Transcriptome of the synganglion and characterization of the cys-loop ligand-gated ion channel gene family in the tick *Ixodes ricinus*.

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Abstract

Ticks represent a major health issue for humans and domesticated animals. Assessing the expression patterns of the tick's central nervous system, known as the synganglion, is an important step in understanding tick physiology and in managing tick-borne diseases. Neuron-specific genes like the cys-loop ligand-gated ion channels (cys-loop LGICs) are important pharmacological targets of acaricides. Here, we carried out the sequencing of transcriptomes of the *I. ricinus* synganglion, for adult ticks in different conditions (unfed males, unfed females, and partially-fed females). The *de novo* assembly of these transcriptomes allowed us to obtain a large collection of cys-loop LGICs sequences. A reference meta-transcriptome based on synganglion and whole body transcriptomes was then produced, showing high completeness and allowing differential expression analyses between synganglion and whole body. Many of the genes upregulated in the synganglion were related to biological processes or functions associated with neurotransmission and located in neurones or the synaptic membrane, including most of the cys-loop LGICs. As a first step of a functional study of cysLGICs, we cloned the predicted sequence of the resistance to dieldrin (RDL) subunit homologue, and functionally reconstituted the first GABA-gated receptor of *Ixodes ricinus* using a heterologous expression approach. A phylogenetic study was performed for the nicotinic acetylcholine receptors (nAChRs) and for other cys-loop LGICs respectively, showing tick-specific expansions of some types of receptors (Histamine-gated, GluClIs).

Introduction

Like the vertebrate brain, the central nervous system (CNS) of arthropods performs essential functions, such as the perception of stimuli, the control of movement and the regulation of many physiological processes. Whether one compares the brain of vertebrates and the CNS of arthropods, or compares the CNS of different arthropod groups, important similarities but also differences are evident, and can be related to the specific biology and evolution of each group¹. In ticks, a group of haematophagous arthropods belonging to the Chelicerata, the CNS is a specialized organ called a synganglion², organized as a single mass of cells. As ticks are potential transmitters of many pathogens and parasites, a better knowledge of the function of this essential organ could help in understanding interactions between ticks and tick-borne-pathogens³, but also provide targets for chemical acaricides⁴. Indeed, to-date, most acaricides have targeted neural functions, but as with

many insect pests, the continued use of these molecules has generated resistance, prompting the need to find new methods or targets⁵.

Knowledge of the neural biology of ticks is scarced compared to other invertebrates², particularly with respect to gene expression changes in the synganglion during blood feeding or after mating⁶. However progress has been made, notably through the sequencing of synganglion transcriptomes of several tick species. For example, a study of expressed-sequence tags of the *Rhipicephalus sanguineus* synganglion has identified different neural receptors⁷. Two companion studies of this study reported the sequencing of synganglion transcriptomes of the American dog tick, *Dermacentor variabilis*^{8,9}. A synganglion transcriptome was also sequenced and annotated for *Ixodes scapularis*⁶. The above studies have identified several genes associated with the neural system and nerve impulse transmission, for example neuropeptides and proteins associated with neurotransmission. The synganglion transcriptome of the cattle tick, *Rhipicephalus microplus*, was investigated with a focus on predicting genes in the G protein-coupled receptor (GPCR) family¹⁰, representing potential targets for new acaricides¹¹. But in each of the above studies, relatively few genes associated with neurotransmission were reported, probably due to relatively low sequencing depth. Further, in these data sets, it is impossible to assess the extent to which these gene collections were synganglion specific, as expression levels were only measured for synganglion samples, and were not compared to other tissues or the whole body. Recently, the *Ixodes scapularis* ISE6 cell line was shown to have neuronal characteristics, which provided a base for investigating neuronal interactions between ticks and pathogens³.

Ligand-gated cys-loop ion channels (cys-loop LGICs) are key receptors mediating neurotransmission both in vertebrates and invertebrates. Ligand-gated ion channels are major pharmacological targets of pesticides or acaricides such as macrocyclic lactones, phenylpyrazoles and neonicotinoids⁵. Yet, their knowledge for ticks is partial in several aspects, including sequence reconstruction and annotation, phylogeny, or function, even though complete genomes of ticks have been sequenced for *Ixodes scapularis*⁴ (32 sequences of cysLGICs were found in this study) or five other tick species¹² (no specific annotation of cysLGICs reported in this study). The cys-loop LGICs form a superfamily of subunits that share a similar membrane topology. Indeed, they comprise one large amino terminal ligand-binding domain and four transmembrane domains, the second of these domains determining the ionic pore. The ionic pore consists of a multimeric assembly of five subunits, which can be identical or different. Subgroups within LGICs are determined by the activating neurotransmitter molecules - acetylcholine, γ -Aminobutyric acid (GABA), glycine, glutamate, or histamine - able to bind and trigger the opening of the channel gate. In Chelicerata, a first extensive reconstruction and in-depth study of the cys-loop LGICs was carried out for the two-spotted spider mite *Tetranychus urticae* (Acari: Acariformes), a phytophagous parasite, on the basis of its genome sequence¹³. On the basis of its transcriptomes, a similar reconstruction and phylogenetic study was carried out for the spider *Cupiennius salei*¹⁴. Similar analysis is lacking for ticks, despite the important implications of this knowledge for a better understanding of tick neurobiology and for providing potential targets for their control. The tick *Ixodes ricinus* is a widespread species in Europe, where it represents an increasing human health concern, being the main potential transmitter of the Lyme disease agent and of other microbes or parasites¹⁵. No transcriptome of the synganglion has been sequenced and no annotation of the cys-loop LGICs has been yet performed for this species, to our knowledge.

To study the interactions between ion channels and pesticides, i.e. between the nicotinic receptors of acetylcholine (nAChRs) and neonicotinoids, and between GABA receptors and other molecules, and to evaluate possible adverse effects on non-target species, systems of *ex-vivo* expression constitute an approach of choice. However the heterologous expression of some ion channels such as cholinergic receptors remains challenging. This has been succesful in some arthropods using the alpha subunits of the target species combined with vertebrate beta subunits (making hybrid receptors), whereas a first non-hybrid arthropod receptor using alpha and beta units from the target

species, was obtained for the salmon louse *Lepeophtheirus salmonis*¹⁶. For arachnids, the first functional characterization of an alpha-type receptor was done for the brown dog tick *Rhipicephalus sanguineus*, but in vitro expression patterns were unreliable¹⁷. In the tick *I. ricinus*, the microtransplantation of synganglion membranes in oocytes of *Xenopus levis* allowed to express nicotinic acetylcholine receptors and to evaluate their sensitivity to insecticides¹⁸, but insofar no cloning of alpha-type receptor has been reported for ticks. A GABA receptor was first identified in *Drosophila melanogaster* (the *Rdl* gene¹⁹, conferring resistance to the insecticide dieldrin), whereas *Rdl* homologs were cloned in the ticks *Dermacentor variabilis*²⁰ and *Rhipicephalus microplus*²¹ respectively.

