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INVESTIGATION OF MECHANICAL AND PHYSICAL PROPERTIES OF GRAPHENE WITH EPOXY MATRIX

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ABSTRACT

Composite materials have become a highly preferred technology nowadays in the industry with their advantages such as superior mechanical performance and weight loss for aerospace applications, especially due to the combination of different components and the formation of new products. In addition, with nanocomposites, this technology skips a step further and allows more effective products to be produced. Nanocomposites are produced using carbon allotropes such as graphene and carbon nanotubes, which have the title of being one of the strongest materials that have been the subject of academic studies in recent years.

Keywords: Graphene, Epoxy Resin, Nanocomposites

1. INTRODUCTION

The world of nanoscience, nanotechnology and nanocomposites has developed in recent years and the importance of this subject has increased with the diversity of automotive, aerospace, electronics, biotechnology, flexible sensors and many other applications. In this context, an important gain has been made with nanocomposites to innovations based on graphene materials.

Graphene, which is the main building block of graphite, nanotube and C60, which makes sp² hybridization in the honeycomb crystal structure, was synthesized and characterized only in 2004. The term "graphene", which is derived from the words "Graphite" and "ene" in English, has found its equivalent in Turkish as graphene. The 2010 Nobel Prize in Physics was awarded to Andre Geim and Konstantin Novoselov "for their groundbreaking experiments on the two-dimensional graphene material".

Graphene is the basic structural unit of carbon materials such as carbon nanotubes, graphite, fullerene and the like [1, 2]. Due to its excellent physical and chemical properties, it has become the research point of materials science and coacervation physics in physics research and practical applications since it was discovered in 2004 [3].

Graphene has a high surface area as well as high adsorption and surface reactions, electron mobility, thermal conductivity and mechanical strength. As the thinnest known material, graphene has high thermal conductivity, quantum hall effect, high Young's modulus (1 TPa), high optical transmittance and high electronic transport, barrier and flame retardant properties.

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With its high surface area compared to any nanomaterial, it increases the interaction between the layers and the polymer material in nanocomposites. It has applications in a variety of fields, including electronics and biomedicine. Graphene-based composite materials attract the attention of materials scientists today. Because structural fabrication is possible, it is expected to lead to new applications. Extensive research and development work has been carried out on graphene-reinforced polymer composites in the last two decades. The addition of graphene as a reinforcing agent in a polymer matrix has resulted in improving the overall performance and properties of such composites. The unique mechanical, thermal, electrical and physicochemical properties of graphene are among the most important reasons why graphene is the subject of much research. In addition to these, it has also been the focus of attention, especially with its structural thinness. Due to these properties, extensive studies have been initiated for the promising graphene technology to be applied in many sectors and fields.

Graphene can be obtained by many different production methods [4]. The main ones of these methods can be listed as micromechanical separation of graphite layers (Exfoliation) [1], chemical vapor deposition method [5], reduction of graphene oxide [4] and epitaxial growth [6].

Potential applications of graphene and its derivatives are mainly focused on specific properties, with the phased production of different graphene materials (graphene oxide (GO), reduced graphene oxide (rGO), functionalized graphene oxide (fGO), functionalized reduced graphene oxide (frGO) and modified graphene (mG)). is directed.

If we talk about the direction that graphene and its derivatives give to the sector in aviation in the world, very light batteries for airplanes at NASA's Langley Research Center (source: Mackey, Paul J. "Graphene Based Ultra-Light Batteries for Aircraft", NASA, https://nari.arc.nasa.gov/node/301), a new solution-free graphene synthesis technique called Holey Graphene (source: Vitug, Eric; (2017), "NASA Langley's Technology Further Enhances Graphene Functions", NASA, https://www.nasa) .gov/langley/business/feature/nasa-langley-s-technology-further-enhances -graphene-functions), high-power capacitors for electric vehicles (source: Kovo, Yael; (2015), "Technology Opportunity: Graphene Composite Materials for Supercapacitor Electrodes", NASA, https://www.nasa.gov/ames partnerships/technology/technology-opportunity-graphene-composite-materials-for supercapacitor-electrodes.) and protective material development for spacecraft.

In this study, a nanomaterial was reinforced into a thermosetting resin curing at 180 °C and conforming to aerospace standards. Mechanical/thermal properties and microstructure were analyzed outside the process.

2. METHODS

2.1. Materials

HexFlow RTM6 (resin transfer molding), an aerospace grade epoxy resin purchased from Hexcel, was used as the matrix material in the experiments. Graphene nanoplatelets with an average thickness of 6-8 nm and an average lateral size of 25 μ m were used. Graphene nanoplatelets (GNP) powder used as reinforcement was provided by Nanografi Co. Inc. and used as received. Three different types of nanomaterial were used. The difference is in the surface area (150, 500 and 800 m²/g). LOCTITE brand 700-NC was used as a mold release agent.

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2.1. Procedure

GNP reinforcement content was mixed in epoxy resin with the dispersion process, which is one of the nanocomposite production methods, which has been recently studied in the literature (Figure 1). In this method, three roll mill system is used. By making use of centrifugal force, the rollers rotate clockwise and counterclockwise, passing the resin through 3 cylinders and making it homogeneous. During the experimental studies, wt% 0.25, 0.50, 1, 2 and 4 GNP nanomaterials were added to the resin and subjected to the three roll mill process.





