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Ameliorating and Tailoring The Morphological, Structural, and Dielectric Characteristics of SiO₂ /NiO Futuristic Nanocomposites Doped PVA-PEG for Nanoelectronic and Energy Storage Applications

Waleed Khalid Kadhim¹ · Majeed Ali Habeeb²

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Abstract

The current investigation inquiry involves silicon dioxide (SiO₂) and nickel oxide (NiO) nanoparticles to enhance the structural and dielectric properties of a polyvinyl alcohol (PVA) with polyethylene glycol (PEG) combination for use in flexible pressure sensors and nanoelectrical devices. Solution casting was used to fabricate PVA-PEG-SiO₂/NiO nanocomposites at various weight percentages of (SiO₂/NiO) N.Ps (0, 2, 4, 6 and 8) wt%. The structural properties of PVA-PEG-SiO₂/NiO nanocomposites were studied by X-ray diffraction (XRD), and the amorphous state of the mixture consisting of polyvinyl alcohol (PVA) and polyethylene glycol (PEG) was revealed. Furthermore, the characteristic peak of the original polymers was much smaller at higher doping concentrations. According to field emission scanning electron microscopy (FE-SEM), when the weight percentage approaches 8%, the top surface of the (PVA-PEG-SiO₂/NiO) N.Cs films exhibits homogenous and cohesive clumps or fragments dispersed randomly. Optical microscopy made it possible to observe that nanoparticles (SiO₂/NiO) generate an integrated network inside the matrix of polymers, unlike the pure film of (PVA-PEG). The electrical properties of alternating current illustrate that as the frequency of the applied electrical field increases, the dielectric constant and dielectric loss of nanocomposites decline. Also, on the contrary, these values increase in conjunction with the increase in the concentration of nanoparticles, and the highest value is at a frequency of 100 Hz at a concentration of 8%. The (PVA-PEG) blend's dielectric constant and A.C. electrical conductivity were improved by almost 300% and 112%, respectively, at the highest addition rate (8 wt.%). The findings obtained revealed that the structural and AC electrical conductivity were enhanced by doping (PVA-PEG) with (SiO₂/NiO) NPs. Findings indicated that the (PVA-PEG-SiO₂/NiO) nanostructures would be excellent materials for a range of nanoelectronics industries. The results obtained showed an increase in parallel capacity. It reached 400 pf with an increase in applied pressure, as well as an increase in sensitivity to pressure of about 77.2 with the biggest percentage of weight addition of nanoparticles.

Keywords Structural properties \cdot Dielectric properties \cdot Pressure sensor \cdot SiO₂/NiO nanoparticles \cdot Sensitivity

1 Introduction

The quantity of plastic garbage that is everywhere these days is getting to the point that it is polluting the entire planet. There are an infinite amount of applications for

 Majeed Ali Habeeb pure.majeed.ali@uobabylon.edu.iq
Waleed Khalid Kadhim welidkalid37@gmail.com

- ¹ Education Babylon, Ministry of Education, Hillah, Iraq
- ² Department of Physics, College of Education for Pure Sciences, University of Babylon, Hillah, Iraq

polymers, thus finding ecologically and human-friendly polymers is imperative [1, 2]. One of the various methods available today for producing polymeric materials is mixing polymers. This procedure combines at least two different kinds of polymers to produce a novel sort of polymeric matrix [3, 4]. Two methods to increase the AC conductivity of polymeric materials are plasticizers and polymer mixing. It is easy to produce polymer blend complexes from PVA and PEG because of their tunable chemical and physical characteristics. Toughness, nontoxic, biodegradable, and heat stability are some of these characteristics [5, 6]. Over the past many years, polymer nanocomposites have garnered significant interest from both academia and industry. They are now an essential part of creating new, cutting-edge materials for a variety of uses, notably electrical engineering [7, 8]. There are several options within the field of nanotechnology to evolve nanomaterials with better qualities; this will allow to be utilized across a range of sectors. The general definition of nanotechnology is the manipulation of materials at the atomic or molecular level through procedures including consolidation, deformation, and separation [9, 10]. Our way of life may be enhanced by nanotechnology in many ways, including quicker electronics, massive memory, and more affordable energy through more efficient transformation of energy, and heightened safety with the creation of nanoscale [11, 12]. Despite all technological specialties, nanotechnology is one of the newest favored fields for study and advancement. With the use of nanotechnology, innovative materials will be created that may be used to construct and create new structures and properties with greater functioning, decreased upkeep expenses, and increased performance [13, 14]. Given that they are considered to be essential technologies for the 21st-century, a large sum of research has been done in this area. The particular chemical, biological, and physical features of nanoscale structures in relation to their macroscopic counterparts may soon lead to new uses [15, 16]. An innovative method for producing materials made from polymer is to profit from polymer blends, which merge two or more different polymers to create composite materials. One of the blend's most notable qualities is its adaptability to many uses [17, 18]. Due to their remarkable effectiveness and variety of features, which are achieved by adding a small plenty of nanoparticles to the polymer mixture / polymer nanocomposites have generated a lot of curiosity. When nanoparticles and polymer are combined, nanocomposites perform noticeably better at far lower loadings than when polymer is used alone [19, 20]. Polymer electrolytes are used in sensors, electrochemical devices, and rechargeable batteries; they have attracted a lot of interest lately. A number of beneficial characteristics of biopolymer materials include their affordability, non-toxicity, natural availability, biodegradability, and ecological advantages [21, 22]. Polyvinyl alcohol, or PVA, is widely used in many important uses and fields and is recognized as one of the most important polymers. Batteries, electro chromic, detectors, and the medical sectors are a few instances of these industries and uses. PVA has superior durability, resistance to corrosion, and high thermal properties compared to other kinds of polymers [23, 24]. Because of its exceptional electrical characteristics, high dielectric strength, and remarkable storage capacity, PVA shows great promise as a material. This adaptable material has several important uses; including medicine systems for distribution, packing, and polymer reuse [25].

