Conservation of Silk Genes in Trichoptera and Lepidoptera

Naoyuki Yonemura · Kazuei Mita · Toshiki Tamura · František Sehnal

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Abstract Larvae of the sister orders Trichoptera and Lepidoptera are characterized by silk secretion from a pair of labial glands. In both orders the silk filament consists of heavy (H)- and light (L)-chain fibroins and in Lepidoptera it also includes a P25 glycoprotein. The L-fibroin and H-fibroin genes of Rhyacophila obliterata and Hydropsyche angustipennis caddisflies have exon/intron structuring (seven exons in L-fibroin and two in H-fibroin) similar to that in their counterparts in Lepidoptera. Fibroin cDNAs are also known in Limnephilus decipiens, representing the third caddisfly suborder. Amino acid sequences of deduced L-fibroin proteins and of the terminal H-fibroin regions are about 50% identical among the three caddisfly species but their similarity to lepidopteran fibroins is <25%. Positions of some residues are conserved, including cysteines that were shown to link the L-fibroin and H-fibroin by a disulfide bridge in Lepidoptera. The long internal part of H-fibroins is composed of short motifs arranged in species-specific repeats. They are extremely uniform in R. obliterata. Motifs (SX)₆, GGX, and GPGXX occur in both Trichoptera and Lepidoptera. The trichopteran H-fibroins further contain charged amphiphilic motifs but lack the strings of alanines or alanine-glycine dipeptides that are typical lepidopteran motifs. On the other hand, sequences composed of a motif similar to ERIVAPTVITR surrounded by the (SX)₄₋₆ strings and modifications of the GRRGWGRRG motif occur in Trichoptera and not in Lepidoptera.

Keywords Silk evolution · Trichoptera · Lepidoptera · Fibroin · DNA repeats · Protein polymers · Insect genes

Introduction

Several groups of terrestrial arthropods produce proteins that polymerize into fibers known as silk (Sehnal and Akai 1990; Craig 1997). Silk secretion from a pair of labial glands is characteristic for the larvae of several clades of Holometabola and reaches perfection in the supraorder Amphiesmenoptera, which includes Trichoptera (caddisflies) and Lepidoptera (moths and butterflies) (Akai et al. 2003). Studies on the major commercial silk producer, the domestic silkworm Bombyx mori, revealed that proteins secreted in the posterior section of each gland polymerize into a core silk filament; the pair of filaments is sealed into a single fiber by a layer of sericins derived from the middle section of the glands. The filaments consist of three proteins known as heavy- and light-chain fibroins (H- and L-fibroins, respectively) and the P25 protein or fibrohexamerin (Tanaka et al. 1999a). Disulfide linkage between the high molecular H-fibroin (>350 kDa) and the L-fibroin (~25 kDa) proved indispensable for the secretion of both components (Takei et al. 1987). P25 occurs as two differentially glycosylated moieties of ca. 27 and 31 kDa. Their interaction with the H-fibroin N-terminus is believed to facilitate storage of the highly insoluble H-fibroin/L-fibroin dimer in the form of a gel in the silk gland lumen, as well as gel conversion into the solid filament during spinning (Inoue et al. 2004). Identification of homologous silk components in other Lepidoptera (reviewed by Fedič et al. 2009).
including the ancient superfamilly Yponomeutoidea (Yonemura and Sehnal 2006), indicated that this composition of the silk filament occurs in most Lepidoptera. It was probably secondarily lost in the evolutionary advanced family Saturniidae, whose silk filament is made of H-fibroin homodimers without participation of either L-fibroin or P25 (Tamura et al. 1987; Tanaka and Mizuno 2001).

The existence of distinct secretions from the posterior and the middle silk gland sections was demonstrated by histological staining in the caddisfly larvae. By analogy with Lepidoptera, the product from the posterior section was referred to as “fibroins” and that from the middle section as “sericins” (Zaretschnaya 1966; Engster 1976). A systematic search for silk components homologous to the lepidopteran silk proteins was undertaken in the caddisflies Limnephilus decipiens and lepidopteran silk proteins was undertaken in the caddisflies Limnephilus decipiens and Hydropsyche angustipennis (Yonemura et al. 2006). The first species was chosen as a representative of the suborder Integripalpia (the tube case-makers), which use silk rather sparsely for stitching portable larval cases from the pieces of plants, sand, or other foreign materials. Larvae of the second species (suborder Annulipalpia, the retreat-makers) produce large amounts of silk to spin food-collecting nets and retreat tunnels. Both species proved to express mRNAs encoding proteins homologous to the lepidopteran H-fibroin and L-fibroin in the posterior silk gland section. The search for a P25 homologue was unsuccessful at both the nucleic acid and the protein level. It was tentatively concluded that the H-fibroin and L-fibroin genes originated in ancestral Amphiesmenoptera, whereas P25 evolved in Lepidoptera after their separation from Trichoptera. However, no data were available on the silk composition in the third caddisfly suborder, the cocoon-makers Spicipalpia, which use silk mainly for cocoon construction, similarly to many Lepidoptera. The present paper describes H-fibroin and L-fibroin transcripts in Rhyacophila obliterata from the Spicipalpia and shows that the H-fibroin and L-fibroin genes in this species and in H. angustipennis are homologous to those of Lepidoptera. The P25 silk component is apparently lacking in Trichoptera.

Materials and Methods

Insect Collection and Sample Preparation

Fully grown larvae of Rhyacophila obliterata (suborder Spicipalpia, family Rhyacophilidae) and Hydropsyche angustipennis (suborder Annulipalpia, family Hydropsychidae) were collected in the Czech Republic, at about 49°N, 13°E. Those of R. obliterata occurred in a pristine river in the Sumava Mountains (about 1000 m a.s.l.) in July to August, and larvae of H. angustipennis were taken from a small brook in the vicinity of the town České Budějovice (about 400 m a.s.l.) in April to June. Both species pupated in cases attached to stones as small domes made of sand grains. Inside the case the larva spun a cocoon that was a thin mesh of silk fibers in H. angustipennis and a more solid structure composed of cross-linked and mutually fusing fibers in R. obliterata.

