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## Preparation and characterisation of graphene oxide containing block copolymer worm gels†

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This paper reports a generic method for preparing reinforced nanocomposite worm-gels. Aqueous poly(glycerol monomethacrylate)-*b*-poly(2-hydroxypropyl methacrylate) (PGMA–PHPMA) and methanolic poly(glycerol monomethacrylate)-*b*-poly(benzyl methacrylate) (PGMA–PBzMA) worm gels were prepared by RAFT-mediated polymerisation-induced self-assembly (PISA). The former system undergoes a reversible worm-to-sphere degelation transition upon cooling to 5 °C whilst the latter system undergoes the same transition on heating to 56 °C. This transition allows these copolymer dispersions to be readily mixed with graphene oxide (GO) whilst in a low viscosity state and form nanocomposite gels on returning to room temperature via a sphere-to-worm transition. Various quantities of GO were added to the studied copolymer dispersions at a fixed copolymer content of 15% w/w. A general trend was observed whereby relatively small quantities of GO caused the gel strength of the nanocomposite gel to be higher than that of the pristine worm-gel, as determined by oscillatory rheology. Additional quantities of GO resulted in gel weakening or prevented gel-reformation altogether. For instance, 15% w/w PGMA<sub>52</sub>–PHPMA<sub>130</sub> worm gels had a storage modulus ( $G'$ ) of approximately 1.5 kPa. The addition of 1.5% w/w GO based on the copolymer caused  $G'$  to increase to approximately 4.0 kPa but >1.5% w/w GO resulted in gel strengths <1.0 kPa. A combination of aqueous electrophoresis and transmission electron microscopy measurements were used to investigate the mechanism of nanocomposite gel formation. It was observed that the PGMA-based copolymers readily absorb onto the surface of GO. Thus, the role of GO is both to strengthen the worm-gels when an optimal concentration of GO is used, but also prevent worm-reformation if too much copolymer becomes absorbed on the surface of the sheets.

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## Introduction

Composite hydrogels are an interesting class of materials which have become widely investigated in recent years.<sup>1–6</sup> For example, in 2002 Haraguchi *et al.*<sup>7</sup> reported a high-strength nanocomposite hydrogel obtained by *in situ* polymerisation of *N*-isopropylacrylamide (NIPAm) in a suspension of Hectorite nanosheets. The reported tensile strength of these gels was approximately 10 times that of traditional hydrogels. Subsequent studies have shown that the preparation of nanocomposite gels can be relatively simple, economical and environmentally friendly, and

result in hydrogels with significantly improved mechanical properties and swelling behaviour.<sup>8–10</sup> At present, numerous types of nanocomposite gel have been reported including graphene oxide-based composite gels,<sup>11–16</sup> carbon nanotube composite gels,<sup>17–21</sup> nanocellulose composite gels,<sup>22–26</sup> inorganic clay composite gels<sup>7,27–32</sup> and other inorganic nanoparticle-containing composite gels.<sup>33–37</sup>

Graphene oxide (GO) is a water-dispersible 2D material which has the planar structure of graphene but has a number of oxy-functional groups present.<sup>11,38–40</sup> The oxy functional groups are hydroxyl and epoxy groups on the basal plane, with smaller amounts of carboxyl, carbonyl, phenol, lactone, and quinone at the sheet edges.<sup>39,41</sup> The presence of these functional groups provides potential advantages for using GO in numerous applications, particularly due to the enhanced water dispersibility of GO in comparison to graphene.<sup>42,43</sup> Furthermore, it is known that the incorporation of 2D materials such as graphene oxide into polymeric hydrogels can drastically modify their mechanical,<sup>44,45</sup> biocompatibility<sup>46–49</sup> and optical properties.<sup>50,51</sup> For instance, Liu<sup>52</sup> used GO as a cross-linker and acrylamide (AM) as a monomer to prepare PAM-GO nanocomposite gels by *in situ* polymerisation. Compared with other

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† Electronic supplementary information (ESI) available: Synthetic schemes; <sup>1</sup>H NMR spectra and GPC chromatograms and digital photographs of (co)polymers; variable temperature DLS and rheology studies on copolymers; TEM images of copolymers, GO and composites; aqueous electrophoresis studies of (co)polymer/GO mixtures; critical gelation temperature recorded over several heating-cooling cycles for G<sub>52</sub>–H<sub>130</sub>–1.5% GO; rheology data for 15% w/w G<sub>60</sub>–B<sub>70</sub> with various GO concentrations in methanol. See DOI: 10.1039/d2sm00045h



cross-linked PAM hydrogels, it had improved mechanical properties. For example, breaking elongation reached more than 3000% and the observed tensile strength was nearly 390 kPa. In addition, photothermally sensitive GO-NIPAM composite gels showed optical sensitivity, as the phase transition of the gels could be controlled by an infrared laser.<sup>50</sup> Bai *et al.*, reported supramolecular GO containing gels using different copolymers as a crosslinker. In this case, they indicated that hydrogen bonding,  $\pi$ -stacking and hydrophobic effects between copolymer and GO sheets, as well as the lateral dimensions of the GO sheets influenced gel formation. In addition, gel formation could be adjusted by modulating the attraction between the GO sheets and copolymer.<sup>53</sup>

Block copolymer self-assembly is generally used to prepare nanoparticles and structures in the length scale of 10 nm to 1  $\mu$ m. This length-scale can greatly affect the physical, chemical and biological properties of these materials.<sup>54–56</sup> Block copolymer self-assembly in a dispersion can result in a variety of ordered structures such as spherical micelles (spheres), worm-like micelles (worms), and polymersomes (vesicles).<sup>57–61</sup> The self-assembled structures formed can be adjusted by changing *e.g.*, the solvent or relative molar mass of each block of the copolymer. Relatively recently, polymerisation-induced self-assembly (PISA) *via* reversible addition–fragmentation chain-transfer (RAFT) polymerisation has been demonstrated to be a versatile and efficient route towards the synthesis of various types of block copolymer nano-objects.<sup>55,62</sup> The particles formed can have a range of functionalities, both in their core and corona.<sup>63</sup> RAFT aqueous dispersion polymerisation can be used to directly prepare block copolymer nanoparticles in water with a range of morphologies. For example, poly(glycerol monomethacrylate)-*b*-poly(2-hydroxypropyl methacrylate) (PGMA–PHPMA) prepared *via* RAFT aqueous dispersion polymerisation readily forms so-called ‘worm gels’ which have been widely studied and reported on.<sup>58–60,64</sup> These PGMA–PHPMA worm-gels are free-standing at room temperature but become free-flowing liquids when cooled to 4 °C. This reversible de-gelation transition is attributed to worm-to-sphere order-order transitions which cause a loss of worm entanglement.<sup>59,65,66</sup> Due to this thermo-responsive property, these gels can be used as sterilisable gels *via* ultrafiltration at low temperatures. Other systems have been reported which show a similar transition when exposed to a suitable stimulus such as temperature or pH change.<sup>59,67,68</sup> However, RAFT aqueous dispersion polymerisation is only suited to a limited range of formulations such as the PGMA–PHPMA system described above and *e.g.*, poly(ethylene glycol)-*b*-poly(4-hydroxybutyl acrylate).<sup>69</sup> Therefore, research into RAFT-mediated dispersion polymerisation in a wider range of media such has proliferated.<sup>70–72</sup> Interestingly, reversible gel transitions can also be triggered by heating for some block copolymer worm gel systems. For instance, poly(lauryl methacrylate)–poly(benzyl methacrylate) (PLMA–PBzMA) copolymers showed a worm-to-sphere-to-worm morphological transition when heating above 50 °C and returning to room temperature.<sup>66</sup> This behaviour has also been demonstrated for PGMA–PBzMA

copolymers prepared in alcoholic media.<sup>73</sup> Thus these reversible worm gel transitions provide an opportunity to load these materials with fillers such as GO and study how self-assembly and the resulting properties are affected.