In the present study, our principal aims were i) to characterize genes that are specifically expressed (or up-regulated) in the synganglion of the tick *Ixodes ricinus* ii) to undertake a test of ion channel functionality, using the *Rdl* gene as a model iii) to carefully reconstruct the sequences of the *Ixodes ricinus* cys-loop LGICs, and to study their phylogeny including homologs in Arachnida and other arthropods.

Material & methods

Tick sampling

Ticks (*Ixodes ricinus*) were collected under two conditions, unfed or partially fed. Unfed adult female and male ticks were collected by the flagging method on vegetation in Carquefou (Loire-Atlantique, France). Partially fed adult females (half-fed) were collected in Chizé (Deux-Sèvres, France), from roe deers captured in the framework of a long-term monitoring programme of these mammals in the Chizé Forest, a biosphere Reserve.

Ethics statement

Roe deers were captured and released alive, following a protocol that complies with Directive 2010/63/EU of the European Parliament and of the council of 22 September 2010 on the protection of animals used for scientific purposes. This protocol was approved by the Ethics Committee in Animal Experimentation of the University Claude Bernard Lyon 1 (CEEA-55; DR2014-09). The capture of roe deer was carried out with minimal stress and pain to animals, which were released alive at the same site and on the same day of capture.

Sample preparation

Ticks were dissected under a stereomicroscope within 2-3 days of collection. Whole ticks were washed in RNase-ExitusPlus™ plus water. Synganglia were excised, rinsed in PBS buffer 1X, and dried by placing them gently on a clean sheet of paper. Synganglia were then placed in RNase-free microfuge tubes containing RNA extraction buffer (RA1 buffer, Nucleospin RNA XS kit, Macherey Nagel) and stored at -80°C until use. For RNA extraction, the tubes were gently thawed and tissues were mechanically disrupted with decontaminated pestles (3%H₂O₂) and eluted in 200 microliters of RA1. Samples were prepared for three conditions, unfed females, males, and partially fed females, each with three replicates (Table 1). Samples were made with pools of synganglia (n=25 for unfed females or males, n=10 for partially fed females). RNA was extracted using the Nucleospin RNA XS kit (Macherey Nagel, Düren, Germany), adapted for small samples, with a 5U DNase treatment.

RNA dosage and quality

RNA purity was assessed with a NanoDrop (Thermo Fisher Scientific Inc., Waltham, MA, USA). The commonly used threshold to consider samples as sufficiently pure is A280/A260 >1.80. Three out of nine samples were just below that threshold (F03, M01 and M02) but were kept for further analyses. Quantity was assessed with a Qubit (Invitrogen, Thermo Fisher Scientific Inc., Waltham, MA, USA): quantities were low in the three samples mentioned above (they were not even dosable), but ranged between 68 and 304 ng for other samples. Finally, a further assessment of quantity and quality was conducted with the 2100BioAnalyzer instrument (Agilent Technologies Inc., Santa Clara, CA, USA): quality was determined by peak profiles, which was good for all samples, with a single, clear peak corresponding to rRNA, but low quantities for three samples, in particular M01 and M02. We decided to pool the latter two samples to make a single library, named M01_02.

Library preparation and sequencing

RNA-Seq library preparations were carried out from 60ng total RNA, or the total volume when not quantifiable (for F03, M01_02). We used the NEBNext Ultra II directional RNA library prep kit (NEB, Ipswich, MA, USA), which allows mRNA strand orientation (sequence reads occur in the same orientation as anti-sense RNA). Briefly, poly(A)+ RNA was selected with oligo(dT) beads, chemically fragmented and converted into single-stranded cDNA using random hexamer priming. Then, the second strand was generated to create double-stranded cDNA. cDNA were then 3'-adenylated, and Illumina adapters were added. Ligation products were PCR-amplified. Ready-to-sequence Illumina libraries were then quantified by qPCR using the KAPA Library Quantification Kit for Illumina Libraries (KapaBiosystems, Wilmington, MA, USA), and libraries profiles evaluated with an Agilent 2100 Bioanalyzer (Agilent Technologies, Santa Clara, CA, USA). Each library was sequenced using 151 bp paired end reads chemistry on a Illumina HiSeq 4000 sequencer, producing a minimum of 40 million paired-reads for each library.

Read cleaning and assembly.

Short Illumina reads were bioinformatically post-processed as in ²² to filter out low quality data. First, low-quality nucleotides (Q < 20) were discarded from both read ends. Then remaining Illumina sequencing adapters and primer sequences were removed and only reads ≥ 30 nucleotides were retained. These filtering steps were done using in-house-designed software based on the FastX package (<https://www.genoscope.cns.fr/fastxtend>). Finally, read pairs mapping to the phage phiX genome were identified and discarded using SOAP aligner ²³ (with default parameters) and the Enterobacteria phage PhiX174 reference sequence (GenBank: NC_001422.1). In addition, ribosomal RNA-like reads were detected using SortMeRNA ²⁴ and filtered out. This set of cleaned reads were deposited to EBI under the BioProject accession PRJEB40724. The quality of cleaned reads was evaluated with fastQC, using MultiQC for visualization²⁵, all parameters being satisfactory. Potential duplicates were removed with the tool fastuniq (<https://sourceforge.net/p/fastuniq/>). The assembly was done with Trinity (version 2.3.2)²⁶ with the options normalize-reads and RF (to account for strandedness). Calculations were done on the Genotoul Bioinformatics platform, using 100GB of memory. We made two separate assemblies, for respectively unfed ticks (females and males) and feeding ticks (females only). This choice was motivated by the expectation of different profiles of transcription in both conditions²⁷ and of a reduced risk of producing chimeric contigs compared to a single assembly of all the reads.

Construction of a high quality reference transcriptome

The two assemblies produced for the synganglion transcriptomes were combined with two previously published transcriptomes of the tick whole body - assembly accessions GFVZ²⁸ and GIDG²⁹ in the TSA division of Genbank - to produce a reference meta-transcriptome. The rationale

for mixing different sources of transcribed sequences was to obtain the most complete transcriptome possible, in order to better evaluate the specificity of transcripts in the synganglion versus all tissues combined (the whole body). We did not use other transcriptomes published for specific organs of *I. ricinus*, because they have been sequenced a relatively low coverage and show comparatively low completeness²⁸. To obtain the meta-transcriptome, we used the DRAP software (RunMeta)³⁰, a dedicated tool able to fuse contigs of different assemblies in a unique representative contig set. This tool allows to reduce redundancy within and between sets, and to eliminate low quality contigs with low read support. Completeness of the different transcriptome assemblies was assessed with BUSCO (version 4.0.2)³¹, using the arachnida_odb10 set of n=2,934 conserved genes.

