

Developing A 3D Printable Electret Material For Sensing Applications.

O. A. Omoniyi, B. Tiller, R. O'Leary, J. F.C. Windmill

Centre for Ultrasonic Engineering, Electronic & Electrical Engineering Dept.,
Bioacoustics Group, University of Strathclyde
Glasgow, Scotland, United Kingdom
Oluwaseun.omoniyi@strath.ac.uk

I. SUMMARY AND MOTIVATION

The introduction of three-dimensional (3D) printing technology has opened up a world of rapid product development in different fields. 3D printing of piezoelectric materials has been growing in popularity in recent years showing increasing promise in the design of miniature sensors for acoustic devices, diagnostics devices, and energy harvesters. In this study, we demonstrate that voided polymer electret material can be printed into 3D structures using Stereolithography (SL) printing technology. A test sample was fabricated by printing a thin membrane of voided polymer electret sandwiched between two base structures. The polymer electret is formed by incorporating dry expanded Expancel microspheres into a photoactive polymer solution of polyethylene glycol diacrylate (PEGDA). The average d_{33} coefficient of the material were determined as 58pm/V. The results obtained showed the potential of using a voided polymer composite for 3D-printing functional electret based sensing devices.

II. ADVANCES OVER PREVIOUS WORKS

The piezoelectric properties of certain polymers have been studied in depth and much work has been done to exploit this advantage [1-2]. Polymers such as polytetrafluoroethylene (PTFE), polyvinylidene fluoride (PVDF), and Polypropylene (PP) have been shown to possess good piezoelectric properties and enhanced charge retention [3]. However, it has been shown that a new class of polymer electrets, the voided cellular polymer electrets, offers a number of advantages [6]. An electret is a dielectric material with quasi - permanent polarization. Voided polymer electrets are polymer materials consisting of internal voids, which are electrically charged. These materials have been reported to have higher piezoelectric constants due to their soft internal structure, when compared to other piezoelectric polymers, and offer better acoustic impedance matching in water and air coupled ultrasonic applications [4-5].

The possibility of fabricating miniature devices with enhanced operational capabilities and complex geometries with reduced cost using piezocomposites has been shown in [1].

In this study, we show that a sensing device can be fabricated through a combination of 3D printing with cellular polymer electret technology

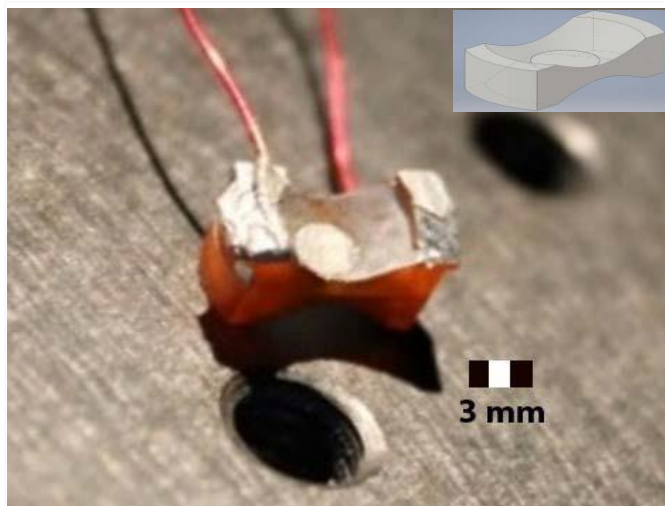


Fig. 1: Shows the 3D printed device (top) and its CAD model. The electret membrane is scaled to size.

III. MATERIALS AND METHODS

The materials used in this experiment include commercially available Expancel microspheres DE with particle size 30 μm (Akzo Noble pulp, Sundsvall Sweden), polyethylene glycol diacrylate (PEGDA), phenylbis (2,4,6-trimethylbenzoyl), phosphine oxide (Irgacure 819), and Sudan I (Sigma Aldrich).

A. Material Synthesis and 3D printing

The material synthesis was carried out in two steps. Two different polymer-based materials were synthesized to 3D-print the sample reported in this work. The first step consisted of mixing PEGDA with Irgacure 819 at 1wt%. Irgacure 819 is a photoinitiator, necessary to start the polymerization process of PEGDA when exposed to ultraviolet (UV) light. The mixture was stirred using a magnetic mixer for 24 hours to ensure a homogenous mix. After the mixing process, 0.1 wt% of SUDAN I was added into the mixture. SUDAN I is a UV blocker that allows an improvement of the resolution of the 3D-printed part. To ensure a homogenous composition, the mixture was put in a THINKY AER-250 mechanical mixer, mixing the composite for 3 minutes at 1500 rpm, and de-foaming it for 2 minutes at 1200 rpm. The second step

involved mixing Expancel microspheres (EP μ S), with PEGDA at a ratio of 1 to 15. Thus, 4 grams of EP μ S were added into 60 grams of PEGDA, and the mixture was mechanically mixed following the aforementioned process. After mechanical mixing a control PEGDA:EP μ S sample was made using a screen printing technique, with a layer thickness of 80 μ m, using an RK control coater. Every layer was cured under UV light for two minutes using an Intertronics IU250 Hand Lamp (Intertronics, Kidlington, England, UK).