Herein, the preparation of a fascinating class of 3D composite hydrogels comprising 1D block copolymer worm-like micelles and 2D graphene oxide sheets is reported using two previously reported worm-gel formulations, aqueous PGMA–PHPMA and methanolic PGMA–PBzMA.<sup>58,59,64,74,75</sup> The reversible de-gelation of the copolymer worm-gels on cooling or heating allows facile mixing of GO with the copolymer dispersions (Fig. 1). On returning to room temperature composite gels are formed and subsequently investigated using a combination of transmission electron microscopy and rheology.

## Experimental

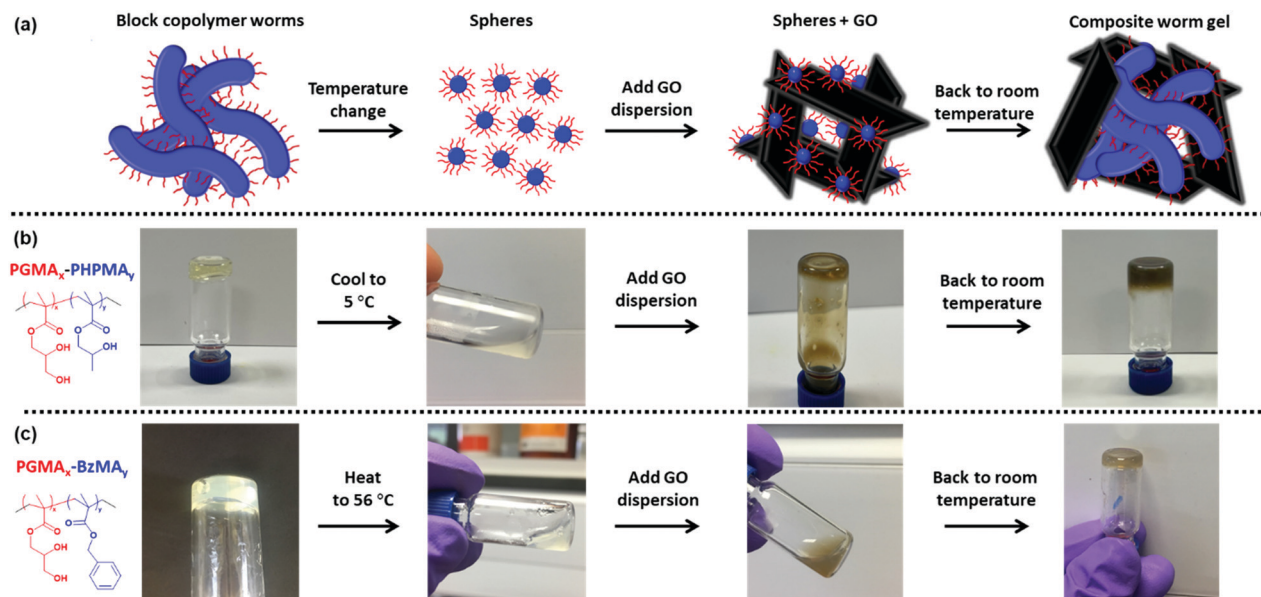
### Materials

All reagents were purchased from Sigma-Aldrich (UK) and used as received, unless otherwise noted. Glycerol monomethacrylate (GMA) was kindly donated from GEO Specialty Chemicals (UK), 2-Hydroxypropyl methacrylate (HPMA, 97%) was purchased from Alfa Aesar (UK). Benzyl methacrylate (BzMA) was purchased from Alfa Aesar (UK) and passed through a column of activated basic alumina to remove inhibitors and impurities before use. Methanol (>99.9%) and ethanol (95%) were purchased from Fisher Scientific (UK) and used as received. Graphene oxide aqueous dispersion was purchased from Graphena (Spain) and purified before use. 4-Cyano-4-(2-phenylethane sulfanylthiocarbonyl) sulfanylpentanoic acid (PETTC) was prepared in-house using previously published methods.<sup>76,77</sup> Deionised water was used in all experiments.

### Synthesis of poly(glycerol monomethacrylate) (PGMA) *via* RAFT solution polymerisation

The preparation of PGMA by RAFT solution polymerisation has been reported extensively in the literature.<sup>59</sup> In this work, polymerisation of GMA was conducted in ethanol at 70 °C. This protocol afforded PGMA macromolecular chain transfer agents (macro-CTAs; denoted G<sub>n</sub>) with narrow molecular weight distributions at high yield. A typical protocol is as follows. For a target degree of polymerisation of 50 (G<sub>50</sub>), GMA monomer (20 g, 124 mmol) and PETTC RAFT agent (0.8476 g, 2.5 mmol) were weighed into a 250 mL round-bottomed flask and purged with N<sub>2</sub> for 30 min. 4,4-Azobis(4-cyanovaleric acid) (ACVA) initiator (0.14 g, 0.5 mmol, PETTC/ACVA molar ratio = 5:0) and anhydrous ethanol (20 g, previously purged with N<sub>2</sub> for 20 min) were then added, and the resulting yellow solution was degassed for a further 10 min while stirring to form a homogeneous solution. The flask was subsequently sealed and immersed in an oil bath set at 70 °C. After 120 min, the polymerisation was quenched by immersion in an ice bath and opening to air. The final degree of polymerisation (DP) was 52, as determined by <sup>1</sup>H-NMR analysis (Fig. S3, ESI<sup>†</sup>) using D<sub>2</sub>O. The polymer was purified by dialysis





**Fig. 1** (a) Scheme for preparing graphene oxide containing block copolymer worm gels. (b) PGMA<sub>x</sub>-PPHMA<sub>y</sub> copolymer dispersions are cooled to 5 °C, mixed with graphene oxide (GO) and allowed to return to room temperature to form free-standing nanocomposite gels. (c) PGMA<sub>x</sub>-PBzMA<sub>y</sub> copolymer dispersions are heated to 56 °C, mixed with graphene oxide and allowed to return to room temperature to form free-standing nanocomposite gels.

