

## Effect of graphene nanoplatelets on flame retardancy and corrosion resistance of epoxy nanocomposite coating

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### Abstract

Various concentrations of graphene nanoplatelets (GNP) i.e. 0.2, 0.4, 0.6, 0.8 and 1.0 wt. % were incorporated into the epoxy resin by sonication technique and mechanical agitation process. Limiting oxygen index (LOI) test and thermogravimetric analysis (TGA) indicated that the presence of GNP greatly enhanced the flame retardancy properties of the epoxy coating. Salt spray results obtained suggested that the addition of GNP enhanced corrosion performance and reducing the water absorption in comparison with pristine epoxy coating. Adhesion (cross-cut test) revealed that the presence of GNP showed great adhesion to substrates. Incorporation of 0.8 wt. % GNP exhibited the superior anticorrosion performance, great adhesion to subtract, and the lowest water uptake among other samples.

**Keywords:** Epoxy, nanocomposites, coatings, graphene nanoplatelets

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## INTRODUCTION

Epoxy coatings have been developed over the years due to their good barrier properties that can protect metals from corrosion over long periods of time (Pour-Ali *et al.*, 2014; Atta *et al.*, 2015). However, pristine epoxy coatings have low durability as anticorrosion coating due to the presence of holes and defects which are permeable to oxygen, water, and corrosive ions such as Cl<sup>-</sup>, H<sup>+</sup>, and SO<sub>2</sub><sup>-4</sup> (Popović *et al.*, 2005). Therefore, numerous types of additives have been employed to improve the barrier performance of these polymeric coatings (Zhang *et al.*, 2007; Shao *et al.*, 2009; Chen *et al.*, 2017). The incorporation of fine particles that are miscible into epoxy resin can occupy cavities by reducing the porosity and zigzagging the diffusion path for deleterious species and consequently enhance the integrity and durability of coatings (Shi *et al.*, 2003; Hartwig *et al.*, 2005).

Graphene related material have attracted tremendous attention in various potential applications including batteries (Raccichini *et al.*, 2015), capacitors (Xia *et al.*, 2015), nanocomposites (Mittal *et al.*, 2015), sensors (Vashist *et al.*, 2015), and coatings for corrosion prevention (Chen *et al.*, 2017; Prasai *et al.*, 2012). These applications are possible due to its unique structures and extraordinary electronic, thermal, mechanical, and barrier properties. Previous studies have indicated that graphene-based coatings possess excellent anticorrosion barriers for metals (Kirkland *et al.*, 2012; Chang *et al.*, 2012; Zhou *et al.*, 2013; Yu *et al.*, 2014). Kirkland *et al.* (2012) reported that the electrochemical response of graphene-coated nickel and copper, where the graphene layer was able to improve the anticorrosion performance of pure metals. The capability of graphene to enhance

oxidation resistance was demonstrated by Chen *et al.* (2011). It helps to create an air-tight “balloon” that presumably increase barrier properties and impermeability to oxygen (Bunch *et al.*, 2008).

To the best of our knowledge, not many studies investigate the effect of graphene on corrosion performance and flame retardancy of epoxy nanocomposite coating. One related study by Alhumade *et al.* (2016) revealed the effect of different graphene nanosheets (GNS) loading on corrosion performance of stainless-steel Type 304. They found that low graphene loading up to 0.5 wt. % boost up the life cycle of stainless-steel substrates due to the barrier property of graphene, which limits the diffusion of corrosive agents to reach the metal substrates. Nevertheless, the effect of GNS on flame retardancy was not discussed and highlighted in their study.

In this work, epoxy resin type BE-188 and GNP nanofiller were prepared by stirring and bath sonication before coated on the mild steel plate. The anticorrosion performance and flame retardancy properties of epoxy filled GNP nanocomposite coatings were evaluated by salt spray and LOI test, respectively.

## EXPERIMENTAL

### Materials

The mild steel plate which was readily sandblasted was cut into dimensions of 100 mm × 50 mm × 1.5. For salt spray and adhesion tape test, they were coated with epoxy-filled graphene nanocomposite coating. The diglycidyl ether of bisphenol A (DGEBA), epoxy resin (BE-188), and polyamine hardener (H2310 polyamine amide) were supplied by Mc-Growth Chem. GNP was purchased from XG Science

Inc., East Lansing, USA with the specification of 6-10 layer graphene, 3.4-7.0 nm and 50  $\mu\text{m}$  in thickness and diameter, respectively.