Caddisfly larvae were brought to the laboratory in shallow containers cooled by ice. Dissected silk glands were mostly frozen in liquid nitrogen and used for nucleic acid analysis. Some freshly dissected glands were ruptured in a small volume of chilled distilled water (50 μl per a gland pair) and the silk gel flown out within 30 min was collected into a pipette. The gel and water admixture was briefly vortexed with an equal volume of 10 mM Tris buffer, pH 7.0, containing 8 M urea, 2% SDS, and 5% 2-mercaptoethanol. The solution was left overnight at 4°C, then centrifuged, and the supernatant was taken for polyacrylamide electrophoresis. The spun-out silk used for stitching the sand grains into protective domes was analyzed in R. obliterata. The silk fraction solubilized in 8 M guanidinium isothiocyanate was also subjected to electrophoresis.

Silk Gland-Specific cDNA Library

RNA was prepared from 34 pairs of R. obliterata silk glands that were pulverized in a mortar under liquid nitrogen. The powder was extracted with 750 μl Isogen-LS reagent (Nippon Gene Co. Ltd., Tokyo), proteins and genomic DNA were removed by partitioning with 250 μl chloroform, and total RNA was precipitated from the remaining watery phase with an equal volume of 2-propanol. The precipitate was rinsed with 70% ethanol, dried briefly at room temperature, dissolved in 0.5% SDS with 20 mM sodium acetate (pH 5.3), and stored at −80°C. The cDNA library was constructed commercially by Takara Bio Inc. (Ohtsu City, Japan) with the ZAP-cDNA Synthesis Kit (Stratagene, La Jolla, CA, USA). The kit employs StrataScript reverse transcriptase for first-strand and Pfu DNA polymerase for complementary-strand cDNA synthesis; EcoRI and XhoI restriction sites are inserted for further cloning.

Randomly chosen clones were sequenced in both directions with the T3 or T7 primers on the ABI PRISM 3730 Genetic Analyzer (Applied Biosystems). Sequences (typically exceeding 700 nt) were edited with ABI PRISM Sequencing Analysis software, version 3.3, and those 95% identical over a stretch of more than 300 nt were assembled in clusters. The validity of cDNA clustering and the mutual relationships of different clusters were checked with respect to both the nucleotide sequence and the nature of
encoded proteins with software used in the *Bombyx* genome project (Mita et al. 2003). Contigs were assembled from the clustered tags with the software ABI PRISM AutoAssembler, version 2.0. Clustering might be incorrect in the case of cDNAs containing long repeats with reiterated motifs that could be aligned in more than one modality. To verify faithfulness of the assembly, each 5′ end sequenced. Since the inserts were of different lengths, the overlapping array of their 5′ ends had to match the contig, which was based on the assembly of sequences obtained from automatic analysis of the cDNA clones.

Targeted cDNA Analysis and Sequencing

5′ RACE PCR was employed to identify the 5′ end of the long cDNAs that were incomplete in the library. Total RNA was prepared from silk glands with the Isogen-LS reagent (Nippon Gene Co.) and used for 5′ amplification with the BD SMART RACE cDNA Amplification Kit (BD Biosciences); reverse primers (Table 1) were derived from sequences identified in the cDNA library. PCR products were either sequenced directly or cloned into the T-vector pCR4-TOPO or pCR-XL-TOPO (Invitrogen). Sequencing employed BigDye Terminator version 3.1 and was done on ABI Genetic Analyzer 3100 or 3130 (Applied Biosystems).

Protein Analysis

Deduced amino acid sequences of the contigs were used as queries for the BLAST search in public protein databases and in our files of the silk gland cDNAs identified in the caddisflies *H. angustipennis* and *L. decipiens* (Yonemura et al. 2006). Proteins were regarded as similar if they were more than 30% identical over a region longer than 100 amino acid residues. The presence of expected proteins in the silk of *R. obliterata* was verified by electrophoretic analysis of the silk extracts in 8 M urea (Yonemura and Sehnal 2006). Blots of selected proteins in the Immobilon-P membrane (Millipore, Bedford, MA, USA) were sent to the commercial facility at the Medical College of Wisconsin for N-terminal sequencing.

Analysis of Genomic DNA

Last-instar larvae of *H. angustipennis* and *R. obliterata* were individually pulverized in mortars under liquid nitrogen. The powder was suspended in 2 ml G2 buffer (800 mM guanidine HCl, 30 mM Tris- HCl, pH 8.0, 30 mM EDTA, pH 8.0, 5% Tween-20, 0.5% Triton X-100) supplemented with 400 μg RNase A and 150 mU proteinase K. The suspension was incubated overnight at 50°C and centrifuged (20,000 g at 4°C for 10 min). The supernatant was loaded on a buffer-equilibrated genomic-tip column (Qiagen). Eluted DNA was precipitated with 2-propanol (70% of the elute volume), washed with 70% ethanol, dried briefly, and resuspended in 50 μl 10 mM Tris-HCl, pH 8.5. Aliquots containing 20 ng genomic DNA were taken for PCR with primers deduced from the cDNA and later also from the genomic sequences (Table 1). PCR analyses of genomic DNA preparations were done with Advantage 2 polymerase mix (BD Biosciences) and high-fidelity PCR with DNA polymerases KOD FX (Toyobo, Osaka, Japan) or *Pfu* Ultra II (Stratagene). PCR products were separated by agarose gel electrophoresis; short products were visualized with ethidium bromide under UV light, and long products susceptible to UV damage were stained with Crystal Violet. DNA was extracted from agarose with the MiniElute Gel Extraction kit (Qiagen) and 10-μl eluates in 10 mM Tris-HCl, pH 8.5, were either taken for direct sequencing with the PCR primers or cloned into pCR4-TOPO or pCR-XL-TOPO vectors. This TA cloning required 3′ adenine overhang, which was present in PCR products amplified with the Advantage 2 polymerase mix, while DNA products obtained with polymerase KOD FX or *Pfu* Ultra II had to be incubated with 2.5 U ExTaq (Takara), 1 μM dATP, and 1 × ExBuffer for 15 min at 72°C.

