	A3 79.63	B4 0.1038	C5 -0.1959	HYM	60	<i>Vanhornia eucnemidarum</i>	A4 80.11	B3 0.0857	C2 -0.3282
COL	19	<i>Rhagophthalmus ohbai</i>	A3 79.15	B4 0.1185	C4 -0.2256	LEP	61	<i>Acraea issoria</i>	A3 79.76	B1 -0.0234	C4 -0.2352
COL	20	<i>Rhopaea magnicornis</i>	A3 75.59	B2 0.0050	C4 -0.2380	LEP	62	<i>Adoxophyes honmai</i>	A4 80.39	B1 -0.0010	C5 -0.1964
COL	21	<i>Sphaerius</i> sp.	A4 80.68	B2 0.0104	C6 -0.1482	LEP	63	<i>Antheraea pernyi</i>	A4 80.16	B1 -0.0214	C4 -0.2163
COL	22	<i>Tetraphalerus bruchi</i>	A1 66.99	B6 0.2477	C3 -0.2515	LEP	64	<i>Antheraea yamamai</i>	A4 80.29	B1 -0.0221	C4 -0.2199
COL	23	<i>Trachypachus holmbergi</i>	A3 79.45	B2 0.0205	C5 -0.1536	LEP	65	<i>Artogeia melete</i>	A3 79.78	B2 0.0121	C4 -0.2218
COL	24	<i>Tribolium castaneum</i>	A2 71.68	B4 0.1091	C2 -0.3053	LEP	66	<i>Bombyx mandarina</i>	A4 81.68	B3 0.0547	C4 -0.2131
DIP	25	<i>Aedes aegypti</i>	A3 79.00	B2 0.0169	C4 -0.2109	LEP	67	<i>Bombyx mori</i>	A4 81.32	B3 0.0587	C4 -0.2162
DIP	26	<i>Aedes albopictus</i>	A3 79.54	B2 0.0079	C5 -0.1812	LEP	68	<i>Coreana raphaelis</i>	A4 82.66	B1 -0.0474	C5 -0.1578
DIP	27	<i>Anopheles gambiae</i>	A3 77.56	B2 0.0322	C5 -0.1540	LEP	69	<i>Diatraea saccharalis</i>	A4 80.02	B2 0.0213	C3 -0.2575
DIP	28	<i>Anopheles quadrimaculatus A</i>	A3 77.36	B2 0.0406	C5 -0.1814	LEP	70	<i>Eriogyna pyretorum</i>	A4 80.82	B1 -0.0305	C4 -0.2047
DIP	29	<i>Bactrocera carambolae</i>	A2 73.55	B3 0.0656	C4 -0.2237	LEP	71	<i>Lymantria dispar</i>	A3 79.88	B2 0.0160	C4 -0.2473
DIP	30	<i>Bactrocera dorsalis</i>	A2 73.58	B3 0.0675	C4 -0.2283	LEP	72	<i>Manduca sexta</i>	A4 81.79	B1 -0.0053	C5 -0.1811
DIP	31	<i>Bactrocera oleae</i>	A2 72.63	B3 0.0884	C3 -0.2802	LEP	73	<i>Ochrogaster lunifer</i>	A3 77.84	B2 0.0301	C2 -0.3175
DIP	32	<i>Bactrocera papayae</i>	A2 73.52	B3 0.0662	C4 -0.2263	LEP	74	<i>Phthonandria atrilineata</i>	A4 81.02	B2 0.0066	C5 -0.1921
DIP	33	<i>Bactrocera philippinensis</i>	A2 73.63	B3 0.0656	C4 -0.2244	LEP	75	<i>Saturnia boisduvalii</i>	A4 80.62	B1 -0.0240	C4 -0.2170
DIP	34	<i>Ceratitis capitata</i>	A3 77.48	B2 0.0211	C5 -0.1851	NEU	76	<i>Ascaloptynx appendiculatus</i>	A3 75.57	B3 0.0676	C4 -0.2059
DIP	35	<i>Chrysomya putoria</i>	A3 76.70	B2 0.0204	C5 -0.1701	NEU	77	<i>Ditaxis biseriata</i>	A3 79.79	B2 0.0154	C5 -0.1793
DIP	36	<i>Cochliomyia hominivorax</i>	A3 76.90	B2 0.0344	C4 -0.2067	NEU	78	<i>Libelloides macaronius</i>	A2 74.5	B3 0.0719	C5 -0.1767
DIP	37	<i>Culicoides arakawae</i>	A3 77.36	B2 0.0092	C4 -0.2369	NEU	79	<i>Polystoechotes punctatus</i>	A3 78.96	B1 -0.0287	C5 -0.1612
DIP	38	<i>Cydistomyia duplonotata</i>	A3 77.93	B2 0.0031	C5 -0.1771	MEG	80	<i>Corydalus cornutus</i>	A2 74.90	B2 0.0140	C3 -0.2620
DIP	39	<i>Dermatobia hominis</i>	A3 77.82	B2 0.0429	C4 -0.2267	MEG	81	<i>Protohermes concolorus</i>	A3 75.83	B1 -0.0111	C3 -0.2539
DIP	40	<i>Drosophila littoralis</i>	A3 76.24	B2 0.0124	C5 -0.1826	MEG	82	<i>Sialis hamata</i>	A3 78.32	B2 0.0145	C5 -0.1711
DIP	41	<i>Drosophila mauritiana</i>	A3 77.71	B2 0.0094	C6 -0.1348	RPH	83	<i>Mongoloraphidia harmandi</i>	A4 80.31	B2 0.0230	C4 -0.2277
DIP	42	<i>Drosophila melanogaster</i>	A4 82.16	B2 0.0167	C6 -0.1504	MCP	84	<i>Neopanorpa pulchra</i>	A3 76.38	B1 -0.0139	C5 -0.1676

ORD, order; N, identification number in Figure 3; COL, Coleoptera; DIP, Diptera; HYM, Hymenoptera; LEP, Lepidoptera; NEU, Neuroptera; MEG, Megaloptera; RPH, Raphidioptera; MCP, Mecoptera. AT%, A+T content expressed in percent. A1 (65.00 < AT% < 70.00); A2 (70.00 < AT% < 75.00); A3 (75.00 < AT% < 80.00); A4 (80.00 < AT% < 85.00); A5 (85.00 < AT% < 90.00). B1 (-0.050 < AT-skew ≤ 0.000); B2 (0.000 < AT-skew ≤ 0.050); B3 (0.050 < AT-skew ≤ 0.100); B4 (0.100 < AT-skew ≤ 0.150); B5 (0.150 < AT-skew ≤ 0.200) B6 (0.200 < AT-skew ≤ 0.250). C1 (-0.400 < GC-skew < -0.350); C2 (-0.300 < GC-skew < -0.250); C3 (-0.300 < GC-skew < -0.250); C4 (-0.250 < GC-skew < -0.200); C5 (-0.200 < GC-skew < -0.150); C6 (-0.150 ≤ GC-skew < -0.100). Within each order taxa are listed alphabetically.

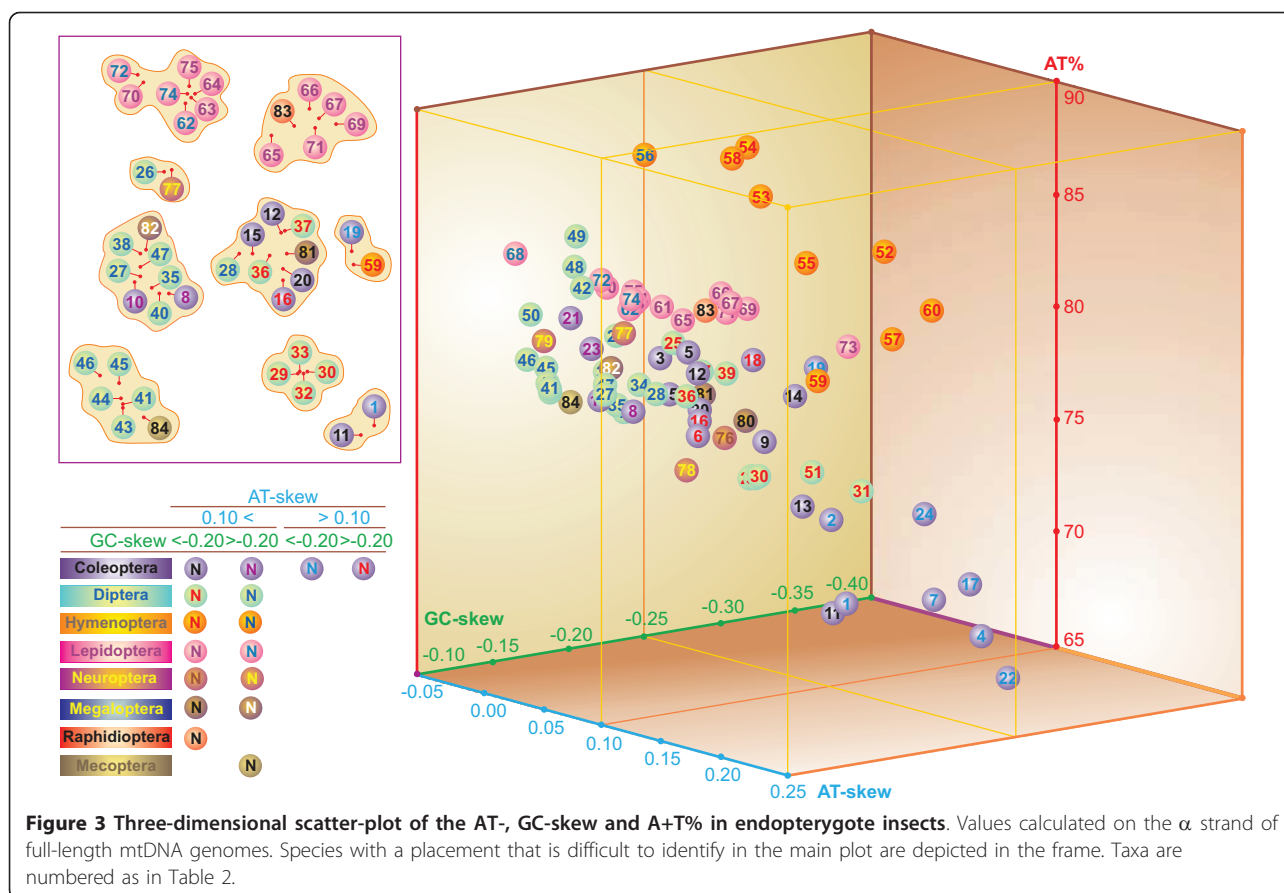


Figure 3 Three-dimensional scatter-plot of the AT-, GC-skew and A+T% in endopterygote insects. Values calculated on the α strand of full-length mtDNA genomes. Species with a placement that is difficult to identify in the main plot are depicted in the frame. Taxa are numbered as in Table 2.

parameter distribution followed a continuum range of variation, most of the sequenced endopterygote mtDNAs exhibited A+T contents in the range 70%-80%, with the highest density in the interval 75%-80%; AT-skews in the range 0.003-0.010; and GC-skews in the interval -0.300-0.150. This behavior can be represented by the cluster formula: A2/A3,B2/B3,C3-C5. Most neuropterid mtDNAs are included in this formula, with the exceptions of *P. punctatus* and *P. concolorus* (B1).

While no genome exhibited extreme values for any of the three parameters, some mtDNAs markedly departed from the most represented ranges for the behavior of one or two parameters. Very low AT content was coupled with high AT-skew in some coleopteran mtDNAs. Very high A+T contents (>84%) were restricted to a small group of hymenopterans and dipterans. Highly negative GC-skews (< -0.300) were found mostly in hymenopterans, plus the moth *O. lunifer* and the beetle *T. castaneum*.

The comparisons performed at the intra-ordinal level showed that in several cases the extremes of the ranges of variation of the analyzed parameters were occupied by the same mtDNAs. This behavior could be the result of limited taxon sampling. Thus, further data are

necessary to verify if the observed patterns are consistent over a broader taxon sampling.

Generally, the A+T content, AT-skew, and GC-skew exhibited complex behaviors that do not appear to be tightly linked, at least at the order level.

Start/stop codons in neuropterid PCGs

The inferred start/stop codons for neuropterid PCGs are listed in Table 3. ATN, GTG, TTG, and GTT are accepted canonical mitochondrial start codons for invertebrate mtDNAs and most of PCGs exhibited these start codons [64].

The *cox1* start codon for *L. macaronius* was inferred to be the triplet ACG, a non-canonical start codon for mtDNA genes. Despite the fact that *cox1* is one of more conserved mitochondrial genes, the identification of its start codon has been problematic (see [13]). An ATC codon located 27 bp upstream of ACG, within the *trnY* gene that is encoded on the β strand, could be a start codon for *L. macaronius cox1*. However, amino acid sequence comparison with a broad selection of endopterygote insects led us to the more probable conclusion that the ACG triplet was the start codon for *L. macaronius cox1*. An ACG start codon for *cox1* has also been

Table 3 Start/stop codons in neuropterid mitochondrial genes

Taxon	<i>nad2</i>		<i>cox1</i>		<i>cox2</i>		<i>atp8</i>		<i>atp6</i>		<i>cox3</i>		<i>nad3</i>	
	ST	EN	ST	EN	ST	EN	ST	EN	ST	EN	ST	EN	ST	EN
<i>Ascalopteryx appendiculatus</i>	ATT	TAA	TTA	Taa	ATG	Taa	ATT	TAA	ATG	TAa	ATG	TAA	ATA	TAA
<i>Ditaxis biseriata</i>	ATT	TAA	ATT	Taa	ATG	Taa	ATT	TAA	ATG	TAa	ATG	TAA	ATT	TAA
<i>Libelloides macaronius</i>	TTG	TAA	ACG	Taa	ATG	Taa	ATT	TAA	ATG	TAa	ATG	TAA	ATA	TAA
<i>Myrmelon immaculatus</i>	ATA	TAA	ACG	Taa	ATG	Taa	ATT	TAA	ATG	TAG	ATG	n.a.	n.a.	n.a.
<i>Polystoechotes punctatus</i>	ATT	TAA	TCG	Taa	ATG	Taa	ATT	TAA	ATG	TAA	ATG	TAa	ATT	TAA
<i>Corydalus cornutus</i>	ATT	TAA	ATT	TAa	ATG	Taa	ATT	TAA	ATG	TAa	ATG	Taa	ATT	Taa
<i>Protohermes concolorus</i>	ATT	TAa	ATT	TAA	ATG	TAA	ATT	TAA	ATG	TAa	ATG	Taa	ATT	Taa
<i>Sialis hamata</i>	ATT	TAA	ATT	Taa	ATG	Taa	ATC	TAA	ATG	TAa	ATG	TAA	ATT	TAA
<i>Agulla sp.</i>	n.a.	n.a.	CTG	Taa	ATT	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.
<i>Mongoloraphidia harmandi</i>	ATT	TAa	CTG	Taa	ATA	Taa	ATC	Taa	ATG	TAa	ATG	TAa	ATC	Taa

Taxon	<i>nad5</i>		<i>nad4</i>		<i>nad4L</i>		<i>nad6</i>		<i>cob</i>		<i>nad1</i>	
	ST	EN	ST	EN	ST	EN	ST	EN	ST	EN	ST	EN
<i>Ascalopteryx appendiculatus</i>	ATA	Taa	ATG	TAa	ATG	TAA	ATC	TAA	ATG	Taa	ATA	TAG
<i>Ditaxis biseriata</i>	ATT	Taa	ATG	Taa	ATG	TAA	ATT	TAA	ATG	Taa	TTG	TAA
<i>Myrmelon immaculatus</i>	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	Taa	n.a.	TAG
<i>Libelloides macaronius</i>	ATT	Taa	ATG	TAA	ATG	TAA	ATA	TAa	ATG	Taa	ATA	TAG
<i>Polystoechotes punctatus</i>	ATT	Taa	ATG	Taa	ATG	TAA	ATC	TAa	ATG	Taa	TTG	TAG
<i>Corydalus cornutus</i>	ATT	Taa	ATG	Taa	ATG	TAA	ATT	TAa	ATG	Taa	TTG	TAA
<i>Protohermes concolorus</i>	ATT	Taa	ATG	Taa	ATG	TAA	ATT	TAa	ATG	Taa	TTG	TAA
<i>Sialis hamata</i>	ATT	Taa	ATG	Taa	ATG	TAA	ATT	TAa	ATG	Taa	ATG	TAA
<i>Agulla sp.</i>	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	TAA	n.a.	TAA
<i>Mongoloraphidia harmandi</i>	ATA	TAA	ATG	Taa	ATT	TAA	ATA	TAa	ATG	TAA	TTG	TAA

ST, start codon; EN, end (stop) codon; TAa and Taa, stop codon incomplete; n.a., not available.

proposed for *M. immaculatus* [13]. The non-canonical start codons TTA and TCG have previously been inferred for *cox1* in *A. appendiculatus* and *P. punctatus* [13]. The remaining neuropterid *cox1* genes exhibited an ATT start codon. Non-canonical start codons in *cox1* have been inferred repeatedly for endopterygote insects (e.g., [12,65,66]).

