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Rheological Studies of Injectable Thermoresponsive Biodegradable Hydrogels and Porcine Ocular Tissues

Chandana Reddy Damera
University of Tennessee Health Science Center

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Research Advisor
Tao L. Lowe, Ph.D.

Committee
David Hamilton, D.V.M. George Huang, D.Sc. Bernd Meibohm, Ph.D.

ORCID
http://orcid.org/0000-0002-2768-1948

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Rheological Studies of Injectable Thermoresponsive Biodegradable Hydrogels and Porcine Ocular Tissues

A Thesis
Presented for
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Health Science Center

In Partial Fulfillment
Of the Requirements for the Degree
Master of Science
From The University of Tennessee

By
Sri Chandana Reddy Damera
May 2018
DEDICATION

Dedicated to my loving family.
ACKNOWLEDGEMENTS

First and foremost, I would like to express my sincere gratitude to my advisor, Dr. Tao Lowe for her continuous support and guidance throughout my graduate studies and related research. I would like to thank her for her patience, motivation, and immense knowledge. Her office was always open whenever I ran into a problem or had a question about my research or writing. Her writing and editing skills are superior to many and she was an irreplaceable mentor throughout the writing process. Her undeniable devotion to her students, research, and the future of science is inspiring and did not go unnoticed. Besides my advisor, I would like to thank Dr. George Huang for providing me with dental pulp stem cells and Dr. Rajashekar Gangaraju for providing me with adipose-derived stem cells for my project and for their scientific discussions.

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ABSTRACT

This research has evaluated the rheological properties of two types of materials which include (i) semi-synthetic polymer-based injectable hydrogels [P(NIPAAm-co-DEX-Lactate HEMA) and P(NIPAAm-co-HA-AEMA)], and (ii) biological tissues (porcine ocular tissues).

A series of thermoresponsive and biodegradable in situ hydrogel based on N-isopropylacrylamide (NIPAAm) monomer and hydrolytically degradable oligolactate Dextran-lactate-HEMA or Hyaluronic acid (HA)-AEMA macromer were investigated as encapsulation matrices (scaffolds) for DPSCs/ADSCs and assess their applicability in dental/retinal tissue engineering. The rheological properties of the hydrogels were strongly dependent on the reaction conditions and composition of the hydrogels. Both the hydrogels exhibited linear viscoelasticity, and the values of storage modulus (G’) at 1 Hz appeared in the range of 15-35 Pa for P(NIPAAm-co-DEX-Lactate HEMA) gels and 20-40 Pa for P(NIPAAm-co-HA-AEMA) gels at 15 mg·mL⁻¹ concentration. No significant differences were found for the hydrogels prepared with the HA or dextran at the same concentration and monomer to macromer ratio (19:1). The hydrogels were nontoxic to DPSCs and ADSCs at a concentration up to 15 mg·mL⁻¹, and in vitro survival studies using MTT and DAPI indicated that both DPSCs and ADSCs have favorably adhered and proliferated within the dextran and HA-based hydrogels. Survival of the DPSCs/ADSCs in the hydrogels, growth and morphology make these scaffolds a viable option for using as a biological scaffold for tissue engineering.

Since the functionality of the ocular tissues closely depends on their biomechanical properties, the rheological properties of the ocular tissues at different stages of development (preterm, neonatal and adult stage) is of therapeutic interest and has not been thoroughly investigated. This study has investigated the shear properties of porcine cornea, sclera and vitreous under strain and frequency sweep. All three tissue types at different stages have shown linear viscoelastic region under strain sweep. The G’ of cornea and vitreous weakly varied between neonatal and preterm, however, the adult cornea (~1.3-fold) and vitreous (~1.8-fold) showed significant higher moduli. On the other hand, the G’ of the sclera increased significantly with the developmental stage of the tissue, (preterm < neonatal (~1.9-fold) < adult (17.4-fold)). Thus, the rheological properties of the porcine ocular tissues varied with the developmental stage of the tissue.
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<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Full Form</th>
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<tbody>
<tr>
<td>ADSCs</td>
<td>Adipose-Derived Stem Cells</td>
</tr>
<tr>
<td>AEMA</td>
<td>Aminoethyl methacrylate</td>
</tr>
<tr>
<td>CaCl2</td>
<td>Calcium Chloride</td>
</tr>
<tr>
<td>CDI</td>
<td>N,N’-carbonyl diimidazole</td>
</tr>
<tr>
<td>DAPI</td>
<td>(4’,6-diamidino-2-phenylindole)</td>
</tr>
<tr>
<td>DMAP</td>
<td>4-(N,N-dimethylamino) pyridine</td>
</tr>
<tr>
<td>DMSO</td>
<td>Dimethyl Sulfoxide</td>
</tr>
<tr>
<td>DP</td>
<td>Degree of Polymerization</td>
</tr>
<tr>
<td>DPSCs</td>
<td>Dental Pulp Stem Cells</td>
</tr>
<tr>
<td>DS</td>
<td>Degree of Substitution</td>
</tr>
<tr>
<td>ECM</td>
<td>Extra Cellular Matrix</td>
</tr>
<tr>
<td>ESB</td>
<td>European Society of Biomaterials</td>
</tr>
<tr>
<td>G”</td>
<td>Loss modulus</td>
</tr>
<tr>
<td>G’</td>
<td>Storage Modulus</td>
</tr>
<tr>
<td>HA</td>
<td>Hyaluronic Acid</td>
</tr>
<tr>
<td>HA</td>
<td>Hydroxyapatite</td>
</tr>
<tr>
<td>HEMA</td>
<td>2-hydroxyethyl methacrylate</td>
</tr>
<tr>
<td>KCl</td>
<td>Potassium Chloride</td>
</tr>
<tr>
<td>MgCl2</td>
<td>Magnesium Chloride</td>
</tr>
<tr>
<td>MTT</td>
<td>3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide</td>
</tr>
<tr>
<td>NaCl</td>
<td>Sodium Chloride</td>
</tr>
<tr>
<td>NaH2PO4</td>
<td>Sodium dihydrogen phosphate</td>
</tr>
<tr>
<td>NaHCO3</td>
<td>Sodium Bicarbonate</td>
</tr>
<tr>
<td>PCL</td>
<td>Polycaprolactone</td>
</tr>
<tr>
<td>PCLLA</td>
<td>Poly(caprolactone-co-lactide)</td>
</tr>
<tr>
<td>PLA</td>
<td>Polylactide</td>
</tr>
<tr>
<td>PLG</td>
<td>Polyglycolide</td>
</tr>
<tr>
<td>PLGA</td>
<td>Poly lactide-co-glycolide</td>
</tr>
<tr>
<td>PNIPAAM</td>
<td>Poly-N-isopropylacrylamide</td>
</tr>
<tr>
<td>RGCs</td>
<td>Retinal Ganglion Cells</td>
</tr>
<tr>
<td>SnOct2</td>
<td>Stannous octoate</td>
</tr>
<tr>
<td>TCP</td>
<td>Tri-calcium Phosphate</td>
</tr>
<tr>
<td>TEMED</td>
<td>Tetramethylethlyenediamine</td>
</tr>
<tr>
<td>THF</td>
<td>Tetrahydrofuran</td>
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CHAPTER 1. INTRODUCTION

This research has evaluated the rheological properties of two types of materials which include (i) semi-synthetic polymer-based injectable hydrogels [P(NIPAAm-co-DEX-Lactate HEMA) and P(NIPAAm-co-HA-AEMA)], and (ii) biological tissues (porcine ocular tissues).

PNIPAAm-Based (Thermoresponsive) Injectable Hydrogels

Among the injectable hydrogels, in situ hydrogels formed by thermoresponsive phase transition have gained extensive attention because the gelation can be triggered as the temperature rises above the lower critical solution temperature (LCST) of the polymer. LCST can be designed below body temperature so that the sol-gel phase transition of injectable systems get triggered at the physiological condition (37°C). PNIPAAm is a typical thermosensitive polymer used in the synthesis of various hydrogels. Below LCST, PNIPAAm is hydrophilic and remains in a soluble state in aqueous solution, but with an increase in the temperature above LCST it undergoes a phase transition and forms a hydrogel. Several studies have shown that the phase behavior and mechanical properties of PNIPAAm hydrogels can be modified by the addition of hydrophobic or hydrophilic monomers to impart desired properties into the material. Various composite hydrogel systems were reported in the literature by modifying the PNIPAAm with poly(ethylene glycol), chitosan, hyaluronic acid, chondroitin sulfate or other polymers to alter its physicochemical/biological/mechanical properties such as to preserve the viability and phenotypic morphology of the cells, improve proliferation, differentiation and extracellular matrix secretion of the cells entrapped within the hydrogel, provide mechanical properties for providing ambient environment for supporting the cell growth.

Although promising, the thermoresponsive NIPAAm homopolymer are not biodegradable. It is more advantageous to introduce degradable property into the PNIPAAm-based scaffolds as degradation of the hydrogel matrix not only circumvents removal of the empty device but also can be used to modulate space in response to the tissue regeneration and nutrient diffusion. Also, when the polymer slowly degrades, the cells will be able to produce their own ECM and generate successful cell-polymer graft. Therefore, in this study, we designed and synthesized injectable thermoresponsive and biodegradable hydrogels by copolymerizing NIPAAm monomer with dextran-lactate-2-hydroxyethyl methacrylate (DEX-Lactate-HEMA) or HA-aminoethylmethacrylate (HA-AEMA) macromer using the synthetic technique. Because of the existence of a hydrolyzable/enzymatically degradable DEX-Lactate/HA in the prepolymer mixture, biodegradable crosslinks can be formed within the PNIPAAm hydrogel systems, resulting in biodegradable thermoresponsive systems. Dextran and HA used in the study for modifying PNIPAAm are naturally occurring biodegradable polymers. Dextran is a natural polysaccharide that is enzymatically degradable by dextranases [12]. Dextran is commercially available in wide range of MWs with relatively narrow MW distribution. It
has been successfully used for various tissue engineering applications because of its excellent biocompatibility, biodegradability, hydrophilicity, stability under slightly acidic and basic conditions, and surface hydroxyl functionality for conjugation [13-16]. Several reports have been published on altering the dextran properties to make various dextran graft polymers via various chemical modifications for various applications [12]. HA is one of the major glycosaminoglycan polysaccharides in the ECM of the newly developed dentin, pulp tissues, and vitreous humor, and is enzymatically degradable by hyaluronidase. The biocompatible structure, biodegradation rate of HA and the flexibility to modify structurally and chemically has suggested its suitability as a scaffold for DPSCs/ADSCs in tissue engineering applications [5, 17-20].

**Hypothesis and Specific Aims**

The long-term goal of the project is to develop biomaterials for dental and retinal tissue engineering. The short-term goal of this project was to design and develop injectable thermoresponsive and biodegradable hydrogels P(NIPAAm-co-DEX-Lactate-HEMA) and P(NIPAAm-co-Hyaluronic acid AEMA) as scaffolds for encapsulating DPSCs/ADSCs for dental and retinal tissue engineering applications. Also, to investigate whether the scaffold material properties can be controlled by polymer modification and whether cell viability and growth could be improved. The hypothesis of this research was that the injectable biodegradable P(NIPAAm-co-DEX-Lactate-HEMA) and P(NIPAAm-co-HA-AEMA) hydrogels were nontoxic to DPSCs and ADSCs cells and promote their growth inside the matrices.