Transcriptome annotation

Annotation of the reference transcriptome was done following the Trinotate pipeline (<https://github.com/Trinotate/Trinotate.github.io>)³². First, genes were predicted with TransDecoder.Predict (<https://github.com/TransDecoder/TransDecoder/wiki>), a prediction guided by PFAM-A searches with hmmscan (to recognize conserved domains) and a homology search with Diamond³³ blastp (<https://github.com/bbuchfink/diamond>) between long peptides and the uniprot_sprot database. Predicted peptides were then searched with SignalP (to determine the presence of signal peptides), tmhmm (to determine if transmembrane domains are present) and rnammer (to identify rRNA contigs). We also performed homology searches against the UniRef90 database, with Diamond blastp (for predicted peptides) and Diamond blastx (for transcripts). Finally an annotation report was generated to combine all results, including retrieved Gene Ontology terms, KEGG, Egglog and Pfam domains.

Annotation of cys-loop LGIC genes in the reference transcriptome.

A collection of reference cys-loop LGIC proteins comprising genes from *D. melanogaster* and the mite *T. urticae*¹³ (Acari, Acariformes) was established, to be used as tblastn queries on the reference meta-transcriptome of *I. ricinus*. A careful examination of the matches was then performed, to remove any remaining redundancy. Based on similarities within contigs and preliminary phylogenetic analyses, we identified events of chimerism among different gene sequences, which occurred in the GABA and nAChR families respectively. For these genes, we extracted matching reads, which were re-assembled with CAP3³⁴, using stringent parameters. This allowed to disentangle chimerisms and reconstructing complete sequences within gene families. Once we achieved the annotation of cys-loop LGICs and occasional correction of the initial contigs, these genes were re-integrated in the meta-transcriptome: to avoid redundancy, the edited and curated sequences were used and substituted to the corresponding initial contigs, and were annotated with the same Trinotate pipeline.

Read counts and Differential Expression Analyses

Alignment of reads with the meta-transcriptome including the curated cys-loop LGIC sequences were performed with Bowtie2(v2.4.1)-RSEM(v1.3.1), using default parameters. Expression levels were measured using the "count per million" metric produced by RSEM. Levels of expression were measured for eight libraries corresponding to the synganglion (produced for this study), but also, for comparison, for part of the "whole body" libraries used to build the reference meta-transcriptome as described above. For example, libraries from the GIDG transcriptome were not used because they were not replicated by condition. Read counts from the BioProject PRJNA395009 (GFVZ TSA assembly) were used, for all libraries except four shown to be of lesser quality in a previous study: library names and their accession are given in Table 1. First, to determine which transcripts were

more expressed in the synganglion, we made a differential expression analysis (DE) comparing synganglion and whole body libraries, with edgeR and limma^{35,36}. Second, to compare the patterns of expression in the synganglion of unfed and partially fed ticks (hereafter "unfed" and "fed" conditions), we used the read counts from the synganglion libraries only. The same methods were used in both DE analyses. We used the function `gene_to_trans_map` of edgeR, which assigned read counts to the same gene for contigs defined as isoforms based on their Trinity name (i.e. contigs that differed only by the final `_i` suffix). This resulted in 63194 unique genes. The statistics of each gene (read count, log-fold change, significance, and annotation) were visually explored with the Glimma package³⁷. Criteria for defining an up- or down-regulated gene were an absolute log-fold change >2 and an adjusted P-value (adj-P-value) < 0.05). The topGO package was used for the analysis of GO enrichment in genes up- or down-regulated in the synganglion (the whole body being taken as the reference) and for the analysis of enrichment in genes up- or down-regulated in the synganglion-"fed" libraries (the "synganglion-unfed" condition being taken as the reference) - (Alexa A, Rahnenfuhrer J (2021). *topGO: Enrichment Analysis for Gene Ontology*. R package version 2.44.0).

Functional expression of tick neuroreceptors

The coding sequence of *Rdl* was PCR-amplified using specific primers designed based on the synganglion transcriptome assembly (Iri-RDL-F0 GGTCAAGGAGGTCGCTTGCC; Iri-RDL-R0 ACGACAACCTTTAAAGGCGAATGC, FuIri-RDL-ptb-XhoF AGCGATGGCGTTCAGTTGCTG; FuIri-RDL-ptb-ApaR GACCGTGTGCACTATTCGTGCG). The *Rdl* PCR-product was subcloned into the transcription vector pTB-207³⁸ using the In-Fusion® HD Cloning Kit (Clontech™) and cRNAs were synthesized with the T7 mMessage mMachine kit (Ambion™). Expression of GABA-Rdl in *Xenopus laevis* oocytes and two-electrode voltage clamp electrophysiology were carried out as described previously³⁹. Briefly, 36 nL of 700 ng/μL *Rdl* cRNAs were micro-injected into defolliculated *Xenopus* oocytes (Ecocyte Bioscience) using a Drummond Nanoject II microinjector and incubated 3 days at 19°C to allow subunit expression. Electrophysiological recordings were performed using an Oocyte Clamp OC-725C amplifier (Warner Instruments) under voltage clamp at -60 mV and analyzed using the pCLAMP 10.4 package (Molecular Devices).

Phylogenetic analyses of the cys-loop LGICs

Because nAChRs constitute a divergent group within cysLGICs, we analysed separately this group of sequences. Thus, for respectively nAChRs and all other cysLGICs, we made alignments with MUSCLE, cleaned these alignments with GBLOCKS and further edited manually the alignments (with slight corrections to preserve conserved blocks). Phylogenetic maximum likelihood trees were obtained with IQ-TREE⁴⁰ the best model of substitution being determined with Model Finder⁴¹, and branch support being assessed with 1000 ultrafast bootstrap replication⁴². Phylogenetic trees were formatted graphically with ITOL⁴³.

Results

Transcriptome assembly and reference transcriptome

After sequencing a total of 69.7 Gb, the reads of libraries corresponding to synganglions of ticks in "fed" and "unfed" conditions were assembled separately. Both assemblies were combined with two more assemblies recently obtained for whole ticks transcriptomes, to obtain a reference meta-transcriptome. Detailed assembly statistics for the meta-transcriptome and for its four components are given in Table 2. Numbers of contigs were >600K for the synganglion assemblies. The percentage of predicted complete BUSCO genes were relatively even among these four assemblies,

ranging between 90.9 and 92.4%. The meta-transcriptome contained 70,125 contigs, a sharp decrease compared to the synganglion transcriptomes, which is explained by the filtering steps of Drap and the selection of contigs with a minimum read support, here 1 fpkm (Fragments Per Kilobase Million). The BUSCO completeness of the reference transcriptome (96.6%) was higher than any of the component assemblies, and the number of fragmented genes was very low (only 8).

