

Journal of Intelligent Material Systems and Structures

<http://jim.sagepub.com/>

Effect of the γ -radiation on the properties of epoxy-based shape memory polymers

Jinsong Leng, Fang Xie, Xuelian Wu and Yanju Liu

Journal of Intelligent Material Systems and Structures published online 20 September 2013

DOI: 10.1177/1045389X13504474

The online version of this article can be found at:

<http://jim.sagepub.com/content/early/2013/09/16/1045389X13504474>

Published by:



<http://www.sagepublications.com>

Additional services and information for *Journal of Intelligent Material Systems and Structures* can be found at:

Email Alerts: <http://jim.sagepub.com/cgi/alerts>

Subscriptions: <http://jim.sagepub.com/subscriptions>


Reprints: <http://www.sagepub.com/journalsReprints.nav>

Permissions: <http://www.sagepub.com/journalsPermissions.nav>

>> [OnlineFirst Version of Record](#) - Sep 20, 2013

[What is This?](#)

Effect of the γ -radiation on the properties of epoxy-based shape memory polymers

Journal of Intelligent Material Systems and Structures
0(0) 1–8
© The Author(s) 2013
Reprints and permissions:
sagepub.co.uk/journalsPermissions.nav
DOI: 10.1177/1045389X13504474
jim.sagepub.com


Jinsong Leng¹, Fang Xie², Xuelian Wu^{1,3} and Yanju Liu²

Abstract

In this article, the epoxy-based shape memory polymers were exposed to simulated γ -radiation environments up to 140 days for an accelerated irradiation. The influence of γ -radiation on thermal and mechanical properties was evaluated by differential scanning calorimetry, dynamic mechanical analysis, and tensile test. The glass transition temperatures (T_g) determined by differential scanning calorimetry and dynamic mechanical analysis decreased no more than 10%, and the shape recovery speed became a little faster after γ -radiation of 1×10^5 Gy. The tensile strength and elastic modulus could, respectively, maintain 26 MPa and 1.36 GPa after being irradiated by 1×10^6 Gy radiation, showing great potential in aerospace structural materials.

Keywords

Shape memory polymer, epoxy-based shape memory polymer, γ -radiation

Introduction

Shape memory polymers (SMPs) are materials responding to external stimulus to change their shapes after a programmed process (Behl and Lendlein, 2007). The shape memory effect (SME) of SMPs can be triggered by heat, electric field, light, magnetic field, moisture, and even some chemical solutions (Gall et al., 2005; Huang et al., 2005; Luo and Mather, 2010; Mohr et al., 2006; Small et al., 2005). In recent researches, the mechanisms of the SME have been investigated deeply. Several deformation constitutive models were developed to describe the thermally triggered SME (Ge et al., 2012; Qi et al., 2008), thermo- or chemo-responsive SME (Huang et al., 2012), multi-SMEs (Yu et al., 2012), and so on. The mechanical properties of SMPs vary over a wide range due to their diverse and tunable chemical structures. Main chemistries raw materials of SMPs include polyurethane (Tobushi et al., 1992), polystyrene (Xu et al., 2010), epoxy (Leng et al., 2009b; Rousseau and Xie, 2010), polycaprolactone (PCL) (Alvarado-Tenorio et al., 2011), poly(methyl methacrylate) (PMMA) (Zhao et al., 2011), and perfluorosulfonic acid (PFSA) (Xie, 2010; Xie et al., 2010). SMPs have common advantages of traditional polymers, such as lower cost and weight and less complicated processing and programming. Compared with shape memory alloys, SMPs possess higher deformation capability and higher recoverable strain levels (Lendlein and Kelch, 2002). The

fast programming process and biological compatibility of some kinds of SMPs have enabled their applications in medical devices (Lendlein and Langer, 2002; Nardo et al., 2011). Potential application for SMPs also lies in smart textiles (Hu et al., 2009), sensors (Kunzelman et al., 2008), actuators (Ma et al., 2013), and the deployable space structures (Barrett et al., 2007; Lan et al., 2009).

As an emerging class of smart structure and material, SMPs and their adaptability in the space environment attract more and more attention. Materials used for the deployable space structures must have good adaptability to the space environment (e.g. high vacuum, ionizing radiation, ultraviolet (UV) radiation, atomic oxygen atmosphere, plasma, meteoroids, debris, thermal cycles, and electrostatic charging.). Among recent research about SMP performances in the simulated in-service environments, several works have reported the effect of

¹Centre for Composite Materials and Structures, Harbin Institute of Technology, Harbin, P.R. China

²Department of Astronautical Science and Mechanics, Harbin Institute of Technology, Harbin, P.R. China

³School of Mechanical Engineering, Jiangsu University, Zhenjiang, P.R. China

Corresponding author:

Jinsong Leng, Centre for Composite Materials and Structures, No. 2 Yikuang Street, Science Park of Harbin Institute of Technology, Harbin, P.O. Box 3011, P.R. China.
Email: lengjs@hit.edu.cn

UV radiation (Fulcher et al., 2010; Lu et al., 2011; Tandon et al., 2010; Xu et al., 2011), xenon arc radiation (Tandon et al., 2009), electron beam radiation (Youssef, 2009), and so on. So far, some works have already been carried out to investigate the durability of the SMPs under γ -radiation, for example, on the thermoplastic bioabsorbable polymer-based SMPs in the medical level (Paakinaho et al., 2012; Yakacki et al., 2008). However, the SMPs have not fully exerted their technological potential applications in spatial deployment structure. In this case, there are numerous works to be done on γ -radiation performances of thermosetting SMP use in the space environment.