(MWCO = 3500 g mol<sup>-1</sup>) against deionised water and freeze-dried to form a yellow powder. DMF GPC analysis indicated an  $M_n$  of 4700 g mol<sup>-1</sup> and an  $M_w/M_n$  of 1.17 (Fig. S2, ESI<sup>†</sup>).

#### Preparation of PGMA-PPHMA worm gels via RAFT aqueous dispersion polymerisation of HPMA

The preparation of PGMA-PPHMA worm-gels by RAFT dispersion polymerisation has been reported extensively in the literature.<sup>59</sup> A typical protocol for the synthesis of a PGMA<sub>52</sub>-PPHMA<sub>130</sub> (G<sub>52</sub>-H<sub>130</sub>) worm gel is as follows. PGMA<sub>52</sub> macro-CTA (5.0 g, 0.566 mmol) and HPMA monomer (11.4 g, 79.274 mmol; target DP = 130) were weighed into a 100 mL round bottomed flask and purged with N<sub>2</sub> for 20 min. ACVA was added (31.74 mg, 0.101 mmol, CTA/ACVA molar ratio = 5.0) and purged with N<sub>2</sub> for a further 5 min. Deionised water (65.84 mL, producing a 20.0% w/w aqueous solution), which had been previously purged with N<sub>2</sub> for 30 min, was then added and the solution was purged for a further 5 min prior to immersion in an oil bath set at 70 °C. The reaction was stirred for 3 h before the polymerisation was quenched by exposure to air. The product was a soft free-standing gel. The absence of signals owing to the vinyl protons of the HPMA monomer in the <sup>1</sup>H NMR spectrum indicated that the polymerisation had attained more than 99% conversion (Fig. S3a, ESI<sup>†</sup>).

#### Preparation of PGMA-PBzMA worm gels via RAFT alcoholic dispersion polymerisation of BzMA

A typical protocol for the synthesis of a PGMA<sub>60</sub>-PBzMA<sub>70</sub> (G<sub>60</sub>-B<sub>70</sub>) worm gel is as follows.<sup>73</sup> PGMA<sub>60</sub> macro-CTA (1.0 g, 0.096 mmol) and BzMA monomer (1.23 g, 7.0 mmol; target DP = 70) were weighed into a 25 mL round bottomed flask and purged with N<sub>2</sub> for 15 min. ACVA was added (31.74 mg,

0.101 mmol, CTA/ACVA molar ratio = 5.0) and purged with N<sub>2</sub> for a further 5 min. Methanol (8.958 g, producing a 20.0% w/w solution), which had been previously purged with N<sub>2</sub> for 30 min, was then added, and the solution was purged for a further 5 min prior to immersion in an oil bath set at 64 °C and reflux for 24 h. The reaction was quenched by exposure to air. The product was a soft free-standing gel. The absence of signals at 6.15 ppm and 5.57 ppm owing to the vinyl protons of the BzMA monomer in the <sup>1</sup>H NMR spectrum indicated that the polymerisation had attained high monomer conversion (Fig. S3b, ESI<sup>†</sup>).

#### Preparation of graphene oxide-containing block copolymer worm gels

All GO dispersions were purified by dialysis against water for 5 days and sonicated in a bath sonicator for 20 min before use. The mean intensity-average particle diameter for the GO dispersion after bath sonication was ~1900 nm. TEM studies indicated that the dispersion contained a broad distribution of sheet sizes, with a significant population of sub-μm sheets (see Fig. 2 and 3 and Fig. S7, S8, S12 and S13 in the ESI<sup>†</sup>).

For PGMA<sub>52</sub>-PPHMA<sub>130</sub>-x% GO composite worm-gels, GO dispersion (4 mg mL<sup>-1</sup>, pH 5.1) and 20% w/w G<sub>52</sub>-H<sub>130</sub> copolymer worm gel were cooled to approximately 5 °C until the copolymer dispersion was in a free-flowing state (pH 6.3). Depending on the target GO concentration in the final mixture, appropriate quantities of the pre-cooled GO dispersion and/or deionised water were added to cooled copolymer dispersion to yield a copolymer concentration of 15% w/w (measured dispersion pH 5.9 to 6.2). The samples were mixed thoroughly for 10 s whilst cool using a vortex mixer and subsequently allowed to return to room temperature.



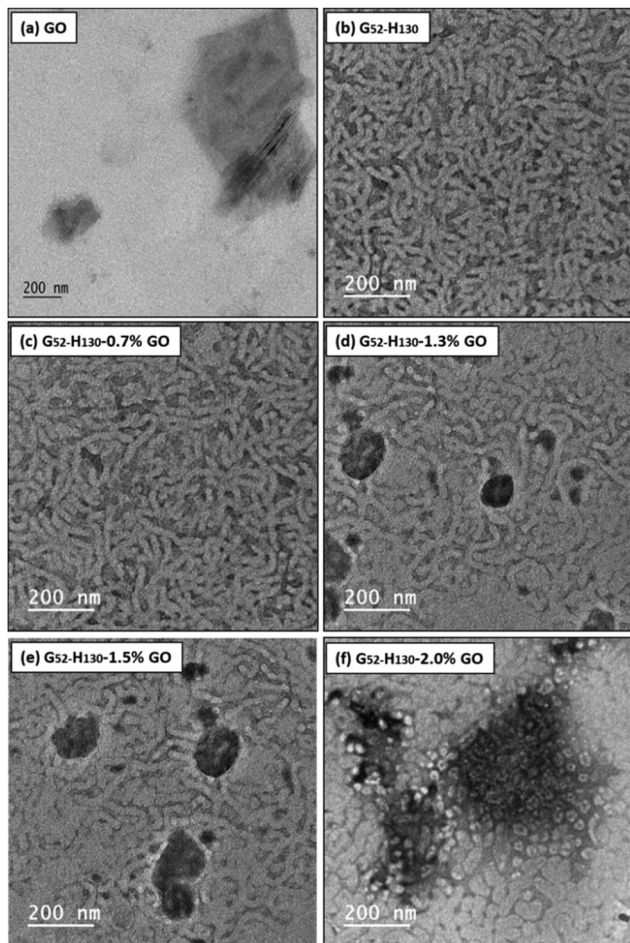


Fig. 2 TEM images for (a) GO; (b) 15% w/w  $G_{52}$ - $H_{130}$ . (c)  $G_{52}$ - $H_{130}$ -0.7% GO. (d)  $G_{52}$ - $H_{130}$ -1.3% GO; (e)  $G_{52}$ - $H_{130}$ -1.5% GO and (f)  $G_{52}$ - $H_{130}$ -2.0% GO. Samples were diluted to 0.01% w/w before being deposited on to carbon-coated TEM grids at room temperature.