Clustering of libraries based on read counts

We observed a strong clustering of libraries that corresponded to the same combination of tissue and feeding status, suggesting that other factors (stage, sex) influenced comparatively less the levels of expression (MDS plot, Figure 1). Indeed, the first axis of variation (axis 1, weight of 34%) clearly separated synganglion and whole body samples, whereas the second factor (axis 2, weight of 17%) separated the unfed and the partially fed conditions, respectively. Of note, expression levels were much less influenced by the feeding status in synganglion libraries, compared to whole body libraries. We also observe that the "fed-synganglion" libraries were somewhat less differentiated from whole-body libraries than the "unfed-synganglion" libraries (see Discussion). Given our focus on specific patterns of expression of the synganglion, we decided to make a differential expression analysis in two steps: first, we compared all synganglion libraries on one side versus all whole body libraries on the other side. Second, for the synganglion libraries only, we made a comparison between the "fed" and "unfed" samples, to evaluate the effect of the feeding status on expression levels in this tissue.

Differential expression analysis, comparison Synganglion vs Whole Body.

A total of 8483 genes were up-regulated in the synganglion (Syn+ genes) while 5040 genes were down-regulated (Syn- genes), the remaining 22592 genes showing no significant difference between the synganglion and whole body samples. Therefore ~37% of genes were found to be differently expressed among tissues. Many GO term associated with up-regulated genes for the Biological Process (BP) and Molecular Function (MF) categories were consistent with neuronal functions (Table 3). This is the case of the terms GO:0098632 (cell-cell adhesion mediator activity), GO:1904315 (transmitter-gated ion channel activity involved in regulation of postsynaptic membrane potential, a GO term associated with cys-loop LGICs). As for the localization of expression (Cellular component, CC), many of the enriched terms were consistent with post-synaptic membranes or neurons. We also note the importance of terms associated with a large family of membrane-associated proteins, the G protein-coupled receptors (GPCRs). An MD plot also allowed to visualize genes with a significantly change of expression level in both tissues (Figure 2). Among genes with the highest over-expression in the synganglion, we note the cases of three genes of interest (Putative apoptosis-promoting rna-binding protein tia-1/tiar rrm superfamily, Putative beta-amyloid-like protein, Microtubule-associated protein tau). Most of the cys-loop LGICs were characterized by a strong over-expression in the synganglion (Figure 3): this included the GABA-1 (*Rdl*) receptor, and the eight nAChRs. We note that three of the nAChRs (beta1, alpha1 and alpha2) had very high and roughly equal levels of expression in the synganglion, suggesting a possible co-expression and that these three genes could form heteromeric receptors. By contrast, the levels of expression of some of the histamine receptors were not significantly different between the synganglion and whole body samples respectively, as was the case of Ig1, the homolog of CG7589, CG6927, and CG11340 (forming the clade Insect group 1, in the house-fly). For the MF category, GO terms enriched in genes down-regulated in the synganglion included several linked terms (associated to the same genes): the first three categories (Iron ion binding, heme binding, aromatase activity) were indeed often found for genes belonging to the cytochrome P450 family (Supplementary Table 1).

Comparison Fed versus Unfed for the synganglion libraries

A total of 2243 genes were up-regulated in the fed condition (Fed+ genes), whereas 2305 genes were down-regulated (Fed- genes) - for the remaining 35667 genes, the expressions levels did not differ significantly between the fed and unfed samples. This amounted to ~11% of genes differently expressed among feeding conditions. The functional profiles of Fed+ genes showed notably an enrichment in genes associated with the metabolism of chitin and with the cuticle, whereas Fed- genes were enriched in a variety of GO terms, several of them being also enriched in the Syn+ category defined in the Synganglion vs Whole body comparison (e.g. terms associated with the synapse and with the plasma membrane). Among Syn+ genes, 1362 were Fed- whereas only 79 were Fed+.

*Functional expression of Rdl in *Xenopus laevis* oocytes*

A full-length cDNA was obtained by RT-PCR, using the primers designed using the *Rdl* contig. The corresponding cRNA was then microinjected into *Xenopus laevis* oocytes in order to monitor the potential functional expression of the RDL receptor. When challenged with 100 μ M GABA, the micro-injected *Xenopus* oocytes responded with robust currents in the μ A range (Figure 4A), providing a functional evidence of the expression of RDL. To further characterize this effect, the GABA concentration–response relationship was obtained by applying increasing concentrations of GABA (1 – 300 μ M). The GABA concentration-response curve revealed an EC₅₀ value of 19.95 μ M (log EC₅₀ = -4.7 ± 0.035 , $n = 18$), with current amplitudes normalized to the maximal response obtained to 300 μ M GABA (Figure 4B).

Phylogeny of AChRs from ticks and other Arachnida

The ML phylogenetic study of tick nAChRs, including outgroups from other Arachnida and from an insect, *D. melanogaster*, allowed us to define eight clades with strong bootstrap support (Figure 5). Because genes of this family have been named and numbered independently in each species, and due to duplication or gene loss in different branches, a consistent naming system valid for all species is impossible, so we numbered clades based on orthogroups in Arachnida. Four clades named α 1-4 were related to four genes in *D. melanogaster* (Dmel α 1, Dmel α 2, Dmel α 3 and Dmel β 2), but there was no one-to-one orthology between the genes from Arachnida and from *D. melanogaster* respectively. For example the α 1 and α 2 clade were co-orthologous to Dmel α 1 whereas the α 3 clade of Arachnida was orthologous to the pair of co-orthologs Dmel α 2 and Dmel α 3. The α 6 and β 1 clades were orthologous to *D. melanogaster* co-orthologs Dmel α 5-6-7 and Dmel β 1, respectively. A second beta-type clade of Arachnida genes, named β 2, was present both in ticks and other Arachnida but had no homologs in the genome of the house-fly nor in other insect genomes. This was also the case of a clade of alpha-type copies, that we named α 5. Last, we identified a group of divergent nAChR genes (characterized by very long branches) comprising both the sequence Dmel β 3 and sequences found in the spider *Parasteatoda tepidariorum*, but apparently without orthologs in ticks.