Stop codons for the 13 neuropterid PCGs were almost invariably complete (TAA) or incomplete TAa/Taa. The only exception was observed in *nad1*, where the two ascalaphids *L. macaronius* and *A. appendiculatus* and *P. punctatus* exhibited TAG as stop codon.

Codon usage and amino acids abundance

All analyzed neuropterid mtDNAs use the standard invertebrate mitochondrial genetic code. This result was determined using the server GenDecoder [67,68]. The behavior of codon families in PCGs was investigated and the results are presented in Figures 4, 5. First codons, as well as stop codons, complete and incomplete, were excluded from the analysis to avoid biases due to unusual putative start codons and incomplete stop codons. The abundance of codon families and relative synonymous codon usage (RSCU) were calculated together with the other statistics listed below.

The A+T content, AT-skew, G+C content and GC-skew, which were calculated in pooled- $\alpha + \beta$, pooled- α and pooled- β PCGs, exhibited behaviors identical, similar to or contrasting with those observed for the whole genomes (see Additional file 2, Table S2).

The total number of codons was different in analyzed species (Figure 4; Additional file 2, Table S2). The codon families exhibiting more than 50 codons per thousand cds varied (7-9) and the majority of them were two-fold degenerate in codon usage and AT-rich. In almost all mtDNAs, the four most represented codon families were Leu2, Ile, Phe and Met with Leu2 being the most common. The only exception was *P. punctatus*, in which Ser2 was the fourth most abundant codon family, while Met family occupied only the seventh position. Leu, Ile, Ser and Phe were the most abundant amino acids, and their sum varied from 42.75% (*L. macaronius*) to 46.62% (*M. harmandi*) of the total number of amino acids.

Four-fold degenerate codon usage was A/T biased in the third position, and T was the preferred nucleotide, except in the Arg, Gly and Ser1 codon families, where A was the most common nucleotide in the third position. Two-fold degenerate codon usage showed a similarly biased pattern, with A/T favored over G/C in the third

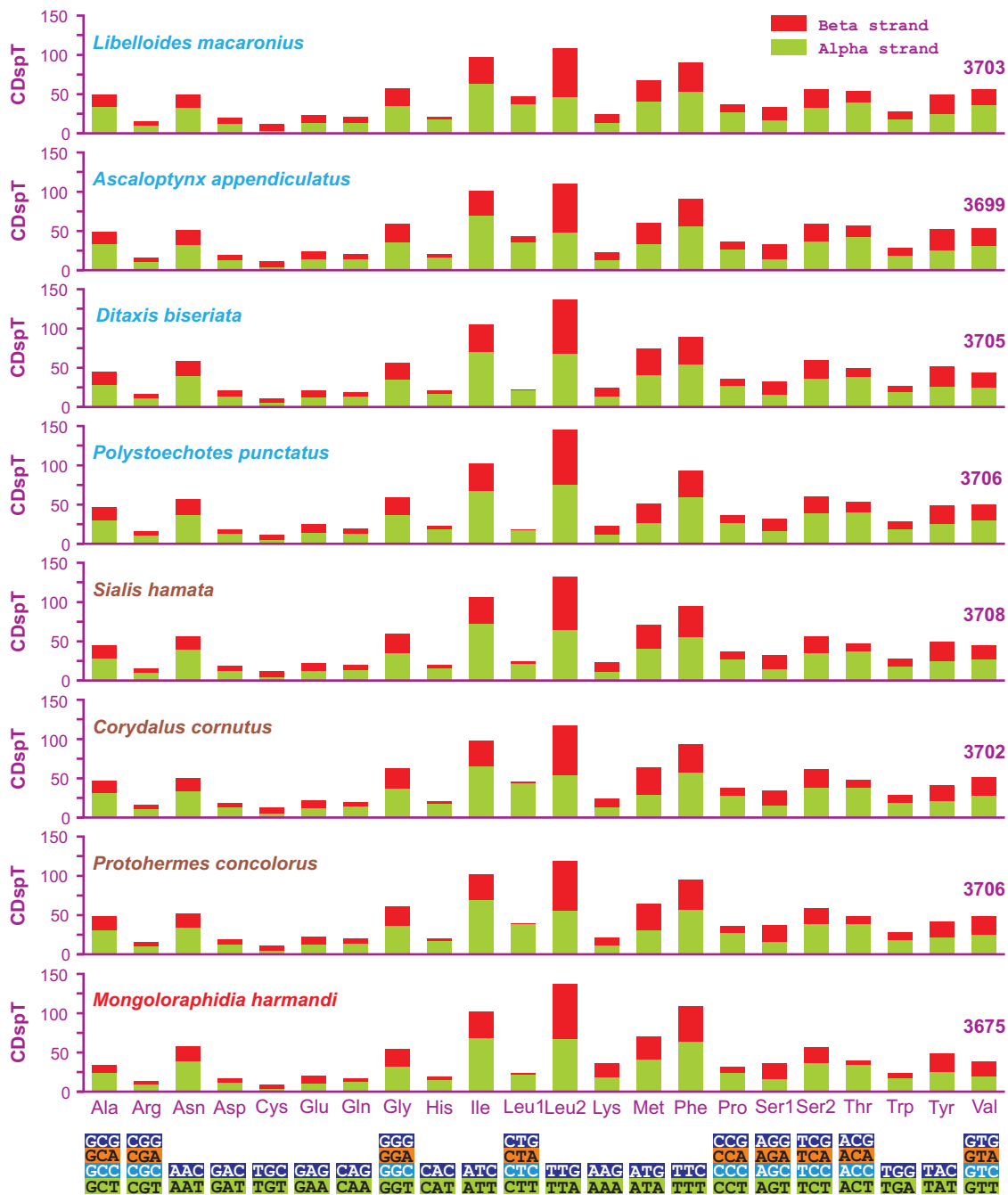


Figure 4 Codon distribution in neuropterid mtDNAs. Numbers to the left refer to the total number of codons. CDspT, codons per thousand codons. Codon families are provided on the x-axis. Within each family, the percentage of codons located, respectively, on the α strand or β strand is colored blue or red.

position (Figure 5). Both patterns were in agreement with the AT-biased content exhibited by pooled- α + β PCGs.

In pooled- α PCGs, the four-fold degenerate codon usage followed a pattern similar to that of the whole PCG set, with the exceptions of Ala, Pro and Val codon families (Figure 5 vs. Additional file 3, Figure S1).

In the pooled- β PCGs, the four-fold degenerate codon usage partly mirrored the pattern observed for pooled- α + β PCGs. Among the discrepancies, the loss of synonymous GC-rich codons was the easiest to detect (Figure 5 vs. Additional file 4, Figure S2).

The behavior of pooled- α + β PCGs in different species varied with respect to the usage of the whole set (62) of

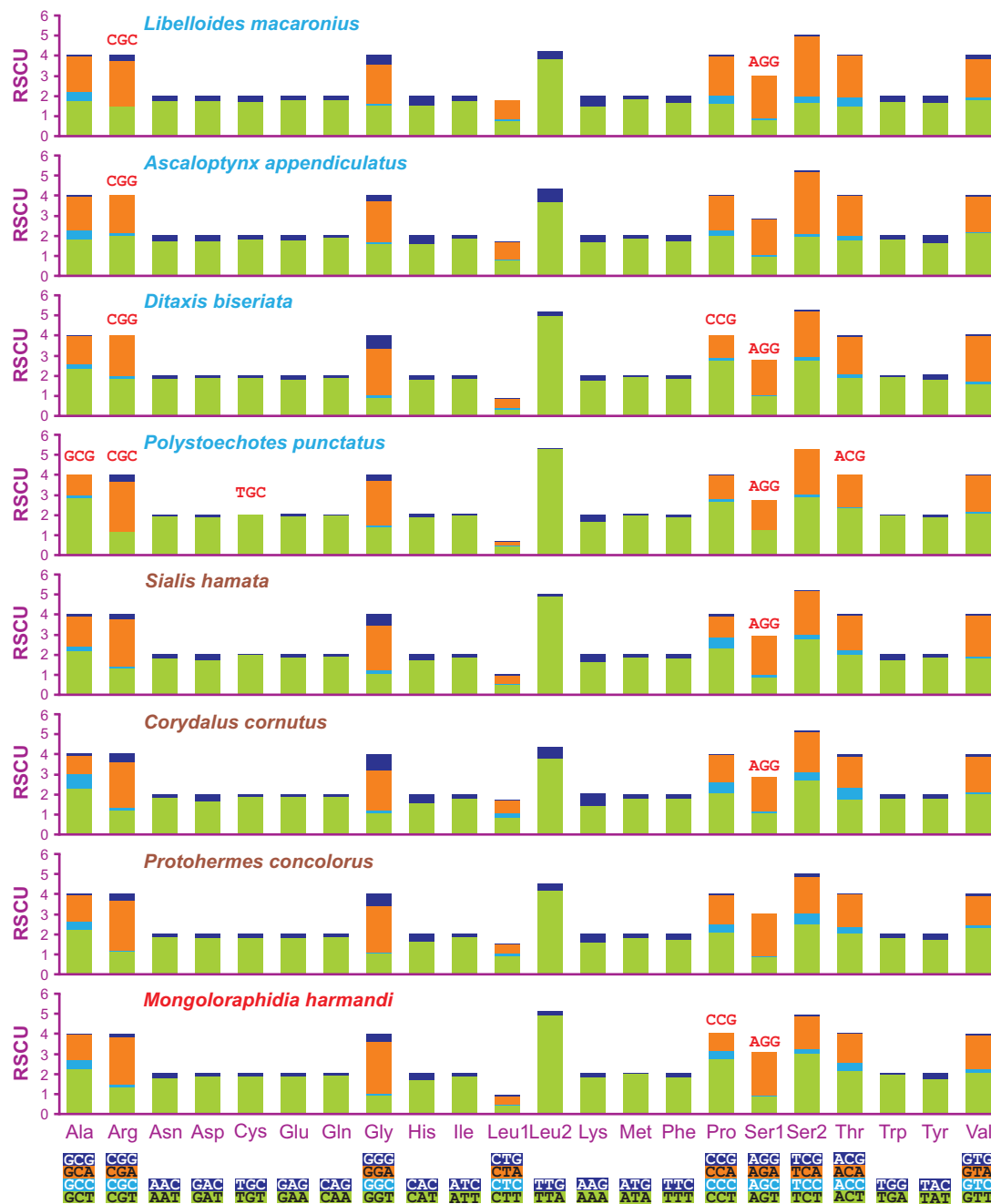


Figure 5 Relative synonymous codon usage (RSCU) in neuropterid mtDNAs. Codon families are provided on the x-axis. Red codon, codon not present in the genome.

codons encoding amino acids. All missed codons were GC-rich. Codon usage could not be directly linked to total number of codons nor to A+T content (Figure 5; Additional file 2, Table S2). The reduction in codon usage could be possibly linked to AT/GC-skews. Indeed, *P. punctatus*, which used the smallest number of codons, also exhibited the most negative AT-skew_{α+β} and the highest GC-skew_{α+β}. The number of used

codons decreased when PCGs encoded on a single strand were analyzed (Additional file 2, Table S2; Additional file 3, Figure S1; Additional file 4, Figure S2). The pattern observed for pooled-α+β PCGs was mostly mirrored by pooled-α PCGs. On the β strand, codon reductions appeared to be species specific. The lost codons belonged to GC-rich and comparatively A-rich codon-families. Finally, on the β strand, the pattern of codon

Table 4 Total number of α -codons vs. total number of β -codons ratio $_{\alpha/\beta}$

codon family	avg-ratio $_{\alpha/\beta}$	STD
oavg	1.59	0.004
Ala	1.94	0.14
Arg	1.81	0.08
Asn	2.01	0.18
Asp	2.03	0.28
Cys	0.56	0.08
Glu	1.40	0.09
Gln	2.22	0.21
Gly	1.50	0.05
His	4.41	0.25
Ile	2.06	0.11
Leu1	13.22	7.46
Leu2	0.89	0.10
Lys	1.10	0.12
Met	1.18	0.24
Phe	1.54	0.12
Pro	3.01	0.22
Ser1	0.83	0.10
Ser2	1.66	0.17
Thr	3.67	1.06
Trp	2.18	0.28
Tyr	1.02	0.03
Val	1.37	0.30

Oavg, overall (o) average(avg) ratio $_{\alpha/\beta}$; avg-ratio $_{\alpha/\beta}$, average-ratio $_{\alpha/\beta}$; STD, standard deviation

loss appeared to be linked to the single/combined effects of GC content and AT-/GC-skews.

The total number of α -codons vs. the total number of β -codons (ratio $_{\alpha/\beta}$) was calculated for every species (Table 4). The overall (o) average (avg) ratio $_{\alpha/\beta}$ ($1.59 \pm 4 \times 10^{-3}$) was used as a measure to identify possible distributional biases of different codon families. Several codon families exhibited ratio $_{\alpha/\beta}$ values close to oavg-ratio $_{\alpha/\beta}$, suggesting that their distribution is not markedly diverse in the PCGs coded on different strands. Codon families His, Leu1, Pro, and Thr exhibited ratio $_{\alpha/\beta}$ values strongly biased toward the α strand. Conversely, codon families Cys, Leu2 and Ser1 presented ratio $_{\alpha/\beta}$ values clearly β strand-oriented.

The most obvious bias was due to structural/functional requirements of PCGs. This point is well represented by the distribution of Cys codon family. Cys was the rarest amino acid in mtDNA proteins and was more abundant in polypeptides (NADH subunits) encoded on the β strand than in the proteins encoded on the α strand. The Cys residues can produce intra- and inter-chain disulfide bridges (S-S) that play a fundamental structural role. Disulfide bonds and interactions between pairs of Cys residues have been postulated to have an important structural role in NADH subunits of primate

mtDNA [69]. The abundance of Cys residues in neuropterid NADHs, and in general in endopterygotes, appears to agree with findings derived from primates. Several marked departures from oavg-ratio $_{\alpha/\beta}$ can be explained in terms of structural/functional requirements of single polypeptides and are not explored here.

A compositional bias effect was evident in the behavior of Leu1 (CTN) and Leu2 (TTR) codon families. As mentioned above, Leu was the most abundant amino acid in the PCGs. The pooled superfamily Leu = Leu1 + Leu2 exhibited an avg-ratio $_{\alpha/\beta} = 1.28 \pm 0.11$. This result supports a moderate β strand bias in the distribution of Leu. However, Leu1 and Leu2 codon families exhibited an opposite bias in term of strand distribution. These strand biases can be linked to the AT-skew and GC content shown by pooled- α and pooled- β PCGs.