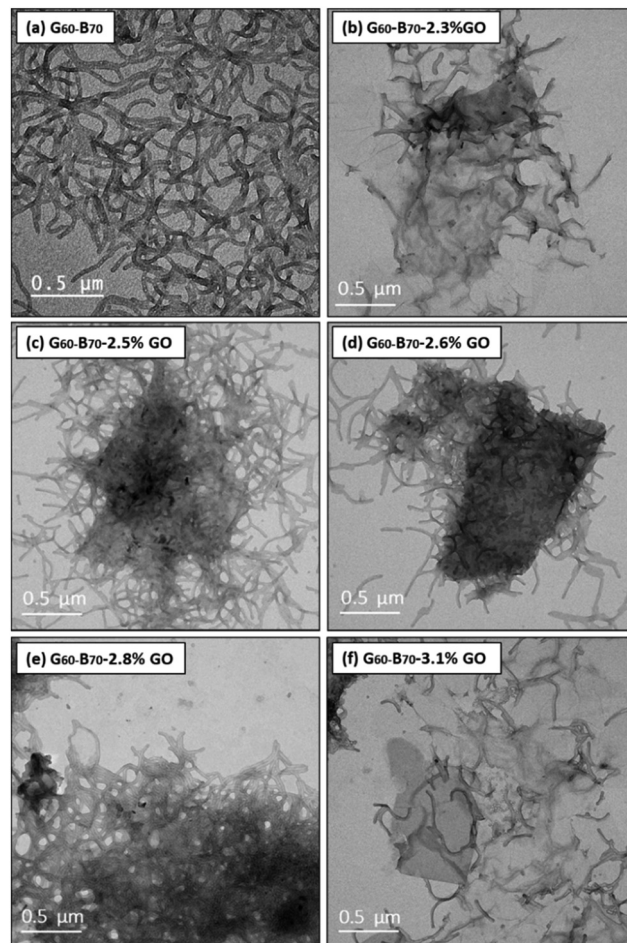


Fig. 3 TEM images for (a) 15% w/w  $G_{60}$ - $B_{70}$ ; (b)  $G_{60}$ - $B_{70}$ -2.3% GO, (c)  $G_{60}$ - $B_{70}$ -2.5% GO, (d)  $G_{60}$ - $B_{70}$ -2.6% GO; (e)  $G_{60}$ - $B_{70}$ -2.8% GO and (f)  $G_{60}$ - $B_{70}$ -3.1% GO. Samples were diluted to 0.01% w/w before being deposited on to carbon-coated TEM grids at room temperature.

For  $PGMA_{60}$ - $PBzMA_{70-x\%}$  GO gels,  $G_{60}$ - $B_{70}$  worm gels (pH 6.7) were heated to approximately 56 °C and allowed to equilibrate for ~10 minutes. On heating, the worm gels transformed into a viscous but free-flowing liquid. Two types of continuous phase were used to prepare  $G_{60}$ - $B_{70}$ - $x\%$  GO worm gels: methanol and water/methanol mixtures. For water/methanol mixtures, different concentrations of pre-heated aqueous GO dispersion and/or deionised water were added to the methanolic copolymer dispersion. For methanol only formulations, GO was freeze-dried and dispersed in methanol using bath sonication for 40 minutes prior to heating and addition to the heated copolymer dispersion. The samples were mixed thoroughly for 10 s whilst warm using a vortex mixer and subsequently allowed to return to room temperature. In all cases the copolymer content of the final mixture was fixed at 15% w/w (measured dispersion pH 6.4 to 6.6).

### Characterisation

**$^1H$  NMR spectroscopy.** All  $^1H$  NMR spectra were recorded on either a Bruker Avance III 400 MHz spectrometer with 128 scans

averaged per spectrum at 25 °C. PGMA was dissolved in  $D_2O$  and  $PGMA$ - $PHPMA$  diblock copolymers were freeze-dried and dissolved in  $DMSO-d_6$  prior to  $^1H$  NMR analysis.

**Gel permeation chromatography (GPC).** 0.50% w/w copolymer solutions were prepared in DMF containing DMSO ( $10 \mu L mL^{-1}$ ) as a flow-rate marker. GPC measurements were conducted using HPLC-grade DMF eluent containing 10 mM LiBr at 60 °C at a flow rate of  $1.0 mL min^{-1}$ . An Agilent Technologies 1260 Infinity GPC/SEC system fitted with two Polymer Laboratories PL gel  $5 \mu m$  Mixed C columns connected in series, and a refractive index detector was used to assess molar mass distributions using polystyrene calibration standards.

**Dynamic light scattering (DLS) and aqueous electrophoresis.** A Malvern Zetasizer Nano ZS instrument was used to measure particle size and zeta potential. The instrument is equipped with a He-Ne solid-state laser operating at 633 nm and detects back-scattered light at a scattering angle of 173°. All size measurement data were averaged over three consecutive runs comprising thirteen measurements each. For Zeta potential



measurements, the same instrument was used. The solution pH was initially adjusted to 10 using 0.1 M KOH in the presence of 1 mM KCl and then manually lowered from 10 to 4 using 0.01 M HCl.

**Transmission electron microscopy (TEM).** Diblock copolymer dispersions were diluted to 0.20–0.15% w/w at 20 °C prior to staining. 3  $\mu$ L was then placed onto 400 mesh carbon-coated copper grids for 90 min and carefully blotted with filter paper to remove excess dispersion. The samples were stained in the vapour space above RuO<sub>4</sub> solution for 7 min at room temperature.<sup>66</sup> The mean nanoparticle diameter was determined by ImageJ software by counting over 200 randomly selected particles for each sample. Imaging was performed using a Philips CM 20 instrument connected to a Gatan 1 k CCD camera at an accelerating voltage of 200 kV.

**Rheology measurements.** An AR-G2 rheometer (TA instruments) equipped with a variable temperature Peltier plate and 60 mm steel parallel plate was used for all experiments. An oscillatory mode was used to measure storage modulus ( $G'$ ) and loss modulus ( $G''$ ) as a function of angular frequency. Percentage strain amplitude as a function of temperature was used to assess critical gelation temperature and gel strengths. Percentage strain amplitude sweeps were conducted between 0.01 and 130 rad s<sup>-1</sup> at a constant temperature of 25 °C, with a frequency of 10 Hz. Temperature sweeps were conducted using applied strain amplitude of 1.0% at an angular frequency of 10 Hz.