Phylogeny of the non-AChR cysloop-LGICs

The numbers of different cysLGICs in *I. ricinus* and other representative arthropods are summarised in Table 4 : the total was 21 in the house-fly, while 54 genes were found in the house spider (*P. tepidariorum*) and 46 in *I. ricinus*. Based on our phylogenetic tree supported by high bootstrap values, we could define eight major clades of non-AChR cysloop-LGICs in Arachnida, comprising ticks and predatory mites (Parasitiformes, e.g. *Galendromus occidentalis*), plant-feeding mites (Acariformes, e.g. *Tetranychus urticae*) and spiders (Figure 6):

- i) GABA-group: five different sub-groups were found, including GABA-1, corresponding to the "Resistant to dieldrin" (Rdl) homologs, GABA-2, corresponding to "ligand-gated chloride ion channel homolog 3" (lcch3), GABA-3 which was represented in several Arachnida groups, but not in *D. melanogaster*, GABA-4, corresponding to homologs of the CG8916 gene in *D. melanogaster*, and GABA-5, representing "GABA and glycine-like receptor" (grd) homologs. Of note, Rdl in *I. ricinus* was characterized by three alternative isoforms: their sequences were identical except in a stretch of 154 amino acids, which did align well but showed several substitutions, suggesting the presence of alternative exons (alignment shown in Suppl. Figure 1).
- ii) pH-sensitive chloride channels (pHCl): a single homolog was identified for *I. ricinus*, compared to three in *P. tepidariorum*.
- iii) "insect group 1": this clade comprised three copies in *D. melanogaster*, with one homolog in *I. ricinus* and two in *P. tepidariorum*.
- iv) Glutamate gated chloride channels: within this group, we could distinguish one "conserved" subclade - one homolog in *I. ricinus* (Glu1) but multiple copies in *P. tepidariorum* (2) or *T. urticae* (5) - and one "divergent" subclade represented in *I. ricinus* (5 copies) and *P. tepidariorum* (3 copies).
- v) Glycine receptors were represented by multiple copies in several groups of Arachnida, with 5 copies in *I. ricinus*.
- vi) Histamine-gated chloride-channels (HisCl) : three copies in *I. ricinus* (His1, His2 and His3) were found to be homologous of the genes HisCl1 and HisCl2 in *D. melanogaster*.
- vii) other clades: we found three more clades represented only in Arachnida, or only in ticks, all characterized by long branches, i.e. by divergent sequences. This was the case of a clade with one copy in *I. ricinus* (His 12): this clade grouped robustly with the Histamine clade, so we named it "Divergent histamine 1". A second clade, related to these genes, but with lower bootstrap support (<80) comprised genes in *I. ricinus* (His16-17-18), and also three genes in *P. tepidariorum* : we named it "Divergent histamine 2". The third clade, characterized by particularly divergent sequences, comprised as many as 14 copies in *I. ricinus* (named His4 to His15) for which we found no homolog in other Arachnida. In our phylogeny, this group branched with the gene CG12344 from *D. melanogaster*, but with low bootstrap support. Based on the genomic organization in *I. scapularis* (Supplementary Figure 2) we found that eight of these copies (His4-His11) formed a cluster of tandem copies immediately adjacent to genes of the conserved HisCl clade (*I. scapularis* sequences homologous to His1, His 2 and His3). This very close physical proximity suggest that all of these copies probably emerged through tandem duplication, and that the digergent clade (His4 to His 11 and His 13 to His15) and His-gated subunits have a common ancestor. We then named it "Divergent histamine 3".

Discussion

A meta-transcriptome of high completeness, enriched by synganglion transcriptome sequencing.

Our work is based on the high throughput sequencing of a synganglion transcriptome, a first for the tick *Ixodes ricinus*, but also on the combination of these data with two previous assemblies of tick whole body transcriptomes. This allowed us to obtain a meta-transcriptome of very high completeness (>96% BUSCO completeness), an essential step for comparing expression levels in the synganglion and in the whole body. This approach eliminated a large part of the noise associated with *de novo* RNA-Seq assemblies (our two assembled synganglion transcriptomes each had >600K contigs), as we obtained a compact reference assembly with ~70K contigs only. This reduction, due to the filtering of contigs with very low read support, could be associated with a loss of information, but high completeness metrics of the meta-transcriptome indicate that this risk was reduced. Overall, it appears that our newly acquired transcriptomic data for the synganglion has greatly improved the level of completeness of the meta-transcriptome, thanks to the inclusion of many transcripts that are highly specific to the synganglion.

High expression specificity and neuronal signature of the synganglion. The DE study comparing synganglion and whole body samples showed a strong association between genes up-regulated in the synganglion and neuronal/neurotransmission functions. Still, a possible limitation of our approach is that the tissues we dissected and determined to be the synganglion could have been contaminated by adjacent tissues or cells, for example due to the particular organisation of this tissue which is crossed by the oesophagus and surrounded by the fat body. Although it is difficult to rule out this possibility entirely, the high specificity and distinct functional profile of the synganglion libraries indicate that contamination must have been limited. Enriched categories in the synganglion include neuropeptides, ion channels, etc., many of which were either completely absent or represented by partial sequences in transcriptomes obtained from the whole body or other tissues. A large majority of cysloop-LGICs were also upregulated in the synganglion, which is expected for these ion channels known to be expressed in neurons, which are particularly concentrated in the CNS. This overexpression concerns even divergent sequences of the cysLGICs (in the GluCl_s, or His-gated groups expanded clades), suggesting that these subunits are active mainly in the tick CNS. For the His-gated group, however, we note the exception of His12 ("divergent Histamine-1" clade) which was downregulated in the synganglion. A few of the cysLGIC genes were not differentially expressed (Synganglion vs Whole body comparison), including the homolog of 'Insect group 1' (a group comprising CG7589, CG6927, and CG11340 in *D. melanogaster*). This is consistent with a study that showed that IG1 is non-neuronal in this species⁴⁴. Although our study focused primarily on one class of ion channels, the cys-loop LGICs, our assemblies (the synganglion assemblies as the meta-transcriptome) provide rich material for gene discovery in many functional groups associated with neurotransmission or neuronal functions in general. For example, we found that several of the genes with the highest log-fold variation in expression in the synganglion have vertebrates homologues which have been extensively studied for their association with Alzheimer's Disease (AD): this is the case for the APP-like peptide (APPL) and of tau protein⁴⁵, as well as a homolog of TIA-1, a gene associated with the development of tau protein⁴⁶. Interestingly, a putative tau homolog was also identified in the synganglion transcriptome of another tick species, *Dermacentor variabilis*, where it has been reported to be strongly up-regulated upon initiation of blood feeding (unfed versus partially-fed ticks)⁸. In addition to this, in our study, a contig identified as Cathepsin B was among the most strongly downregulated genes in the synganglion. Cathepsin B has been shown to have complex interactions with AD and to be involved in the degradation or production of Abeta, as reviewed in⁴⁷. In *D. melanogaster*, APPL has been shown to be restricted to the nervous system, and the authors hypothesized that a neural-specific function encoded by the APP gene has been selectively maintained along evolution⁴⁸. In the tick synganglion, the strong up-regulation (tau, APP, TIA1) or down-regulation (CathB) of genes known to be involved in AD pathology in humans illustrates the interest of arthropod models to study the expression profiles of these genes, their interactions and their evolution.

