Supramolecular Modification of a Sequence-Controlled Collagen-Mimicking Polymer

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ABSTRACT: Structurally and functionally well-deﬁned recombinant proteins are an interesting class of sequence-controlled macromolecules to which different crosslinking chemistries can be applied to tune their biological properties. Herein, we take advantage of a 571-residue recombinant peptide based on human collagen type I (RCPhC1), which we functionalized with supramolecular 4-fold hydrogen bonding ureido-pyrimidinone (UPy) moieties. By grafting supramolecular UPy moieties onto the backbone of RCPhC1 (UPy-RCPhC1), increased control over the polymer structure, assembly, gelation, and mechanical properties was achieved. In addition, by increasing the degree of UPy functionalization on RCPhC1, cardiomyocyte progenitor cells were cultured on “soft” (~26 kPa) versus “stiff” (~68–190 kPa) UPy-RCPhC1 hydrogels. Interestingly, increased stress ﬁber formation, focal adhesions, and proliferation were observed on stiffer compared to softer substrates, owing to the formation of stronger cell–material interactions. In conclusion, a bioinspired hydrogel material was designed by a combination of two well-known natural components, i.e., a protein as sequence-controlled polymer and UPy units inspired on nucleobases.

INTRODUCTION

The extracellular matrix (ECM) acts as a natural scaffold for cells and is important for the regulation and maintenance of cellular fate in terms of tissue development and pathology.1–3 Many researchers have studied the inﬂuence of different ECM properties and the effect on cellular behavior for improved cell-based tissue regeneration.4 Hydrogels are a class of biomaterials that resemble natural tissues with respect to their viscoelastic behavior. As such, hydrogels are commonly used as instructive microenvironments for cells.5–7 Hydrogel-based biomaterials should display (i) optimal physical and mechanical properties that mimic the in vivo environment, (ii) sufﬁcient adhesion sites to allow cells to adhere and migrate, and (iii) allow for cell-induced degradation.8 In general, naturally derived hydrogels or materials, such as matrigel,9 fibrin,10 or collagen,11 are used. These hydrogels contain high degrees of bioactive sites and therefore seem inexorable for the development of natural matrices.12,13 However, these naturally based hydrogels have low controllability and often contain mixtures of different components and/or polymers with different lengths. Moreover, the synthesis of sequence-controlled synthetic polymers remains challenging in the field of polymer chemistry.14

It is proposed that recombinantly produced proteins are ideal candidates to develop biomaterials with full control of bioactive properties and polymer sequence, length and structure.15–18 This is achieved using genetically modiﬁed organisms such as yeast or bacteria.19 Additionally, protein-engineered biomaterials can be further functionalized to tune the biological and mechanical properties. Heilshorn et al., developed elastin-like polypeptide (ELP) hydrogels, in which the stiffness of the hydrogel network and arginine–glycine–aspartate (RGD) ligand density could be tuned independently.20 This was done using tris(hydroxymethyl) phosphine (THP) to cross-link the poly(ethylene glycol) bis(amine) linker with speciﬁc amines located on the ELP. By changing the THP concentration, different storage moduli were obtained, ranging from 0.01 to 2.5 kPa. Encapsulated ﬁbroblast cells showed more spread morphologies on the softer and compliant gels (~1.3 kPa) compared to stiffer gels (~2.5 kPa), as a result of the smaller hydrogel mesh size and increased crosslink density.20 Another example of protein-engineered hydrogels is based on recombinant silk-based polypeptides.18 Repeating units of glycine, alanine, and serine, in the backbone of silk-based polypeptides, induce the formation of β-sheets and consequently hydrogel networks can be formed.21 With the high degree of control in molecular weight, polarity, and mechanical properties, recombinantly synthesized silk-based polypeptides are suitable to create biocompatible protein-engineered hydrogels.22–24 These examples illustrate the

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fit of recombinantly synthesized hydrogels for regenerative medicine purposes.

Recombinant peptides based on human collagen type I (RCPhC1), commercially known as Cellnest (FujiFilm Manufacturing Europe BV), are protein-engineered macro-molecules with controlled lengths and specific amino acid sequences. Moreover, repeating amino acid sequences based on the integrin-binding peptide, RGD, have been engineered into the backbone of RCPhC1, to enhance cellular adhesion.25 Another advantage of the application of this recombinant collagen as the polymer is due to its defined length, controlled sequence, and low immunologic risk.26 RCPhC1 was previously used as a scaffold for the delivery of adipose-derived cells,25 as a synthetic extracellular matrix for the development of bone tissue,27,28 and as a delivery vehicle for bone-morphogenic protein-2.29,30 Depending on the application or cell type to be cultured, the RCPhC1 sequence can be designed and adapted with different cross-links, e.g., methacrylates,31 to create protein-engineered hydrogels.