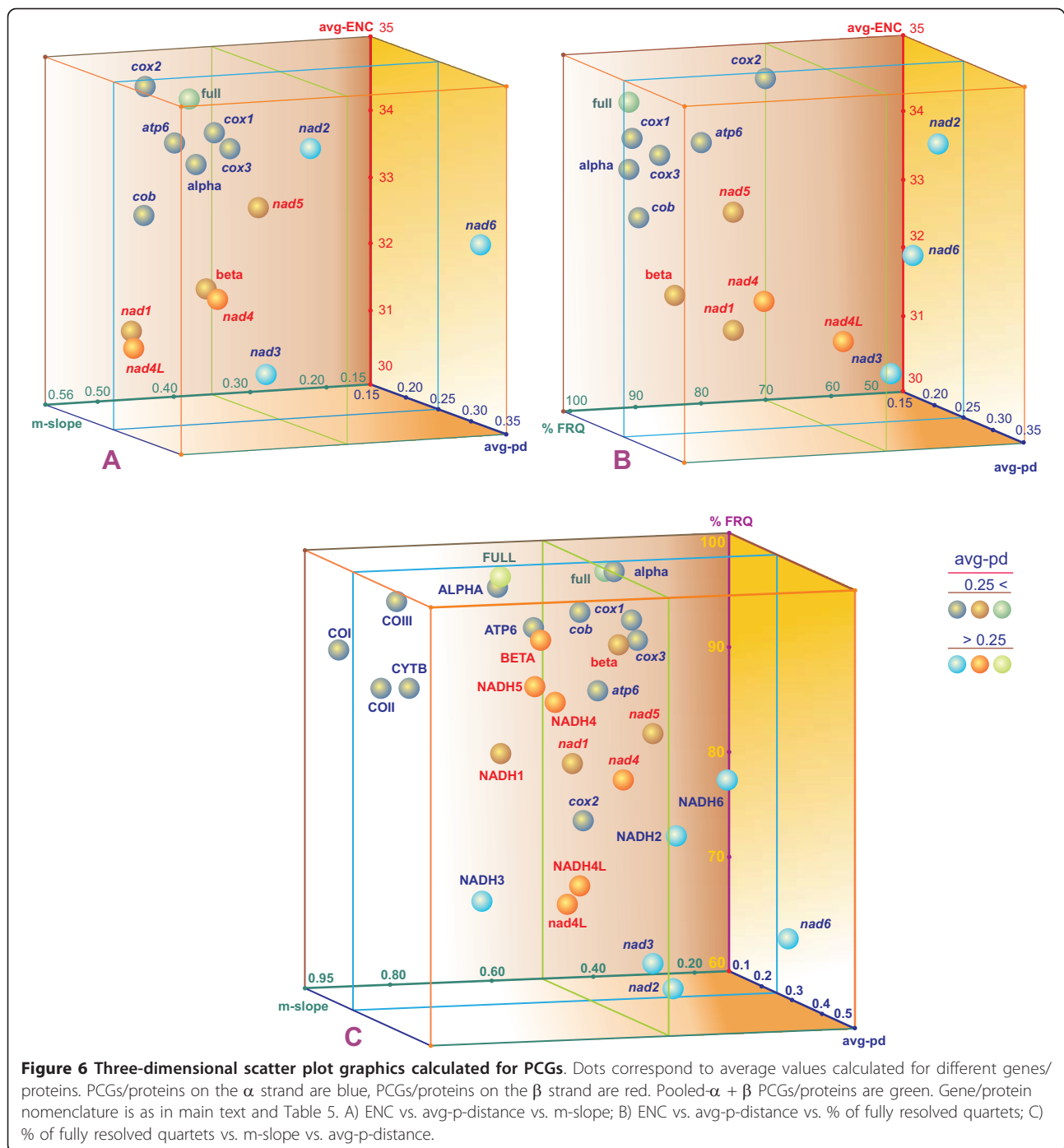
Behavior of PCGs in neuropterid mtDNAs

Different parameters have been used to evaluate the performance of mitochondrial single/combined PCGs for phylogenetic, population genetics, and taxon molecular identification purposes (e.g., [11,12]). The suitability of different PCGs is linked to, influenced by the evolutionary history of each molecular marker. The calculation of p-distance values has been used to test the level of divergence within each PCG (e.g., [11]). Recently, codon usage has also been investigated to study the differentiation of mitochondrial PCGs [12]. In the present paper, the evolutionary behavior of PCGs, at both the nucleotide and amino acid levels, was studied in a combined analysis based on the calculation/estimation of p-distances, effective number of codon usage (ENC), phylogenetic signal and the saturation of the substitution process present in single/combined PCGs.

The percent of fully resolved quartets (%FRQ), obtained in a maximum likelihood mapping analysis, was used as a measure of phylogenetic signal [70]. The saturation of the substitution process was established by calculating the m-slope of the regression line passing through the origin in a scatter plot analysis of p-distances vs. GTR+I+G - mtREV24/JTT + G distances (see Methods for further details).

Parameters were estimated for pooled- $\alpha+\beta$, pooled- α , pooled- β , and single PCGs at both the nucleotide and amino acid levels, with the exception of *atp8*. This gene was not analyzed individually because it contains a limited number of codons (< 55) to provide reliable estimations of ENC [71]. Conversely, *atp8* was included in the pooled- $\alpha+\beta/-\alpha$ sets. The results of the global analysis are summarized in Figure 6 and Table 5.

As a general rule, proteins, individually or pooled, exhibited levels of saturation of the substitution process lower than those observed in the nucleotidic counterparts. This result can be explained in terms of the



substitution process, which permits more alternatives in the amino acid replacements. The m-slope values show that all PCGs/proteins experienced some degree of saturation of the substitution process. This phenomenon was particularly marked at the third codon positions of some PCGs, where the GTR+I+G-distance values largely exceeded 1 (data not shown), a result strongly supporting a complete saturation of the substitution process [72].

Pooled- α + β PCGs/proteins and pooled- α PCGs/proteins showed the best phylogenetic signals and clearly surpassed pooled- β PCGs/proteins. Because pooled- α + β and pooled- α proteins also exhibited a less pronounced saturation of the substitution process than their nucleotide counterparts, they represented the first choice as markers to investigate the phylogenetic relationships among Neuropterida at the inter-order taxonomic level.

Table 5 Phylogenetic signal, saturation of substitution process, p-distances, and effective codon usage in neuropterid PCGs

pcg/PROTEIN	%FRQ	m-slope	avg-pd	avg-ENC
pooled-($\alpha+\beta$) (full)	100	0.466	0.247 \pm 0.034	34.758 \pm 2.640
POOLED-($\alpha+\beta$) (FULL)	100	0.684	0.262 \pm 0.062	n.a.c.
pooled- α (alpha)	100	0.458	0.248 \pm 0.030	33.783 \pm 2.111
POOLED- α (ALPHA)	98.6	0.676	0.238 \pm 0.055	n.a.c.
pooled- β (beta)	92.86	0.442	0.246 \pm 0.040	31.898 \pm 1.603
POOLED- β (BETA)	94.3	0.625	0.297 \pm 0.073	n.a.c.
<i>nad2</i>	61.43	0.392	0.346 \pm 0.037	34.321 \pm 3.279
NADH2	77.1	0.436	0.430 \pm 0.067	n.a.c.
<i>cox1</i>	94.28	0.384	0.189 \pm 0.023	33.991 \pm 1.584
COI	91.4	0.908	0.105 \pm 0.035	n.a.c.
<i>cox2</i>	75.71	0.491	0.209 \pm 0.036	34.840 \pm 4.054
COII	88.6	0.867	0.174 \pm 0.076	n.a.c.
<i>atp6</i>	88.57	0.482	0.244 \pm 0.034	34.105 \pm 2.770
ATP6	94.3	0.591	0.215 \pm 0.061	n.a.c.
<i>cox3</i>	98.86	0.390	0.220 \pm 0.032	33.890 \pm 2.895
COIII	97.1	0.845	0.191 \pm 0.059	n.a.c.
<i>nad3</i>	62.86	0.394	0.280 \pm 0.032	30.720 \pm 4.398
NADH3	70	0.751	0.313 \pm 0.068	n.a.c.
<i>nad5</i>	84.28	0.377	0.249 \pm 0.042	33.086 \pm 2.306
NADH5	89.9	0.639	0.300 \pm 0.078	n.a.c.
<i>nad4</i>	80.00	0.436	0.254 \pm 0.044	31.771 \pm 2.270
NADH4	88.6	0.608	0.315 \pm 0.075	n.a.c.
<i>nad4L</i>	68.57	0.551	0.262 \pm 0.044	31.150 \pm 4.875
NADH4L	71.4	0.571	0.336 \pm 0.067	n.a.c.
<i>nad6</i>	65.71	0.171	0.349 \pm 0.045	32.722 \pm 3.411
NADH6	82.8	0.360	0.468 \pm 0.090	n.a.c.
<i>cob</i>	95.71	0.498	0.214 \pm 0.030	32.932 \pm 2.162
CYTB	88.6	0.639	0.182 \pm 0.059	n.a.c.
<i>nad1</i>	81.43	0.516	0.217 \pm 0.038	31.225 \pm 2.100
NADH1	82.9	0.673	0.246 \pm 0.072	n.a.c.

Pcg-PROTEIN, protein coding gene/relative protein; %FRQ, percent of fully resolved quartets obtained in a maximum likelihood mapping analysis [70]; m-slope, slope of the regression line passing through the origin in a scatter plot analysis of p-distances vs. GTR+I+G - mtREV24/JTT + G + G distances distances; avg-pd, average p-distance; avg-ENC, average-ENC (effective number of codon usage); n.a., not applicable calculation.

The single genes exhibiting the best phylogenetic signals at the inter-order taxonomic level were *cob*, *cox1* and *cox3* (%FRQ > 94.00). Intermediate values were observed in *atp6*, *cox2*, *nad4*, *nad5* and *nad1* (89.00 < %FRQ > 75.00) The lowest signals were present in *nad2*, *nad3*, *nad4L* and *nad6* (%FRQ < 70.00).

Pooled and single PCGs having the best phylogenetic signals usually exhibited a combination of high avg-ENCs, small avg-pds, and high m-slopes in their protein counterparts (e.g., *cox1*), but exceptions occurred. Conversely, PCGs with low phylogenetic signals usually showed high avg-pds and low values of m-slope at both the nucleotide and amino acid levels (e.g., *nad2*).

Reduction of phylogenetic signal with respect to the best-performing PCGs can be explained in terms of three different processes/properties: a) accelerated evolution; b) retarded evolution; and c) possession of an intrinsically lower phylogenetic signal.

A marked accelerated evolution mechanism can be invoked for *nad2*, *nad3* and *nad6*. These PCGs showed avg-pds, at both the nucleotide and amino acid levels, which were clearly higher than those observed in *cox1*, *cox3* and *cob*. Furthermore, their m-slope values favored the occurrence of a more pronounced saturation of the substitution process, a finding corroborated by the lowest values of fully resolved quartets. All of these results are compatible only with an acceleration of the substitution process. Thus, *nad2*, *nad3* and *nad6* should be excellent markers for studies performed at medium/very low taxonomic levels (i.e., family, genus, species).

The mechanism of accelerated evolution, but less pronounced than that described above, can be invoked also for *atp6*, *nad5* and *nad4*. These genes showed avg-pds higher than *cox1*, *cox3* and *cob*, at both the nucleotide and amino acid levels. They exhibited phylogenetic signals higher than *nad2*, *nad3* and *nad6* but lower than the best-performing PCGs. Finally, their m-slope values were comparable to those of *cox1*, *cox3* and *cob*. All these findings suggest that *atp6*, *nad5* and *nad4* evolved slightly faster than the best-performing PCGs and that these genes can be optimally used at the order or lower taxonomic levels.

The *nad1* and *cox2* genes had avg-pds similar to or even smaller than *cox1*, *cox3* and *cob*, while their levels of saturation of the substitution process appeared to be less marked than PCGs exhibiting the highest %FRQ values. These results suggest that the reduction of phylogenetic signal in *cox2* and *nad1* was probably due to a slower rate of divergence of these two genes in comparison with *cox1*, *cox3* and *cob*. Thus, they should be considered favored markers at high/very high taxonomic ranks.

The behavior of *nad4L* provided contradictory evidence. The low %FRQ and relatively high avg-pd value suggested that this is a fast evolving gene and that the loss of phylogenetic signal was due to an accelerated rate of multiple substitutions. However, the m-value was the highest obtained in the analysis and did not favor a strong saturation of the substitution process in the evolution of this gene. Thus, it could be that even evolving in a way not dissimilar to *cox1*, *cox3* and *cob*, *nad4L* has an intrinsically lower phylogenetic signal due to its reduced size.

Taken together, these data indicate that the 13 PCGs encoded by mtDNA exhibit complex evolutionary trajectories that can be investigated using the combination of

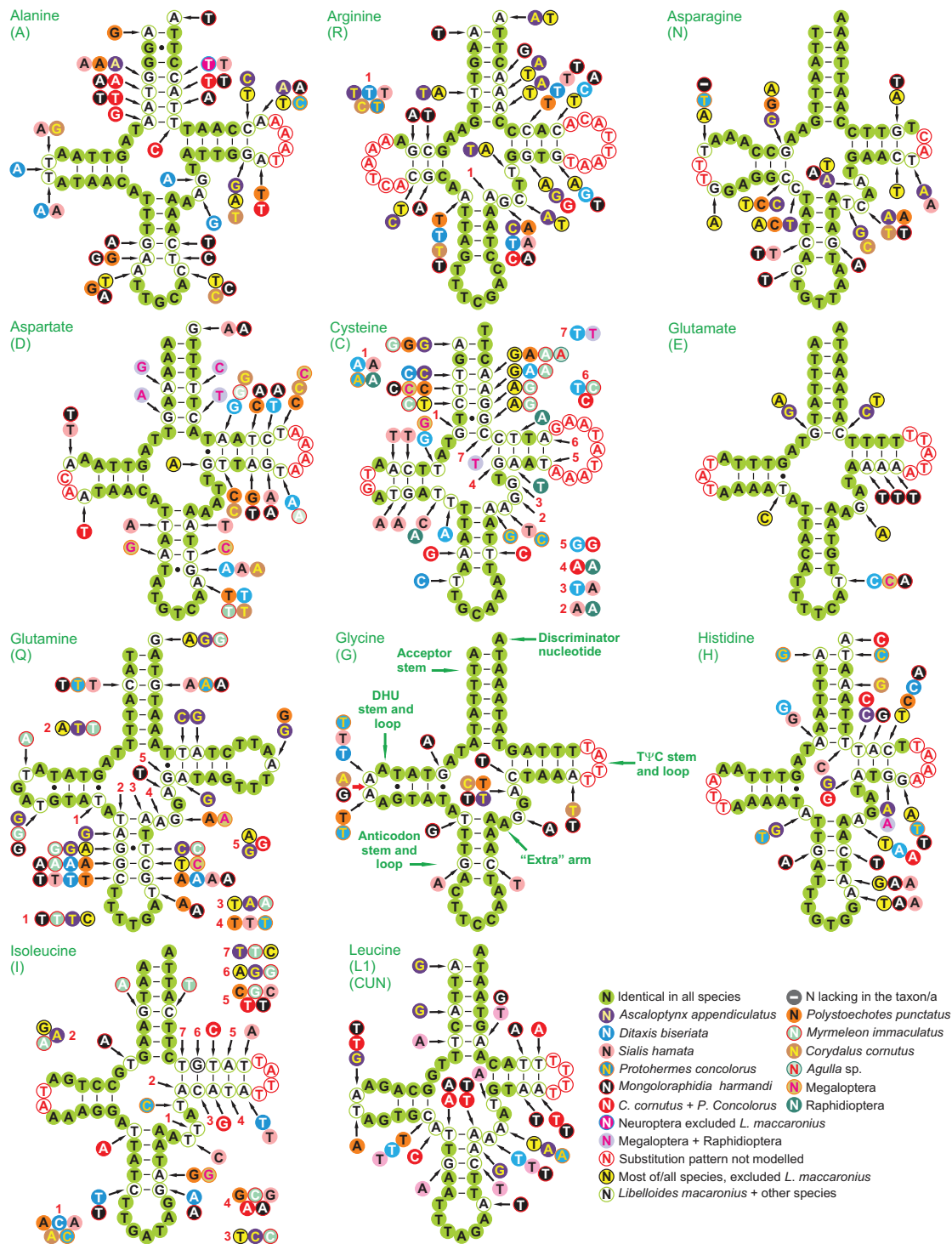


Figure 7 Secondary structure of tRNA families (*trnA-trnL1*) in neuropterid mtDNAs. The nucleotide substitution pattern for each tRNA family was modeled using as reference the structure determined for *L. maccaronius*.

