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IMPACT BEHAVIOUR OF GLASS FIBRE /EPOXY COMPOSITES WITH NANO-ENHANCED RESIN AFTER WATER EXPOSURE

ABSTRACT

Impact behaviour of glass fibre /epoxy composites with nano- SiO_2 modified resin was studied in terms of low velocity impact after water exposure. Nanocomposites with 1%, 2%, 3% 5% 7% nano- SiO_2 (Nanopox- Evonic) were investigated. Peak impact load and impact damage area as a function of nanoparticle contents were compared for dry specimens and for samples exposed to water (0.7 %wt. 1.7% water absorbed) at 1J, 2J 3J impact energies. For unmodified composite peak force was higher than for 3% modified specimens and higher for dry specimens than those exposed to water. Impact damage areas were plotted as a function of water contents for modified and unmodified samples. Failure modes were illustrated using SEM micrographs. Numerous matrix cracks were the dominating failure mode in dry specimens both unmodified and the modified. Fibre fracture was observed at 3J impact energy in all dry unmodified samples, however water exposure prevented early fibre fracture in nanocomposites. The proposed energy absorption mechanism is nanoparticles debonding.

Keywords: Polymer composites, nano-composites, low velocity impact

INTRODUCTION

In the recent years there has been a rapid growth in the use of fibre reinforced composite materials in engineering applications. However, the poor tolerance to accidental low velocity impacts of composite laminates is yet a limitation to their use in many industrial applications [1]. According to the some reports [1, 2] adding low concentrations of nanoparticles into polymers is a good solution to improve their mechanical and impact performance without compromising on density, toughness or manufacturing process. Hosur et al [3] showed that the addition of nanoclay in carbon/epoxy composites decreases the impact damage as a consequence of higher stiffness and resistance to damage progression of the nanophase-reinforced laminates. The potential of nanoparticles as reinforcement of epoxy resins is clear due to the very large surface of interaction with polymer matrix. However, nanoparticles form agglomerates which are difficult to disperse even after intense mixing. Therefore, development of nano-products in which SiO_2 nanoparticles are evenly distributed in basic resin and ready to be mixed with the matrix resin was an important step forward in nanoparticle materials processing. SiO_2 nanoparticles are chemically synthesized from aqueous sodium silicate solution. In this process the binding agent is not damaged in contrast to the processes in which powdered fillers are dispersed with dissolvers or some equipment using high shear energy. The spherical silica is distributed agglomerate free in the matrix resin. This results in a very low viscosity of the dispersion with SiO_2 contents of up to 40wt.%

[4]. Zheng et al. used 15wt.% SiO₂ nanoparticles as Nanopox (Evonic) to modify epoxy resin in carbon epoxy laminate which resulted in 20-30% improvement of fracture toughness G_{Ic} , and obtained 40% increase in Young's modulus in glass fibre reinforced laminate using 15% SiO₂ nanoparticles as epoxy matrix modifier [5]. The positive SiO₂ nanoparticles effect on the flexural strength of similar material was demonstrated in [4] using VARTM laminate processing technique.

Since the authors were previously involved in the study of water immersion effect on impact behaviour of fibre reinforced/ epoxy composites [6] and laminates with SiO₂ nanoparticles – modified surface layers [7] the aim of the present study was to determine the effect of SiO₂ nanoparticles on combined water immersion and impact behaviour in reinforced-glass fibre epoxy laminates processed by hand lay up technique still used in small ship construction companies.

EXPERIMENTAL PROCEDURE

Glass fibre-reinforced polymer (GFRP) laminates were investigated in this study with Hexion L285 DGEBA epoxy resin as matrix (curing agent 285). 0%, 1%, 2%, 3%, 5%, 7% wt. SiO₂ nanoparticles (Nanopox A410 supplied by Evonic) were added to the matrix. Nanopox A 410 is a suspension of SiO₂ 20 nm i diameter (40%wt) in Basic epoxy resin DGEBA to be mixed with epoxy resins [4]. The reinforcement was orthogonal glass fibre fabric 620 g/m² (5 plies 80 g/m² and two plies 110 g/m²). 6 sheets were formed using hand lay-up method, and cured for 24 hours in ambient temperature then postcured in 70°C for 24 hours. Specimens 100x100x1mm were conditioned in water at 50°C up to ca. 2.2% wt., water contents. Next low velocity impact tests were performed using conditioned and dry samples. Impact tester was drop weight Tower Instron CEAST 9310 with 20 mm spherical tip. Dynamic impact behaviour was recorded using DAS 64 K-S.C. recorder. Impact energy was: 1J, 2J, 3J. Next the macrophotographs of the specimens surfaces were taken and impact damage area was assessed by image analysis software Multiscan. SEM (Philips XL 30 ESEM) micrographs were taken of impacted specimens' sections so as to illustrate impact damage.

