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Journal

Biomacromolecules, 15(1)

ISSN

1525-7797

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Publication Date

2014-01-13

DOI

10.1021/bm401598z

Peer reviewed

Structural Proteins from Whelk Egg Capsule with Long Range Elasticity Associated with a Solid-State Phase Transition

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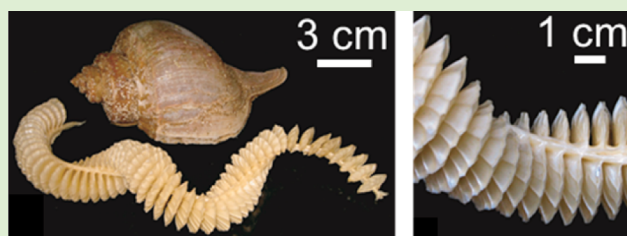
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Supporting Information

ABSTRACT: The robust, proteinaceous egg capsules of marine prosobranch gastropods (genus *Busycotypus*) exhibit unique biomechanical properties such as high elastic strain recovery and elastic energy dissipation capability. Capsule material possesses long-range extensibility that is fully recoverable and is the result of a secondary structure phase transition from α -helical coiled-coil to extended β -sheet rather than of entropic (rubber) elasticity. We report here the characterization of the precursor proteins that make up this material. Three different proteins have been purified and analyzed, and complete protein sequences deduced from messenger ribonucleic acid (mRNA) transcripts. Circular dichroism (CD) and Fourier transform infrared (FTIR) spectroscopy indicate that the proteins are strongly α -helical in solution and primary sequence analysis suggests that these proteins have a propensity to form coiled-coils. This is in agreement with previous wide-angle X-ray scattering (WAXS) and solid-state Raman spectroscopic analysis of mature egg capsules.



1. INTRODUCTION

Rubbery proteins such as elastin, resilin, and abductin are well studied from a variety of organisms in which they endow specialized tissues such as aorta, insect wings, and scallop hinge ligaments with low modulus, high extensibility (>100%), high resilience (>90%), and entropy-driven elastic recovery.^{1–4} Accordingly, the entropy of a rubbery protein chain at rest is much greater than of a chain pulled taut, hence its spontaneous recovery upon unloading. Whereas these structures are mostly disordered, α -helical proteins are also found as building blocks of a diverse variety of extra-organismal load-bearing scaffolds. The keratins found in mammalian hair, wool, nails, horns, claws, and hooves⁵ are familiar examples, whereas the hagfish slime^{6,7} is a more esoteric one. Unlike the proteinaceous egg capsules of other oviparous species, such as dogfish and spiders which produce collagen and silk-based capsule proteins, respectively,^{8,9} the whelk egg capsule is predominantly an α -helical biomaterial. This molecular design was also recently reported in the egg case of the praying mantis.¹⁰

Recently, there has been growing evidence that certain proteins or protein assemblies that feature significant long-range order as detected by X-ray scattering can undergo a phase transition during uniaxial stretching, from one crystalline structure such as α -helix to another such as β -sheet,¹¹ according to a mechanism initially suggested by Flory.¹² This paradigm

differs significantly in that recovery from large extensions is driven mostly by internal energy rather than entropy variations as a function of strain. Aculeate silk was the first fiber shown by wide-angle X-ray scattering (WAXS) to undergo an α -to- β transition, but the transformation was irreversible.¹³ Wool keratin fiber was then later shown to also undergo an α -to- β transition during extensions of up to 40% but its recovery was slow and incomplete.^{5,14–16} In contrast, the egg capsule of the prosobranch gastropod mollusks or whelks of the genus *Busycotypus* (formerly known as *Busycon*) exhibits a completely and spontaneously reversible structural transition during large extensions from α -helical coiled-coils to β -sheets¹⁷ (or extended β -strands according to recent investigations¹⁸). Other coiled-coil fibrous proteins that are susceptible to an α -to- β molecular transition include myosin, fibrin, and intermediate filaments (IFs) from the cell cytoskeleton, and a comparison of their stress–strain behaviors at various length scales has been recently reviewed.¹¹ The ability of coiled-coil fibers to undergo an α -to- β transition under mechanical stress is also well supported by molecular dynamic simulations carried out on various IFs¹⁹ and fibrinogen.²⁰

Received: August 12, 2013

Revised: December 6, 2013

Published: December 18, 2013

The *Busycotypus* egg case material consists of proteins that encapsulate embryos during their extended development period.¹⁷ Meter-long strings of hundreds of interconnected egg capsules (colloquially known as mermaid's necklaces, Figure 1) are routinely deposited on the seabed, and rely chiefly

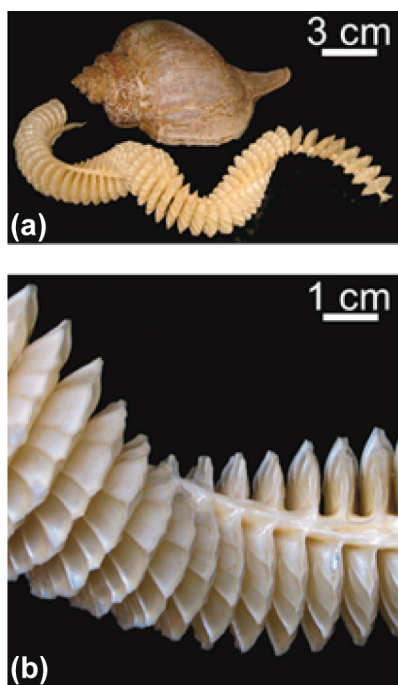


Figure 1. Photographs of (a) *Busycotypus canaliculatus* with its string of egg capsules and (b) capsules at higher magnification.

on the protein casing to protect the developing embryos from strong hydrodynamic forces associated with wave velocities in excess of 25 m sec^{-1} , abrasion from sand and other particulate debris, predation, and harmful ultraviolet (UV) light.^{21–23} Fully processed, or mature, whelk egg capsule is a highly extensible, self-assembled, proteinaceous material which possesses remarkable self-healing or shape-memory properties. The egg capsules approach the extensibility of the biological rubbers such as elastin and resilin, but have an initial stiffness 2 orders of magnitude higher. Strain energy absorption capacity of the capsule material is more than five times greater while its resilience is two-thirds that of elastin²⁴ (Table 1). Furthermore, even after the material is extended beyond its yield point, upon relaxation it returns to its original size and shape and within seconds recovers its high initial stiffness²⁵ (Figure 2). This is a rare and highly desirable combination of properties of polymer fibers.²⁶

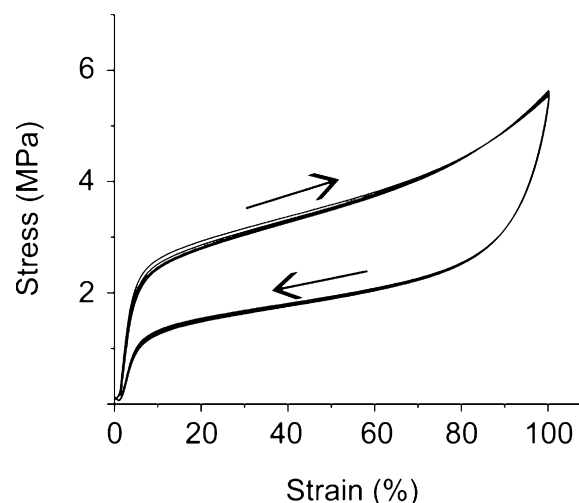


Figure 2. Typical engineering stress–strain curve of whelk egg capsule material under uniaxial tension. Sample was cyclically strained multiple times at a rate of 5 mm/min with no rest between cycles. Recovery of initial stiffness was complete and rapid.

Previous studies have detailed the tensile properties^{25,32} and structural conformation^{33,34} of the capsule material, as well as modeled the thermodynamics of its self-healing/shape-memory behavior.¹⁸ However, little is known about the protein chemistry of the egg capsule material. It is therefore of critical importance to characterize and wherever possible, reconcile the biochemistry with the observed tensile properties of the constituent proteins of this material, hereafter referred to as the *Busycotypus canaliculatus* capsule proteins (Bc-CPs).