Polyethylene glycol (PEG) is an additional artificial polymer with a broad molecule weight range that is solvent in water. Blending it with stiff polymers to create novel materials with specific qualities is a suitable option because of its elastic chain [26]. Due to its special physicochemical characteristics, polyethylene glycol (PEG) is a polyether compound that finds use in a wide range of scientific fields and enterprises. PEG is widely used in biomedicine, pharmacology, and other sectors because of its chemical solubility in water and organic solvents, as well as its non-toxic nature [27]. Nanoparticles of silicon dioxide (SiO₂) are utilized as reinforcements in electronic packaging and thermoplastic polymers. Known for its substantial particular area of surface, strong thermal properties, and developed mechanical properties, silica is a white powder that may be found in nature. SiO_2 is environmentally friendly, amorphous, and may be molded into an optoelectronic nanocomposite material by filling it with polymers that have nanopores [28]. The SiO_2 particles serve as a solid plasticizer, enhancing the mechanical, chemical, and dimensional integrity of the composite polymer. in the field of materials science, SiO₂ NPs are employed as fillers or reinforcements in composite materials for bettering mechanical properties, such as strength and toughness, their small size and huge surface area enable effective dispersion within matrices, leading to boosted material performance [29]. Amidst the plethora of available nanomaterials, nickel oxide (NiO) nanoparticles have garnered significant attention mainly because of their remarkable stability and excellent magnetic, electrical, optical, and catalytic properties. These properties also account for their many applications, including gas detectors, electro chromic components, fuel cells, batteries, solar cells sensitive to dyes, solar energy absorbers, and magnetic cameras [30]. In addition, compared to other metal oxide nanoparticles, NiO nanoparticles are incredibly affordable, environmentally friendly, and highly fixed conductive materials. Among the medical applications for these nanoparticles were biological identification, imaging, drug delivery, and antibacterial. Apart from their previously mentioned applications, (NiO) NPs have the ability to effectively remove both organic and inorganic pollutants; hence, they are crucial for maintaining the integrity of the environment [31]. Furthermore, the drugs, catalysis, and dye manufacturing sectors are more possible uses for silica nanoparticles. Likewise, silica may be used to reinforce polymer combinations in thermoplastic polymers, volatile flavorings, compound polymer gel solutions, coatings for preventing corrosion, and plastic containers [32]. The most common materials employed by pressure sensors contain silicon, sputtered thin films, polysilicon thin films, bonded metal foils, and inkjet-printed films. Piezoresistive pressure sensors are the most widely utilized technique within the **Fig. 1** Depicts the XRD patterns of (PVA-PEG -SiO₂-NiO) NCs with varying amounts of (SiO₂-NiO) NPs where (**A**) represents the SiO₂ phase and (**B**) represents the NiO phase

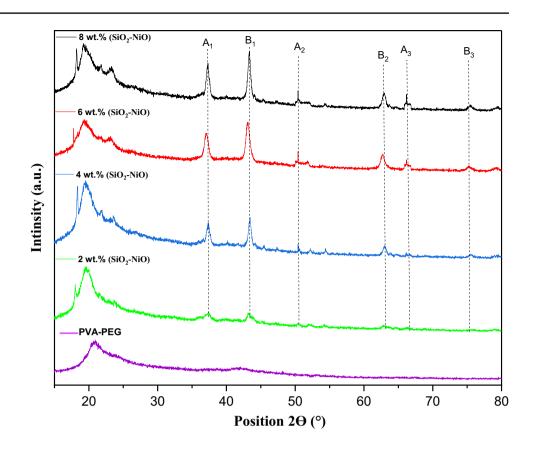


Table 1 The mean crystallite size for "SiO2/NiO" nanoparticlesdetermined from patterns of XRD utilizing the Scherer equation

Material	2 0 (°)	hkl	FWHM (rad)	Crystallite size (nm)
SiO ₂				
A_1	37.2463	110	0.3023	28.98
A_2	50.4078	210	0.501	18.31
A ₃	66.2561	310	0.5821	17.03
NiO				
B_1	43.3726	200	0.285	31.34
B_2	62.872	220	0.5334	18.24
B ₃	75.4065	311	0.7246	14.48

pressure sensor industry because of its high sensitivity and inexpensive cost [7]. Based on whether behavior is more suited in a particular case, the piezoresistive or pseudocapacitive behavior of the nanocomposite pressure sensors drives their operation. Important performance parameters like pressure sensitivity, pressure range, ability to detect small pressures, and endurance of larger forces set apart various types of pressure sensors [33]. This study addresses the manufacturing of (PVA-PEG-SiO₂-NiO) nanocomposites (NCs) for optoelectronics applications, with an aim of offering a low-cost and straightforward process.