RESULTS AND DISCUSSION

Fig. 1 compares dynamic force –time impact characteristics at 3J impact energy for unmodified specimens (Fig. 1a) and specimens with 3%wt. nanoparticles (Fig. 1b) both in “dry” condition or with 0.7% and 1.7% water absorbed. Smooth curve profile for “dry” and 0.7% water (Fig. 1a) indicates that only minor impact damage resulted in these specimens whereas at higher water contents (1.7%) more severe fibre damage resulted. Addition of 3% SiO₂ (Fig. 1b) changes the failure mode of “wet (1.7% water) specimens compared to the dry (3% SiO₂) and to the unmodified (0% SiO₂). At 3J impact energy, dry (and 0.7% water) 3% SiO₂ modified specimens are subjected to fibre breakage whereas in wet condition (1.7% water) smooth load-time curve is observed which is indicative of minor damage (matrix cracks) and a positive nano-particle effect. Fig 2a illustrates slightly decreased peak impact load for SiO₂ modified specimens. This property remains practically unchanged after water exposure (up to 2% water absorbed). However failure modes are different. Energy absorbed (Fig. 2b) decreased with increased water amount and absence of nanoparticle effect was observed.

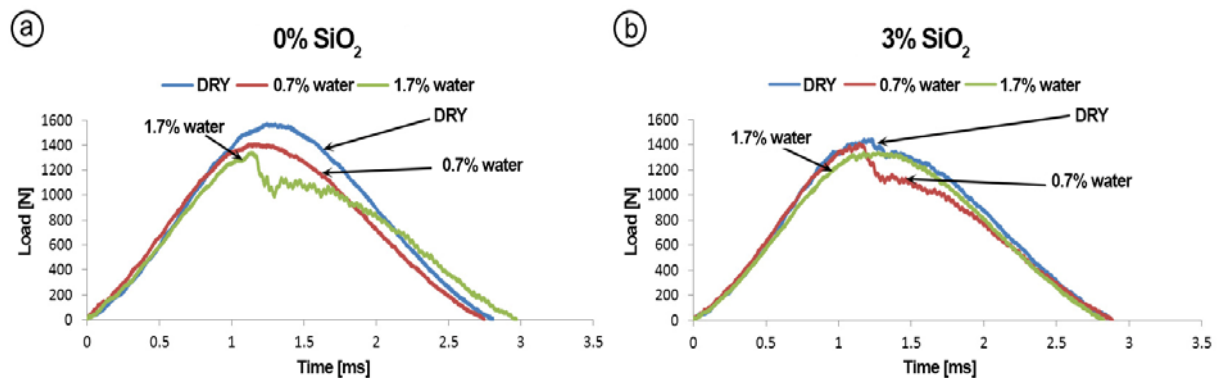


Fig. 1. Impact load-time characteristics for (a) unmodified, (b) 3% nanoparticle modified specimens in dry and wet conditions (with variable water contents). Impact energy 3J