Capsule proteins undergo at least two separate post-translational processing steps in the female whelk before the mature end product is formed. First, soluble precursor capsule proteins self-assemble into a jelly like structure which possesses no macroscopic load-bearing capability. Then, this jelly like structure is covalently cross-linked before the egg capsule is released and abandoned on the ocean floor.³² As a consequence of the covalent cross-linking or sclerotization, it is extremely difficult to extract full-length, intact protein from the mature capsule. However, the precursor proteins are readily extractable from gravid female nidamental glands for subsequent purification. Previous work has determined that the precursors extracted from the gland tissue are the same as the proteins present in the immature capsules.³² Using nidamental gland-derived precursors, we determined the protein molecular masses, amino acid compositions, primary amino acid sequences, secondary structures in solution and self-assembly behavior. From these data, as well as data from previous work,

Table 1. Mechanical Properties of the Whelk Egg Capsule Compared with Other Fibrous Protein Materials^{a,5,24,27–31}

protein material	E_i (MPa)	yield stress (MPa)	breaking stress (MPa)	yield strain (%)	breaking strain (%)	strain energy to failure (MJ m^{-3})	resilience (%)
whelk egg capsule	60	2.5	10.0	5	169	8.5	49
wool keratin	3000	30	150	5	52	37.4	55
resilin	2	NA	4	NA	190	4	92
elastin	1	NA	2	NA	150	1.6	90
tendon collagen	1200	20	39	1.5	12	4.1	90
dragline silk	10000	160	1100	4	30	160	35

^aStrain rates for egg capsule, keratin, collagen and silk are all comparable, but are unavailable for resilin and elastin.

plausible structure–function models for these proteins are proposed.

2. MATERIALS AND METHODS

2.1. Materials. All materials were purchased from Sigma-Aldrich or Fisher Scientific unless otherwise stated.

2.2. Capsule Proteins Purification. Live whelks (*Busycotypus cancellatus*) were purchased from the Marine Biological Laboratories (Woods Hole, Massachusetts, U.S.A.). Females were euthanized by immersion in a cold ethanol bath and were immediately dissected to remove the nidamental gland which was then placed at -80°C . Crude extract was prepared by treating the gland material with 6 M urea in 5% acetic acid and 50 mM tris-(2-carboxyethyl)phosphine (TCEP) to reduce disulfide bonds. Reverse phase high pressure liquid chromatography (RP-HPLC) was performed using a C8 column (Perkin-Elmer Brownlee Aquapore Octyl C8 or Agilent Technologies Zorbax 300SB-C8 with a Varian ProStar or Agilent Technologies 1260 Infinity HPLC system, respectively). Crude extract and purified protein fractions were assayed via sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) using 15% polyacrylamide gels.

2.3. Mass Spectrometry. The mass-to-charge ratio (m/z) was determined by matrix-assisted laser desorption/ionization with time-of-flight (MALDI-ToF) mass spectrometry using a PerSeptive Biosystems Voyager DE mass spectrometer (AB Biosystems, Foster City, California, U.S.A.). Measurements were also carried out on an AXIMA Performance mass spectrometer (Shimadzu, Kyoto, Japan). In both cases, protein solutions were dried on a sample plate, over which a saturated solution of sinapic acid in 50% acetonitrile and 0.1% trifluoroacetic acid was subsequently added and dried.

2.4. Amino Acid Composition Analysis. Lyophilized RP-HPLC fractions were hydrolyzed in vacuo using 6 N HCl at 110°C for 24 h. Following hydrolysis, the samples were washed with high purity water and methanol, twice each, and dried. Amino acid compositions were determined using a Beckman Coulter 6300 amino acid analyzer (Brea, California, U.S.A.) with ninhydrin-based post column derivatization as described elsewhere.²⁴ Additional measurements were also acquired using a Sykam S433 amino acid analyzer (Eresing, Germany) with identical sample preparation procedures.

2.5. Edman Sequencing. Lyophilized RP-HPLC fractions were subjected to Edman sequencing reactions using a Porton Instruments PI 2020 as described elsewhere.³⁵ For CP-2 and CP-3, N-terminal Edman sequencing was also performed by transferring protein from SDS-PAGE gel to Immobilon-P polyvinylidene difluoride (PVDF) membrane (Qiagen, Valencia, California, U.S.A.), excising bands and submitting them to the Protein Facility of Iowa State University (Ames, Iowa, U.S.A.) for analysis.

2.6. Reverse Transcription Polymerase Chain Reaction (RT-PCR) with Rapid Amplification of cDNA Ends (RACE) and Sequencing. RNA was purified from fresh (nonfrozen) nidamental gland tissue immediately following dissection with TRIzol Reagent following the manufacturer's instructions. Total RNA was reverse transcribed to RACE-ready cDNA (cDNA) using a GeneRacer Kit (Invitrogen, Carlsbad, California, U.S.A.), primers were purchased from Integrated DNA Technologies (Coralville, Iowa, U.S.A.) and PCR reagents were purchased from Novagen (Darmstadt, Germany). PCR was carried out using a Mastercycler Gradient thermocycler (Eppendorf, Hauppauge, New York, U.S.A.). PCR products were purified using 1.1% agarose gels, visualized by ethidium bromide, and bands were excised and purified with a Qiagen Gel Extraction Kit. PCR products were directly sequenced by GENEWIZ (La Jolla, California, U.S.A.).

From the partial N-terminal sequence, degenerate primers were designed for RT-PCR. First, 5' RACE-PCR was performed to give the sequence of the 5' untranslated region. Next, 3' RACE-PCR was performed to give cDNA sequences of the signal peptide to the start methionine and beyond for each transcript.

Degenerate primers used for the Bc-CPs:

5' RACE. CP1-a: 5'-GAYTNGAYTTYGAYGTNCCNAAR-GARGCN-3'; CP1-b: 5'-GCNYTNGAYTTYGAYGTNACNAAR-GARGCN-3'; CP2/3: 5'-TTYCARGGNTTYAAAYTTYGGNAARAA-RGAY-3', where Y = C, T; R = A, G; N = A, C, G, T.

3' RACE. CP1-a: 3'-CTRRANCTRAARCTRCANGGNTTYCTYCGH-5'; CP1-b: 3'-CGNRANCTRAARCTRCANTGNTTYCTYCGN-5'; CP2/3: 3'-AARGTYCCNAARTTRAARCCNNTTYTTYCTR-5', where Y = C, T; R = A, G; N = A, C, G, T.

From the results of the 5' and 3' RACE-PCR, new sets of nondegenerate primers were designed for 3'-RACE to amplify full-length transcripts.

2.7. Circular Dichroism (CD) Spectroscopy. Far UV (190–260 nm) CD measurements were carried out using an Olis RSM 1000 spectrophotometer (Olis, Bogart, Georgia, U.S.A.) with a 0.5 mm quartz cell at 1 nm resolution. Purified protein solutions were prepared in 50 mM acetate buffer, pH 4.0. A total of 10 spectra were taken at room temperature and the results were averaged. Spectra are presented in mean residue ellipticity with protein molecular weight determined by MALDI-ToF, and protein concentration estimated by quantitative ninhydrin-based amino acid analysis (see above) as described elsewhere.³⁶ Secondary structure analysis was performed using K2D3 software;³⁷ each spectrum was scaled linearly until a best fitting curve was found.

2.8. Fourier Transform Infrared (FTIR) Spectroscopy. Infrared absorption measurements were performed using a VERTEX 70 or 70v FTIR spectrometer (Bruker, Karlsruhe, Baden-Württemberg, Germany), respectively, equipped with a PIKE MIRacle or Harrick Horizon attenuated total reflection (ATR) measuring unit, each containing a ZnSe ATR crystal (angle of incidence $\theta = 45^{\circ}$). Purified protein solutions were prepared in 50 mM acetate buffer, pH 4.0. The sample chamber was continuously purged with dry carbon dioxide-free air, and the total reflected IR beam intensity was measured using a liquid nitrogen-cooled photovoltaic mercury cadmium telluride (MCT) detector. Spectra were recorded with a spectral resolution of 4 cm^{-1} in double-sided acquisition mode, and the mirror velocity was set to 80 kHz. At least 1000 scans were taken for each spectrum, which was calculated using a 3-term Blackman-Harris apodization function. Spectra of proteins were obtained by subtracting IR absorption due to acetate buffer. Secondary structure analysis was performed using the OPUS 6.5 software (Bruker, Karlsruhe, Baden-Württemberg, Germany).