**Specific Aim 1**

To synthesize and characterize P(NIPAAm-co-DEX-Lactate-HEMA) and P(NIPAAm-co-HA-AEMA) hydrogels by UV polymerization and FTIR. A series of biodegradable P(NIPAAm-co-DEX-Lactate-HEMA) and p(NIPAAm-co-HA-AEMA) hydrogels were synthesized with varying monomer/macromer ratios, initiator concentration, UV intensity, and reaction times. The hydrogels were characterized for sol-gel transition and rheological properties.

**Specific Aim 2**

To evaluate in vitro cytotoxicity and encapsulation capabilities of P(NIPAAm-co-DEX-Lactate-HEMA) and P(NIPAAm-co-HA-AEMA) hydrogels. In vitro cytotoxicity of hydrogels to DPSCs and ADSCs was tested at various concentrations of the hydrogel in cell growth medium by MTT assay. The growth of DPSCs/ADSCs inside the hydrogel matrix was examined using MTT and DAPI assays.
Rheological Studies of Porcine Ocular Tissues

The proper vision of the eye strongly depends on its ability to maintain optical clarity and structural integrity under routine variations in pressure, changes in humidity and temperature, constant muscular stress/strain and rapid sudden movements. Thus the proper functioning of the eye is dependent on its physical and mechanical properties, and altered chemical and mechanical states of ocular tissues could lead to abnormalities and may result in complete loss of the functionality. Various age-related changes in composition and structure of the eye tissues affect the biomechanical properties and performance of the eye tissues and could contribute to the predisposition of various eye abnormalities [21-26]. For example, aging results in stiffening of the cornea tissue which could be because of the additional age-related non-enzymatic cross-linking affecting the stromal collagen fibrils [24, 25]. It is reported in the literature that age-related variations affect the biomechanical characteristics of the sclera [21], and aging results in the microstructural changes in the sclera and contribute to the predisposition to primary open-angle glaucoma [22]. With age, collagen networks in the vitreous begin to break down and pockets of liquid form in the eye, the process of liquefaction in which the liquid volume increases and gel volume decreases [23, 26]. These age-related changes could lead to structural and mechanical differences between vitreous at various stages of development, and in about 25-30% of the population, these age-related degenerative changes eventually result in posterior vitreous detachment [27]. Given the importance of the viscoelastic properties of the ocular tissues, to its function and to pathology, detailed rheological characterization of these tissues is crucial to appreciate the normal functions and quantitative understanding of disease pathophysiology of these tissues at various stages of development. The objective of this project was to characterize the rheological properties of the ocular tissues (cornea, sclera and vitreous) at different stages (preterm, neonatal and adult) of development. Though several measurements could be measured in vivo, such as topography, thickness distribution, but the attempts to investigate the biomechanical properties of the tissues still depend on ex-vivo experiments involving human or other animal tissues [28]. Because of the difficulties in obtaining human donor tissues in sufficient numbers for the research and experimental needs, it had been necessary to use animal ocular tissues as approximate models for human ocular tissues for performing various evaluations including the mechanical property characterization studies. Several investigations used porcine tissues for ocular related experiments [28-30], although other animal’s rabbit and bovine tissues a common practice. Therefore, the current rheological characterization also contributes to the literature for identifying and assessing the important similarities and significant differences between the rheological behaviors of porcine ocular tissues and human and other animal tissues and evaluating the suitability of the porcine eye tissues as a model of the human tissues.

Hypothesis and Specific Aims

The hypothesis of this project was that the storage and loss modulus of the porcine ocular tissues differ at different stages of development (preterm, neonatal and adult). There were two specific aims:
Specific Aim 1

To conduct the shear-sweep experiments and identify the linear viscoelastic region of the porcine ocular tissues (cornea, sclera, vitreous) at different stages of development (preterm, neonatal, adult).

Specific Aim 2

To conduct the frequency sweep experiment and analyze the gel-like behaviors of the porcine ocular tissues (cornea, sclera, vitreous) at different stages of development (preterm, neonatal, adult).
CHAPTER 2. SYNTHEThESIS AND CHARACTERIZATION OF INJECTABLE THERMORESPONSIVE AND BIODEGRADABLE HYDROGELS FOR STEM CELL-BASED TISSUE ENGINEERING

Introduction

Stem-Cell Based Therapy

Organ failure disorders and several of neurodegenerative diseases characterized by death of cells are leading to the irreversible loss of tissues and functions. For many of the neurodegenerative diseases, particularly in those conditions where an extensive cell loss was occurred prior to molecular level diagnosis of their gene defects, just the drug and/or gene therapy may not be enough to resolve the issues [31]. In such instances, the strategies were focused on cellular or tissue replacement, and based on recent studies, it was found that these strategies were beneficial [31-33]. Cell-based therapy involves the use of exogenous cells obtained from either autologous or allogenic source [34]. In such cases, use of stem/progenitor cells presents major advantages over differentiated cells as they can proliferate to the required amounts before or after transplantation [34].

Even though the stem-cell based therapies are promising, still they need an effective transplantation system. Current direct delivery methods could result in massive cell loss and limited cellular integration following transplantation [31]. For example, studies indicated that after bolus injection of photoreceptor cells, less than 0.01% of transplanted cells survive and even fewer actually integrate within the host retina [35, 36]. Therefore, there is a need for efficient delivery methods that could minimize the cell loss and assist in the growth and cell integration into the host system, and tissue engineering scaffolds are the alternative option that has shown to help in addressing the above-mentioned issues [37-39].

Tissue Engineering

Since its first use as a term in the mid-1980s, tissue engineering has moved from ‘science-fiction’ concept to ‘science fact’ and emerged as a significant multidisciplinary field in the biotechnological area, with the research-oriented acceptance of its potential to regulatory approvals and market availability for the treatment [40]. Especially, with a more significant advancement of biomaterials, tissue engineering is emerging as a great potential alternative or complementary solution for restoring, maintaining or improving tissue or organ functions [41]. A general tissue engineering paradigm involves the use of a highly porous artificial extracellular matrix or ‘scaffold’ as a temporary support matrix for cell growth, differentiation, and matrix production, and the resulting tissue engineering construct is then grafted back in vivo into the subject. These constructs are either fully functional or they grow into the required functionality [41, 42]. Thus, tissue
engineering has the promising potential to revolutionize methods in medical treatment and can offer a long-term solution for many age-related conditions.

In simple terms, scaffolds are porous artificial materials functioning as an extracellular matrix that accommodates and guides the growth and proliferation of cells in the 3D environment [39, 42]. The native extracellular matrix (ECM) not only provides physical support or appropriate architecture to the cells but also functions as a substrate for cell adhesion, migration and controls cellular proliferation and supplies with various growth factors and nutrients. In order to guide the formation of tissues with a required structural and functional characteristics, it is essential to provide a dynamic in vitro microenvironment with a proper quantity of physical strength such as shear and mechanical forces that promote their development [43]. Therefore, the key objective of a scaffold is to closely mimic the native ECM environment in terms of physical structure, chemical composition, and biological functions and play a similar role for promoting tissue regeneration in vitro as native ECM does in vivo [41]. Overall, the desirable physical characteristics of biomaterial scaffold for tissue engineering applications include nontoxic/non-inflammatory, hydrophilicity, high porosity with uniformly distributed interconnected pores throughout the matrix, large surface area, large pore size, and the ability to degrade in response to tissue repair and growth progresses, ideal shear and mechanical support as native environment [44-46].

**Dental and Retinal Tissue Engineering**

Data on tooth loss showed that at least 20 million people in the US have missed all of their teeth, and more than 100 million have lost 11–15 teeth [47]. Dental tissue engineering, to regenerate the dental pulp or dentin tissues in the pulp space of teeth, has grown tremendously as a new option in the area of endodontics for preserving and restoring the teeth [34, 48]. Although dentine can regenerate, the pulp has a limited capability for regeneration as it is encased in dentine and has limited apical blood supply [49]. Dental pulp stem cells (DPSCs) are an accessible multipotent stem cell source with therapeutic applicability in repair and regeneration of damaged tissues [50]. Dr. Huang’s group has demonstrated that DPSCs are a promising source for dental tissue regeneration (dentin-pulp like tissues, bioroot) [51]. Also because of their angiogenic and angioinductive property, they are a good source for non-dental (bonelike tissues, neurogenic, angiogenic) tissue engineering applications as well [34]. Studies conducted on dental pulp tissue regeneration, utilizing the DPSCs seeded PLGA-based synthetic preformed scaffolds, have shown the formation of pulp-like tissue in both in vitro and in vivo models. However, due to the smaller size and encasement of the dental pulp inside the root canal, utilizing an injectable scaffold that can be administered into the enclosed area could complement the efficiency of the DPSCs-based therapy for engineering the dentine-pulp complex by minimizing the cell loss and assisting in the growth and cell integration into the host system upon administration.

Retinal degenerations are a group of conditions that can lead to permanent visual loss and affect millions of people worldwide [52]. Clinically, there are no neuroprotective
nor axogenic therapies are available that can restore lost visual functionality in retinal degenerative diseases [53]. Current therapeutic strategies such as anti-angiogenic drugs, anti-inflammatory, and gene therapy can delay the disease progression but not effective in permanent cure. Several studies are in progress aiming to regenerate the diseased retina by stem-cell transplantation for the replacement of lost retinal ganglion cells (RGCs) and photoreceptors. Currently, there are many ongoing clinical trials which aim to test the safety and efficacy of stem cell transplantation in the eye [52]. Several reports in the literature have suggested that the retinal tissue can be replaced and some degree of functional recovery can be regained following the delivery of stem-cells [54]. Adipose-derived stem cells (ADSCs) are progenitor cells that can easily be isolated from adipose tissue [55]. They have the capability of self-renewal and multilineage differentiation and can be transplanted to either autologous or allogeneic body safely with less implant migration and foreign body reaction [20]. The ADSCs have functional and phenotypic overlap and plays a key role in providing vascular protection [20, 56, 57]. ADSCs are also known to produce angiogenic and antiapoptotic factors [55]. Recent findings from Dr. Gangaraju’s group on the regenerative therapeutic potential of ADSCs and showed that a single direct intravitreal injection of ADSCs significantly limited the development of diabetic ocular complications [55]. Intravitreal injection of ADSCs has shown to stabilize the retinal microvasculature, thereby enabling repair and regeneration of damaged capillaries in vivo [58]. The results suggested that ADSCs have great potential to be developed into a cell-based therapy to address both retinal capillary and neurodegeneration [55]. Earlier retinal transplantation studies involved direct injection of the cells into the vitreous or subretinal space with the assumption that the injected cells would migrate to the degenerated site of the retina, integrate within the host retina and restore vision functionality [52]. Despite promising outcomes of cell-based therapies to regenerate the degenerating retina, they also demonstrated that the direct bolus injection of stem cells in vivo could result in disorganized and poorly localized tissue grafts [59]. The resulted outcomes of low rates of cell survival due to poor donor cell integration and injection reflux are primarily related to the transplantation procedure [59]. Use of injectable hydrogel scaffolds for seeding and delivering the stems cells into the retina could be an attractive alternative to direct injection as they can potentially limit donor cell death due to injection reflux and help in improving the integration of the donor cells into host system [60].