Inverse PCR

Samples of 100 ng genomic DNA prepared as described above were digested with 10 U BanI, HaeII, and *XhoI*, respectively, or with a mix of EcoRI and *MunI* (all from Takara Bio Inc.) for 3 h. The digest was incubated overnight at 4°C with 100 U T4 ligase (New England Biolabs). Self-ligated circular fragments were precipitated and washed with ethanol, dried briefly, and dissolved in distilled water. PCR products amplified with the Advantage 2 polymerase mix were cloned into the pCR4-TOPO vector.

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1 Amplification and cloning of DNA sequences up to about 5000 bp were most conveniently done with the Advantage 2 polymerase mix. According to the User Manual for TA cloning, this mix introduces 25 errors per 100,000 bp after 25 PCR cycles. We found a slightly higher error rate, about 1 per 3000. Toyobo Co. reported that KOD FX DNA polymerase exhibited 19 errors per 144,535 bp, which is 11 times less error rate, about 1 per 3000. Toyobo Co. reported that KOD FX DNA polymerase exhibited 19 errors per 144,535 bp, which is 11 times less accurate than the *Pfu* DNA polymerase. According to the technical support section of Stratagene, the 25 × 10^7 figure indicates the occurrence of 1 mutation in 2,500,000 colonies of the amplified lac gene; *Taq* accuracy in a similar test was about 1.25 × 10^7. The DNA polymerase *Pfu* Ultra II Fusion HS is, in our experience, about 3 times more accurate than the standard *Pfu* polymerase and 20 times better than the *Taq* polymerase.
Table 1 Key primers referred to in the text

| Primer | Sequence | GenBank reference | Transcript | Use |
|--------|----------|------------------|------------|-----|
| HaH-F42 | 5’GC GGCAATTCTCTCTGATCTTTATCTGG 3’ | AB354591/1465 | 31 | PCR for intron detection |
| HaH-R41 | 5’GCCGCGCAACCACTCCTACGATCTCTC 3’ | - / 1944 | 510 | PCR for intron detection |
| HaH-F51 | 5’GAAGTCCTGGATGGTTGTGCAGCCGC 3’ | - / 1919 | 485 | Inverse PCR of 5’ region |
| HaH-R51 | 5’GAATAGATGAGAGATTTGGCAGGGG 3’ | - / 1488 | 54 | Inverse PCR of 5’ region |
| HaH-F40 | 5’CTTCAACCTAGTGCTCTCCGCTCC 3’ | - / 65 | -1370 | PCR of 5’ gene end |
| HaH-R40 | 5’CGGCTGATACCTCCTCCAGAGGCCACT 3’ | - / 2910 | 1476 | PCR of 5’ gene end |
| HaH-F46 | 5’GGTTGGTAATGCTGCAAGCTTTAAGCGG 3’ | AB214507/2277 | ? | Inverse PCR of 3’ region |
| HaH-R46 | 5’AACTCTGGCTGGCAACTACCATGAGACCTG 3’ | - / 2161 | ? | Inverse PCR of 3’ region |
| HaL-F21 | 5’GCCGCTACTCTAGGCTGATGACACTGG 3’ | AB354593/534 | 5 | Most of the gene amplified |
| HaL-R21 | 5’TATACAAATGCAGCGCATATCCCGC 3’ | - / 3162 | 2633 | Most of the gene amplified |
| HaL-R25 | 5’TGTCCAATACGGCTACAGAAGAAACC 3’ | - / 2375 | 1846 | 5’ genome walking |
| HaL-R32 | 5’CGGACCTTTTACGACATTTG 3’ | - / 1462 | 933 | 5’ genome walking |
| HaL-R31 | 5’CCGAACTACATCATAGACCATG 3’ | - / 648 | 119 | 5’ genome walking, |
| HaL-R24 | 5’TCTACGCCCTCGTGCACACTAC 3’ | - / 2002 | 1473 | 3’ genome walking |
| HaL-F25 | 5’AACGCAGCTGTAGTGCAAGAGCAGTC 3’ | - / 2809 | 2280 | 3’ genome walking |
| HaL-R03 | 5’GTGACAGCTGGCTACAGGAGGATATG 3’ | - / 1320 | 2591 | 3’ genome walking |
| HaL-F33 | 5’GCTCGTAGAGAGAAGTAGCAAGTTAGGA 3’ | - / 22 | -508 | Whole gene PCR |
| HaL-R33 | 5’AACTACAGCTGGAAGACCTGCTTCG 3’ | - / 3907 | 3378 | Whole gene PCR |
| RoH-F02 | 5’TGTAGCTTCTTCTGKGTGKCATCAGC 3’ | AB354689/3592 | 42 | PCR of 5’ cDNA region |
| RoH-R01 | 5’CAAGATCACCCTGACGAGGAG 3’ | - / 7205 | 3655 | PCR of 5’ cDNA region |
| oH-F21 | 5’GGGCAATTCTTCTCCTACAGTCTAGTCTGTC 3’ | - / 3580 | 30 | PCR for intron detection |
| RoH-R11 | 5’TGTGATTTACCTGCTGCTTTGTGTTG 3’ | - / 6567 | 3017 | 5’ cDNA RACE |
| RoH-F31 | 5’CAGGACCTAGTGGAAGACGTACGG 3’ | - / 3806 | 256 | Inverse PCR for 5’ end |
| RoH-R31 | 5’GAGCGAGCTATTGACGAGGATGAGATTGCC 3’ | - / 3610 | 60 | Inverse PCR for 5’ end |
| RoL-F11 | 5’ATACCGACCTGTCCAGGAGATGCGCTC 3’ | AB354690/1717 | 14 | PCR of 5’ gene end |
| RoL-R05 | 5’CAGTGACAAACTTCACTCCCTAGTTGCGCTGT 3’ | - / 8507 | 6804 | PCR of 5’ gene end |
| RoL-F02 | 5’CGCATTTCCTCGCTATGGGAGCCACT 3’ | - / 8292 | 6589 | PCR of 3’ gene end |
| RoL-R01 | 5’AGGTTAATGCTGCTGACAGTCGCCTGC 3’ | - / 11642 | 9939 | PCR of 3’ gene end |
| RoL-F18 | 5’ACTGACGGCACAGCCTCTTGGAAT 3’ | - / 7907 | 6204 | Inverse PCR, gene 5’ end |
| RoL-R15 | 5’GCACGGATTTGGGCAACCCCTC 3’ | - / 1835 | 132 | Inverse PCR, gene 5’ end |
| RoL-F05 | 5’CATTTCCATTCCAGACAGACAGACAAAG 3’ | - / 10844 | 9141 | Inverse PCR, gene 3’ end |
| RoL-R11 | 5’AACTGGTGGCTGCCATTTAGGTCTTAC 3’ | - / 9424 | 7721 | Inverse PCR, gene 3’ end |
| RoL-F20 | 5’CGCAAATCTGTAATCATTACCTCCCGACACT 3’ | - / 735 | -969 | Whole gene PCR |
| RoL-R57 | 5’TTCACAGTCTGATTATTAGAAGCGCTTGTTGTT 3’ | - / 12247 | 10544 | Whole gene PCR |