2 Experimental Part

The casting approach was utilized to create nanocomposites from polyvinyl alcohol (PVA)/Polyethylene glycol (PEG) and silicon dioxide (SiO₂), nickel oxide (NiO) NPs. It involved creating a more homogenous solution by dissolving pure PVA and PEG in 45 ml of water distillation for 50 min while stirring with a magnetic stirrer at seventy degrees Celsius. Silicon dioxide (SiO₂) and nickel oxide (NiO) NPs were included to the polymer mixture at weight percentages of 2%; 4%; 6%; and 8%, respectively. After casting it, it was left to dry for 6 days at room temperature and the polymer nanocomposites were formed. Square pieces of (PVA-PEG-SiO₂/NiO) nanocomposites were taken for different measurements. Using field emission scanning electron microscopy (fesem), (model/ InspectTM F50—DETAILS\1.2 nm at 10 kV; pressure 3.11e-3 par country/ Holland), was used to examine the structural properties of nanocomposites. Olympus offers the Optical Microscope (OM) (Top View Model Number: Nikon-73346) with a camera for microscopic images with a magnification of 20x, in alongside employing a diffract meter with a radiation wavelength of 1.5418 Å to measure X-ray diffraction (XRD). Utilizing an LCR meter (HIOKI; 3532-50; LCR HI TESTER), the dielectric properties of nanocomposites were examined between

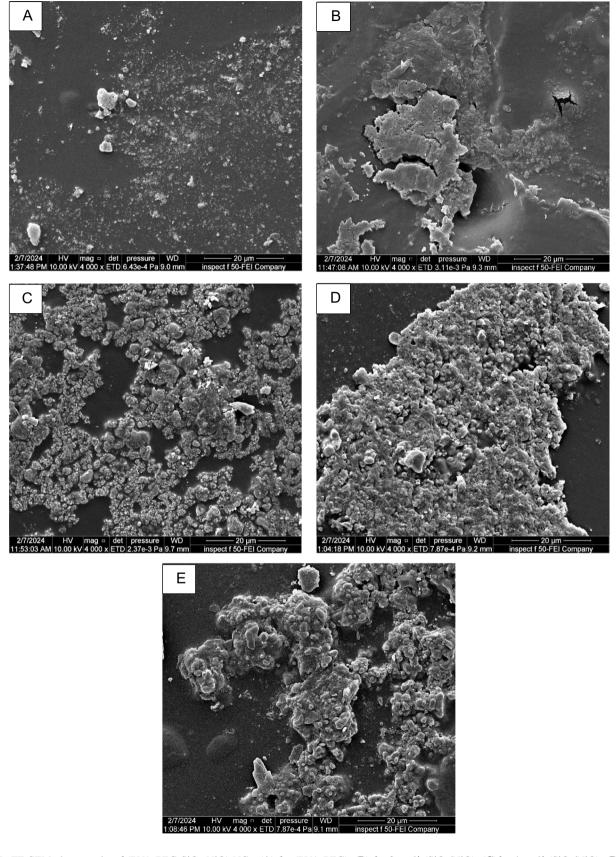


Fig. 2 FE-SEM photographs of (PVA-PEG-SiO₂-NiO) NCs: (**A**) for (PVA-PEG), (**B**) for 2 wt.% (SiO₂/NiO), (**C**) for 4 wt.% (SiO₂/NiO), (**D**) for 6 wt% (SiO₂/NiO), (**E**) for 8wt% (SiO₂/NiO)

