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Elastic proteins and elastomeric protein alloys Behnaz Aghaei-Ghareh-Bolagh^{1,2,3}, Suzanne M Mithieux^{1,2,3} and Anthony S Weiss^{1,2,3}

The elastomeric proteins elastin and resilin have been used extensively in the fabrication of biomaterials for tissue engineering applications due to their unique mechanical and biological properties. Tropoelastin is the soluble monomer component of elastin. Tropoelastin and resilin are both highly elastic with high resilience, substantial extensibility, high durability and low energy loss, which makes them excellent candidates for the fabrication of elastic tissues that demand regular and repetitive movement like the skin, lung, blood vessels, muscles and vocal folds. Combinations of these proteins with silk fibroin further enhance their biomechanical and biological properties leading to a new class of protein alloy materials with versatile properties. In this review, the properties of tropoelastin-based and resilin-based biomaterials with and without silk are described in concert with examples of their applications in tissue engineering.

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Current Opinion in Biotechnology 2016, 39:56–60

This review comes from a themed issue on Nanobiotechnology

Edited by Michael Nash and Oded Shoseyov

<http://dx.doi.org/10.1016/j.copbio.2015.12.020>

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Introduction

An elastomer is a rubber-like material with high resilience, extensibility and effective energy storage [\[1,2](#page--1-0)]. Elastomeric materials undergo large elastic deformations, and energy used for their deformation can be recovered after removal of the applied force [[2\]](#page--1-0). There are multiple elastomeric proteins in nature including elastin, resilin, wheat gluten, abductin and flagelliform silk but of these elastin is the only mammalian elastomer. These materials have been reviewed previously [[1,3,4](#page--1-0)]. Pure elastin comprises assembled tropoelastin monomers. In this review we focus on tropoelastin and resilin as elastomeric proteins that have been extensively investigated for the fabrication of materials for tissue

engineering applications. Tropoelastin-silk and resilinsilk composite materials are also reviewed as an emerging, versatile class of biomaterials with tunable mechanical and biological properties for engineering of multiple tissue types.

Tropoelastin

Tropoelastin is the soluble protein precursor of elastin. Elastin, a highly insoluble, durable and long lasting protein polymer, is responsible for the elasticity of different tissues in the body including, skin, blood vessels and lung [\[5](#page--1-0)–7]. Elastin's monomer tropoelastin is a biologically active molecule secreted by elastogenic cells after which it self-assembles and organizes in association with a fibrillin-rich microfibrillar structure to form elastin [\[8,9](#page--1-0)]. Synthetic elastin does not require the microfibrillar component to form as it is fabricated. Tropoelastin is composed of alternating hydrophobic and hydrophilic domains. At low temperatures tropoelastin is a soluble monomer. With increasing temperature, the hydrophobic domains interact such that the molecules begin to aggregate into a non-soluble viscoelastic phase termed a coacervate. The coacervate is an early step towards elastin fibre formation [\[6,7\]](#page--1-0). These coacervated tropoelastin molecules then align with each other. Covalent crosslinking of hydrophilic domains, particularly those rich in Lys and Ala ensues resulting in the formation of a stabilized mature elastic protein polymer network [[6,8,10](#page--1-0)]. The rubber-like properties of elastin are due to repetitive modules such as VPGXG [[3,11](#page--1-0)]. With a Young's modulus of \sim 3 kPa, the tropoelastin molecule is significantly more elastic than other extracellular molecules [[9,12](#page--1-0)]. Naturally cross-linked tropoelastin has a Young's modulus of 300– 600 kPa [[9\]](#page--1-0). Tropoelastin can be extended to 8 times of its resting length in a fully reversible process [\[12](#page--1-0)] due to the dynamic nature of the hydrophobic domains and the reversible organization of solvent molecules around these domains [\[13](#page--1-0)]. This interaction with water confers an entropic mechanism of elasticity, where stretching elastin decreases bulk entropy due to the increased organization of water around these domains; thus the maximum entropic level is restored when the protein recoils back to its original state [\[7,12\]](#page--1-0). The glass transition temperature (T_{σ}) of elastin depends on its hydration. Dehydrated elastin has T_g of 200 °C whereas 30% hydration decreases T_g to $30 °C$ [\[14](#page--1-0)].

Tropoelastin is produced by bacterial overexpression [\[15](#page--1-0)]. Recombinant tropoelastin is used to generate materials that restore the function of damaged or diseased

elastic tissues. The fabrication of elastin-based biomaterials generally involves coacervation and crosslinking of the precursor molecule. For example, tropoelastin proceeds to gel-sol transition under alkaline conditions, where tropoelastin forms a stable hydrogel without the need for further chemical crosslinking [[16\]](#page--1-0). There are multiple alternative paths to stabilize tropoelastin hydrogels including chemical, physical, pH-based, photochemical and enzymatic crosslinking, to ensure its stability, durability and utility as a biomaterial for tissue engineering. Tropoelastin assemblies have been used in the fabrication of a range of biomaterials in diverse forms and shapes including sponges, hydrogels, electrospun mats and tubes [17–[24\]](#page--1-0).