In this study, we used RCPhC1 as the polymer backbone for the introduction of supramolecular 2-ureido-4[1H]-pyrimidine (UPy) hydrogen bonding units, to achieve control over hydrogel formation, mechanical properties, and cellular response. Supramolecular UPy moieties were previously used to modify different synthetic polydisperse polymers such as hydrophilic poly(ethylene glycol) or polyesters such as polycaprolactone and hydrophobic amorphous prilast polymers, which illustrates the versatility of UPy-based supramolecular biomaterials.32–34 Compared to covalently cross-linked hydrogels, UPy-modified hydrogels allow for the possibility to tune the material properties, such as stiffness, degradability, and bioactivity. Here, the degree of UPy substitution onto the backbone of RCPhC1 was varied and the structure, molecular assembly, and gelation are studied and characterized. By increasing the amount of grafted UPy groups, it is hypothesized that the stiffness of the hydrogel is increased. The applicability of UPy-modified RCPhC1 hydrogels as the two-dimensional cell culture matrix is tested using human cardiac progenitor cells (CPCs).35 These cells are ideal candidates for cardiac regeneration applications due to their self-renewal, ECM production capacity, and potential to differentiate into cardiomyocytes.36–38 Finally, the relationship between molecular interactions and assembly versus the observed cellular response is discussed.

■ EXPERIMENTAL SECTION

Materials. Chemicals and reagents were purchased from commercial sources at the highest purity available and used without further purification. All solvents were purchased from Sigma-Aldrich unless stated otherwise. Cellnest, a recombinant peptide based on human collagen type I (RCPhC1), was a gift from Fujifilm.
Manufacturing Europe B.V. and was used without further purification (Figure S1). Phosphate buffered saline (PBS) tablets were purchased from Sigma-Aldrich (pH 7.2–7.6). Trypsin–ethylenediaminetetra-acetic acid (EDTA) solution was purchased from Sigma (0.5 g/L porcine trypsin and 0.2 g/L EDTA in Hank’s balanced salt solution with phenol red). All compound concentrations were determined by weight.

Methods. Analytical Techniques and Equipment. Reverse-phase high-performance liquid chromatography—mass spectrometry (RP-HPLC–MS) was performed with a Thermo scientific LCQ fleet spectrometer. Proton nuclear magnetic resonance (1H NMR) spectra were recorded on a 400 MHz NMR (Varian Mercury Vx or Varian 400MR) operating at 400 MHz to determine the degree of functionalization. Proton chemical shifts are reported in ppm downfield from tetramethylsilane using the resonance of the deuterated solvent as the internal standard. Samples for NMR were prepared in D2O/KOD with pH ≈ 11 at a concentration of 25 mg/mL. The purity of RCPHC1 and UPy-RCPHCl derivatives was determined with Waters Xevo G2 Quadrupole Time-Of-Flight liquid chromatography—mass spectrometry equipped with an Agilent Polaris C18A reverse-phase column (12.0 mm, length 100 mm). Derivatives were dissolved in H2O (0.5 mg/mL) and flowed (0.3 mL/min) over the column using a 15–75% water/acetonitrile gradient with 0.1% formic acid prior to analysis in the positive electrospray ionization (ESI) mode in the mass spectrometer. Dynamic light scattering (DLS) was performed on a TA Instruments Discovery Hybrid 3 shear rheometer with a 25 mm Sandblasted Peltier plate. To prevent water evaporation, an oil trap based on silicone oil (Rhodorsil) was used with a 25 mm Sandblasted Peltier plate. To prevent water evaporation and concentrated under vacuum to yield 1.01 g (1.69 mmol, 99%) of a white waxy solid. 1H NMR (CDCl3/MeOD) δ (ppm): 5.86 (s, 1H), 3.28 (2H, 2H), 3.11 (m, 6H), 2.26 (s, 3H), 1.59 (m, 2H), 1.44 (s, 1H), 1.36 (m, 4H), 1.25 (m, 18H) (Figure S3). LC–MS (ESI) m/z calculated [C15H26N2O4] 393.43; found 394.42 [M + H]+.

Synthesis of 12-(Ureido-pyrimidinone-hexyl-urea)-1-dodecyl-boc (2). A round bottom flask was charged with ureido-pyrimidinone-hexyl-isocyanate (499 mg, 1.70 mmol, 1 equiv), monoprotonated amine 1 (314 mg, 1.71 mmol, 1.01 equiv) and dissolved in CHCl3 (30 mL) and MeOH (10 mL), N,N-Diisopropylethylamine (0.5 mL, 2.91 mmol, 1.7 equiv) was added to the stirring solution. After 4 h of stirring, the reaction mixture was purified on a short plug of silica eluting with CHCl3/MeOH (1:1) to remove excess of amine 1. The solvents were removed by rotary evaporation and concentrated under vacuum to yield 1.01 g (1.69 mmol, 99%) of 2, 5, and 8 UPy derivatized hydrogels were reconstituted in 10 wt/vol % (100 mg/mL) hydrogels were reconstituted in PBS tablets with pH = 12.7 (for 2, 5, and 8 UPy functionalities between 0.7–10 wt/vol %). Different amounts of grafted UPy functionalities between 0.7–10 wt/vol % were used to prepare rates of drug release.

