Controlled Functionalization of Poly(4-methyl-1-pentene) Films for High Energy Storage

<u>G. Wang¹</u>, M. Zhang², Z Xu¹ and L. Zhang^{1,3,*}

¹Electronic Materials Research Laboratory, Key Laboratory of the Ministry of Education and International Center for Dielectric Research, Xi'an Jiaotong University, Xi'an 710049, China.

Center for Dielectric Research, XI an Jiaolong University, XI an /10049, China.

²College of Materials Science and Engineering, Beijing Institute of Petrochemical Technology,

Beijing, 102617, China

²Department of NanoEngineering, University of California, San Diego, La Jolla, CA 92093-0448, USA. *Lin Zhang: zhanglin.materials@gmail.com

Dielectric materials with a high electric energy density and a low dielectric loss play a very important role in modern electrical power systems, e.g., hybrid electric vehicles, medical defibrillators, filters and switchedmode power supplies. Compared to batteries and supercapacitors, dielectric capacitors offer many unique advantages, such as fast charge and discharge characteristics, a high power density, a high output voltage, and a wide operating temperature range. Compared to other dielectric capacitors, polymer film capacitors are more attractive for the applications mentioned above due to their higher breakdown strength and a more graceful failure mechanism. In addition, the development of dielectric-polymer-based capacitors has been motivated by the increasing demand for compact, low-cost, light-weight and flexible energy storage devices, which makes polymer films the material of choice for the next generation of high pulse capacitors

A new family of poly(4-methyl-1-pentene) ionomers [PMP-(NH₃)_xA-y] (x = 1, 2, 3 and A = Cl⁻, SO₄²⁻, PO₄³⁻, y = NH₃ content) modified $(NH_3^+)_x A^{x-}$ ionic groups has been synthesized. The ionomers were synthesised using either a traditional Ziegler-Natta or a metallocene catalyst for the copolymerisation of 4-methyl-1-pentene and bis(trimethylsilyl)amino-1-hexene. A systematic study was conducted on the effect of the subsequent work-up procedures that can prevent undesirable side reactions during the synthesis of the [PMP-(NH₃)_xA-y] ionomers. The resulting PMP-based copolymers were carefully monitored by a combination of nuclear magnetic resonance (NMR), gel permeation chromatography (GPC), differential scanning calorimetry (DSC), mechanical property, dielectric properties, and electric displacement-electric field (*D-E*) hysteresis loop measurements. Our results reveal that the [PMP-(NH₃)_xA-y] ionomer films show a significantly enhanced dielectric constant (~5) and higher breakdown field (~ 612 MV/m) as compared with pure PMP films. Additionally, these PMP-based films show good frequency and temperature stabilities (up to 160 °C). A reliable energy storage capacity above 7 J/cm³ can be obtained, and is twice the energy storage capacity of state-of-the-art biaxially oriented polypropylene films, which can be attractive for technological applications for the energy storage devices.