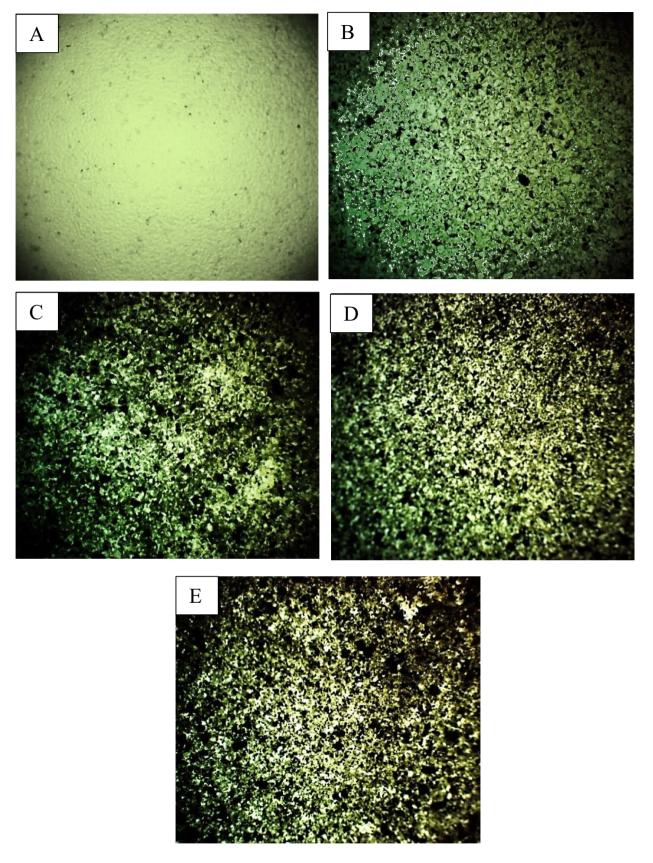


Fig. 3 Optical microscope photographs of (PVA-PEG-SiO₂-NiO) NCs at a magnification (10x):- **A** (PVA-PEG); **B** 2 wt.% (SiO₂/NiO); **C** 4 wt.% (SiO₂/NiO); **D** 6 wt.% (SiO₂/NiO); **E** 8 wt.% (SiO₂/NiO)

100 Hz and 5 MHz. By means of LCR meter to calculate the parallel capacitance between both poles above and below the test piece over a range of pressures (80–160 bar), the effectiveness of the pressure sensor of (N.Cs) was determined.

The dielectric constant (ϵ') is calculated from the following function [34]:

$$\epsilon' = C_P / C_0 \tag{1}$$

Capacitance is indicated by C_p , while a vacuum capacitor is indicated by (C_o) . We find the dielectric loss (ϵ ") using [35]:

$$\epsilon'' = \epsilon' \, \mathrm{D} \tag{2}$$

(D) is displacement. Here is the formula for calculating AC electrical conductivity [36]:

$$\sigma_{\rm a.c} = \omega \epsilon'' \epsilon_0 \tag{3}$$

where (ω) is the angular frequency. The following is a definition of pressure sensor sensitivity [37, 38]:

$$S_{AP}(\%) = (1 - C_b/C_n) \times 100\%$$
 (4)

When blends and nanocomposites' capacitances are represented by the symbols C_b and C_n , respectively.

3 Results and Discussions

3.1 XRD Analysis

The XRD of the PVA-PEG mixture and nanocomposites is displayed in Fig. 1. The structure of PVA-PEG blends and nanocomposites with different concentrations of nanoparticles (SiO₂/NiO) was investigated using X-ray diffraction (XRD). The nanocomposites and (PVA-PEG)

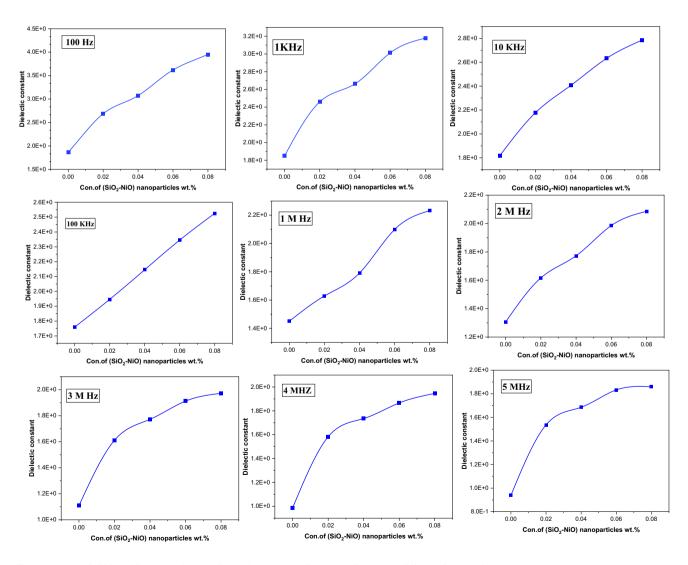


Fig. 4 Impact of (SiO₂/NiO) NPs ration on dielectric constant of PVA-PEG blend at different frequencies