The γ -radiation is an electromagnetic radiation of high frequency, and consequently high energy. The electron and proton radiation degradation of materials are the primary factors to be concerned in addition to atomic oxygen and UV radiation. The nominal orbital environmental parameter of the space station truss structure for 30 years electron radiation is 10^4 to 10^5 Gy (Tmopking et al., 1988), and the corresponding radiation rate is around 6.34×10^{-4} to 6.34×10^{-3} Gy/min. Accelerated irradiation test of up to 140 days exposure to γ -radiation (up to 1×10^6 Gy) was conducted on the epoxy-based SMPs in this study to investigate the effect of γ -radiation on them. Differential scanning calorimetry (DSC), dynamic mechanical analysis (DMA), tensile test, and a “U”-shape bending test were carried out to measure the relevant thermomechanical properties and shape memory behaviors of the epoxy-based SMP before and after γ -radiation. All the radiation tests were conducted at room temperature in this current research. Given the complex and volatile space environments, it is only a baseline assessment of the durability of epoxy-based SMPs being considered as structural material substrates. The SMP behaviors under other individual factor (e.g. extremely high- or low-temperature condition) or the synergetic and additive effect of several factors will be conducted in our future work.

Experimental

The polymer matrix consists of an epoxy base resin, a curing hardener, and an active linear epoxy monomer. The linear monomer is composed of a long linear chain of C–O bonds. The epoxy resin and the linear monomer have an epoxy group number ratio of 1:1 in one molecule, a molecular weight ratio of 2:5, and a density ratio of 1:1. The resin and hardener are mixed with a weight ratio of 1:0.8, and linear monomer contents of 0, 5, 10, and 15 wt% were used for specimens SMP1, SMP2, SMP3, and SMP4, respectively. To obtain a bubble-free prepolymer, mixtures of epoxy resin, hardener, and linear epoxy monomer were degassed at a vacuum oven for 15 min and then injected into preheated glass molds

composed of two slides separated by a silica gel rod spacer. After that, a three-step curing condition was performed at 80°C for 3 h, 100°C for 3 h, and 150°C for 5 h. After the curing process, the thermoset cross-linked network of the SMP products was obtained. The composition and preparation of the epoxy-based SMPs similar to this work have been reported in our previous study (Leng et al., 2009b).

The γ -irradiation is a highly penetrating technique that uses γ -rays from a radioactive Cobalt 60 (^{60}Co) isotope. In this study, the accelerated weathering conditioning was performed in a ^{60}Co isotope exposure chamber under a limited atmospheric condition. During irradiation, the temperature in the exposure chamber was controlled to room temperature (25°C), and dosages of irradiation were 1×10^5 Gy and 1×10^6 Gy. Due to the limitation of the experimental time, the radiation rate is set to 5 Gy/min for accelerated irradiation, which is much higher than the nominal value. After exposure, the specimens of epoxy-based SMPs were examined for any visible damage and apparent color changes. Irradiation was performed by Heilongjiang Academy of Sciences (Harbin, China).

The DSC (NETZSCH DSC 204, Selb, Germany) measurements were employed to characterize the changes in thermal properties of the SMPs from 0°C to 200°C at a heating or cooling rate of 10°C/min. The glass transition temperatures (T_g) were identified as the shift in the baseline of the DSC curves.

The DMA (NETZSCH DMA Q800) experiments were conducted in a tensile mode. Samples with dimensions of $30 \times 5 \times 1 \text{ mm}^3$ were prepared. All runs were performed at 5 Hz, and at temperature ranges from 25°C to 250°C at a heating rate of 5°C/min. The glass transition temperatures (T_g) were identified at the tan delta ($\tan \delta$) peak values in the DMA tests.

Isothermal quasi-static tension test was carried out on a Zwick Z010 (Zwick GmbH, Ulm, Germany) equipped with an extensometer at room temperature (20°C). Standard samples (ASTM D638, Type IV) for tensile test were cut by a laser cutter from the prepared SMP sheet and stretched at an elongation rate of 1 mm/min.