Table 1. Degree of Functionalization of the UPy-RCPHCl Derivatives Based on 1H NMR

| UPy-RCPHCl derivative | RCPHCl- NH2/UPy-CDI molar ratio | integral at 5.7–5.8 ppm (%) | integral at 0.8–1.0 ppm (%) | theoretical molecular weight (kg/mol) |
|-----------------------|---------------------------------|----------------------------|-----------------------------|-------------------------------------|
| UPy-RCPHCl-12         | 1:0.25                          | 85.6                       | 264                         | 52.2                                |
| UPy-RCPHCl-5          | 1:0.5                           | 90.4                       | 264                         | 53.8                                |
| UPy-RCPHCl-8          | 1:1                             | 99.3                       | 264                         | 55.4                                |
| UPy-RCPHCl-12         | 1:1                             | 81.2                       | 264                         | 57.4                                |
| UPy-RCPHCl-16         | 1:1                             | 77.4                       | 264                         | 59.5                                |
the different UPy-RCPhC1 polymers, different pH values were used to improve solubility. To readjust the pH and increase UPy–UPy association, r-glucono-δ-lactone (GDL) powder was mixed with the UPy-RCPhC1 solution. An amount of 10 mg/mL (for 2, 5, and 8 UPy functionalities) or 20 mg/mL (for 12 and 16 UPy functionalities) of GDL powder was added for each hydrogel. Due to the hydrolysis of GDL to gluconic acid, the pH slowly reduces to pH ~5 – 6 after 24 h and hydrogels are formed.

**Dynamic Light Scattering.** Samples for DLS measurements were prepared by dissolving 2 mg/mL of each compound in PBS and annealing for 1 h at 70 °C. The samples were filtered before measurement with a Whatman poly(vinylidene difluoride) filter, 0.45 μm. The temperature was set at 20 °C and decreased to 5 °C with steps of 5 °C, equilibrating for 500 s before measuring.

ζ-Potential. The samples were dissolved in 4-(2-hydroxyethyl)-1-piperazinethanesulfonic acid (1 mM, pH ~7.6) at a concentration of 0.1 mg/mL and filtered with a 0.45 μm collagen filter. A DTS1070 cuvette was used for measuring the ζ-potential. The samples were measured in triplo, at room temperature, with a 30 s equilibration time. Measurement duration was automated and automatic attenuation and voltage selection were turned on.

**Circular Dichroism.** Samples for CD measurements were prepared by dissolving 0.5 mg/mL of the compounds in ultrapure water and stirring for approximately 30 min. The higher functionalized compounds (UPy-RCPhC1-1, UPy-RCPhC1-16) were annealed for 30 min at 30 °C. Measurements were performed with a scan speed of 100 nm/min, data pitch of 0.25 nm, a response time of 0.5 s, a bandwidth of 2, and a path length of 0.1 cm. The spectra were measured from 170 to 300 nm. The signal was processed with "Adjacent Averaging", 5 points of window. Experiments were conducted at a concentration of 0.5 mg/mL in a 0.1 cm Hellma quartz cell. For each measurement, the temperature was increased initially to 60 °C and decreased with steps of 1 °C/min to 5 °C. The molar residual ellipticity was determined using the following equation:

$$[\theta] = \frac{\theta \times m}{c \times l / n_i}$$

where θ is the ellipticity in millidegrees, m is the molecular weight in g/mol, c is the concentration in mg/mL, l is the path length in cm, and n_i is the number of amino acids in the peptide. Graphs are shown from 180 to 280 nm, due to the higher noise ratio observed below 200 nm and due to the wavelength absorption of water.

**Cryogenic Transmission Electron Microscopy.** Cryogenic transmission electron microscopy was performed using samples with a concentration of 0.5 or 5 mg/mL. Vitriﬁed ﬁlms were prepared using a computer-controlled vitriﬁcation robot (FEI Vitrobot Mark III, FEI Company) at 22 °C, and at a relative humidity of 100%. In the preparation chamber of the "Vitrobot", 3 μL sample was applied on a LaceY ﬁlm (LC200-CU, Electron Microscopy Sciences). These ﬁlms were surface plasma treated just prior to use, with a Cressington 208 carbon coater operating at 5 mA for 40 s. Excess sample was removed by blotting using ﬁlter paper for 3 s at ~3 mm, and the thin ﬁlm thus formed was plunged (acceleration about 3 g) into liquid ethane just above its freezing point. Vitriﬁed ﬁlms were transferred into the vacuum of a Cryo-ETTAN equipped with a ﬁeld emission gun that was operated at 300 kV, a post-column Gatan energy ﬁlter, and a 2048 × 2048 Gatan CCD camera. Micrographs were taken at low dose conditions, starting at a magniﬁcation of 6500 times with a deconvoluted setting of 40 μm, and at a magniﬁcation of 24 000 times with deconvoluted settings of 10 and 15 μm. The sizes of the observed micelles were measured manually using Fiji software.