## Results and discussion

### Copolymer worm-gel preparation

PGMA<sub>52</sub> ( $M_w/M_n = 1.17$ ) and PGMA<sub>60</sub> ( $M_w/M_n = 1.26$ ) were synthesised *via* RAFT solution polymerisation in ethanol at 70 °C. PGMA<sub>52</sub> was block-extended with HPMA *via* RAFT aqueous dispersion polymerisation at 20% w/w and 70 °C (see Fig. S1a, ESI†). The target DP of the core forming PHPMA was 130 to obtain a free-standing worm-gel (Fig. S4d, ESI†).<sup>78</sup> GPC analysis confirmed a relatively narrow molecular weight distribution ( $M_w/M_n = 1.06$ ) (Fig. S2a, ESI†) and the thermal response of the G<sub>52</sub>-H<sub>130</sub> copolymer dispersion was investigated by variable temperature DLS measurements (Fig. S4a, ESI†) on 0.1% w/w aqueous dispersions. The initial particle diameter reported by DLS was approximately 190 nm at 40 °C. This reduced to approximately 30 nm at 2 °C confirming the expected worm-to-sphere morphological transition.<sup>59</sup> On heating, a constant diameter of  $\sim$ 30 nm was recorded, indicating an irreversible transition at low copolymer concentrations. Oscillatory rheology (Fig. S4c, ESI†) conducted on the copolymer at 20% w/w demonstrated fully reversible de-gelation with a critical gelation temperature (CGT), which is defined as crossover temperature for  $G'$  and  $G''$  curves, of 10 °C on cooling and 11 °C on heating. The gel strength ( $G'$ ) of the 20% w/w G<sub>52</sub>-H<sub>130</sub> copolymer gel was nearly 1.3 kPa (Fig. S4b, ESI†), as was expected from previous work.<sup>59</sup>

BzMA was polymerised in the presence of PGMA<sub>60</sub> *via* RAFT alcoholic dispersion polymerisation at 20% w/w in methanol

(Fig. S1b, ESI†).<sup>73</sup> The target DP of the core forming PBzMA was 70 to obtain a free standing gel and TEM confirmed that a worm morphology was obtained (Fig. S5c, ESI†). GPC analysis confirmed a relatively narrow molecular weight distribution ( $M_w/M_n = 1.27$ ) (Fig. S2b, ESI†) and oscillatory rheology studies indicated that  $G'$  of the 20% w/w PGMA<sub>60</sub>-PBzMA<sub>70</sub> gel was approximately 1.1 kPa (Fig. S5b, ESI†).<sup>73</sup> Variable temperature DLS measurements examined the thermal response of the G<sub>60</sub>-B<sub>70</sub> dispersions. For 0.1% w/w aqueous dispersions, the Z-average particle diameter was approximately 900 nm at 25 °C (Fig. S5a, ESI†). After heating to 60 °C, the particle diameter reduced to  $\sim$ 90 nm indicating that the worm-to-sphere morphological transition occurred. On cooling back to 25 °C, DLS confirmed that the transition was irreversible at low copolymer concentration, as observed for G<sub>52</sub>-H<sub>130</sub>.<sup>66</sup>

### Preparation of G<sub>52</sub>-H<sub>130</sub>-GO composite gels

To enable mixing with GO dispersions, the aqueous G<sub>52</sub>-H<sub>130</sub> diblock copolymer dispersions necessarily required dilution. On reducing the solids content from 20% w/w to 15% w/w, a free-standing gel was retained (Fig. S6a, ESI†) and the copolymer morphology was unchanged (Fig. 2b). Diluting the copolymer reduced the gel strength from  $\sim$ 1.5 kPa at 20% w/w (Fig. S4b, ESI†) to  $\sim$ 1.0 kPa at 15% w/w (Fig. 4b). However, when the copolymer was diluted further to 10% w/w, the copolymer was no longer a free-standing gel (Fig. S6a, ESI†). Therefore, all subsequent dilutions of G<sub>52</sub>-H<sub>130</sub> with GO dispersions were conducted at a fixed final copolymer concentration of 15% w/w.

As shown in Fig. 1b the 15% w/w G<sub>52</sub>-H<sub>130</sub> worm gel dispersion undergoes de-gelation on cooling to form a free-flowing liquid. The low viscosity of this cooled copolymer dispersion (at 5 °C) facilitates mixing with various concentrations of pre-cooled GO dispersion. Therefore, GO was mixed with cooled copolymer to yield a GO content between 0.7 to 2.2% w/w based on copolymer, whilst keeping the total copolymer content fixed at 15% w/w. At low temperature, all samples appeared as brown free-flowing liquids (Fig. 1b). On warming the mixtures to room temperature, most samples returned to being a gel, as judged by tube inversion tests (Fig. 1b). However, gelation did not occur for samples containing relatively high concentrations of GO ( $\leq$ 1.5% w/w based on copolymer, Fig. 4a).

Oscillatory rheology studies of the G<sub>52</sub>-H<sub>130</sub>-GO composite gels at room temperature clearly show differences between respective gel strengths with increasing GO loading (Fig. 4). For example, increasing the GO content from 0 to 1.5% w/w based on copolymer increases the measured gel strength from  $\sim$ 1.5 to  $\sim$ 4.0 kPa (Fig. 4b). TEM images of the composite gels (Fig. 2c–e) demonstrate that the copolymer worms re-assemble after the cooling-mixing-heating cycle and allow the gels to reform. Thus, the observed gel strengthening effect is likely a result of attractive polymer-GO interactions and strengthening of the composite/worm network by the incorporated GO flakes. To support the hypothesised interaction between these PGMA-based copolymers and GO, aqueous electrophoresis studies



