

The Structural Analysis of PVDF/WS₂ Polymer Nanocomposite films by TEM and Small-Angle X-Ray Scattering method

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In recent years, there has been growing interest by academics, scientific community, and industry in the development of polymer composites containing organic, inorganic, or hybrid fillers. This interest is primarily due to the potential to enhance thermal, mechanical, and dynamic-mechanical properties through interfacial interactions between the polymer matrix and the filler [1, 2]. The polyvinylidene fluoride (PVDF) polymer matrix is tough, flexible, and easy to process into films, allowing for the creation of lightweight, bendable, and wearable shields. The high surface area of WS₂ nanosheets enhances the formation of a conductive network at low loadings, making the shield efficient and lightweight. Based on the search results and current scientific literature, the application of PVDF+%WS₂ nanocomposite films for radiation protection is an emerging and promising area of research, though not as well established as their use in energy harvesting. These composites show potential primarily for shielding against electromagnetic (EM) radiation and, to a lesser extent, ionizing radiation.

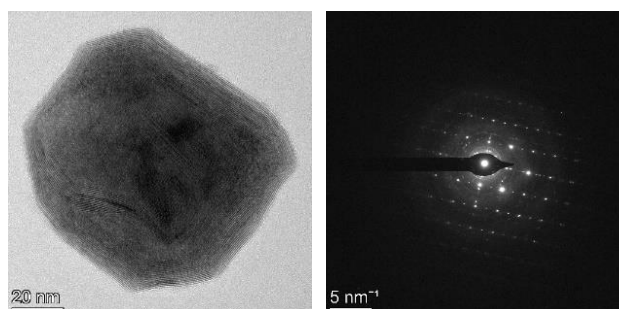


Figure 1. TEM micrograph and electron diffractogram of WS₂ nanosheets

In the current work a polymeric matrix samples of pure PVDF and PVDF/WS₂ composite films irradiated with different gamma-ray doses (100, 300, 500 and 700 kGy) were analyzed using small-angle and wide-angle X-ray scattering. The SAXS/WAXS measurements were carried out at the Frank Laboratory of Neutron Physics, Joint Institute for Nuclear Research (JINR), Dubna, Russia [3]. SAXS experiments showed WS₂ nanosheets mainly distributed in polyvinylidene fluoride matrix as aggregates system [4]. WS₂ disperses well in the range of 0.1-0.5%, and an average distance of ~9-10 nm between particles is observed within the PVDF matrix. As the concentration increases, surface roughness increases (2.05 → 2.30). 0.5% is the optimal concentration: This sample exhibits both an ordered structure (inter-particle distance ~9.2 nm) and favorable surface roughness ($D_s \approx 2.30$) for interfacial activity. The percolation threshold is crossed between 0.5% and 10%. At 10%, the "broad peak" disappears, and a dual-level fractal structure forms instead. This indicates that WS₂ has undergone strong aggregation with ($D_s \approx 2.92$) indicating much roughness than that of pure WS₂ ($D_s \approx 2.68$), it is expected that PVDF chains nucleate on the WS₂ surfaces, creating additional roughness.

1. Nabiye¹ A.A., Olejniczak A., et al. Composite Films of HDPE with SiO₂ and ZrO₂ Nanoparticles: The Structure and Interfacial Effects // *Nanomaterials*. 2021. V. 11. N 10. P. 2673.
2. Nabiye¹ A.A., Mustafayev I.I. et al. Post- γ -irradiation effects in nano-SiO₂ particle reinforced high-density polyethylene composite films: Structure–property relationships, thermal stability and degradation // *Polymer Composites*. 2025. V. 46. P. S44-S62.

3. Nabiyev A.A., Ivankov O.I. et al. High-dose gamma irradiation effects on HDPE/SiO₂ nanocomposite films: Structure, crystallinity, defects, radiation endurance, dispersion, and interfacial behavior // Polymer Degradation and Stability. 2026. V. 245. P. 111851.