**Nanoindentations.** Nanoindentation tests were applied to measure the mechanical properties of UPy-RCPhC1 hydrogels, which is based on a probe that is in contact with the surface of the material and is pushed through. Depending on the load (P), spherical tip radius (R), displacement (h), and stiffness of the cantilever, the effective Young’s modulus (E_{eff}) can be determined via the following formula considering the Hertzian contact model:

$$E_{eff} = \frac{P \times 3/4}{\sqrt{R_i \times h_0^{3/2}}}$$

where the effective Young’s modulus is derived from a certain percentage of the elastic–plastic loading regime, displacement h₀, of the loading curve. Hydrogels with volumes of 50 μL were formed in the cap of a 200 μL Eppendorf tube, which was glued inside a small Petri dish. Before measuring the mechanical properties, the Petri dishes were filled with serum-free media (M199, Gibco) to cover the hydrogels. Indentations were performed using a cantilever with a stiffness of 0.53 N/m and a tip radius of 24 μm or a cantilever with a stiffness of 4.71 N/m and a tip radius of 21 μm for hydrogels based on UPy-RCPhC1-5/UPy-RCPhC1-8 and UPy-RCPhC1-12/UPy-RCPhC1-16, respectively. Calibrations were performed before measurements and indentation profiles were identical for each hydrogel. The effective Young’s modulus (E_{eff}) was determined using the Hertzian contact model which is fitted through 20–60% of the loading curves using DataViewer (Piuma, OPTICS11).

**Rheological Analysis.** Hydrogels were prepared as stated previously. Following the addition of GDL, the gelation was followed at a constant shear rate of 1 rad/s, strain amplitude of 1%, and temperature of 25 °C for 1 h until complete gelation. Temperature sweeps were performed from 45 to 5 °C with increments of 4 °C and at each temperature step, hydrogels were soaked for 5 min to allow equilibration of temperature and rearrangement of molecules.

**Cell Experiments.** L929 cardiomyocyte precursor cells (CPCs) were immortalized by lentiviral transduction of I-Tert and BMI-1 (L9TB).35 CPCs were cultured in SP+ growth medium containing M199 (Gibco), which uses a bicarbonate buffer system, and EGM-2 BulletKit (Lonza) in a 3:1 volume ratio, supplemented with 10% fetal bovine serum, 1% penicillin/streptomycin (Lonza), and 1% nonessential amino acids (Gibco) at physiological pH. CPCs were routinely cultured on 0.1% gelatin-coated PS, passed at 80–90% confluence and seeded at a concentration of 3.1 × 10^5 cells/cm². Hydrogels were washed with SP+ before seeding cells to remove any excess of gluconic acid.

**Immunofluorescence Staining.** CPCs cultured on UPy-RCPhC1 hydrogels were first washed with PBS, fixed in 3.7% formaldehyde (Merck) for 10 min, washed twice with PBS, and permeabilized with 0.5% Triton X-100 (Merck) for 10 min. Non-specific binding of antibodies was minimized by incubating in 2% horse serum in PBS for 20 min. Cells were then incubated with primary antibodies in 10% horse serum in PBS for 2 h at 4 °C. Subsequently, the cells were first washed with PBS and incubated with a secondary antibody and phalloidin–FITC for 1 h in PBS followed by incubation with 4′,6-diamidine-2-phenylindole (0.4 μg/mL) in PBS for 5 min. Finally, the samples were washed and mounted on cover glasses with Mowiol (Sigma). Information regarding primary and secondary antibodies are listed in the Supporting Information (Table S1). The samples were imaged with a confocal laser scanning microscope (Leica TCS SP5X).

**Cell Proliferation Assay.** CPCs cultured on UPy-RCPhC1 hydrogels were first washed with PBS and the culture plate was frozen at ~80 °C. A commercially available CyQuant Assay was used to measure the ﬂuorescence of a dye that binds to nucleic acids. Based on a standard curve of known cell numbers (Figure S21), the fluorescence could be translated into cell numbers for each sample.

**Statistical Analysis.** Data are presented as mean ± standard deviation (SD). These data consisted of nanoindentations, cell numbers, and particle size measured with cryo-transmission electron microscopy (TEM). All statistical differences were determined using a nonparametric Kruskal–Wallis test with Dunn’s post hoc test. Probabilities of p < 0.05 were considered as significantly different. All statistical analyses were performed using GraphPad Prism 5 Software (GraphPad Software, Inc.).