An ASIGA PicoPlus27 3D printer (ASIGA, Anaheim Hills, California, USA) was used to print the device using the materials created in both steps above. Both of them were 3D-printed at a slice thickness resolution of 10 μ m, 5 sec exposure time and 100 sec burn in time. The support structure was 3D-printed first using the initial mixture of only PEGDA. By pausing the 3D printing process at the correct build layer, it was possible to switch the build fluid to print a membrane of PEGDA: EP μ S of 0.21 mm. By pausing the 3D printing process at the correct build layer, it was possible to switch again to the PEGDA fluid and print the second part of the support structure. The final membrane consisted of a disk of 0.21 mm thickness and a circle of 3 mm in diameter as shown in Fig. 1. After 3D-printing, Thin film coating of the sample was done using silver as the contact electrode. The membrane was poled using the corona discharge technique, which involved charging the sample under a pin electrode at a distance of 2 cm by applying a constant voltage of positive polarity at 12 kV. This process was carried out for 60mins.

IV. EXPERIMENTAL RESULTS

A. Structural Analysis

The microstructure of electret membrane was investigated using a table-top scanning electron microscope at an accelerating voltage of 15 kV (*SEM, Hitachi TM1000, Krefeld, Germany*). The microstructure of the electret was investigated to determine the degree of bonding between the voids and matrix and the dispersion of the voids since this greatly affects the poling result and thus piezoelectric and mechanical properties of composites [10].

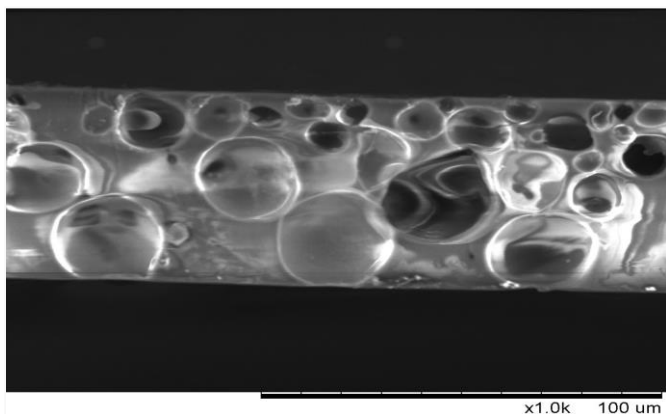


Fig. 2: Shows the cross-section of the device membrane. The internal voids are clearly visible and appears smooth and homogenous. Although voids sizes of 60 μ m were used, smaller sized voids are clearly seen. Spaces observed in the micrographs appear to be as a result of the mixing technique used.

B. Piezoelectric properties

The piezoelectric properties of the device were investigated using a microscanning laser Doppler vibrometer (Polytec MSA100-3D, Waldbronn, Germany) technique [11]. The voltage stimulus was generated by the MSA100-3D internal data acquisition board. The analysis of the device's velocity and displacement was determined with the laser vibrometer's control PC and data management system. A sinusoidal AC voltage of 3V_{pp} was applied across the axis of charging and the resulting displacement along the poling axis was measured. A frequency sweep stimulus was applied from 100 Hz to 25 kHz in order to obtain the characteristic frequencies at which the device will respond, as shown in Fig. 3. The average d_{33} coefficient of the material was determined as 58 pm/V.

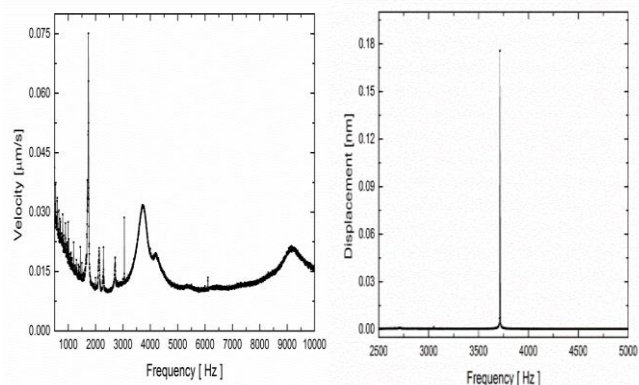


Fig. 3. Shows the response of the 3D printed membrane to a frequency sweep, and the magnitude of mechanical displacement (peak to peak) in response to a sinusoidal voltage of 3V_{pp} at a frequency of 3.7 kHz.

V. CONCLUSION

In conclusion this work demonstrates the possibility to 3D-print a cellular polymer electret. The 3D-printing process employed allowed the fabrication of a device with different material layers, and good resolution. The device consists of a 3D-printed voided polymer electret thin membrane sandwiched between a solid base of 3D-printed polymer. From the analysis obtained, it was shown that the voided polymer electret membrane is piezoelectric and can be employed in the fabrication of MEMS devices. Furthermore, the device provided a good response at a broader range of frequencies. This work also gives an insight into the creation of functional 3D-printed sensors through a cheaper and more time-efficient process.

Further investigation is still required to verify the volume of the voids and the charge distribution during poling as these are assumed to play a very important role in the electro-mechanical properties of the new electret material. Mechanical properties and Finite element modelling of the material would also be carried out to understand the physics of the material and its properties.

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