RSC Advances



PAPER

View Article Online
View Journal | View Issue



Cite this: RSC Adv., 2025, 15, 15218

High β -phase PVDF composite thin films filled with metal (M = Ni, Ag, Co) phosphate-based particles: advanced materials for energy harvesting applications†

Chaymae Bahloul,^a Adil Eddiai,^b Omar Cherkaoui,^c M.'hammed Mazraoui,^b Fatima-Zahra Semlali (10) ** and Mounir El Achaby (10) ** a

This study demonstrates a systematic approach to developing high-performance poly(vinylidene fluoride) (PVDF) nanocomposites through the strategic incorporation of metal phosphate nanostructures synthesized via distinct methodologies. Ni-P particles were synthesized using hydrothermal processing and Aq-P via a precipitation technique, while Co-P and Co-Pn were prepared through co-precipitation and solvothermal routes, respectively. These diverse synthetic approaches yielded particles with controlled crystallinity, morphology, and surface properties, as verified through comprehensive XRD, FTIR spectroscopy, Raman and SEM. The integration of these nanostructures into the PVDF matrix significantly promoted electroactive β-phase formation, with improvements ranging from 68% to 96%, with optimal transformation typically achieved at 3 wt% loading. Mechanical characterization revealed remarkable property enhancements, with PVDF/3Co-Pn exhibiting an unprecedented 181.83% increase in tensile strength and a 184.76% improvement in Young's modulus. Thermogravimetric analysis demonstrated substantial thermal stability enhancement, with PVDF/7Ni-P showing a 24.92 °C increase in onset degradation temperature. Ferroelectric measurements indicated that PVDF/7Co-P composites achieved superior remnant polarization with a 200% increase, while PVDF/3Co-P demonstrated optimal maximum polarization with a 75% improvement. Importantly, this research establishes clear correlations between synthesis methodology, resultant nanoparticle characteristics, and composite performance, providing critical insights into structure-property relationships in metal phosphate-PVDF systems. These findings advance the fundamental understanding of interface engineering in polymer nanocomposites and establish design principles for developing advanced materials for flexible electronics, energy storage devices, and sensing technologies.

Received 5th March 2025 Accepted 27th April 2025

DOI: 10.1039/d5ra01605c

rsc.li/rsc-advances

Introduction

Two of the most pressing challenges facing humanity are energy scarcity and environmental pollution, both undermining sustainable development.¹ These challenges necessitate the development of environmentally friendly, affordable, and sustainable energy technologies.^{2,3} Energy harvesting (EH) has emerged as a promising approach, converting ambient energy

into electricity from sources such as light, radio frequency radiation, heat, and mechanical vibrations.^{4,5}

Among these diverse energy sources, biomechanical energy presents significant untapped potential due to its ubiquitous availability. The human body dissipates over 100 watts of energy daily, while wearable devices typically require less than 1 watt, a gap that continues to widen with technological advancement.⁶ Harnessing even a fraction of this biomechanical energy could power personal electronic devices, offering an eco-friendly alternative to traditional batteries.^{6,7} To effectively capture this biomechanical energy, materials with specific conversion capabilities are essential.⁸

Piezoelectric materials are well-suited for biomechanical energy harvesting due to their intrinsic capability to convert mechanical energy into electrical energy. These materials mainly include single crystals, ceramics, polymers, and polymer composites. Among them, polymers and their composites have demonstrated significant potential for energy harvesting

^aMaterials Science, Energy and Nanoenineering Department (MSN), Mohammed VI Polytechnic University (UM6P), Lot 660 – Hay Moulay Rachid, 43150 Ben Guerir, Morocco. E-mail: mounir.elachaby@um6p.ma; fatimazahra.semlali@um6p.ma; Tel: +212662010620; +212661100919

^bCondensed Matter Physics Laboratory, Ben M'sick Faculty of Sciences, Hassan II University of Casablanca, Morocco

REMTEX Laboratory, Higher School of Textile and Clothing Industries (ESITH), Casablanca, Morocco

[†] Electronic supplementary information (ESI) available. See DOI: https://doi.org/10.1039/d5ra01605c

Paper

applications, particularly where flexibility is essential. Although

their electrical performance is typically lower than that of ceramics, their mechanical flexibility makes them highly attractive for wearable and flexible energy harvesting devices.10,11

Within the category of piezoelectric polymers, poly(vinylidene fluoride) (PVDF) stands out as the most extensively researched thermoplastic due to its affordability, processability, and exceptional functional characteristics.12 PVDF exists in five crystalline phases $(\alpha, \beta, \gamma, \delta, \text{ and } \varepsilon)$, with the β -phase being most critical for piezoelectric applications due to its polar structure. 13 Various approaches have been developed to enhance the crucial β-phase content in PVDF, including solvent casting, copolymerization, and notably, the incorporation of fillers into the polymer matrix. 12,14,15 The selection of appropriate fillers is therefore a key consideration for optimizing PVDF's piezoelectric performance.

A wide range of fillers have been explored to enhance the electroactive performance of PVDF, including ceramic nanoparticles such as BaTiO₃, PZT, ZnO, and carbon-based materials like graphene and carbon nanotubes.16-19 These fillers are known to promote β-phase nucleation and improve dielectric properties by inducing interfacial polarization and acting as nucleating agents. However, many of these materials face key limitations, including brittleness, environmental toxicity is particularly for lead-containing ceramics, and challenges in achieving uniform dispersion within the polymer matrix. Ferrite-based fillers have also shown promise, 20-22 but they often suffer from poor compatibility with PVDF, requiring additional surface modification steps to improve their dispersion and interfacial bonding. These drawbacks have driven the search for alternative fillers that are not only functional and effective but also environmentally benign, structurally tunable, and easy to process. In this context, metal phosphate-based particles (MP-Ps) offer a compelling solution, combining chemical stability, morphological versatility, and potential for strong interaction with the PVDF chains to form next-generation, sustainable piezoelectric composites.

Recent advances in materials science have highlighted MP-Ps as particularly promising fillers due to their natural abundance, environmental friendliness, low cost, chemical stability, and structural versatility.23,24 These compounds can be synthesized in various morphologies ranging from one-dimensional structures to complex three-dimensional architectures, conferring unique properties including high chemical stability, mechanical flexibility, and large surface areas.25 Their synthesis methods, including ball milling, chemical precipitation, and hydrothermal treatments, significantly influence morphology, surface properties, and reactivity,26,27 thereby offering considerable potential for tailoring their interaction with polymer matrices like PVDF.

This potential synergy between MP-Ps and PVDF matrices represents an emerging research direction with significant implications for energy applications. To date, only a few studies have explored the potential of MP-Ps based fillers to enhance the functional properties of PVDF. For instance, Biswas et al.28 developed Ca₃(PO₄)₂ nanorod-incorporated PVDF films with

enhanced piezoelectricity, while other researchers have investigated PVDF composites with α-zirconium phosphate29 and NASICON-type LiSnZr(PO₄)₃ ceramic³⁰ for specific applications. However, despite these promising initial results, a comprehensive study addressing the interactions between MP-Ps and PVDF polymer, including their effects on structural, thermal, mechanical, and ferroelectric properties, remains absent from literature.

To address the growing demand for flexible, sustainable, and biocompatible materials for energy harvesting and storage, this study explores the integration of MP-Ps, specifically Ni, Ag, and Co-based phosphates, into a PVDF matrix. By tailoring particle morphology, size, and surface characteristics through various synthesis methods, we systematically investigate how these parameters influence the nucleation of the electroactive β-phase in PVDF. The resulting nanocomposites exhibit a remarkably high β-phase content exceeding 96%, a value significantly higher than those typically reported in the literature. This enhancement is attributed to strong interfacial interactions between the MP-Ps and PVDF chains, facilitated by the specific electronegativity and morphology of the metal phosphate fillers. In addition to promoting β phase formation, the incorporation of MP-Ps leads to simultaneous improvements in mechanical strength, thermal stability, and ferroelectric performance, demonstrating a synergistic enhancement across multiple functional properties. Importantly, the use of leadfree, phosphate-based fillers offers an environmentally friendly and potentially biocompatible alternative to traditional piezoceramics, thereby contributing to the development of safe and efficient energy harvesting systems for wearable and portable electronics. This work not only introduces a novel class of functional composites but also advances the fundamental understanding of phosphate-polymer interfaces and their role in optimizing multifunctional performance.

2. Materials and experimental details

2.1. Materials

All chemicals were obtained from commercial sources and used without further purification. Nickel(II) acetate tetrahydrate $(Ni(CH_3COO)_2 \cdot 4H_2O, \geq 98\%)$, silver nitrate (AgNO₃, $\geq 99.8\%$), cobalt(II) nitrate hexahydrate (Co(NO₃)₂·6H₂O₂, \geq 99.0%), and cobalt (II) acetate tetrahydrate (Co(CH₃COO)₂·4H₂O, \geq 98%) were purchased from Sigma-Aldrich for the synthesis procedures. Supporting reagents including ammonium dihydrogen phosphate (NH₄H₂PO₄), potassium hydroxide (KOH), sodium hydroxide (NaOH), ethylene glycol, and ethanol (all with purity >98.99%) were also obtained from Sigma-Aldrich. For composite preparation, polyvinylidene fluoride (PVDF) with a melting point of 170-180 °C and density of approximately 1.78 g cm⁻³ was sourced from Sigma-Aldrich, while dimethylformamide (DMF, 99% purity) was acquired from Merck KGaA.

2.2. Synthesis of nickel phosphate Ni₃(PO₄)₂

Hydrated nickel phosphate was synthesized by via a hydrothermal reaction with the supporting steps Fig. 1(a). Typically,

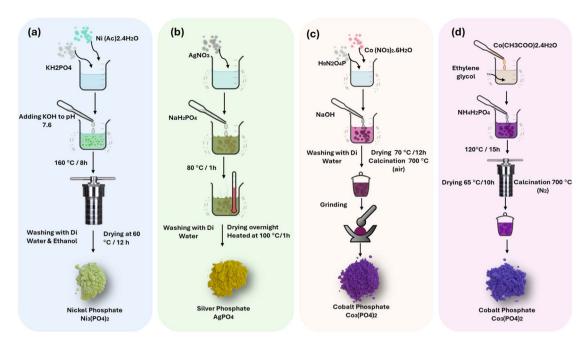


Fig. 1 Schematic illustration of the as-synthesized particles using different methods (a) Ni-P, (b) Ag-P, (c) Co-P and (d) Co-Pn.

a 1.4 g of Ni(Ac) $_2$ ·4H $_2$ O was dissolved in distilled water. The mixture then dropped with a solution of 0.52 g KH $_2$ PO $_4$ dissolved in 20 ml of solvent while the mixture was continuously stirred. The pH of the resulting solution was carefully adjusted to 7.6 using 2 M KOH solution. After mixing thoroughly, the suspension was placed in a Teflon-lined autoclave and held at 160 °C for 8 hours to accelerate the hydrothermal process. Once the reaction is complete, the product is removed from the mixture by centrifugation (Fig. 1). The obtained products were centrifuged, washed with absolute alcohol, and deionized water, and then dried at 60 °C for 12 h. These particles are denoted as Ni–P.

2.3. Synthesis of silver phosphate Ag₃PO₄

The Ag_3PO_4 particles were obtained *via* a precipitation method Fig. 1(b). A typical procedure involves dissolving 0.005 mol phosphate salt NaH_2PO_4 in 100 ml distilled water. This phosphate solution is then added dropwise to a 0.1 M $AgNO_3$ solution, prepared by dissolving 0.01 mol $AgNO_3$ in 100 ml distilled water. The addition is carried out with continuous stirring, resulting in the formation of a yellow suspension. The suspension is heated to 80 °C for 1 hour to promote reaction and ensure homogeneity. After heating, the mixture is filtered to separate the solid material, which is then rinsed three times with distilled water to remove any impurities. The washed particles are then dried overnight and heated at 100 °C for 1 h to ensure complete removal of moisture. These particles are denoted as Ag-P.

