Tailored interface and enhanced elastic modulus in epoxy-based composites in presence of branched poly(ethyleneimine) grafted multiwall carbon nanotubes

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In this study, branched poly(ethyleneimine), BPEI, was synthesized from carboxylic acid terminated multi-walled carbon nanotubes (c-MWNTs) and characterized using FTIR, TEM and TGA. The BPEI was then chemically grafted onto MWNTs to enhance the interfacial adhesion with the epoxy matrix. The epoxy composites with c-MWNTs and the BPEI-g-MWNTs were prepared using a sonication and mechanical stirring method, followed by curing at 100 °C and post-curing at 120 °C. The dynamic mechanical thermal analysis showed an impressive 49% increment in the storage elastic modulus in the composites. In addition, the nanoindentation on the composites exhibited significant improvement in the hardness and decrease in the plasticity index in the presence of the BPEI-g-MWNTs. Thus, epoxy composites with BPEI-g-MWNTs can be further explored as self-healing materials.

Covalent functionalization of CNTs is an effective tool to improve interfacial adhesion between CNTs and epoxy. For instance, when amino functionalized nanotubes were subjected to dicarboxylic acid acyl peroxide treatment, they revealed an impressive 70% increase in the ultimate strength and storage modulus. Non-ionic surfactant (Triton) treated CNTs showed bridging effects between nanotubes and epoxy because of the hydrophilic and hydrophobic parts of the surfactant. TETA covalently modified CNTs improved the interfacial interaction, and thus the mechanical and thermal properties of epoxy-based composites were improved. Yoonessi et al. covalently modified CNTs with aromatic, aliphatic, and aliphatic ether diamines. It was observed that aromatic diamin functionalized CNTs significantly enhanced the glass transition temperature and shear moduli. In another strategy, the epoxy chains were attached to CNTs using poly(acryloyl chloride) as a bridging agent. These composites showed higher glass transition temperature and improved mechanical properties.

Recently, nanoindentation has proved to be an important tool to measure the localized mechanical properties, i.e., modulus, stiffness, hardness, and elastic-plastic behaviour. It also serves as an efficient tool to study the interface between the thermoset and thermoplastic systems and the diffusion of one phase into another. In a recent review, the mechanical behaviour of thermoset polymers with different functionalized CNTs has been presented and the factors involved, such as, dispersion and the concentration of filler, have been highlighted.

In the present work, an epoxy matrix has been reinforced with carboxylic acid functionalized MWNTs (c-MWNTs) and

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branched poly(ethyleneimine) grafted MWNTs (BPEI-g-MWNTs) to achieve better dispersion and enhanced interfacial adhesion with the epoxy matrix. BPEI is a macromolecule rich in amine and imine groups, which can interact with the epoxy resin.\textsuperscript{21} In one of the earlier studies, carbon nanofibers were amine functionalized using PEI and made into nanopapers to reinforce an epoxy matrix.\textsuperscript{22} Schaubroeck \textit{et al.}\textsuperscript{23} studied the adhesion properties of PEI anchored epoxy resin coupled via cyanuric chloride on the surface of metals. In a recent work, PEI functionalized CNTs based epoxy composites were fabricated to study their thermal conductivity.\textsuperscript{24} Kim \textit{et al.}\textsuperscript{25} described the synthesis of a BPEI-GO conjugate for gene delivery applications. Although some studies that highlighted the improved properties of epoxy matrix in the presence of BPEI exist, a detailed analysis has not received much attention. Therefore, this work focuses on understanding the mechanical behaviour of epoxy composites in the presence of c-MWNTs and BPEI-g-MWNTs by dynamic mechanical thermal analysis and nanoindentation. BPEI was covalently grafted on the surface of c-MWNTs using EDC chemistry.\textsuperscript{26} The resulting product was further characterized using FTIR, TEM, TGA and AFM. Epoxy composites were prepared with c-MWNT and BPEI-g-MWNT by sonication and mechanical mixing followed by curing and post-curing cycles. The state of dispersion of MWNTs was studied using SEM. The mechanical properties, i.e., storage modulus, hardness and elastic-plastic behaviour, were studied using DMA and nanoindentation.

\section*{Experimental}

\subsection*{Materials and method}
Epoxy resin (bisphenol-F-epichlorohydrin, EPOLAM 8052) and amine based hardener (2,2′-dimethyl-4,4′-methylene bis(cyclohexylamine)) were kindly provided by Axson technologies (France). The branched poly(ethyleneimine) \( M_w = 25\,000 \) was procured from Sigma Aldrich. The carboxylic acid functionalized multi-walled carbon nanotubes (c-MWNTs, NC3151) were obtained from Nanocyl, Belgium. Ethanol, 1-ethyl-3-[3(dimethylamino)propyl] carbodiimide hydrochloride (EDC), N-hydroxysuccinimide (NHS) and triethylamine (TEA) were procured commercially.