### Preparation of nanocomposites

GNP filler was dispersed in ethanol (1:250 ratio) and subsequently sonicated using a sonicator ultrasonic Bilon-1200Y for 15 minutes. The suspension was then kept in the fume hood to evaporate the ethanol in order to reduce the formation of an air bubble on the coating surface. After that, the dispersion was added gradually into the epoxy resin and slowly stirred at  $\sim 200$  rpm by a mechanical stirrer for 15 minutes. Finally, a stoichiometric amount of hardener with a weight ratio of 2:1 was added into the mixture and continuously stirred until the mixture was uniformly mixed. Various formulations of the epoxy-graphene nanocomposite are tabulated in Table 1. Well-dispersed epoxy/nanoparticles filler mixture was then coated onto mild steel plates to form  $\sim 0.25$  mm thickness of coating samples using a doctor blade film applicator.

**Table 1** Formulation of the EP-GNP nanocomposite coatings and their corresponding LOI values.

Sample	Composition (wt. %)			LOI (%)
	BE188	Hardener	GNP	
EG0	66.7	33.3	0.0	21.0
EG1	66.7	33.3	0.2	22.0
EG2	66.7	33.3	0.4	22.5
EG3	66.7	33.3	0.6	22.5
EG4	66.7	33.3	0.8	23.0
EG5	66.7	33.3	1.0	24.0

### Measurements

Flame retardant properties of epoxy nanocomposites coatings were measured by limiting oxygen index (LOI) tests. LOI was measured according to ASTM D 2863 using Dynisco Instruments, USA). The specimens used for the test were of dimensions  $80 \times 10 \times 3$  mm<sup>2</sup>.

The thermal gravimetric analysis (TGA) measurement was conducted by the aid of a Perkin Elmer TGA/DSC at a thermal rate of 20 °C/min in the range of 20 - 500 °C. About 4-10 mg of the samples were heated from room temperature to 900 °C at heating rate 10 °C/min under air temperature.

Corrosion study was observed by the salt spray test on scratched samples for 300 and 500 h exposure to an accelerated environment of salt fog chamber according to ASTM B117.

The interfacial adhesion between the coating and mild steel substrate plate was measured according to the ASTM-D3359 standard using an adhesion tape kit (PA-2000, Paul N. Gardner Company Inc., Florida, USA) with a standard blade (PA-2253 thick coatings).

The water absorption test was conducted according to the specifications of ASTM D570-98. The water uptake ( $M_t$ ) of the specimens was calculated by using the following formula:

$$M_t = \frac{W_t - W_0}{W_0} \times 100\% \quad (1)$$

$M_t$ ,  $W_t$  and  $W_0$  are the moisture content at a given time, weight of the sample at the time of the measurement and initial weight, respectively. Fick's equation (Eq. 2) was used to calculate the coefficient of diffusion,

$$\frac{\Delta M_t}{M_o} = \frac{\Delta M_\infty}{M_o} \sqrt{\frac{16D}{\pi}} \frac{\sqrt{t}}{h} \quad (2)$$

where  $\Delta M_t$  (g) is the amount of water absorbed by the film at time  $t$  (s),  $M_o$ (g) is the initial film mass,  $\Delta M_\infty$ (g) is the amount of water absorbed at equilibrium,  $D$  is the diffusion coefficient, and  $h$  is the film thickness.

The distribution of nanoparticles fillers in the epoxy matrix was examined with an HRTEM 120 KV Hi-Tech Instruments.