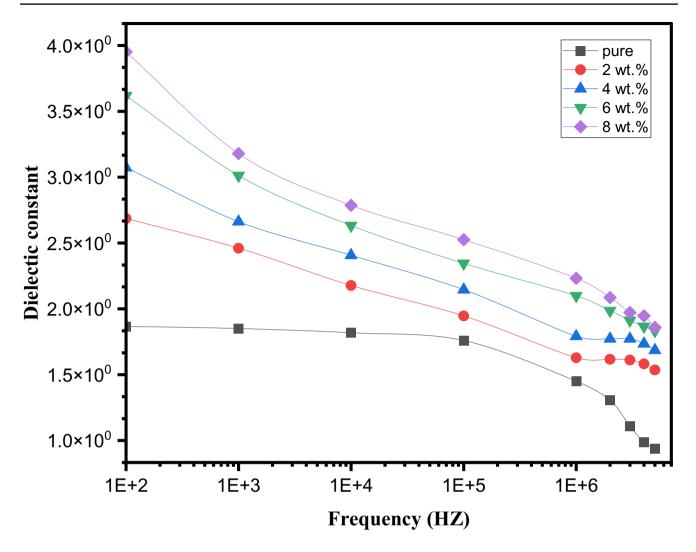


Fig. 5 Change of ϵ' with a frequency of (PVA-PEG-SiO₂/NiO) NCs

blend display a broad band located at $2\theta = 20.26^{\circ}$, indicating the amorphous nature of PVA and PEG, according to XRD analysis [7]. The primary mechanism responsible for the preservation of PVA and PEG amorphous nature is between molecules hydrogen bonding, different monomer structures and the molecules inside each component form this link [39]. A reduction in the polymer blend's degree of amorphous form was seen upon insertion of (SiO₂/NiO) nanoparticles. The results confirm that there is variation in the electrostatic forces that occur between (SiO₂/NiO) NPs and the polymer blend. As the (SiO₂-NiO) concentration in the polymer nanocomposite samples varies as a result of this change, the polymer blend's structures vary as well, eventually raising the degree of crystallization of the nanocomposites. Ion mobility is accelerated and conductivity is raised in the absence of a well-defined structure. The crystalline state of nanoparticles (SiO₂/NiO) is responsible for the steep peaks seen at $(2\theta = 37^{\circ}, 43^{\circ}, 63^{\circ})$ and 75°) which are linked to the increasing concentration of nanoparticles (SiO₂/NiO) in the nanocomposites. The findings demonstrate that the specimen's structural features may be altered by the addition of (SiO₂/NiO) nanoparticles [40, 41], as indicated in Table 1.

3.2 Field Emission Scanning Electron Microscopy (FE-SEM) Measurements of (PVA-PEG-SiO₂/NiO) NCs

Representative (FE-SEM) photographs of (PVA-PEG-SiO₂/NiO) films with and without varying SiO₂ and NiO nanoparticle concentrations are shown in Fig. 1, which provides a good idea of their size and shape. It is discovered that the polymer picture (A) is more coherence and homogeneous. The surface morphology of (PVA-PEG-SiO₂/NiO) nanocomposites alters noticeably when (SiO₂/NiO) nanoparticles are introduced (see figures B; C; D;

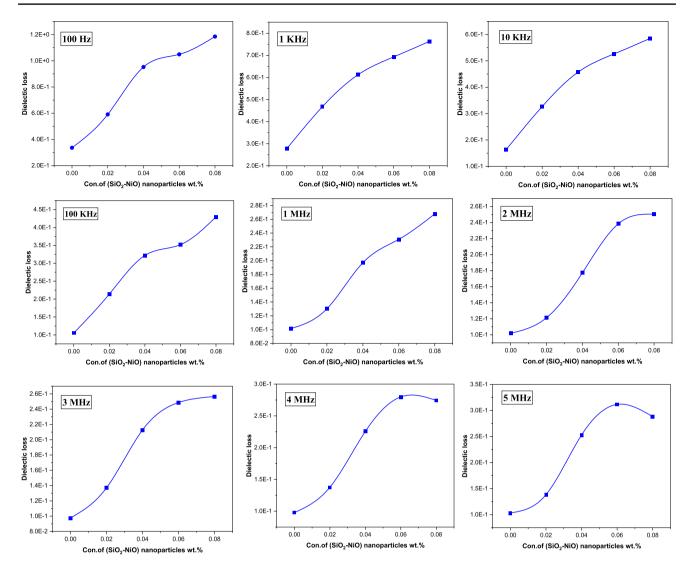


Fig. 6 Impact of (SiO₂/NiO) NPs ration on dielectric loss of PVA-PEG blend at different frequencies

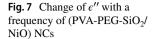
and E). The photographs also demonstrate that the granules particles' (SiO_2/NiO) nanoparticle concentration is increasing. The upper surface of (PVA-PEG-SiO_2/NiO) nanocomposites has a lot of clumps or particles dispersed on the uppermost surface. Nanoparticles in (PVA-PEG-SiO_2/NiO) nanocomposite films are uniformly dispersed and have a tendency to cluster; the (FE-SEM) images also show the formation of paths and a connected network within the nanocomposites that allow the passage of charge carriers [42, 43] (Fig. 2).