Therefore, new biomaterials, injectable thermoresponsive and biodegradable hydrogel scaffolds, have been evaluated in this project for seeding DPSCs/ADSCs for dental/retinal tissue engineering applications.

Injectable Hydrogel Biomaterials

Biomaterials have been proved to significantly affect the proliferation and differentiation potential of stem cells by controlling physicochemical properties (e.g., surface charge, source, mechanical properties and microstructural patterns) [61]. In the recent years, significant efforts have been made to develop several biocompatible scaffolds for tissue engineering based on synthetic and natural polymers and studied to
improve the cell attachment, viability/proliferation, differentiation and growth of the DPSCs/ADSCs [44, 46, 62-65]. The synthetic polymers (polyglycolide (PLG) [66, 67], polylactide (PLA)[68, 69], polylactide-co-glycolide (PLGA) [70-72], polycaprolactone (PCL) [73], PNIPAAm-based possess much flexibility in terms of tailoring the architecture (scaffold shape, porosity, mechanical properties), controlling the degradation rate and have shown much success [74-76]. But the limitations include the risk of rejection rate due to cell and tissue necrosis due to degradation products of polymers [44, 77]. Unlike synthetic polymer-based scaffolds, natural polymers (polysaccharides (e.g., chitin [78], chitosan [79-81], hyaluronic acid (HA) [17-19], alginate [82-85]) proteins (collagen [17, 86, 87], gelatin [84], fibrin [76]) are bioactive and promote cell growth and cell adhesion. Further, they are biodegradable thus can allow the host cells to grow and produce their own ECM and replace the degraded scaffold over time. Thus advantages of these single type of polymers have motivated the researchers to investigate combinatorial systems/composite scaffolds comprising multiple types of biomaterials [44]. Such examples include combining synthetic polymers with natural polymers [88, 89], for enhancing their biological capacity. These combinational systems can have controlled degradation, have a wide range of mechanical properties that enable them to be used in various tissue engineering applications. Since human tissue ranges in terms of mechanical properties, control of mechanical properties of the materials using a composite scaffold of synthetic and natural polymers will be greatly advantageous.

Among various biomaterials, hydrogels represent an important class of scaffolds that have been explored for tissue engineering applications. These are 3-dimensional-crosslinked hydrophilic polymeric networks that absorb and retain large amounts of water or biological fluids [90]. They possess several unique properties such as 3-dimensional network, tissue compatibility and similarity to native ECM, support cell proliferation and migration, controlled release of growth factors and nutrient diffusion [91]. They have gained significant attention for tissue engineering applications because of their attractive physicochemical properties such as hydrophilicity, water imbibing nature, softness, nutrient diffusion, and synthetic flexibility [91-93]. However, the traditional pre-formed hydrogel scaffold needs invasive procedures for administration and may have improper adaptation to the defect site which could lead to scaffold failure. Therefore, injectable biodegradable hydrogel biomaterials, which are liquids before injection and transform to a gel at the site of injection in response to external stimuli, have been extensively investigated as cell scaffolds owing to their advantages such as ease of administration with minimal invasiveness, less discomfort and less scarring, complete filling of the defect area with better adaptation and good permeability [94]. In addition, cells can be easily incorporated in the solution by mixing prior to injection under appropriate conditions, and after injection, they can take the shape of the cavity and fill irregular defects [92, 93]. Eliminating highly invasive procedures could lower the healthcare costs and improve the recovery time for the patients.

Various physical and chemical mechanisms can be used to trigger this phase transition [94, 95]. Various types of injectable hydrogels have been investigated for the regeneration of i) cartilage tissues (using chondrocytes in: collagen [96], chitosan-hyaluronic acid (HA) [97], methacrylated glycol chitosan-HA [98], gelatin–hydroxy
phenylpropionic acid [99], HA-PEG [100], PEG-based [101], dextran-tyramine [13], HA-g-dextran-tyramine [15], ii) intervertebral discs (using chondrocytes in: amidic alginate [102]; using mesenchymal stem cells in: HA–PNIPAAm [103], PHEMA-co-aminopropyl methacrylamide (APMA)-graft-poly(amide amine) [104], HA-PEG [105]), iii) bone (using calvarial osteoblasts in: alginate hydrogels along with poly(aldehyde glucoronate) [106], RGD peptide modified alginate [107]; using human mesenchymal stem cells in: PCLA–PEG–PCLA [108], acrylated HA-PEG [109]; using osteoblasts in: RGD modified PEG hydrogel [110]; using marrow mesenchymal stem cells in: chitosan [111], chitin–PCL–nanoHAp [111], chitosan–alginate [112]); iv) adipose tissue (using adipose-derived stem cells in: PNIPAAM-HA [5], modified PEG [5], alginate-O-carboxymethyl chitosan–fibrin [113]; using preadipocytes in: alginate hydrogels linked with RGD peptide [114, 115]), v) muscle tissues (using skeletal myoblasts in: fibrin [116, 117], poly(d-valerolactone) (PVL)–PEG–PVL [118]; using cardiosphere derived cells in combination polymers PNIPAAm-PCL-HEMA-DBLA, Dex-PCL-HEMA/PNIPAAm [119, 120], using satellite cells in photo-crosslinkable HA [121], chitosan/beta-glycerolphosphate/collagen [122]; using mesoangioblasts in: PEG-fibrinogen [123]; using ischaemic progenitor cells in: collagen, alginate, HA, chitosan [124, 125]); v) dental (using DPSCs in Puramatrix [126]).

### PNIPAAm-Based (Thermoresponsive) Injectable Hydrogels

Among the injectable hydrogels, in situ hydrogels formed by thermoresponsive phase transition have gained extensive attention because the gelation can be triggered as the temperature rises above the lower critical solution temperature (LCST) of the polymer [1]. LCST can be designed below body temperature so that the sol-gel phase transition of injectable systems get triggered at the physiological condition (37°C) [2]. PNIPAAm is a typical thermosensitive polymer used in the synthesis of various hydrogels [3]. Below LCST, PNIPAAm is hydrophilic and remains in a soluble state in aqueous solution, but with an increase in the temperature above LCST it undergoes a phase transition and forms a hydrogel [4]. Several studies have shown that the phase behavior and mechanical properties of PNIPAAm hydrogels can be modified by the addition of hydrophobic or hydrophilic monomers to impart desired properties into the material [2, 5, 6]. Various composite hydrogel systems were reported in the literature by modifying the PNIPAAm with poly(ethylene glycol), chitosan, hyaluronic acid, chondroitin sulfate or other polymers to alter its physicochemical/ biological/mechanical properties such as to preserve the viability and phenotypic morphology of the cells, improve proliferation, differentiation and extracellular matrix secretion of the cells entrapped within the hydrogel, provide mechanical properties for providing ambient environment for supporting the cell growth [5-11].

Although promising, the thermoresponsive NIPAAm homopolymer are not biodegradable. It is more advantageous to introduce degradable property into the PNIPAAm-based scaffolds as degradation of the hydrogel matrix not only circumvents removal of the empty device but also can be used to modulate space in response to the tissue regeneration and nutrient diffusion. Also, when the polymer slowly degrades, the
cells will be able to produce their own ECM and generate successful cell-polymer graft. Therefore, in this study, we designed and synthesized injectable thermo-responsive and biodegradable hydrogels by copolymerizing NIPAAm monomer with dextran-lactate-2-hydroxyethyl methacrylate (DEX-Lactate-HEMA) or HA-aminoethylmethacrylate (HA-AEMA) macromer using the synthetic technique. Because of the existence of a hydrolyzable/enzymatically degradable DEX-Lactate/HA in the prepolymer mixture, biodegradable crosslinks can be formed within the NIPAAm hydrogel systems, resulting in biodegradable thermo-responsive systems. Dextran and HA used in the study for modifying PNIPAAm are naturally occurring biodegradable polymers. Dextran is a natural polysaccharide that is enzymatically degradable by dextranases [12]. Dextran is commercially available in wide range of MWs with relatively narrow MW distribution. It has been successfully used for various tissue engineering applications because of its excellent biocompatibility, biodegradability, hydrophilicity, stability under slightly acidic and basic conditions, and surface hydroxyl functionality for conjugation [13-16]. Several reports have been published on altering the dextran properties to make various dextran graft polymers via various chemical modifications for various applications [12]. HA is one of the major glycosaminoglycan polysaccharides in the ECM of the newly developed dentin, pulp tissues, and vitreous humor, and is enzymatically degradable by hyaluronidase. The biocompatible structure, biodegradation rate of HA and the flexibility to modify structurally and chemically has suggested its suitability as a scaffold for DPSCs/ADSCs in tissue engineering applications [5, 17-20].

**Hypothesis and Specific Aims**

The long-term goal of the project is to develop biomaterials for dental and retinal tissue engineering. The short-term goal of this project was to design and develop injectable thermo-responsive and biodegradable hydrogels P(NIPAAm-co-DEX-Lactate-HEMA) and P(NIPAAm-co-Hyaluronic acid AEMA) as scaffolds for encapsulating DPSCs/ADSCs for dental and retinal tissue engineering applications. Also, to investigate whether the scaffold material properties can be controlled by polymer modification and whether cell viability and growth could be improved. The hypothesis of this research was that the injectable biodegradable P(NIPAAm-co-DEX-Lactate-HEMA) and P(NIPAAm-co-HA-AEMA) hydrogels were nontoxic to DPSCs and ADSCs cells and promote their growth inside the matrices.

**Specific Aim 1.** To synthesize and characterize P(NIPAAm-co-DEX-Lactate-HEMA) and P(NIPAAm-co-HA-AEMA) hydrogels by UV polymerization and FTIR. A series of biodegradable P(NIPAAm-co-DEX-Lactate-HEMA) and p(NIPAAm-co-HA-AEMA) hydrogels were synthesized with varying monomer/macromer ratios, initiator concentration, UV intensity, and reaction times. The hydrogels were characterized for sol-gel transition and rheological properties.

**Specific Aim 2.** To evaluate in vitro cytotoxicity and encapsulation capabilities of P(NIPAAm-co-DEX-Lactate-HEMA) and P(NIPAAm-co-HA-AEMA) hydrogels. In vitro cytotoxicity of hydrogels to DPSCs and ADSCs was tested at various concentrations
of the hydrogel in cell growth medium by MTT assay. The growth of DPSCs/ADSCs inside the hydrogel matrix was examined using MTT and DAPI assays.