Note: GenBank reference: the accession number of the deposited sequence and the primer position in that sequence. Transcript: the primer position in relation to the transcription start

Genome Walking

Samples of 1 µg genomic DNA of H. angustipennis were digested with 30 U each Kpn1 and SacI (Takara Bio Inc.) for 2 h and further analyzed with the TOPO Walker Kit (Invitrogen). DNA digest was dephosphorylated, purified by phenol:chloroform extraction and ethanol precipitation, and used as a template for extension of the 5’ and 3’ gene regions with the Advantage 2 polymerase mix and several primers (Table 1).

Results

R. obliterata Light-Chain Fibroin cDNA and Protein

The silk gland-specific cDNA library of R. obliterata contained 114 related cDNAs (~5% of all clones), which apparently represented several alleles of one gene. A contig assembled from 32 clones was designated allele rhs004.12 and deposited in GenBank under accession number AB354590. It began with the monamer CATAGGCCG and...
H. angustipennis (Fig. 2a) yielded the sequence AIQPALIEAT, which from the silk used to glue sand grains into protective domes residues that were conserved in all L-fibroins (Fig. 1). Site and a shorter gap between the second and third Cys 16–19 residues shortly after the signal peptide cleavage were distinguished from those of Lepidoptera by a gap of 24.0%–26.8% similarities. The L-fibroins of Trichoptera Yponomeuta evonymella, Galleria mellonella, and Bombyx mori disclosed 24.0%–26.8% similarities. The L-fibroins of Trichoptera were distinguished from those of Lepidoptera by a gap of 16–19 residues shortly after the signal peptide cleavage site and a shorter gap between the second and third Cys residues that were conserved in all L-fibroins (Fig. 1). Caddisfly L-fibroins contained additional Cys residues (three in L. decipiens, four in R. obliterata, and six in H. angustipennis).

N-terminal sequencing of a 26-kDa protein extracted from the silk used to glue sand grains into protective domes (Fig. 2a) yielded the sequence AIQPALIEAT, which matched residues 20–29 of the deduced L-fibroin sequence. Analyzed protein was obviously secreted L-fibroin; the first 19 residues of L-fibroin deduced from the cDNA represented the signal peptide. A 26-kDa fraction was also detected among proteins extracted from the silk gland lumen of larvae several weeks before pupation (Fig. 2b).

The L-Fibroin Gene of R. obliterata

Two overlapping portions of the gene were amplified from the genomic DNA of a single larva designated Ro03 by PCR with the Advantage 2 polymerase mix. A 7-kb part of the 5' end was obtained with primers RoL-F11 and RoL-R05, and a 3-kb 3' end with primers RoL-F02 and RoL-R01, respectively (Table 1). Several clones were sequenced with a number of primers and the resulting continuous sequence of 9925 nt was compared with the L-fibroin cDNA. The comparison revealed that the genomic sequence began at position 13 from the transcription start, included five introns, and extended to position 388 upstream from the polyadenylation signal at the 3' end. Introns occupied most of the sequence (Fig. 3). The sequence of the coding region differed slightly from the rhsg004.12 cDNA contig (GenBank accession no. AB354590). Differences included transitions of C to T at position 288, A to C at 420, T to C at 459, C to T at 522, and C to T at 765. Transversions occurred at three other

| Residue | Ro | Ld | Ha | Ye | Gm | Mp |
|--------|----|----|----|----|----|----|
| 1 | H | L | M | C | L | M |
| 2 | L | L | L | C | L | M |
| 3 | L | L | L | C | L | L |
| 4 | L | L | L | C | L | L |
| 5 | L | L | L | C | L | L |
| 6 | L | L | L | C | L | L |
| 7 | L | L | L | C | L | L |
| 8 | L | L | L | C | L | L |
| 9 | L | L | L | C | L | L |
| 10 | L | L | L | C | L | L |
| 11 | L | L | L | C | L | L |
| 12 | L | L | L | C | L | L |
| 13 | L | L | L | C | L | L |
| 14 | L | L | L | C | L | L |
| 15 | L | L | L | C | L | L |
| 16 | L | L | L | C | L | L |
| 17 | L | L | L | C | L | L |
| 18 | L | L | L | C | L | L |
| 19 | L | L | L | C | L | L |
| 20 | L | L | L | C | L | L |
| 21 | L | L | L | C | L | L |
| 22 | L | L | L | C | L | L |
| 23 | L | L | L | C | L | L |
| 24 | L | L | L | C | L | L |
| 25 | L | L | L | C | L | L |
| 26 | L | L | L | C | L | L |
| 27 | L | L | L | C | L | L |
| 28 | L | L | L | C | L | L |
| 29 | L | L | L | C | L | L |

Fig. 1 Deduced L-fibroin sequences of the caddisflies Rhychophylca obliterata (Ro), Limnephilus decipiens (Ld), and Hydropsycha angustipennis (Ha) and the moths Yponomeuta evonymella (Ye), Galleria mellonella (Gm), and Bombyx mori (Bm). Amino acid residues are numbered from the translation start; those conserved at least in two caddisfly species are highlighted in gray. Cys residues in boldface, and those with known function in Lepidoptera are underlined. The first residue of secreted L-fibroin (after the signal peptide cleavage) is underlined with double straight lines when confirmed by protein sequencing and with a single wavy line when based on prediction. GenBank entries: Ro, AB354590; Ld, AB214510; Ha, AB214508; Ye, AB195977; Gm, S77817; Bm, X17291.
positions: replacement of T with C at position 307 changed Phe$_{103}$ to Leu, replacement of A with C at position 454 changed Asn$_{152}$ to His, and replacement of A with G at position 776 changed His$_{259}$ to Arg. The larval Ro03 obviously contained an \textit{L-fibroin} allele different from rhsg004.12.