Profiles of downregulated genes in the synganglion. For a complete understanding of tick neurobiology, it is also interesting to assess which genes are downregulated in the synganglion. Genes in this category showed enrichment in terms associated with iron binding, heme binding, and aromatase activity, in particular. In a previous study²⁸ where we investigated expression in the whole body of the tick, comparing fed and unfed conditions, genes downregulated during feeding (i.e. significantly more expressed in the whole body of unfed ticks than of fed ticks) were particularly enriched in the same first three terms. We found that these terms were co-occurring in the same genes, most of which of the cytochrome P450 (cytP450) family. This suggests that one group of cytP450 tends to be particularly expressed in the whole body of unfed ticks, but is expressed at a much lower level both in the whole body of partially fed ticks and in the synganglion of ticks of all conditions.

Differentiation between the "fed" and "unfed" state. Our design allowed to compare the expression levels in the synganglion of respectively unfed ticks (males or females) and partially fed female ticks. The transcriptomes of these two conditions were comparatively less differentiated than the synganglion and whole body samples (11% and 36% of DE genes in the two comparisons, respectively). Up-regulated genes upon feeding (fed+ genes) were enriched in chitin-associated proteins, a result that appears surprising given the absence of cuticle in this tissue. In a previous transcriptome study of the tick whole body²⁸, cuticle associated proteins were also the most up-regulated genes upon feeding, suggesting that an increased expression of cuticle-related products upon feeding is a general pattern. To determine the intersection of changes upon feeding (in the synganglion) and of genes of expression among tissues (synganglion vs whole body), we counted the genes that were both differentially expressed in the fed vs unfed comparison and Syn+: with this filtering, up-regulated genes in the fed condition were down to 79 (from 2243) whereas down-regulated genes were down to 1362 (from 2305). This suggests that extremely few genes up-regulated in the synganglion were Fed+, and then that the 'cuticle' signature of Fed+ genes is not specific to the synganglion, but is a global pattern in the tick body. By contrast, a relatively large fraction of genes were at the same time Syn+ and downregulated in the synganglion upon feeding. Consistently, Fed- genes had functional profiles relatively similar to Syn+ genes (which can be seen comparing Table 4 and Suppl. Table 3). Overall, this indicates that the expression profile of the synganglion of unfed ticks has a more distinct profile (compared to the whole body) than the synganglion of feeding ticks. We observed this precise pattern for most of the cys-loop LGIC gene (most of them being Syn+), since their expression levels were often comparatively lower in the fed samples than in the unfed samples, as was the case of GABA-1 (Rdl), Glu-6, PhCl for example. We propose that the synganglion overall activity could be lowered when ticks are fixed on a host and start to feed, due to the down-regulation of several functions associated with mobility or sensorial activities related to host-search, for example. This can be compared to another study of the synganglion of unfed, part-fed and replete ticks⁸: in this study, expressions profiles of the latter two stages were relatively similar, whereas the unfed state had the most dissimilar profile, and the "replete" stage showed a decline in expression and specificity of expression, which was linked to a possible shift from active stages of the tick life-cycle (host search, feeding, mating) to late stages (oviposition, senescence and death).

Completeness of cys-loop LGICs. Our synganglion transcriptomes allowed us, thanks to a careful manual curation, to obtain a remarkably large and complete collection of cys-loop LGICs compared to previous studies, even when based on complete genome sequences: for example, a compilation of cys-LGICs in the *I. scapularis* genome⁴ reported a total of 32 genes, less than the 44 found in our study. In the recent study of five tick complete genomes¹², no specific compilation of cys-LGICs was reported, but our phylogenetic studies and the alignments we obtained indicate that for all these species, the cysLGIC record is less complete than in our study, with either some genes entirely missing or having only partial sequences, possibly due to annotation issues. This difference may be explained by difficulties associated with assembling and annotating the complex genomes of ticks, and illustrates the high value of transcriptome data for the tick CNS.

Conclusions: our new transcriptome data for the tick synganglion, thanks to its quality and high coverage, allowed us to build a high quality reference transcriptome for *Ixodes ricinus*, strongly enriched in transcripts specific or at least strongly over expressed in neurones. This allowed us to reconstruct a large catalogue of the tick LGICs, showing that for the nAChRs and for the GABA group, this tick species has a repertoire comparable to other arthropods, while it shows marked expansions of other cysLGICs, especially the GluClIs and the Histamine gated-like group. For GABA-Rdl, an internal duplication (triplication of an exon) allows the production of three alternative transcripts, either with or without a mutation conferring resistance to dieldrin. Based on these synganglion transcriptomic ressources, we achieved the first functional reconstitution of a

GABA-gated channel for *Ixodes ricinus*, made of the RDL subunit. Future studies will be needed to evaluate the pharmacological and biophysical properties of RDL and of other neuroreceptors, including sensitivity to insecticides.

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Availability of data : Read sequences of the synganglion transcriptomes have been published to EBI as BioProject PRJEB40721. The nucleotide sequences of cysLGICs have been deposited to GenBank as MZ027281-MZ027288 (n=8 nAChRs) and MZ099581-MZ099616 (other cysLGICs, n=36 transcripts, n=34 different genes). In addition, the gly1 and gly2 sequences correspond to accessions GIDG01020391.1 (gly1, coding sequence 1-1978) and GIDG01020392.1 (gly2, coding sequence 1-1993). Contig sequences of the meta-transcriptome, their annotation table, interactive plots of the Synganglion-vs-Whole body differential expression analysis, non-aligned and aligned protein sequences of the cysLGICs are available in a public repository (<https://data.inrae.fr/>): Rispe, Claude, 2021, "Transcriptome of the the tick synganglion", <https://doi.org/10.15454/FGTIHR>".

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Table 1: List of libraries used for the comparison of expression levels in the synganglion and in the whole body. Accession: Accession numbers of reads (synganglion libraries, generated for the present study, were deposited as BioProject PRJEB40724, whereas whole body samples correspond to BioProject PRJNA395009, produced in a previous work of our group²⁸). Feeding condition: ticks were either unfed (collected on the vegetation) or fed (semi-engorged state of repletion, ticks collected on roe deers for the synganglion libraries and on mice or rabbits for the whole body libraries). Stage/sex: indicates if ticks were female or male (for adults), or nymphs (the sex of nymphs could not be determined). Tissue: libraries were obtained from the dissected synganglion (this study) or from tick whole bodies (already published sequence data).