## RESULTS AND DISCUSSION

**Synthesis and Characterization of UPy-RCPhC1 Derivatives.** A 1,1′-carbonyldimidazole (CDI)-activated UPy synthon composed of a urea group and a 12-carbon
alkyl spacer 4 was designed and can be easily reacted with the RCPbCl protein. The reaction involving CDI is fast and selective and thereby circumvents the need for any catalyst.41

First, 1,12-dodecadiamine was protected with a boc-group yielding 1 (yield = 46%). Subsequently, 1 was reacted with UPy-hexyl-isocyanate, resulting in 2 (yield = 99%). Next, compound 2 was deprotected yielding amine-terminated UPy synthon 3 (yield = 78%). The CDI-activated UPy synthon 4 was synthesized by reacting amine-terminated UPy group 3 with CDI in the presence of a base. Finally, the CDI-activated UPy synthon 4 was reacted with nucleophilic amines on RCPbCl, which resulted in the formation of an additional urea group (Scheme 1).3H NMR was used to determine the amount of grafted UPy functionalities were observed, which are likely due to the presence of impurities in the CDI-activated UPy synthon 4 (Table 1). With this approach, a library with different degrees of UPy substitutions, i.e., UPy-RCPbCl-1, UPy-RCPbCl-5, UPy-RCPbCl-8, UPy-RCPbCl-12, and UPy-RCPbCl-16, was achieved by varying the ratio of 4 to the amount of RCPbCl (Figure S13). Based on RP-HPLC, broadening of the product is observed compared to pristine RCPbCl, which indicates increased polydispersity and multiple degrees of functionalization for UPy-RCPbCl compounds (Figure 1C). Additionally, an increase in retention time is observed, which is likely due to the presence of an increased amount of hydrophobic alkyl spaced UPy functionalities (Figure 1C). Higher retention time was observed for UPy-RCPbCl-12 compared to UPy-RCPbCl-16, which could be due to the collapse of UPy-RCPbCl-16 in H2O and a decrease in the interaction with the column of the chromatogram. The molecular weights of the UPy-RCPbCl derivatives were determined using mass spectrometry for UPy-RCPbCl-1, -5, and -8 (Figures S8–S10). Due to the high degree of UPy conjugation and decrease of protonated amines, it was

Figure 1. Characterization of supramolecular UPy-modified recombinant collagen peptide derivatives. (A) 1H NMR graphs showing characteristic peaks of the protons on alanine residues (δ-shift = 0.7–0.9 ppm), which are used as the reference, and peaks of the alkylidene proton of the UPy groups (δ-shift = 5.7–5.8 ppm) for RCPbCl and each UPy-RCPbCl derivative. (B) Schematic representation of protons used to determine UPy conjugation, which is the alkylidene proton of the UPy group (UPy-H) and the protons on the methyl group on alanine groups (Ala-CH3). (C) Chromatogram of UPy-RCPbCl derivatives in H2O (arrow indicates an increase in UPy grafting).
impossible to determine the molecular weights of UPy-RCPhC1-12 and UPy-RCPhC1-16 (Figures S11 and S12). Nevertheless, a library containing different degrees of UPy substitutions was successfully synthesized (Figure S13). Additionally, due to the presence of grafted UPy functionalities, the supramolecular assembly can be studied based on the combination of hydrogen bonding and hydrophobic interactions (Figure S13).

Structure of RCPhC1 and UPy-RCPhC1 Derivatives.
RCPhC1 is a sequence controlled, monodisperse polymer with a molecular weight of 51 kDa (Figure S7). In PBS solution, it assembles into particles with a size of 6.9 ± 1.0 nm, observed with Cryo-TEM (Figures 2A,C and S14). UPy-RCPhC1-8 shows a similar particle size of 6.6 ± 1.0 nm (Figures 2B,D and S14). These results show that the conjugation of at least 8 UPy groups to RCPhC1 does not have a large influence on the structural properties of pristine RCPhC1.
It is known that RCPCh1 does not form organized triple helical structures typically seen for natural collagen type I. However, due to the presence of proline residues in the amino acid sequence of RCPCh1, some secondary structures, categorized as "random coils", are formed similar to gelatin and can be detected with circular dichroism (CD). Secondary structures formed by pristine RCPCh1 show a minimum at ~195 nm and a maximum at ~220 nm (Figure 3A). Upon cooling RCPCh1 to 5 °C, intermolecular interactions, based on hydrogen bonding, ionic and hydrophobic interactions, are stabilized and result in an increased CD effect, which is in agreement with other collagen-based peptides or proteins found in the literature (Figure 3A). Moreover, after conjugating UPy groups to RCPCh1 a typical "random coil" structure and a small shift of the minima was observed (UPy-RCPCh1-2: ~198 nm; UPy-RCPCh1-5: ~202 nm; UPy-RCPCh1-8: ~200 nm; UPy-RCPCh1-12: ~198 nm; UPy-RCPCh1-16: ~205 nm) (Figure 3B–F). This could simply be the result of small variations in the CD spectrum for the different UPy-RCPCh1 polymers, which indicate minimal differences in the secondary structure following covalent conjugation of UPy functionalities to residual amines. Interestingly, an increase of the CD effect was observed for all UPy-RCPCh1 derivatives upon cooling to 5 °C (Figure 3A–F). This effect illustrates that enhanced CD effects as a result of cooling and stabilization of intermolecular interactions are maintained following UPy conjugation. Moreover, functionalization of RCPCh1 with 2, 5, and 8 UPy groups resulted in an increase of the maxima at ~220 nm, which was higher compared to pristine RCPCh1 (at 220 nm; RCPCh1: [θ] = 0.59; UPy-RCPCh1-2: [θ] = 1.27; UPy-RCPCh1-5: [θ] = 8.65; UPy-RCPCh1-8: [θ] = 5.14) (Figure 3A–D). It is proposed that a stabilization effect occurs as a result of UPy–UPy interactions, which enhances the CD effect and conceivably the secondary structure of RCPCh1.