Inverse PCR was used to verify and extend terminal gene sequences. A 2-kb product was amplified with primers RoL-R15 and RoL-F18 (Table 1) from the self-ligated \textit{Ban}II digest of the Ro03 larval DNA. The product contained a sequence extending upstream to $-1703$ from the transcription start. A 4.5-kb stretch of the 3' gene region was obtained from a DNA sample self-ligated after digestion with \textit{Xba}I and subjected to inverse PCR with primers RoL-R11 and RoL-F05. The sequence extended beyond the poly(A) tail attachment site and allowed design of primers for amplification of the whole \textit{L-fibroin} gene. A product of 11,513 nt was amplified from genomic DNA using \textit{Pfu}Ultra II polymerase with primers RoL-F20 and RoL-R57, cloned into the pCR-XL-TOPO vector, and sequenced. In combination with the overlapping products of inverse PCR the known sequence began at $-1703$ and included 13,142 nt. It was deposited in GenBank (accession no. AB354690) as \textit{Rhyacophila obliterata L-fibroin} allele 03a (abbreviated Ro03LFa).

The \textit{L-Fibroin} Gene of \textit{H. angustipennis}

A considerable part of the gene was amplified from genomic DNA of a larva designated Ha23 by PCR with KOD FX polymerase. Primers HaL-F21 and HaL-R21 (Table 1), which were deduced from the cDNA analyzed earlier (Yonemura et al. 2006), amplified a 2.6-kb product that was cloned into the pCR4-TOPO vector and sequenced. As expected from the primer positions, the sequence began at $+5$ from the transcription start and ended 110 nt upstream from the first polyadenylation signal.

Fig. 2 \textit{Silk proteins of Rhyacophila obliterata} separated by polyacrylamide electrophoresis from silk fibers used to construct shelters for pupation (a) and from dope collected from the silk gland lumen (b). The arrow in A indicates the fraction proven to be \textit{L-fibroin}. Note that proteins of similar size occur also in the silk dope.

Fig. 3 \textit{L-Fibroin} gene structure in \textit{H. angustipennis} (GenBank accession no. AB354593), \textit{R. obliterata} (AB354690), and \textit{B. mori} (M76430); exons are shown as boxes, and introns as lines. Numbering of sequences begins at the transcription start. Sequences deposited in GenBank include longer upstream regions than shown here. Positions of TATA boxes, translation initiation codons, termination codons, and polyadenylation signals are indicated.
signal. Comparison with the cDNA (Yonemura et al. 2006) disclosed the positions of five introns (Fig. 3).

The 5' and 3' parts of the gene were identified with the aid of the TOPO walker kit. Products obtained with primers HaL-R25 and HaL-F24, respectively (Table 1), were ligated and taken for PCR with the LinkAmp primer 1 paired with primer HaL-R32 for 5' and with HaL-F25 for 3' analyses. Subsequent nested PCR with primers HaL-R32 and HaL-R31 amplified 700 bp of the upstream region. Three products, of 300, 900, and 1100 bp, of the 3' end were amplified with primers HaL-F25 and HaL-F03. Sequences of the 700- and 1100-bp products permitted design of a primer pair HaL-F33 and HaL-R33 (Table 1) that amplified a product of 3886 nt extending from position –529 to position +3378 of the gene. A contig of 4229 nt, which was based on independent sequencing of the 5' region (0.7 kb), a major part of the gene (3.8 kb), and the 3’ region (1.1 kb), was deposited in GenBank as allele 23 of the H. angustipennis L-fibroin gene (Ha23LF) (accession no. AB354593).

Heavy-Chain Fibroin cDNA in R. obliterata

Thirty clones identified in the R. obliterata cDNA library were of different lengths but the identity of their 3’ end indicated that they were derived from a single gene. The sequence of 2974 nt based on 17 clones encoded 908 amino acid residues of a truncated protein that was homologous to the previously described H-fibroins of caddisflies (Yonemura et al. 2006). The cDNA sequence apparently corresponded to the 3' end of the R. obliterata H-fibroin gene. It was deposited in GenBank as allele RoHF3’a (accession no. AB354588). Most of the sequence consisted of highly conserved blocks of 408 nt. The first one was incomplete and included only 373 nt; the complete blocks began at positions 374, 782, 1190, 1598, and 2006, respectively. They differed by only four transitional point mutations. The regularity of repeats dissipated at nt 2413, which was 315 nt upstream from the stop codon. The polyadenylation signal was located 227 nt after the stop codon and a poly(A) tail began 25 nt farther downstream.

All H-fibroin cDNAs detected in our library were truncated at the 5'end. PCR search for the 5' region was launched with the degenerate primer RoH-F02 (degeneracy index = 384), which corresponded to nt 42–67 (counted from the transcription start) of H. angustipennis H-fibroin cDNA (Yonemura et al. 2006), and the specific reverse primer RoH-R01 (Table 1), which matched repetitive sequence close to the 5' end of the RoHF3’a fragment. A 1-kb PCR product contained a continuous open reading frame. Reverse primer RoH-R11, derived from the 5' end of this product, was used for 5'RACE PCR. The established sequence apparently began with the transcription start that was identified in one clone as the first, and in two clones as the second, nucleotide of the consensus CATCAGTCA. Contig RoHF5’, which combined results of the initial PCR with those of 5' RACE, included 1010 nt and was deposited in GenBank under accession no. AB354587. At its 3’ end, the contig contained one full and one partial repeat of 408 nt.