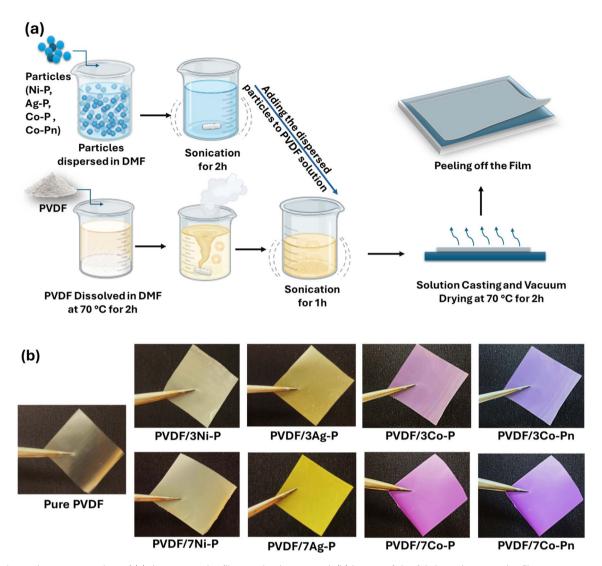
2.4. Synthesis of cobalt phosphate Co₃(PO₄)₂

Cobalt phosphate was synthesized using a co-precipitation method Fig. 1(c). To begin, stoichiometric volumes of $\text{Co(NO}_3)_2 \cdot 6\text{H}_2\text{O}$ (4.37 g), $\text{H}_9\text{N}_2\text{O}_4\text{P}$ (1.7 g), and NaOH (1.0 g) were

separately dissolved in 25 mL of deionized water. After that, these solutions were mixed to create a homogenous mixture with a volume of 75 ml. To ensure full reaction, the mixture was then stirred for hours at room temperature. After that a precipitate was formed, then collected and washed with deionized water until it reached a pH of 7. After washing, the particles were dried for 12 hours at 70 °C in an electric oven, and then the final step was annealing the dried product for 5 h at 700 °C. These particles are denoted as Co–P.

2.5. Synthesis of cobalt phosphate Co₃(PO₄)₂

Cobalt phosphate flakes were produced through a solvothermal approach Fig. 1(d). For a typical procedure, 0.75 g of Co (CH₃-COO)2·4H2O was first dissolved in 10 ml of ethylene glycol solution and stirring was maintained for 30 minutes. A 30 mL solution of 0.2 M NH₄H₂PO₄ was added to the mixture and stirred for an additional 2 hours to achieve a uniform mixture. The resulting solution was transferred to a Teflon-lined stainless-steel autoclave and heated at 120 °C for 15 hours. The resulting particles were collected, thoroughly washed with deionized water and ethanol, and dried at 65 °C for 10 hours. Then heat-treated at 350 °C for 5 hours in a nitrogen environment (N_2) at a heating rate of 3 °C min⁻¹. These particles are denoted as Co-Pn. To explore the effect of particle morphology on composite properties, cobalt phosphate (Co₃(PO₄)₂) was synthesized using two different methods: co-precipitation and solvothermal synthesis. These methods yield distinct particle morphologies, which are expected to influence the dispersion within the PVDF matrix and enhance β-phase formation. In contrast, nickel phosphate and silver phosphate were each synthesized using a single method (hydrothermal and precipitation, respectively), as these approaches produced the desired phase and morphology suitable for composite fabrication.



Schematic representation of (a) the composite film synthesis protocol, (b) image of the fabricated composite films.

Composite preparation

For composite preparation (Fig. 2), a solution was prepared by dissolving PVDF powder in DMF. The desired amounts of MP-Ps (Ni-P, Ag-P, Co-P, and Co-Pn) were dispersed separately in DMF using ultrasound for 2 hours to ensure uniform dispersion and minimize agglomeration. Simultaneously, PVDF was dissolved in DMF under continuous stirring at 70 °C for 2 hours until a homogeneous and clear solution was obtained, using a weight-to-weight (w/w) ratio of PVDF to DMF of 1:4. The predispersed particle solution was then added to the PVDF/DMF solution and continuously stirring for 1 hour to ensure thorough mixing. The mixture was further homogenized by ultrasonic treatment for an additional 1 hour to achieve complete dispersion of the particles in the polymer matrix. The resulting solution was spread onto a clean glass substrate using the Doctor blade technique to control the films thickness. The films were dried at 70 °C for 2 hours to evaporate the solvent and then carefully peeled off the substrate. The prepared films were labeled based on their composition: pure PVDF was designated

as "PVDF," while composites containing Ni-P, Ag-P, Co-P, and Co-Pn were labeled as "PVDF/xNi-P," "PVDF/xAg-P," "PVDF/ xCo-P," and "PVDF/xCo-Pn," respectively. Here, x represents the particle loading in the nanocomposites, which was either 3 or 7 wt%. The particles content is limited to 7 wt% to prevent negative effects on both the composite's mechanical properties and the β-phase formation. Higher concentrations can lead to particle agglomeration, which disrupts the uniformity of the composite and reduces the β-phase content, essential for piezoelectric performance.

3. Characterization techniques

3.1. Structural characterization X-ray diffraction (XRD)

A Bruker D8 Discover was used to perform X-ray diffraction (XRD) characterization utilizing Cu Ka radiation ($\lambda = 1.54184$ nm) in the 2θ range of 10–70° for particle analysis and in the range of 15–25° for elaborated composite films. Whereas the current and voltage were maintained at 100 mA and 40 kV, respectively.

3.2. Fourier transforms infrared spectroscopy

FTIR spectra of MP-Ps as well as neat PVDF, and PVDF composite films were conducted on FTIR, PerkinElmer Spectrum 2000 equipped with ATR accessory for the films, however KBR accessory for the powder. Each spectrum was obtained in $400-4000~\rm cm^{-1}$ range in the case of nanocomposites meanwhile in the range of $600-4000~\rm cm^{-1}$ for the powder of MP-Ps. The samples were analyzed with an accumulation of 16 scans and a resolution of 4 cm⁻¹.

3.3. Raman spectroscopy

The vibrations of the synthesized MP-Ps were investigated using Raman scattering spectroscopy using a HORIBA LABRAM-HR Evolution device under green laser illumination in the wavenumber range of $150-1000~\rm{cm}^{-1}$.

3.4. Scanning electron microscope

In order to characterize the shape of MP-Ps and assess their distribution inside the polymeric film matrix, a Zeiss EVO 10 scanning electron microscope (Carl Zeiss Microscopy, GmbH, Jena, Germany) was employed. To obtain a precise and clean fracture surface, the films were cryo-fractured using liquid nitrogen. Prior to scanning, the samples were made conductive by applying a gold coating using a sputtering process.

3.5. Thermogravimetric analysis

The thermal stability and degradation behaviors of synthesized MP-Ps, neat PVDF as well as the composites films were conducted by thermographic analysis using TA instruments (Discovery TGA). Under a nitrogen atmosphere with a gas flow of 20 ml to prevent oxidation effects and heated between 25 and 700 °C at a heating rate of 10 °C min⁻¹.

3.6. Differential scanning calorimetry

The melting and crystallization behavior of elaborated films were investigated using Differential scanning calorimetry (DSC) (Discovery DSC, TA instruments). During the analysis process, the samples were gradually heated from 20 to 200 °C at a rate of 10 °C min $^{-1}$. Then, they were held at 200 °C for three minutes to destroy any preceding thermal or mechanical effects. Next, the samples were cooled from 200 °C to -80 °C, also at a rate of 10 °C min $^{-1}$, to further assist in observing their crystallization behavior. After a stabilization period of three minutes at -80 °C, the samples were heated back from -80 to 200 °C at the similar rate as before to examine their melting properties.

The degree of crystallinity (X_c) was calculated according to the following equation:

$$X_{\rm c}(\%) = \frac{\Delta H_{\rm m}}{F \times \Delta H m_0} \times 100$$

where F is the weight fraction of PVDF and $\Delta H_{\rm m}$ is the enthalpy of melting of 100% crystalline PVDF and was taken as 104 J g⁻¹.

3.7. Tensile test

The tensile characteristics of developed films were assessed using a universal testing machine (SHIMADZ, Kyoto, Japan) fitted with a 1 KN load cell in accordance with the ASTM D882-00 standard. With a measured gauge length of 50 mm and a moving clamp speed of 5 mm min $^{-1}$, the film samples were cut into 10 \times 50 mm² rectangles. No fewer than five samples were used for each test, and the findings are average.

3.8. Ferroelectric analysis

P–E hysteresis loops were obtained by Radiant Precision Workstation ferroelectric testing system at room temperature equipped with a high-voltage source of 10 kV. The Vision5 data management software was used to obtain results at a frequency of 5 hertz with the normal dipolar mode, which produced a triangular signal of high voltage and enabled us to measure the subsequent polarization of PVDF composite films.

Results and discussions

4.1. Characterization of synthesized metal phosphate-based particles (MP-Ps)

4.1.1. Structural characteristics. X-ray diffraction (XRD) analysis was employed as a key characterization technique to investigate the crystalline phases and structures of the synthesized particles. This method provided essential information, including phase identification, sample purity, crystallite size, and, in certain cases, morphology. To verify the successful development of the expected crystalline phases and ensure the structural integrity and purity of the produced particles, the diffraction patterns were analyzed.32 The XRD patterns of the prepared Ni-P, Ag-P, Co-P and Co-P nano-flakes are shown in Fig. 3. The X-ray diffraction (XRD) patterns of the synthesized MP-Ps confirm their distinct crystal structures and phase purities. The X-ray diffraction analysis revealed a monoclinic crystal structure characterized by multiple well-defined reflections. The diffractogram exhibited its maximum intensity at 2θ = 13.35° , which was indexed to the (020) crystallographic plane. The structural characterization was further confirmed by additional reflections observed at various diffraction angles: 11.22° (110), 18.40° (200), 19.68° (-101), 20.78° (011), 22.03° (130), 23.36° (101), 24.75° (-211), 27.03° (040), 28.29° (031), 30.57° (-301), 32.77° (240), 33.51° (-321), 34.46° (330), 36.01° (141), $37.49^{\circ}(301), 39.40^{\circ}(-112), 41.16^{\circ}(-341), 41.96^{\circ}(-251), 44.32^{\circ}$ (350), and 47.78° (132). This diffraction pattern is consistent with the characteristic signature of the monoclinic phase.33,34 For Ag-P, the X-ray diffraction analysis revealed a well-defined cubic crystal structure characterized by eight distinct diffraction peaks. The diffractogram exhibited reflections at $2\theta = 20.9^{\circ}$ (110), 29.7° (200), 33.3° (210), 36.6° (211), 47.8° (222), 52.7° (320), 55.0° (321), and 71.9° (520). This pattern of crystallographic planes consists with the characteristic signature of the cubic structure of silver phosphate, confirming the successful synthesis of phase-pure consistent with the JCPDS Card No. 06- $0505.^{35,36}$ The XRD analysis revealed that both Co–P and Co–Pn samples crystallized in the hexagonal phase of Co₃(PO₄)₂, with

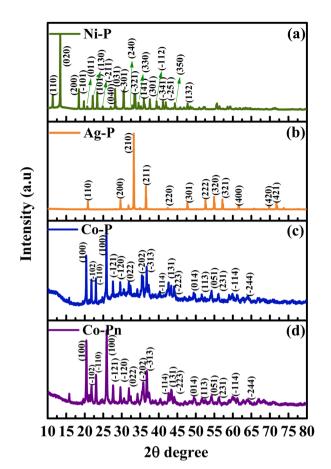


Fig. 3 XRD diffractograms of synthesized (a) Ni-P, (b) Ag-P, (c) Co-P, (d) Co-Pn.

diffraction patterns matching JCPDS Card No. 70-1795. The observed crystallographic planes were indexed according to previously reported literature values for this structure.^{37,38} Overall, the XRD analysis confirms that all synthesized MP-Ps exhibit high crystallinity and phase purity, with no indications of impurity phases, providing valuable insights into their structural properties.

