Proteome of Hydra Nematocyst*

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Prakash G. Balasubramanian 1, Anna Beckmann 2, Uwe Warnken 3, Martina Schnölder 2, Andreas Schüler 4, Erich Bornberg-Bauer 5, Thomas W. Holstein 6, 7, and Suat Özbek 1, 2

From the 1 Department of Molecular Evolution and Genomics, Centre for Organismal Studies, University of Heidelberg and 5 Functional Proteome Analysis, German Cancer Research Center (DKFZ), 69120 Heidelberg and the 7 Institute for Evolution and Biodiversity, University of Münster, D48149 Münster, Germany

Background: Nematocysts are the most sophisticated organelles in the animal kingdom and a hallmark of the phylum Cnidaria.

Results: We present the first complete protein map of the Hydra nematocyst.

Conclusion: The nematocyst proteome is highly complex and includes novel structural proteins.

Significance: The proteome data reveal a eukaryotic origin of the nematocyst and point to common molecular features with the extracellular matrix.

Stinging cells or nematocytes of jellyfish and other cnidarians represent one of the most poisonous and sophisticated cellular inventions in animal evolution. This ancient cell type is unique in containing a giant secretory vesicle derived from the Golgi apparatus. The organelle structure within the vesicle comprises an elastically stretched capsule (nematocyst) to which a long tubule is attached. During exocytosis, the barbed part of the cuticle structures (8–10). The molecular composition of the nematocyst, only a small fraction has hitherto been defined. These include mainly structural proteins like the large family of mini-collagens (13–15), the C-type lectin NOWA (16), and spinalin, which is part of the tubule spines (17). Here, we present a proteome analysis of Hydra nematocysts, showing an unusual composition of venomous and structural proteins forming one of the most sophisticated organelles in the animal kingdom.

EXPERIMENTAL PROCEDURES

Venoms are produced in probably all animal phyla (2, 3) and represent a mixture of effector proteins often aimed at tissue integrity by targeting extracellular matrix (ECM) 3 and receptor molecules. Intriguingly, many toxins share close structural and functional properties with ECM proteins, indicating a common evolutionary origin or co-evolution (4). It was hitherto unknown whether this process also contributed to ECM formation in providing efficient organ or organelle structures for toxin delivery.

Nematocysts or cnidae are highly complex projectile organelles used for capture of prey and defense in all cnidarians (5, 6). They are produced in specialized stem cell derivatives called nematocytes by a continuous secretory process of proteins into a giant post-Golgi vesicle. Nematocysts are composed of a cylindrical capsule body to which an extended tubule is attached that is often armed with spines. The capsule morphology varies among different cnidarian species, with a general tendency toward higher complexity in medusozoans compared with anthozoans (7). In Hydra, four types of nematocysts are present: the small desmonemes, with a tightly coiled tubule used for prey attachment; the holotrichous and spineless atrichous isorhizas; and the large stenoteles, with a prominent stylet apparatus at the tubule base employed for piercing solid cuticle structures (8–10). The molecular composition of the capsule comprises a highly dense collagenous polymer charged with an osmotic pressure of 150 bars (11, 12). Nematocyst discharge, triggered by the mechanosensory cnidocil apparatus, is one of the fastest processes in biology and creates an impact on the harpoon-like expelled tubule comparable with technical bullets (1). Of the protein components constituting the nematocyst, only a small fraction has been hitherto defined. These include mainly structural proteins like the large family of mini-collagens (13–15), the C-type lectin NOWA (16), and spinalin, which is part of the tubule spines (17).

Animals—Hydra magnipapillata was used for all experiments. Animals were cultured in Hydra medium at 18 °C and fed two to three times a week with freshly hatched Artemia salina nauplii. Animals used for the experiments were starved
for 24 h. Intact nematocysts were isolated from whole Hydra tissue as described (18).