Deduced H-Fibroin Protein in R. obliterata

Putative translation product of the RoHF5’ fragment contained a predicted signal peptide of 20 amino acid residues followed by 460 residues of the secreted H-fibroin. In the optimized alignment of the nonrepetitive N-termini, the sequences of R. obliterata and H. angustipennis H-fibroins contained 56% residues in comparable positions (Fig. 4a). The similarity to lepidopteran H-fibroins was rather low; only 20% of amino acid positions were conserved in the nonrepetitive regions of the R. obliterata and G. mellonella H-fibroins. However, the spacing of certain residues was retained and all H-fibroins contained in their N-terminus a motif rich in Lys and Glu in combination with the nonpolar residues Val and Ile (residues 121–129 in R. obliterata H-fibroin). H-fibroin similarities between compared species ended shortly after this motif (Fig. 4a).

Beginning by residue 159, the N-terminal sequence deduced from RoHF5’ became very similar to the repeats encoded by most of the RoHF3’a fragment (Fig. 4b). The N-terminal sequence ended and the C-terminal one began with highly conserved repetitive domains of 136 amino acid residues (corresponding to the blocks of 408 nt). The unidentified central part of the H-fibroin most likely consisted of such domains. In Fig. 4, each domain was divided into repeats RD, RF, and RE to accentuate similarities to the LD, LE, and LF repeats previously identified in the
H-fibroin of *L. decipiens* and the HD repeat found in *H. angustipennis* (Yonemura et al. 2006). The repeats were composed of a few distinct motifs. A stretch of residues of different properties, which was flanked on both sides by dodecamers composed of hydrophobic and charged residues. The terminal motif GKKHG of this repeat was similar to the GRRGW of the RE repeats.

The repetitive region of *R. obliterata* H-fibroin ended with a truncated and degenerate repeat RF1 at –104 to –77 (Fig. 4b). The nonrepetitive stretch comprising 78 C-terminal residues with known function in Lepidoptera are in boldface. Dashed-underlined stretches represent terminal parts of the central repetitive regions (the species differ greatly in the length of the nonrepetitive C-termini).
The **H-Fibroin Gene of R. obliterata**

The *H-fibroin* gene of Lepidoptera contains a large intron not far from the start of the coding sequence. The existence of a similar arrangement in the *H-fibroin* of caddisflies was probed by PCR based on primers from the nonrepetitive 5′ cDNA region. A 3-kb fragment was amplified from the genomic DNA of *R. obliterata* larva designated Ro01 with primers RoH-F21 and RoH-R11 (Table 1). The sequence of this fragment revealed overlap with the cDNA sequence and disclosed the presence of an intron 2645 bp long (Fig. 5a). To determine the upstream gene region, the Ro01 DNA was digested with a mix of *Eco*RI and *Mun*I, self-ligated, and used for inverse PCR with primers RoH-R31 and RoH-F31 (Table 1). A 3.5-kb PCR product extended from a *Mun*I restriction site in the intron (+425 from the transcription start) to a *Mun*I site at −3550. A combination of this sequence with that of the 3′-terminal 5-kb PCR product yielded a contig that spanned from −3550 to +3017 and, in the regions 1–42 (first exon) and 2713–3655 (part of the second exon), matched precisely the cDNA sequence established by 5′ RACE. Since the cDNA contained an additional 638 nt, the 5′ part of the *H-fibroin* gene (*Ro03HFS*) deposited in GenBank under accession no. AB354689 covers the region from −3550 to +3655. The 3′ end of the *R. obliterata* *H-fibroin* gene (Fig. 5b) was analyzed only at the cDNA level.

### Table 1

| Species     | GenBank Accession | Sequence Length |
|-------------|-------------------|-----------------|
| R. obliterata| AB354689          | 2645 bp         |

The **H-Fibroin Gene of H. angustipennis**

Analysis of the *H-fibroin* gene in the *H. angustipennis* larva Ha23 by means of PCR with the cDNA-derived primers HaH-F42 and HaH-R41 (Table 1) yielded a 0.5-kb genomic product. Primers HaH-R51 and HaH-F51 derived from this product were used for inverse PCR with the DNA fragments that had been self-ligated after digestion with the *Hae*II restrictase. A product of 1.5 kb extended the known *H-fibroin* sequence in both directions. Another product of 2.8 kb was obtained from the Ha23 genomic DNA by PCR employing primers HaH-F40 and HaH-R40 (Table 1). A contig combining sequences of the 0.5-, 1.5-, and 2.8-kb products spanned from −1434 to +1476, with a 77-bp intron beginning at +67. The sequence was registered in GenBank as *Ha23HFS* (accession no. AB354591). The region from about −60 to +70 (end of the first exon) exhibited a high sequence similarity to the *H-fibroin* of *R. obliterata* and, to lesser extent, to that of the moths (Fig. 5a). The distances between the TATA box, transcription start, translation initiation, and the end of the first exon were remarkably similar in all species.

The 3′ region of the *H-fibroin* gene was amplified with the Advantage 2 polymerase mix from the Ha23 genomic DNA self-ligated after *Ban*II digestion. A 3-kb fragment was obtained in a PCR with primers HaH-R46 and HaH-F46. It was cloned and sequenced. The sequence matched

### Fig. 5

Fig. 5 Similarities in the *H-fibroin* gene among Trichoptera and Lepidoptera (nucleotides conserved in at least two caddisfly species are highlighted in gray); no sequence homologies were detected upstream or downstream from the depicted regions. Exons are in capital letters. a The *H-fibroin* 5′ region in *R. obliterata* (Ro; GenBank accession no. AB354689), *H. angustipennis* (Ha; AB354591), *G. mellonella* (Gm; AF095239), and *B. mori* (Bm; AF226688). Sequences are numbered from the transcription start, the TATA box is underlined with a straight line, the initiator consensus sequence (AyCAgyyy) with a wavy line, and the translation initiation codon is double-underlined. Intron donor (gtaagt) and acceptor (gtcttatttttag) sites are underlined in the Ha sequence. b Alignment of the *H-fibroin* 3′ region in *R. obliterata* (Ro; GenBank accession no. AB354588), *H. angustipennis* (Ha; AB214507), *L. decipiens* (Ld; AB214509), *Y. evonymella* (Ye; AB195978), *G. mellonella* (Gm; AF095240), and *B. mori* (Bm; AF2266880). The termination codon is in boldface.
the previously identified cDNA (Yonemura et al. 2006) but continued with 1819 nt beyond its poly(A) attachment site. The genomic sequence representing the 3’ end of H-fibroin allele 23 (Ha23HF3) was assigned GenBank accession no. AB354592. The region with similarities to the 3’end of the H-fibroin gene of caddisflies and a few moths is shown in Fig. 5b.