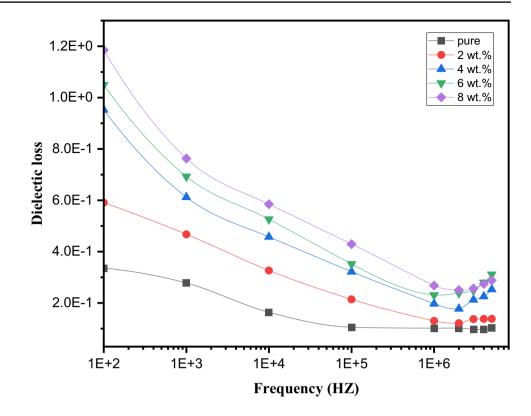
By comparing the results of XRD and SEM, it was clear from the X-ray examination that PVA and PEG are in amorphous form and the degree of amorphous decreases with the increase of nanoparticles in the polymer blend. Also, the crystalline state of the nanoparticles (SiO₂/NiO) is responsible for the appearance of sharp peaks. As a result, the structural properties of the

models change. As for the SEM examination, it showed the presence of clusters and aggregates on the surface of the nanocomposites when adding the filler, they tend to aggregate and show paths for the charge carriers to move through them, unlike the image of the more cohesive polymer blend.

3.3 The Optical Microscope of (PVA-PEG-SiO₂/NiO) N.Cs

The microscopic image of PVA-PEG-SiO₂/NiO nanocomposites (at magnification $10 \times$) is displayed in Fig. 3. A distinct variance among the samples can be seen in microscopic pictures as the amounts of (SiO₂/NiO) nanoparticles climb. As can be seen in the photos (A-B-C-D and E), the lack of any discernible holes or fractures, demonstrates the efficient interfacial contact between the





blending matrix and the (SiO₂/NiO) nanoparticle composition. Nanocomposites have noticeable form uniqueness. At a percentage of 8% by weight for (SiO₂/NiO) nanoparticles, an ongoing network of nanoparticles forms. When multiple spherical particle pieces or clusters create on the surface of nanocomposites, it means that a homogeneous growth process is present. During which the welldispersed and aggregating nanoparticles in the PVA-PEG blend establish an ongoing network inside the polymers. As the quantities of (SiO₂/NiO) nanoparticles increase, microscopy results clearly demonstrate the distinction among the samples, as seen in the photos (A-B-C-D and E). Hence, charge carriers are permitted to pass through specific channels in this network, which lowers the composite material's resistance (PVA-PEG) [44, 45].

3.4 The A.C electrical Properties of (PVA-PEG-SiO₂/ NiO) NCs

At room temperature, Fig. 4 illustrates the alteration in the real dielectric constant with nanoparticles concentration at frequencies ranging from 100 Hz to 5 MHz. The dielectric constant of the nanocomposites (PVA-PEG-SiO₂/NiO) is high at low frequencies, as the Fig. 5 shows, since the electric dipoles have enough relaxation time to be maintained by the field. Consequently, the dielectric constant is large. Nevertheless, by increasing the frequency, the dipoles are not given enough

time to be maintained by the field, resulting in a decrease in relaxation time and dielectric constant, as seen in Fig. 5.

We notice that the behavior of the dielectric constant increases with increasing proportions of (SiO_2/NiO) nanoparticles for all frequencies, as is clear in Fig. 4 [46, 47].

There is a group of polarizations that occur in nanocomposites, including: directional polarization, which results from the effect of the applied electric field that makes the dipoles align with their direction, and ionic polarization, which occurs due to an exterritorial electric field that affects the ions and is opposite, and these polarizations determine the behavior of the dielectric constant by altering added nanoparticles and their concentration [48, 49].

At ambient temperature, Fig. 6 illustrates the alteration in the dielectric loss with nanoparticle concentration at frequencies ranging from 100 Hz to 5 MHz. The graph displays that when the amount of nanoparticles increases, dielectric loss rises as well. Dielectric loss depends on the electrical conductivity, as well as the distance and internal polarization factors of the (PVA-PEG-SiO₂/NiO) nanocomposites. Also the nanocomposite forms an ongoing network of particles when the concentration of these nanoparticles approaches 8 weight percent. Then the dielectric loss becomes the highest possible [50, 51]. Polarization causes a change in dielectric loss, including: interfacial polarization resulting from the non-homogeneity of materials, and dipole polarization. The fluctuation of (PVA-PEG-SiO₂/NiO) nanocomposites dielectric loss with frequency at room temperature is demonstrated

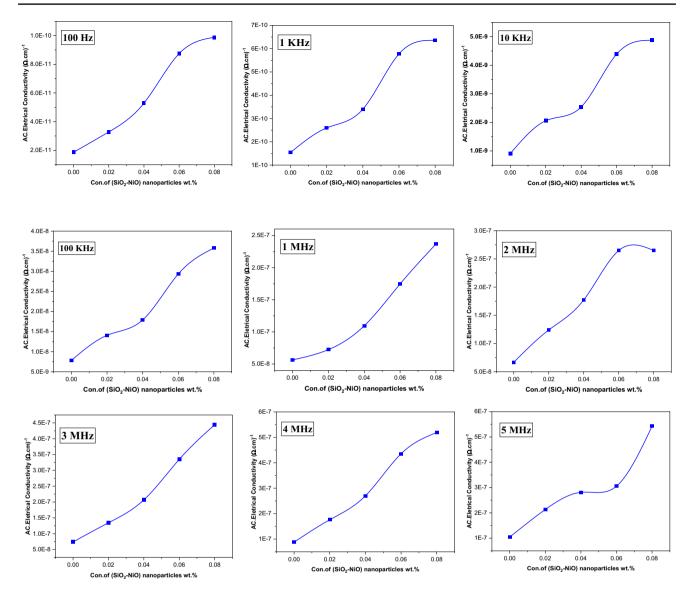


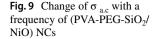
Fig. 8 Impact of the percentage of (SiO₂-NiO) NPs on (σ_{ac}) of the PVA/PEG blend at different frequencies