Fourier Transform Infrared (FTIR) spectroscopy is an essential technique for characterizing the as-prepared particles, offering detailed insights into their molecular structure. By identifying unique vibrational modes, FTIR reveals the functional groups, bonding arrangements, and chemical composition of the particles. This information is crucial for confirming the successful synthesis and understanding the interactions within the material.39 The FTIR spectra of MP-Ps presented in Fig. 4 was used to further confirm their bonding arrangement. All spectra indicate the presence of characteristic bands at 882 cm⁻¹, 741 cm⁻¹, 597 cm⁻¹, 580 cm⁻¹, 557 cm⁻¹, 556 cm⁻¹ and 450 cm⁻¹ referring to literature the vibrational bands occur within this range is (ν_3) (F_2) PO₄³⁻ ascribed to the metal oxygen bond.40-42 Furthermore, the vibrational bands $1005~{\rm cm}^{-1}$, $1049~{\rm cm}^{-1}$, $1031~{\rm cm}^{-1}$ and $1050~{\rm cm}^{-1}$ are due to the triply degenerated (ν_3) asymmetric stretching vibrations of P-O bonds.43 For the samples Ag-P, Ni-P there is the presence of a vibration bonds at 1665 cm⁻¹ and 1622 cm⁻¹ respectively, are

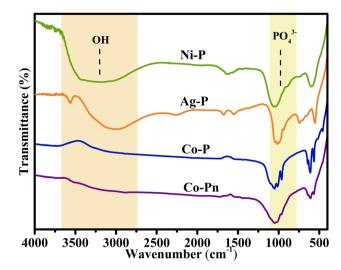


Fig. 4 FTIR spectra of the synthesized MP-Ps

ascribed to the H–O–H bending vibration from the structural water $^{44-46}$ and also a broad peaks ranging 3500 cm $^{-1}$ to 3000 cm $^{-1}$ stretching vibration and O–H stretching in crystallization water. Furthermore, it is observed from the FTIR spectra of the Ag–P particles the presence of a bond at 3555 cm $^{-1}$ related to hydroxyl group –OH adsorbed on the surface of the Ag–P particle νia surface hydrolysis and protonation. 48,49

Raman spectroscopy is a highly reliable and widely used tool for characterizing materials, as it provides valuable information about the structural quality and can also reveal the presence of dopants or other modifications.50 Fig. 5 represents the Raman spectrum of MP-Ps. The internal modes in Raman spectroscopy typically involve the displacement of atoms within molecular units, such as the oxygen atoms in tetrahedral anions like $(PO_4)^{3-}$. These modes exhibit frequencies that are closely related to those of the free anion. For example, the free (PO₄)³⁻ ion displays a characteristic symmetric stretching mode (A₁₀) around 938 cm⁻¹. In nickel phosphate, similar vibrational frequencies can be observed, reflecting the internal dynamics of the phosphate groups, the intense peak of 908 cm⁻¹ is attributed to the terminal oxygen vibrational stretching of the PO₄ group for the silver phosphate.51 The Raman peaks around 1000 cm⁻¹ for the cobalt phosphate are from the triply degenerate n3 antisymmetric stretching vibrations $(A_{1g} + E_g)$ of the phosphate ligands.52

4.1.2. Morphological and chemical characteristics. The combined use of scanning electron microscopy (SEM) and energy-dispersive X-ray spectroscopy (EDX) is a crucial technique for the characterization of the synthesized particles, providing detailed insights into their size, shape, elemental composition, mixing state, surface area estimates and surface features.⁵³ The SEM images of synthesized particles reveal distinct morphological characteristics due to the varied synthesis methods employed. Each material displays unique structural features, ranging from particle size and shape to surface texture, all of which are critical to understanding their

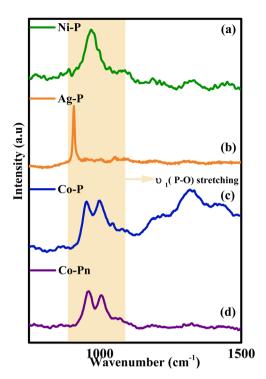


Fig. 5 Raman spectra of the synthesized (a) Ni-P, (b) Ag-P, (c) Co-P, (d) Co-Pn.

interaction with the PVDF matrix. The grain sizes and dimensions reported for the synthesized particles were estimated using ImageJ software by analyzing the SEM images. Fig. 6(a) shows that Ni-P synthesized through hydrothermal method exhibits a sphere like shape with practically homogeneous distribution and an average particles size of 39.42 nm. From Fig. 6(b) the synthesized Ag-P via precipitation method shows a cube-like shape indicating a single crystalline morphology of the particles with relatively smooth surface and bimodal particle size, presenting a population of small particles averaging 2.68 µm and big particles averaging 9.430 µm. Fig. 6(c) shows the morphology of Co-P synthesized using Coprecipitation method, it displays irregular rod-like, the particle size distribution also shows two distinct groups: larger particles with average of 1.43 µm with moderate variability, and smaller particles with average of 220 nm that are more uniform. This may be attributed to the wet chemical reaction employed for synthesizing these particles, as the slow reaction kinetics can result in a non-uniform nucleation rate.54 Meanwhile, Co-Pn synthesized using solvothermal reaction, as depicted in Fig. 6(d) shown a well-defined flakes-like morphology, with almost homogeneous distribution with practically smooth surface. On average, they have a thickness of 337 nm, a length of 16.38 μ m, and a width of 1.69 μ m.

For all the particles, the EDX spectrum confirmed the predictable chemical composition, while the elemental mapping images showed a uniform distribution of elements, indicating consistent synthesis (Fig. S1, ESI†). In summary, the diverse morphologies and scales of the synthesized materials, directly influenced by the different synthesis methods, are

expected to play a crucial role in determining the overall structural, thermal, and electrical properties of the resulting composites. Furthermore, the particle size distribution and morphology are expected to impact on the dielectric and ferroelectric properties of the composites. This study is therefore vital in assessing how these MP-Ps can improve the ferroelectric behavior and β -phase content of PVDF, leading to high-performance materials.

4.1.3. Thermal characteristics. Thermal characterization of the synthesized particles was performed using thermogravimetric analysis (TGA). The weight changes observed during the temperature variations provide insights into the composition of the particles and their thermal stability.55 TGA curves depicted in Fig. 7 revealed distinct degradation behaviors for the four synthesized MP-Ps (Ni-P, Ag-P, Co-P, and Co-Pn). The Ni-P particles exhibited two stages of degradation. The first stage, occurring below 30 and 120 °C, was attributed to the loss of physically adsorbed water,55 while the second stage occurring between 120 and 340 °C corresponded to Fig. 8 the decomposition of chemically bound crystalline water. 56 In contrast, Ag-P displayed a stable thermal profile with no observable degradation stages, indicating excellent thermal stability, a behavior similarly observed for Co-P, which also showed no signs of thermal degradation. On the other hand, Co-Pn demonstrated two degradation stages. The first accruing bellow 100 °C was due to the loss of adsorbed water or residual solvent. Notably, the second stage involved a 2% weight loss at approximately 479 °C, reflecting the breakdown of surface-bound organic residues, likely stemming from the ethylene glycol used as a solvent and surface modifier.57 These findings underline the varying thermal behaviors of the particles, influenced by their composition and synthesis conditions, with Ag-P and Co-P showing high thermal resilience, while the degradation of Ni-P and Co-Pn highlights the impact of water and organic additives on their thermal stability.

4.2. Characterization of composite films

4.2.1. Structural characteristics. Fourier Transform Infrared spectroscopy (FTIR) plays a vital role in the characterization of the β-phase in PVDF, as it effectively distinguishes the electroactive β -phase from the non-electroactive α -phase through its unique vibrational modes. This technique provides precise and reliable phase identification, making it indispensable for analyzing PVDF's phase composition and validating the presence of electroactive phases in polymeric systems. 58 Fig. 9 represents the FTIR spectrum of composite materials-based MP-Ps. The spectrum of pure polyvinylidene fluoride (PVDF) for all composition exhibits distinct absorbance bands typical of nonpolar α-phase. These include bands at 488 cm⁻¹ (CF₂ wagging), 532 cm $^{-1}$ (CF $_2$ bending), 615 cm $^{-1}$ and 764 cm $^{-1}$ (CF $_2$ bending and skeletal bending), and 796 cm⁻¹ and 976 cm⁻¹ (CH₂ rocking). A slight peak at 838 cm⁻¹ (CH₂ rocking, CF₂ stretching, and skeletal C-C stretching) is also present, suggesting a slight presence of β or $\beta + \gamma$ crystals, as supported by previous research findings.58 The lack of a characteristic absorbance band for γ crystals around 1234 cm⁻¹ indicates that

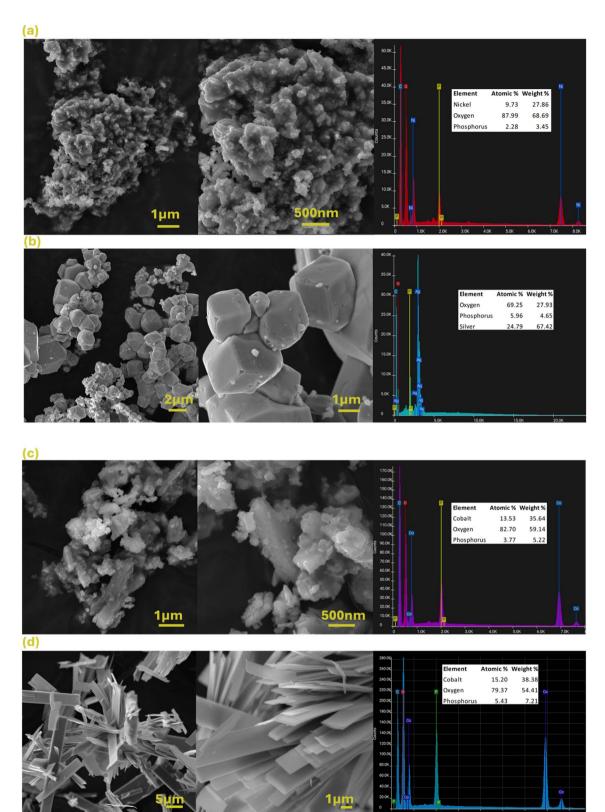


Fig. 6 SEM images and EDX spectrum of the as-synthesized particles: (a) Ni-P, (b) Ag-P, (c) Co-P and (d) Co-Pn.

the minor peak around 840 cm^{-1} is likely due to the $\beta\text{-phase,}$ signifying some polymer chain alignment in the TTTT conformation within the pure PVDF film.58 In contrast, the composite

samples exhibit a complete disappearance of the characteristic peaks associated with nonpolar α -phase. Prominent absorbance bands emerge at 445 cm⁻¹ (CF₂ and CH₂ rocking), 479 cm⁻¹

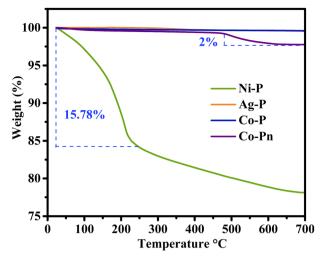


Fig. 7 TGA curves of synthesized MP-Ps.

(CH₂ deformation), 510 cm⁻¹ (CH₂ stretching), 600 cm⁻¹ (CH₂ wagging), 838 cm⁻¹, and 1274 cm⁻¹. The presence of bands at 510 cm⁻¹, 838 cm⁻¹, 445 cm⁻¹, and 1274 cm⁻¹, alongside the absence of the γ crystal band at 1234 cm⁻¹ support the presence of electroactive β -phase in PVDF film which suggests that even a minimal filler concentration of 3 wt% can entirely suppress

the α -phase while encouraging the formation of the electroactive β -phase. This aligns with the findings previously confirmed by the XRD. Beside to that, an absorbance band have been detected at 960 cm⁻¹ observed in higher particle loading of composite PVDF/xAg-P and PVDF/xCo-P related directly to the ionic phosphate group PO₄^{3-,59} is likely ascribed to the incorporation of MP-Ps, providing further evidence of their presence and complexation within the host polymer matrix.³⁰ This validates the possible interaction between these particles and PVDF, which is further supported by the XRD findings.

The fraction of β -phase in different samples is displayed through quantitative measurement, which confirms this conversion even further. The following formula is used to determine β -phase percentage in PVDF:

$$F(\beta) = \frac{A_{\beta}}{\left(\frac{k_{\beta}}{k_{\alpha}}\right) A_{\alpha} + A_{\beta}} \tag{1}$$

4.2.2. Morphological characteristics. The morphological alteration of the prepared films was assessed using the SEM. From Fig. 10 it is seen that for the pure PVDF surface, a globular structure with a porous region between the globes has been detected. However, a more smooth and dense morphology is observed from the cross-section. Surface globular structure has

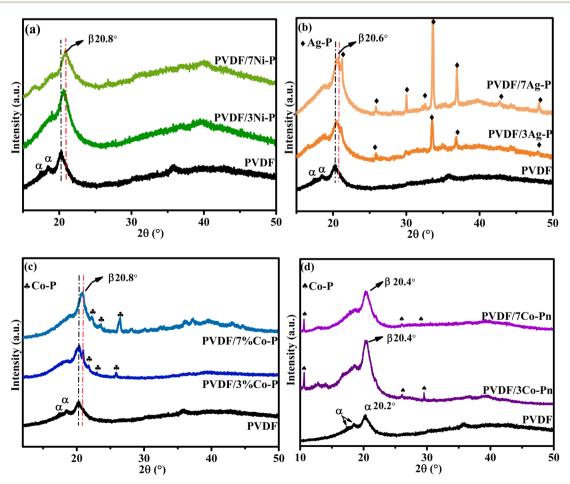


Fig. 8 XRD patterns of composite films (a) PVDF/Ni-P, (b) PVDF/Ag-P, (c) PVDF/Co-P and (d) PVDF/Co-Pn.