\subsection*{Characterization}

The covalent functionalization of BPEI onto MWNTs was confirmed using Fourier transform infrared spectroscopy (FTIR) on a Perkin-Elmer instrument. The morphology of the synthesized BPEI-g-MWNT was characterized by transmission electron microscopy (TEM) on FEI Tecnai F30 S-TWIN and the % grafting was evaluated using Netzsch STA 409 PC Thermogravimetry Analyzer (TGA). The state of dispersion and the fracture morphology of the epoxy composites were observed by scanning electron microscopy (SEM) on FEI Sirion XL30 FEG. The mechanical behaviour of the composites was studied on Hysitron TI 900 TribolIndenter Nanoindentator with Berkovich indenter and 3-point bending clamp of TA Q800 Dynamic Mechanical Analyzer (DMA). Thermal decomposition analysis was done using Netzsch STA 409 PC Thermogravimetry Analyzer (TGA) at the rate of 10 °C min\textsuperscript{-1} under oxygen atmosphere.

\section*{Results and discussion}

\subsection*{Characterization BPEI-g-MWNT}

Amidation reaction between BPEI and carboxylic acid functional groups present on the surface of c-MWNT was characterized using spectroscopic techniques; Fig. 1a shows FTIR spectra, where the characteristic peak for carboxylic group in c-MWNNT is evident at 1700 cm\textsuperscript{-1}, while it has completely
disappeared for BPEI-g-MWNT. In addition, a sharp and intense peak at 1640 cm\(^{-1}\) appeared in the spectra of BPEI-g-MWNT. These two observations clearly demonstrate the successful grafting of BPEI onto MWNTs. TGA was used to evaluate the % grafting of BPEI.\(^{27}\) From the TGA scans (Fig. 1b), the % residual mass of c-MWNTs at 550 °C was observed to be ca. 82 wt%, while for BPEI-g-MWNTs, it was observed to be ca. 59%; rapid decomposition of BPEI was observed above 200 °C. Relative weight loss was calculated, which corresponded to the weight percent of BPEI grafted onto MWNTs. It was observed that ca. 28% of BPEI was grafted on the surface of MWNT. The morphology of the synthesized BPEI-g-MWNT was studied using TEM (Fig. 1c and d). The diameter of the c-MWNT was in the range of 8–10 nm, whereas that of BPEI-g-MWNT was found to be between 14–18 nm. Moreover, c-MWNTs were characterized by sharp edges, whereas in the case of BPEI-g-MWNTs, a layer along the axis of the MWNT was clearly observed from the TEM micrograph. Both the spectroscopy and the microscopic techniques support the successful grafting of BPEI on the surface of MWNTs. The size of the grafted BPEI chains was calculated using the following expression, where \(R_g\) is the radius of gyration and \(M_n\) is number average molecular weight of the grafted polymer.\(^{28}\) This clearly shows a single layer of BPEI on the surface of MWNTs.

\[
R_g = 0.028 M_n^{0.5} = 4.43 \text{ nm}
\]

State of dispersion of MWNTs and the fracture surface morphology of epoxy composites

Fig. 2a–e show the SEM micrographs of the cryo-fractured morphologies of neat epoxy (Fig. 2a) and the various epoxy composites. Fig. 2b and c clearly show smaller clusters of c-MWNTs due to the lack of interaction with the matrix as well as due to van der Waals’ attractive forces.\(^{29}\) However, after
Functionalization with BPEI, no such features were observed. The BPEI-g-MWNT interacts via amine and imine linkages with the epoxy resin leading to better dispersion of MWNTs (Fig. 2d and e). At higher magnification (Fig. 2e), the MWNTs are well-dispersed in the matrix. In addition, individual MWNTs were observed in the cryo-fractured morphology of the BPEI-g-MWNT–epoxy composites (as highlighted). Peng et al. reported that silane grafted CNTs were dispersed efficiently in epoxy.\(^{30}\) In another study, the fluorination of SWNTs resulted in a good dispersion of SWNTs in epoxy.\(^{31}\)

**Fig. 2** SEM micrographs of fractured surfaces of epoxy composites: (a) neat epoxy, (b) and (c) 1 wt% c-MWNT–epoxy composites, (d) and (e) 1 wt% BPEI-g-MWNT–epoxy composites.
Dynamic mechanical analysis of epoxy composites

Fig. 3a depicts the storage moduli and tan delta curves for the various composites. It is observed that the addition of different MWNTs improved the mechanical properties of the epoxy resin. For example, 1 wt% c-MWNTs improved the storage modulus by 20%, while 1 wt% BPEI-g-MWNTs further enhanced the modulus by 49% with respect to the neat epoxy resin. This can be attributed to the better dispersion and improved interfacial adhesion between BPEI-g-MWNTs and epoxy resin. The BPEI grafted onto MWNTs allows better interaction with the epoxy matrix, and thus allows more cross-linking. This network formation between grafted polymer and matrix assists efficient load transfer.\(^\text{32}\) In addition, it is clearly evident that the storage modulus for c-MWNT–epoxy composite was lower than neat epoxy in the rubbery region, while BPEI-g-MWNT–epoxy composite showed a higher storage modulus. Therefore, it can be inferred that BPEI-g-MWNT–epoxy composites demonstrate better mechanical properties even in the rubbery region.