One-dimensional Electrophoresis and Orbitrap™ MS—Nematocyst proteins were separated on a one-dimensional NuPAGE 4–12% BisTris gel (pH 6.4) (Invitrogen) and stained with colloidal Coomassie Blue. The resulting gel was cut into 27 pieces. In-gel trypic digestion and extraction were performed as described previously (19). Peptide separation was achieved using a nano-ACQUITY UPLC® system (Waters). The nano-UPLC system was coupled on-line to an LTQ Orbitrap XL mass spectrometer (Thermo Fisher). Data-dependent acquisition using Xcalibur 2.0.6 (Thermo Fisher) was performed by one Fourier transform MS scan with a resolution of 60,000 and a m/z range from 370 to 2000 in parallel with six MS/MS scans of the most intense precursor ions in the ion trap. The mgf files were used for database searches with the Mascot search engine (Matrix Science, London, United Kingdom) against Hydra Hma2 protein models (20).

Two-dimensional Electrophoresis and Electrospray Ionization Quadrupole TOF-MS—Protein separation was performed with the PROTEAN system (Bio-Rad). The protein samples were applied to PROTEAN ReadyStrip immobilized pH gradient (IPG) 17-cm pH 3.0–10.0 strips (Bio-Rad). Isolated caps were applied to PROTEAN ReadyStrip immobilized pH gradient Quadrupole TOF-MS using the PROTEAN system (Bio-Rad). The protein samples were run on the gel in 30 mM Tris-HCl (pH 8.8) and 0.2% SDS at a constant current mode of 17 mA/gel for 12 h with running buffer (2.5 mM Tris-HCl (pH 8.3), 19.2 mM glycine, and 0.01% SDS) in a Bio-Rad PROTEAN II xi cell system. To visualize the protein spots on the gel, the colloidal Coomassie Blue staining procedure was employed. From the two fractions together, 186 spots were visualized, excised, and digested with trypsin as described (19). Nanoscale LC-electrospray ionization MS/MS analysis of the protein extract was performed using the CapLC capillary LC system (Waters), which was coupled to a Q-Tof Ultima mass spectrometer. Data acquisition was controlled by MassLynx™ system (Waters). The nano-UPLC system was coupled on-line to an LTQ Orbitrap XL mass spectrometer (Thermo Fisher). Data-dependent acquisition using Xcalibur 2.0.6 (Thermo Fisher) was performed by one Fourier transform MS scan with a resolution of 60,000 and a m/z range from 370 to 2000 in parallel with six MS/MS scans of the most intense precursor ions in the ion trap. The mgf files were used for database searches with the Mascot search engine (Matrix Science, London, United Kingdom) against Hydra Hma2 protein models (20).

Proteome Annotation—The annotation of the nematocyst proteome was performed manually for all protein sequences using the EMADT (extensive manual annotation(s) by domain(s) and taxonomy) protocol developed heuristically for this analysis. The protein sequences were analyzed by PSI-BLAST (21) with default algorithm parameters. The E value cutoff was set at 1e−05. The protein domains including the number of repeats identified by the Conserved Domains Database were noted. Domains were also analyzed in parallel using InterProScan, and signal peptide prediction was performed using the SignalP 3.0 server. The redundancy of identified domains was checked, and the annotation was manually curated if necessary. Multidomain proteins were analyzed for every single domain in the sequence. Functional organization of the proteome was done based on the domains identified.

Comparative Genomics—Predicted peptide data sets were retrieved from the DOE Joint Genome Institute (www.jgi.doe.gov/); Nematostella vectensis, Monosiga brevicollis, and Trichoplax adhaerens), from NCBI (ftp.ncbi.nih.gov/genomes/; Strongylocentrotus purpuratus), and from the Ensembl Database (Branchiostoma floridae, Caenorhabditis elegans, Drosophila melanogaster, and Homo sapiens) as of June 10, 2010. BLASTp from the NCBI blastall package (21) was used to scan all nematocyst proteins against all reference proteomes mentioned above for the identification of orthologs. BLASTp was run with an E value cutoff of 10−5 and the parameters −F"m S" and −sT for soft filtering of low complexity regions and locally optimal Smith-Waterman alignments, respectively (parameter −F"m S" refers to soft filtering of low complexity regions, and parameter −sT refers to locally optimal Smith-Waterman alignments). The BLAST output was further filtered to include only hits, which correspond to query sequence coverage of at least 30%. Protein domains were annotated based on InterPro (22) domain definitions. Overlapping domain annotations were resolved by removing the annotation with the less significant E value. Putative nematocyst venom proteins were identified based on similarity to known venom proteins stored in the ToxProt Database (23). A BLAST search of all nematocyst proteins versus all sequences in the ToxProt Database yielded 54 putative nematocyst venom proteins (E < 10−5).