Discussion

Conservation of the L-Fibroin and H-Fibroin Silk Genes in Trichoptera and Lepidoptera

Analysis of the silk gland-specific cDNAs in R. obliterata complements earlier investigations in H. angustipennis and L. decipiens (Yonemura et al. 2006) and demonstrates that transcripts homologous to the lepidopteran L-fibroin and H-fibroin genes occur in all three suborders of Trichoptera. This paper shows that the organization of the L-fibroin and H-fibroin genes in R. obliterata and H. angustipennis is similar to that of the corresponding lepidopteran genes. Finally, the expression of both genes is restricted to the posterior silk gland section (Yonemura et al. 2006) like in Lepidoptera. These data suggest that ancestral H-fibroin and L-fibroin genes evolved about 250 mya (million years ago), prior to the separation of Trichoptera and Lepidoptera, and were retained in both orders. Fossil Trichoptera date back to ~235 mya (Sukacheva 1968, 1973), and fossil Lepidoptera to ~203 mya (Whalley 1985).

B. mori is the only lepidopteran in which the L-fibroin gene was analyzed (GenBank accession no. M76430) and found to contain six introns (Fig. 3). The L-fibroin gene of caddisflies R. obliterata and H. angustipennis (GenBank accession nos. AB354690 and AB354593, respectively) contains five introns that occur at positions comparable to those of introns 1–4 and intron 6 of B. mori, indicating that the fifth intron of B. mori was inserted into the gene after the moths had split from the caddisflies. The total length of introns is about 12,000 bp in B. mori, 9000 in R. obliterata, and <1500 in H. angustipennis. The first intron is much longer than any other and the fourth is the shortest in all species. Donor sequences at the exon/intron boundaries are close to the AGGTRAGT motif (exon sequence underlined), while the acceptor motif YYYYYYYYNCAG is in some cases modified to YYYYYYYYNTAG. Sequences TTAATCT, TTAATCT, TTGCTT, and TTGATG within the introns can be regarded as intron branch points.

The TATA box was identified in the L-fibroin gene of H. angustipennis at position –29 relative to the transcription start, similarly to B. mori (–31). R. obliterata contains a TATA box at –286 followed by a putative transcription initiation sequence at –255, but no corresponding transcript was detected. The first Met codon ATG follows 29 nt after the transcription start in H. angustipennis, 27 nt in R. obliterata, and 42 nt in B. mori. The lengths of exons are alike in all species except for an insertion of 71 bp into the second exon and a split of the original fifth exon into two in B. mori and a terminal extension of exon 6 in H. angustipennis. The sequence of the first exon is about 57% identical in the caddisfly species and different from that in B. mori. The last exon in all species contains a putative polyadenylation signal within 20 bp after the termination codon (not shown in Fig. 3) but signals in more distant positions are obviously used for the poly(A) attachment (Fig. 3). The L-fibroin gene of H. angustipennis harbors three putative signals in the region 2450–2530 (only one of them is indicated in Fig. 3) and the poly(A) chain is attached at 2617. Additional polyadenylation signals occur at 2946 and 2998 and their use is indicated by the finding of cDNAs with a poly(A) tail attached at 2974 and 3032, respectively.

In the 5’ region the H-fibroin gene of R. obliterata and H. angustipennis includes (GenBank accession nos. AB354689 and AB354591, respectively) a short first and a long second exon, similar to the H-fibroin gene of the moths, for example, G. mellonella and B. mori (GenBank accession nos. AF095239 and AF226688). A low but distinct sequence similarity between the H-fibroins of caddisflies and moths begins in the 5’ region about 60 nt upstream from the transcription start and increases around the TATA box at –31 in R. obliterata and at –30 in the other species (Fig. 5a). Maximal similarity is found in the first exon, which is nearly identical in the moths analyzed (Yonemura and Sehnal 2006). The nucleotide sequence (but not the encoded peptide) of this exon is more diversified among the caddisflies; some regions of the first exon in H. angustipennis resemble the H-fibroin genes of moths more than that of R. obliterata. The nucleotide sequence similarity among and between Trichoptera and Lepidoptera is retained in the exon/intron boundary but rapidly dissipates in the intron. The donor and acceptor sites are conserved and close to the consensus sequences.

Most of the H-fibroin second exon is occupied by species-specific repetitive blocks in both Lepidoptera (e.g., Mita et al. 1994; Sezutsu and Yuukihiro 2000; Zhou et al. 2001; Žurovec and Sehnal 2002; Fedić et al. 2003; Yonemura and Sehnal 2006) and Trichoptera (Yonemura et al. 2006; present data). The blocks of 408 nt found in R. obliterata are exceptionally uniform. At the end of the coding regions the blocks are replaced by nonrepetitive sequences that exhibit some similarities among the caddisflies and, to a lesser extent, also in the moths (Fig. 5b). The untranslated 3’ terminus contains a single polyadenylation signal 143 nt after the termination codon in L. decipiens and 226 nt after in R. obliterata; the H-fibroin
The linkage of L-fibroin and H-fibroin proteins by a disulfide bridge between Cys22 of the H-fibroin (Tanaka et al. 1999b) and Cys170 of the L-fibroin (Yamaguchi et al. 1989) proved indispensable for the secretion of both components in B. mori (Takei et al. 1987). Silk filament formation in most Lepidoptera further involves a P25 glycoprotein. For B. mori it was shown that noncovalent interaction of P25 with the H-fibroin N-terminus is important for gel/filament processing of the hydrophobic H-fibroin/L-fibroin dimers (Tanaka et al. 1999a; Inoue et al. 2000), while L-fibroin protects attachment of the sugar moieties that render P25 hydrophilic (Inoue et al. 2004). Our failure to detect any sequence homologous to P25 among 2304 silk gland-specific cDNAs in R. obliterata and the negative results of our previous search in H. angustipennis and L. decipiens (Yonemura et al. 2006) suggest strongly that this type of protein is lacking in caddisfly silk.