IR absorption spectra of solid-state egg capsules were also acquired in the ATR mode under the same conditions.

2.9. Calculation of Protein Dimensions. Based on the results of sequencing and the secondary structure analyses of the FTIR spectra, the average number of amino acid residues per protein chain present in the capsule material was calculated and the relative amounts of α -helices, β -sheets, and random coils were determined. Those numbers were used to compute the maximum end-to-end protein length by summing the length of each type of secondary structure, assuming that they are linearly arranged. The translational rise is 1.5 Å per residue for α -helix, 3.5 Å per residue for parallel β -sheet, and 3.3 Å per residue for antiparallel β -sheet. For the length of random coil, statistical analysis of steric restrictions was taken into account and a characteristic ratio C_{∞} of 9.0 was used as described by Creighton.³⁸ That resulted in a calculation that was dictated by the equation:

$$\langle r^2 \rangle_0^{1/2} = (130n)^{1/2}$$

where the root-mean-square $\langle r^2 \rangle_0^{1/2}$ was taken as the end-to-end length (Å).

3. RESULTS

3.1. Capsule Proteins Purification, Mass Spectrometry, and Amino Acid Composition Analysis. Consistent with previous findings, SDS-PAGE analysis of nidamental gland crude extract indicates a family of proteins around 50 kDa with three distinct bands. In order of increasing molecular mass, the bands are named CP-1, CP-2, and CP-3. Using RP-HPLC, each

individual protein in this family can be purified from the crude extract (Figure 3). In order of increasing retention time, the

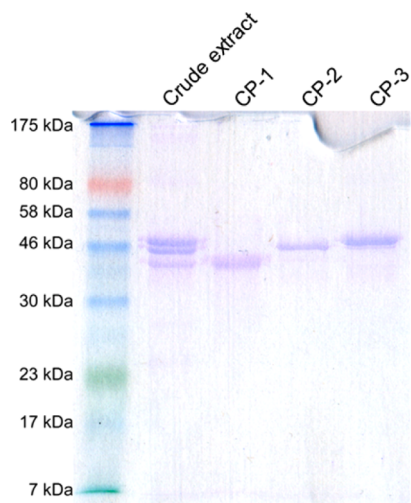


Figure 3. SDS-PAGE of Bc-CPs purified from nidamental gland crude extract.

proteins are CP-1, CP-3, and CP-2. MALDI-ToF mass spectrometry indicates the molecular masses of CP-1, CP-2,

and CP-3 to be 47.5, 51.6, and 52.1 kDa, respectively (Figure 4). The three proteins have amino acid compositions that are similar to one another as determined by ninhydrin-based analysis. They are also similar to hard α -keratins in terms of richness in Asx, Glx, and Leu, but different in two key aspects. First, α -keratins have high Cys, or disulfide, content, whereas Bc-CPs have almost none. Second, Bc-CPs have a much higher Lys content than α -keratins³⁹ (Table 2).

3.2. Protein and cDNA Sequencing. N-terminal amino acid sequence of the first ten residues was obtained by automated Edman reaction involving the RP-HPLC purified fractions. CP-1 has a unique N-terminal sequence whereas CP-2 and CP-3 share the first 10 amino acids. This sequence identity for CP-2 and CP-3 was verified by repeating the Edman reaction with SDS-PAGE membrane transfers of crude extract (Supporting Information, Table S1).

From the partial N-terminal sequence, degenerate primers were designed for RT-PCR. First 5' RACE was performed to give the sequence of the 5' untranslated region. Then, nondegenerate primers were designed for 3' RACE to amplify full-length transcripts. 3' RACE of the CP-1 cDNA transcript produced two PCR products, one at ~1500 bp and another at ~2400 bp, whereas that of the CP-2/CP-3 cDNA transcript produced only one PCR product at ~2400 bp (Supporting Information, Figure S1). These three PCR products were gel-excised and sequenced directly without cloning.

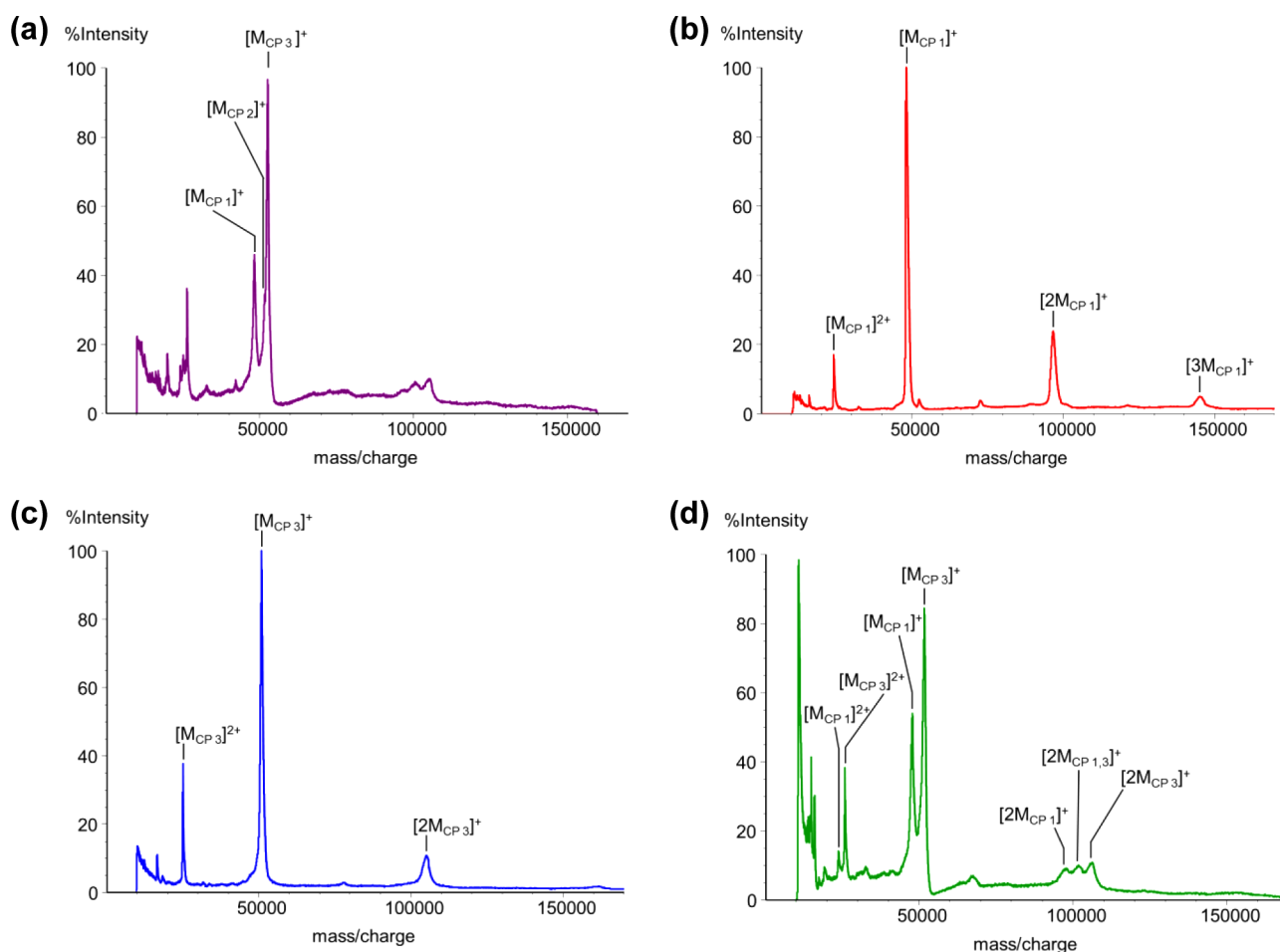


Figure 4. MALDI-ToF mass spectra of (a) nidamental gland crude extract, (b) CP-1, (c) CP-3, and (d) a mixture of CP-1 and CP-3. Observed molecular masses of CP-1, CP-2, and CP-3 are 47.5, 51.6, and 52.1 kDa, respectively.