Next, particle size distribution and information related to the aggregation of RCPCh1 and UPy-RCPCh1 derivatives were studied with dynamic light scattering (DLS) (Figure 3A–F and Table 2). Pristine RCPCh1 had a particle size of 14.5 ± 1.1 nm, which is larger than the size observed with cryo-TEM (Figures 3A' and 2A), an effect that is commonly observed and likely due to the hydration shell of the proteins. No difference in particle size was observed upon cooling pristine RCPCh1 to 5 °C, which was not expected due to the increase in the CD effect, which was observed upon cooling as a result of stabilized intermolecular interactions. However, longer incubation time periods resulted in an increase in the particle size which indicates aggregation of RCPCh1 particles via weak intermolecular interactions (data not shown).

Functionalization of RCPCh1 with UPy groups resulted in a small increase in hydrodynamic particle size for UPy-RCPCh1-2 (21.3 ± 0.3 nm), UPy-RCPCh1-5 (21.6 ± 0.1 nm), and UPy-RCPCh1-8 (21.6 ± 0.1 nm) and similar particle size for UPy-RCPCh1-12 (16.1 ± 0.1 nm) and UPy-RCPCh1-16 (15.2 ± 0.4 nm) (Table 2). Similar to pristine RCPCh1, cooling of UPy-RCPCh1-12 and UPy-RCPCh1-16 to 5 °C did not change the particle size and dispersity index (Figure 3A′,E,F). For UPy-RCPCh1-2, 5, and 8, a clear increase in the hydrodynamic particle size and dispersity index is observed upon cooling the samples to 5 °C (Figure 3B′–D′). An increase of the dispersity index is the result of an increased dispersity of particle size within the sample, which is the result of aggregation and was only seen for intermediate UPy conjugations (2, 5, and 8). For UPy-RCPCh1-2, two distinct populations were observed at 20 °C (Figure 3B′). It is likely due to the presence of both grafted and unmodified RCPCh1, which results in different types of aggregation mechanisms and in the formation of larger and smaller particles. These results could be partially explained by the balance between intra- and intermolecular interactions and the polarity between the different UPy-RCPCh1 derivatives (Figure S16). By increasing the degree of functionalization of UPy groups, a decrease in polarity (Figure 1C) and a decrease of the ζ-potential of each UPy-RCPCh1 derivative was observed (Table 2). This is expected since polar and positively charged amine functional groups are replaced by hydrophobic UPy functionalities, which could influence the aggregation of UPy-RCPCh1 derivatives via their RCPCh1–RCPCh1 or UPy–UPy interactions. It is speculated that molecular packing and self-assembly via RCPCh1–RCPCh1 and/or UPy–UPy interactions are enhanced for the intermediate degree of functionalization (UPy-RCPCh1-2, 5, and 8) due to the presence of sufficient free amines that enhance solubility while allowing for intermolecular interactions with carboxyl groups between UPy-RCPCh1 molecules. Accordingly, higher degrees of functionalization (UPy-RCPCh1-12 and 16) resulted in a decrease in polarity and solubility, which results in more dense molecular packing.