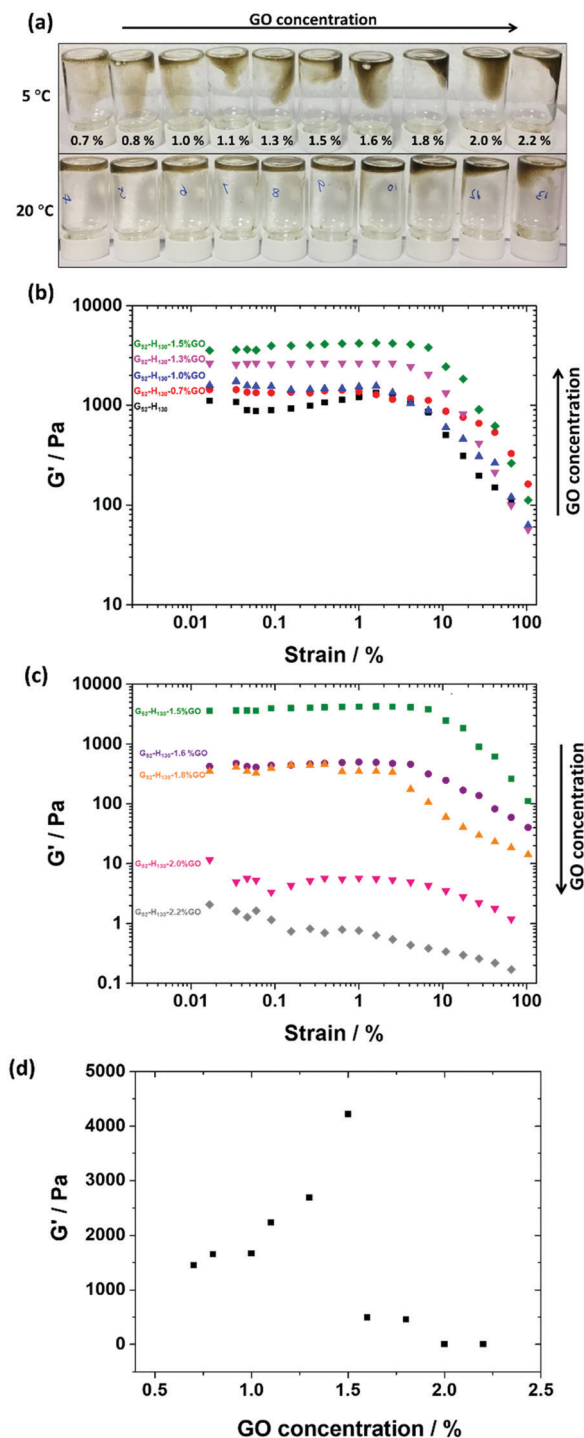


Fig. 4 (a) Photographs of graphene oxide containing PGMA<sub>52</sub>-PHPMA<sub>130</sub> block copolymer dispersions with graphene oxide concentrations between 2.2 and 0.7% w/w. The upper image shows samples after mixing at 5 °C and the lower image shows samples after equilibration for > 2 h at room temperature. Storage modulus ( $G'$ ) versus % strain for (b) 15% w/w G<sub>52</sub>-H<sub>130</sub> with increasing GO concentration (0–1.5% w/w) and (c) 15% w/w G<sub>52</sub>-H<sub>130</sub> with decreasing GO concentration (1.5–2.2% w/w). (d) Average storage modulus measured between 0.01 and 2% strain for different GO-containing G<sub>52</sub>-H<sub>130</sub> copolymer dispersions. All measurements were conducted at a frequency of 10 Hz, a strain of 1.0% and a controlled temperature of 25 °C.

were conducted as function of pH on G<sub>52</sub> macro-CTA, diluted G<sub>52</sub>-H<sub>130</sub> worms, GO dispersion (4 mg mL<sup>-1</sup>), and selected (co)polymer-GO mixtures (0.7%, 1.5% and 2.0% GO) after purification (Fig. S9a and b, ESI<sup>†</sup>). Both the G<sub>52</sub> macro-CTA and the G<sub>52</sub>-H<sub>130</sub> copolymer dispersions exhibited slightly negative Zeta potentials.<sup>60</sup> This anionic character is imparted due to the carboxylic acid end-groups from the PETTC RAFT agent.<sup>79</sup> The GO dispersion was more anionic across the whole pH range and had a Zeta potential less than -30 mV at pH > 5. On addition of either G<sub>52</sub> macro-CTA or G<sub>52</sub>-H<sub>130</sub>, the anionic charge from GO is screened and the Zeta potential becomes less negative. For example, with 2.2% GO based on (co)polymer the Zeta potential is highly anionic at pH > 5. However, with 0.7% GO based on (co)polymer the Zeta potential appears more similar to the pristine (co)polymer. Thus, it is likely that on mixing GO with cooled G<sub>52</sub>-H<sub>130</sub> dispersions the copolymer spheres adsorb onto the surface of the GO sheets, become incorporated homogeneously within the reformed composite worm gel and reinforce the gel upon the application of shear.

On increasing the GO content from 1.5 to 2.2% w/w based on copolymer the measured gel strength significantly decreases (Fig. 4c). This is hypothesised to be due to the higher concentration of GO flakes absorbing more copolymer chains on their surface and thus disrupting copolymer worm formation. As such, there is a lower concentration of polymer able to re-form into worms and a weaker gel is formed. Alternatively, it may be that the higher concentration of GO flakes simply prevents sphere-sphere fusion. In either case the presence of a population of spherical micelles can be observed in the TEM images for 1.8% w/w GO based on copolymer and above (Fig. 2f and Fig. S7, ESI<sup>†</sup>). Thus, there is an optimal concentration of GO for forming composite worm-gels for this particular G<sub>52</sub>-H<sub>130</sub>-GO formulation (Fig. 4d).

#### Influence of graphene oxide concentration on composite gel temperature response

The G<sub>52</sub>-H<sub>130</sub>-GO gels retained their thermo-responsive behaviour. Variable temperature rheology studies were conducted on the composite gels between 2 to 40 °C using an applied strain of 1.0% (Fig. 5). The 15% w/w G<sub>52</sub>-H<sub>130</sub> copolymer had a critical gelation temperature (CGT) of 14 °C on cooling and 15 °C on heating, as judged by the cross-over temperature for  $G'$  and  $G''$ .<sup>78</sup> The addition of GO initially decreased the CGT of the gels on cooling to 9 °C for 0.7% GO (Fig. 5b) and 6 °C for 1.5% GO (Fig. 5c). For both of these formulations, re-gelation was observed upon heating, with a CGT similar to that on cooling. This observation further supports that the reversible worm-to-sphere transition was retained in the presence of GO. The depression of the CGT can be attributed the association between the GO and copolymer making the gels more temperature stable.

With GO concentrations > 1.5% based on copolymer, the determined gel strengths were significantly decreased (Fig. 4c) and the CGT increased to approximately 20 °C on cooling and 24 °C upon re-heating (Fig. 5d). The higher CGT values and increased hysteresis supports that high GO contents hinder the degree of sphere-sphere fusion on heating but also result in