Table 2. Amino Acid Composition, in Mole Percent, of Purified Bc-CPs and Egg Capsule, Showing Large Differences in Cys and Lys Content Compared to α -Keratin^{a,39}

amino acid	CP-1	CP-2	CP-3	bulk egg capsule	α -keratin
Ala	9.0	6.4	6.3	7.5	7.7
Arg	4.7	4.2	4.1	3.9	7.9
Asx	10.1	10.8	10.7	13.5	9.6
Cys	0.0	0.0	0.0	0.0	6.0
Glx	18.6	16.1	16.7	15.6	16.9
Gly	5.1	5.9	6.3	8.0	5.2
His	2.2	3.4	3.7	1.5	0.6
He	5.9	5.4	6.1	4.9	3.8
Leu	9.5	9.1	8.6	9.5	10.2
Lys	12.0	12.2	12.6	9.4	4.1
Met	0.0	0.0	0.0	0.8	0.6
Phe	2.4	4.1	5.6	4.6	2.0
Pro	0.9	2.9	1.6	1.9	3.3
Ser	7.2	7.9	7.5	8.0	8.1
Thr	5.0	4.7	3.5	5.0	4.8
Tyr	0.3	0.5	0.0	0.4	2.7
Val	7.1	6.4	6.7	5.5	6.4
Total	100	100	100	100	100

^aThe maximum standard deviation is ~5% of the values shown.

All three transcripts contain an open reading frame, which translates to a full length gene with start (methionine) and termination codon. Despite the difference in the sizes of the PCR products, both CP-1 transcripts translate to proteins of identical length comprising 446 amino acid residues and have 95% sequence identity. Hereafter, these variants are referred to as CP-1a and CP-1b. However, unless otherwise stated, when a protein is described as “CP-1”, it is a mixture of CP-1a and CP-1b.

The transcript for CP-2 and CP-3 translates to a single protein comprising 479 amino acid residues (Figure 5). The fact that this transcript is responsible for the production of two different proteins can be explained by potential early termination in the translation process. Since sequencing was performed directly on gel-purified PCR products without cloning, multiple variants may be present, resulting in different nucleotide sequences at the same position in the transcript. The primary codon at position 476 is AGA (arginine). However, in the raw sequencing chromatogram there is a smaller, but still significant, thymine peak in the first position. This would result in TGA, a termination codon, which reduces the length of the protein by four amino acid residues. These four residues have a combined weight of 578.6 Da. This accounts for the observed molecular mass difference between CP-2 and CP-3. The computed molecular masses (excluding signal peptides) of the cDNA-deduced protein sequences are consistent with the actual protein molecular masses observed by MALDI-ToF mass spectrometry (Figure 4). Amino acid compositions of the purified proteins also agree with the compositions predicted from cDNA-deduced protein sequences (Table 3).

In addition, all three transcripts code for signal peptides, indicating that these proteins are secreted via secretory vesicle pathways. Cleavages between G-21 and D-22 in CP-1a, G-21 and A-22 in CP-1b, as well as G-18 and L-19 in both CP-2 and CP-3 are predicted using SignalP (ExpASY) and match the N-terminal sequences derived by Edman sequencing. Sequence

analysis was performed using BLAST (NCBI), which did not produce a match with E -value $< 10^{-9}$ for CP-1a and CP-1b, and E -value $< 10^{-5}$ for CP-2 and CP-3 in the database of sequences from eukaryotic species, indicating that the Bc-CPs are not homologous to any known eukaryotic protein.

3.3. CD and FTIR Spectroscopy. Both far-UV CD spectroscopy and mid-IR FTIR spectroscopy were used to examine the secondary structures of Bc-CPs in solution. Lyophilized RP-HPLC purified Bc-CPs were resuspended in 50 mM acetate buffer, pH 4.0 (lyophilized proteins have limited solubility at more neutral pH), and spectra were collected at room temperature.

The CD spectra of both CP-1 and CP-2 (Figure 6) were characteristic of single coil α -helical structure,⁴¹ exhibiting two minima at 208 and 222 nm with a $[\theta]_{222}/[\theta]_{208}$ ratio of < 1 . Secondary structure analysis by K2D3 showed that, while both CP-1 and CP-2 are dominated by α -helices ($> 50\%$), there are small amounts of β -sheets ($\sim 6\%$). The existence of β -sheets is confirmed by FTIR spectroscopy, as described below.

The overall FTIR absorption spectra of CP-1 and CP-3 in the ATR mode (Figure 7c,d) show sharp intense bands centered at ~ 1650 and ~ 1540 cm^{-1} , corresponding to Amide I and Amide II bands, respectively, with a clear presence of methylene peaks between 2800 and 3000 cm^{-1} . In order to determine the various vibration modes of the peptide backbone and obtain the relative amounts of the different types of secondary structures, the Amide I peak was curve-fitted using the Bruker OPUS 6.5 software. A 80% Gaussian/20% Lorentzian function was employed to deconvolute the Amide I band into its constituent vibration modes. The positions of the constituent bands were held fixed, while the intensities and bandwidths were allowed to vary. From the area under the fitted curves, the relative fractions of α -helices, β -sheets, and random coils were determined. Assignment of Amide I components to the various secondary structures were based on literature.⁴² All samples are dominated by a high amount of α -helices, ranging from 70 to 75% for the pure Bc-CPs. The β -sheet content, although lower, remains significant at 15 to 20%, while unordered regions account for about 10% of the secondary structure. FTIR spectroscopy was also performed on an equimolar mixture of CP-1 and CP-3. The mixture exhibits a significant decrease in the relative fraction of α -helices ($\sim 60\%$), together with an increase in both β -sheets ($\sim 23\%$) and unordered regions ($\sim 15\%$), thus, resulting in an overall lower α -helix/ β -sheet ratio compared to the individual Bc-CPs.

FTIR spectra of pure Bc-CPs that were subjected to freeze-drying from a highly concentrated state show significant differences in the shape of the Amide I band in comparison to Bc-CPs that were freeze-dried from diluted solutions (Supporting Information, Figure S4). For both proteins, a strong decrease in α -helical content is observed (40 and 20% for CP-1 and CP-3, respectively) and accompanied by an increase in β -sheet content (up to 50% for CP-1). In other words, in going from a state of diluted to concentrated solution, the secondary structure of the Bc-CPs changes from an α -helix to a β -sheet dominated one. As a result, the proteins also became less soluble.

In addition, mid-IR FTIR spectroscopy was used to examine the secondary structure of the *solid-state* native egg capsule in the ATR mode. The overall spectrum (Figure 7a) similarly shows a clear presence of Amide I and II bands, as well as methylene peaks. Curve-fitting (Figure 7b) revealed a high α -helical content, but it is significantly lower compared to the

Table 3. Amino Acid Compositions and Molecular Masses of Bc-CPs^a

	CP-1a		CP-1b		CP-2		CP-3	
	observed	predicted	observed	predicted	observed	predicted	observed	predicted
Ala	9.0	8.5	9.0	8.9	6.4	5.7	6.3	5.6
Arg	4.7	4.5	4.7	4.5	4.2	3.1	4.1	3.3
Asx	10.1	11.5	10.1	11.6	10.8	12.9	10.7	11.8
Cys	0.0	0.7	0.0	0.7	0.0	0.4	0.0	0.4
Glx	18.6	16.2	18.6	16.9	16.1	14.8	16.7	14.8
Gly	5.1	4.9	5.1	4.9	5.9	5.9	6.3	5.9
His	2.2	1.9	2.2	2.1	3.4	3.7	3.7	3.7
He	5.9	7.5	5.9	6.8	5.4	6.6	6.1	6.5
Leu	9.5	9.6	9.5	9.9	9.1	9.4	8.6	9.3
Lys	12.0	10.6	12.0	10.4	12.2	11.4	12.6	11.5
Met	0.0	0.7	0.0	0.7	0.0	0.7	0.0	0.7
Phe	2.4	2.1	2.4	2.1	4.1	5.9	5.6	6.3
Pro	0.9	0.5	0.9	0.2	2.9	0.9	1.6	0.9
Ser	7.2	8.2	7.2	7.8	7.9	7.9	7.5	8.8
Thr	5.0	5.4	5.0	5.6	4.7	3.9	3.5	3.9
Trp		0.0		0.0		0.2		0.2
Tyr	0.3	0.0	0.3	0.0	0.5	0.0	0.0	0.0
Val	7.1	7.1	7.1	6.8	6.4	6.6	6.7	6.5
total	100	100	100	100	100	100	100	100
mass (Da)	47461	46880	47461	46962	51550	51437	52129	52016

^aObserved values are obtained with ninhydrin-based analysis and MALDI-ToF mass spectrometry, whereas predicted values are derived from translated cDNA sequences. For the observed values, the maximum standard deviation is ~5% of the values shown.

conformation when the load is removed, whereas the hagfish slime is irreversibly deformed. Hydrated α -keratins recover slowly, dissipating energy, but are not as extensible as the egg capsule. What is most remarkable is the ability of the capsule material to rapidly and fully recover its initial stiffness despite having gone through multiple cycles of tensile strain.¹⁷ Biochemical differences between keratins, hagfish, and *Busycotypus* egg cases are likely to explain the differences in their mechanical properties (Figure 8).