It is speculated that for higher UPy conjugations, more stable intramolecular interactions are formed based on hydrogen bonding and hydrophobic interactions of the UPy moieties, which results in a decrease in electrostatic interactions of free amines and carboxyl groups between UPy-RCPCh1 molecules (Figure S16). This could explain the decrease in the formation of larger aggregates in dilute solutions (Figure 2E′,F′) of UPy-RCPCh1-12 and UPy-RCPCh1-16. Since the aggregation behavior of UPy-RCPCh1 derivatives is dictated via both RCPCh1–RCPCh1 and UPy–UPy interactions, it remains difficult to elucidate the true effect of different degrees of UPy functionalization. For this reason, the pH was first increased (pH > 12) to dissociate UPy–UPy interactions and consequently decreased to also
study the aggregation behavior at different temperatures. At elevated pH, RCPhC1 and UPy-RCPhC1 derivatives have a similar particle size; however, some subpopulations are observed for intermediate degrees of functionalization UPy-RCPhC1-2, -5, and -8 (Figure S15). Upon decreasing the temperature to 5 °C, no change in particle size is observed, which indicates a decrease in intermolecular interactions as a result of elevated pH. Upon lowering the pH, different particle size distribution and larger particle sizes were observed for RCPhC1 and all UPy-RCPhC1 derivatives. In addition, decreasing the temperature to 5 °C resulted in a wider distribution of the particle size (Figure S15). Moreover, for high degrees of UPy functionalizations, UPy-RCPhC1-12 and UPy-RCPhC1-16, larger aggregates formed after adjusting the pH. These results show the complexity of molecular aggregation as a result of changes in pH or temperature which is due to the presence of both RCPhC1–RCPhC1 and UPy–UPy interactions.

Hydrogel Formation and Mechanical Properties. The effect of increasing the degree of functionalization with UPy groups was also studied in concentrated solutions of UPy-RCPhC1 in PBS (100 mg/mL). The temperature was lowered to determine the liquid-gel cross-over of these solutions (Figure S17). In general, concentrated solutions of pristine RCPhC1 show a cross-over of the G' and G" and gel formation at ~10 °C. Interestingly, upon functionalization of RCPhC1 with on average 2 UPy groups and 5 UPy groups, an increase in cross-over temperature was observed, 14 and 16 °C, respectively (Figure S17). For UPy-RCPhC1-8, -12, and -16, a hydrogel was observed at all temperatures. These results clearly indicate the influence of UPy functionalities on the increase of cross-link formation and faster gelation.

Next, mechanical properties were measured using a nanoindenter, since loads (micronewton range) and scales (10–20 μm) are similar to what cells are able to sense. Interestingly, robust and stable hydrogels were formed through the modification of RCPhC1 with UPy functionalities (100 mg/mL, 20 °C) (Figure 4A). Hydrogels based on UPy-RCPhC1-5, 8, 12, and 16 remained intact at room temperature, however, an increase in opacity was observed as the degree of UPy functionalization increased (Figure 4A). This could be due to the presence of more UPy groups that cause the formation of larger aggregates within the hydrogel network, which resulted in decreased transparency. Interestingly, an increase in effective Young's modulus was observed upon increasing the degree of functionalization (Figure 4C). This is due to the presence of larger amounts of cross-links when more UPy groups are coupled to the backbone of RCPhC1. Indeed, the load–displacement curves recorded for each hydrogel show a local increase in the stiffness and maximum load (Figure 4B) and thereby confirm that the macroscopic properties we observe originate from our grafting strategy at the molecular scale. Here, it was chosen to study the response of mechanically sensitive cells, i.e., cardiac progenitor cells, on different UPy-RCPhC1 hydrogels as a culture platform. The stiffness of cardiac tissue typically ranges between 1–2 and 10–20 kPa from cardiac development up to mature cardiac tissue, respectively. Additionally, following a myocardial infarction, a fibrotic scar is formed that typically displays a higher order of magnitude mechanical stiffness with Young’s moduli of 35–70 kPa. It was shown that embryonic cardiomyocytes respond differently on substrates with these mechanical rigidities. For this reason, it is speculated that cardiac progenitor cells would also respond differently on “soft” UPy-RCPhC1-8 hydrogels (Eeff = 26 ± 19 kPa) compared to more rigid UPy-RCPhC1-12 and UPy-RCPhC1-16 hydrogels (Eeff = 68 ± 51 and 190 ± 118 kPa, respectively). Unfortunately, UPy-RCPhC1-2 and -5 did not form hydrogels at a concentration of 10 wt % at 37 °C.

