

# OFFICIAL ABSTRACT and CERTIFICATION

## Synthesis and Analysis of a Novel Biodegradable Polyester Fiber Scaffold Derived from Poly(glycerol sebacate)

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Poly(glycerol sebacate) (PGS) is a biodegradable elastomer with elastic properties advantageous for use as a scaffold in soft tissue engineering. However, it has been unable to be electrospun into fibrous scaffolds—uncured PGS is a viscous liquid at room temperature and is therefore unable to hold any fiber shape, while cured PGS, though solid at room temperature, isn't easily dissolved, a requirement for electrospinning—and requires copolymerization that confers undesirable properties on the copolymer. The molecular weight of PGS has been shown to be 1110 g/mol, but fibrous structures are formed in other polymers above over 13000 g/mol.

The aim of this study was to increase the molecular weight of PGS by modification of the synthesis reaction in order to allow for electrospinning. PGS was copolymerized with 1,8-octanediol in monomer ratios of 1:1:2, 1:3:4, and 1:4:5 (glycerol:1,8-octanediol:sebacic acid) and catalyzed by the enzyme Novozym 435. The resulting poly(1,8-octanediol-glycerol sebacate) (POGS) polymers exhibited molecular weights 46 to 74.5 times greater and melting temperatures greater than PGS and above body temperature. POGS was electrospun but did not form fibers visible to the naked eye under the first set of parameters used. Altering applied voltage, the solvent, polymer concentration, or other parameters may form definitive fibers.

The improvement in molecular weight and thermal properties demonstrates potential, and as properties varied with monomer ratio, this approach could be used to fine tune the properties of the resulting polymers for the formation of elastic tissue scaffolds.

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