Materials and Methods

Chemicals and Reagents

Dextran (MW = 15,000 g·mol⁻¹) was purchased from Polysciences, Inc., Warrington, PA. Hyaluronic Acid (HA) was provided by Bloomage Freda Biopharm Co., Ltd., Jinan, China. The following materials were obtained from Sigma–Aldrich, Inc (St. Louis, MO): N-isopropylacrylamide (NIPAAm), 2-hydroxyethyl methacrylate (HEMA), 4-(N,N-dimethylamino) pyridine (DMAP), N,N'-carbonyl diimidazole (CDI), L-lactide, stannous octoate (SnOct₂), tetrahydrofuran (THF), irgacure 2959, dimethyl sulfoxide (DMSO), 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT), 4',6-diamidino-2-phenylindole (DAPI). Dialysis tubing of 50,000 molecular weight cut off (MWCO) was purchased from Spectrum Laboratories. DPSCs were obtained from Dr. Huang’s Lab and ADSCs were obtained from Dr. Gangaraju Rajashekar’s Lab. All the chemicals were used as received. Deionized distilled water was used in all the experiments.

Synthesis of DEX-Lactate-HEMA and HA-AEMA Macromers

DEX-Lactate-HEMA macromer with a theoretical degree of polymerization 4 (DP) and degree of substitution 10% (DS) was synthesized according to the previously published procedure [4]. Briefly, at first, L-lactide (4.32 g, 0.02 mol) and HEMA (1.8 g, 0.01 mol) were reacted in the presence of nitrogen and catalyst SnOct₂ (0.05 g, 0.00013 mol) at 110 °C for 1 h. HEMA-lactate product was collected by dissolving the cooled reaction mixture in THF, precipitating in ice-cold water, dissolving in ethyl acetate, drying over MgSO₄, and concentrating under reduced pressure. The yield of the above reaction was 35%. Next, CDI (2.59 g, 0.015 mol) was dissolved in THF (100 mL) in a nitrogen atmosphere and reacted with the above HEMA-lactate (4.454 g, 0.01 mol) at room temperature for 16 h. The resulting HEMA-lactate-imidazolyl carbamate (HEMA-lactateCI) was obtained by solvent evaporation under reduced pressure, and the reaction yield was 95%. Finally, dextran (4 g, 0.024 mol of glucopyranose residues) was dissolved in DMSO (90 mL) in a nitrogen atmosphere. After the dissolution of DMAP (2.0 g), HEMA- lactate-Cl (1.62 g, 0.002 mol) was added, and the mixture was stirred at room temperature for 4 days. The DEX-Lactate-HEMA product was obtained by precipitating the reaction mixture in a large excess volume of cold dry 2-propanol, washed several times with 2-propanol, and dried in a vacuum for at least 24 h. The yield of the final reaction was 85%.

HA-AEMA macromer of molecular weight 18k and 25% AEMA substitution used in thesis work was synthesized by previous lab member Mr. K.B. Compton and
reported in his thesis [127].

**Synthesis of P(NIPAAm-co-DEX-Lactate-HEMA) and P(NIPAAm-co-HA-AEMA) Hydrogels**

P(NIPAAm-co-DEX-Lactate-HEMA or DEX-based gels) and P(NIPAAm-co-HA-AEMA or HA-based gels) hydrogels were synthesized by photopolymerization of NIPAAm monomer and hydrolytically degradable oligolactate 2-hydroxyethyl methacrylate (DEX-Lactate-HEMA) macromer or HA-AEMA macromer. The variables that were studied and optimized during the synthesis of hydrogels include: NIPAAm to macromer ratio, the concentration of the pre-polymer mixture, initiator concentration, UV intensity, reaction time, distance of the tip of the probe from the surface of the liquid, and dialysis time. Different weight ratios of NIPAAm:DEX-Lactate-HEMA or NIPAAm:HA-AEMA (19:1~16:4, w:w) were used for the synthesis. Irgacure 2959, a biocompatible photoinitiator was used in different amounts relative to prepolymer mixture weight, 0.0125, 0.025, 0.05 w% for the initiation of the crosslinking. Different UV intensities (100, 250, 375, 500, 750, 1000 mW·cm\(^{-2}\)), and different reaction time (3, 10 min) are used for synthesizing the hydrogels. DEX-Lactate-HEMA (or) HA-AEMA macromers were dissolved in 25% ethanol. The concentration of the pre-polymer mixture used was 0.5, 1, 2 mg·mL\(^{-1}\). To the dissolved macromer solution, NIPAAm was added and vortexed to get a uniform solution. Initiator in different amounts (0.05, 0.025, 0.0125 w%) were added to the above solution. 50 µL of the pre-polymer mixture was placed into a 96 well plate and exposed to UV at different intensities of 100-750 mW·cm\(^{-2}\) for 3 (or) 10 min. The distance of the tip of the probe from the surface of a mold used was 0.5 or 0 cm. The obtained hydrogels were dialyzed against DI water for 4-6 h with changing the water every half an hour in a 2 L beaker. After dialysis, the contents were collected and freeze-dried to get the dried polymer, and characterized by FTIR. The yield of the DEX-based gels was 52, 75, 91 % with 0.0125, 0.025, 0.05 wt% initiator respectively and the yield of HA-based gels was 55, 79, 94 % with 0.0125, 0.025, 0.05 wt% initiator, respectively.

**Rheological Properties of the Hydrogels**

The rheological properties of hydrogels were tested using an AR-G2 Rheometer (TA Instruments, New Castle, Delaware) with 8mm parallel plate geometry. Specifically, the hydrogels were subjected from 0.01 to 100 percent strain, while maintaining a normal force to 0.01 - 0.03N, the temperature at 37°C, and a constant 1 Hz frequency of oscillation to determine linear viscoelastic region. 50 µL of hydrogel was used in each run and the gap of 500 µm was maintained between the top plate and bottom Peltier plate. The software for the rheometer was set up to take 15 sample points per decade of strain, for the strain sweep and 10 sample points per decade of frequency in a frequency sweep, and a resulting graph of the storage modulus over the range of percent strain or frequency was created at the termination of the procedure. Strain sweep was conducted to determine the linear viscoelastic region of the gels where G’ and G” are independent of % strain. For
the temperature ramp, $G'$ and $G''$ were measured with an increase in temperature from 15 to 45 °C at the rate of 1 °C/min.

**Sol-Gel Transition**

The dried DEX-Lactate-HEMA was dissolved at concentration 10 or 20 mg/mL in the dental pulp stem cell (DPSC) medium and transferred to Eppendorf tubes. Then the tubes were heated to 37°C from room temperature to test the gelation.

**Cell Culture**

The DPSCs were cultured in DPSCs medium containing a-MEM 500 mL, FBS 55 mL (final content: 10%-15%), 2mM L-glutamine 5.5 mL (final: 200mM), 1% antibiotics and 100 µg·mL-1 b-FGF 20 µL (final: 4 ng·mL-1) at 37°C in a humidified incubator containing 5% CO2. The ADSCs were cultured in DMEM/F-12 medium containing 10% FBS (Fetal Bovine Serum) and 1% antibiotics.

**In Vitro Cytotoxicity of Hydrogels to Stem Cells**

The cytotoxicity of P(NIPAAm-co-DEX-Lactate-HEMA) and P(NIPAAm-co-HA-AEMA) against DPSCs were performed by MTT assay. The cells were seeded in 96-well plate at a density of 7500 cells/well. After the incubation in 150 µL of DPSCs medium for 48 h, with replacing 100 µL of medium after 24 h, the medium was replaced with an assigned amount of p(NIPAAm-co-DEX-Lactate-HEMA) dissolved in DPSCs medium, respectively, and the cells were incubated at 37 °C and 5% CO2 for another 24 h. Hydrogel solution was added to each of the test wells at different concentrations (0.5, 1, 10 mg·mL⁻¹). After 24 h of incubation, 20 µL of MTT solution (5 mg·mL⁻¹ in PBS) was added to each well and the cells were further incubated for 4 h. A solubilizing solution, dimethyl sulfoxide (DMSO) is added to dissolve the insoluble purple formazan product into a colored solution and shaken for 10-15 minutes. The absorbance of this colored solution was quantified by measuring at a wavelength of 570 nm using a microplate reader (BioTek Instruments, Inc., Winooski, VT, USA). The relative percent cell viability was plotted.

**DPSCs (or) ADSCs Cell Survival Study by MTT and DAPI**

The materials used for this study was synthesized from macromer DEX-Lactate-HEMA (or) HA-AEMA (18k-25), NIPAAm:macromer was used in the ratio of 19:1 (w:w). Irgacure 2959 was used as initiator at the concentration of 0.05 w% of total monomer and macromer. No gap was maintained between the tip of UV probe and mold. UV intensity at 750 mW·cm⁻² for 10 min was used. The product obtained was then dialyzed against 4L of distilled water for 6h by changing the water every half an hour and
freeze-dried. DPSCs (or) ADSCs were grown to confluence and encapsulated in hydrogels with cell seeding density of $5 \times 10^4$ cells·mL$^{-1}$. Incubated at 37 °C, 5% CO$_2$ for 24 h. Concentrations of the hydrogels used were 10, 15 and 20 mg·mL$^{-1}$. After 24 h the hydrogels with the cells were centrifuged and the hydrogels were replaced with the medium. Then the medium with cells is seeded in a 96 well plate (200 µL/well) and 24 well plate (250 µL/well) for MTT and DAPI assays respectively and incubated at 37 °C, 5% CO$_2$ for 24 h. Then the following procedure was followed for the cell survival study by MTT. 26 µL of 5 mg·mL$^{-1}$ MTT was added to each well and incubated at 37 °C, 5% CO$_2$ for 4 h. Then 200µL of DMSO was added to each well, and shaken for 15 minutes. The absorbance was measured at 570 nm. For the cell survival study by DAPI assay, the following procedure was followed. After incubation for 24 h at 37 °C, 5% CO$_2$, the cells were washed with PBS 2-3 times and fixed by using 4% paraformaldehyde for 15 min. Then the cells were washed with PBS for 2-3 times and incubated with DAPI at concentration 1µg·mL$^{-1}$ for 15 - 20 min. After incubation, the cells were again washed with PBS for 2-3 times and then the images were taken at 10x.

**Statistical Analysis**

Data were reported as the mean ± standard deviation (SD) or mean ± standard error (SE) from at least n=3. Two-tailed Student’s t-test was used to analyze the differences between groups. A statistically significant difference was reported if p=0.05 or less.