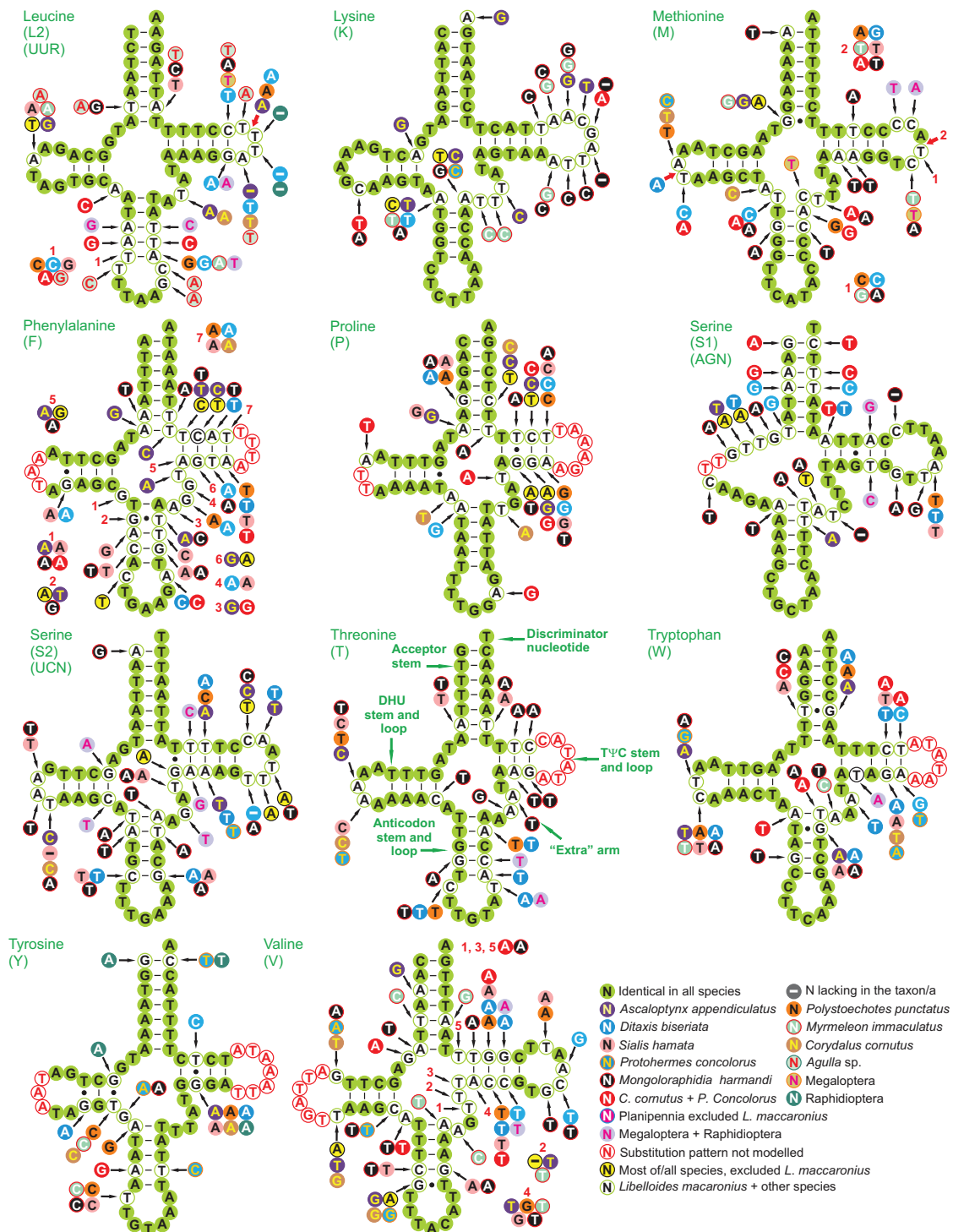


Figure 8 Secondary structure of tRNA families (*trnL2-trnV*) in neuropterid mtDNAs. The nucleotide substitution pattern for each tRNA family was modeled using as reference the structure determined for *L. macaronius*.

the parameters considered above. No single parameter seems able to fully describe the behavior of PCGs.

Comparison of tRNA structure in neuropterid genomes

The mtDNA of *L. macaronius* contained all 22 tRNAs genes found in other neuropterid mtDNAs and typical of animal mitochondrial genomes [1]. All neuropterid mtDNAs had 14 α -strand tRNAs and 8 β -strand tRNAs (Figures 1, 2). Among the 22 tRNAs, only *trnS1* did not exhibit the common cloverleaf structure, due to the absence of DHU stem. Loss of this stem in *trnS1* is a feature common to many insect mtDNAs (e.g., [12]). The results of comparative analyses on secondary structures of neuropterid tRNAs are provided in Figures 7, 8. Multiple alignments of tRNAs genes were produced, and the percent of identical nucleotides (%INUC) was calculated for each group of orthologous sequences (Additional file 5, Table S3).

The pattern of nucleotide conservation was markedly α strand-biased. Indeed, *trnG*, *trnK*, *trnN*, *trnM* and *trnE*, which showed the highest levels of nucleotide conservation (%INUC \geq 70), were all located on the α strand.

TrnL2, *trnP*, *trnQ*, *trnS1*, *trnS2*, *trnT*, *trnW* and *trnY* showed 60 < %INUC < 70. Only three of them were located on the β strand (Figure 2). Seven remaining tRNAs exhibited 50% < %INUC < 60. Four of them were located on the α strand, i.e., *trnI*, *trnA*, *trnD*, and *trnR*, while three were on the β strand, i.e., *trnL1*, *trnH* and *trnV*. %INUC values < 50% characterized the β -strand genes *trnF* and *trnC*. While the level of conservation was positively α strand-biased, no clear pattern could be identified for single tRNAs located at various points along the genome. For example, *trnK*, one of the most conserved tRNAs, was located immediately upstream of *trnD* (Figure 2), a much less conserved gene (see above).

The tRNAs closest to the control region (i.e., *trnV* and *trnI*), which is one of the start points of mtDNA replication, were not among most/less conserved. The same applied to *trnS2*, immediately upstream of the s2 spacer, where another center of mtDNA replication was located (see below). The abundance of codon families did not appear to be linked to the level of tRNA conservation. Indeed, none of the most abundant codon families (Leu2, Ile and Phe) exhibited the highest %INUCs.

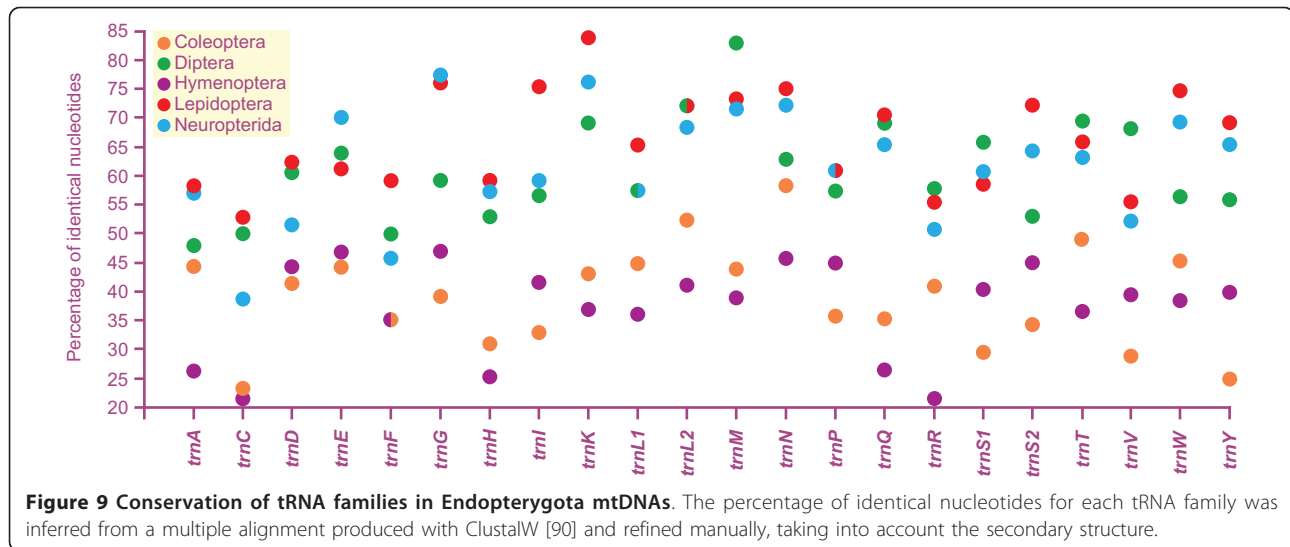
Irrespective of the level of conservation, some tRNAs presented mismatched pairs in stems (e.g., A-A in the anticodon stem of *trnK*; T-T in the acceptor stem of *trnM*). These mismatches are common in arthropod mtDNAs and can be corrected through editing processes (e.g., [73]) or may represent unusual pairings [74]. Further research at the transcript level is necessary to better characterize this feature in neuropterid mtDNAs.

In the most conserved tRNAs the nucleotide substitutions are largely restricted to T Ψ C and DHU loops and extra arms (Figures 7, 8), with changes on acceptor and anticodon stems reduced to 0-3 fully compensatory base changes (cbcs) (e.g., G-C vs. A-T in the anticodon stem of *trnG*) and/or hemi-cbc (e.g., G-T vs. A-T on the acceptor stem of *trnM*) [75]. Note that in *trnG*, the most conserved tRNA, it was not possible to model the substitution pattern in the T Ψ C loop due to a high level of variation among orthologous sequences (Figure 7). With the increased variation, the number of cbcs and hemi-cbcs increased in stems, usually with a more marked substitution process in the T Ψ C stem (e.g., *trnI*, *trnV*). In several tRNAs it was not possible to model properly the substitution patterns in T Ψ C and DHU loops due to the high level of divergence among sequences. Cbcs and hemi-cbcs were restricted to a single species or characterized taxa at a higher taxonomic rank (family/order). An example of the first type is the G-C pair found in the *trnH* acceptor arm of *P. concolorus*, which was mirrored by A-T in all other neuropterids (Figure 7). An example of a full cbc characterizing a unique family is the G-C pair found in the acceptor stem of *trnEs* of family Ascalaphidae (*A. appendiculatus* and *L. macaronius*), while other taxa exhibited the A-T pair (Figure 7). A substitution pattern involving two full cbcs characterizing the *trnS1* acceptor arm of *C. cornutus* and *P. concolorus* (Corydalidae megalopterans) is another example of high-taxonomic rank cbcs (Figure 8). Figures 7, 8 depict several more examples.

The presence of hemi-cbcs and, particularly, full cbcs characterizing taxa at different taxonomic levels underscores the potential phylogenetic value of tRNA sequences, especially when secondary structures are taken into account. Until now, tRNAs have been rarely considered in the study of insect phylogenetic relationships [76,77]. Our results suggest that these markers deserve much more attention and should be more routinely used, as demonstrated by analyses performed in other animal groups (e.g., [78]).

The study of the level of conservation in tRNA families was extended to other endopterygotan orders in which at least two fully sequenced mtDNAs exist. The results of this analysis are summarized in Figure 9.

The pattern of %INUC variation exhibited by codon families was not consistent among different endopterygotan taxa. The highest level of conservation was observed in lepidopteran *trnK* (%INUC = 84.51), while the minimum was found in hymenopteran *trnC* (%INUC = 20.55). Lepidoptera had nine codon families with %INUC \geq 70, whereas five such families were found in Neuropterida, two in Diptera, and none in Coleoptera or Hymenoptera. Lepidopteran codon families were the most conserved (avg-%INUC = 66.25



± 8.49), immediately followed by neuropterid codon families (avg-%INUC = 61.46 ± 9.84). Dipteran codon families showed slightly more divergence (avg-%INUC = 60.84 ± 8.59), while coleopteran (avg-%INUC = 38.92 ± 8.84) and hymenopteran (avg-%INUC = 37.19 ± 8.22) codon families exhibited much lower levels of conservation.

Neuropterid %INUCs were in some cases very similar (e.g., *trnA*, *trnH*, *trnM*) or even surpassed (*trnE*, *trnG*, *trnS1*) their lepidopteran counterparts (Figure 9). This finding is striking because it indicates that the level of variation among the three neuropterid orders is very low, taking into account that Lepidoptera have very conserved tRNAs among Insecta [11]. Conversely, Hymenoptera and Coleoptera had diverging codon families. Indeed, all hymenopteran tRNAs showed %INUC values ≤ 47.14 , while no coleopteran tRNA exhibited %INUC > 58.21 . The %INUC scores indicate a clear-cut discrepancy between the morphological diversity observed in neuropterid orders and the low rate of substitution that characterizes the evolution of their codon families. If the level of divergence is considered a measure of taxonomic diversity, then the %INUC values would suggest that Neuroptera, Megaloptera and Raphidioptera should be treated as members of a single order.

Comparison of neuropterid *rrnS* and *rrnL* structures

The inferred secondary structure model for *rrnS* of *L. macaronius* is provided in Figure 10. This is the first prediction for a neuropterid insect. The overall structure, including three domains and 34 helices (progressively numbered in Figure 10), is largely in agreement with those proposed for other endopterygote orders (i.e. Coleoptera, Diptera, Hymenoptera, and Lepidoptera) [11,17,74,79]. A limited number of non-canonical

pairings (e.g., G-A on helix H23-434) were present in the *rrnS* of *L. macaronius*. The multiple alignment of neuropterid *rrnS*s spanned 812 positions and contained 353 conserved (43.47%) and 459 variable (56.53%) positions, respectively. Nucleotide variability among domains and helices was unevenly distributed (Figure 10).

The inferred secondary structure model for the *rrnL* of *L. macaronius* is provided in Figure 11. This is the first prediction for a neuropterid insect. The structure of this *rrnL* largely overlaps with previously published structures for endopterygote insects (i.e. Coleoptera, Diptera, Hymenoptera, and Lepidoptera) [11,17,74,79]. It presents the five canonical domains (I-II, IV-VI) of insect *rrnL*s that do not contain domain III [74]. In the present paper domain I was fully modeled. This is the first time that a secondary structure has been provided for this domain, which in previous studies has been left unmodeled (e.g., [8,11,56,74,79]). The predicted secondary structure of domain I included eight helices and is consistent among all neuropterid taxa (Additional file 6, Figure S3). The proposed structure can also be fitted to *rrnL* of *D. melanogaster* (Additional file 6, Figure S3), further extending the taxonomic range. Domain I contained a limited number of non-canonical pairings (Additional file 6, Figure S3). More studies will be necessary to corroborate the validity of the new structure presented here. Taking into account the secondary structure of domain I, *L. macaronius rrnL* includes 50 helices. The multiple alignment of neuropterid *rrnL*s extended over 1376 positions and contains 591 conserved (42.95%) and 785 variable sites (57.05%). Conserved nucleotides were unevenly distributed throughout the *rrnL* secondary structure, with highest level of invariable positions located on domain IV and lowest level observed in domains I-II (Figure 11).