After the dispersion process is completed, the resin mixture is placed in a vacuum at 80 °C for 30 minutes for degassing. It is then cured at 180 °C for 2 hours.

The morphologies of the obtained samples were examined with a Quanta 400 F Field Emission scanning electron microscope. Before analysis, the surfaces of the samples were coated with gold/palladium.

Three-point bending (3PB) tests were carried out to determine the mechanical properties of the samples. The three-point bending test was performed using the Shimadzu Universal Tester according to the ISO 178 standard.

Thermal gravimetric analyzes were carried out to determine the thermal properties of the composites. Thermal degradation behavior was investigated with TA Instruments Thermal Gravimetric Analyzer with a heating rate of 10 C^o/min under nitrogen atmosphere. Decomposition temperatures at maximum weight loss were determined from the first derivative of the weight loss - temperature curves.

3. RESULTS & DISCUSSION

Scanning electron microscopy (SEM), thermogravimetric analysis (TGA), and three point bending tests were performed on the samples obtained as a result of experimental studies.

The temperatures at 5% weight loss in TGA analysis, the temperatures at maximum weight loss obtained from the first derivative of the weight loss versus temperature curve curves, and the amount of ash remaining after complete degradation of the material, including residual carbon at 900 °C, of epoxy-graphene based composites are given in Table 1.

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Nanomaterials	Td%5 [°C]	Tdmax [°C]	Char [%]
Epoxy	351	404	10.79
%0.25-G150	343	400	12.83
%0.5-G150	341	388	13.77
%1-G150	340	390	10.42
%2-G150	341	390	14.32
%4-G150	338	393	12.72
%0.25-G500	337	395	12.16
%0.5-G500	338	397	11.18
%1-G500	343	399	12.23
%2-G500	339	397	15.78
%4-G500	350	397	15.09
%0.25-G800	338	389	11.32
%0.5-G800	340	397	11.15
%1-G800	341	388	12.51
%2-G800	340	397	13.02
%4-G800	340	394	15.03

 Table 1. Temperatures at 5% weight loss, temperatures at maximum weight loss and char yield of epoxy-graphene composites.

According to Table 1, the addition of graphene in epoxy matrix decreased the temperature at 5 wt. % loss of the epoxy. This can be due to the high thermal conductivity of graphene particles than the epoxy matrix which may act as heat dissipating source during the earlier steps of the thermal degradation. The temperature at maximum weight loss of the epoxy-graphene composites is also lower than the neat epoxy. Addition of a carbon based material in the matrix may accelerate the thermal degradation of the epoxy composites. Wang et al. and Song et al. also found lowered thermal stability with the addition of graphene into epoxy matrix and explained this with the presence of thermally unstable chemicals, which on decomposition, lowered T_d as compared to monolithic epoxy. Generally, higher percent addition of nanocomposites increased the char yield of the epoxy. This can be due to the high surface area of the graphene nanocomposites which enhanced the catalytic effects of the charring reaction. In general, this trend is observed regardless of the surface area of the nanomaterial.

The bending strength of the samples was measured with a three point bending test according to ASTM D790 and ISO 178 tests for each doping ratio. Considering the results, the flexural strength of the pure epoxy resin (0% GNP by volume) was measured at an average of 117 mPa. It was also observed that the mechanical properties were related to the surface area of the GNPs according to their volume percent doping ratio. As the surface area increased, a decrease in flexural strength was observed at the same doping ratio in the sample. This jump in flexural strength is indicated in Figure 2. The reason for this interaction is the increased surface area and the possibility of agglomeration, which negatively affects the mechanical properties.

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Figure 2. Mechanical test results and standard deviation

The dispersion of GNPs in epoxy after grinding was used SEM for nanocomposite suspension. Distribution is the most important factor to consider. When replacing a polymer matrix, the agglomeration of the nanofill can affect the stress concentration. SEM images of the surfaces of epoxy nanocomposites as taken are shown (Figure 3).



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Figure 3. (a) Neat epoxy and, respectively, agglomerations in epoxy resin doped with 1% pure GNP (b) 150, (c) 500 and (d) 800.

Moreover, SEM analysis found that the nanoparticles were homogeneously dispersed in the epoxy resin and in a small fraction of the wrinkled GNPs, as seen in Figure 3. These findings show that it is possible to detect agglomerations formed at a given doping rate. It has been observed that the agglomeration problem has decreased significantly, especially at 1% and below.

4. CONCLUSION

The aim of this study is to improve the physical and mechanical properties of polymers with nanomaterial reinforcement. In the experimental studies, various combinations of GNP and Epoxy resin systems were studied at different doping rates. Approximately 20 samples were produced to reach the optimum result. It was observed that the yields of the samples with low doping ratio were higher. At the same time, their mechanical performance and thermomechanical results are quite good compared to polymers with high doping ratios. Considering the results obtained from the study, it is aimed that the structural parts will be nanodoped composite, that is, nanocomposite, by combining the resin and GNP mixture fiber, which will give the most optimum result in the future, in accordance with aviation standards.

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CONFLICT OF INTEREST

The authors stated that there are no conflicts of interest regarding the publication of this article.

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