Analysis of Orphan Proteins—The 117 nematocyst orphan proteins (without hits in the reference genomes and without InterPro domains) were clustered based on sequence similarity using BLASTclust (21). Proteins with 25% sequence identity in a region spanning at least 35% of the sequence length were clustered together. This procedure yielded two clusters of sizes 10 and 9, respectively. Both clusters were aligned using T-Coffee (24), and a hidden Markov model based on those alignments was scanned against the NCBI non-redundant protein database using HMMER. This scan yielded significant hits in proteins containing the sweet tooth domain for the cluster of size 3. The three proteins in the nematocyst-specific cluster were predicted to have nucleotide-binding activity based on an analysis with the I-TASSER server (25). Furthermore, the conserved region in the three proteins of the Hydra-specific cluster was predicted to be globular (instead of disordered or fibrous) (26).

Cloning and Recombinant Expression of Cnidoin—Preparation of whole RNA was performed using an RNeasy mini kit (Qiagen) according to the manufacturer’s instructions. The isolated RNA was transcribed by reverse transcriptase into cDNA. PCR for the Cnidoin coding sequence was performed using primers ATGTCTCGATTACTACTTC (forward) and TTATCCTTTTTACAAAGCTCC (reverse). The purified PCR product was ligated into the pGEM-T vector (Promega).
and confirmed by sequencing. Recombinant expression in Escherichia coli BL21(DE3) cells was performed with a pET-21b vector (Novagen), which introduces a C-terminal polyhistidine tag. Cnidoin protein was exclusively found in inclusion bodies and purified under denaturing conditions (8 M urea) using nickel-nitrilotriacetic acid beads.

Immunostaining—H. magnipapillata animals were relaxed in 2% urethane in Hydra medium and then fixed in freshly prepared 4% paraformaldehyde in Hydra medium for 30 min. The fixative was removed by three 10-min washing steps with PBS and 0.1% Triton X-100. The antibody was diluted 1:250 in PBS and 1% BSA and incubated overnight at 4 °C. In the case of co-staining with two antibodies, both antibodies (anti-minicollagen-1, 1:500 dilution) were incubated simultaneously overnight at 4 °C. To remove unbound antibodies, three 10-min washing steps with PBS and 0.1% Triton X-100 were performed. Incubation with the secondary antibodies was performed for 2 h at room temperature. For detection of Cnidoin, an Alexa 568-conjugated goat anti-guinea pig antibody was used; for detection of minicollagen-1, an Alexa 488-conjugated goat anti-rabbit antibody was used. The secondary antibodies were diluted 1:400 in PBS and 1% BSA. To remove unbound antibodies, the animals were washed three times with PBS and then mounted on object slides with PBS and 90% glycerol.

Microscopy—Fluorescent images were captured with a Nikon A1R confocal laser scanning microscope at the Nikon Imaging Center at the University of Heidelberg. In situ images were captured with a Nikon Eclipse 80i microscope.