## RESULTS AND DISCUSSION

### Limiting oxygen index

The Limiting Oxygen Index (LOI) values of EP-GNP nanocomposite coatings are given in Table 1. Neat epoxy coating exhibited the LOI value of 21.0% and there was a gradual increase for EP-GNP samples which considered unlikely to support burning in atmospheric air (Bajaj, 1992). Epoxy resin has a tendency to spread flame away from a fire source. Liu *et al.* (2014) reported that the incorporation of very low content (0.5 wt. % GNS) can effectively decrease the melt flow and inhibit the flammable drips of epoxy resin during combustion as well as can change the path of thermal degradation of epoxy resin. The present study indicated that the flame retardancy increased with the increase of GNP content. The LOI value reached 24.0 (increase by 14 % relative to neat epoxy coating) with the incorporation of 1.0 wt. % of GNP indicating that their flammability was remarkably reduced due to the addition of GNP. This similar phenomenon was also observed in the previous report on PET/PP/GNP nanocomposite (Inuwa *et al.*, 2014). The char formations from the LOI test are depicted in Fig. 1. It is apparent that the additional GNP produced more char when compared to neat epoxy coating.

### Thermal gravimetric analysis (TGA)

The thermal stability of different EP-GNP nanocomposite coatings was investigated by TGA curve (Fig. 2a). It can be seen that all samples have a similar one-stage weight loss, indicating that they have a similar thermal degradation mechanism. The curves of all samples were similar at an initial temperature between 100 and 350 °C with weight loss less than 14 % at 320 °C. It can be seen that the inclusion of GNP has slightly shifted the degradation stage to a low temperature and induced the more stable char layer at high temperature. The residual weight of neat epoxy, GNP 0.2, 0.6, and 1.0 wt. % at 800 °C were 8.1, 13.6, 13.8, and 14.0 wt. %, respectively. The higher residual weight indicated that the incorporation of GNP induces an important thermal stabilization, resulting in good fire protection performance. This enhancement probably ascribed to the so-called "tortuous" path effect, which limits the entry of oxygen thus delaying the escape of volatile degradation products and also char formation (Prolongo *et al.*, 2013, Wang *et al.*, 2012, Kuilla *et al.*, 2010). As illustrated in DTG curves (Fig. 2b), the thermal degradation of neat epoxy and EP-GNP nanocomposite coating are characterized by a single stage, indicating the addition of GNP has no influence on decomposition process.

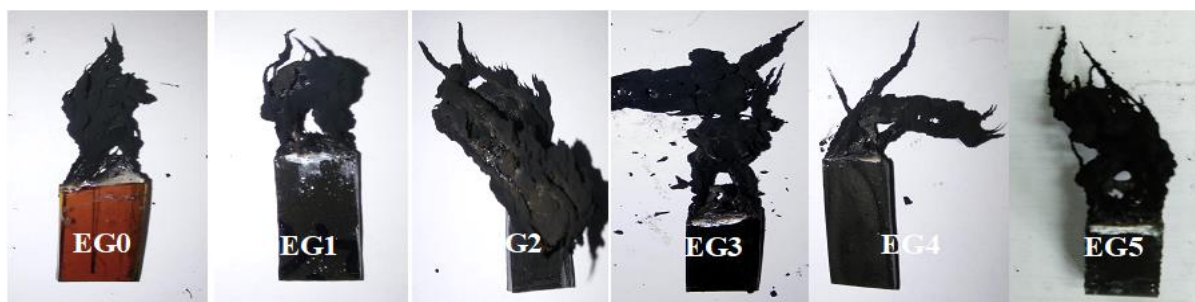


Fig. 1 Char residue of neat and EP-GNP samples after the LOI test.