Interestingly, this overall stress-strain behavior has been observed or predicted at the single-molecule level in other coiled-coil fibrous proteins. In myosin, AFM single-molecule force spectroscopy experiments yielded similar force-extension curves, consisting of an initial linear response, followed by a plateau region, and restiffening at large elongations, and the process was reversible.⁴⁴ More recently, the same behavior was predicted by molecular dynamic simulations of force-induced unfolding of fibrinogen,²⁰ where β -sheets were formed at high extensions, thus suggesting a somehow "universal" mechanism of α - β transition in coiled-coil load-bearing proteins.

4.2. *Busycotypus* Egg Case Chemistry and Coiled-Coil Predictions. Proteins comprise over 90% of the dry weight of the whelk egg capsule^{17,25} and the Bc-CPs characterized in this work represent the most abundant of these.³² The amino acid composition of the Bc-CPs is highly similar to the capsule material and is predominantly made up of amino acids, such as Ala, Leu, Met, Lys, and Glx (Gln or Glu), that have a high propensity to form α -helix, with relatively few amino acids, such as Val, Ile, Phe, Tyr, Cys, and Trp, that have the propensity to form β -sheet. This composition is also very similar to the α -keratin filaments, but with a striking difference in the Cys and Lys contents. Cys is the primary cross-linking residue in the keratins,⁵ and previous studies have suggested Lys as the potential cross-linker in the egg capsule.^{45,46} As seen from the results of sequencing (Figure 5 and Supporting Information, Figure S2), CP-2 has the same sequence as CP-3 and is only

four amino acid residues shorter at the C-terminal than CP-3. Since the sequences of CP-2 and CP-3 are more than 99% similar, they are likely to exhibit identical structural properties and are therefore treated as the same protein, henceforth, unless otherwise stated.

Although there are striking similarities between the whelk egg capsule material and IF materials in terms of mechanics, amino acid compositions, and the ability of the precursor proteins to self-assemble into filaments in vitro without accessory molecules, primary sequence analyses indicate that the Bc-CPs are very different from the IF proteins. BLAST (NCBI) analyses produced *E*-values of $<10^{-9}$ for CP-1a and CP-1b and $<10^{-5}$ for CP-2 and CP-3, indicating that they are not homologous to every known eukaryotic IF protein in the database. Each of the Bc-CPs contains a signal peptide sequence at the N-terminus. These short sequences, 21 residues long in CP-1a and CP-1b, and 18 residues long in CP-2 and CP-3, indicate that the Bc-CPs are exported from the cell via a controlled vesicle based secretory pathway. This is in direct contrast to the hard α -keratins and the hagfish slime proteins, which remain in the intracellular space due to cornification or enter the extracellular region by apoptosis of the specialized cells in which they are accumulated.^{47,48}

As was with IF materials, X-ray scattering studies have shown that the α -helical filaments of the whelk egg capsule are in the form of a coiled-coil structure.⁵ This well-known structure is the result of a heptad repeat of amino acid residues *a-g*, in which apolar residues at positions *a* and *d* form hydrophobic interactions that bring and hold two or more α -helices together, while the flanking charged residues *e* and *g* provide electrostatic stabilization.⁴⁹⁻⁵¹ Thus, we strongly expected that sequence analysis of CP-1 and CP-3 would predict for coiled-coil structures. CP sequences were subjected to coiled-coil analysis using the SCORER 2.0 software⁴⁰ which predicts for coiled-coil propensity, while the canonical heptad repeat *abcdefg* of coiled-coil domains was assigned based on the MARCOIL software.⁵²

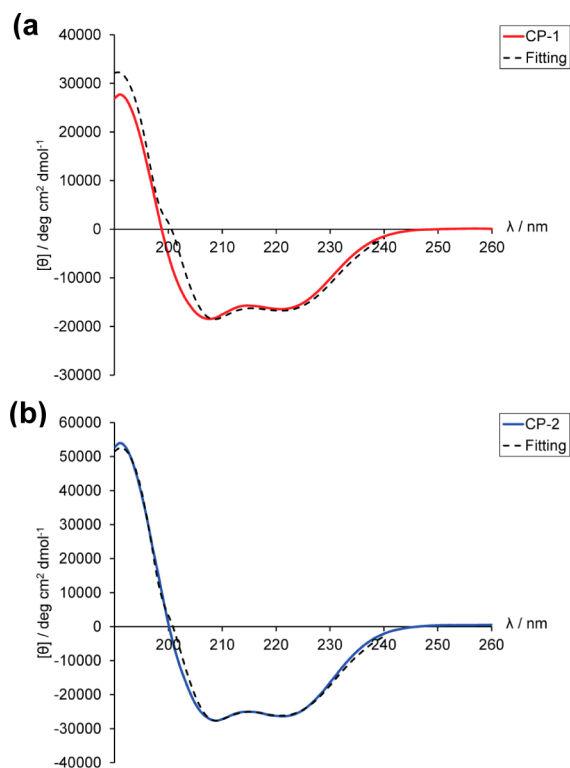


Figure 6. CD spectra of purified (a) CP-1 and (b) CP-2 in acetate buffer, pH 4.0. Minima at 208 and 222 nm and a $[\theta]_{222}/[\theta]_{208}$ ratio of <1 are representative of single coil α -helix. Concentrations of CP-1 and CP-2 were ~ 0.8 and ~ 1.2 mg/mL, respectively. K2D3 curve-fitting³⁷ for both CP-1 and CP-2 revealed secondary structure compositions that are dominated by α -helices ($>50\%$), with small amounts of β -sheets ($\sim 7\%$). CD spectroscopy was also performed on mixtures of CP-1 and CP-2 in different molar ratios at pH 4.0. This also produced similar α -helical spectra with a $[\theta]_{222}/[\theta]_{208}$ ratio of <1 (Supporting Information, Figure S3).

The results are shown in Figure 5, where the heptad domains are highlighted in yellow and blue. Not surprisingly, large coiled-coil domains are unambiguously predicted for both CP-1 and CP-3. Using a 50% confidence level, both proteins are predicted to contain $\sim 50\%$ coiled-coils, with domains as large as 25 heptads long. Additional corroboration of coiled-coil predictions was also obtained via the Lupas method⁵³ (data not shown). Furthermore, the SCORER algorithm predicts that coiled-coil domains are more likely to form trimers, in contrast to the dimers found in hagfish slime and α -keratin materials, and the same prediction was obtained using Berger's method.⁵⁴ Based on the strong suggestion of coiled-coil trimers, the capsule proteins may be more closely related to fibrinogen, which is a trimeric coiled-coil protein that also has load-bearing capability, than the α -keratins or hagfish slime filament structural measurements.⁵⁵ A further inspection of heptad assignment revealed the presence of stutters and heptad discontinuities involving deletions of 2 residues (Figure 5). These coiled-coil irregularities may play a critical role in the α - β transition, as discussed in section 4.3.

Interestingly, while there is little difference in the hydrophobic and charged residue content (Supporting Information, Figure S6) in the coiled-coil domains of the Bc-CPs compared to other known coiled-coil proteins in the CC+ database,⁵⁶ there is a relatively large amount of Gly and polar amino acid

residues (Ser, Thr, Asn, Gln). On the other hand, the Bc-CPs contain relatively fewer aromatic residues such as Phe, Trp and Tyr (Figure 9). These differences are notably evident within the hydrophobic core at heptad positions *a* and *d*. Very likely, with significantly fewer bulky aromatic groups, the coiled-coil Bc-CPs are able to undergo easy and quick uncoiling and recoiling, during loading and unloading, respectively, due to less steric hindrance. In addition, the larger presence of polar groups across all heptad positions in the Bc-CPs is likely to yield hydrophilic interactions with water (considering that the native egg case serves to function in a marine environment), thereby assisting water in its role as a plasticizer that would enable smooth and fast structural transitions both upon loading and unloading by increasing chain flexibility.