RESULTS

Global Annotation of Nematocyst Proteome—We isolated intact undischarged nematocysts (stenoteles, desmonemes, and atrichous and holotrichous isorhizas) (14) from H. magnipapillata by density gradient centrifugation (18). The purity of the preparation was confirmed by light and electron microscopy (Fig. 1, C and D). The nematocyst preparation was solubilized by DTT/SDS treatment and separated by one-dimensional gel electrophoresis. Gel slices (n = 27) covering the whole molecular mass range (Fig. 1E) were subjected to Orbitrap MS analysis. Fig. 1F shows that the nematocyst proteins exhibited a Gaussian distribution regarding their molecular masses, with the majority ranging between 10 and 40 kDa. One-dimensional gels guaranteed higher protein yields than two-dimensional gels, which were analyzed in parallel (supplemental Fig. 1). Annotation of the obtained nematocyst protein sequences was performed by a manual procedure applying PSI-BLAST and InterProScan analyses in parallel (EMADT; see “Experimental Procedures” for details). By this approach and the application of a MASCOT protein score threshold of 50 to exclude insignificant hits, we identified a total number of 410 unique protein sequences (supplemental Table 1). An additional 25 sequences from the full proteome analysis were categorized as contaminants and included histones, ribosomal proteins, cytoskeletal components, and cytoplasmic enzymes. The fraction of cellular contaminants in the complete proteome was low as estimated from the intensity of the actin spot in the two-dimensional resolution of the nematocyst sample.

FIGURE 1. Nematocyst morphology and protein spectrum. A, H. magnipapillata. B, close-up of undischarged nematocysts assembled in battery cells of the tentacles. C, light microscopic view of isolated nematocyst capsules. D, scanning electron microscopic image of isolated nematocyst. E, one-dimensional SDS-PAGE of a nematocyst sample. The gel slices used for mass spectrometry are indicated. F, molecular mass distribution of the nematocyst proteome. Scale bars = 0.5 mm (A), 15 μm (B and C), and 2 μm (D).
A comparison with homologous sequences in the genome of the starlet sea anemone *N. vectensis* showed that, in total, 44% of the *Hydra* nematocyst proteins were represented here (Fig. 3A). Although most of the metabolic proteins are conserved between both species, the number of novel and structural proteins is markedly pronounced in *Hydra*, which might indicate a large diversity between nematocyst proteins of different taxa or a higher morphological complexity of hydrozoan nematocysts (7). We categorized the novel proteins into four groups: sequences 1) unique to the proteome of *Hydra* nematocysts, 2) unique to *Hydra*, 3) unique to *Hydra* and *Nematostella*, and 4) not exclusive to cnidarians (supplemental Table 1).

Fig. 3B extends this comparison and shows the evolutionary distribution of the proteome among metazoans and eukaryotes. Overall, *Hydra* nematocysts comprise a gene repertoire of which 5% are cnidarian-specific with homologs exclusively in *Nematostella*, and 18% have homologs in Eumetazoa and 31% in Eukaryota. An additional 18% have no detectable homologs in other proteomes but contain InterPro domains. 28% of the sequences were classified as orphan genes. Among the proteins with detectable InterPro domains, several structural and enzymatic domains, e.g. α-collagen, AbfB, and von Willebrand factor, are strongly overrepresented (Fig. 3C). Among the remaining orphan proteins, several can be grouped together based on conserved common motifs (see “Experimental Procedures”). Ten proteins in the largest cluster contain the “sweet tooth” domain, which was reported to be among the most abundant extracellular protein domains in the *Hydra* genome but is not integrated in InterPro yet (30). The three orphan proteins of the second largest cluster share very high sequence identity (>80%) and are predicted to be globular and to have a nucleotide-binding propensity (see “Experimental Procedures”). A hidden Markov model created from their alignment (Fig. 3D) was scanned against the NCBI non-redundant database but returned no significant hit. Consequently, this protein structure seems to be unique for the proteome of the *Hydra* nematocyst and deserves further functional analysis.

Potential horizontal gene transfer events of bacterial origin comprise three proteins involved in poly-γ-glutamate biosynthesis: CapA, its isoform CapA2, and Hma2.206651, a protein with predicted enzymatic function. Poly-γ-glutamate biosynthesis is a crucial feature of nematocytes, and so far, cnidarians represent the only metazoan organisms with this capacity (29). Recently, Denker *et al.* (30) characterized CapA as a nematocyte-specific gene of defined bacterial origin, but the low percentage of horizontal gene transfer candidates in our analysis rejects a bacterial origin of the nematocyst.