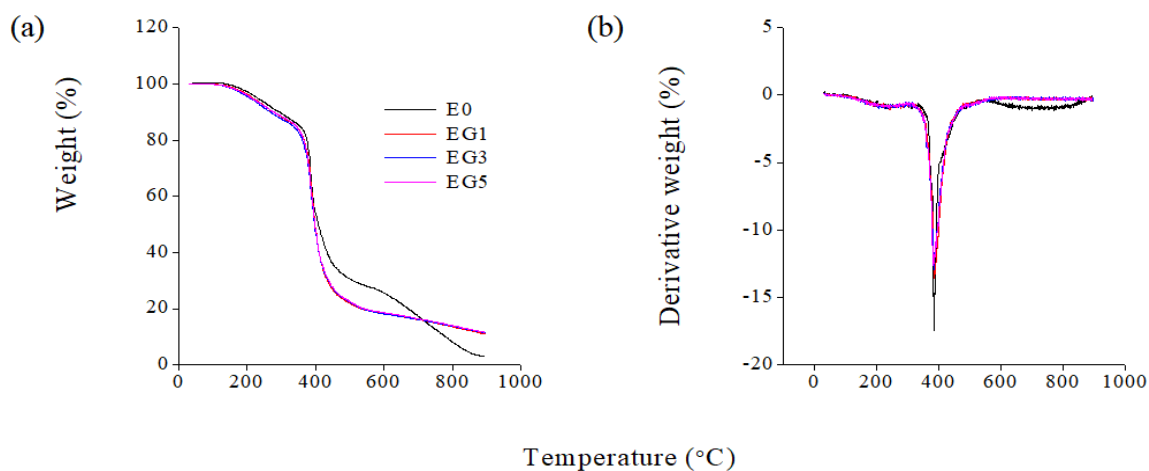


Fig. 2 (a) TGA and (b) DTG thermogram of neat and EP-GNP samples.

### Salt spray test

The salt spray test was conducted as an accelerated corrosion testing method by evaluating the rust, blister, cracking, and delamination along the coating's surface on the mild steel substrate. Fig. 3 shows the appearance of the surfaces of the EP-GNP nanocomposite coatings samples after 200 and 500 hours of salt spray for different GNP loading. A mild steel plate coated with neat epoxy system underwent severe corrosion compared to EP-GNP nanocomposite coating. Large blisters appeared nearby the scratches and over the whole coated surface in the neat epoxy coating sample with a brown adherent corrosion product (Fig. 3 (a)). Some rusting and blisters were observed on EG1, EG2, and EG3 coating surface after 200 hours of exposure, while, no blister was found in EG4 and EG5 samples. A previous study by Liu et al. (2016) revealed that superior anticorrosive performance was exhibited by coating with 0.5 wt. % graphene. They concluded that the well-dispersed graphene blocks the coating pores and reduce the electrolyte diffusion toward the substrate, thus reduces the corrosion activity after exposure to salt spray.

Serious rusting appeared along the surface after 500 hours of exposure to salt spray test for EG0, EG1, and EG2 coatings with a diameter of rusts almost 2.0-2.5 mm (Fig. 3 (g), (h) and (i)). The yellow-brown rust and blistering appeared near the scratches and the bottom of the samples due to the poorest protection of the cut edges of the sample. Few rusts could be observed for EG3 and EG4 coatings (Fig. 3 (j) and (k)). Although the corrosion process has started to take effect on this topcoat as well, the blisters formed in this sample were very small (diameter between 1.0-1.5 mm). At 1.0 wt. % EP-GNP content, anticorrosion performance started to decrease (Fig. 3 (l)). Graphene tends to agglomerate in the form of layers, by layer stacking due to the van der Waals interactions (Terrones et al., 2011) which then limit the barrier effect of nanofiller.

### Adhesion test

Table 4 is tabulated to represent the adhesion test data for mild steel plate coated with neat and EP-GNP nanocomposite coatings after exposure to 5.0 wt. % NaCl solution. As expected, incorporation of

GNP improved the adhesion (described as 5B) before immersion. The adhesion of neat epoxy coating showed 5B adhesion strength and declined to 0B after 200 h of exposure. The result showed that EG1 and EG2 samples were reduced in adhesion from 5B to 1B, while EG3 and EG5 experienced a reduction to 2B. This might happened due to the penetration of electrolyte through coating pores and reach the metal surface which results in adhesion failure and initiating the electrochemical process on a metal surface (Hosseini et al., 2009; Zhou, 2011). In the present study, EG4 (0.8 wt. % GNP) demonstrated the best adhesion properties with about 5 to 15 % surface affected area of the lattice (adhesion at scale No.3B). Increasing graphene content in the epoxy matrix beyond an optimum loading (0.5 wt. %) resulted in the loss of interfacial adhesion bond (Alhumade et al., 2016) and the degree of dispersion was proportional to the graphene loading. Graphene is easily to aggregate together due to their strong van der Waals forces and high specific surface areas, which makes it difficult to achieve a homogeneous dispersion of graphene in a polymer matrix (Gu et al., 2015).