Library name	Accession	Feeding Condition	Stage/sex	Tissue
F01	SAMEA7383684	Unfed	females	Synganglion
F02	SAMEA7383685	Unfed	females	Synganglion
F03	SAMEA7383686	Unfed	females	Synganglion
M01_02	SAMEA7383691	Unfed	males	Synganglion
M03	SAMEA7383687	Unfed	males	Synganglion
FSG01	SAMEA7383688	Fed	females	Synganglion
FSG02	SAMEA7383689	Fed	females	Synganglion
FSG03	SAMEA7383690	Fed	females	Synganglion
N1	SRR5839703	Unfed	nymphs	Whole body
N2	SRR5839701	Unfed	nymphs	Whole body
M1	SRR5839697	Unfed	males	Whole body
M2	SRR5839696	Unfed	males	Whole body
F1	SRR5839704	Unfed	females	Whole body
F2	SRR5839708	Unfed	females	Whole body
F3	SRR5839707	Unfed	females	Whole body
NG1	SRR5839700	Fed	nymphs	Whole body
NG2	SRR5839699	Fed	nymphs	Whole body
NG3	SRR5839698	Fed	nymphs	Whole body
FG	SRR5839710	Fed	females	Whole body

Table 2: Statistics on transcriptome assemblies for *I. ricinus*, including two previously published datasets (transcriptomes of tick whole bodies), two transcriptome assemblies obtained for the tick synganglion (this study), and for the reference meta-transcriptome we obtained combining these assemblies, with Drap. Assembly short name (TSA accession for whole body transcriptomes), Tissue, stages, Read type (length and strandedness), Total sequenced bases, Number of contigs, N50 metric of each assembly, Maximum contig size, Busco completeness metrics, using the Arachnida odb10 data set (S, single and complete, D, duplicated and complete, F, fragmented, M, missing, %C, percentage of complete genes).

Assembly	Tissue	Stages/Conditions	reads type	bases (Gb)	#contigs	N50	max size	Busco completeness (Arachnida, odb10)				
								S	D	F	M	%C
GFVZ	Whole Body	males, nymphs (fed or unfed), adult females (fed or unfed)	oriented, 2x125 bp	20.10	179316	875	20233	2213	473	54	194	91.5
GIDG	Whole Body	eggs, larvae, nymphs (fed or unfed) and adult females (fed or unfed)	non oriented, 2x50 bp	21.52	55046	1979	24178	1852	816	38	228	90.9
SYN-NG	Synganglion	males and adult females, unfed	oriented, 2x150 bp	37.08	695933	523	21882	1518	1195	56	165	92.4
SYN-FSG	Synganglion	adult females, fed	oriented, 2x150 bp	32.59	602593	466	33016	1484	1203	58	189	91.6
meta-transcriptome 1fpkm	-	-	-	111.29	70125	1915	24178	2314	519	8	93	96.6

Table 3: Top 20 enriched Gene Ontology terms among genes significantly up-regulated in the synganglion, compared to the whole body. Columns, GO IDs and terms for Molecular Function, Biological Process and Cellular localization; Annotated : number of transcripts with each GO; Significant: number of genes annotated with this term in the enriched category. Expected: expected number of genes. Weight01: level of significance of the enrichment.

GO.ID	Term	Annotated	Significant	Expected	weight01
Molecular Function					
GO:0005515	protein binding	6664	1677	1263.4	3.5e-19
GO:0098632	cell-cell adhesion mediator activity	49	37	9.3	1.3e-17
GO:0005509	calcium ion binding	569	177	107.9	1.2e-12
GO:1904315	transmitter-gated ion channel activity involved in regulation of postsynaptic membrane potential	56	38	10.6	5.2e-11
GO:0043565	sequence-specific DNA binding	830	252	157.4	2.3e-10
GO:0001228	DNA-binding transcription activator activity, RNA polymerase II-specific	167	66	31.7	4.8e-10
GO:0001227	DNA-binding transcription repressor activity, RNA polymerase II-specific	127	53	24.1	2.5e-09
GO:0004930	G protein-coupled receptor activity	212	115	40.2	1.1e-08
GO:0008528	G protein-coupled peptide receptor activity	71	46	13.5	1.6e-07
GO:0005544	calcium-dependent phospholipid binding	34	20	6.5	3.0e-07
GO:0000981	DNA-binding transcription factor activity, RNA polymerase II-specific	486	180	92.1	5.3e-07
GO:0004971	AMPA glutamate receptor activity	11	10	2.1	5.4e-07
GO:0005244	voltage-gated ion channel activity	110	61	20.9	5.4e-07
GO:0004984	olfactory receptor activity	15	12	2.8	5.4e-07
GO:0015277	kainate selective glutamate receptor activity	20	14	3.8	9.2e-07
GO:0008188	neuropeptide receptor activity	35	26	6.6	1.3e-06
GO:0030552	cAMP binding	18	13	3.4	1.3e-06
GO:0004383	guanylate cyclase activity	26	16	4.9	2.0e-06
GO:0000978	RNA polymerase II proximal promoter sequence-specific DNA binding	257	82	48.7	2.1e-06
GO:0005096	GTPase activator activity	246	77	46.6	2.1e-06
Biological Process					
GO:0007156	homophilic cell adhesion via plasma membrane adhesion molecules	136	77	26.48	2.80E-22
GO:0007186	G protein-coupled receptor signaling pathway	442	215	86.06	6.10E-20
GO:0007268	chemical synaptic transmission	612	286	119.16	1.40E-17
GO:0045944	positive regulation of transcription by RNA polymerase II	771	239	150.12	1.80E-17
GO:0007411	axon guidance	385	167	74.96	7.50E-17
GO:0007165	signal transduction	3588	1106	698.62	2.10E-16
GO:0007218	neuropeptide signaling pathway	47	34	9.15	5.70E-15
GO:0070593	dendrite self-avoidance	75	43	14.6	4.40E-13
GO:0034765	regulation of ion transmembrane transport	228	105	44.39	8.00E-13
GO:0007155	cell adhesion	816	300	158.88	7.80E-12
GO:0042391	regulation of membrane potential	260	107	50.62	8.30E-12
GO:0050877	nervous system process	782	290	152.26	2.30E-10
GO:0000122	negative regulation of transcription by RNA polymerase II	563	164	109.62	1.10E-08
GO:0035556	intracellular signal transduction	1969	602	383.38	1.40E-08
GO:0048149	behavioral response to ethanol	47	26	9.15	5.20E-08
GO:0043113	receptor clustering	60	30	11.68	6.00E-08
GO:0045892	negative regulation of transcription, DNA-templated	867	255	168.81	7.50E-08
GO:0008045	motor neuron axon guidance	90	39	17.52	1.90E-07
GO:0007612	learning	131	57	25.51	2.40E-07
GO:0110111	negative regulation of animal organ morphogenesis	27	17	5.26	3.50E-07
Cellular Component					
GO:0005886	plasma membrane	3677	1195	734.53	< 1e-30
GO:0030054	cell junction	1061	387	211.95	< 1e-30
GO:0045211	postsynaptic membrane	281	154	56.13	6.00E-24
GO:0005887	integral component of plasma membrane	935	368	186.78	1.00E-23
GO:0030425	dendrite	515	207	102.88	4.40E-16
GO:0043005	neuron projection	1224	489	244.51	5.20E-15
GO:0030424	axon	686	283	137.04	5.30E-15
GO:0098839	postsynaptic density membrane	57	38	11.39	8.60E-15
GO:0014069	postsynaptic density	279	129	55.73	2.10E-12
GO:0098978	glutamatergic synapse	206	87	41.15	2.20E-12
GO:0043204	perikaryon	138	62	27.57	2.50E-11
GO:0042734	presynaptic membrane	121	61	24.17	4.10E-10
GO:0031594	neuromuscular junction	120	57	23.97	4.90E-10
GO:0008076	voltage-gated potassium channel complex	27	20	5.39	2.00E-09
GO:0043025	neuronal cell body	483	183	96.49	3.50E-09
GO:0016021	integral component of membrane	4100	1093	819.03	3.80E-09
GO:0043195	terminal bouton	99	52	19.78	4.20E-08
GO:0005604	basement membrane	90	40	17.98	1.20E-07
GO:0099056	integral component of presynaptic membrane	37	22	7.39	1.60E-07
GO:0043194	axon initial segment	16	13	3.2	2.40E-07