**Results and Discussion**

**Synthesis and Characterization of Hydrogels**

UV initiated photo emulsion polymerization, using Irgacure 2959 as an initiator, offered the flexibility of fabricating the hydrogels under mild conditions such as low temperature (room temperature) and short reaction times (<10 mins). Also, the initiator Irgacure 2959 solution has been widely used for *in situ* polymerization *in vivo* and does not cause irritation [128]. The synthesis was successful and confirmed by FTIR spectra of hydrogels 19:1 - NIPAAm to macromer are shown in Figure 2-1. The hydrogels contain characteristic peaks from both PNIPAAm and DEX-Lactate-HEMA or HA-AEMA macromer, namely, the absorbance of the carbonyl group at 1640 cm$^{-1}$, N-H bending at 1538 cm$^{-1}$ and N-H stretching at 3300 cm$^{-1}$ are the characteristic peaks of NIPAAm. The divided bands of symmetric C-H bending from the -CH(CH$_3$)$_2$ group (1369 and 1367 cm$^{-1}$) are attributed to PNIPAAm, C-OH stretching (∼1020 cm$^{-1}$) due to the abundant hydroxyl groups of dextran backbone and C-H asymmetric stretching at 2974 cm$^{-1}$. All the OH and NH stretching were confined b/w 3000 -3500 cm$^{-1}$. The presence of more hydroxyl groups in dextran gives rise to the high-intensity peaks in the spectrum of DEX-Lac-HEMA.
Figure 2-1. FTIR spectrum of hydrogels

Notes: N-H stretching at 3287 cm$^{-1}$, C-H asymmetric stretching at 2974 cm$^{-1}$, absorbance of amide-carbonyl group at 1640 cm$^{-1}$, N-H bending at 1538 cm$^{-1}$, symmetric C-H bending from the -CH(CH$_3$)$_2$ group at 1367 cm$^{-1}$ and C-OH stretching at 1021 cm$^{-1}$
The early stage of optimization of the variables was primarily focused on P(NIPAAm-co-DEX-Lactate-HEMA) hydrogels. The endpoints or the observations that were monitored during the synthesis optimization include: i) State or physical appearance of the reaction mixture after photopolymerization if it is a liquid or solid gel, ii) State or physical appearance of the reaction mixture after heating at 37°C. The ideal characteristics of the gel include: being liquid at room temperature and turning into solid/gel upon heating to 37°C. The observations/comparisons of the formation of gels with the changes in parameters and the trends are mentioned in Table 2-1. When the DEX-Lactate-HEMA macromer concentration was increased from 5 to 20%, strong insoluble gels were formed, and this could be attributed to higher crosslinking in case of 16:4 hydrogels compared to 19:1.

The polymerization is proportional to the monomer concentration and inversely related to the initiator concentration. The higher the concentration of the polymer, stronger the polymer with higher molecular weight and more insoluble it could become. Similar results were observed when the prepolymer mixture concentration was increased, the gelation increased and the gel becomes more insoluble and resulted in a solid gel. However, the gels formed at 1mg·mL\(^{-1}\) concentration showed favorable results as they exhibited viscous liquid behavior and turned solid upon heating. Table 2-1 also presents the gelation process of the hydrogels with an increase in the initiator concentration as well. The size of the solid gels formed was same for 0.05 and 0.025 irgacure w% but it decreased with the 0.0125 irgacure w%, however, the yield was less with the decrease in the initiator concentration to 0.025 and 0.0125 wt%. Overall, with the decrease in the initiator (irgacure), the size of the solid gels obtained decreased and this could be attributed to the formation of insoluble higher chain length polymer at higher initiator concentration. The optimum concentration was found to be 0.05 w%. Based on the results from Table 2-1, it can also be observed that the gelation process increased with increase in UV intensity. 500 mW·cm\(^{-2}\) was selected and continued to characterize further.

*In vitro* cytotoxicity testing of the P(NIPAAm-co-DEX-Lactate-HEMA) gels (synthesized at 1 mg·mL\(^{-1}\), 500 mW·cm\(^{-2}\), 0.5 cm distance, 3 min reaction time) showed toxicity and had low percentage cell viability values (below 50%). In case of 19:1, the gels made at 0.0125 wt% initiator showed higher cytotoxicity (1.6 fold) than gels made at 0.05 and 0.025 wt% initiator. However, 16:4 gels showed a different behavior where the gels made at 0.05 and 0.0125 wt% were toxic than that of gels made at 0.025 wt%. Overall, all the gels were cytotoxic to the DPSCs, which could be attributed to lack of complete polymerization of the monomer/macromer mixture, and presence of any unreacted monomer/macromers that were toxic to the cells. In order to complete the polymerization, the reaction time (time of UV exposure) was increased from 3 to 10 min. and the resulted polymeric gels were compared with raw materials (PNIPAAm and DEX-Lactate-HEMA macromer) in terms of cytotoxicity to DPSCs. The raw materials were found to be toxic (%cell viabilities 40-80) to the DPSCs at all the studied concentrations 0.1, 10, 20 mg·mL\(^{-1}\). However, after polymerization P(NIPAAm-co-DEX-Lactate-HEMA) gels were found to be non-toxic at 0.1 mg·mL\(^{-1}\) concentration, still, the cell
Table 2-1. Effect of NIPAAm/DEX-Lactate-HEMA macromer ratio, UV intensity, precursor concentration, initiator concentration on the formation of P(NIPAAm-co-DEX-Lactate-HEMA) gels.

<table>
<thead>
<tr>
<th>Precursor concentration (mg/µL)</th>
<th>UV intensity (mW·cm⁻²)</th>
<th>Macromer*</th>
<th>NIPAAm/macromer (w/w)</th>
<th>Irgacure (w%)</th>
<th>Distance between tip of UV probe and surface of liquid (cm)</th>
<th>Product appearance</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>100</td>
<td>DEX</td>
<td>19:1</td>
<td>0.05</td>
<td>0.5</td>
<td>Liquid</td>
</tr>
<tr>
<td>1</td>
<td>250</td>
<td>DEX</td>
<td>19:1</td>
<td>0.05</td>
<td>0.5</td>
<td>Liquid</td>
</tr>
<tr>
<td>1</td>
<td>375</td>
<td>DEX</td>
<td>19:1</td>
<td>0.0125</td>
<td>0.5</td>
<td>Liquid + small soft gel</td>
</tr>
<tr>
<td>1</td>
<td>375</td>
<td>DEX</td>
<td>19:1</td>
<td>0.025</td>
<td>0.5</td>
<td>Liquid + soft gel</td>
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<tr>
<td>0.5</td>
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<td>DEX</td>
<td>19:1</td>
<td>0.05</td>
<td>0.5</td>
<td>Soft gel</td>
</tr>
<tr>
<td>1</td>
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<td>DEX</td>
<td>19:1</td>
<td>0.05</td>
<td>0.5</td>
<td>Liquid + soft gel</td>
</tr>
<tr>
<td>2</td>
<td>375</td>
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<tr>
<td>1</td>
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</tr>
<tr>
<td>1</td>
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</tr>
<tr>
<td>1</td>
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<td>16:4</td>
<td>0.025</td>
<td>0.5</td>
<td>Liquid + small soft gel</td>
</tr>
<tr>
<td>0.5</td>
<td>375</td>
<td>DEX</td>
<td>16:4</td>
<td>0.05</td>
<td>0.5</td>
<td>Liquid + soft gel</td>
</tr>
<tr>
<td>1</td>
<td>375</td>
<td>DEX</td>
<td>16:4</td>
<td>0.05</td>
<td>0.5</td>
<td>Soft gel</td>
</tr>
<tr>
<td>2</td>
<td>375</td>
<td>DEX</td>
<td>16:4</td>
<td>0.05</td>
<td>0.5</td>
<td>Soft gel</td>
</tr>
<tr>
<td>1</td>
<td>750</td>
<td>DEX</td>
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<td>0.05</td>
<td>0</td>
<td>Soft gel</td>
</tr>
<tr>
<td>1</td>
<td>750</td>
<td>HA</td>
<td>19:1</td>
<td>0.05</td>
<td>0</td>
<td>Soft gel</td>
</tr>
<tr>
<td>1</td>
<td>750</td>
<td>HA</td>
<td>19:1</td>
<td>0.05</td>
<td>0</td>
<td>Soft gel</td>
</tr>
<tr>
<td>1</td>
<td>1000</td>
<td>DEX</td>
<td>19:1</td>
<td>0.05</td>
<td>0</td>
<td>Strong soft gel</td>
</tr>
<tr>
<td>1</td>
<td>1000</td>
<td>DEX</td>
<td>19:1</td>
<td>0.05</td>
<td>0</td>
<td>Strong soft gel</td>
</tr>
<tr>
<td>1</td>
<td>1000</td>
<td>DEX</td>
<td>19:1</td>
<td>0.05</td>
<td>0</td>
<td>Strong soft gel</td>
</tr>
<tr>
<td>1</td>
<td>1000</td>
<td>HA</td>
<td>19:1</td>
<td>0.05</td>
<td>0</td>
<td>Strong soft gel</td>
</tr>
<tr>
<td>1</td>
<td>1000</td>
<td>HA</td>
<td>19:1</td>
<td>0.05</td>
<td>0</td>
<td>Strong soft gel</td>
</tr>
<tr>
<td>Notes: Macromer*: DEX: DEX-Lactate-HEMA; HA: HA-AEMA; NIPAAm: Macromer = 19:1 (or) 16:4 (w:w), Concentration: 0.5,1,2 mg·mL⁻¹; UV intensity: 100, 250, 375, 500, 750, 1000 mW/cm² Distance between tip of the UV probe and mold: 0, 0.5 cm</td>
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</table>
viability values decreased with increased polymer concentration. Increasing the reaction time alone did not result in reduced cytotoxicity of the hydrogels. The synthetic conditions used might have resulted in partial or incomplete polymerization and presence of any unreacted starting materials, because of these the gels were found to be toxic. Therefore, the gel synthesis parameters needed more optimization.

To further optimize the gels to improve the polymerization and to lower the cytotoxicity, the variables that are altered during the synthesis include i) distance between the UV probe and the tip of the liquid surface. Instead of using 0.5 cm distance, the gap between the UV probe and the mold surface was completely eliminated and kept at 0 cm distance; ii) the UV intensity was increased from 500 mW·cm$^{-2}$ to 750 mW·cm$^{-2}$; and iii) the dialysis time was increased from 4 h to 6 h to remove any unreacted raw materials. The gels obtained were exposed to DPSCs to estimate the in vitro cytotoxicity. Figure 2-2 illustrates the cell viability of gels (synthesized at different initiator levels) to DPSCs at three different concentrations 0.5, 1 and 10 mg·mL$^{-1}$. Independent of the initiator percentages (0.05, 0.025, 0.0125 wt%) used in the synthesis, elimination of gap, increase in UV intensity and dialysis time had shown a drastic improvement in the cell viability values of the gels. The cell viabilities improved from 40-60% to 90-100%. Removing the gap and increasing the UV intensity might have increased the polymerization, and increasing the dialysis time might have led to the removal of any unreacted starting materials, and this could have resulted in the improved cell viability of the gels. Overall, the gels made with 0.05 w% initiator concentration has shown relatively higher yield when compared with the 0.025 and 0.0125 w%. Increased initiator concentration could lead to higher polymer chain length and as result, the hydrogels with 0.05 w% initiator concentration could have formed the gels with higher yield when compared with lower initiator concentration.

**Synthesis and Comparison of Hydrogels Based on HA-AEMA Macromer**

After optimizing the synthetic parameters of the injectable hydrogels, another macromer based on hyaluronic acid was investigated for synthesizing the hydrogels and compared the properties with dextran-based hydrogels. The P(NIPAAm-co-HA-AEMA) hydrogels were synthesized at NIPAAm/Macromer of 19:1 (w:w) with prepolymer concentration of 1 mg·mL$^{-1}$ by direct exposure to UV probe (without gap) at two different intensities (750 and 1000 mW·cm$^{-2}$) and three different initiator concentrations (0.05, 0.025, 0.0125 w%). The gels synthesized were observed for their physical appearance and compared with P(NIPAAm-co-DEX-Lactate-HEMA). The observations are mentioned in Table 2-1. When observed for the overall trend, both the gels formed at higher intensity resulted in a stronger solid gel formation before dialysis. However, all the gels (polymerization mixture) after dialysis yielded a viscous gel. No other significant differences were found in terms of the physical appearance of the gels.
Figure 2-2. Cell viability of the P(NIPAAM-co-DEX-LactateHEMA) gels to DPSCs.