4.3. Egg Case Protein Secondary Structure by CD and FTIR Spectroscopy. In agreement with the helix-favoring amino acid composition and coiled-coil predictions, both CD and FTIR data indicate that the Bc-CPs are strongly α -helical in solution. In order to obtain some insights about post-translational self-assembly and processing, it is instructive to compare the relative secondary structure content obtained by FTIR in the precursor proteins in solution and in the fully processed, mature capsule in a hydrated state. In the solid-state capsule material, the α -helical content is less pronounced, and concomitantly, there are relatively more β -sheets and random coils. In situ tensile experiments using WAXS¹⁷ and Raman spectroscopy¹⁸ have confirmed the dominance of α -helices in the unstrained capsule material, but β -sheets, on the other hand, were not detected. While Raman and FTIR spectroscopy are complementary techniques, FTIR has a higher sensitivity than Raman, and is more suitable for semiquantitative analysis.⁵⁷ In Raman spectroscopy, the signal intensity of a specific band does not vary linearly with the relative abundance of the secondary structure. It is thus very likely that a structure present in significant amount can be buried in the background, especially when its vibration mode has a weak polarizability. Such a signal would show up only when the structure becomes dominant in the sample. In the previous measurements, the extended β -structure (which is also clearly detected by X-ray) was detected at high strain but its intensity was low in comparison to other Raman bands. Hence, it is likely that in these previous investigations, the initial β -sheet content was too low to be detected. On the other hand, the current data obtained by ATR-FTIR spectroscopy indicates a significant presence of β -sheets in the native material.

In the capsule material, the α -helix/ β -sheet ratio is lower than the individual CP-1 or CP-3 but is similar to the equimolar mixture of them. In addition, the α -helix/random coil and β -sheet/random coil ratios are lower than the individual CP-1 or CP-3 but are similar to the equimolar mixture of them. These ratios suggest that assembly and cross-linking of the precursor proteins into the mature egg capsule is associated with β -sheets and random coils formation at the expense of α -helices. Because the relative content of α -helices and β -sheets in the equimolar mixture of CP-1 and CP-3 is also significantly closer to the capsule material than the individual CP-1 or CP-3 is, these data suggest that during capsule processing, CP-1 and CP-3 associate to form localized regions of interchain β -sheets. However, it is unknown if they would associate and form dimeric or trimeric coiled-coils. Furthermore, it is also unknown if the coiled-coils of the egg capsule are homomeric or heteromeric. The propensity of the Bc-CPs for β -sheet formation is supported by FTIR measurements of individual

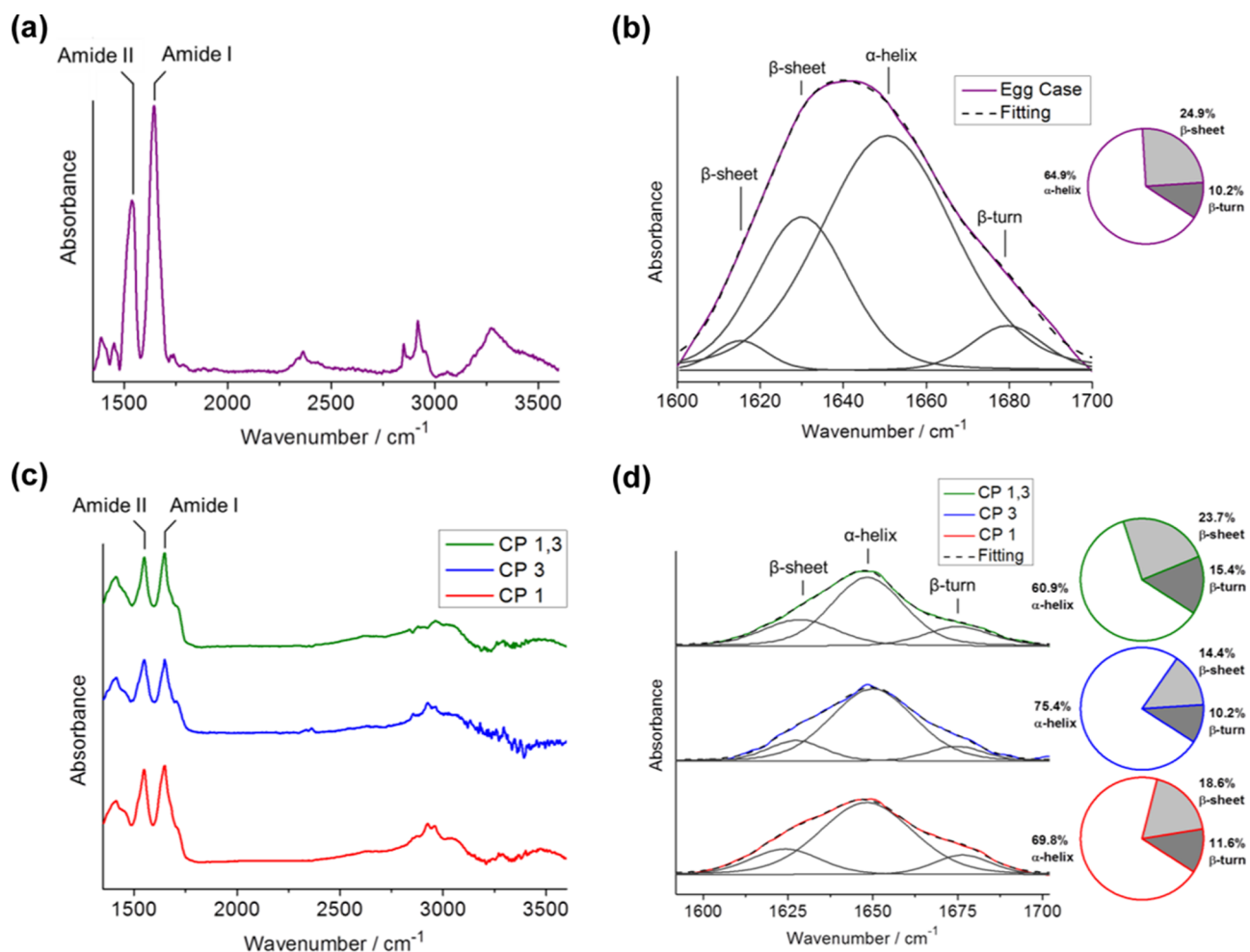


Figure 7. ATR-FTIR spectra of native (hydrated) egg case and purified egg case proteins. (a) Native egg capsule and (b) curve-fitted Amide I spectra of the egg case. (c) Purified Bc-CPs and (d) curve-fitted Amide I of purified Bc-CPs showing secondary structure compositions that are dominated by α -helices (>50%). The α -helix/ β -sheet ratio is ~ 2.5 in the egg capsule and in the mixture of CP-1 and CP-3 and is significantly lower than that in CP-1 or CP-3 individually.

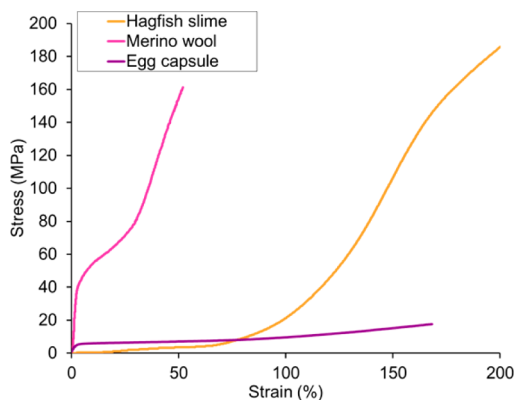


Figure 8. Engineering stress–strain curves of the whelk egg capsule material, hagfish slime, and merino wool keratin.²⁸ Figure adapted from ref 28 with permission from The Royal Society.