From the peak in tan\(\delta\) plot, shown in Fig. 3b, the glass transition temperature, \(T_g\), which is the temperature at which transition from the glassy to the rubbery state occurs, can be estimated.\(^\text{15,16,13}\)

Interestingly, the \(T_g\) of c-MWNTs–epoxy composites was lowered with respect to that of neat epoxy by 7 \(^\circ\)C possibly due to the formation of CNT agglomerates in the matrix, which eventually reduced the cross-linking density. Interestingly, in the presence of BPEI-g-MWNTs, the \(T_g\) increased by 2 \(^\circ\)C, indicating a reduced macromolecular mobility in the presence of exfoliated CNT.

Hardness and elasticity assessment by nanoindentation

Fig. 4a and b illustrate the load-time cycle (loading, holding and unloading) conditions adopted in this work and the

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**Fig. 3** (a) Storage modulus and (b) Tan delta data for epoxy composites (neat, 1 wt% c-MWNT, 1 wt% BPEI-g-MWNT) obtained from dynamic mechanical analyzer.

**Fig. 4** Representative plots for (a) load as a function of time and (b) load as a function of displacement. (c) Load as a function of displacement for different samples.
corresponding load-displacement graphs. Fig. 4c shows the load as a function of displacement for different samples. Table 1 summarizes the hardness value for the various epoxy composites. The hardness, \( H \), is calculated from eqn (1), where \( P_{\text{max}} \) is the maximum applied load and \( A \) is the contact area.

The area for the Berkovich indentor is given by eqn (2). 20 The nanoindentation curve describes the mechanical behaviour of a polymer during loading and unloading cycles, and hence it reveals critical insights into the elastic-plastic behaviour, which is also known as the plasticity index, \( \psi \) (eqn (3)). 34,35

\[
H = \frac{P_{\text{max}}}{A}
\]

\[
A = 24.5h_p^2
\]

\[
\psi = \frac{A_p - A_e}{A_p}
\]

It is evident that the addition of different MWNTs enhanced the hardness of the epoxy resin. This observation is in line with the fact that stiffer and stronger nanoparticles enhance the hardness of the polymer matrix. 34 Therefore, it can be concluded that during the indentation process, the indenter is in contact with stiffer particles and undergoes little plastic deformation. The BPEI-g-MWNTs–epoxy composites showed the highest hardness value among the different composites studied here. As mentioned above, the plasticity index sheds light on the mechanical properties of a material. Moreover, it is a parameter used to analyse the self-healing ability of a material. 34 From Table 2, it can be inferred that there is an improvement of ca. 9% in the plasticity index of BPEI-g-MWNTs–epoxy composite compared to neat epoxy matrix. Thus, epoxy–BPEI-g-MWNTs shows self-healing behaviour under the given load conditions and can further be explored as a self-healing material.

**Thermal properties**

TGA studies were carried out to study the thermal degradation behaviour in the composites. Fig. 5 depicts the TGA profile for different epoxy composites. It is observed that in all the composites, the degradation started around 300 °C. Therefore, these composites can also be explored for high temperature application. Epoxy composites with BPEI-g-MWNTs showed a small amount of degradation below 300 °C, which can be attributed to the decomposition of BPEI.

**Conclusions**

In this study, BPEI-g-MWNTs was synthesized using EDC chemistry, which were used to improve the interfacial adhesion, and thus the mechanical behaviour of epoxy matrix was improved using an amine and imine rich molecule. Epoxy composites with 1 wt% carboxylic acid modified MWNT and BPEI-g-MWNTs were prepared using sonication with mechanical stirring followed by curing and post-curing cycles. The fracture surface morphology of the composites showed better dispersion in the case of BPEI-g-MWNTs in epoxy than in the case of c-MWNTs. The storage elastic modulus of BPEI-g-MWNTs–epoxy composites showed ca. 49% improvement with respect to neat epoxy samples and a marginal improvement of 2 °C in \( T_g \). The nanoindentation results displayed increase in the hardness, and hence a lower plastic deformation of epoxy in the presence of 1 wt% BPEI-g-MWNTs. From the plasticity indices value, it was concluded that BPEI-g-MWNTs–epoxy composites show potential self-healing ability to a certain extent under the given loading conditions. Thermogravimetric analysis showed that the decomposition temperature of all the composites studied here is above 300 °C.

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