in Fig. 7. The graph displays that dielectric loss lowers with increasing frequency. This phenomenon can be explained by a failure of dipoles made of composite molecules to rotate in parallel, which causes an interval between the dipole's frequency and the electric field's frequency. In other words, dipoles cannot keep up with the frequency of the applied electric field. As a result, neither charge accumulation nor ion dispersion occurs [52, 53].

Figure 8 illustrates the alteration in the A.C electrical conductivity with nanoparticle concentration at frequencies ranging from 1×10^2 Hz to 5×10^6 Hz. SiO₂/NiO nanoparticle concentrations rise are accompanied by a growth in conductivity, as indicated in Fig. 8. A rise in electric charge results from to configure of saturated

nanoparticles. The findings may be explained by the effect of space charge brought about by the increase in charge carriers and their even distribution throughout the polymer matrix [54, 55].

The A.C. electrical conductivity efficiency of the (PVA-PEG-SiO₂/NiO) NCs as an indicator of frequency is depicted in Fig. 9. The electrical conductivity of (PVA-PEG-SiO₂/NiO) nanocomposite samples is found to increase with a rise in the applied electric field frequency. The polarization of free charge, which happens at extremely low frequencies in combination with the jumping mobility of charge carriers, is responsible for the aforementioned phenomena [56, 57]. Table 2 shows the sample values of (ϵ' , ϵ'' , and σ a.c.) for (PVA-PEG-SiO₂/NiO) NCs at 1kHz.



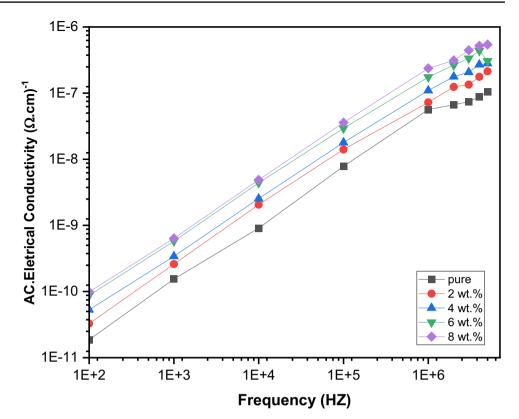


Table 2 Values of dielectric [constant—loss] and A.C. electrical conductivity ($\sigma_{a,c}$) for (PVA-PEG-SiO₂/NiO) N.Cs at 10³ Hz

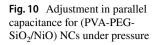
Con. Of SiO ₂ /NiO (wt.%)	ε'	ε"	$\sigma_{a.c}$ (S/ cm)
0	1.84992	1.866	1.54222E-10
2	2.46034	2.685	2.59807E-10
4	2.66204	3.346	3.40286E-10
6	3.01212	3.945	5.77556E-10
8	3.17872	4.518	6.36E-10

3.5 Utilizing (PVA-PEG-SiO₂-NiO) NCs for Pressure Sensor Applications

The parallel capacitance fluctuation in (PVA-PEG-SiO₂/NiO) nanocomposites at various proportions of weight of (SiO₂/NiO) nanoparticles during applied pressure is displayed in Fig. 10. There is a direct correlation between the value of capacitance and the load pressure's scale, as pressure rises, the capacitance increases because of the crystal's several interconnecting domains that contain both positive and negative charges, as the graph depicts [58, 59]. These symmetric regions inside the crystal

structure lead to the absence of any net charge in the crystal. This symmetry is disrupted when the crystal is under stress. To bring symmetry back, these domains reposition, generating a current and raising the capacitance in this step [60].

Figure 11 illustrates the effect of (SiO₂/NiO) nanoparticles on the electrical capacitance (Cp) for (PVA-PEG-SiO₂/NiO) NCs at 80 bars. This graph makes it evident that when the concentration of (SiO₂/NiO) nanoparticles rises, so does the electrical capacitance of nanocomposites. This might be caused by the increase in charge carriers density in nanocomposites [61]. The effectiveness of the nanostructures (PVA-PEG-SiO₂/NiO) is strongly influenced by the concentration of (SiO₂/NiO) nanoparticles, establishes the pressure sensing range, leading to a range of applications that call for greater forces. The impact of (SiO₂/NiO) nanoparticles on (PVA-PEG-SiO₂/NiO) NCs sensitivity is seen in Fig. 12 and Table 3. This graph makes it evident that when the concentration of (SiO₂/NiO) nanoparticles climbs, so does the sensitivity of nanocomposites. This is due to an internal dipole moment [62].