Tropoelastin-based hydrogels can be prepared by chemical crosslinking using bis-sulfosuccinimidyl suberate (BS3) and glutaraldehyde [[18,25,26](#page--1-0)] that benefit from the availability of 35 juxtaposed lysine residues acrossthe molecule. Physical crosslinking has been used to produce elastic hydrogels. For example, methacrylated tropoelastin is formed through photo-initiated cross-linking. Using this approach, a highly stretchable and porous hydrogel with a high extensibility of 400% and tensile modulus of 2.8 ± 0.6 to 14.8 ± 1.9 kPa is produced [[19](#page--1-0)]. This method allows for the fabrication of a range of biomaterials with different pore size, swelling and mechanical performance based on tropoelastin concentration and degree of methacrylation [\[19\]](#page--1-0). Methacrylated tropoelastin has also been used to generate micropatterned elastin-based hydrogels suitable for engineering of tissues like cardiac tissues where elasticity and organization of cells plays a central role $[27\degree]$ $[27\degree]$ $[27\degree]$. In contrast, the solubility of tropoelastin in organic solvents like hexafluoroisopropanol facilitates the development of 3-dimensional elastomeric biomaterials using electrospinning techniques. Electrospun elastic mats and tubes made of tropoelastin have been stabilized chemically by crosslinkers selected from hexamethylene diisocyanate $[20-22]$ $[20-22]$, glutaraldehyde $[18,23]$ $[18,23]$ $[18,23]$ and disuccinimidyl suberate [[24](#page--1-0)].

These tropoelastin-based constructs not only provide mechanical integrity and elasticity to multiple tissue types but also confers biological interactions including cell binding and signalling cues [[5,17](#page--1-0)]. They can recruit and support the growth of diverse attachment cell types including dermal fibroblasts [[18\]](#page--1-0), stem cells [\[28](#page--1-0)] and cardiomyocytes $[27^{\bullet\bullet}]$ $[27^{\bullet\bullet}]$ $[27^{\bullet\bullet}]$. On this basis, tropoelastin-based constructs have been investigated for engineering of vascular grafts, wound dressing materials, skin replacement and stem cell delivery for wound healing [[18,20](#page--1-0)– [24\]](#page--1-0). Tropoelastin and elastin-based materials have been recognised as non-thrombogenic and non-inflammatory biomaterials [\[29](#page--1-0)]. Bio-functionalization of metallic coronary stents with tropoelastin decreases thrombogenic responses to implanted stents [[30\]](#page--1-0).

Resilin and resilin-like polypeptides (RLPs)

Resilin, first described in locusts by Weis-Fogh, is present in the tissues of insects that require rapid elastic responses [[3,31,32](#page--1-0)]. Thus resilin is important for insect jumping [\[33](#page--1-0)] and flight [[32\]](#page--1-0) and in organs that demand highly repetitive movement such as their specialized sound-producing organs. For this reason, resilin responds elastically at higher frequencies than elastin; in contrast, elastin typically operates around the heartbeat frequencies of mammals. The flexibility of resilin is due to the presence of repetitive amino acid sequences rich in proline and glycine. Natural resilin is cross-linked by the formation of diand tri-tyrosine between tyrosine residues connecting resilin polypeptides [\[13,34,35\]](#page--1-0).

Resilin is a highly elastomeric, rubber-like protein with high elasticity and resilience [[13,31](#page--1-0)]. The cross-linked protein can be extended up to 3–4 times its size and can return to its original state [\[36](#page--1-0)]. Natural resilin has an elastic modulus of 600–2000 kPa [[32,37,38\]](#page--1-0). Like elastin, resilin is brittle and stiff when dry but after hydration, it becomes soft and elastic. The elasticity of resilin is directly proportional to the level of hydration [[39\]](#page--1-0). Resilin is stable when heated up to $125\textdegree C$ and can withstand vibrations with the frequency of 4 kHz [[3,32,40\]](#page--1-0). It is insoluble in water and nonpolar solvents [[32\]](#page--1-0).

The resilin precursor CG15920 in Drosophila melanogaster consists of a signal peptide sequence and three main exons [\[41](#page--1-0)]; after removal of the signal peptide, the proresilin is secreted to the extracellular space. Pro-resilin is uncross-linked and is composed of three main domains, N-terminal and C-terminal elastic domains encoded by exon 1 and exon 3 respectively and the chitin-binding domain encoded by exon 2 [[34,41,42\]](#page--1-0). Recombinant production of resilin regions in bacteria yields 15– 450 mg of protein [[34\]](#page--1-0). Proteins produced by exon 1 showed higher elasticity than those produced from exon 3 and better mimic the elastic properties of the natural resilin [[42\]](#page--1-0). Rec1-resilin (N-terminal domain of resilin) encoded by exon 1 of *D. melanogaster* is a water-soluble protein that has been produced in *Escherichia coli*. This protein can be cross-linked by formation of di-tyrosine crosslinking either chemically (peroxidase) or physically (ruthenium-mediated photo-crosslinking) resulting in stable elastomeric biomaterials. The cross-linked material is resilient and can be stretched up to 300% of its original length [\[13,39\]](#page--1-0). A photo-Fenton reaction is also used to crosslink the proteins produced by exon 1 or exon 3 of the CG15920 gene, which results in a rubber like and highly adhesive materials [[42\]](#page--1-0).

In order to improve the biological properties of resilin, RLPs are produced by the addition of other sequences to the resilin consensus sequences. Charati et al. has reported RLP12, which contains 12 repeats of resilin consensus sequences from *D. melanogaster*, additional Download English Version:

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