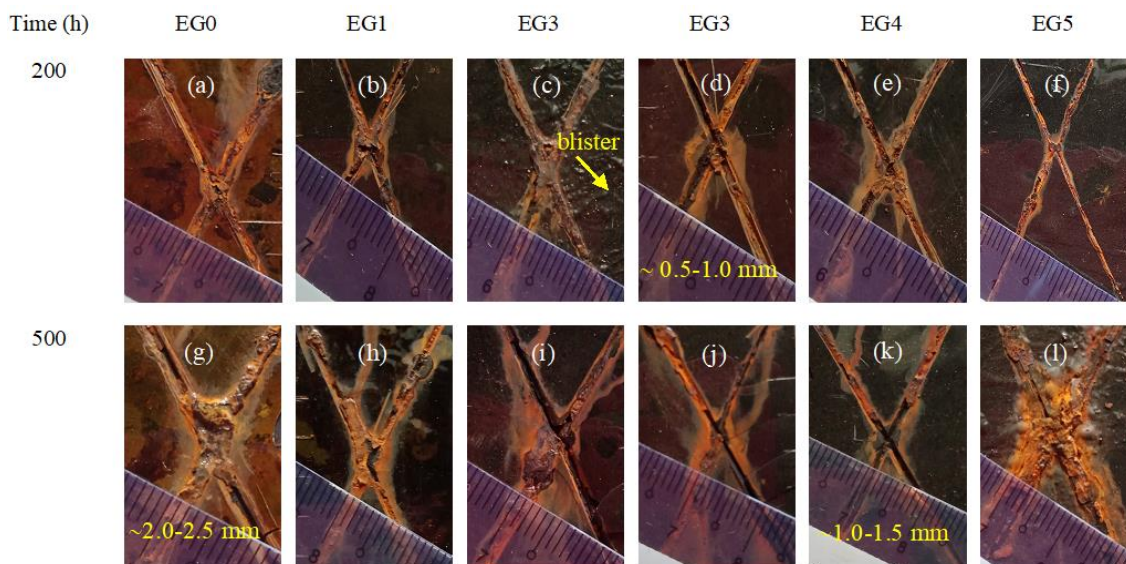
Table 2 Adhesion test result for neat and EP-GNP nanocomposite coatings after 200 h exposure to 5.0 wt. % NaCl solution according to ASTM D3359.

Samples	Adhesion strength	
	Before exposure	After 200 h exposure
EG0	5B	0B
EG1	5B	1B
EG2	5B	1B
EG3	5B	2B
EG4	5B	3B
EG5	5B	2B