Libelloides macaronius *rrnS*

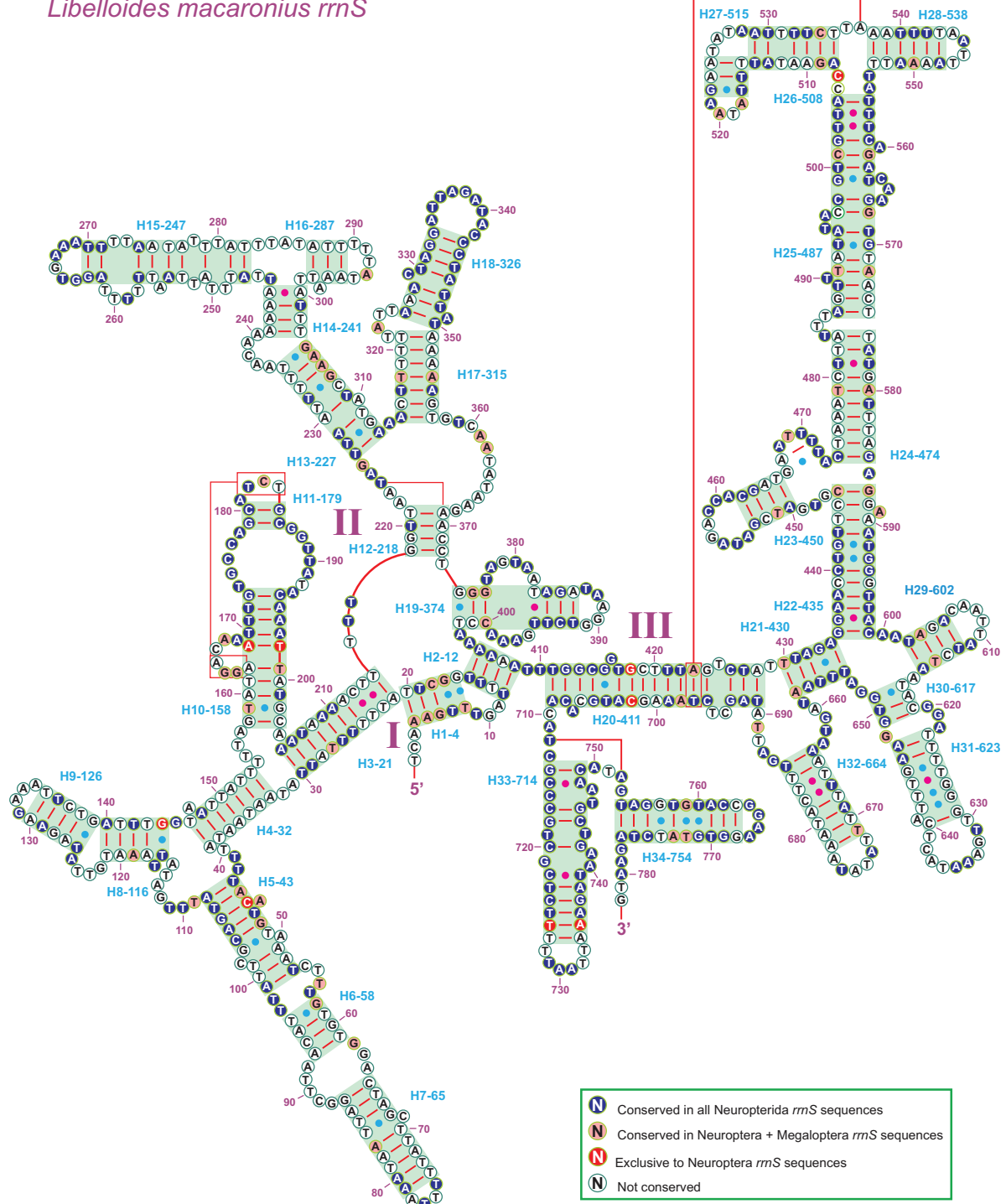


Figure 10 Secondary structure of *L. macaronius* *rrnS*. Each helix is numbered progressively from the 5' to the 3' end together with the first nucleotide belonging to the helix itself. Domains are labeled with Roman numerals. Tertiary structures are denoted by boxed bases joined by solid lines. Watson-Crick pairs are joined by dashes. GT pairs are joined by a blue dot, while other non-canonical pairs are connected by a red dot.

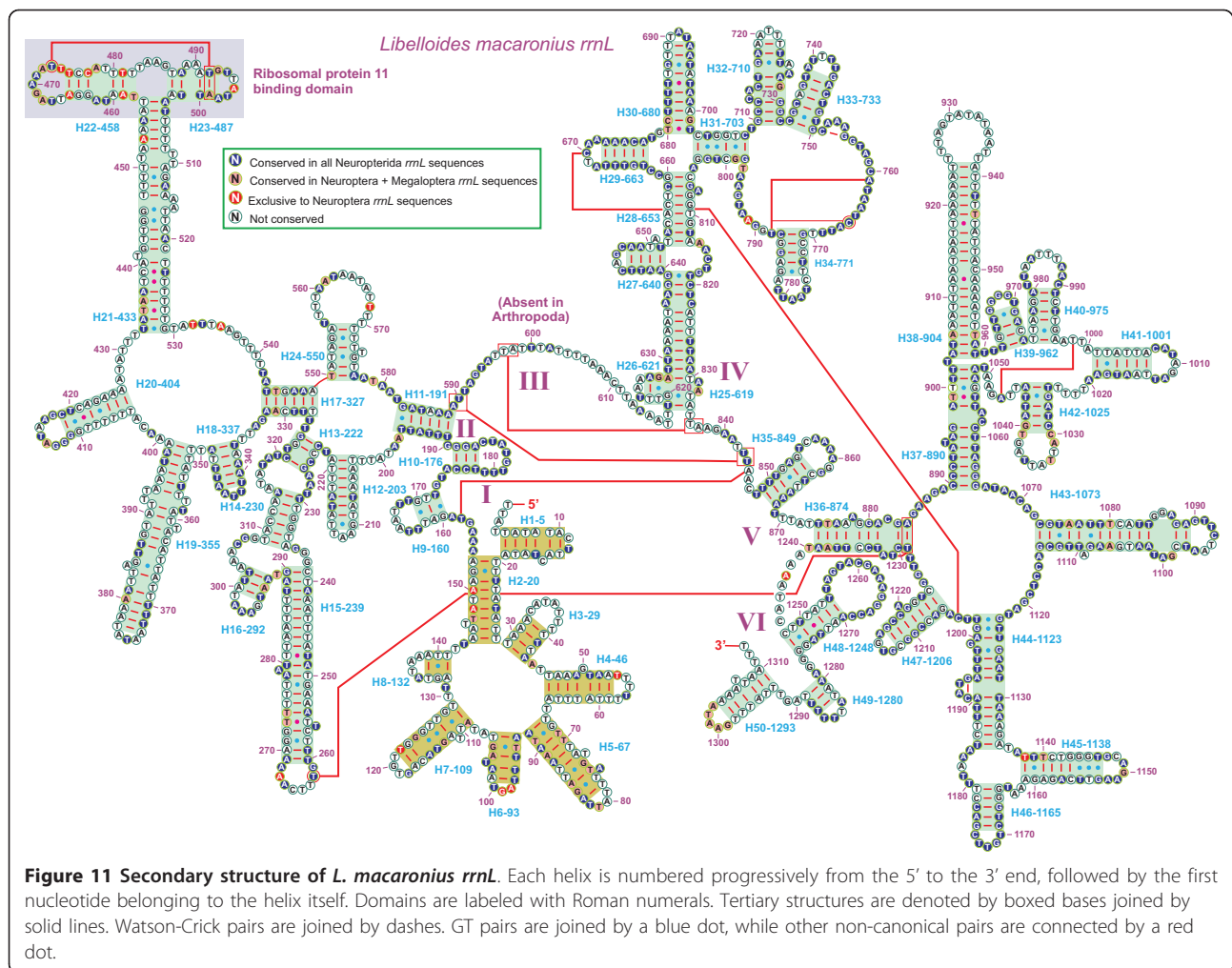


Figure 11 Secondary structure of *L. macaronius rrnL*. Each helix is numbered progressively from the 5' to the 3' end, followed by the first nucleotide belonging to the helix itself. Domains are labeled with Roman numerals. Tertiary structures are denoted by boxed bases joined by solid lines. Watson-Crick pairs are joined by dashes. GT pairs are joined by a blue dot, while other non-canonical pairs are connected by a red dot.

Comparison of neuropterid mtDNA genomic spacers

The *L. macaronius* mtDNA contained 14 non-coding portions (ncps), extending from 1 to 1049 nucleotides, interspersed through PCGs, tRNAs and *rrnL* and *rrnS* genes (Figure 2). Three of them spanned more than 15 bp and are labeled as spacers s1-s3 in Figures 2 and 12. More or less similar patterns, but not fully overlapping, were observed in other neuropteran mtDNAs, where the ncps varied from a maximum of 15 (*D. biseriata*) to a minimum of 10 (*P. punctatus*), with 13 non-coding portions found in *A. appendiculatus*. The dobsonflies *C. cornutus* and *P. concolorus* exhibited the same number (10) of ncps even if their distributional patterns did not coincide. Conversely, the other megalopteran, *S. hamata*, had 12 ncps. Finally, the raphidiopteran *M. harmandi* exhibited the most compact mtDNA, having only six ncps (Figure 12).

The spacers of *L. macaronius* are described more in detail below. The s1 spacer was located between *trnI* and *trnQ* and spanned 55 bp. Spacer s1 was present in

all partly/fully sequenced mtDNAs available for Neuroptera, i.e., *A. appendiculatus*, *P. punctatus*, *M. immaculatus* and *D. biseriata* [13,14], present paper)]. It was also found in the megalopteran *S. hamata*, while it was absent in the other megalopterans *C. cornutus* and *P. concolorus* and in the raphidiopteran *M. harmandi* (Figure 12) [13-15]. The *L. macaronius* s1 can be aligned confidently only with the counterpart found in *A. appendiculatus*.

The s1 sequences of *L. macaronius* and *A. appendiculatus* shared a CCCCCC repeat located near the boundary with *trnI* and the CAA(A/G)TTAA(A/C)TAAAT (TA/GT)A(C/T)GCA motif adjacent to *trnQ*. *A. appendiculatus* and *L. macaronius* belong to the same family, Ascalaphidae. Thus, it seems that s1 sequences, when present, diverge very fast among different families of the same order.

The *L. macaronius* s2 was 20 bp long and was located between *trnS2* and *nad1*. The s2 spacer was present in all partly or fully sequenced neuropterid mtDNAs

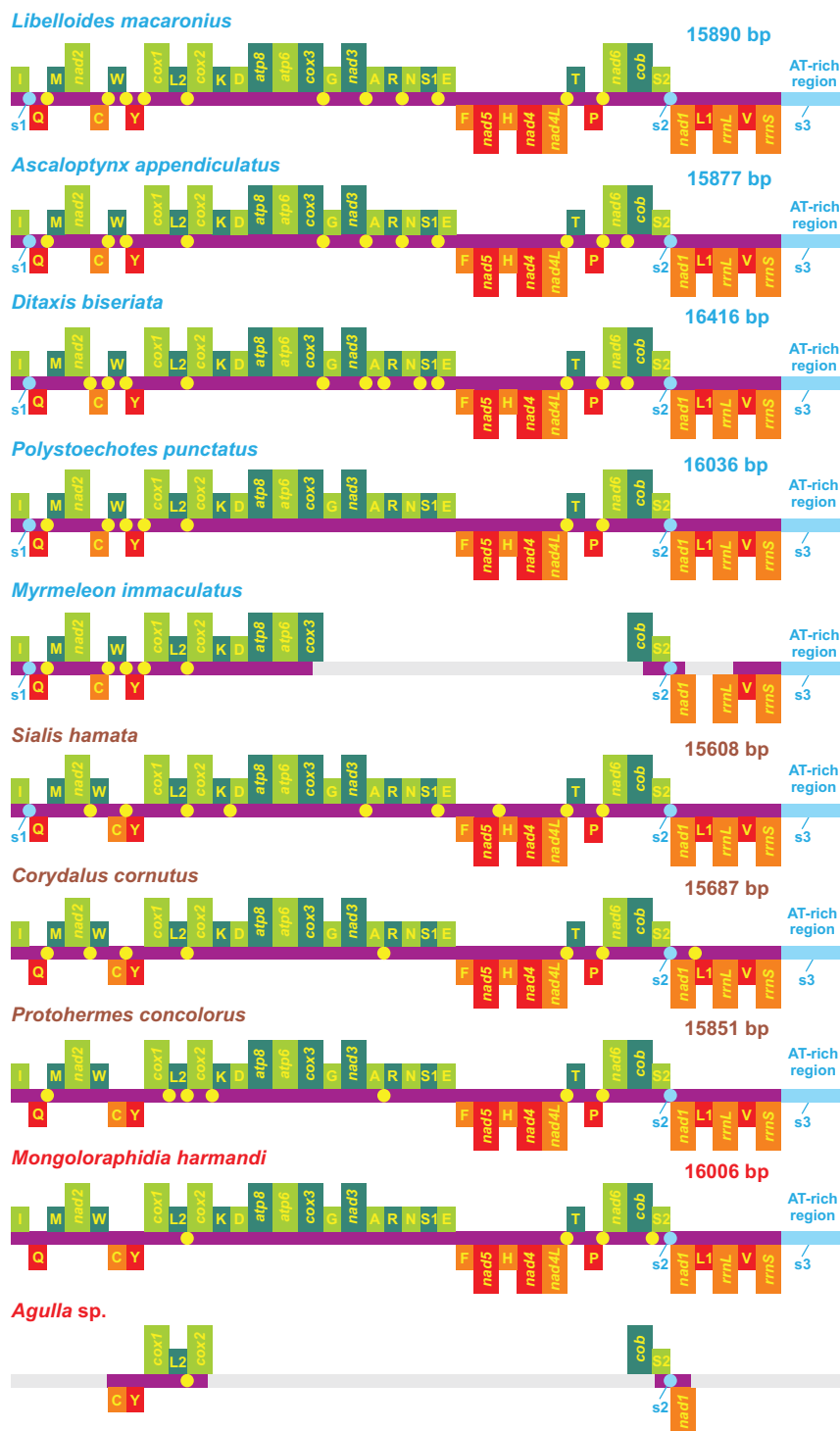


Figure 12 Distribution of non-coding portions in neuropterid mtDNAs. The non-coding portions are presented as cyan/yellow dots. s1-s3, spacers 1-3. Gene nomenclature is as in Figures 1 and 2. Genomes, genes and non-coding portions are not drawn to scale, for reasons of clarity. The unknown portions of partial mtDNAs are gray.

(Figure 12) [[13-15], present paper]. The s2 spacer is supposed to contain [13] the binding site for the DmTFF bidirectional transcription termination factor [80]. Spacer s2 is a common feature of insect mtDNAs (e.g., [12]).

The *L. macaronius* s3 spacer was composed of a 1049-bp AT-rich region. The A+T content was 84.46%. This finding implies that s3 contained 10% more A+T than the whole α strand. Conversely, s3 was much less AT-skewed than the complete α strand (0.029 vs. 0.072). Strings of repeated single nucleotides (T)₃₋₈, (A)₃₋₁₁, (C)₂₋₅, (G)₂₋₅ characterized the s3 sequence. Furthermore, the abundance of A/T was mirrored by the presence of repeated motifs containing mostly/only A/T, a common feature of insect control regions [81]. These motifs were represented by shorter identical strings as well as longer consensus patterns sharing 75% of identical positions. The motif AT was the most abundant, occurring 192 times. ATA occurred 70 times, ATAT 29, AATAATTA 5, and ATATAAATA 6. Finally, five distinct A/T-rich 12-mers were repeated twice each (data not shown). When a 75% minimum identity was allowed among repeated motifs, two strings were identified, each containing 37 nucleotides and extending, respectively, from base 551 to base 591 and from 811 to 847. Their consensus sequence was ATATTA (C/T) ATAT (GT/AA) ATA (AT/TG) TA (T/A) TTATTA (A/T) TATATAAAT. Thus, the *L. macaronius* AT-rich region contains several repeated motifs of different sizes. A similar trend also occurs in other neuropterids (data not shown).

Pair-wise as well as multiple alignments between/among neuropterid AT-rich regions were performed by applying very relaxed alignment parameters (i.e., gap-opening = 1 vs. standard 15; and gap-extending = 3 vs. standard 6.6). This strategy was applied to maximize the matches of these fast-evolving genomic portions (see below). This approach allowed us to align quite diverse sequences but did not guarantee that the positional homology principle was retained when the more diverging sequences were compared. The obtained results are summarized in Table 6.

What is immediately evident is that the level of conserved positions dropped rapidly when comparisons were extended above the taxonomic rank of family. Indeed, *L. macaronius* and *A. appendiculatus*, both members of the family Ascalaphidae, shared 71.59% nucleotide identity and a common GC-rich motif TCCCCGCCCCCAGGAT located 96 bp downstream of the 5'-end of *rrnS* in *L. macaronius* mtDNA. This motif could be a molecular signature for the whole family, but a better taxon sampling is necessary to assess this point.