Notes: Macromer: DEX: DEX-Lactate-HEMA; NIPAAm: Macromer = 19:1 (w:w), Concentration : 1mg·mL⁻¹; UV intensity : 500, 750 mW/cm²; Distance between tip of the UV probe and mold : 0 cm.
Rheological Properties

The rheological property of hydrogels needs to be evaluated for their rheological behavior within their range of linear viscoelastic region to ensure their stability [129]. Figure 2-3 illustrates the linear viscoelastic region of the P(NIPAAm-co-DEX-Lactate-HEMA) and P(NIPAAm-co-HA-AEMA) gels under oscillatory strain ramp. The $G'$ of hydrogels remains unaffected by strain amplitude (linear viscoelastic region) for $2-20\%$ and $2-100\%$ (Figure 1-3) for every Dextran-based and HA-based gels respectively. The gels exhibited similar behavior at both $10$ and $15\, \text{mg}\cdot\text{mL}^{-1}$ concentrations. In case of strain sweep, when the concentration of the hydrogels was increased from $10$ to $15\, \text{mg}\cdot\text{mL}^{-1}$, the $G'$ was increased about $2.4$ fold for dextran-based hydrogels and $3.2$ fold for HA-based hydrogels. At $15\, \text{mg}\cdot\text{mL}^{-1}$, the value of $G'$ at $1\, \text{Hz}$ appears in the range of $20-40\, \text{Pa}$ for HA-based gels and $15-35\, \text{Pa}$ for dextran-based gels. Since both the hydrogels have shown linear viscoelasticity at $10\%$ strain, in order to ensure that measurements are done within the linear viscoelastic region, all subsequent rheological characterization was performed at $10\%$ strain. Frequency sweep studies were conducted to determine the characteristic gel behavior ($G'>G''$). Figure 2-4 shows the $G'$ and $G''$ of the hydrogels as a function of frequency. The $G'$ was always higher than $G''$ suggesting that the elastic behavior dominates the viscous behavior in these hydrogels, which is the typical characteristic of a hydrogel. After about $2\, \text{Hz}$ frequency, both the $G'$ and $G''$ increased linearly with increasing frequency, suggesting that both types of hydrogels were in the glass transition stage and became stiffer with increasing frequency. In case of HA-or Dextran-based hydrogels, increase in concentration from $10$ to $15\, \text{mg}\cdot\text{mL}^{-1}$ has resulted in a significant increase in $G'$ and $G''$ values ($2-4$ fold). No significant differences were found for the hydrogels prepared with the HA or dextran at the same concentration, as the majority of the polymer is NIPAAm (95%). The individual figures of strain sweep, frequency sweep and temperature ramp for each hydrogel are given in the Appendix (Figure A-1 through Figure A-12).

Sol-Gel Transition of Hydrogels

The feasibility of P(NIPAAm-co-DEX-Lactate-HEMA) hydrogels for using as injectable in situ hydrogels for tissue engineering, a sol-gel transition study was conducted to evaluate their behavior in presence of stem cells. The solutions of gels ($10\, \text{mg/mL}$ and $20\, \text{mg/mL}$) encapsulated with DPSCs were immediately transformed (gelation time $<2\, \text{min}$) into solid gels when exposed to $37^\circ\text{C}$ (Figure 2-5). The temperature-dependent transition of hydrogels was studied by monitoring the storage modulus values as a function of temperature on a rheometer. Followed by a plateau regime, rapid buildup regime was observed in $G'$ (Figure 2-6), and this fast increase in $G'$ was attributable to gelling (sol to gel transition) or 3D crosslink network formation of the polymeric chains. Because of the strong hydrophobic interactions between the PNIPAAm chains above LCST, the PNIPAAm exhibit a phase transition or thermoresponsive behavior. The midpoint temperature ($\sim35\, ^\circ\text{C}$) of the transition is about the same for all samples.
Figure 2-3. Representative shear storage modulus (G’) and shear loss modulus (G’’') as a function of oscillatory strain sweep for the hydrogels.

Figure 2-4. Representative shear storage modulus (G’) and shear loss modulus (G’’') as a function of frequency for the hydrogels.
Figure 2-5. The thermoresponsive sol-gel transition of the hydrogels.

Figure 2-6. Thermoresponsive phase transition and representative change in shear storage modulus ($G'$) and shear loss modulus ($G''$) as a function of temperature ramp for the hydrogels.
Cell Survival in Hydrogels

To assess the cytotoxicity of DPSCs and ADSCs encapsulated in Dex and HA-based hydrogels in vitro, the MTT assay was utilized. The DPSCs & ADSCs viability in Dex-based hydrogels with a concentration of 10, 15, 20 mg·mL⁻¹ was about 117±7.4, 106±7.8, 79±5.3 & 110±7.8, 99±6.6, 80±7.3 % respectively (Figure 2-7). The DPSCs & ADSCs viability in HA-based hydrogels with a concentration of 10, 15, 20 mg·mL⁻¹ was about 122±6.6, 103±12.8, 91±7.1 & 143±9.7, 132±9.1, 90±9.4 % respectively (Figure 2-8). There was a statistically significant decrease in the cell viability for the Dex-based hydrogels between concentrations of 10 and 15, 15 and 20 mg/mL for the ADSCs (p<0.05). DPSCs cell viabilities in Dex-based hydrogels decreased significantly between 15 and 20 mg/mL (p<0.01). In HA-based hydrogels the ADSC viabilities decreased significantly between the concentrations 10 and 15 (p<0.05), 15 and 20 mg/mL (p<0.01). In addition to the cell viability by MTT, the percentage live cell number of the DPSCs and ADSCs after 24hrs of encapsulation in the hydrogels was determined using DAPI (Figure 2-9 and Figure 2-10). The DPSCs & ADSCs percent cell number compared to control in Dex-based hydrogels with a concentration of 10, 15, 20 mg·mL⁻¹ was about 112±1.58, 102±1.22, 80±0.71 & 121±2.92, 97±1.41, 89±0.71 % respectively (Figure 2-11). The DPSCs & ADSCs percent cell number in HA-based hydrogels with a concentration of 10, 15, 20 mg·mL⁻¹ was about 127±3.39, 107±0.71, 82±1.22 & 134±3.67, 97±0.71, 89±1.87 % respectively (Figure 2-12). There was a statistically significant decrease in the ADSCs number for the Dex-based hydrogels between 10 & 15, 15 & 20 mg/mL (p<0.01). In Dex and HA-based hydrogels the DPSCs number decreased significantly between the concentrations 10 & 15 (p<0.05), 15 & 20 mg/mL (p<0.01). In HA-based hydrogels the ADSC viabilities decreased significantly between 10 and 15 mg/mL (p<0.01). Cell survival studies based on MTT and DAPI live cell counting demonstrated that both DEX-based and HA-based based hydrogels showed promising cell viability and cell number values (~100% and above) at concentrations of 10 and 15 mg/mL for DPSCs and ADSCs. The results from the MTT assay and cell number studies by DAPI staining were in accordance with each other and confirm that the Dex & HA hydrogels are suitable for the encapsulation of ASDSCs & DPSCs at 10 and 15 mg/ml concentrations.

Conclusions

A series of biodegradable hydrogels were synthesized by photopolymerization of NIPAAm monomer and hydrolytically degradable oligolactate DEX-Lactate-HEMA or HA-AEMA macromers. The thermoresponsive, rheological and biodegradable properties of the hydrogels were strongly dependent on the reaction conditions and composition of the hydrogels. The higher the amount of the Irgacure 2959 initiator, the higher the UV intensity, the longer the reaction time, and the higher amounts of crosslinker used for the synthesis, the higher the yield of the synthesized hydrogels. The hydrogels were not cytotoxic to DPSCs and ADSCs at a concentration up to at least 15 mg·mL⁻¹, and have appropriate mechanical and biodegradable properties to promote DPSCs and ADSCs growth. Based on in vitro studies it was observed that DEX-gels and HA-gels appear to
Figure 2-7. Cell viability of DPSCs & ADSCs in injectable NIPAAm-DEX-Lactate-HEMA hydrogels using MTT assay.

Notes: NIPAAm:DEX-Lactate-HEMA - 19:1, Initiator – 0.05%, UV - 750mW·cm−2, UV exposure time - 10 min, Distance of UV probe from the mold – 0 cm. (Mean ± SD, n=3; *: p-value < 0.05, #: p-value < 0.01)

Figure 2-8. Cell viability of DPSCs & ADSCs in injectable NIPAAm-HA-AEMA hydrogels using MTT assay.

Notes: NIPAAm:HA-AEMA - 19:1, Initiator – 0.05%, UV -750mW·cm−2, UV exposure time - 10 min, Distance of UV probe from the mold – 0 cm. (Mean ± SD, n=3; *: p-value < 0.05, #: p-value < 0.01)
Figure 2-9. DPSCs growth and morphology in injectable P(NIPAAm-co-DEX-Lactate-HEMA) and P(NIPAAm-co-HA-AEMA) hydrogels.

Notes: P(NIPAAm-co-DEX-Lactate-HEMA) – represented as DEX in Figure (Synthetic conditions: NIPAAm:DEX-Lactate-HEMA-19:1, Initiator – 0.05%, UV – 750 mW·cm², UV exposure time - 10 min, Distance of UV probe from the mold – 0 cm; Cells staining – DAPI);
P(NIPAAm-co-HA-AEMA) – represented as HA in Figure (NIPAAm:HA-AEMA-19:1, Initiator – 0.05%, UV – 750 mW·cm², UV exposure time - 10 min, Distance of UV probe from the mold - 0cm; Cells staining – DAPI)
Figure 2-10. ADSCs growth and morphology in injectable P(NIPAAm-co-DEX-Lactate-HEMA) and P(NIPAAm-co-HA-AEMA) hydrogels.

Notes: P(NIPAAm-co-DEX-Lactate-HEMA) – represented as DEX in Figure (Synthetic conditions: NIPAAm:DEX-Lactate-HEMA-19:1, Initiator – 0.05%, UV – 750 mW·cm$^{-2}$, UV exposure time - 10 min, Distance of UV probe from the mold - 0cm; Cells staining – DAPI);
P(NIPAAm-co-HA-AEMA) – represented as HA in Figure (NIPAAm:HA-AEMA-19:1, Initiator – 0.05%, UV – 750 mW·cm$^{-2}$, UV exposure time - 10 min, Distance of UV probe from the mold - 0cm; Cells staining – DAPI)
Figure 2-11. Cell count of DPSCs & ADSCs in injectable NIPAAm-DEX-Lactate-HEMA hydrogels by DAPI staining.