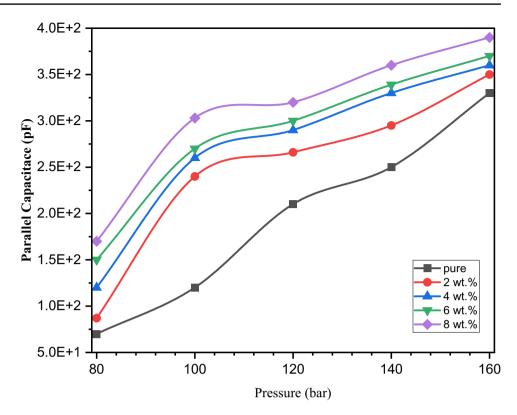
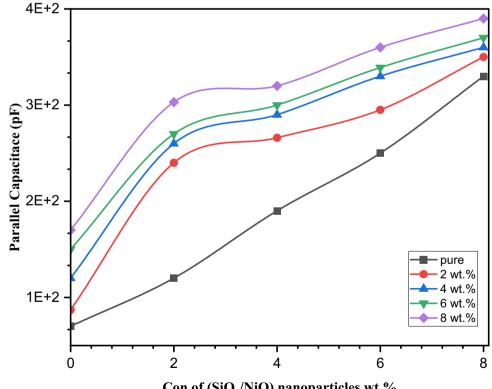
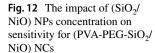


Fig. 11 The impact of concentration of (SiO₂/NiO) NPs on parallel capacitance for (PVA-PEG-SiO₂/NiO) NCs at 80 bar



Con.of (SiO₂/NiO) nanoparticles wt.%



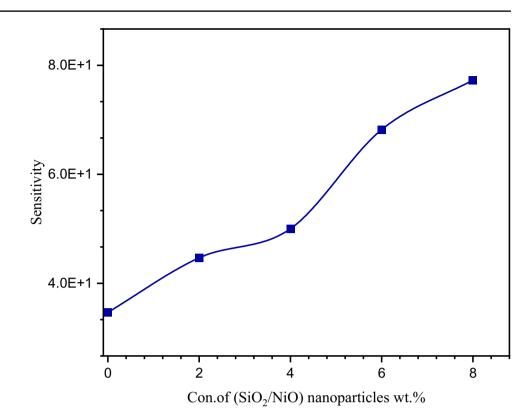


Table 3 Sensitivity values with concentrations of (SiO $_2$ /NiO) NPs for (PVA-PEG-SiO $_2$ /NiO) NCs

Con.of (SiO ₂ -NiO) NPs (wt. %)	Sensitivity (%)
0	34.65347
2	44.66667
4	50
6	68.18182
8	77.27273

4 Conclusions

In this work, (PVA-PEG-SiO₂/NiO) NCs were created by the solution casting approach. X-ray diffraction (XRD) revealed that the structure of the PVA/PEG mixture is amorphous, mainly due to the hydrogen bond between the molecules. Sharp peaks appear when nanoparticles are added, due to the crystalline state of the nanoparticles and the change in the structural properties of the samples. FE-SEM has been used to examine the surface morphology of the films formed from (PVA-PEG-SiO₂/NiO) NCs. The findings show that a wide variety of aggregates or pieces are present and are dispersed randomly over the top surface. The combination of silicon dioxide (SiO_2) and nickel oxide (NiO) nanoparticles at a concentration of 8% weight percent leads to the creation of an integrated network inside the polymer mix, according to optical microscope (O.M.) photographs. The dielectric

properties of (PVA-PEG-SiO₂/NiO) nanocomposites were investigated, and the outcomes showed that a rise in the concentration of (SiO₂/NiO) NPs was associated with an increase in the dielectric constant, dielectric loss, and A.C. electrical conductivity of the (PVA-PEG) blend to (3.17872, 4.518, and 6.36E-10 S/ cm)respectively. Even though the A.C. electrical conductivity rises, the dielectric constant and dielectric loss decline in proportion to an increase in the frequency of the electric signal. The (PVA-PEG-SiO₂/NiO) nanostructures are well-suited for use in a variety of nanoelectronics technologies that value flexibility, cost-effectiveness, high energy storage, as well as low loss due to their advantageous structure and dielectric characteristics. The (PVA-PEG-SiO₂/NiO) nanocomposites are lightweight, malleable, and very efficient, but they also show great pressure sensitivity. The highest reported sensitivity of nanocomposites is 77.27% at 8 w.t % concentration.

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Author Contributions All authors contributed to the study's conception and design. Material preparation, data collection and analysis were performed by Majeed Ali Habeeb, Waleed Khalid Kadhim. The first draft of the manuscript was written by Majeed Ali Habeeb and all authors commented on previous versions of the manuscript. All authors read and approved the final manuscript.

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Data Availability No datasets were generated or analysed during the current study.

Declarations

Ethics Approval The research is not involving the studies on human or their data.

Consent to Participate Consent.

Consent for Publication Consent.

Competing Interests The authors declare no competing interests.

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