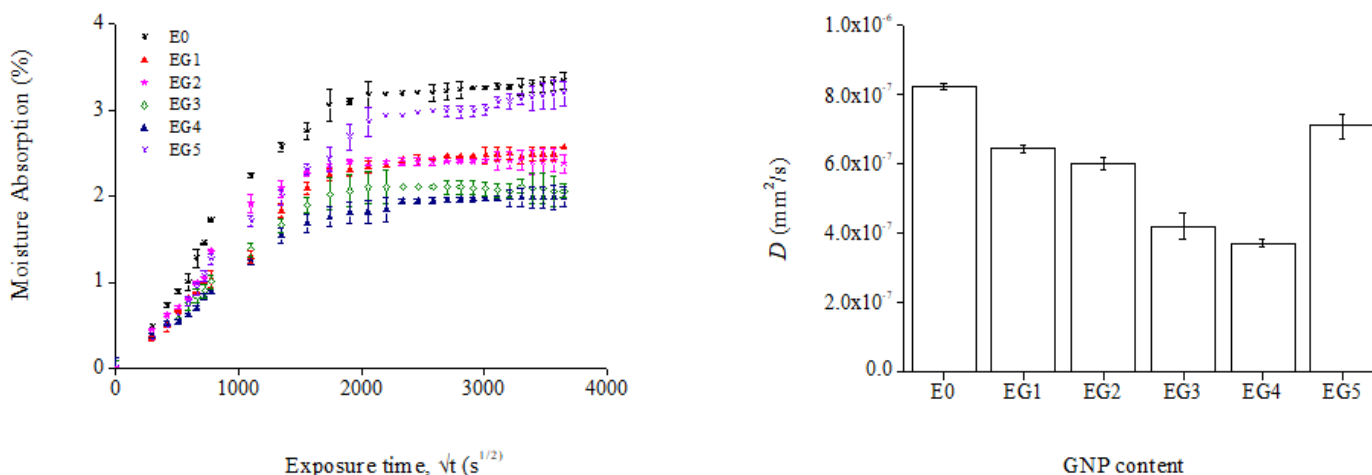
**Water uptake test**

Corrosion performance of coating material is strongly affected by the water uptake which usually diminishes the adhesion of the coating and causes the delamination. Results indicated that the maximum water content absorbed of an EG0 coating was clearly higher compared to the other samples. This might be corresponded to the water molecules interact with polar groups (hydroxyl, amino) of epoxy network and the availability of microvoids (Nogueira et al., 2001) (Fig. 4a). Fig. 4b demonstrated that the diffusion coefficient of

water (*D*) after 154 days of immersion was decreased by incorporation of GNP due to the hydrophobic behavior of EP-GNP that denoted a higher chemical strength in humid environmental. In addition, the *D* value increased abruptly from about  $3.723 \times 10^{-8}$  to  $7.098 \times 10^{-7}$  mm<sup>2</sup>/s for EG4 and EG5, respectively which might be associated with the poor dispersion and/or agglomeration of nanofillers in EG5 sample. This was in accordance with the result found in adhesion tape test.



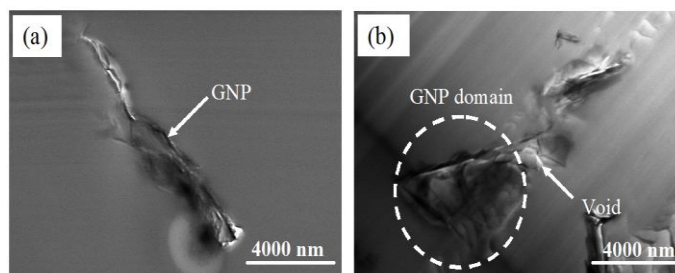
**Fig. 3** Surfaces appearance of mild steel plate coated with neat and EP-GNP coatings at 200 and 500 hours of salt spray test.



**Fig. 4** (a) Water uptake (b) Diffusion coefficient of water (*D*) for neat and EP-GNP coatings in water immersion at room temperature.

**Transmission electron microscopy (TEM)**

The dispersion of GNP fillers in epoxy nanocomposite coatings at the lowest and highest content is as shown in Fig. 5a and 5b. The presence of voids in the stacked nanoplatelets structures and large aggregate domains are clearly seen at higher GNP content (Fig 5b). Agglomerated GNP tend to form obstacles which hindered the flow of polymer into the agglomerates, hence resulting in the formation of holes and voids between GNP and epoxy (King et al., 2008; Zhang et al., 2013; Yue et al., 2014).



**Fig. 5** TEM images of EP-GNP nanocomposite coatings: (a) 0.2 wt. % GNP (b) 1.0 wt. % GNP.

## CONCLUSION

The present study indicated that the incorporation of GNP nanoparticles filler enhanced the flame retardancy and expanded the durability of anticorrosion performance. LOI results showed that GNP enhanced the flammability properties and improved the char yield at 800 °C as well as decreasing the maximum weight loss rate, suggesting improved thermal stability. Results obtained from salt spray test revealed that additional GNP provided corrosion resistance against electrolyte diffusion with the greatest increase in corrosion resistance was observed for 0.8 wt. % GNP content. The water uptake and coefficient diffusion confirmed that GNP might act as reservoirs for the storage and prolonged lease of corrosion inhibitor.