Notes: NIPAAm: DEX-Lactate-HEMA - 19:1, Initiator – 0.05%, UV – 750mW·cm⁻², UV exposure time - 10 min, Distance of UV probe from the mold – 0 cm. (Mean ± SD, n=4; *: p-value < 0.05, #: p-value < 0.01)

Figure 2-12. Cell count of DPSCs & ADSCs in Injectable NIPAAm-HA-AEMA Hydrogels by DAPI staining.

Notes: NIPAAm: HA-AEMA - 19:1, Initiator – 0.05%, UV – 750mW·cm⁻², UV exposure time - 10 min, Distance of UV probe from the mold – 0 cm. (Mean ± SD, n=4; *: p-value < 0.05, #: p-value < 0.01)
be suitable for encapsulation of DPSCs/ADSCs at optimum hydrogel concentration. The data confirmed that both DPSCs and ADSCs favorably adhered and proliferated within the biomaterials in vitro. Additionally, the current study confirmed that DPSCs and ADSCs have superior growth properties in HA-AEMA hydrogels compared to DEX-Lactate-HEMA hydrogels. The developed hydrogels have shown promising potential for encapsulating DPSCs/ADSCs for tissue engineering applications. However, further optimization is needed to have appropriate mechanical and degradation properties to cater as per DPSCs or ADSCs growth inside their matrices, and successful completion of this project will have a significant impact on dental and retinal tissue engineering fields.
CHAPTER 3. RHEOLOGICAL PROPERTIES OF THE PORCINE OCULAR TISSUES AT DIFFERENT STAGES OF DEVELOPMENT

Introduction

The eye is a complex and delicate organ being divided into two anatomical segments called the anterior (comprises of cornea, conjunctiva, iris, lens, ciliary body, and the anterior part of the sclera) and posterior (comprises of vitreous humor, retina, choroid, and posterior part of the sclera) segments [130]. Cornea, sclera and vitreous humor are the three tissues that were investigated in this study for the rheological properties. The cornea is a specialized transparent tough tissue covering the front portion of the eye and protects the sensitive internal contents of the eye and is responsible for refracting the light [131]. The tissue is under constant stress because of various forces such as external (eye rubbing, poking, eyelid closure) and internal (intraocular pressure because of aqueous/vitreous flow dynamics). Cornea’s mechanical behavior, fundamental for maintaining its dimensional stability, refractability and vision, depends on the thickens, topography and biomechanical properties of the tissue [28]. The extracellular matrix of the cornea called the stroma forming the bulk of the corneal thickness primarily contributes to its biomechanical properties [131]. The sclera is a dense connective tissue covering the five-sixths of the eye. The sclera is a strong and firm tissue and maintains the shape of the globe. It helps in the maintenance of intraocular pressure, provides attachment sites for the external eye muscles, and provides the internal eye structures from injuries and mechanical displacement. The vitreous body/vitreous humor is a transparent and gelatinous substance that fills the inner posterior cavity (space between the lens and the retina) of the eye. Vitreous humor is highly hydrated and comprised of 99% of water, avascular, and nearly acellular. The unique extracellular matrix of the vitreous humor, composed of a complex network of collagen fibers and hyaluronic acid (glycosaminoglycan), significantly contributes to the viscoelastic nature of the vitreous [132]. Vitreous plays a significant role in metabolic nutrient supply to the lens, coordinating eye growth (developmental) allows light to reach the retina (optical), providing support to the retina, maintain the shape of the eyeball and protects eye tissues from high-frequency stresses (mechanical) [132, 133].

The proper vision of the eye strongly depends on its ability to maintain optical clarity and structural integrity under routine variations in pressure, changes in humidity and temperature, constant muscular stress/strain and rapid sudden movements. Thus the proper functioning of the eye is dependent on its physical and mechanical properties, and altered chemical and mechanical states of ocular tissues could lead to abnormalities and may result in complete loss of the functionality. Various age-related changes in composition and structure of the eye tissues affect the biomechanical properties and performance of the eye tissues and could contribute to the predisposition of various eye abnormalities [21-26]. For example, aging results in stiffening of the cornea tissue which could be because of the additional age-related non-enzymatic cross-linking affecting the stromal collagen fibrils [24, 25]. It is reported in the literature that age-related variations affect the biomechanical characteristics of the sclera [21], and aging results in the
microstructural changes in the sclera and contribute to the predisposition to primary open-angle glaucoma [22]. With age, collagen networks in the vitreous begin to break down and pockets of liquid form in the eye, the process of liquefaction in which the liquid volume increases and gel volume decreases [23, 26]. These age-related changes could lead to structural and mechanical differences between vitreous at various stages of development, and in about 25-30% of the population, these age-related degenerative changes eventually result in posterior vitreous detachment [27]. Given the importance of the viscoelastic properties of the ocular tissues, to its function and to pathology, detailed rheological characterization of these tissues is crucial to appreciate the normal functions and quantitative understanding of disease pathophysiology of these tissues at various stages of development. The objective of this project was to characterize the rheological properties of the ocular tissues (cornea, sclera and vitreous) at different stages (preterm, neonatal and adult) of development. Though several measurements could be measured in vivo, such as topography, thickness distribution, but the attempts to investigate the biomechanical properties of the tissues still depend on ex-vivo experiments involving human or other animal tissues [28]. Because of the difficulties in obtaining human donor tissues in sufficient numbers for the research and experimental needs, it had been necessary to use animal ocular tissues as approximate models for human ocular tissues for performing various evaluations including the mechanical property characterization studies. Several investigations used porcine tissues for ocular related experiments [28-30], although other animal’s rabbit and bovine tissues a common practice. Therefore, the current rheological characterization also contributes to the literature for identifying and assessing the important similarities and significant differences between the rheological behaviors of porcine ocular tissues and human and other animal tissues and evaluating the suitability of the porcine eye tissues as a model of the human tissues.

**Hypothesis and Specific Aims**

The hypothesis of this project was that the storage and loss modulus of the porcine ocular tissues differ at different stages of development (preterm, neonatal and adult). There were two specific aims:

**Specific Aim 1**

To conduct the shear-sweep experiments and identify the linear viscoelastic region of the porcine ocular tissues (cornea, sclera, vitreous) at different stages of development (preterm, neonatal, adult).

**Specific Aim 2**

To conduct the frequency sweep experiment and analyze the gel-like behaviors of the porcine ocular tissues (cornea, sclera, vitreous) at different stages of development (preterm, neonatal, adult).
Materials and Methods

Materials

Sodium Chloride (NaCl), potassium chloride (KCl), calcium chloride (CaCl₂), magnesium chloride (MgCl₂), sodium dihydrogen phosphate (NaH₂PO₄), sodium bicarbonate (NaHCO₃), sodium hydroxide (NaOH) and Hydrochloric acid (HCl) were bought from Sigma Aldrich. Porcine eyes tissues were obtained from Dr. Steven Davison and Dr. Cameron Fili.

Sample Preparation

Fresh porcine eyeball tissues were provided by Dr. Steven Davison and Dr. Cameron Fili. A circular trephine punch from Katena was used to punch specimens of diameter 8 mm from the central region of the cornea and from the region below optic nerve in the sclera. AR-G2 TA Instruments Rheometer with accompanying Rheology Advantage Instrument Control software was used to perform the experiments. Prior to testing, the specimens were equilibrated in GBR solution (bicarbonate-Ringer's solution (BR) containing reduced glutathione) (NaCl, 6.2 g/L; KCl, 0.358 g/L; CaCl₂, 0.115 g/L; MgCl₂, 0.159 g/L; NaH₂PO₄, 0.103 g/L; NaHCO₃, 2.454 g/L; glucose, 0.90 g/L; reduced glutathione, 0.92 gm./L and pH was adjusted to 7.4 using Sodium hydroxide (NaOH) and Hydrochloric acid (HCl) [134], which has been shown to maintain endothelial function for up to 6 hr., was added to the immersion cell bathing either side of the cornea tissue. The thickness of the specimens was measured using screw gauge. The sample was placed on the lower geometry, and then the upper geometry was lowered to contact the sclera. The gap between the parallel plates was maintained at the thickness of the sample tissues used and the experiments were done. In this experimental method, circular specimens of sclera and cornea tissues are sheared between two parallel plates. In order to characterize the shear viscoelastic properties, two types of oscillatory tests, i.e. the strain and frequency sweep experiments, were conducted. Although, it is known that the cornea and sclera tissues have an inhomogeneous microstructure through the thickness, and the present study did not consider these inhomogeneities and its findings should only be considered as the macroscale (effective) shear properties of the tissue. However, care was taken while sample preparation to be consistent. For characterizing the range of linear viscoelasticity, the strain sweep experiments (4 samples) were done at frequency 1Hz and force 0.01 N. For frequency sweep experiments (4 samples), the frequency of 0.01–100 Hz and a shear strain magnitude of 0.05 % (which was within the region of linear viscoelasticity) were selected. The age and weight of pigs are mentioned in Table 3-1.
Table 3-1. Age and weight of pigs used in the study.

<table>
<thead>
<tr>
<th>Pig</th>
<th>Age</th>
<th>Weight</th>
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<tr>
<td>Preterm Pig</td>
<td>10 days prior</td>
<td>0.5 - 1.5 Kg</td>
</tr>
<tr>
<td>Neonatal Pig</td>
<td>20 days</td>
<td>2.5 - 3.5 Kg</td>
</tr>
<tr>
<td>Adult Pig</td>
<td>1 year</td>
<td>80 - 100 Kg</td>
</tr>
</tbody>
</table>
**Rheological Properties**

Strain sweep oscillation was performed to determine the limit of linear viscoelasticity. Four preconditioned specimens were tested at 1 Hz frequency. For the determination of linear viscoelastic region (where $G'$ and $G''$ remain constant), strain sweep oscillation was performed at constant frequency with varying strain amplitude from 0.01 to 100%. Unlike the conventional solid mechanical definition, strain here is defined as the excursion of the circumference of the disk-shaped specimen divided by its thickness. Rheological properties such as $G'$ and $G''$ remain constant within the limit of linear viscoelasticity. After the determination of tissue’s viscoelastic region by a strain sweep, the tissues were further characterized by a frequency sweep at a strain in the linear viscoelastic region to understand the gel behaviors of the tissues ($G'>G''$). The viscosity of the whole vitreous was measured at a shear rate 0.1 to 1000 s$^{-1}$ under continuous flow mode.

**Results and Discussion**

**Figures 3-1, 3-2 and 3-3** show the range of linear viscoelasticity of the porcine cornea, sclera and vitreous as a function of strain sweep (at 1 Hz frequency) at different ages. These figures confirm that the studied tissues, like other soft tissues, showed a viscoelastic behavior. The strain sweep experiments on all the tissues at different ages showed that the measured storage modulus was nearly constant over small amplitude shear strain (%) and decreased nonlinearly with increasing percent strain after a turnover point, called critical shear strain level. Two different regions were clearly differentiated: a linear viscoelastic region where $G'$ and $G''$ were practically constant (small deformation), and a nonlinear region, in which $G'$ and $G''$ began to diminish with increasing strain (large deformations) [135]. Within the range of linear viscoelastic response, the viscoelastic parameters are independent of the shear strain amplitude. The strain independent region indicates that extent of strain applied is small enough not to cause any perturbation in the structure of the material, and in biological samples, these linear region includes normal physiological loads and represent normal *in vivo* response [136]. The individual figures of strain sweep, frequency sweep and viscosity for each tissue or vitreous are given in the **Appendix** (Figure A-13 through Figure A-33).