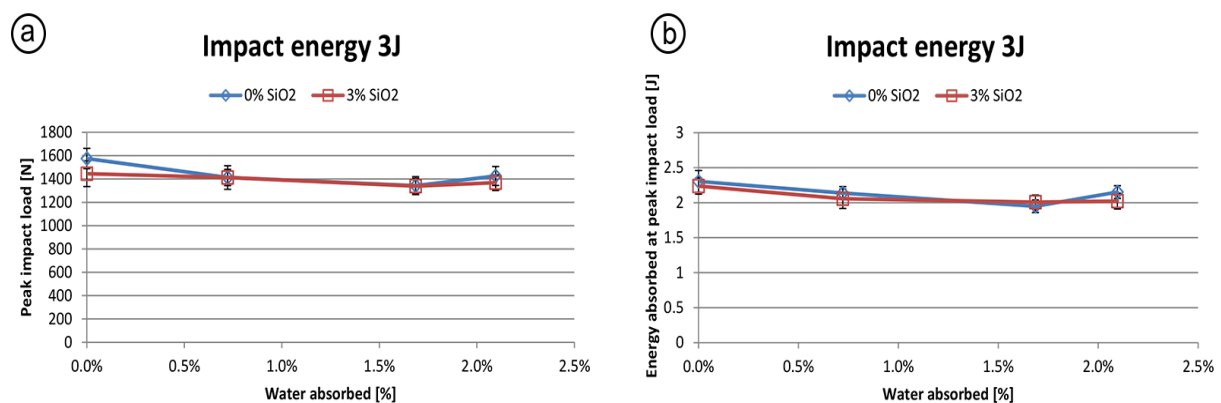


Fig. 2. a) Peak impact load as a function of amount of water absorbed by the specimens with 0% and 3% nanoparticles, b) energy absorbed at peak impact load as a function of amount of water absorbed by the specimens with 0% and 3% nanoparticles. Impact energy 3J

Fig. 3 illustrates impact damage in laminates modified with 1-7% nanoparticles in dry (upper row) and wet (0.5% water absorbed). The shape of impacted zone differs slightly due to different failure modes, however impact damage area (Fig. 4) is 10-15% lower for unmodified specimens. The improvement in the behaviour of 3% SiO₂ modified specimens is observed in terms of impact damage area for “wet” (1-2% water absorbed) specimens.

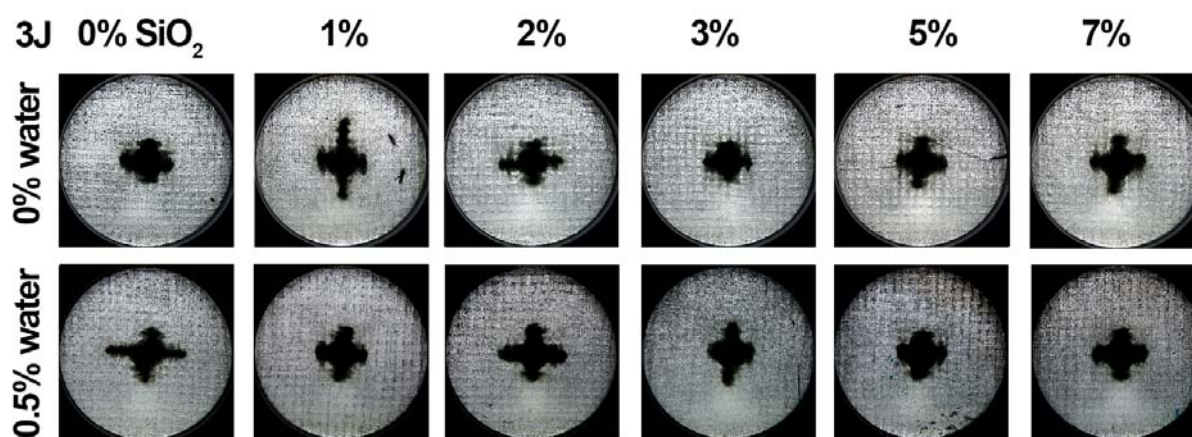


Fig. 3. Impact damage in laminates modified with 1-7% nanoparticles in dry (upper row) and wet (0,5% water absorbed)