CPC Behavior on UPy-RCPhC1 Hydrogels. Previous studies have determined the relationship between the mechanical stiffness of the environment and the biological response of cells. Moreover, in the field of cardiac development and regeneration, the effect of the substrate stiffness on the adhesion, proliferation, and differentiation of cardiac progenitor cells have been thoroughly studied previously. It was shown that matrix stiffness influences the genetic expression of cardiac progenitor cells (CPCs) via
their mechanotransduction pathways.\textsuperscript{36,54} Here, CPCs were cultured on top of UPy-RCPhC1-8, -12, and -16 hydrogels and both adhesion and proliferation were studied (Figure 5). The polymer concentration (10 wt %) and thereby the RGD concentration (each RCPhC1 molecule contains a fixed number of 12 RGD) were kept constant to study only the effect of substrate stiffness as a result of changing the UPy conjugation. Intriguingly, CPCs showed increased spreading and decreased clustering on stiffer UPy-RCPhC1-12 and -16 substrates compared to UPy-RCPhC1-8, which was coupled with an increase in stress fiber and zyxin formation after 1 day of culture (Figure 5A). Zyxin is a phosphoprotein which is located at the focal adhesions and thereby indicates a strong interaction between cells and their extracellular matrix.\textsuperscript{55} Increased amounts of zyxin spots on UPy-RCPhC1 hydrogels could be due to increased stiffness, whereas on softer UPy-RCPhC1-8 surfaces cells favor cell clustering and cell–cell interactions due to the low mechanical rigidity (Figure 5A). Next, CPC proliferation was studied on UPy-RCPhC1-18, -12, and -16 after 3 days of culture (Figure 5B,C). For UPy-RCPhC1-8, CPCs showed low proliferation and cell numbers (Figure 5B) and low expression of proliferation marker ki-67 was observed (Figure 5C). For stiffer substrates based on UPy-RCPhC1-12 and -16, an increase in cell number (Figure 5B) and ki-67 expression was observed (Figure 5C).

These results indicate that the proliferation of CPCs is inhibited on softer substrates. These findings are in agreement with previous research; however, different rigidities and cell types were studied.\textsuperscript{47,56} The mechanical stiffness of the substrate may also direct the cellular fate of CPCs or cardiac stem cells.\textsuperscript{36} Therefore, we further studied the effect of stiffness of the different substrates on the behavior of CPCs. To this end, the intracellular Yes-associated protein (YAP) was stained and imaged (Figures S18 and S20). YAP is a downstream effector protein involved in the Hippo pathway that is a key player in sensing substrate stiffness and consequently driving cellular fate.\textsuperscript{57–59} The mechanism is based on the translocation of YAP from the cytoplasm to the nucleus and results in downstream signaling related to proliferation and differentiation. Accordingly, immunofluorescence images show a minor increase of the YAP signal located at the nucleus compared to the cytoplasm on UPy-RCPhC1-18 hydrogels compared to UPy-RCPhC1-8 and UPy-RCPhC1-16 (Figure S18). However, due to the high degree of clustering, it remains inconclusive to relate the mechanical rigidity of the substrate with the YAP translocation, since it was shown that YAP mechanosensing is correlated to cell density regardless of the substrate rigidity.\textsuperscript{60} Taken together, different degrees of CPC adhesion, proliferation, and YAP location was observed on the different UPy-RCPhC1 hydrogel substrates, which is speculated to be due the difference in crosslinking density and consequently the mechanical rigidity (Figure S19).

**Figure 5.** Cell adhesion and proliferation on UPy-RCPhC1 hydrogels. (A) Immunofluorescence images showing CPC distribution, f-actin stress fiber formation, and zyxin protein localization on (top-to-bottom) UPy-RCPhC1-8, UPy-RCPhC1-12, and UPy-RCPhC1-16. For each hydrogel, the individual channels, merged and zoomed images are shown from left-to-right, the nuclei (blue), f-actin stress fibers (green), zyxin (red), merged image and zoomed images (the dotted area is seen on the merged image). Scale bar is equal to 75 and 25 μm. (B) Graph showing the cell numbers after 3 days of culture on UPy-RCPhC1-8, UPy-RCPhC1-12, and UPy-RCPhC1-16. The dotted line represents seeded cells on day 0. (C) Immunofluorescence images showing ki-67 protein expression located at the nuclei after 3 days of culture on UPy-RCPhC1-8, UPy-RCPhC1-12, and UPy-RCPhC1-16. Ki-67 is shown in white, f-actin in green, nuclei in blue, and the scale bar is equal to 100 or 75 μm.
CONCLUSIONS
In this work, collagen-mimicking peptides were successfully modified with different degrees of supramolecular UPy groups. It was shown that the assembly and folding of UPy-RCPC1 were dependent on the degree of functionalization and resulted in an increased stability of intermolecular interactions based on UPy–UPy interactions. Additionally, increased control over hydrogel formation and mechanical properties compared to pristine RCPhC1 hydrogels was achieved. The applicability of this hydrogel was shown by a clear difference in focal adhesion formation and proliferation of C2C12, as a consequence of changing the mechanical rigidity. This work illustrates the versatility of modifying biomaterials with supramolecular UPy-based crosslinks for tissue engineering applications. In future work, UPy-RCPhC1 derivatives can be used as pH-sensitive injectable hydrogels (Figure S22) and as cellular carriers that act as synthetic extracellular matrices that used to enhance cellular-based tissue regeneration therapies.

ASSOCIATED CONTENT
1 Supporting Information
The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.bioconjmat.9b00353.

1H NMR spectra, LC–MS spectra, mass spectra, schematic illustrations, cryo-TEM images, DLS, rheology, standard curves, and immunofluorescence images (PDF)

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