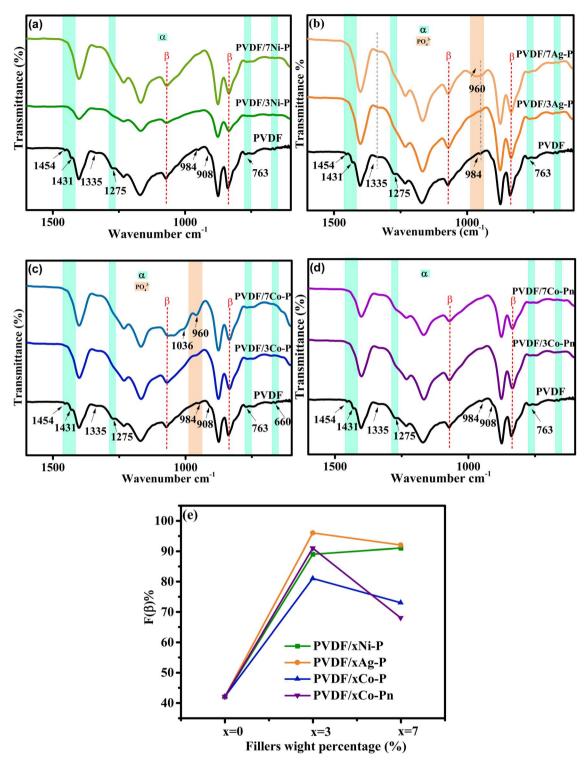


Fig. 9 FTIR spectrum (a)–(d) along with β -phase percentage (e) of the elaborate composite films.

seen to be smoother with low percentage addition of the Ni–P nanoparticles for the composite PVDF/3Ni–P. Along the cross-section it is observed a homogeneous distribution of the Ni–P nanoparticles with no aggregates. Compared to the surface morphology of the Neat PVDF, a relevant change it is noted suggests that the Ni–P particles affect the morphology of PVDF/xNi–P composites. With the percentage of 7 wt% of the Ni–P same

surface morphology is appearing but with reduced pore size and deformed globule shape. From the cross section homogeneous distribution of the Ni–P is observed without relative agglomeration proposing excellent dispersion within the polymer matrix. Indicating that the filler phase acts as a crystal nucleating agent affecting the size and the perfection of the polymer crystal, 60 confirming the previous results from FTIR and XRD.

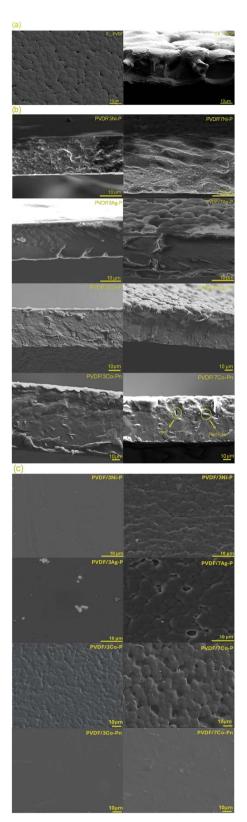


Fig. 10 SEM images of (a) neat PVDF, (b) cross-sectional view of composite films, and (c) surface morphology of composite films.

For the composite PVDF/xAg-P, following a similar trend with the addition of 3 wt% of Ag-P as low percentage the surface shows to be smooth compared to pure PVDF with some

minimal aggregation of particles in the surface, also from the cross section the surface is dense with relatively no agglomerations. Furthermore, the addition of 7 wt% of Ag–P with a high aspect ratio showing no obvious aggregation either on the surface nor in the cross-sectional part but with a globular structure and larger pores on the surface. Demonstrating a change in the crystalline structure. The shift from a globular structure to a smoother one can be explained by the fact that small blocks of lamella are pulled away from the original lamellae to form a fibrillar structure of crystallites, this microstructural alteration is linked to the α - to β -phase changeover. This aligns perfectly with the previously discussed findings from XRD and FTIR.

For the composite PVDF/xCo-P, a globular structure is observed at both low (3 wt%) and high (7 wt%) filler concentrations. At the lower concentration, the surface appears denser with minimal voids, while larger pores become evident at higher filler concentrations. This structural change is attributed to matrix modification. At higher filler levels, some particles are partially embedded in the surface, appearing both exposed and immersed, which increases surface roughness compared to pure PVDF. Cross-section reveals a dense surface with no visible particle agglomeration. The absence of gaps between the particles and the matrix suggests strong interactions between the polymer and the particles.⁶²

In contrast, for the composite PVDF/xCo–Pn, smooth surfaces are observed at both filler concentrations, with the particles fully embedded within the film surface. Cross-sectional examination indicates that the particles are well incorporated into the polymer matrix without observable gaps. However, minor voids are noted, potentially caused by particle pull-out during fracture. 62 This phenomenon is more prominent at higher filler concentrations and could be linked to the initiation of agglomeration, which may hinder interactions between the polymer and the fillers. These observations align with previous findings, where the β -phase content decreases at higher filler concentrations, indicating reduced interactions between the polymer matrix and the fillers.

This leads us to the conclusion that the structure and interactions within the host polymer matrix are significantly impacted by the morphology of the filler particles, which leads to modifications in the overall properties of the composite. ⁶³ Dispersion of the fillers is another important issue that can have a big impact on the performance of the composite. Particle agglomeration can produce stress concentration points, which is well recognized in composite materials and may weaken the mechanical properties of the composite. ⁶⁴ Achieving good dispersion, as seen in most of the composites, plays a critical role in improving their mechanical performance.

4.2.3. Thermal characteristics. Thermogravimetric analysis (TGA) is a widely used analytical technique that has been employed to give essential information about the thermal behavior of polymers and polymer composites. The TGA curves for the pure PVDF film and its composites are shown in Fig. 11, providing insights into their weight loss profiles. Complementary information is provided in the derivative thermogravimetric (DTG) curves, displayed in Fig. S2 of the ESI,†

which further elucidates the degradation patterns and maximum decomposition rates of the materials. And Table 1 summarizes the values of initial degradation temperature (T_{onset}) , maximum degradation rate temperature (T_{max}) , obtained through the peak of the mass loss derivative, and residue content, corresponding to the remaining mass at 700 °C.

From the current data it is observed that the pure PVDF film is stable up to 340 °C, after which follows a one-step degradation. The process of breakdown seen in pure PVDF involves the carbon-hydrogen bond breaking, which produces hydrogen fluoride, and then more hydrogen fluoride loss throughout the polymer chains.66 The incorporation of MP-Ps into the PVDF matrix resulted in a notable increase in the onset degradation temperature of the composites, indicating a significant enhancement in the thermal stability of PVDF when combined with MP-Ps.67 Moreover, the thermal stability further improved with increasing filler content in the composites. The first derivative peak temperature of pure PVDF was observed at 484 $^{\circ}$ C. For composites containing Ni–P filler, the peak temperatures were 500 °C and 505 °C for 3 wt% and 7 wt%, respectively. Similarly, PVDF/xAg composites exhibited peak temperatures of 503.31 °C and 505.77 °C for 3 wt% and 7 wt%, respectively.

Table 1 Thermogravimetric results of pure PVDF and composites

Samples	$T_{ m onset}$ (°C)	T_{max} (°C)	Char content%
PVDF neat	470.44	484.30	23.21
PVDF/3Ni-P	490.79	499.46	21.34
PVDF/7Ni-P	495.36	503.93	23.15
PVDF/3Ag-P	489.66	503.31	25.33
PVDF/7Ag-P	490.42	505.77	26.88
PVDF/3Co-P	473.99	490.69	25.92
PVDF/7Co-P	483.19	498.70	29.36
PVDF/3Co-Pn	476.06	499.46	24.45
PVDF/7Co-Pn	489.64	503.77	19.08

Composites with Co-P particles showed peak temperatures of 490.69 °C and 498.70 °C for 3 wt% and 7 wt%, respectively, while composites with Co-Pn particles displayed peak temperatures of 499.46 °C and 503.77 °C for 3 wt% and 7 wt%, respectively.

The thermal behavior of the PVDF composites can be directly linked to the thermal characteristics of the incorporated MP-Ps depicted in Fig. 7. The improved thermal stability observed in the composites is influenced by the high thermal resilience of

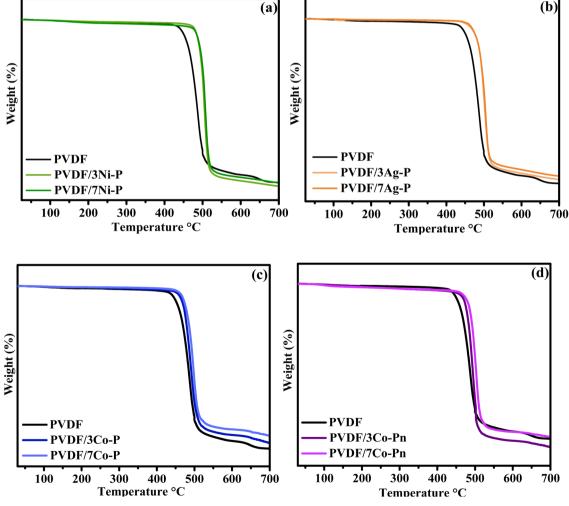


Fig. 11 TGA curves of PVDF and composite films (a) PVDF/Ni-P, (b) PVDF/Ag-P, (c) PVDF/Co-P and (d) PVDF/Co-Pn.

MP-Ps particles like Ag-P and Co-P, which exhibit no significant degradation in TGA analysis. These stable particles enhance the overall thermal performance of the PVDF matrix. Conversely, the degradation stages of Ni-P and Co-Pn, associated with water loss and organic residue decomposition, suggest that their influence on the composite's stability is partly due to these factors. Nonetheless, the uniform dispersion of MP-Ps within the PVDF matrix likely mitigates these effects, contributing to the observed increase in the degradation temperatures, especially at higher filler concentrations.⁶⁸

Understanding the effect of these fillers on the enhanced thermal stability of the composite requires first examining the degradation mechanism of PVDF. This will help determine whether the improvements are linked to phase transitions and changes in the crystalline structure, as previously noted in other analyses. Hence, to create C-C double bonds, the PVDF begins with the generation of free radicals by first scissoring the -C-H bond and then the -C-F bond.69 The elimination of hydrogen fluoride is subsequently spread across the chains of polymers. Thus, mass loss happens in two stages: HF elimination and the polymer chain's disintegration. Meanwhile, there are other factors involved in this enhancement. MP-Ps can function as radical scavengers, effectively inhibiting chain transfer reactions and preventing the decomposition of polymer chains. As a result, the nucleation rate is reduced, requiring more time for volatiles to accumulate to a critical concentration.70 This leads to an improvement in the thermal stability of the composite compared to neat PVDF.

Notably, the char content shows an increase for composites containing Ag-P and Co-P, while a decrease is observed for composites containing Ni-P and Co-Pn. The improvement in char content for composites containing Ag-P and Co-P can be attributed to the inherent thermal stability and high thermal resilience of these particles. Both Ag-P and Co-P exhibit minimal or no significant degradation during TGA analysis, as they lack water content and organic residues that contribute to weight loss. This stability ensures that these particles form a robust, thermally resistant residue at higher temperatures, thereby increasing the char content in the composites. In contrast, composites containing Ni-P and Co-Pn show relatively lower char content due to the thermal behavior of these particles. Ni-P undergoes significant weight loss in two degradation stages, primarily due to the loss of physically adsorbed water and decomposition of chemically bound crystalline water. Similarly, Co-Pn exhibits degradation related to the loss of water and the breakdown of surface-bound organic residues. These processes reduce the amount of stable residue remaining at higher temperatures, resulting in a lower char content in the composites.