When all Neuroptera were considered, the identical positions diminished to 30.15% in the multiple alignment. Thus, a substantial drop in nucleotide identity

Table 6 Pairwise and multiple alignments of neuropterid control regions

ORD		idNP	% ID	ALNL
NEU	<i>L. macaronius</i> vs. <i>A. appendiculatus</i>	809	71.59%	1130
NEU	<i>L. macaronius</i> vs. <i>P. punctatus</i>	765	63.80%	1199
NEU	<i>L. macaronius</i> vs. <i>D. biseriata</i>	845	55.78%	1515
NEU	<i>L. macaronius</i> vs. <i>M. immaculatus</i>	721	65.13%	1107
NEU	<i>A. appendiculatus</i> vs. <i>M. immaculatus</i>	731	65.56%	1115
NEU	<i>A. appendiculatus</i> vs. <i>D. biseriata</i>	863	56.85%	1518
NEU	<i>A. appendiculatus</i> vs. <i>P. punctatus</i>	787	64.88%	1213
NEU	<i>P. punctatus</i> vs. <i>D. biseriata</i>	948	62.00%	1529
NEU	<i>P. punctatus</i> vs. <i>M. immaculatus</i>	748	63.02%	1187
NEU	<i>D. biseriata</i> vs. <i>M. immaculatus</i>	813	53.88%	1509
NEU	Neuroptera multiple alignment	487	30.15%	1615
MEG	<i>C. cornutus</i> vs. <i>P. concolorus</i>	762	63.50%	1200
MEG	<i>C. cornutus</i> vs. <i>S. hamata</i>	632	62.27%	1015
MEG	<i>P. concolorus</i> vs. <i>S. hamata</i>	658	57.52%	1144
MEG	Megaloptera multiple alignment	561	45.91%	1222
	Neuropterida orders multiple alignment	273	14.80%	1845

ORD, order; idNP, identical nucleotide in the alignment position; % ID; percentage of identical nucleotides; ALNL, length of the alignment; NEU Neuroptera, MEG, Megaloptera.

shared by AT-rich regions occurs at the order level, even permitting very permissive gap costs.

Similarly, *C. cornutus* and *P. concolorus*, both members of the family Corydalidae, shared 63.50% nucleotide identity in their control regions. Conversely, the identity level decreased to 45.91% when all megalopteran AT-rich regions were compared. Finally, the alignment at the interordinal taxonomic rank of neuropterid control regions produced only 14.80% identical positions. Provided that gaps are quite numerous, this result leads to the conclusion that the AT-rich region is a fast-evolving genomic region. This behavior of neuropterid control regions is a common feature of insect AT-rich regions (e.g., [12]) and is shared with other animal groups. In this respect, a well-documented group is represented by mammals (e.g., [82,83]).

The utility of the AT-rich region as a phylogenetic marker should be most effective at low taxonomic rank (family level and below).

Conclusions

The mtDNA of *L. macaronius*, the fourth genome sequenced for the order Neuroptera, exhibits the peculiar translocation of the *trnC* gene with respect to the ancestral gene order of insects. This structural modification represents an exclusive feature of partly or fully sequenced neuropteran mtDNAs and could be a peculiar genomic marker characterizing the entire order Neuroptera.

The analysis of the AT%, AT-skew and GC-skew parameters, which were performed on a set of 84

holometabolous insect mtDNAs, shows that these characteristics exhibit complex patterns of variations. Such patterns can be linked or completely unlinked in their variation process.

The neuropterid mtDNAs sequenced to date use the standard invertebrate mitochondrial genome. The distribution of codon families in the PCGs located on the α and β strands is influenced by both structural/functional requirements of the corresponding proteins and by the base composition and AT-/GC-skews. The comparison among orthologous neuropterid PCGs shows that different genes have been subject to different rates of molecular evolution and form a pool of markers suitable for different phylogenetic purposes.

The evolution of neuropterid tRNAs seems to have been variable both in terms of sequence conservation and nucleotide substitution processes.

Neuropterid *rrnL* and *rrnS* structures, as demonstrated by the models produced for *L. macaronius*, appear similar to those determined for other insects. In their structural domains, they show diverse levels of conservation, influenced by different rates of substitution.

An important advance in our comprehension of the structure of *rrnL* is the production of a model for its domain I. To our knowledge, this is the first time that such a structural modeling has been attempted for an arthropod *rrnL*.

Neuropterid mtDNAs are punctuated by non-coding portions highly variable in size. Among them, the most extended segment is the control region (AT-rich region), which appears to be a fast-evolving genomic region characterized by short to medium-size repeated motifs/AT-rich patterns.

Methods

Sample origin and DNA extraction

The *L. macaronius* specimen used to determine the mtDNA was collected in June 2007 by EN on arid prairies (45°47'35.75"N 13°35'50.63"E, 98 m elevation) located in the Triestine karst along the road connecting San Giovanni al Timavo to Medeazza (Friuli Venezia Giulia region, northeastern Italy).

Total DNA was purified using the Invisorb DNA extraction kit (Invitec). The quality of DNA was assessed through electrophoresis in a 1% agarose gel and staining with SYBR-safe DNA gel stain (Invitrogen).

PCR amplification and sequencing of *L. macaronius* mtDNA

PCR amplification was performed using a mix of insect universal primers and primers specifically designed against the *L. macaronius* sequences [84,85]. The complete list of successful primers is available upon request. The PCR products were visualized by electrophoresis in

a 1% agarose gel and staining with SYBR-safe DNA gel stain. Each PCR product represented by a single electrophoretic band was purified with the ExoSAP-IT kit (Amersham Biosciences) and directly sequenced. Sequencing of both strands was performed at the BMR Genomics service (<http://www.bmr-genomics.it/>) on automated DNA sequencers mostly employing the primers used for PCR amplification.

Sequence assembly and annotation

The mtDNA final consensus sequence was assembled using the SeqMan II program from the Lasergene software package (DNASTar, Madison, WI). Gene and strand nomenclature used in this paper were described by Negrisola et al. [86].

Sequence analysis was performed as follows. Initially, the mtDNA sequence was translated *in silico* into putative proteins using the Transeq program available at the EBI web site. The true identity of these polypeptides was established using the BLAST program available at the NCBI web site [87,88]. Gene boundaries were determined as follows. The 5' ends of PCGs were inferred to be at the first legitimate in-frame start codon (ATN, GTG, TTG, GTT) in the open reading frame that was not located within the upstream gene encoded on the same strand [64,89]. The only exceptions were *atp6* and *nad4*, which overlap with their upstream genes (*atp8* and *nad4L*, respectively) in many mtDNAs (e.g., [13]). The PCG terminus was inferred to be at the first in-frame stop codon encountered. When the stop codon was located within the sequence of a downstream gene encoded on the same strand, a truncated stop codon (T or TA) adjacent to the beginning of the downstream gene was designated the termination codon. This codon was thought to be completed by polyadenylation to a complete TAA stop codon after transcript processing. Finally, pair-wise comparisons with orthologous proteins were performed with ClustalW program to better define the limits of PCGs [90].

Irrespective of the real initiation codon, formyl-Met was assumed to be the starting amino acid for all proteins, as previously shown for other mitochondrial genomes [91,92].

The transfer RNA genes were identified using the tRNAscan-SE program or recognized manually as sequences having the appropriate anticodon and capability of folding into the typical cloverleaf secondary structure [64,93].

The boundaries of the ribosomal *rrnL* gene were assumed to be delimited by the ends of the *trnV-trnLI* pair. The 3' end of the *rrnS* gene was assumed to be delimited by the start of *trnV*, while the 5' end was determined through comparison with orthologous genes of other previously sequenced neuropterid insect mtDNAs.

The published neuropterid genomes were re-annotated using the criteria listed above. This approach led us to propose different start/end positions for some genes.

Genomic analysis

Nucleotide composition was calculated with the EditSeq program included in the Lasergene software package. GC-skew = $(G-C)/(G+C)$ and AT-skew = $(A-T)/(A+T)$ were used to measure the base-compositional difference between the different strands or between genes coded on the alternative strands [94]. The prediction of the genetic code used by each mtDNA was performed using the GenDecoder web server [67,68].

Relative synonymous codon usage (RSCU) values were calculated with MEGA 4 program [95].

Codon usage in neuropterid mtDNAs was studied by calculating the effective number of codon used index (ENC) with the INCA 2.1 program [71,96]. Multiple alignments of genes were produced with the ClustalW algorithm implemented in MEGA 4 [90,95]. Both p-distances and the numbers of different nucleotides in pair-wise comparisons were calculated with the PAUP* program [97].

Sequence motifs in the AT-rich region were identified using the Spectral Repeat Finder program [98]. Not only simple motifs were searched but also longer consensus patterns with a minimum match of 75% among different strings.

Testing for phylogenetic signal and saturation of the substitution process

An a priori estimation of the phylogenetic signal present in the multiple alignments was performed by maximum likelihood mapping [70]. The phylogenetic signal was evaluated using the TREEPUZZLE 5.2 program [99].

For the nucleotide sequences, the level of saturation of the substitution process was estimated by plotting uncorrected p-distances (based on observed substitutions) against GTR+I+G estimated distances for multiple alignment of single PCGs as well as concatenated PCGs. In the case of amino acid sequences, the level of saturation was estimated by plotting uncorrected p-distances against mtREV24/JTT + G models implemented in MEGA4 and TREEPUZZLE 5.2. After fitting a regression line, its slope (m) was used as a measure of saturation. If $m = 1$, no saturation is inferred, while for $m \ll 1$, a high level of saturation has occurred.

rrnL and *rrnS* homology modeling

Secondary structures of *rrnL* and *rrnS* were produced through homology modeling using as templates published structures of *Apis mellifera*, *Drosophila melanogaster* and *Drosophila virilis* [74,79].

Additional material

Additional file 1: Table S1: Subdivision of endopterygotan mtDNAs in clusters based on A+T content (T1a), AT-skew (T1b), and GC-skew/T1c9 respectively.

Additional file 2: Table S2: A+T%, AT-skew, G+C%, GC-skew, for whole genome α strand; pooled $\alpha + \beta$ strands PCGs; pooled- α strand PCGs and pooled- β strand PCGs.

Additional file 3: Figure S1: Relative Synonymous Codon Usage (RSCU) in neuropterid pooled α -strand protein-coding genes. Codon families are provided on the x axis. Red-colored codon, codon not present in the pooled genes.

Additional file 4: Figure S2: Relative Synonymous Codon Usage (RSCU) in neuropterid pooled β -strand protein-coding genes. Codon families are provided on the x axis. Red-colored codon, codon not present in the pooled genes.

Additional file 5: Table S3: Summary of multiple alignments of tRNA families in neuropterid mtDNAs.

Additional file 6: Figure S3: Domain I of *rrnL* in neuropterid species and in *Drosophila melanogaster*. Green background, conserved nucleotide in the pair-wise alignment with *L. macaronius*.

Abbreviations

mtDNA: mitochondrial DNA; *atp6* and *atp8*: ATP synthase subunits 6 and 8; *cob*: apocytochrome b; *cox1-3*: cytochrome c oxidase subunits 1-3; *nad1-6* and *nad4L*: NADH dehydrogenase subunits 1-6 and 4L; *rrnS* and *rrnL*: small and large subunit ribosomal RNA (rRNA) genes; *trnX*: transfer RNA (tRNA) genes; where X is the one-letter abbreviation of the corresponding amino acid; s1-s3: mitochondrial genomic spacers; A+T region: the putative control region; PCG: protein-coding gene; RSCU: relative synonymous codon usage; ENC: effective number of codons used; nt: nucleotides; bp: base pairs.

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Authors' contributions

EN and TP designed and coordinated all experiments. EN and MB conducted the molecular experiments. EN performed the genomic analyses and wrote the first draft of the manuscript. All authors contributed the final version of the manuscript and approved it.

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Supporting Table S1a. Clusters of A+T%

ORD	N	TAXON	A+T%	CL	n	ORD	N	TAXON	A+T%	CL	n
COL	22	<i>Tetrapterus bruchi</i>	66,99	A1	5	DIP	38	<i>Cydistomyia duplonotata</i>	77,93	A3	
COL	11	<i>Lucanus mazama</i>	67,11			DIP	44	<i>Drosophila simulans</i>	77,93		
COL	4	<i>Apatides fortis</i>	67,19			COL	12	<i>Macrogyrus oblongus</i>	78		
COL	1	<i>Acmaeodera</i> sp	68,41			COL	3	<i>Anoplophora glabripennis</i>	78,31		
COL	17	<i>Pyrophorus divergens</i>	69,44			MEG	82	<i>Sialis hamata</i>	78,32		
COL	7	<i>Chrysochroa fulgidissima</i>	69,92	DIP	45	<i>Drosophila yakuba</i>	78,59				
COL	24	<i>Tribolium castaneum</i>	71,68	NEU	77	<i>Polystoechotes punctatus</i>	78,96				
COL	13	<i>Mordella atrata</i>	71,94	DIP	25	<i>Aedes aegypti</i>	79				
DIP	31	<i>Bactrocera oleae</i>	72,63	COL	5	<i>Chaetosoma scaritides</i>	79,04				
COL	2	<i>Adelium</i> sp	72,71	DIP	46	<i>Haematobia irritans</i>	79,07				
DIP	32	<i>Bactrocera papayae</i>	73,52	COL	19	<i>Rhagophthalmus ohbai</i>	79,15				
DIP	29	<i>Bactrocera carambolae</i>	73,55	COL	23	<i>Trachypachus holmbergi</i>	79,45				
DIP	30	<i>Bactrocera dorsalis</i>	73,58	DIP	26	<i>Aedes albopictus</i>	79,54				
DIP	33	<i>Bactrocera philippinensis</i>	73,63	COL	18	<i>Rhagophthalmus lufengensis</i>	79,63				
DIP	51	<i>Trichophthalma punctata</i>	73,96	LEP	61	<i>Acraea issoria</i>	79,76				
NEU	76	<i>Libelloides macaronius</i>	74,5	LEP	65	<i>Artogeia melete</i>	79,78				
MEG	80	<i>Corydalus cornutus</i>	74,9	NEU	79	<i>Ditaxis biseriata</i>	79,79				
COL	9	<i>Cyphon</i> sp.	75,17	LEP	71	<i>Lymantria dispar</i>	79,88				
NEU	78	<i>Ascaloptynx appendiculatus</i>	75,57	LEP	69	<i>Diatraea saccharalis</i>	80,02				
COL	20	<i>Rhopaea magnicornis</i>	75,59	HYM	60	<i>Vanhornia eucnemidarum</i>	80,11				
MEG	81	<i>Protohermes concolorus</i>	75,83	LEP	63	<i>Antheraea pernyi</i>	80,16				
HYM	59	<i>Orussus occidentalis</i>	76,21	LEP	64	<i>Antheraea yamamai</i>	80,29				
DIP	40	<i>Drosophila littoralis</i>	76,24	RPH	83	<i>Mongoloraphidia harmandi</i>	80,31				
MCP	84	<i>Neopanorpa pulchra</i>	76,38	LEP	62	<i>Adoxophyes honmai</i>	80,39				
COL	14	<i>Priasilpha obscura</i>	76,5	HYM	52	<i>Abispa ephippium</i>	80,61				
COL	15	<i>Psacotheta hilaris</i>	76,63	LEP	75	<i>Saturnia boisduvalii</i>	80,62				
DIP	35	<i>Chrysomya putoria</i>	76,7	COL	21	<i>Sphaerius</i> sp.	80,68				
COL	6	<i>Chauliognathus opacus</i>	76,86	LEP	70	<i>Eriogyna pyretorum</i>	80,82				
COL	8	<i>Crioceris duodecimpunctata</i>	76,89	DIP	50	<i>Simosyrphus grandicornis</i>	80,84				
DIP	36	<i>Cochliomyia hominivorax</i>	76,9	LEP	74	<i>Phthonandria atrilineata</i>	81,02				
COL	10	<i>Hydroscapha granulatum</i>	77,29	LEP	67	<i>Bombyx mori</i>	81,32				
DIP	28	<i>Anopheles quadrimaculatus A</i>	77,36	LEP	66	<i>Bombyx mandarina</i>	81,68				
DIP	37	<i>Culicoides arakawae</i>	77,36	LEP	72	<i>Manduca sexta</i>	81,79				
COL	16	<i>Pyrocoelia rufa</i>	77,41	HYM	55	<i>Cephus cinctus</i>	81,95				
DIP	34	<i>Ceratitis capitata</i>	77,48	DIP	42	<i>Drosophila melanogaster</i>	82,16				
DIP	27	<i>Anopheles gambiae</i>	77,56	LEP	68	<i>Coreana raphaelis</i>	82,66				
DIP	43	<i>Drosophila sechellia</i>	77,57	DIP	48	<i>Mayetiola destructor</i>	84,12				
DIP	47	<i>Lucilia sericata</i>	77,61	HYM	53	<i>Apis mellifera ligustica</i>	84,86				
DIP	41	<i>Drosophila mauritiana</i>	77,71	DIP	49	<i>Rhopalomyia pomum</i>	85,15				
HYM	57	<i>Evania appendigaster</i>	77,77	HYM	58	<i>Melipona bicolor</i>	86,72				
DIP	39	<i>Dermatobia hominis</i>	77,82	HYM	54	<i>Bombus ignitus</i>	86,78				
LEP	73	<i>Ochrogaster lunifer</i>	77,84	HYM	56	<i>Diadegma semiclausum</i>	87,41				