CP-1 and CP-3, which were freeze-dried from concentrated solutions in 5% acetic acid (Supporting Information, Figure S4). A clear increase in β -sheet content was noticed with higher concentrations and was accompanied with a lower protein solubility. However, the Bc-CPs very likely misfolded into non-

native amyloid-like structures, as is commonly seen with many other proteins at high concentrations.⁵⁸

The SDS-PAGE of immature capsules and crude nidamental gland extract indicated a trace presence of proteins other than the Bc-CPs. Therefore, although TEM showed that the purified Bc-CPs are capable of self-assembling into fibers (Supporting Information, Figure S7), an equally plausible scenario that could contribute to the increase in the β -sheets and random coils content during the natural assembly process is an incorporation of additional proteins, which may serve to bundle the coiled-coil filaments into bigger fibers that make up the mature egg capsule. The presence of such proteins in the mature capsule is further suggested by a significantly high Gly content that cannot be solely attributed to the Bc-CPs.

CD experiments were carried out at pH 4 in order to keep CP-1 and CP-2 in solution. At this low pH, CD spectra of individual Bc-CPs suggest that they preferentially adopt single-coil rather than coiled-coil conformations, as seen from the $[\theta]_{222}/[\theta]_{208}$ ratios, which are < 1 ^{59,60} (Figure 6). Given that both CP sequences unambiguously predict for coiled-coil structures and the clear coiled-coil signature of mature capsules by WAXS, it is reasonable to assume that single-coil helices assemble into coiled-coils upon secretion into the higher pH extracellular environment (seawater, pH ~ 8). CD spectra at

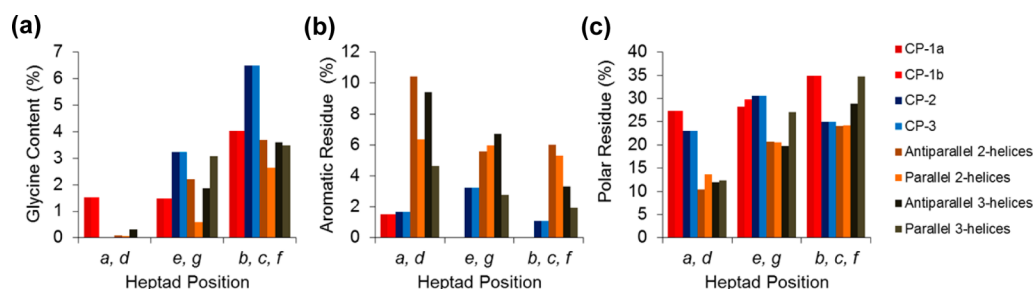


Figure 9. Relative amounts of (a) glycine, (b) aromatic, and (c) polar amino acid residues at various heptad positions in the predicted coiled-coil domains of Bc-CPs, and other known coil-coiled proteins in the CC+ database.⁵⁶ In the Bc-CPs, the glycine and polar residue content at the heptad positions a and d is notably higher, whereas the aromatic residue content is significantly lower.

seawater pH or even neutral pH could not be obtained since both proteins would not solubilize at pH above 5.5.

A critical question we sought to answer involves understanding the molecular mechanisms by which the α -helix to β -sheet transition (and its associated unique mechanical properties) is initiated in the mature capsule. We propose two possible mechanisms by which the phase transition can be initiated by means of nucleation, namely, (i) by elongation of pre-existing β -sheets and (ii) by local unwinding of the coiled-coil at regions of instabilities due to stutters or where the heptad is incomplete (Figure 10). In the first case, while the β -sheet content does not

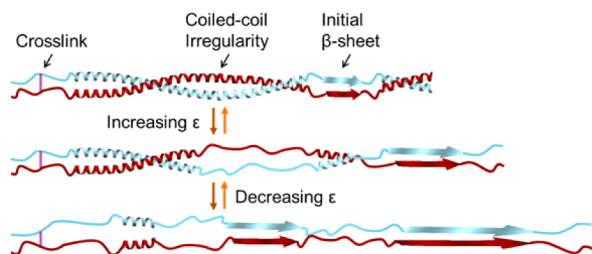


Figure 10. Proposed model of α - β transition, revised after ref 17. Regions of coiled-coil irregularity, such as stutter or stammer, and the ends of pre-existing β -sheets can both serve as stress concentration points where extension and unraveling of coiled-coils occur to form new β -sheets during uniaxial tension. (Coiled-coils are arbitrarily chosen to be dimeric for the purpose of the illustration.)

vary significantly between the solution states and the solid egg capsule, its presence in the unstrained capsule material is unlikely to be fortuitous and may have a significant role in the strain-induced α -helix to β -sheet phase transition. Thermodynamically, small domains of β -sheets can serve as nucleation sites for the strain-induced α -helical coiled-coil to β -sheet transformation detected by WAXS. β -Sheets are extended crystalline structures and are inherently stiffer than other secondary structures, as is well-established in silks⁶¹ and other IFs that undergo α -helix to β -sheet transition.⁶² Hence, when a series of stiff β -sheets and compliant coiled-coils are pulled axially, a stress concentration occurs at the coiled-coil/ β -sheet interface. This creates an ideal driving force for the coiled-coils or random coils at the interfaces to transform into new β -sheets that add to the amount of pre-existing β -sheets. The presence of pre-existing interchain β -sheets in a coiled-coil of CP-1 and CP-2 would then mean that the new β -sheets are formed between the chains of CP-1 and CP-2 as the coiled-coil extends and unfolds. In other words, much like impurities are well-known to help initiate liquid-to-solid phase transitions by lowering the activation energy barrier, initial β -sheet crystals

would be able to locally initiate the large-scale strain-induced phase transition by lowering the energy barrier for the transition. The second mechanism postulates that the transition starts occurring at coiled-coil instabilities. Stutters are found in all the egg case proteins and were also identified in the capsular proteins of a related species, *Pugilina cochilidum*.⁶³ Critically, stutters have been found by comprehensive molecular dynamic simulations to be the preferred site for the initiation of the α - β transition in the IF protein vimentin^{64,65} and their presence is thus viewed as a necessary condition to induce the transition. Both mechanisms could possibly be concurrently active, but further experimental work is necessary before more definitive conclusions can be drawn.

4.4. Proposed Micromechanical Model of α - β Transition. Based on an equimolar amount of CP-1 and CP-3, the maximum end-to-end distance of the average capsule protein in the solid-state unstrained egg capsule is 93.3 or 91.0 nm, depending on whether the pre-existing β -sheets are parallel or antiparallel, respectively. When the capsule material is stretched, α -helices unravel and form β -sheets. If these newly formed β -sheets are parallel, the end-to-end lengths increase to 153.4 and 151.1 nm, corresponding to an increase of 64.4 and 66.0%, respectively. If the new β -sheets are antiparallel, the lengths would increase to 147.3 and 145.0 nm, which correspond to increments of 57.9 and 59.3%, respectively. All these values are close to the 60% post-yield engineering strain that was observed with the bulk egg capsule material. From the amino acid composition analyses, the average Lys content of CP-1 and CP-3 is \sim 3% higher than the bulk material. This difference has been thought to be due the formation of Lys-Lys cross-links when the immature egg capsule is processed under the ventral pedal gland (VPG) of the snail. Lys-Lys cross-links are resistant to acid hydrolysis and so would account for the lower Lys content in the mature capsule, as measured by amino acid analysis. As the distribution of Lys is fairly constant throughout both the primary sequences of CP-1 and CP-3 (Figure 11), the α -helices, β -sheets, and random coils in the mature egg capsule are, respectively, 1.5, 0.6, and 0.9% cross-linked on average. It has been hypothesized that disulfide linkages prevent unraveling of the 1B and 2B helical domains¹⁶ in the hard α -keratin materials, but compared to the high cross-link density of up to 15% in the hard α -keratins, the low cross-link density in the α -helical domains of the egg capsule is unlikely to have a huge effect on the unraveling and hence extensibility of the helices. Indeed, the extensibilities of the α -keratin materials and the egg capsule material are vastly different in terms of the rate and extent of recovery, which could very well be due to the stark differences in the nature and densities of the cross-links.