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## REFERENCES

- S. Pour-Ali, C. Dehghanian, A. Kosari. 2014. In situ synthesis of polyaniline–camphorsulfonate particles in an epoxy matrix for corrosion protection of mild steel in NaCl solution. *Corrosion Science*, 85, 204-214.
- A. M. Atta, A. M. El-Saeed, G. M. El-Mahdy, H. A. Al-Lohedan. 2015. Application of magnetite nano-hybrid epoxy as protective marine coatings for steel. *RSC Advances*, 5, 101923-101931.
- M. Popović, B. Grgur, V. Mišković–Stanković. 2005. Corrosion studies on electrochemically deposited PANI and PANI/epoxy coatings on mild steel in acid sulfate solution. *Progress in Organic Coatings*, 52, 359-365.
- X. Zhang, F. Wang, Y. Du. 2007. Effect of nano-sized titanium powder addition on corrosion performance of epoxy coatings. *Surface and Coatings Technology*, 201, 7241-7245.
- Y. Shao, C. Jia, G. Meng, T. Zhang, F. Wang. 2009. The role of a zinc phosphate pigment in the corrosion of scratched epoxy-coated steel. *Corrosion Science*, 51, 371-379.
- C. Chen, S. Qiu, M. Cui, S. Qin, G. Yan, H. Zhao, L. Wang, Q. Xue. 2017. Achieving high performance corrosion and wear resistant epoxy coatings via incorporation of noncovalent functionalized graphene. *Carbon*, 114, 356-366.
- G. Shi, M. Q. Zhang, M. Z. Rong, B. Wetzell, K. Friedrich. Friction and wear of low nanometer Si 3 N 4 filled epoxy composites. 2003. *Wear*, 254, 784-796.
- A. Hartwig, M. Sebald, D. Pütz, L. Aberle. 2005. Preparation, characterisation and properties of nanocomposites based on epoxy resins. *Macromolecular Symposia*, 221, 127-135.
- R. Raccichini, A. Varzi, S. Passerini, B. Scrosati. 2015. The role of graphene for electrochemical energy storage. *Nature materials*, 14, 271-279.
- H. Xia, C. Hong, B. Li, B. Zhao, Z. Lin, M. Zheng, S.V. Savilov, S. M. Aldoshin. 2015. Facile synthesis of hematite quantum-dot/functionalized graphene-sheet composites as advanced anode materials for asymmetric supercapacitors. *Advanced Functional Materials*, 25, 627-635.
- G. Mittal, V. Dhand, K. Y. Rhee, S.-J. Park, W. R. Lee. 2015. A review on carbon nanotubes and graphene as fillers in reinforced polymer nanocomposites. *Journal of Industrial and Engineering Chemistry*, 21, 11-25.
- S. K. Vashist, J. H. T. Luong. 2015. Recent advances in electrochemical biosensing schemes using graphene and graphene-based nanocomposites. *Carbon*, 84, 519-550.
- D. Prasai, J. C. Tuberquia, R. R. Harl, G.K. Jennings, K. I. Bolotin. 2012. Graphene: corrosion-inhibiting coating. *ACS Nano*, 6, 1102-1108.
- N. T. Kirkland, T. Schiller, N. Medhekar, N. Birbilis. Exploring graphene as a corrosion protection barrier. *Corrosion Science*, 56 (2012) 1-4.
- C.-H. Chang, T.-C. Huang, C.-W. Peng, T.-C. Yeh, H.-I. Lu, W.-I. Hung, C.-J. Weng, T.-I. Yang, J.-M. Yeh. 2012. Novel anticorrosion coatings prepared from polyaniline/graphene composites. *Carbon*, 50, 5044-5051.
- F. Zhou, Z. Li, G.J. Shenoy, L. Li, H. Liu. 2013. Enhanced room-temperature corrosion of copper in the presence of graphene. *ACS Nano*, 7, 6939-6947.
- Y.-H. Yu, Y.-Y. Lin, C.-H. Lin, C.-C. Chan, Y.-C. Huang. 2014. High-performance polystyrene/graphene-based nanocomposites with excellent anti-corrosion properties. *Polymer Chemistry*, 5, 535-550.
- S. Chen, L. Brown, M. Levendorf, W. Cai, S.-Y. Ju, J. Edgeworth, X. Li, C.W. Magnuson, A. Velamakanni, R. D. Piner. 2011. Oxidation resistance of graphene-coated Cu and Cu/Ni alloy. *ACS Nano*, 5, 1321-1327.