Likewise, the melting and crystallization behaviors of PVDF and PVDF composites are measured using a DSC analyzer. Additional insights into the influence of fillers on polymer crystallinity are obtained through these DSC determinations. The melting temperature ($T_{\rm m}$), enthalpy of melting ($\Delta H_{\rm m}$), crystallization temperature ($T_{\rm c}$), and crystalline ($X_{\rm c}$) were extracted from the DSC curves, as shown in Fig. 12 and Table 2. These parameters provide valuable insights into the thermal

behavior and stability of the materials. The melting temperature of pure PVDF was found to be 162.94 °C. With the addition of the fillers, it was observed that melting endotherms are broader, and their peak temperatures shifted slightly to lower temperatures. Changes in the melting temperature can be attributed to alterations in the crystal structures of PVDF within the composites. This suggests that the formation of double bonds during the reaction may slightly modify the main chains of PVDF, thereby affecting the melting temperature. This phenomenon can confirm the presence of the β-phase, which is known to have a melting point lower than that of the α phase crystal of PVDF, as documented in the literature. 67,71,72 The changes observed in each composite depend on its composition. Variations in melting points suggest differences in particle-PVDF interactions, which align with SEM images showing distinct morphological modifications in each material. Furthermore, in the same tendency the melting enthalpy and crystallinity shows a significant increment, which means that more energy is required to break the crystalline structure which indicates stronger intermolecular interactions confirming the FTIR finding. The presence of strong intermolecular interactions, such as hydrogen bonding or dipole-dipole interactions, within the composite film can contribute to higher melting enthalpy. In the case of PVDF composites, if the β-phase is prevalent, the strong polar interactions between the parallel dipoles in the TTT conformation would be a factor and that could be explained by the previous result from the XRD and FTIR. 13 The inclusion of fillers or additives in the composite film can enhance the crystallization process or reinforce the polymer matrix, contributing to the increased melting enthalpy. These fillers could act as nucleating agents, promoting the formation of crystalline regions. High crystallinity often leads to improved mechanical properties, such as increased stiffness, tensile strength, and toughness. The composite films would be more robust, and durable compared to neat PVDF.

4.2.4. Mechanical characteristics. Research on the mechanical properties of composite materials used in piezo-electric applications is critically important, as these properties directly affect the material's ability to convert mechanical stress into electrical energy. The mechanical performance of piezo-electric composites influences their durability, flexibility, and overall efficiency in energy harvesting and sensing applications, making it essential to optimize these properties for enhanced functionality.⁷³ Significant variables that affect the overall mechanical properties of the composite systems including particles size, shape, distribution, and dispersion within the polymer matrix, as well as their reinforcing and interfacial compatibility.^{74,75}

Therefore, comprehensive tensile test has been carried out to examine the effects of the shape and size of the synthesized particles (Ni–P, Ag–P, Co–P and Co–Pn) and their interactions, particularly their interfacial adhesion within the PVDF host matrix on the overall properties of the prepared composites. The tensile stress–strain graph provides key parameters: Young's modulus, tensile strength, and strain at break. Young's modulus reflects the material's ability to store mechanical energy during elastic deformation.⁷⁴ Tensile strength which is

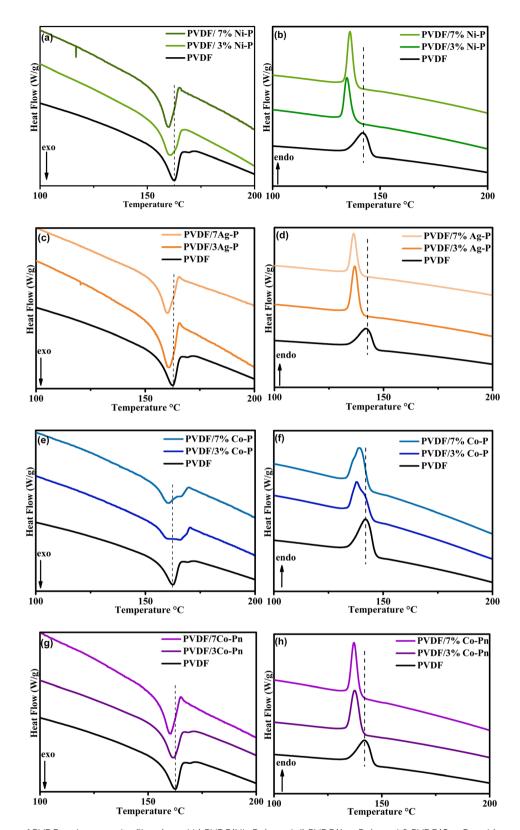


Fig. 12 DSC curves of PVDF and composite films (a and b) PVDF/Ni-P, (c and d) PVDF/Ag-P, (e and f) PVDF/Co-P and (g and h) PVDF/Co-Pn.

known as the maximum stress that a material can resist when subjected to uniaxial tensile. Strain at break indicates ductility, with higher values signifying greater flexibility.²² The findings are shown in detail in Fig. 13.

The results indicate that all the produced composites exhibit a significant improvement in Young's modulus compared to neat PVDF. Composites containing Ni–P and Ag–P demonstrated similar enhancements at both loading percentages

Table 2 Data obtained from DSC analysis

Samples	T_{m} (°C)	$\Delta H_{ m m} \left({ m J g}^{-1} ight)$	X_{c} (%)
PVDF neat	162.49	18.67	17%
			, ,
PVDF/3Ni-P	160.33	21.76	21%
PVDF/7Ni-P	159.76	24.76	25%
PVDF/3Ag-P	160.74	27.89	27%
PVDF/7Ag-P	160.20	26.13	27%
PVDF/3Co-P	159.91	19.60	20%
PVDF/7Co-P	159.91	20.32	21%
PVDF/3Co-Pn	161.44	22.47	22%
PVDF/7Co-Pn	159.88	27.07	28%

(3 wt% and 7 wt%), with their Young's modulus exceeding twice the value of neat PVDF. Meanwhile, composites with Co–P also showed improved Young's modulus, although the enhancement was moderate compared to PVDF/xNi–P and PVDF/xAg–P. In contrast, composites filled with Co–Pn displayed a distinct behavior: a significant improvement in Young's modulus was observed at 3 wt% loading, but the modulus decreased at higher particle loading (7 wt%). It is reported that the Young's modulus is generally insensitive to the particles size and generally increased with increasing particles loading since rigid inorganic MP-Ps generally have a much higher stiffness than polymer matrix.⁷⁴ This increase with an increase in fillers fraction, as expected due to the higher intrinsic strength and stiffness of

these particles compared with the PVDF matrix. Suggesting their excellent reinforcing effect as they make the composites stiffer by restricting the mobility of PVDF macromolecules under tensile stress. Such significant improvements in mechanical properties are largely attributed to the good dispersion of the fillers and their very high specific surface areas that help distribute load from the polymer matrix to the fillers. 15,76 In contrast to the other composites, we find that composite PVDF/7Co-Pn has a noticeably lower Young's modulus. The reason behind this decrement could be directly supported by the prior assumption from SEM observation that there are tiny voids in this composition, most likely brought on by the initial phase of agglomeration. Factors like entangled molecular chains, voids, and other defects usually weaken and decrease stiffness, which has a negative impact on the mechanical properties.76

The effect of different particle loading on the tensile strength of the composite polymer also is discussed. As was discussed previously, tensile test results showed an increase in elastic modulus with increasing powder loading using different MP-Ps (Ni-P, Ag-P, Co-P and Co-Pn) with different size and morphology. In contrast, the tensile strength of the composite generally decreased with increasing particles loading beyond. This refers to a variety of parameters. It is demonstrated that the strength of composites is determined not only by particle size and particle/matrix interfacial adhesion but also by particle

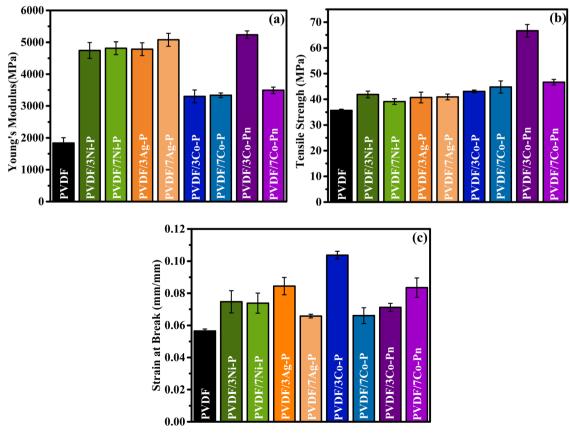


Fig. 13 Tensile test of neat PVDF and the composites: (a) Young's modulus, (b) tensile strength, and (c) strain at break.

loading.⁷⁷ From the available data, it is observed that there is an enhancement in the strength of particularly all the composite materials with Ni-P, Ag-P, Co-P and Co-Pn with both 3 wt% and 7 wt% of loading, this could be interpreted in first class by the excellent adhesion between this particles and the PVDF matrix, related to the interaction acquiring between this latter and the host matrix as it is evident from the previous founding supports by the XRD and FTIR. The tensile strength of polymerbased composites is significantly influenced by the quality of filler dispersion within the host matrix and the interfacial interactions between the fillers and the polymer. For flake-like Co-Pn particles, a 3 wt% loading demonstrates higher tensile strength due to better dispersion and enhanced stress transfer at the interface, resulting in superior reinforcement. In contrast, at 7 wt% loading, the tensile strength decreases, likely due to filler agglomeration, which creates stress concentrations and weakens the polymer-filler interface.74 The planar morphology of flakes allows for reinforcement in two dimensions, offering higher mechanical performance when the fillers are well-aligned and evenly distributed.78 Poor dispersion, however, can lead to low interfacial adherence, compromising the composite's tensile strength.⁷⁹

By evaluating the elongation at the break parameter, the produced films containing Ni-P, Ag-P, Co-P, and Co-Pn particles were further examined for ductility and flexibility. The composites capacity to withstand significant deformation prior to failure is shown by their greatest elongation value.80 The findings indicate that composites containing 3 wt% of fillers exhibit the highest elongation at break. For composites with PVDF/xNi-P, similar elongation at break is observed for both filler loadings (x = 3 wt% and 7 wt%), which can be attributed to the homogeneous dispersion and strong adhesion between the nanoparticles and the PVDF matrix, as previously confirmed by SEM analysis. As discussed, XRD and FTIR result also confirm these strong interactions. Further supported by the enhancement of the β-phase at higher filler percentages. The improved interaction can be linked to the small size and large surface area of the nanoparticles, which increases interfacial adhesion and promotes better compatibility with the polymer matrix. For

PVDF/xAg-P and PVDF/xCo-P composites, a similar trend is observed, with elongation at break being enhanced at 3 wt% filler loading but decreasing at 7 wt% loading. This decrease can be attributed to the initiation of particle aggregation at higher loadings, which directly impacts interfacial adhesion. Consequently, this weakens the matrix-filler interaction, leading to the observed reduction in elongation at break with increased filler content. For composite PVDF/xCo-Pn different behavior is remarqued with enhancement of the it could relate to the planar morphology of these flakes like particles providing large contact area, which resulted in the increase in strain at break providing more flexibility and ductility.⁸¹

To conclude, the study highlights the significant influence of particle type, size, morphology, and loading on the mechanical properties of PVDF composites. The incorporation of MP-Ps (Ni-P, Ag-P, Co-P, and Co-Pn) enhances Young's modulus and tensile strength due to improved filler dispersion, interfacial adhesion, and stress transfer within the matrix. However, excessive filler loading, particularly with flake-like Co-Pn particles, leads to agglomeration, void formation, and reduced mechanical performance. The ductility and flexibility of the composites are optimized at lower filler loadings (3 wt%), making these materials promising for applications in energy harvesting and piezoelectric devices where mechanical robustness and flexibility are critical.

4.2.5. Ferroelectric characteristics. *P–E* hysteresis loops provide critical information regarding a material's polarization behavior and serve as essential evidence for confirming ferroelectric character by demonstrating the nonlinear relationship between polarization and electric field, including key parameters such as remnant polarization (P_r) and maximum polarization (P_{max}), which quantify the material's ability to maintain polarization after field removal and its total polarization capacity, respectively. Fig. 14 shows P–E hysteresis loops for the composites at a voltage of ± 5 kV. Among all prepared composites, only PVDF/Ni–P and PVDF/Co–P were selected for PE-hysteresis loop measurements, as they demonstrated the most promising behaviour in terms of morphology, filler dispersion, and β phase enhancement. This strategic choice was

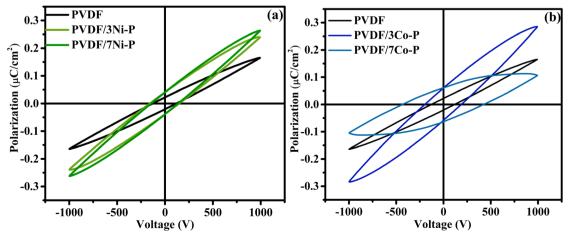


Fig. 14 PE-hysteresis loops of PVDF and composite films: (a) PVDF/Ni-P, (b) PVDF/Co-P.