**Venoms and Enzymes Represent Conserved Feature of Nematocysts**—The primary function of nematocysts is the release of venoms into the tissue of a prey organism or attacking predator. In *Hydra* and other cnidarians, polypeptide venoms are expressed mainly in nematocytes and represent a highly complex array of effector molecules aimed at paralyzing a prey and disintegrating its tissue. Our proteome analysis revealed 55 toxin-related sequences, mostly with homologous toxin proteins in diverse phyla (see “Experimental Procedures”). These
include neurotoxins, cytolsins, toxic phospholipases, many peptidases, and proteins of the SCP_GAPR-1-like family (supplemental Table 1). Their molecular masses range from 25 to 100 kDa, indicating a lack of small peptide toxins affecting Na\(^+\) and K\(^+\) channels identified in many sea anemones (31, 32). Four venom sequences are cnidarian-specific (supplemental Table 1), and most venom proteins are part of the soluble nematocyst proteome (Fig. 2).

**Enzymes with Lytic Function Constitute Unexpectedly High Proportion of Nematocyst Proteome**—The largest group among these is represented by peptidases (supplemental Table 1). This feature is conserved in cnidarians, as 57% of the identified protease genes have a *Nematostella* homolog compared with 52% of the structural proteins (Fig. 3A). The protease proteome includes several major families, with matrix metalloproteases being the most prominent. Matrix metalloproteases can activate toxins or act as toxins themselves (33). In addition, they are essential factors in ECM remodeling and tissue morphogenesis. We assume that their proteolytic activity is used both for the maturation of the nematocyst and for prey tissue disintegration. Other enzyme classes comprise carbohydrate metabolism and breakdown, lipid hydrolysis, and protein folding and modification.

**Nematocyst Capsule Structure Is Specialized ECM**—The largest group of annotated proteins contains molecules with an apparent structural function, including the hitherto identified capsule and spine proteins, with NOWA being one of the most prominent components (supplemental Fig. 1). The structural proteins have well defined domains that are characteristic for multidomain ECM proteins involved in protein-protein and protein-carbohydrate interactions. The identified motifs include von Willebrand factor domains, fibronectin type II and thrombospondin type 1 repeats, LamG domains, and diverse lectins (Fig. 4). As many toxins specifically interact with matrix molecules (4), the similarity of the capsule structure to the ECM might have fostered their storage and delivery. In addition, we cannot exclude evolutionary transitions from venom-related molecules to structural proteins. The SCP domain might serve as an example, as it forms a pathogenesis-related superfamily across the metazoan subkingdom (34) and is part of the major capsule structure protein NOWA.

Minicollagens constitute a well characterized subgroup among nematocyst ECM proteins (6). Our analysis revealed almost all hitherto characterized minicollagen molecules and four new members of this protein family (NColl18–21). Thus, the heterogeneity of the minicollagen family in nematocysts almost equals the collagen repertoire in the vertebrate ECM. Minicollagen detection by MS is hampered by the limited accessibility to proteolysis and the high similarity of the repetitive protein sequences. However, the majority of the minicollagens could be identified after a collagenase digest of nematocyst shells. Minicollagens are unique constituents of the cnidarian nematocyst and comprise a short central Gly-\(Y\)-repeat sequence (8–16 repeats) flanked by polyproline stretches and terminal cysteine-rich domains (CRDs) (Fig. 4), which have a canonical cysteine pattern \((\text{CX}_3\text{CX}_3\text{CX}_2\text{CX}_3\text{CCC})\) (7, 13). The CRD has been shown to be responsible for inter-