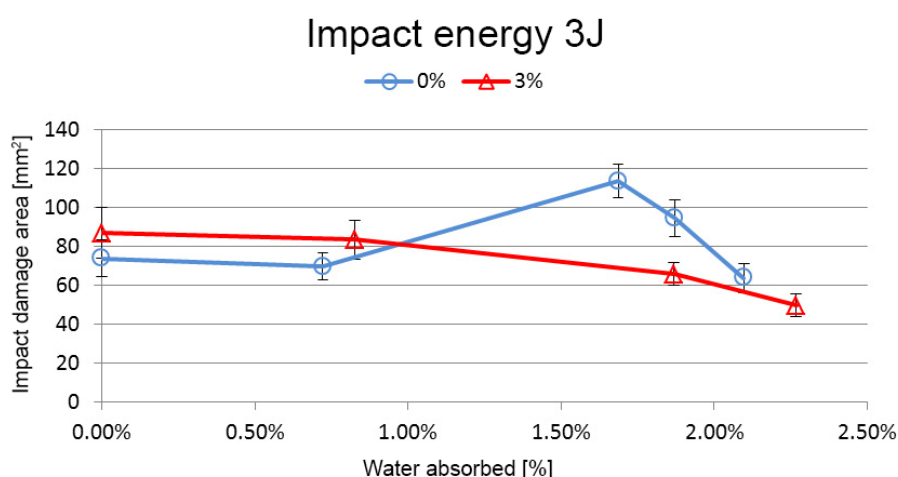


Fig. 4. Impact damage area as a function of percent water absorbed by the specimens with 0% and 3% nanoparticles. Impact energy 3J

Figs. 5ab illustrate SEM micrographs of dry (a) and wet (1.7% water absorbed) specimen modified with 3% nano-SiO₂ after 3J impact. Typical impact failure modes observed in this illustration are numerous matrix cracks (arrows). Nanoparticles cannot be observed at this magnification (100x). In Fig 5b debonded fibres are demonstrated (arrow) which is the effect of water immersion. This effect was observed in the previous study [7]. It appears that the same effect (debondings) could be observed at nanoparticle /matrix interfaces following water exposure which resulted in the change of failure mode observed for impacted specimens in wet condition. The weakest point in the material with up to ca. 2% water absorbed was the matrix with numerous debonded nanoparticles which prevented fibre fracture at 3J impact energy. The presence of debonded nanoparticles accounts for the less severe failure in water affected specimens.

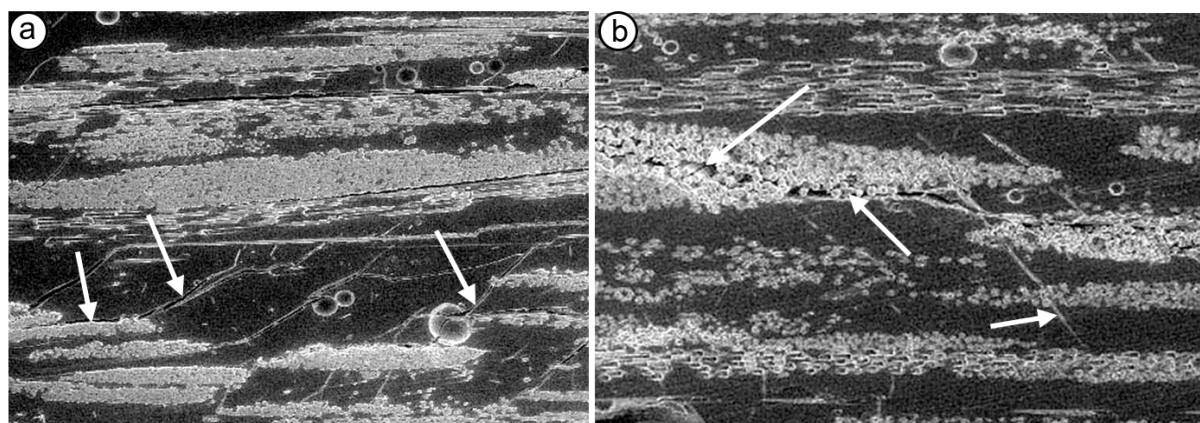


Fig. 5. SEM micrographs of dry (a) and wet (b) (1,7% water absorbed) specimen modified with 3% nano-SiO₂ after 3J impact. Arrows show matrix cracks and debonded fibres

CONCLUSIONS

Impact behaviour of glass fibre /epoxy composites with nano- SiO₂ modified resin was studied in terms of low velocity impact after water exposure. Nanocomposites with 1%, 2%, 3% 5% 7% nano-SiO₂ (Nanopox- Evonic) were investigated. Peak impact load and impact damage area as a function of nanoparticle contents were compared for dry specimens and for samples exposed to water (0.7 %wt. 1.7% water absorbed) at 1J, 2J 3J impact energies. The following conclusions have been drawn:

1. For unmodified composite peak force was higher than for 3% modified specimens and higher for dry specimens than those exposed to water.
2. Matrix cracks were the dominating failure mode in dry specimens both unmodified and the modified.
3. Water exposure prevented early fibre fracture in nanocomposites due to nanoparticles debonding, which is a proposed energy absorption mechanism.

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