The conservation of certain amino acid motifs in the H-fibroin N-terminus in both moths and caddisflies (Fig. 4a) indicates that the role of this region in fibroin processing is not limited to the interaction with P25. The similarity of signal peptide sequences is obviously related to H-fibroin transport into the endoplasmatic reticulum. We cannot appreciate functions of the motifs like IYEEEIIK in R. obliterata (residues 121–129; Fig. 5a) or somewhat similar sequences of the alternating charged and hydrophobic residues (e.g., KGEIIEKII at positions 62–70 in R. obliterata), which occur with slight modifications in all H-fibroins identified so far (Fig. 4a).

The nonrepetitive H-fibroin C-terminus also contains similar features across Trichoptera and Lepidoptera (Fig. 4b). In the caddisfly H-fibroins, a region of about 75 amino acid residues contains relatively high representations of Gly and Pro and a number of single residues or their small groupings, such as GRR, SVYT, HPG, VKVGN, and NGNC, at conserved positions. The spacings of Cys1, Cys22, and a few other residues in the C-terminus are similar to those in the lepidopteran H-fibroins. Cys22 may participate in H-fibroin linkage with the L-fibroin but the absence of Cys4 precludes the formation of the intramolecular disulfide bridge Cys1/Cys4 that is known in B. mori.

The L-fibroins of Trichoptera and Lepidoptera exhibit a similar distribution of many residues with characteristic properties such as size, hydrophobicity, and charge (Fig. 1), including three Cys residues whose significance for L-fibroin function was demonstrated in B. mori (Tanaka et al. 1999b). Specific features of caddisfly L-fibroins include the conservation of regions with alternating groups of hydrophobic and hydrophilic residues, the presence of three to six additional Cys residues (one of them, Cys129 in R. obliterata, is conserved in all three caddisfly species), and the occurrence of short and rather unusual motifs such as ProTrpPro (residues 134–136 in R. obliterata). The features unique to caddisflies might be important for L-fibroin/H-fibroin interaction in the absence of P25.

Filament Polymerization

The total amino acid composition of the repetitive H-fibroin region is similar in all examined caddisflies and distinct from the known lepidopteran silks. In R. obliterata the region contains 28.2% Gly, 15.9% Ser, 9.7% Leu, 8.3% Ile, 6.4% Val, 8.3% Arg, 6.3% His, and 3.2% Pro (Fig. 4). High representation of Gly and Ser, the bulky hydrophobic residues (Leu, Ile, Val), charged residues (Arg, His, Lys, Asp, and Glu), and Pro was found also in H. angustipennis and L. decipiens (Yonemura et al. 2006). Another characteristic feature of caddisflies is the low amount of Ala (1.6% in R. obliterata), in contrast to its high content in the H-fibroin repeats of Lepidoptera (Sehnal and Sutherland 2008).

Filament formation depends on weak molecular interactions between motifs that are arranged in higher-order repeats (Lucas and Rudall 1968). Comparison of reiterated sequences is the first step to understanding the evolution and mechanisms of filament polymerization. H-Fibroins of all three examined caddisfly species contain similar motifs, but their relative representation, their assembly into repeats, and the regularity of repeats are different. To facilitate their comparison, we designate each repeat with two letters: the first is derived from the family name (Hydropsychidae, Limnephilidae, and Rhyacophilidae, respectively), and the second specifies the type of repeat based on the presence of distinct amino acid motifs and the repeat length. The A, B, and C repeat types were recognized in H. angustipennis, the E and F types in L. decipiens and R. obliterata, and the D type in all three species (Fig. 6). The HA, HC, HD, LD, LE, LF, RE, and RF repeats contain a similar stretch of about 11 amino acid residues with a central Trp. Another conserved region of 27–31 residues occurs in the D-type repeats and includes a central amphiphilic region sandwiched between two Ser-rich motifs. It is likely that both conserved regions, which are absent in the H-fibroins of Lepidoptera, evolved prior to the caddisfly separation into three suborders.

Molecular conformation of the repeats determines physical properties of the silk filament. Conformation of some motifs has been demonstrated or proposed on
theoretical grounds. For example, the Ser-rich motifs and the strings of \((SX)_n\) in repeats of types A, C, D, E, and F are likely to form rather rigid \(\beta\)-sheets due to hydrogen bonding via polar zipper interactions (Bini et al. 2004). The GGL motif reiterated in the E repeats provides a 3(1) helix conformation, as shown for LGG triplets (Ashida et al. 2003). The GPGXX motif, which occurs in the HA and RE repeats, is believed to form a \(\beta\)-spiral conferring elasticity to the protein polymer (Hayashi and Lewis 1998). Physical properties of the filament are further affected by repeat regularity that ensures precise registration of the interacting motifs (Sehnal and Žurovec 2004). The length of repeats varies in the H-fibroins of \(H.\) angustipennis and \(L.\) decipiens but the repeats of \(R.\) obliterata form very precise modules of 135 residues.

Bini et al. (2004) suggested that proteins forming water-resistant silk filament must include hydrophilic terminal domains flanking a very long central portion constructed from the hydrophobic blocks alternating with short hydrophilic regions. The strong predominance of hydrophobicity in the major polymerizing part was regarded as essential for water exclusion during \(\beta\)-crystallite formation, while the hydrophilic terminal regions allowed fibroin hydration during storage in the silk gland lumen. These rules may apply to aerial silks spun by caterpillars or spiders but are contradicted by the content and distribution of hydrophilic amino acid residues in the repetitive domain of caddisfly H-fibroins. The amphiphilic plot of \(R.\) obliterata H-fibroin reveals a predominance of hydrophilic regions (Fig. 7), very similar to the situation described for \(H.\) angustipennis and \(L.\) decipiens (Yonemura et al. 2006) and different from that in most Lepidoptera (Sehnal and Žurovec 2004).

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