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**FIGURE 3.** A, homologs of *Hydra* nematocyst proteins identified in the *Nematostella* genome. MMPs, matrix metalloproteases. B, evolutionary distribution of the 410 nematocyst proteins. Nematocyst proteins have been assigned to categories based on the presence of significant BLAST hits and Pfam domains. The “exclusive” distribution refers to a hierarchical classification in which each protein was assigned to only one category. Proteins with significant BLAST hits in basal eukaryotes (*T. adhaerens* or *M. brevicollis*) have been classified as “eukaryotic.” The remaining proteins with significant similarity in *H. sapiens*, *D. melanogaster*, *D. rerio*, *C. elegans*, or *B. floribada* were binned as “eumetazoan.” Proteins that were not classified as either eukaryotic or eumetazoan but with significant hits in the *N. vectensis* genome were classified as “cnidarian.” The remaining proteins were further classified as “InterPro-homologous” if they had a significant hit against the InterPro Database or as “orphans” otherwise. For the “non-exclusive” distribution, each nematocyst protein was assigned to every category where a significant hit could be identified. C, domain distribution of *Hydra* nematocyst proteins with a significant Pfam hit but no significant BLAST hit to eukaryotic, eumetazoan, or other cnidarian genomes. Dark green, total number of occurrences of a domain in the data set; light green, number of proteins in which a domain occurs. \(v\)WA, von Willebrand factor; \(A\)SCP, sperm coat protein. \(D\), alignment of the proteins corresponding to the second largest cluster of orphan proteins (see “Experimental Procedures”). The conserved region is predicted to be globular and thus likely to represent a previously uncharacterized protein domain. MAM, meprin, A-5 protein, receptor protein tyrosine phosphatase mu; Cub, complement C1r/C1s, Uegf, bone morphogenetic protein; Shk7, Stichodactyla toxin domain; Shk, Stichodactyla-like domain.
molecular networks in the capsule structure, which is extremely sensitive to reducing agents (35).

The nematocyst proteome reveals the evolutionary history of the minicollagen family in a broader dimension, as it includes a series of proteins that exhibit minicollagen domains in isolation or combined with unusual motifs (Fig. 4). Non-collagenous proteins containing polyproline stretches and minicollagen CRDs were designated cnidarian proline-rich proteins (CPP) and comprise five members in the capsule proteome (Fig. 4). CRDs are also found autonomously in multiple tandem repeats, often combined with lectin domains as in NOWA (Fig. 4). The CRD thus appears as a widespread linker motif connecting collagenous and non-collagenous structural proteins of the nematocyst.

Capsule Wall Protein Cnidoin Represents Elastic Counterpart of Collagen Network—Hydra nematocysts are unique in storing energy for discharge in the capsule wall, which is elastically stretched by an extreme osmotic pressure (1). The molecular basis of the capsule’s elastic property has hitherto been elusive, as the collagen polymer alone is unlikely to be responsible for it. An intriguing example of a non-collagenous structural protein containing CRDs is Cnidoin (Fig. 4), which exhibits an extended glycine- and glutamine-rich elastic sequence (Fig. 5A) with sequence similarity to the spider silk protein spidroin-2 (36). A common repetitive element in both proteins is GXGQQ, where X is Cys in Cnidoin and Pro in spidroin-2 (37). Spidroins are composed of amorphous glycine-rich sequences providing elasticity to the silk fiber interspersed by polyalanine crystallites, which provide strength. Poly-Ala blocks are missing in Cnidoin, suggesting that it might serve as a purely elastic counterpart of the collagenous wall matrix, providing the molecular basis for the storage of kinetic energy released in the discharge process. Co-immunostaining for Cnidoin and minicollagen-1 showed overlapping signals in the walls of developing nematocysts (Fig. 5, B–D). Cnidoin appears to be more enriched in the inner wall layer (Fig. 5D), suggesting a late incorporation into the nematocyst vesicle. The resulting wall composition is reminiscent of the vertebrate blood vessel matrix, in which the inner elastin layer is stretched against the collagenous connective tissue. Recombinant expression of Cnidoin in E. coli resulted in a highly viscous protein solution, which precipitated to macroscopic sheet-like structures in aqueous solutions (data not shown). In an 8 M urea solution, purified Cnidoin readily assembled to extended fibers of 2–3-μm thickness (Fig. 5E), indicating a tendency for fibrillogenesis as described for other elastic proteins (38).