Table 3 Summary of the structural, morphological, and mechanical properties of PVDF base MP-Ps composites

Composite	β-Phase content (%)	Surface morphology (SEM)	Mechanical performance	Filler dispersion	Remarks
Pure PVDF	~42	Globular, porous surface	Low mechanical strength	No filler	Low β-phase, porous, weak mechanical profile
PVDF/3Ni-P	89	Smoother surface, uniform structure	Improved strength & flexibility	Homogeneous	Good β-phase, good matrix integration
PVDF/7Ni-P	91	Dense, smooth, non- porous morphology	High strength and elasticity	Excellent	Best overall performance; high crystallinity
PVDF/3Ag-P	96	Smooth, minor surface aggregates	Moderate mechanical strength	Fairly uniform	High β-phase, moderate interaction
PVDF/7Ag-P	92	Rougher, large pores	Weakened at higher filler	Acceptable	Surface roughness reduces performance
PVDF/3Co-P	81	Dense with minimal voids	Decent performance	Good	Stable morphology with minor modification
PVDF/7Co-P	73	Larger pores, exposed filler particles	Reduced strength	Partial agglomeration	Surface irregularities affect properties
PVDF/3Co-Pn	91	Smooth, embedded particles	Good at lower loading	Well distributed	Slight voids from pull-out noted
PVDF/7Co-Pn	68	Smooth, minor voids	Lower strength due to voids	Moderate	Onset of agglomeration; reduced β-phase

based on the comprehensive performance summary shown in Table 3. Although the PVDF/Ag–P composite exhibited the highest β phase content, its microstructural features, such as increased surface porosity and larger globules observed via SEM indicated a less favourable morphology for stable ferroelectric domain formation. These characteristics may hinder effective dipole alignment and remnant polarization, limiting its practical ferroelectric response. Therefore, it was excluded from *PE*-loop analysis to focus on the most representative and high-performing systems.

For PVDF/Ni–P composites, the remnant polarization ($P_{\rm r}$) doubles from 0.02 μ C cm⁻² in pure PVDF to 0.04 μ C cm⁻² at 3 wt% loading, while the maximum polarization ($P_{\rm max}$) significantly increases from 0.16 μ C cm⁻² to 0.26 μ C cm⁻². This enhancement can be attributed to the nucleation effect of Ni–P particles on β phase formation, which enables more effective dipole alignment under applied electric fields as previously discussed in the XRD and FTIR section. At 7 wt% Ni–P loading, $P_{\rm r}$ maintains its value (0.04 μ C cm⁻²), but $P_{\rm max}$ shows a slight decrease to 0.23 μ C cm⁻², suggesting that while dipole retention remains stable, the overall polarization capability becomes partially hindered at higher filler concentrations.

The PVDF/Co–P system exhibits distinct behaviour compared to its Ni–P. Remnant polarization increases progressively from 0.02 μC cm $^{-2}$ in pure PVDF to 0.05 μC cm $^{-2}$ at 3 wt% and further to 0.06 μC cm $^{-2}$ at 7 wt%, demonstrating superior dipole stabilization with increasing Co–P content. However, while the maximum polarization initially increases from 0.16 μC cm $^{-2}$ in pure PVDF to 0.28 μC cm $^{-2}$ at 3 wt%, it dramatically decreases to 0.10 μC cm $^{-2}$ at 7 wt%, representing a reduction of approximately 64% compared to the 3 wt% loading and falling below even the pure PVDF value. This substantial drop indicates that while Co–P particles effectively enhance remnant polarization through stabilization of polarized domains, they significantly

impede the total achievable polarization at higher concentrations. The higher $P_{\rm r}$ values observed for Co–P composites suggest stronger interfacial interactions that stabilize the polar β phase, consistent with the higher electronegativity of cobalt compared to nickel.⁸⁵ However, the sharp decrease in $P_{\rm max}$ for PVDF/7Co–P indicates that beyond a critical concentration level, these particles begin to act as defect sites that impede complete domain alignment under applied fields.⁸⁶

The enhanced ferroelectric response in these composites originates from two primary mechanisms. First, increased β phase content with its inherently higher dipole moment per unit cell and also enhanced interfacial polarization at the fillerpolymer boundaries. At moderate filler concentrations (3 wt%), these mechanisms operate synergistically, while at higher loadings (7 wt%), particularly for Co-P, competing effects of polarization enhancement and matrix disruption become evident.87 The SEM observations also support this interpretation, where signs of agglomeration and non-uniform dispersion were observed at higher filler loadings, particularly for the Co-P composite. These microstructural discontinuities likely contribute to the observed decrease in maximum polarization by creating local disturbances in the electric field distribution.87 These results indicate that optimal ferroelectric properties for potential applications in energy harvesting and sensing devices are achieved at moderate filler concentrations (approximately 3 wt% for both systems), where enhancement of β phase content and interfacial polarization operate without significantly disrupting the polymer matrix continuity.

When comparing our results with literature reports, the PVDF/Co–P and PVDF/Ni–P composites demonstrate remarkable ferroelectric performance with relatively low filler loadings. As shown in Table 4, our PVDF/3Co–P composite achieves a remnant polarization of 0.05 μ C cm⁻² with only 3 wt% loading and 81% β phase content, while PVDF/7Ni–P reaches a Pr of 0.04 μ C cm⁻²

Paper RSC Advances

Table 4 Comparison of ferroelectric properties in PVDF based composites

Composite system	Preparation method	Filler content (wt%)	β-Phase (%)	$\text{Pr} \left(\mu \text{C cm}^{-2} \right)$	Reference
PVDF/3Co-P (this work)	Solution casting	3	81	0.05	Present study
PVDF/7Ni-P (this work)	Solution casting	7	91	0.04	Present study
PVDF/CoFe ₂ O ₄	Spin coating	15	\sim 80	0.0006	83
(0.8)PVDF/ (0.2) BaTiO ₃	Solution mixing method	20	_	0.045	88
ϕ BNT + $(1 - \phi)$ PVDF	Solution casting	5	$\sim \! 50$	0.025	89
15PZNT	Drop casting technique	15	88.9	0.035	90

with 7 wt% loading and 91% β -phase content. These values are comparable to or exceed those reported for others research groups despite our significantly lower filler concentrations. These results highlight the exceptional efficiency of MP-Ps as fillers for enhancing PVDF's ferroelectric properties at minimal loading levels, demonstrating remarkable potential for next-generation piezoelectric energy harvesting systems where high performance to weight ratio and cost-effectiveness are crucial design parameters.

Conclusion

In this work, we successfully developed novel PVDF/metal phosphate (MP-Ps) composite materials with significantly enhanced ferroelectric properties, representing a substantial advancement in the field of energy harvesting and flexible electronics. The structural, thermal, mechanical, and ferroelectric properties of the composites were systematically investigated, yielding quantitative and novel findings. Flexible composite films with β -phase content improvements ranging from 68% to 96% and superior ferroelectric responses were fabricated using the solution casting method, demonstrating their potential for high-performance applications.

The SEM analysis revealed exceptional dispersion of MP-Ps within the PVDF matrix, a key factor contributing to the mechanical property enhancements. Strong interfacial interactions enabled by this uniform dispersion resulted in tensile strength improvements of up to 181.83% and Young's modulus increases reaching 184.76% (both in PVDF/3Co-Pn). In addition, the incorporation of MP-Ps led to notable improvements in thermal stability, with onset degradation temperature increases of 24.92 °C (PVDF/7Ni-P) and maximum degradation temperature enhancements of 21.47 °C (PVDF/7Ag-P), underscoring the multifunctional potential of these composites.

The impact of MP-Ps on ferroelectric performance was particularly remarkable. PVDF/7Co–P composites showed a 200% increase in remnant polarization (from 0.02 to 0.06 μC cm $^{-2}$), and PVDF/3Co–P exhibited a 75% increase in maximum polarization (from 0.16 to 0.28 μC cm $^{-2}$). These improvements were driven by the synergistic effects of enhanced β -phase content and homogeneous filler dispersion, emphasizing the pivotal role of particle morphology and matrix interaction in tuning the material's functional performance.

Based on the comprehensive analysis across all formulations (Table 2), PVDF/7Ni-P was identified as the most promising system, offering the best balance of structural integrity, thermal

robustness, mechanical reinforcement, and ferroelectric response. Unlike the Co–P composites, which exhibited agglomeration at higher loadings despite strong ferroelectricity, PVDF/7Ni–P maintained consistent property enhancement with excellent dispersion and structural homogeneity, marking it as the most practically viable candidate.

In summary, the incorporation of MP-Ps into the PVDF matrix proves to be an effective strategy for the development of high-performance composite films with quantifiable, synergistic enhancements across structural, thermal, mechanical, and ferroelectric domains. The significant improvements observed in this study up to 96% β-phase content, 184.76% increase in mechanical strength, 24.92 °C rise in thermal stability, and 200% increase in remnant polarization, underline the transformative potential of metal phosphate nanoparticles in designing next-generation functional materials for flexible energy harvesting systems and advanced electronic applications.

Data availability

The data that support the findings of this study are available from the corresponding author, Fatima-Zahra SEMLALI, upon reasonable request.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We would like to thank the Moroccan Ministry of Higher Education, Scientific Research and Innovation and the OCP Foundation who funded this work through the APRD research program.

References

- 1 Q. Zhu, K. Zhang, D. Li, N. Li, J. Xu, D. W. Bahnemann, *et al.*, Polarization-enhanced photocatalytic activity in noncentrosymmetric materials based photocatalysis: A review, *Chem. Eng. J.*, 2021, **426**, 131681, DOI: **10.1016**/j.cej.2021.131681.
- 2 A. Ahmed, A. Singh and S. Arya, High performance Mn doped ZnO loaded rGO electrode and its practical application as an

- electrode for self-charging supercapacitor device, *J. Energy Storage*, 2025, **105**, 114783, DOI: **10.1016/j.est.2024.114783**.
- 3 S. K. Karan, D. Mandal and B. B. Khatua, Self-powered flexible Fe-doped RGO/PVDF nanocomposite: An excellent material for a piezoelectric energy harvester, *Nanoscale.*, 2015, 7(24), 10655–10666.
- 4 S. Douhi, A. Eddiai, S. Das, B. T. P. Madhav, M. Meddad, O. Cherkaoui, *et al.*, Design of a compact super wideband all-textile antenna for radio frequency energy harvesting and wearable devices, *Opt. Quant. Electron.*, 2023, 55(13), 1–23, DOI: 10.1007/s11082-023-05498-x.
- 5 B. Maamer, A. Boughamoura, A. M. R. Fath El-Bab, L. A. Francis and F. Tounsi, A review on design improvements and techniques for mechanical energy harvesting using piezoelectric and electromagnetic schemes, *Energy Convers. Manag.*, 2019, 199, 111973, DOI: 10.1016/j.enconman.2019.111973.
- 6 M. Liu, F. Qian, J. Mi and L. Zuo, Biomechanical energy harvesting for wearable and mobile devices: State-of-the-art and future directions, *Appl. Energy*, 2022, **321**, 119379, DOI: **10.1016/j.apenergy.2022.119379**.
- 7 S. Douhi, S. Labihi, A. Eddiai, S. Lakrit, M. El Achaby and A. J. A. Al-Gburi, Design, characterization, and electromagnetic performance of a flexible wideband RF antenna using composite materials, *J. Sci. Adv. Mater. Devices*, 2025, **10**(1), 100847, DOI: **10.1016**/j.jsamd.2024.100847.
- 8 A. Ahmed, S. Arora, S. Rasgotra, A. Dubey, A. Singh, R. Singh, et al., Synthesis and characterization of low-density polyethylene (LDPE) bubble wrap-derived reduced graphene oxide for triboelectric nanogenerator electrodes, *Mater. Sci. Eng. B*, 2025, **311**, 117828, DOI: **10.1016/j.mseb.2024.117828**.
- 9 S. Labihi, A. Eddiai, M. El Achaby, M. Rguiti and M. Mazroui, Enhancing β-phase and dielectric properties of BCZT leadfree reinforced in PVDF-HFP composite thick films for ecofriendly energy harvesting, *Ceram. Int.*, 2024, 50(21PA), 40814–40822, DOI: 10.1016/j.ceramint.2024.07.363.
- 10 S. Arya, A. Sharma, A. Singh, A. Ahmed, A. Dubey, B. Padha, *et al.*, Review—Energy and Power Requirements for Wearable Sensors, *ECS Sens. Plus*, 2024, 3(2), DOI: 10.1149/2754-2726/ad54d2.
- 11 E. S. Kadir, R. N. Gayen, R. Paul and S. Biswas, Interfacial effects on ferroelectric and dielectric properties of GO reinforced free-standing and flexible PVDF/ZnO composite membranes: Bias dependent impedance spectroscopy, *J. Alloys Compd.*, 2020, 843, 155974, DOI: 10.1016/j.jallcom.2020.155974.
- 12 S. Barrau, A. Ferri, A. Da Costa, J. Defebvin, S. Leroy, R. Desfeux, *et al.*, Nanoscale Investigations of α- And γ-Crystal Phases in PVDF-Based Nanocomposites, *ACS Appl. Mater. Interfaces*, 2018, **10**(15), 13092–13099.
- 13 A. Gradys and P. Sajkiewicz, Determination of the melting enthalpy of β phase of poly(vinylidene fluoride), *e-Polymers*, 2013, (019), 1–14.
- 14 A. Bhiogade, K. Nagamalleswari, P. Mandal and R. V. K. Mangalam, Improved pyroelectric effect in PVDF/