In the linear viscoelastic region (**Figure 3-1**), the values of $G'$ and $G''$ for all corneal tissues were constant below 2% strain and decreased logarithmically with increasing strain (>2% strain). Linear region for the corneal samples was found to be 0.01 – 2% strain. The consistent linear regions of both $G'$ and $G''$ were observed for all the corneal samples at different ages. Similar to the cornea, scleral tissues (**Figure 3-2**) also behaved as gel-like and exhibited linear viscoelastic region below 3% strain in preterm and neonatal, and 0.1% strain in adult pigs, the critical strain related to the linear viscoelastic region in sclera tissues was different at different ages. Vitreous also exhibited gel-like behavior with a linear viscoelastic region (**Figure 3-3**). It was reported that the gel-like physical properties of vitreous are primarily derived from its chemical composition which includes a hydrated cross-linked network of fibrillar collagens (type)
Figure 3-1. Storage and loss moduli of corneal tissues at preterm, neonatal and adult stages of development under strain-sweep.

Figure 3-2. Storage and loss moduli of scleral tissues at preterm, neonatal and adult stages of development under strain-sweep.
Figure 3-3. Storage and Loss moduli of vitreous humor at preterm, neonatal and adult stages of development under strain-sweep.
and higher-molecular-weight polyanionic hyaluronan macromolecules, which is consistent with the literature [137].

Once the range of linear viscoelastic region was determined, frequency sweep experiments were performed on the samples with a strain amplitude that was within the range of linearity for each sample (Figures 3-4, 3-5 and 3-6). In corneal samples, the moduli increased weakly with increasing frequency (Figure 2-2a), and this trend was similar to the response of corneal stroma [131]. This behavior also related to other tissues such as articular cartilage, meniscus, and intervertebral disk [131, 138-140]. The magnitude of change (increase in storage modulus) increased significantly at a higher frequency (>5 Hz). Adult cornea tissue has shown higher \( G' \) and \( G'' \) compared to the preterm and neonatal corneal tissues. At 1 Hz frequency, the \( G' \) values of preterm, neonatal and adult cornea were ~0.89, 0.97 and 1.22 kPa, respectively. Similar to the cornea, scleral tissues (Figure 3-5) also showed a slight increase in moduli with an increase in the frequency in the low-frequency range (<10 Hz), and the magnitude of increase in the moduli changed significantly at a higher frequency (>10 Hz). The moduli values of sclera increased with increase in the age (preterm<neonatal<adult). At 1 Hz frequency, the \( G' \) values of preterm, neonatal and adult sclera were ~0.96, 1.81 and 16.8 kPa, respectively. In frequency dependence studies, the storage modulus of vitreous varied weakly with frequency up to ~0.4 Hz (Figure 3-6) and raised significantly thereafter. This frequency dependence behavior is characteristic of a lightly crosslinked amorphous network of macromolecules and is consistent with the literature [136]. The storage modulus of the adult vitreous was higher at low frequency (<0.2 Hz) but were similar thereafter to neonatal and preterm samples. At 1 Hz frequency, the \( G' \) values of preterm, neonatal and adult vitreous were ~4.5, 4.8 and 8.8 Pa, respectively.

In frequency dependence studies, all three types of tissues showed a similar behavior of weak variation in moduli with increased frequency. Also, it can be observed that irrespective of the age, all tissue samples showed \( G' > G'' \), which represents that elastic properties of the materials dominate the viscous portion and the samples exhibited solid-like behavior. The difference between \( G' \) and \( G'' \) of corneal and scleral samples is much higher compared to the vitreous humor, and the magnitude of \( G'/G'' \) among the tissues was in the order of cornea > sclera >> vitreous. Thus, the cornea and sclera are more gel-like than the vitreous humor, as the elastic character of the tissue is manifested in the storage modulus.

When the vitreous was ran under shear rate-controlled experiments (Figure 3-7), the viscosity of the vitreous decreased with increasing shear-rate exhibiting a shear-thinning behavior (non-Newtonian) with a value around 39.5±2.7, 40.3±9.1, 54.8±1.2 for a shear rate of 0.93 s\(^{-1}\) and reaching a value around 3.2±0.1, 3.4±0.8, 4.1±0.1 for shear rate of 100 s\(^{-1}\) for preterm, neonatal, and adult porcine vitreous, respectively. At low shear rates (~1 s\(^{-1}\)), the viscosity of the adult vitreous was significantly higher than the neonatal and preterm porcine vitreous. The shear-thinning behavior of the vitreous was consistent with the literature [141]. It is reported that this behavior could be attributed to the presence of HA (primary component) in the liquid portion of the vitreous [141]. HA solutions exhibit this shear thinning behavior [142].
Figure 3-4. Storage and loss moduli of corneal tissues at preterm, neonatal and adult stages of development under frequency-sweep.

Figure 3-5. Storage and loss moduli of scleral tissues at preterm, neonatal and adult stages of development under frequency-sweep.
Figure 3-6. Storage and Loss moduli of vitreous humor at preterm, neonatal and adult stages of development under frequency-sweep.

Figure 3-7. The viscosity of vitreous humor with the shear rate at preterm, neonatal and adult stages of development.
Conclusions

This research presented the data on the effects of shear strain sweep and frequency sweep on rheological properties of the porcine ocular tissues, and compared the differences in tissues at different stages of development. Storage and loss modulus of porcine ocular tissues (cornea, sclera and vitreous) differed at different stages of development (adult, neonatal and preterm), adult tissues being on the higher end of G’ and G”. Nevertheless, future studies are required to fully characterize the mechanical behaviors of the ocular tissues at different stages of development covering the entire age range, and with different diseases. A thorough understanding of the shear properties of the ocular tissues would facilitate in developing better treatments for complex diseases or abnormalities.
A series of biodegradable hydrogels were synthesized by photopolymerization of NIPAAm monomer and hydrolytically degradable oligolactate DEX-Lactate-HEMA or HA-AEMA macromers. The thermoresponsive, rheological and biodegradable properties of the hydrogels were strongly dependent on the reaction conditions and composition of the hydrogels. The higher the amount of the Irgacure 2959 initiator, the higher the UV intensity, the longer the reaction time, and the higher amounts of crosslinker used for the synthesis, the higher the yield of the synthesized hydrogels. The hydrogels were not cytotoxic to DPSCs and ADSCs at a concentration up to at least 15 mg·mL\(^{-1}\), and have appropriate mechanical and biodegradable properties to promote DPSCs and ADSCs growth. Based on in vitro studies it was observed that DEX-gels and HA-gels appear to be suitable for encapsulation of DPSCs/ADSCs at optimum hydrogel concentration. The data confirmed that both DPSCs and ADSCs favorably adhered and proliferated within the biomaterials in vitro. Additionally, the current study confirmed that DPSCs and ADSCs have superior growth properties in HA-AEMA hydrogels compared to DEX-Lactate-HEMA hydrogels. The developed hydrogels have shown promising potential for encapsulating DPSCs/ADSCs for tissue engineering applications. However, further optimization is needed to have appropriate mechanical and degradation properties to cater as per DPSCs or ADSCs growth inside their matrices, and successful completion of this project will have a significant impact on dental and retinal tissue engineering fields.

This research presented the data on the effects of shear strain sweep and frequency sweep on rheological properties of the porcine ocular tissues, and compared the differences in tissues at different stages of development. Storage and loss modulus of porcine ocular tissues (cornea, sclera and vitreous) differed at different stages of development (adult, neonatal and preterm), adult tissues being on the higher end of G' and G". Nevertheless, future studies are required to fully characterize the mechanical behaviors of the ocular tissues at different stages of development covering the entire age range, and with different diseases. A thorough understanding of the shear properties of the ocular tissues would facilitate in developing better treatments for complex diseases or abnormalities.
LIST OF REFERENCES

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Figure A-1. Representative shear storage modulus ($G'$) and shear loss modulus ($G''$) as a function of oscillatory strain sweep for the HA-based 15 mg·mL$^{-1}$ hydrogel.
Figure A-2. Representative shear storage modulus (G’) and shear loss modulus (G’’) as a function of oscillatory strain sweep for the HA-based 10 mg·mL⁻¹ hydrogel.
Figure A-3. Representative shear storage modulus ($G'$) and shear loss modulus ($G''$) as a function of oscillatory strain sweep for the Dex-based 15 mg·mL$^{-1}$ hydrogel.
Figure A-4. Representative shear storage modulus (G’) and shear loss modulus (G’”) as a function of oscillatory strain sweep for the Dex-based 10 mg·mL$^{-1}$ hydrogel.
Figure A-5. Representative shear storage modulus (G') and shear loss modulus (G'') as a function of frequency for the HA-based 15 mg·mL⁻¹ hydrogel.
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Figure A-14. Storage and loss moduli of corneal tissues at neonatal stage of development under strain-sweep.
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Figure A-17. Storage and loss moduli of scleral tissues at neonatal stage of development under strain-sweep.
Figure A-18. Storage and loss moduli of scleral tissues at adult stage of development under strain-sweep.
Figure A-19. Storage and Loss moduli of vitreous humor at preterm stage of development under strain-sweep.
Figure A-20. Storage and Loss moduli of vitreous humor at neonatal stage of development under strain-sweep.
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Figure A-23. Storage and loss moduli of corneal tissues at neonatal stage of development under frequency-sweep.
Figure A-24. Storage and loss moduli of corneal tissues at adult stage of development under frequency-sweep.
Figure A-25. Storage and loss moduli of scleral tissues at preterm stage of development under frequency-sweep.
Figure A-26. Storage and loss moduli of scleral tissues at neonatal stage of development under frequency-sweep.
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Figure A-29. Storage and Loss moduli of vitreous humor at neonatal stage of development under Frequency-sweep.
Figure A-30. Storage and Loss moduli of vitreous humor at adult stage of development under Frequency-sweep.
Figure A-31. The viscosity of vitreous humor with the shear rate at preterm stage of development.
Figure A-32. The viscosity of vitreous humor with the shear rate at neonatal stage of development.
Figure A-33. The viscosity of vitreous humor with the shear rate at adult stage of development.
VITA

Mrs. Sri Chandana Reddy Damera was born in Karimnagar District, Telangana, India in 1990. She received her Bachelor of Pharmacy degree from Teegala Ram Reddy College of Pharmacy, Jawaharlal Nehru Technological University in 2011. She received her Masters in Pharmaceutical Chemistry degree in 2013 from Osmania University. After the acceptance of this thesis, she will have earned a Master’s Degree in Pharmaceutical Sciences. During her time at UTHSC, she has presented her work in conferences including AAPS, BMES, and ARVO. She also served as Secretary for AAPS UTHSC Student Chapter (Aug 2016 – May 2017)