ORD, order; N; reference number in Figure 3; A+T%, A+T content expressed in percent; CL cluster name; n, number of taxa in the cluster. COL, Coleoptera; DIP, Diptera; HYM, Hymenoptera; MCP, Mecoptera; MEG, Megaloptera; NEU, Neuroptera; RPH, Raphidioptera. A1 (65.00 < AT% < 70.00); A2 (70.00 < AT% < 75.00); A3 (75.00 < AT% < 80.00); A4 (80.00 < AT% < 85.00); A5 (85.00 < AT% < 90.00)

Supporting Table S1b. Clusters of AT-skew

ORD	N	TAXON	AT-skew	CL	n	ORD	N	TAXON	AT-skew	CL	n
LEP	68	<i>Coreana raphaelis</i>	-0,047476104	B1	13	DIP	34	<i>Ceratitis capitata</i>	0,021159748	B2	44
LEP	70	<i>Eriogyna pyretorum</i>	-0,030596593			LEP	69	<i>Diatraea saccharalis</i>	0,021379589		
NEU	79	<i>Polystoechotes punctatus</i>	-0,028747433			RPH	83	<i>Mongoloraphidia harmandi</i>	0,023027851		
LEP	75	<i>Saturnia boisduvalii</i>	-0,024063307			HYM	57	<i>Evania appendigaster</i>	0,026703233		
LEP	61	<i>Acraea issoria</i>	-0,023439428			COL	10	<i>Hydroscapha granulum</i>	0,02972382		
LEP	64	<i>Antheraea yamamai</i>	-0,022168088			LEP	73	<i>Ochrogaster lunifer</i>	0,030153238		
LEP	63	<i>Antheraea pernyi</i>	-0,021477801			DIP	27	<i>Anopheles gambiae</i>	0,032225579		
HYM	52	<i>Abispa ephippium</i>	-0,018660812			HYM	55	<i>Cephus cinctus</i>	0,034387028		
MCP	84	<i>Neopanorpa pulchra</i>	-0,013994267			DIP	36	<i>Cochliomyia hominivorax</i>	0,034493953		
MEG	81	<i>Protohermes concolorus</i>	-0,011148087			DIP	28	<i>Anopheles quadrimaculatus A</i>	0,040649047		
LEP	72	<i>Manduca sexta</i>	-0,00535855			COL	12	<i>Macrogyrus oblongus</i>	0,04213851		
DIP	50	<i>Simosyrphus grandicornis</i>	-0,003832005			COL	5	<i>Chaetosoma scaritides</i>	0,04274062		
LEP	62	<i>Adoxophyes honmai</i>	-0,001031337			DIP	39	<i>Dermatobia hominis</i>	0,042965989		
HYM	54	<i>Bombus ignitus</i>	0,002874974	COL	8	<i>Crioceris duodecimpunctata</i>	0,044226044				
DIP	38	<i>Cydistomyia duplonotata</i>	0,003159059	DIP	49	<i>Rhopalomyia pomum</i>	0,046801619				
DIP	46	<i>Haematobia irritans</i>	0,004483599	COL	14	<i>Priasilpha obscura</i>	0,052358082				
DIP	45	<i>Drosophila yakuba</i>	0,005004369	LEP	66	<i>Bombyx mandarina</i>	0,054727133				
COL	20	<i>Rhopaea magnicornis</i>	0,005058513	LEP	67	<i>Bombyx mori</i>	0,058721799				
LEP	74	<i>Phthonandria atrilineata</i>	0,006688963	DIP	48	<i>Mayetiola destructor</i>	0,061457914				
DIP	26	<i>Aedes albopictus</i>	0,007921539	DIP	33	<i>Bactrocera philippinensis</i>	0,065619933				
DIP	44	<i>Drosophila simulans</i>	0,008485472	DIP	29	<i>Bactrocera carambolae</i>	0,065698419				
HYM	56	<i>Diadegma semiclausum</i>	0,008674404	DIP	32	<i>Bactrocera papayae</i>	0,066233655				
DIP	43	<i>Drosophila sechellia</i>	0,009226524	DIP	30	<i>Bactrocera dorsalis</i>	0,067543335				
DIP	37	<i>Culicoides arakawae</i>	0,009265859	NEU	76	<i>Ascaloptynx appendiculatus</i>	0,067677946				
DIP	41	<i>Drosophila mauritiana</i>	0,009459924	COL	13	<i>Mordella atrata</i>	0,069147509				
COL	21	<i>Sphaerius</i> sp.	0,010491803	NEU	78	<i>Libelloides macaronius</i>	0,071971617				
COL	15	<i>Psacothia hilaris</i>	0,011439388	COL	9	<i>Cyphon</i> sp.	0,072114983				
COL	3	<i>Anoplophora glabripennis</i>	0,011658031	COL	11	<i>Lucanus mazama</i>	0,074399531				
LEP	65	<i>Artogeia melete</i>	0,012169882	HYM	60	<i>Vanhornia eucnemidarum</i>	0,085787452				
DIP	40	<i>Drosophila littoralis</i>	0,012446774	DIP	31	<i>Bactrocera oleae</i>	0,088455511				
MEG	80	<i>Corydalus cornutus</i>	0,014043748	DIP	51	<i>Trichophthalma punctata</i>	0,091283912				
MEG	82	<i>Sialis hamata</i>	0,014561518	COL	18	<i>Rhagophthalmus lufengensis</i>	0,103881817				
NEU	77	<i>Ditaxis biseriata</i>	0,015497366	COL	16	<i>Pyrocoelia rufa</i>	0,105884066				
HYM	58	<i>Melipona bicolor</i>	0,015591269	COL	24	<i>Tribolium castaneum</i>	0,109100492				
DIP	47	<i>Lucilia sericata</i>	0,015919192	COL	6	<i>Chauliognathus opacus</i>	0,112081768				
LEP	71	<i>Lymantria dispar</i>	0,016082342	COL	1	<i>Acmaeodera</i> sp.	0,114296016				
DIP	42	<i>Drosophila melanogaster</i>	0,016775803	COL	19	<i>Rhagophthalmus ohbai</i>	0,118513155				
DIP	25	<i>Aedes aegypti</i>	0,016949153	COL	2	<i>Adelium</i> sp.	0,136789298				
HYM	59	<i>Orussus occidentalis</i>	0,016949153	COL	17	<i>Pyrophorus divergens</i>	0,16508844				
HYM	53	<i>Apis mellifera ligustica</i>	0,018315547	COL	4	<i>Apatides fortis</i>	0,170439904				
DIP	35	<i>Chrysomya putoria</i>	0,020498889	COL	7	<i>Chrysochroa fulgidissima</i>	0,202640643				
COL	23	<i>Trachypachus holmbergi</i>	0,020574814	COL	22	<i>Tetraphalerus bruchi</i>	0,247764034				

ORD, order; N; reference number in Figure 3; CL cluster name; n, number of taxa in the cluster. COL, Coleoptera; DIP, Diptera; HYM, Hymenoptera; MCP, Mecoptera; MEG, Megaloptera; NEU, Neuroptera; RPH, Raphidioptera. B1 (-0.050 < AT-skew ≤ 0.000); B2 (0.000 < AT-skew ≤ 0.050); B3 (0.050 < AT-skew ≤ 0.100); B4 (0.100 < AT-skew ≤ 0.150); B5 (0.150 < AT-skew ≤ 0.200); B6 (0.200 < AT-skew ≤ 0.250).

Supporting table S1c. Clusters of GC-skew

ORD	N	TAXON	GC-skew	CL	n	ORD	N	TAXON	GC-skew	CL	n
HYM	52	<i>Abispa ephippium</i>	-0,379562044	C1	1	COL	15	<i>Psacotha hilaris</i>	-0,212955466	C4	30
HYM	57	<i>Evania appendigaster</i>	-0,349154254	C2	5	DIP	25	<i>Aedes aegypti</i>	-0,210977702		
HYM	60	<i>Vanhornia eucnemidarum</i>	-0,328267477			COL	12	<i>Macrogyrus oblongus</i>	-0,208083015		
LEP	73	<i>Ochrogaster lunifer</i>	-0,317510854			DIP	36	<i>Cochliomyia hominivorax</i>	-0,206700892		
HYM	59	<i>Orussus occidentalis</i>	-0,307672027			COL	3	<i>Anoplophora glabripennis</i>	-0,206674473		
COL	24	<i>Tribolium castaneum</i>	-0,305314654			NEU	78	<i>Ascaloptynx appendiculatus</i>	-0,205980923		
COL	4	<i>Apatides fortis</i>	-0,29462771	C3	14	LEP	70	<i>Eriogyna pyretorum</i>	-0,204761905	C5	24
COL	17	<i>Pyrophorus divergens</i>	-0,291514413			COL	5	<i>Chaetosoma scaritides</i>	-0,200861273		
HYM	55	<i>Cephus cinctus</i>	-0,285550459			HYM	56	<i>Diadegma semiclausum</i>	-0,198473282		
DIP	31	<i>Bactrocera oleae</i>	-0,28020328			LEP	62	<i>Adoxophyes honmai</i>	-0,196486662		
COL	11	<i>Lucanus mazama</i>	-0,272328549			COL	18	<i>Rhagophthalmus lufengensis</i>	-0,195945946		
HYM	54	<i>Bombus ignitus</i>	-0,271974229			LEP	74	<i>Phthonandria atrilineata</i>	-0,192111527		
HYM	53	<i>Apis mellifera ligustica</i>	-0,268686869			DIP	34	<i>Ceratitis capitata</i>	-0,185102835		
COL	14	<i>Priasilpha obscura</i>	-0,264615385			DIP	40	<i>Drosophila littoralis</i>	-0,182654402		
MEG	80	<i>Corydalus cornutus</i>	-0,26206196			DIP	28	<i>Anopheles quadrimaculatus A</i>	-0,181480423		
LEP	69	<i>Diatraea saccharalis</i>	-0,257512116			DIP	26	<i>Aedes albopictus</i>	-0,181231672		
COL	13	<i>Mordella atrata</i>	-0,255675304			LEP	72	<i>Manduca sexta</i>	-0,181174805		
MEG	81	<i>Protohermes concolorus</i>	-0,253980684			NEU	79	<i>Ditaxis biseriata</i>	-0,179378957		
COL	22	<i>Tetraphalerus bruchi</i>	-0,251592972			DIP	38	<i>Cydistomyia duplonotata</i>	-0,177126918		
HYM	58	<i>Melipona bicolor</i>	-0,251174935			NEU	76	<i>Libelloides macaronius</i>	-0,176702863		
COL	1	<i>Acmaeodera</i> sp	-0,249268007	C4		MEG	82	<i>Sialis hamata</i>	-0,171149867	C6	10
LEP	71	<i>Lymantria dispar</i>	-0,247366741			DIP	35	<i>Chrysomya putoria</i>	-0,170189702		
DIP	51	<i>Trichophthalma punctata</i>	-0,244788007			MCP	84	<i>Neopanorpa pulchra</i>	-0,167620605		
COL	20	<i>Rhopaea magnicornis</i>	-0,238072965			DIP	47	<i>Lucilia sericata</i>	-0,166666667		
COL	7	<i>Chrysochroa fulgidissima</i>	-0,237364044			COL	8	<i>Crioceris duodecimpunctata</i>	-0,163487738		
DIP	37	<i>Culicoides arakawae</i>	-0,236957582			NEU	77	<i>Polystoechotes punctatus</i>	-0,161232958		
LEP	61	<i>Acraea issoria</i>	-0,235255995			COL	16	<i>Pyrocoelia rufa</i>	-0,158472673		
DIP	30	<i>Bactrocera dorsalis</i>	-0,228353949			LEP	68	<i>Coreana raphaelis</i>	-0,157815443		
COL	9	<i>Cyphon</i> sp.	-0,228238866			DIP	27	<i>Anopheles gambiae</i>	-0,154046997		
RPH	83	<i>Mongoloraphidia harmandi</i>	-0,227791878			COL	23	<i>Trachypachus holmbergi</i>	-0,153655514		
DIP	39	<i>Dermatobia hominis</i>	-0,226784238			COL	6	<i>Chauliognathus opacus</i>	-0,153221126		
DIP	32	<i>Bactrocera papayae</i>	-0,22638823			COL	10	<i>Hydroscapha granulum</i>	-0,151916184		
COL	19	<i>Rhagophthalmus ohbai</i>	-0,225648855			DIP	42	<i>Drosophila melanogaster</i>	-0,150488225		
DIP	33	<i>Bactrocera philippinensis</i>	-0,224499523			COL	21	<i>Sphaerius</i> sp.	-0,148249828		
DIP	29	<i>Bactrocera carambolae</i>	-0,223752969			DIP	45	<i>Drosophila yakuba</i>	-0,136443149		
COL	2	<i>Adelium</i> sp	-0,222222222			DIP	44	<i>Drosophila simulans</i>	-0,135249622		
LEP	65	<i>Artogeia melete</i>	-0,221822934			DIP	43	<i>Drosophila sechellia</i>	-0,135102893		
LEP	64	<i>Antheraea yamamai</i>	-0,219980152			DIP	41	<i>Drosophila mauritiana</i>	-0,134892086		
LEP	75	<i>Saturnia boisduvalii</i>	-0,217069892			DIP	50	<i>Simosyrphus grandicornis</i>	-0,132880698		
LEP	63	<i>Antheraea pernyi</i>	-0,216321244			DIP	46	<i>Haematobia irritans</i>	-0,124517088		
LEP	67	<i>Bombyx mori</i>	-0,216290212	DIP	49	<i>Rhopalomyia pomum</i>	-0,123084069				
LEP	66	<i>Bombyx mandarina</i>	-0,213159698	DIP	48	<i>Mayetiola destructor</i>	-0,110068259				

ORD, order; N; reference number in Figure 3; CL, cluster name; n, number of taxa in the cluster. COL, Coleoptera; DIP, Diptera; HYM, Hymenoptera; MCP, Mecoptera; MEG, Megaloptera; NEU, Neuroptera; RPH, Raphidioptera. C1 (-0.400 < GC-skew < -0.350); C2 (-0.300 < GC-skew < -0.250); C3 (-0.300 < GC-skew < -0.250); C4 (-0.250 < GC-skew < -0.200); C5 (-0.200 < GC-skew < -0.150); C6 (-0.150 ≤ GC-skew < -0.100).