- BaTiO₃ composite flexible films mediated by enhanced β PVDF phase formation, *J. Polym. Res.*, 2023, **30**(8), 1–8, DOI: **10.1007/s10965-023-03669-8**.
- 15 A. A. Tarhini and A. R. Tehrani-Bagha, Graphene-based Polymer Composite Films with Enhanced Mechanical Properties and Ultra-high In-plane Thermal Conductivity, Compos. Sci. Technol., 2019, 184, 107797, DOI: 10.1016/ j.compscitech.2019.107797.
- 16 S. Labihi, N. Chakhchaoui, A. Eddiai, M. El Achaby, M. Meddad, O. Cherkaoui, *et al.*, Enhancement of piezoelectric β-polymorph formation and properties of graphene oxide and PZT-incorporated in PVDF-HFP matrix for energy harvesting applications, *Polym. Compos.*, 2023, 44(4), 2296–2304.
- 17 S. Luo, S. Yu, R. Sun and C. p. Wong, Nano Ag-Deposited BaTiO₃ Hybrid Particles as Fillers for Polymeric Dielectric Composites: Toward High Dielectric Constant and Suppressed Loss, ACS Appl. Mater. Interfaces, 2014, DOI: 10.1021/am404556c.
- 18 J. Diani and K. Gall, Finite Strain 3D Thermoviscoelastic Constitutive Model, *Society*, 2006, 1–10.
- 19 S. Dogra, A. Ahmed, A. Dubey, R. Singh, A. Singh, A. K. Sundramoorthy, *et al.*, Synthesis, characterization, and application of PVDF-PZT composite thin film separator for self-charging supercapacitors, *Surf. Interfaces*, 2024, 51, 104593, DOI: 10.1016/j.surfin.2024.104593.
- 20 S. Baayyad, Y. Esshouba, S. Barhoumi, E. K. Hlil, S. Ez-Zahraoui, F. Z. Semlali, *et al.*, High-density polyethylene composites filled with micro- and nano-particles of nickel ferrite: magnetic, mechanical, and thermal properties, *RSC Adv.*, 2024, 14(26), 18750–18763.
- 21 S. Baayyad, F. Z. Semlali, E. K. Hlil, T. Mahfoud, H. El Moussaoui and M. El Achaby, Developing Ni0.5Zn0.5Fe2O4 ferrite with controlled particle size and morphology through optimized processing conditions of low energy solid state reaction, *RSC Adv.*, 2024, 14(49), 36264–36272.
- 22 C. Bahloul, S. Ez-zahraoui, A. Eddiai and O. Cherkaoui, RSC Advances Ferrite-doped rare-earth nanoparticles for nanocomposites, *RSC Adv.*, 2024, **14**, 38872–38887.
- 23 J. X. Chen, J. W. Li, C. C. Cheng and C. W. Chiu, Piezoelectric Property Enhancement of PZT/Poly(vinylidenefluoride-cotrifluoroethylene) Hybrid Films for Flexible Piezoelectric Energy Harvesters, ACS Omega, 2022, 7(1), 793–803.
- 24 G. Magdy, A. H. Hassanin, I. Kandas and N. Shehata, PVDF nanostructures characterizations and techniques for enhanced piezoelectric response: A review, *Mater. Chem. Phys.*, 2024, 325, 129760, DOI: 10.1016/j.matchemphys.2024.129760.
- 25 H. Zhao and Z. Y. Yuan, Insights into Transition Metal Phosphate Materials for Efficient Electrocatalysis, *ChemCatChem.*, 2020, 12(15), 3797–3810.
- 26 Q. Cheng, X. Zhao, G. Yang, L. Mao, F. Liao, L. Chen, et al., Recent advances of metal phosphates-based electrodes for high-performance metal ion batteries, *Energy Storage Mater.*, 2021, 41, 842–882, DOI: 10.1016/j.ensm.2021.07.017.
- 27 P. Govindrao, N. W. Ghule, A. Haque and M. G. Kalaskar, Journal of Drug Delivery Science and Technology Metal

- nanoparticles synthesis: An overview on methods of preparation, advantages and disadvantages, and applications, *J. Drug Deliv. Sci. Technol.*, 2019, **53**, 101174, DOI: **10.1016/j.iddst.2019.101174**.
- 28 P. Biswas, N. A. Hoque, P. Thakur, M. M. Saikh, S. Roy, F. Khatun, et al., Portable Self-Powered Piezoelectric Nanogenerator and Self-Charging Photo-Power Pack Using in Situ Formed Multifunctional Calcium Phosphate Nanorod-Doped PVDF Films, Langmuir, 2019, 35(52), 17016–17026.
- 29 E. Abdulkarem, Y. Ibrahim, M. Kumar, H. A. Arafat, V. Naddeo, F. Banat, *et al.*, Polyvinylidene fluoride (PVDF)-α-zirconium phosphate (α-ZrP) nanoparticles based mixed matrix membranes for removal of heavy metal ions, *Chemosphere*, 2021, **267**, 128896, DOI: **10.1016**/j.chemosphere.2020.128896.
- 30 T. Pareek, S. Dwivedi, S. A. Ahmad, M. Badole and S. Kumar, Effect of NASICON-type LiSnZr(PO4)3 ceramic filler on the ionic conductivity and electrochemical behavior of PVDF based composite electrolyte, *J. Alloys Compd.*, 2020, 824, 153991, DOI: 10.1016/j.jallcom.2020.153991.
- 31 H. Rekik, Z. Ghallabi, I. Royaud, M. Arous, G. Seytre, G. Boiteux, *et al.*, Dielectric relaxation behaviour in semi-crystalline polyvinylidene fluoride (PVDF)/TiO₂ nanocomposites, *Composites, Part B*, 2013, 45(1), 1199–1206, DOI: 10.1016/j.compositesb.2012.08.002.
- 32 C. F. Holder and R. E. Schaak, Tutorial on Powder X-ray Diffraction for Characterizing Nanoscale Materials, *ACS Nano*, 2019, **13**(7), 7359–7365.
- 33 S. Kullyakool, C. Danvirutai, K. Siriwong and P. Noisong, Determination of kinetic triplet of the synthesized Ni₃(PO 4)₂·8H₂O by non-isothermal and isothermal kinetic methods, *J. Therm. Anal. Calorim.*, 2014, **115**(2), 1497–1507.
- 34 S. S. Patil, J. C. Shin and P. S. Patil, Binder free hydrothermally synthesized nickel phosphate hydrate microplates on nickel foam for supercapacitors, *Ceram. Int.*, 2022, **48**(19), 29484–29492, DOI: **10.1016**/j.ceramint.2022.07.050.
- 35 P. Amornpitoksuk, K. Intarasuwan, S. Suwanboon and J. Baltrusaitis, Effect of phosphate salts (Na₃PO₄, Na₂HPO₄, and NaH₂PO₄) on Ag₃PO₄ morphology for photocatalytic dye degradation under visible light and toxicity of the degraded dye products, *Ind. Eng. Chem. Res.*, 2013, 52(49), 17369–17375.
- 36 Q. Liang, W. Ma, Y. Shi, Z. Li and X. Yang, Hierarchical Ag₃PO₄ porous microcubes with enhanced photocatalytic properties synthesized with the assistance of trisodium citrate, *CrystEngComm.*, 2012, **14**(8), 2966–2973.
- 37 H. Liu, X. Peng, X. Liu, G. Qi and J. Luo, Porous Mn-Doped FeP/Co₃ (PO₄)2 Nanosheets as Efficient Electrocatalysts for Overall Water Splitting in a Wide pH Range, *ChemSusChem*, 2019, **12**(7), 1334–1341.
- 38 J. Theerthagiri, K. Thiagarajan, B. Senthilkumar, Z. Khan, R. A. Senthil, P. Arunachalam, *et al.*, Synthesis of Hierarchical Cobalt Phosphate Nanoflakes and Their Enhanced Electrochemical Performances for

- Supercapacitor Applications, *ChemistrySelect*, 2017, 2(1), 201–210.
- 39 C. Baudot, C. M. Tan and J. C. Kong, FTIR spectroscopy as a tool for nano-material characterization, *Infrared Phys. Technol.*, 2010, 53(6), 434–438, DOI: 10.1016/j.infrared.2010.09.002.
- 40 P. Bhanja, Y. Kim, B. Paul, J. Lin, S. M. Alshehri, T. Ahamad, et al., Facile Synthesis of Nanoporous Transition Metal-Based Phosphates for Oxygen Evolution Reaction, ChemCatChem., 2020, 12(7), 2091–2096.
- 41 F. S. Omar, A. Numan, N. Duraisamy, S. Bashir, K. Ramesh and S. Ramesh, Ultrahigh capacitance of amorphous nickel phosphate for asymmetric supercapacitor applications, *RSC Adv.*, 2016, **6**(80), 76298–76306, DOI: **10.1039/C6RA15111F**.
- 42 I. Ahmed, R. Biswas, R. Sharma, V. Burman and K. K. Haldar, Access to carbon nanofiber composite hydrated cobalt phosphate nanostructure as an efficient catalyst for the hydrogen evolution reaction, *Front. Chem.*, 2023, **11**, 1–10.
- 43 V. P. Padmanabhan, P. Sivashanmugam, R. Kulandaivelu, S. Sagadevan, B. Sridevi, R. Govindasamy, *et al.*, Biosynthesised Silver Nanoparticles Loading onto Biphasic Calcium Phosphate for Antibacterial and Bone Tissue Engineering Applications, *Antibiotics*, 2022, 11(12), DOI: 10.3390/antibiotics11121780.
- 44 A. Attia, Q. Wang, X. Huang and Y. Yang, Titanium phosphates as positive electrode in lithium-ion batteries: Composition, phase purity and electrochemical performance, *J. Solid State Electrochem.*, 2012, **16**(4), 1461–1471.
- 45 K. Kadiya, S. B. Vuggili, U. K. Gaur and M. Sharma, Comparative photocatalytic dye and drug degradation study using efficient visible light-induced silver phosphate nanoparticles, *Environ. Sci. Pollut. Res.*, 2021, **28**(34), 46390–46403.
- 46 J. Li, L. Zhang, P. Yang and X. Cheng, Morphological evolution of Co phosphate and its electrochemical and photocatalytic performance, *CrystEngComm*, 2018, 20(43), 6982–6988.
- 47 L. Kumar and D. Ahuja, Preparation and characterization of aliphatic polyurethane and modified hydroxyapatite composites for bone tissue engineering, *Polym. Bull.*, 2020, 77(11), 6049–6062, DOI: 10.1007/s00289-019-03067-5.
- 48 L. C. Chow, M. Markovic, S. A. Frukhtbeyn and S. Takagi, Hydrolysis of tetracalcium phosphate under a nearconstant-composition condition - Effects of pH and particle size, *Biomaterials*, 2005, 26(4), 393–401.
- 49 Y. Li, Y. Wang, Y. Huang, J. Cao, W. Ho, S. Lee, *et al.*, Controllable synthesis of phosphate-modified BiPO4 nanorods with high photocatalytic activity: Surface hydroxyl groups concentrations effects, *RSC Adv.*, 2015, 5(121), 99712–99721.
- 50 A. Samal, D. P. Das and G. Madras, Repercussion of Solid state *vs.* Liquid state synthesized p–n heterojunction RGO-copper phosphate on proton reduction potential in water, *Sci. Rep.*, 2018, 1–19, DOI: 10.1038/s41598-018-21239-7.