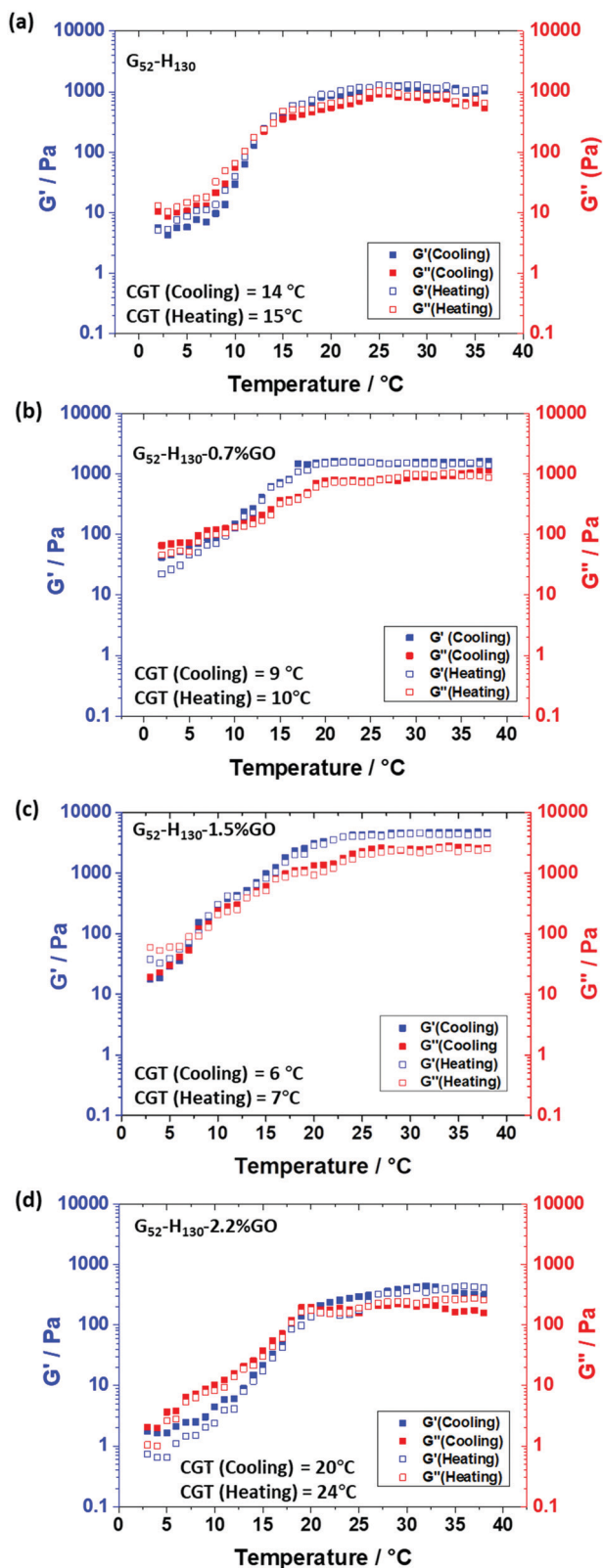


Fig. 5 Temperature-dependent oscillatory rheology studies obtained for aqueous dispersions of: (a) 15% w/w  $G_{52}$ - $H_{130}$ ; (b)  $G_{52}$ - $H_{130}$ -0.7% GO; (c)  $G_{52}$ - $H_{130}$ -1.5% GO; (d)  $G_{52}$ - $H_{130}$ -2.2% GO. The temperature was varied from 40 °C to 2 °C to 40 °C in 1 °C steps with 3 minutes equilibration at each step. All measurements were conducted at an angular frequency of 10 Hz and applied strain amplitude of 1.0%.

less stable gels on cooling due to the lower worm content of these gels.

In addition, the composite gels exhibited reversible de-gelation behavior over a number of heating-cooling cycles. For instance, the CGT of  $G_{52}$ - $H_{130}$  with 1.5% GO based on copolymer remained relatively consistent over eight cooling/heating cycles (Fig. S10, ESI<sup>†</sup>).

### Preparation of $G_{60}$ - $B_{70}$ -GO composite gels

In contrast to aqueous PGMA- $\text{PHPMA}$  copolymer dispersions, which undergo worm-to-sphere de-gelation transitions on cooling, methanolic PGMA- $\text{PBzMA}$  copolymer dispersions undergo the same transition on heating.<sup>66,75</sup> Thus, this alternative system was investigated as a complimentary route towards the formation of composite GO-containing worm-gels.

To facilitate mixing of G-B copolymer dispersions with GO, the dilution of a  $G_{60}$ - $B_{70}$  diblock copolymer dispersion from 20% w/w to 15% w/w was initially studied. When water was added to the methanolic dispersion to reduce the concentration, worm-like micelles were still present (Fig. 3a) and, as expected, the gel-strength was reduced from  $\sim 1.1$  kPa (Fig. S5b, ESI<sup>†</sup>) to  $\sim 0.9$  kPa (Fig. 6b) with a free-standing gel being retained (Fig. S6b, ESI<sup>†</sup>). Interestingly, dilution to 15% w/w with methanol caused the copolymer dispersion to no-longer be a free-standing gel (Fig. S6b, ESI<sup>†</sup>). Given this observation, and that the GO dispersions obtained were water-based, the initial composite  $G_{60}$ - $B_{70}$ -GO gels investigated were therefore methanol/water mixtures.

To prepare the  $G_{60}$ - $B_{70}$ -GO composite gels, the thermo-responsive behaviour of the  $G_{60}$ - $B_{70}$  gels on heating was utilised to facilitate good mixing of the copolymer and GO. As such, 20% w/w methanolic  $G_{60}$ - $B_{70}$  dispersions were heated to approximately 56 °C, where they formed free-flowing liquids (Fig. 1c). The copolymer was then mixed with pre-heated aqueous GO (keeping the copolymer concentration fixed at 15% w/w) and then allowed to cool to room temperature. The range of GO dispersion concentrations studied was initially screened by tube inversion tests. From these simple tests it was determined that the optimal GO concentration for gel re-formation after mixing was between 2.0% w/w and 3.8% w/w based on copolymer (Fig. 6a). As with  $G_{52}$ - $H_{130}$ - $x\%$  GO, re-gelation did not occur for samples containing relatively high GO concentrations ( $\leq 2.8\%$  w/w based on copolymer in this case). Zeta potential measurements were performed on dissolved  $G_{60}$  (Fig. S9c, ESI<sup>†</sup>) and a  $G_{60}$ - $B_{70}$  copolymer dispersion (Fig. S9d, ESI<sup>†</sup>). Due to the carboxylic acid end-groups from the PETTC RAFT agent, the  $G_{60}$  and  $G_{60}$ - $B_{70}$  dispersions showed a slightly negative charge. With the addition of GO the measured zeta showed an increase in negative charge with increasing GO concentration. For instance, the zeta potential measured for 3.1% GO based on (co)polymer is more negative than for 2.3% GO at all pH values studied (Fig. S9c and d, ESI<sup>†</sup>). This again suggests that GO sheets become associated with the (co)polymer during mixing.<sup>73</sup>

The  $G_{60}$ - $B_{70}$ -GO composite gels were investigated using oscillatory rheology. The storage modulus ( $G'$ ) of the prepared gels was