Supporting Table S2. A+T%, AT-skew, G+C%, GC-skew, for whole genome α strand; pooled $\alpha + \beta$ strands PCGs; pooled- α strand PCGs and pooled- β strand PCGs

Taxon	whole genome α strand				pooled $\alpha + \beta$ strands PCGs					
	A+T%	AT-skew	G+C%	GC-skew	CDS	A+T%	AT-skew	G+C%	GC-skew	NLC
<i>Ascaloptynx appendiculatus</i>	75.57	0.06768	24.43	-0.205981	3699	73.91	-0.14289	26.09	-0.021762	1
<i>Ditaxis biseriata</i>	79.79	0.0155	20.21	-0.179379	3705	77.18	-0.13859	22.82	0.015773	3
<i>Libelloides macaronius</i>	74.5	0.07197	25.5	-0.176703	3703	73	-0.11739	27	-0.024341	2
<i>Polystoechotes punctatus</i>	78.96	-0.02875	21.04	-0.161233	3706	77.21	-0.1794	22.79	0.035517	5
<i>Corydalus cornutus</i>	74.9	0.01404	25.1	-0.262062	3702	72.64	-0.17268	27.36	0.001645	1
<i>Protohermes concolorus</i>	75.83	-0.01115	24.17	-0.253981	3706	74.02	-0.16428	25.98	0.004848	0
<i>Sialis hamata</i>	78.32	0.01456	21.67	-0.17115	3708	76.12	-0.15116	23.87	0.022222	1
<i>Mongoloraphidia harmandi</i>	80.31	0.02303	19.69	-0.227792	3675	78.1	-0.14893	21.9	-0.000829	2
average	77.27	0.021	22.72	-0.205	3700.50	75.27	-0.152	24.73	0.004	1.87
standard deviation	2.17	0.035	2.32	0.039	10.68	2.12	0.020	2.13	0.020	1.55

Taxon	pooled- α strand PCGs						pooled- β strand PCGs					
	CDS	A+T%	AT-skew	G+C%	GC-skew	NLC	CDS	A+T%	AT-skew	G+C%	GC-skew	NLC
<i>Ascaloptynx appendiculatus</i>	2270	72.29	-0.07008	27.71	-0.176471	5	1429	76.49	-0.25221	23.51	0.267857	8
<i>Ditaxis biseriata</i>	2275	75.49	-0.10171	24.51	-0.105798	6	1430	79.88	-0.19405	20.12	0.251448	12
<i>Libelloides macaronius</i>	2272	71.13	-0.033	28.87	-0.156504	4	1431	75.98	-0.2428	24.02	0.227934	5
<i>Polystoechotes punctatus</i>	2274	75.59	-0.17704	24.41	-0.077477	9	1432	79.77	-0.18296	20.23	0.252014	18
<i>Corydalus cornutus</i>	2271	70.54	-0.13525	29.46	-0.191829	3	1431	75.96	-0.22784	24.04	0.377907	9
<i>Protohermes concolorus</i>	2274	72.1	-0.14698	27.9	-0.17814	3	1432	77.07	-0.18997	22.93	0.358376	12
<i>Sialis hamata</i>	2279	74.64	-0.11581	25.35	-0.109059	3	1429	78.49	-0.20475	21.51	0.26898	8
<i>Mongoloraphidia harmandi</i>	2261	76.05	-0.11365	23.95	-0.168719	6	1414	81.38	-0.20162	18.62	0.344304	12
average	2272	73.48	-0.112	26.52	-0.145	4.87	1428.5	78.13	-0.212	21.87	0.294	10.5
standard deviation	5.24	2.20	0.045	2.20	0.042	2.10	5.98	2.05	0.026	2.05	0.057	3.93

PCG, protein-coding gene; CDS, number of codons; NLC, number of codon(s) lost in the strand.

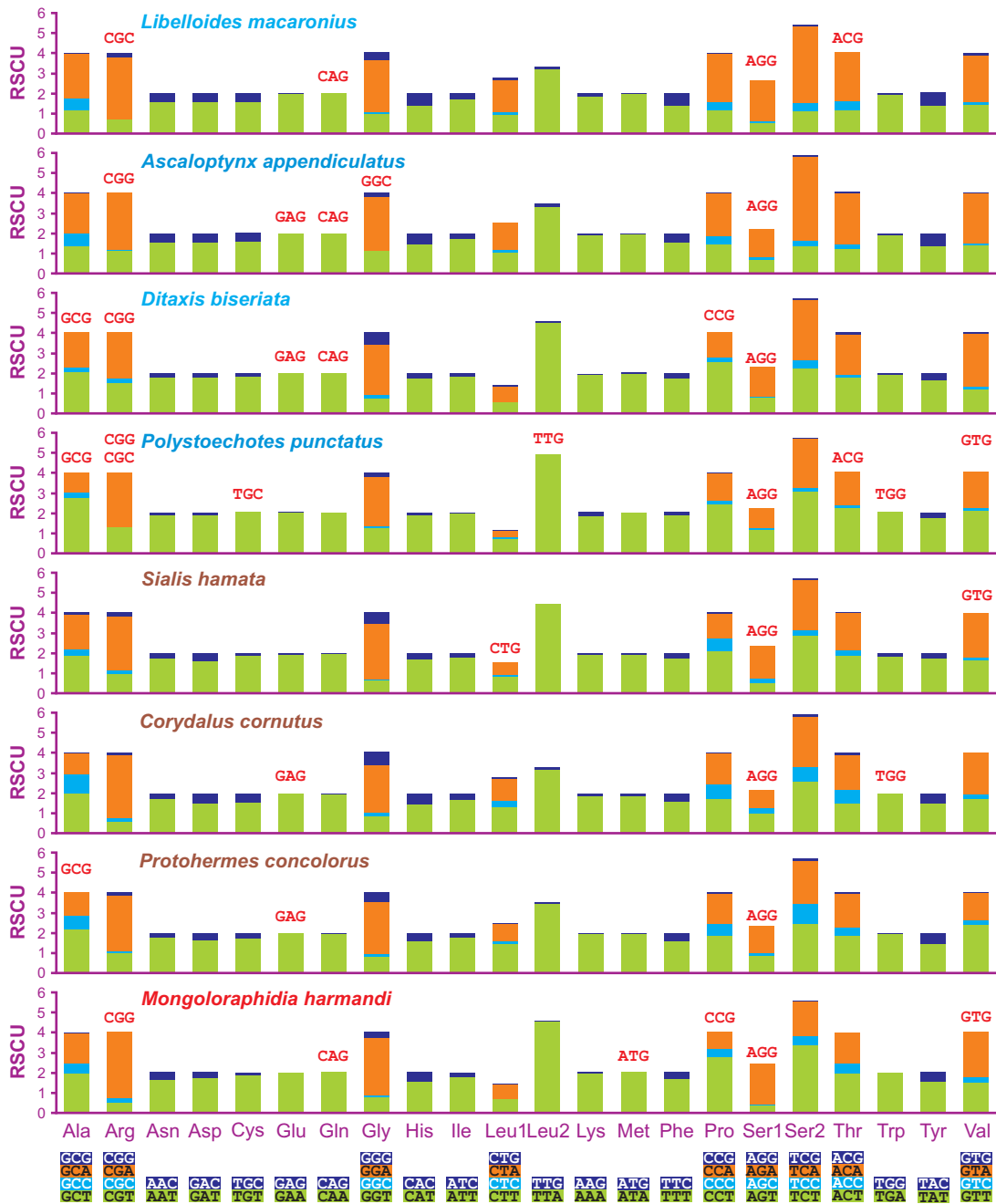


Figure S1. - Relative Synonymous Codon Usage (RSCU) in neuropterid pooled α -strand protein-coding genes .

Codon families are provided on the x axis. Red-colored codon, codon not present in the pooled genes.

Note that the behavior of the families CF_{Ala} , CF_{Pro} and CF_{Val} differs from that observed for the pooled- α + β PCGs . In the first two families the adenine is the most common nucleotide in third positions of ascalaphid *A. appendiculatus* and *L. macaronius* while the thymine is the most abundant in all other neuropterids. In CF_{Val} adenine is the most abundant in third positions with exception represented by *P. punctatus* and *P. concolorus* where it is superseded by thymine.

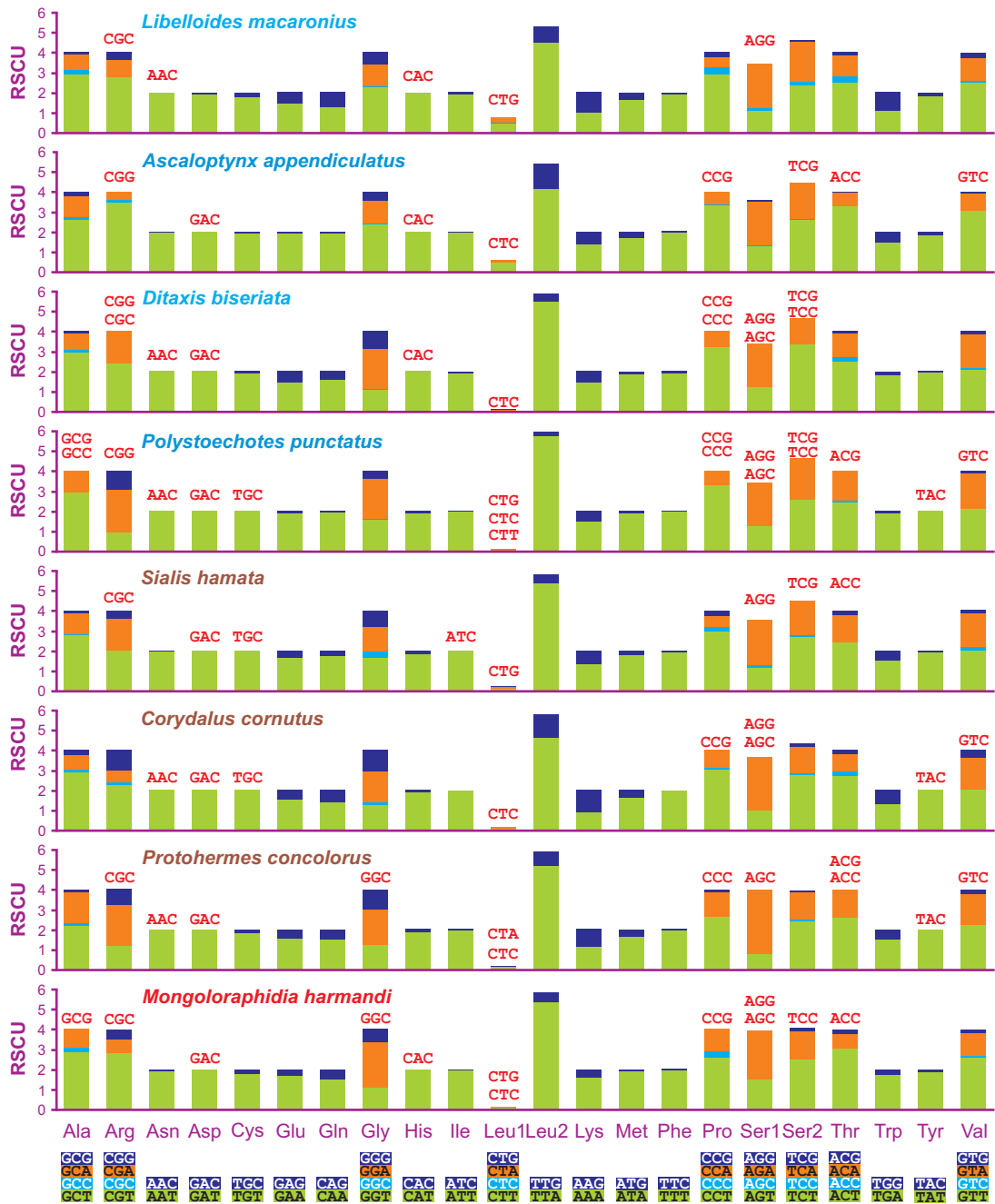


Figure S3. - Relative Synonymous Codon Usage (RSCU) in neuropterid pooled β -strand protein-coding genes .

Codon families are provided on the x axis. Red-colored codon, codon not present in the pooled genes.

Loss of synonymous GC-rich codons is common in fourfold codon families (CFs) (e.g. CF_{Ser1}, CF_{Ser2}). In CF_{Arg} abundance of A vs. T in third positions is very variable ranging from strong bias toward A (e.g. *A. appendiculatus*) up to strong bias toward T (*P. punctatus*). The twofold degenerate CF_{Lys} shows also a variable pattern with G preferred to A in *C. cornutus*, G equaling A (*L. macaronius*) or A favored over G (other species).

Supporting Table S3. Summary of multiple alignments of tRNA genes in neuropterid mtDNAs.

ALN	amino acid	alignment length	identical positions	%INUC
<i>trnA</i>	Alanine	68	39	57.35
<i>trnC</i>	Cysteine	70	27	38.57
<i>trnD</i>	Aspartate	70	40	51.43
<i>trnE</i>	Glutamate	70	49	70.00
<i>trnF</i>	Phenylalanine	68	31	45.59
<i>trnG</i>	Glycine	66	51	77.27
<i>trnH</i>	Histidine	70	40	57.14
<i>trnI</i>	Isoleucine	68	40	58.82
<i>trnK</i>	Lysine	71	54	76.06
<i>trnL1</i>	Leucine (CUN)	68	39	57.35
<i>trnL2</i>	Leucine (UUR)	66	45	68.18
<i>trnM</i>	Methionine	70	50	71.43
<i>trnN</i>	Asparagine	68	49	72.02
<i>trnP</i>	Proline	69	42	60.87
<i>trnQ</i>	Glutamine	69	45	65.28
<i>trnR</i>	Arginine	69	35	50.72
<i>trnS1</i>	Serine (AGN)	68	41	60.29
<i>trnS2</i>	Serine (UCN)	67	43	64.18
<i>trnT</i>	Threonine	65	41	63.08
<i>trnV</i>	Valine	73	38	52.05
<i>trnW</i>	Tryptophan	68	47	69.18
<i>trnY</i>	Tyrosine	72	47	65.28

ALN, alignment name; %INUC, percent of identical nucleotides

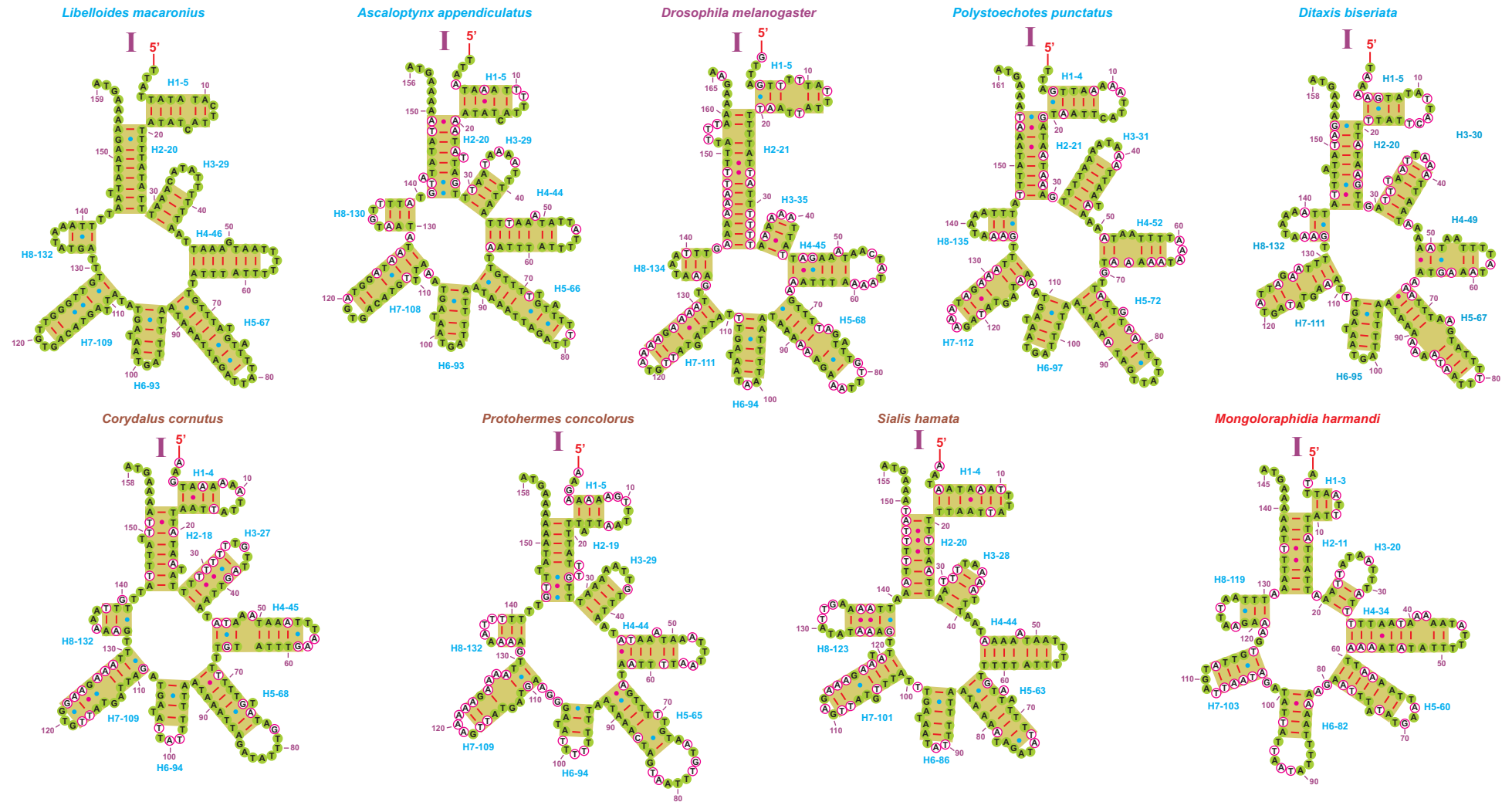


Figure S3. - Domain I of *rrnL* in neuropterid species and in *Drosophila melanogaster*.

Green background, conserved nucleotide in the pair-wise alignment with *L. macaronius*.