- 51 H. H. Agbe, 2O₂ rejuvenation-mediated synthesis of stable mixed-morphology Ag₃PO₄, *Heliyon*, 2018, e00599, DOI: **10.1016/j.heliyon.2018.e00599**.
- 52 J. Qi, Y. p. Lin, D. Chen, T. Zhou, W. Zhang and R. Cao, Autologous Cobalt Phosphates with Modulated Coordination Sites for Electrocatalytic Water Oxidation Oxygen Evolution Reaction Autologous Cobalt Phosphates with Modulated Coordination Sites for Electrocatalytic Water Oxidation, *Angew. Chem., Int. Ed.*, 2020, DOI: 10.1002/anie.202001737.
- 53 A. Brostrøm, K. I. Kling, K. S. Hougaard and K. Mølhave, Complex Aerosol Characterization by Scanning Electron Microscopy Coupled with Energy Dispersive X-ray Spectroscopy, Sci. Rep., 2020, 10(1), 1–15.
- 54 C. H. Lin, S. K. Parthasarathi, S. Bolloju, M. Abdollahifar, Y. T. Weng and N. L. Wu, Synthesis of Micron-Sized LiNi_{0.8}Co_{0.1}Mn_{0.1}O₂ and Its Application in Bimodal Distributed High Energy Density Li-Ion Battery Cathodes, *Energies*, 2022, 15(21), 1–15.
- 55 J. M. Rami, C. D. Patel, C. M. Patel and M. V. Patel, Thermogravimetric analysis (TGA) of some synthesized metal oxide nanoparticles, *Mater. Today: Proc.*, 2020, 43, 655–659, DOI: 10.1016/j.matpr.2020.12.554.
- 56 X. Li, Y. Xiong, M. Duan, H. Wan, J. Li, C. Zhang, et al., Investigation on the adsorption-interaction mechanism of pb(ii) at surface of silk fibroin protein-derived hybrid nanoflower adsorbent, *Materials*, 2020, 13(5), DOI: 10.3390/ ma13051241.
- 57 M. A. Ashraf, Z. Liu and W. X. Peng, Trisaminomethane-cobalt complex supported on Fe₃O₄ magnetic nanoparticles as an efficient recoverable nanocatalyst for oxidation of sulfides and C–S coupling reactions, *Appl. Organomet. Chem.*, 2020, 34(1), 1–15.
- 58 X. Cai, T. Lei, D. Sun and L. Lin, A critical analysis of the α , β and γ phases in poly(vinylidene fluoride) using FTIR, *RSC Adv.*, 2017, 7(25), 15382–15389.
- 59 C. Dayanand, G. Bhikshamaiah, V. Java Tyagaraju, M. Salagram and A. S. R. Krishna Murthy, Structural investigations of phosphate glasses: A detailed infrared study of the *x*(PbO)-(1 X) P₂O₅ vitreous system, *J. Mater. Sci.*, 1996, **31**(8), 1945–1967.
- 60 A. A. Ribeiro, R. F. C. Marques, A. C. Guastaldi and J. S. De Carvalho Campos, Hydroxyapatite deposition study through polymeric process on commercially pure Ti surfaces modified by laser beam irradiation, *J. Mater. Sci.*, 2009, 44(15), 4056–4061.
- 61 V. Sencadas, S. Lanceros-Méndez, I. Sabater, R. Serra, A. Andrio Balado and J. L. Gómez Ribelles, Relaxation dynamics of poly(vinylidene fluoride) studied by dynamical mechanical measurements and dielectric spectroscopy, *Eur. Phys. J. E*, 2012, 35(5), DOI: 10.1140/epje/i2012-12041-x.
- 62 S. Ez-Zahraoui, F. Z. Semlali, M. Raji, F. Z. Nazih, R. Bouhfid, A. E. K. Qaiss, *et al.*, Synergistic reinforcing effect of fly ash and powdered wood chips on the properties of polypropylene hybrid composites, *J. Mater. Sci.*, 2024, 59(4), 1417–1432, DOI: 10.1007/s10853-023-09299-1.

- 63 W. Ji, H. Deng, C. Guo, C. Sun, X. Guo, F. Chen, *et al.*, The effect of filler morphology on the dielectric performance of polyvinylidene fluoride (PVDF) based composites, *Compos. Part A Appl. Sci. Manuf.*, 2019, **118**, 336–343.
- 64 M. Üniversitesi, S. Ersoy and M. Taşdemir, Zinc oxide (ZnO), magnesium hydroxide [Mg(OH)₂] and calcium carbonate (CaCO₃) filled HDPE polymer composites: Mechanical, thermal and morphological properties, *Marmara J. Pure Appl. Sci.*, 2013, 24(4), 93–104, available from: https://dergipark.org.tr/en/pub/marufbd/issue/17878/187433.
- 65 N. Saadatkhah, A. Carillo Garcia, S. Ackermann, P. Leclerc, M. Latifi, S. Samih, *et al.*, Experimental methods in chemical engineering: Thermogravimetric analysis—TGA, *Can. J. Chem. Eng.*, 2020, 98(1), 34–43.
- 66 S. Shetty, A. Mahendran and S. Anandhan, Development of a new flexible nanogenerator from electrospun nanofabric based on PVDF/talc nanosheet composites, *Soft Matter*, 2020, **16**(24), 5679–5688.
- 67 T. Prabhakaran and J. Hemalatha, Flexible films of β-phase poly(vinylidene fluoride)/ZnFe₂O₄ polymer nanocomposite for magnetoelectric device applications, *Sci. Adv. Mater.*, 2014, $\mathbf{6}(7)$, 1313–1321.
- 68 A. M. Alahzm, M. O. Alejli, D. Ponnamma, Y. Elgawady and M. A. A. Al-Maadeed, Piezoelectric properties of zinc oxide/ iron oxide filled polyvinylidene fluoride nanocomposite fibers, *J. Mater. Sci.:Mater. Electron.*, 2021, 32(11), 14610– 14622, DOI: 10.1007/s10854-021-06020-3.
- 69 A. Issa, M. Al-Maadeed, A. Luyt, D. Ponnamma and M. Hassan, Physico-Mechanical, Dielectric, and Piezoelectric Properties of PVDF Electrospun Mats Containing Silver Nanoparticles, *C*, 2017, 3(4), 30.
- 70 J. Yu, X. Huang, C. Wu and P. Jiang, Permittivity, thermal conductivity and thermal stability of poly(vinylidene fluoride)/graphene nanocomposites, *IEEE Trans. Dielectr. Electr. Insul.*, 2011, **18**(2), 478–484.
- 71 I. H. Kim, D. H. Baik and Y. G. Jeong, Structures, electrical, and dielectric properties of PVDF-based nanocomposite films reinforced with neat multi-walled carbon nanotube, *Macromol. Res.*, 2012, 20(9), 920–927.
- 72 F. Liu, M. R. M. Abed and K. Li, Preparation and characterization of poly(vinylidene fluoride) (PVDF) based ultrafiltration membranes using nano γ -Al₂O₃, *J. Membr. Sci.*, 2011, **366**(1–2), 97–103.
- 73 Q. Wu, L. Li, Y. D. Zhang and W. J. Shui, Absorption and mechanical properties of SiCp/PVDF composites, *Composites, Part B*, 2017, **131**, 1–7, DOI: **10.1016**/j.compositesb.2017.07.074.
- 74 S. y. Fu, X. q. Feng, B. Lauke and Y. w. Mai, Effects of particle size, particle/matrix interface adhesion and particle loading on mechanical properties of particulate polymer composites, *Composites, Part B*, 2008, 39, 933–961, DOI: 10.1016/j.compositesb.2008.01.002.
- 75 P. Cheang and K. A. Khor, Effect of particulate morphology on the tensile behaviour of polymer Á hydroxyapatite composites, *Mater. Sci. Eng. A*, 2003, **345**, 47–54, DOI: **10.1016/S0921-5093(02)00284-8**.

Paper

- 76 Y. Pang, J. Yang, T. E. Curtis, S. Luo, D. Huang, Z. Feng, et al., Exfoliated Graphene Leads to Exceptional Mechanical Properties of Polymer Composite Films, ACS Nano, 2019, 13(2), 1097–1106.
- 77 J. Cho, M. S. Joshi and C. T. Sun, Science And Effect of inclusion size on mechanical properties of polymeric composites with micro and nano particles, *Compos. Sci. Technol.*, 2006, 66, 1941–1952, DOI: 10.1016/ j.compscitech.2005.12.028.
- 78 J. Rexer and E. Anderson, Composites with Planar Reinforcements (Flakes, Ribbonsk)-A Review, *Polym. Eng. Sci.*, 1979, **19**(1), 1–11, DOI: **10.1002/pen.760190102**.
- 79 P. W. Balasuriya, L. Ye and Y. Mai, Mechanical Properties of Wood Ake ± Polyethylene Composites. Part I: Effects of Processing Methods and Matrix Melt Ow Behaviour, *Composites, Part A*, 2006, 32, 2001, DOI: 10.1016/S1359-835X(00)00160-3.
- 80 A. Janićijević, S. Filipović, A. Sknepnek, A. Salević-Jelić, R. Jančić-Heinemann, M. Petrović, et al., Structural, Mechanical, and Barrier Properties of the Polyvinylidene Fluoride-Bacterial Nanocellulose-Based Hybrid Composite, Polymers, 2024, 16(8), DOI: 10.3390/polym16081033.
- 81 H. He, G. Fan, F. Saba, Z. Tan, Z. Su, D. Xiong, *et al.*, Enhanced distribution and mechanical properties of high content nanoparticles reinforced metal matrix composite prepared by flake dispersion, *Composites, Part B*, 2023, 252, 110514, DOI: 10.1016/j.compositesb.2023.110514.
- 82 J. F. Scott, Applications of modern ferroelectrics, *Science*, 2007, **315**(5814), 954–959.
- 83 A. Bhiogade, K. Nagamalleswari, P. Mandal and V. K. M. Ramakrishnan, Flexible multiferroic PVDF/CoFe₂O₄ composite films for pyroelectric energy conversion, *J. Mater. Sci.*, 2023, **58**(47), 17805–17815, DOI: **10.1007/s10853-023-09149-0**.

- 84 S. Dash, R. N. P. Choudhary, A. Kumar and M. N. Goswami, Enhanced dielectric properties and theoretical modeling of PVDF-ceramic composites, *J. Mater. Sci. Mater. Electron.*, 2019, 30(21), 19309–19318, DOI: 10.1007/s10854-019-02291-7.
- 85 T. M. Vandana and R. Gupta, Thermal and mechanical energy harvester based on flexible PVDF/PLZT polymer-ceramic composites, *J. Polym. Res.*, 2023, **30**(8), 1–11, DOI: **10.1007/s10965-023-03694-7**.
- 86 Q. Yang, Z. Shi, D. Ma, Y. He and J. Wang, Flexible PbTiO₃-nanowires/P(VDF-TrFE) composite films and their dielectric, ferroelectric and pyroelectric properties, *Ceram. Int.*, 2018, 44(12), 14850–14856, DOI: 10.1016/j.ceramint.2018.05.118.
- 87 S. Dash, H. Sankar, M. Ravikant and A. Kumar, Ferroelectric ceramic dispersion to enhance the β phase of polymer for improving dielectric and ferroelectric properties of the composites, *Polym. Bull.*, 2021, 78(9), 5317–5336, DOI: 10.1007/s00289-020-03372-4.
- 88 M. Sharma, A. Gaur and J. K. Quamara, Effect of 80 MeV O6+ ion irradiation on structural, morphological, dielectric, and ferroelectric properties of (1–*x*)PVDF/(*x*)BaTiO₃ nanocomposites, *Ionics*, 2020, **26**(1), 471–481.
- 89 L. K. Pradhan, R. Pandey, S. Kumar and M. Kar, Lead free Bi0.5Na0.5TiO3 (BNT) and polyvinylidene fluoride (PVDF) based nanocomposite for energy storage applications, *AIP Conf. Proc.*, 2018, **1953**, 1–5.
- 90 S. Pratihar, A. Patra, A. Sasmal, S. K. Medda and S. Sen, Enhanced dielectric, ferroelectric, energy storage and mechanical energy harvesting performance of ZnO-PVDF composites induced by MWCNTs as an additive third phase, *Soft Matter*, 2021, 17(37), 8483–8495.