- J. S. Bunch, S. S. Verbridge, J. S. Alden, A. M. Van Der Zande, J. M. Parpia, H. G. Craighead, P. L. McEuen. 2008. Impermeable atomic membranes from graphene sheets. *Nano Letters*, 8, 2458-2462.
- H. Alhumade, A. Yu, A. Elkamel, L. Simon, A. Abdala. 2016. Enhanced protective properties and UV stability of epoxy/graphene nanocomposite coating on stainless steel. *Express Polymer Letters*, 10(12), 1034-1046.
- P. Bajaj. 1992. Fire-retardant materials. *Bulletin of Materials Science*, 15, 67-76.
- S. Liu, H. Yan, Z. Fang, H. Wang. 2014. Effect of graphene nanosheets on morphology, thermal stability and flame retardancy of epoxy resin. *Composites Science and Technology*, 90, 40-47.
- I. M. Inuwa, A. Hassan, D.-Y. Wang, S. A. Samsudin, M. K. Mohamad Haafiz, S. L. Wong, M. Jawaid. 2014. Influence of exfoliated graphite nanoplatelets on the flammability and thermal properties of polyethylene terephthalate/polypropylene nanocomposites. *Polymer Degradation and Stability*, 110, 137-148.
- S. G. Prolongo, A. Jimenez-Suarez, R. Moriche, A. Ureña. 2013. In situ processing of epoxy composites reinforced with graphene nanoplatelets. *Composites Science and Technology*, 86, 185-191.
- X. Wang, W. Xing, P. Zhang, L. Song, H. Yang, Y. Hu. 2012. Covalent functionalization of graphene with organosilane and its use as a reinforcement in epoxy composites. *Composites Science and Technology*, 72, 737-743.
- T. Kuilla, S. Bhadra, D. Yao, N.H. Kim, S. Bose, J. H. Lee. 2010. Recent advances in graphene-based polymer composites. *Progress in Polymer Science*, 35, 1350-1375.
- S. Liu, L. Gu, H. Zhao, J. Chen, H. Yu. 2016. Corrosion Resistance of Graphene-reinforced waterborne epoxy coatings. *Journal of Materials Science & Technology*, 32, 425-431.
- M. Terrones, O. Martín, M. González, J. Pozuelo, B. Serrano, J. C. Cabanelas, S. M. Vega-Díaz, J. Baselga. 2011. Interphases in Graphene polymer-based nanocomposites: Achievements and challenges. *Advanced Materials*, 23, 5302-5310.
- M. G. Hosseini, M. Raghbi-Boroujeni, I. Ahadzadeh, R. Najjar, M. S. Seyed Dorraji. 2009. Effect of polypyrrole–montmorillonite nanocomposites powder addition on corrosion performance of epoxy coatings on Al 5000. *Progress in Organic Coatings*, 66, 321-327.
- X. Zhou. 2011. *Graphene oxidation barrier coating*. An Undergraduate Honor's Thesis, University of Colorado at Boulder.
- L. Gu, S. Liu, H. C. Zhao, H. B. Yu. 2015. Facile preparation of water-dispersible graphene sheets stabilized by carboxylated oligoanilines and their anticorrosion coatings. *ACS Applied Materials & Interfaces*, 7, 17641-17648.
- P. Nogueira, C. Ramirez, A. Torres, M. J. Abad, J. Cano, J. Lopez, I. López-Bueno, L. Barral. 2001. Effect of water sorption on the structure and mechanical properties of an epoxy resin system. *Journal of Applied Polymer Science*, 80, 71-80.
- J. A. King, R. L. Barton, R. A. Hauser, J. M. Keith. 2008. Synergistic effects of carbon fillers in electrically and thermally conductive liquid crystal polymer-based resins. *Polymer Composites*, 29, 421-428.
- X. Zhang, O. Alloul, Q. He, J. Zhu, M. J. Verde, Y. Li, S. Wei, Z. Guo. 2013. Strengthened magnetic epoxy nanocomposites with protruding nanoparticles on the graphene nanosheets. *Polymer*, 54, 3594-3604.
- L. Yue, G. Pircheraghi, S. A. Monemian, I. Manas-Zloczower. 2014. Epoxy composites with carbon nanotubes and graphene nanoplatelets-Dispersion and synergy effects. *Carbon*, 78, 268-278.