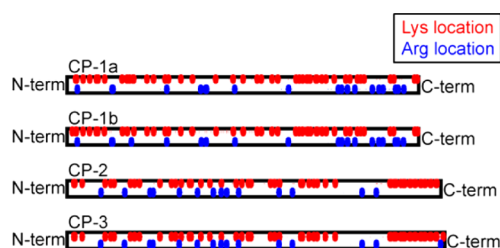


Figure 11. Distribution of Lys and Arg residues in the primary sequences of the Bc-CPs. There are distinct clusters of not only Lys but also Arg, which preliminary data suggests may contribute to cross-linking.

4.5. Egg Case Cross-Linking and Open Questions.

During the egg laying process, Bc-CPs first assemble into filaments in the immature egg capsule before they are cross-linked under the VPG.³² Similar to purified IF proteins in solution, the egg capsule filament assembly can occur in vitro without the aid of additional accessory proteins or molecules. Based on the similar dimensions of the fibers that are created in vitro, observed here by TEM, and those that occur naturally in the egg capsule, observed previously by SEM,¹⁷ the purified Bc-CPs form the filaments that the mature egg capsule material is composed of. While the secondary structure analyses by FTIR suggest that the egg capsule filament is composed of a self-assembling mixture of CP-1 and CP-2, separate solutions of CP-1 and CP-2 are equally capable of forming filaments. Further studies are thus required to determine if filaments of CP-1 or CP-2 are present in the mature egg capsule. On the other hand, IF proteins such as α -keratins are known to form heterodimeric coiled-coils in vivo but not in vitro unless both proteins are present in the solution.⁶⁶ Furthermore, as IF proteins are known to form coiled-coil tetramers, and sometimes octamers, in solution before assembling into filaments,^{66,67} a more thorough investigation into the self-assembly of the Bc-CPs will be necessary to ascertain whether they form coiled-coil multimers in solution prior to filament assembly.

After assembly, cross-linking occurs as the final processing step before the egg capsules mature and are deposited on the ocean floor. It is only after cross-linking that the egg capsules exhibit unique and robust mechanical properties. Previous work on both the egg capsule material⁴⁵ and the VPG where this final processing step occurs⁴⁶ suggested that the cross-links are Lys-derived. As seen in the amino acid composition analyses, all the Bc-CPs have high quantities of Lys which are quite fairly distributed throughout the length of the primary sequences. Based on the heptad repeat assignment by SCORER, Lys residues can be found in positions both within and outside the core of the coiled-coil. This translates to a possibility for interchain cross-links to form both between and within coiled-coils. In CP-2 and -3 specifically, there is a high proportion of Lys residues in the C-terminal regions. CP-2 and CP-3 have 53 and 54 Lys residues out of a total of 475 and 479 amino acid residues, respectively, but 20 and 21 of those Lys residues are located between position 413 and the C-terminus (Figure 11). These regions of the primary sequences seem to be responsible for end-to-end cross-linking of the Bc-CPs, but until a chemical structure of the cross-link is determined, nothing definitive can be said about Lys-based cross-links.

There is a variety of unanswered questions regarding the whelk egg capsule post-translation assembly and processing.

The mature capsule material is over 90% protein by dry weight, but the composition of the remaining mass is unknown. Furthermore, SDS-PAGE of the immature capsules and crude nidamental gland extract indicate that traces of additional proteins, other than the Bc-CPs, are present in this material. They are in far lower quantities, but their structure and function are yet to be investigated. They are potentially matrix proteins that bundles the egg capsule filament proteins into higher hierarchical fibrous structures. α -Keratin materials have globular, high-sulfur content matrix proteins, which contribute significantly to the mechanical properties,^{5,16} whereas hagfish slime does not.²⁸

At the moment, precise molecular-scale knowledge of the steps leading to the extended β -sheet conformation and their time-dependent mechanisms remain unknown. Elucidation of the crystal structures of the Bc-CPs would provide key information to further understand the solid-state phase transformation.

5. CONCLUSIONS

In this study, the precursors of the primary structural proteins of the whelk egg capsules were purified and characterized. Their amino acid compositions are very similar to IF proteins with the exception of Lys and Cys, which are significantly higher in the egg capsule and the IFs, respectively. CD and FTIR data indicate that, much like the IF proteins, the Bc-CPs are strongly α -helical in solution and are capable of self-assembling into filaments without the assistance of accessory molecules.

Using reverse transcription techniques, the primary amino acid sequences of 4 Bc-CPs were deduced from their cDNA. The primary sequences are unique as BLAST did not produce any significant match to known proteins. Analyses of these sequences indicated that the proteins are likely to form coiled-coils, which is consistent with previous X-ray scattering studies. In comparison to coiled-coil proteins from an established database, Bc-CPs appear to have a higher amount of Gly and smaller amounts of aromatic residues in their hydrophobic cores. Furthermore, the terminal regions of the sequences are densely populated by Lys and Arg, both of which are suspected to play a role in the covalent cross-linking of the egg capsule fibers.

This work opens further interesting questions, especially regarding self-assembly and the molecular mechanisms of the α - β transition, which is an emerging theme for a variety of bioelastomeric materials.¹¹ First, the prediction of a trimeric coiled-coil in the egg case capsule that is reminiscent of fibrin needs to be tested further. Second, there are uncertainties whether the coiled-coils are homo-oligomeric or hetero-oligomeric. Finally, the exact nature of the initiation of α - β transition, which is critical in driving the reversible transformation, remains as an intriguing question. ATR-FTIR data indicates a small, yet significant β -sheet content in both the purified proteins and in the mature capsule. These β -sheets "nuclei" could act as local sites where β -sheets further initiate their expansion. Alternatively, the transition could be initiated at coiled-coil instabilities such as the stutters. Elucidation of these mechanisms is currently underway in our laboratory using recombinant and site-directed mutagenesis of the precursor proteins.

■ ASSOCIATED CONTENT

■ Supporting Information

Table S1: N-termini Edman sequencing results of Bc-CPs from individual SDS-PAGE bands of crude extract. Figure S1: 1% agarose gel of RT-PCR with 3' RACE products. Figure S2: Complete amino acid sequences of (a) CP-1b and (b) CP-2, as deduced from cDNA. Figure S3: CD spectra of mixtures of CP-1 and CP-2 in various ratios. Figure S4: ATR-FTIR spectra of CP-1 and CP-3, showing the preferential formation of β -sheets over α -helices when the proteins are freeze-dried from concentrated, compared to diluted, solutions. Figure S5: (a) ATR-FTIR spectra of egg case, showing a loss of Amide II at $\sim 1525\text{ cm}^{-1}$ after hydrogen is substituted by deuterium; (b) curve-fitted Amide I of dehydrated egg case and (c) egg case in D_2O , confirming the presence of β -sheets along with high α -helical content (>50%). Figure S6: Relative amounts of (a) hydrophobic and (b) charged amino acid residues at various heptad positions in the predicted coiled-coil domains of Bc-CPs and other known coil-coiled proteins in the CC+ database. Figure S7: Transmission electron micrographs of purified Bc-CPs. This material is available free of charge via the Internet at <http://pubs.acs.org>.

■ Accession Codes

The sequence data of Bc-CP 1a, Bc-CP 1b, Bc-CP 2, and Bc-CP 3 have been deposited in the NCBI database with the accession numbers GU205809, GU205810, GU205811, and GU205812, respectively.

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■ Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

The authors would like to thank Dr. Krystyna Brzezinska (UCSB) for assistance with the CD spectrometry, Dr. Paul Guerette (NTU) for many useful discussions, Rebekah Ashley (UCSB) for assistance with cDNA sequencing, and Prof. Robert Shadwick (University of British Columbia, Vancouver, Canada) for insightful advice throughout the project. We thank Tan Kang Hong Delford and Loke Jun Jie for assistance with protein purification. This work was supported by a University of California Biotechnology Research and Education Program (UC BREP) Graduate Research and Education in Adaptive Biotechnology (GREAT) Grant (#2007-02), an NIH Grant #R01DE018468 (J.H.W.), and the Singapore National Research Foundation (NRF) through a NRF Fellowship (A.M.). The work made use of MRL Central Facilities supported by the MRSEC program of the NSF under Award #DMR05-20415. C.N. and A.S. acknowledge the partial support of the Austrian Federal Ministry for Transport, Innovation and Technology (GZ BMVIT-612.166/0001-III/I1/2010) and the ZIT (Center of Innovation and Technology of Vienna).

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