DISCUSSION

Nematocysts constitute one of the most complex cellular organelles in the animal kingdom. They represent an exclusive feature of cnidarians, and nematocyst proteins like minicollagens serve as a phylogenetic marker for this phylum as exemplified by a recent report on minicollagens in myxosporidians (39). The identification of the nematocyst proteome of Hydra for the first time allows a view into the evolutionary history of this organelle. We can clearly rule out a major transfer event by stable cellular integration of an endosymbiont, as in the case of mitochondria and chloroplasts, although proteins of bacterial origin such as those involved in poly-γ-glutamate synthesis have essentially contributed to nematocyst functions (31).

Instead, our data suggest an evolution of nematocysts that was tightly coupled with the invention of ECM-like structural proteins from exocytic vesicles of early eukaryotes specialized in venom secretion (Fig. 6). A conserved portion of the venom-related proteome comprises lytic enzymes and is of a general digestive nature. Extracellular digestion might therefore have been a transitory state in nematocyst evolution leading to phagotrophy and internal digestion (Fig. 6) (40). The close molecular relation of the nematocyst structure to ECM points to a premetazoan role of ECM molecules associated with predatory functions such as entangling prey cells in a fibrous mesh or a more efficient absorption of cellular material by the secretion of adhesive molecules (Fig. 6). We assume that the vesicular environment was optimized for ECM assembly as a basis for an accelerated adaptation (41) under the pressure of a beginning predatory phase among eukaryotes that, in turn, has led to a variety of extrusive organelles exhibiting sophisticated discharge mechanisms (Fig. 6) (42). The existence of nematocyst-like organelles in dinoflagellates (43) and other protozoans argues for an evolutionary origin of nematocysts predating...
multicellularity (Fig. 6). The nematocyst proteome can therefore shed light into the primordial state of the ECM to which the nematocyst’s biomechanical requirements probably contributed.

Minicollagens are a main molecular feature of nematocysts, but their evolutionary origin has hitherto been elusive. The CPPs constitute a family of non-collagenous proteins with distinct minicollagen features, and our data indicate a scenario of how CPPs might have evolved to minicollagens. CPP-1 shows a perfect minicollagen domain organization with a central polyproline stretch flanked by CRDs instead of the collagen triple helix (Fig. 4). In the C-terminal CRD of minicollagens, mutation of a conserved proline causes a deviated three-dimensional structure (44). In CPP-1, both CRDs have the N-terminal fold shared with all other CRDs outside the minicollagen family. We therefore propose that an early recombination event between polyproline and CRD domains might have produced a founder molecule for minicollagens.

Proline-rich proteins are found in diverse biological contexts and can be extremely polymorphic concerning post-translational modifications. They often fulfill structural functions similar to extensins in plant cell walls and adopt a polyproline II helix structure as the chains in collagen triple helices (45). Interestingly, the sialomes of salivary glands of diverse species, including humans, contain a large fraction of proline-rich proteins supposed to function in tannin binding and interaction with bacteria (46). Proline-rich proteins share sequence similarities and amino acid compositions with collagens but lack repetitive Gly-X-Y sequences (47, 48). A recent analysis of the black fly sialome identified a family of short collagen-like molecules with 7–15 Gly-X-Y repeats supposed to interact with ECM components (49). This opens the attractive perspective that CPPs evolved from secretory proteins with digestive venom-related function to structural proteins, i.e. to minicollagens. Thus, the formation of a complex ECM within the nematocyst vesicle might have been fostered by an extensive domain rearrangement within the secretome of nematocytes. We speculate that this process involved proteins representing a primitive state of the ECM and uncoupled their evolution from those constituting the cellular ECM.

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