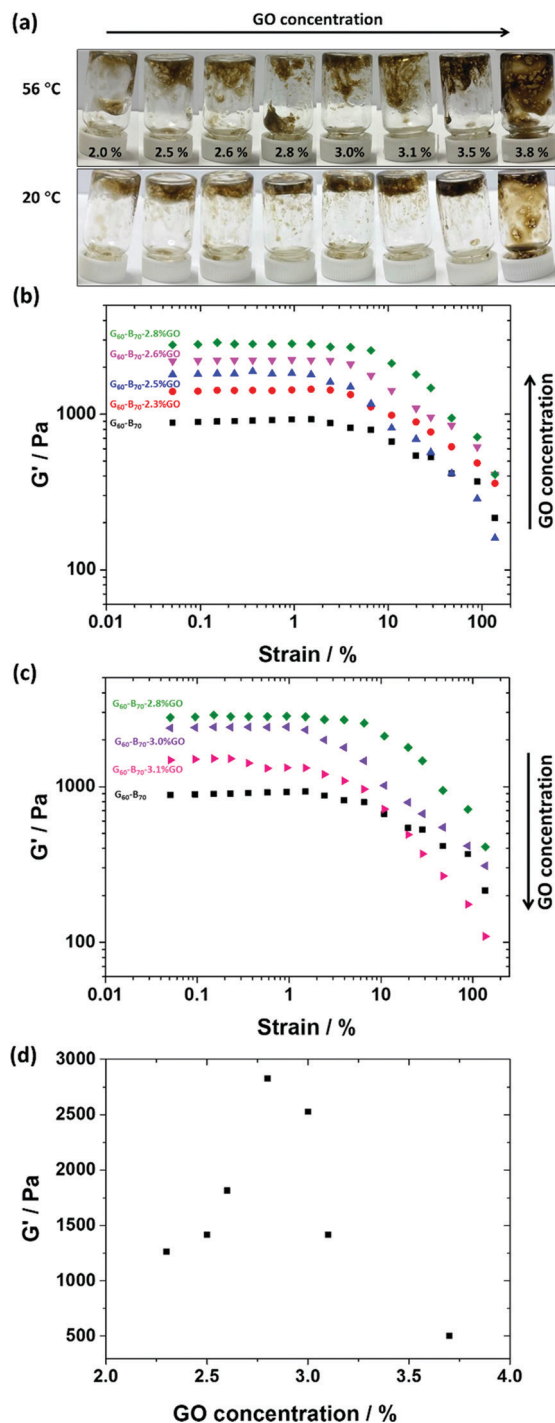


Fig. 6 (a) 15% w/w PGMA<sub>x</sub>-PBzMA<sub>y</sub> block copolymer dispersions after mixing with aqueous graphene oxide between 2.0 and 3.8% w/w based on copolymer. The upper image in (a) shows samples after mixing at 56 °C and the lower image shows samples after equilibration overnight at room temperature. Storage modulus ( $G'$ ) versus % strain for (b) 15% w/w G<sub>60</sub>-B<sub>70</sub> with increasing GO concentration (0–2.8% w/w) and (c) 15% w/w G<sub>60</sub>-B<sub>70</sub> with decreasing GO concentration (2.8–3.1% w/w). (d) Average storage modulus measured between 0.01 and 2% strain for different GO-containing G<sub>60</sub>-B<sub>70</sub> copolymer dispersions. All measurements were conducted at a frequency of 10 Hz, a strain of 1.0% and a controlled temperature of 25 °C.

measured as a function of % strain with varying GO contents (Fig. 6b–d). The moduli in the plateau region are strain independent. As the GO concentration increased from 0 to 2.8% w/w based on copolymer,  $G'$  gradually increased to a maximum of approximately 2.7 kPa, which is nearly 3 times greater than for 15% G<sub>60</sub>-B<sub>70</sub> without the addition of GO (Fig. 6b).<sup>80</sup> TEM images of the G<sub>60</sub>-B<sub>70</sub>-GO composite gels (Fig. 3b–e) show that after the heating-mixing-cooling cycle, copolymer worms are clearly present and that they are associated with the relatively large GO flakes present in the image. As with the G<sub>52</sub>-H<sub>130</sub>-GO composites, the association between GO and the G<sub>60</sub>-B<sub>70</sub> copolymer worms is probably providing reinforcement of the re-formed worm-gels.

Increasing the GO content further for led to  $G'$  dramatically decreasing (Fig. 6c). For example, G<sub>60</sub>-B<sub>70</sub>-3.1% GO had a gel strength of ~1.4 kPa compared to 2.7 kPa G<sub>60</sub>-B<sub>70</sub>-2.8% w/w GO. TEM images for higher GO loadings (Fig. 3) indicate the formation of shorter worms and a small population of spheres in comparison to lower GO loadings. Thus, the prevention of sphere-sphere fusion with higher GO contents is the proposed reason as to why the gels with relatively high GO loadings are weaker. These observations correlate with those made for G<sub>52</sub>-H<sub>130</sub>-GO and thus this seems to be a general mechanism of gel reinforcement/weakening for worm-gel/GO mixtures.

To investigate whether the presence of water was a major factor in the observations made, wholly methanolic copolymer/GO composites were investigated. To facilitate this, freeze-dried GO powder was homogeneously dispersed in methanol using sonication for 40 min. Pre-heated methanolic GO dispersions were then added to G<sub>60</sub>-B<sub>70</sub> at approximately 56 °C whilst keeping the copolymer concentration at 15% w/w.

The addition of GO allowed free-standing gels to form on cooling to room temperature, with an optimal GO concentration between 3.2% w/w and 4.1% w/w based on tube inversion tests (Fig. S11a, ESI†). It is noteworthy that the presence of GO allows gel reformation on cooling and in fact allows the formation of free-standing gels at 15% w/w copolymer for the dispersion diluted with GO dispersed in methanol (which is not the case in the absence of GO). The measured gel strength (Fig. S11b and c, ESI†) showed a similar trend to the formulations described above, with the expected increase and subsequent decrease in gel strength with increasing GO concentration. However, the recorded values for the wholly methanolic system were less than for G<sub>60</sub>-B<sub>70</sub>-GO prepared using aqueous GO. As expected, the recorded TEM images (Fig. S12, ESI†) clearly show an association between the copolymer worms and the GO flakes, indicating the interactions between these particles is not only observed in water. Furthermore, there is clear evidence for the limited amount of worm-reformation with high GO loadings, shown by a high number density of ~30 nm copolymer spheres and lack of long worms in the TEM image for G<sub>60</sub>-B<sub>70</sub>-4.6% GO (Fig. S12d, ESI†).

## Conclusions

Block copolymer worm-gels containing GO were prepared. For PGMA<sub>52</sub>-PHPMA<sub>130</sub> worm gels, cooling induces a worm-to-



sphere de-gelation transition which allows facile mixing with aqueous GO dispersions. On re-heating to room temperature, GO-containing worm gels are formed for 15% w/w copolymer and GO contents up to  $\sim 1.8\%$  w/w based on copolymer. A maximum gel strength of  $\sim 4$  kPa was recorded at a GO content of 1.5% w/w. Aqueous electrophoresis and TEM measurements indicate that there is absorption of the copolymer onto the GO flakes which then allows reinforcement of the gel on worm-reformation. However, too much GO disrupts the worm reformation process, causing a lower measured gel strength and thus an optimal GO concentration. Similar observations were made for PGMA<sub>60</sub>-PBzMA<sub>70</sub> copolymer worms prepared in methanol. In this case, heating the copolymer dispersions induces the worm-to-sphere transition, with composite gels forming on mixing with either aqueous or methanolic GO and cooling to room temperature. For this system the optimal GO concentration was slightly higher in comparison to the aqueous PGMA<sub>52</sub>-PHPMA<sub>130</sub>-GO composite gels. Nevertheless, the same mechanistic process for gel-strengthening and subsequent weakening with increasing GO loadings is observed. Thus, this generic process allows reinforced nanocomposite block copolymer worms gels to be readily prepared by the addition of relatively low concentrations of GO. It is likely that this route is not specific to GO as a re-enforcing nanomaterial and it is expected that this novel class of composite gel has potential applications in many areas of materials science.<sup>81–83</sup>

## Conflicts of interest

There are no conflicts to declare.

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