Table 4: Numbers of cys-loop LGICs in *D. melanogaster* and in three Chelicerata, the house spider *P. tepidariorum*, the mite spider *T. urticae* and the tick *I. ricinus*.

	<i>D. melanogaster</i>	<i>P. tepidariorum</i>	<i>T. urticae</i>	<i>I. ricinus</i>
<i>Nicotinic Acetylcholine receptors (nAChRs)</i>				
alpha1-2-3-4 group	4	9	3	4
beta1 group	1	2	1	1
alpha6 group	3	4	3	1
beta2 group	0	2	0	1
acr26-like group	0	1	1	1
Divergent nAChRs	1	5	2	0
<i>GABA-gated ion channels</i>				
Rdl	1	1	3	1
Grd	1	1	0	1
Lcch3	1	2	0	1
CG8916	1	1	0	1
Other	0	2	0	1
<i>Other Cys-loop LGICs</i>				
Histamine-gated	2	10	4	18
Group 12344	1	0	4	0
Glutamate-gated	1	5	6	6
pHCl	1	3	1	1
Insect group1	3	2	0	1
Glycine group	0	4	1	5
Total cys-loop LGICs	21	54	29	44

Figure 1: MDS plot, based on the read counts of libraries used for a differential expression analysis. Plot shown for dimensions 1 and 2, the two axes with the highest weight. Synganglion libraries were produced in this study, whole body libraries were from a previous study (as described in Table 1). Synganglion of unfed ticks (open diamonds), synganglion of fed ticks (filled diamonds), whole body of unfed ticks (open circles), whole body of fed ticks (filled circles).

Figure 2: MD plot of all genes in the reference transcriptome, in a comparison of expression among synganglion and whole body libraries. In y-axis, log-fold change between the expression level in the synganglion and in the whole body. In x-axis: average log count per million of reads (averaged across all libraries, indicating the overall level of expression). Dots in red correspond to genes that are significantly up-regulated in the synganglion (logfold change >2 and P-adjusted value <0.05); dots in blue correspond to genes that are significantly down-regulated in the synganglion (logfold change <2 and P-adjusted value <0.05); dots in gray are genes with an expression level not significantly different among tissues. Three outliers with among the highest values of logfold-change are distinguished (blastx description between parentheses): 1 (Putative apoptosis-promoting rna-binding protein tia-1/tiar rrm superfamily), 2 (Putative beta-amyloid-like protein), 3 (Microtubule-associated protein tau).

Figure 3: Comparison of expression counts, measured in counts per million reads (cpm) with RSEM, for the cys-loop LGICs. Expression counts were averaged for all synganglion libraries (y-axis) and for all whole body libraries (x-axis). Colors of dots indicate if genes were found to be significantly up-regulated in the synganglion (red), down-regulated in the synganglion (blue) or with no sig. difference (grey). The dotted-line corresponds to $y=x$. For both axes, a logarithmic scale was used.

Figure 4 : Concentration–response relationship of GABA on the *I. ricinus* RDL receptor expressed in *Xenopus* oocytes. A. Representative current traces of a single oocyte microinjected with *Iri-rdl* cRNA perfused with increasing concentrations of GABA for 10 seconds (short bars). The concentration of GABA (μ M) is indicated above each trace. B. Concentration–response curve for GABA on the Iri-RDL channel. All current responses are normalized to 300 μ M and shown as the mean $\pm$ SEM.

Figure 5: Maximum-likelihood phylogeny of nAChRs, including eight sequences identified in the synganglion transcriptome of the tick *I. ricinus* and their homologs in Arachnida (other tick species, other Parasitiformes, Acariformes) or in *Drosophila melanogaster*. Labels indicate the accession numbers of protein sequences, and species name (abbreviated as D for *D. melanogaster*). Labels for *I. ricinus* sequence are in blue. Filled circles on branches indicates bootstrap support (support increases with circle width, ranging from 80 to 100).

Figure 6: Maximum-likelihood phylogenetic tree of cys-loop LGICs subunit protein sequences, excluding nAChRs, in the tick *I. ricinus* and representative arthropods: *P. tepidariorum* (house spider), Acariformes and Parasitiformes, *D. melanogaster*. Labels indicate accession numbers of protein sequences and species name. Labels for *I. ricinus* sequence are in blue. Filled circles on branches indicates bootstrap support (support increases with circle width, ranging from 80 to 100).

A

B

Tree scale: 1

pH-sensitive chloride channels

GABA
group

**glutamate-gated
chloride channels**

**glycine
receptors**

histamine-gated chloride channels

**divergent
histamine-2**

**divergent
histamine-1**

divergent
histamine-3