The shape memory behaviors of the SMPs before and after the irradiation were demonstrated through a “U”-shape bending test in the same condition. To compare shape memory behaviors before and after γ -radiation, the samples were cut from thin polymer sheets into rectangular strips (about $80 \times 10 \times 1 \text{ mm}^3$), using it as the permanent shape. The straight bar-shaped samples were heated up to $T_g + 40^\circ\text{C}$ and held for 20 min for full heating, and then samples became elastic and were deformed to a “U”-shape easily under an external force. The deformed samples were cooled to room temperature to achieve a stiff state of SMPs, so that the temporary shape could be fixed when the samples were unloaded. After this programming

process, the sample maintained the “U” shape without loading. And the shape recovery process under different actuating temperatures (T_g , $T_g + 10^\circ\text{C}$, $T_g + 20^\circ\text{C}$, $T_g + 30^\circ\text{C}$, $T_g + 40^\circ\text{C}$) was recorded by a video recorder. Shape recovery angle is determined by measuring the angle between the straight ends of the bent specimen. The shape recovery ratio (R_r) was calculated as $R_r = \theta_r/\theta_b$, where θ_b represents the bending angle after cooling and θ_r the recovered angle. In addition, shape recovery speed was evaluated by the full recovery time T_r .

Results and discussion

Effect of γ -radiation on surface profile

As shown in Figure 1, the SMP color changes from transparent to yellow-green and dark-brown when exposed to increasing γ -radiation, but there is no visible damage to all the irradiated specimens.

Effect of γ -radiation on thermal properties

The equilibrium properties of epoxy-based SMPs determined by DSC and DMA tests are listed in Table 1 for a clear comparison. The glass transition temperatures (T_g) were identified as the shift in the baseline of the

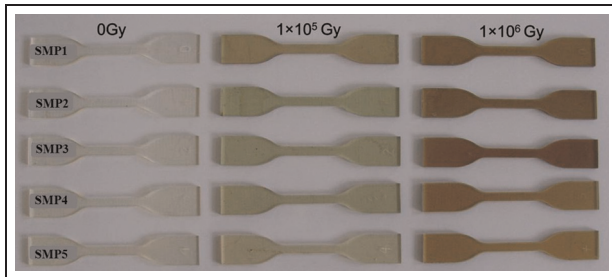


Figure 1. Photographs of epoxy-based SMPs before and after γ -radiation.

SMP: shape memory polymer.

DSC curves. The T_g of the samples confirmed by the DSC curves decreased by 7°C – 15°C with an increase in irradiation dose. Since T_g represents the onset of cooperative segmental motion, widely spaced cross-links will produce only a slight restriction on this motion. As the cross-linking density is increased, the restriction on molecular mobility becomes substantial, and much higher energy will be required to induce segmental motion (T_g increases) (Ebewele, 2000). So, we could speculate that the cross-linking density of the networks decreased after a complex chemical reaction process by the irradiation of γ -ray.

Effect of γ -radiation on dynamic mechanical properties

DMA is a key technique to relate change in molecular motion with macro-properties for polymer materials. Figure 2 presents the storage modulus and $\tan\delta$ as a function of temperature. The relationship between the dynamic stress and the corresponding strain provides the complex modulus, a real part (storage modulus) and imaginary part (loss modulus), and $\tan\delta$ presents the ratio of the loss modulus to the storage modulus. It could be seen clearly that these SMP samples had two different elastic states with the change in temperature: glassy state and rubbery state. The region among two states is glass transition process. A large and sharp decrease in the storage modulus around glass transition could be seen in Figure 2 both before and after γ -radiation, which is crucial for a good shape memory material. The elastic ratio that is defined as the glass-to-rubber modulus ratio (E_g/E_r) is often quoted to estimate the magnitude of the change in modulus when the polymer undergoes shape recovery. A large elastic modulus ratio is beneficial for showing good shape memory capability, since it gives larger shape fixity upon cooling and realizes a larger strain with a small stress at high temperature. As is shown in Figure 2,

Table 1. Equilibrium properties of epoxy-based SMPs determined by DSC and DMA tests.

Samples	Total radiation dosage (Gy)	T_g by DSC ($^\circ\text{C}$)	T_g by $\tan\delta$ ($^\circ\text{C}$)	Amplitude of $\tan\delta$ at T_g
SMP1	0	125.3 ± 2.3	162.8 ± 2.1	1.31
	1×10^5	119.6 ± 1.8	160.0 ± 1.1	1.32
	1×10^6	115.5 ± 1.9	150.3 ± 0.9	1.26
SMP2	0	113.8 ± 0.6	150.7 ± 1.3	1.22
	1×10^5	110.7 ± 1.2	147.8 ± 0.6	1.27
	1×10^6	106.4 ± 1.3	142.1 ± 0.8	1.59
SMP3	0	101.9 ± 0.9	137.9 ± 0.8	1.17
	1×10^5	100.3 ± 1.0	133.5 ± 1.9	1.34
	1×10^6	94.8 ± 2.5	131.5 ± 0.7	1.60
SMP4	0	73.1 ± 1.4	110.3 ± 1.5	1.26
	1×10^5	69.3 ± 1.7	105.3 ± 1.2	1.34
	1×10^6	58.2 ± 0.8	100.0 ± 0.7	1.75

SMP: shape memory polymer; DSC: differential scanning calorimetry; DMA: dynamic mechanical analysis.

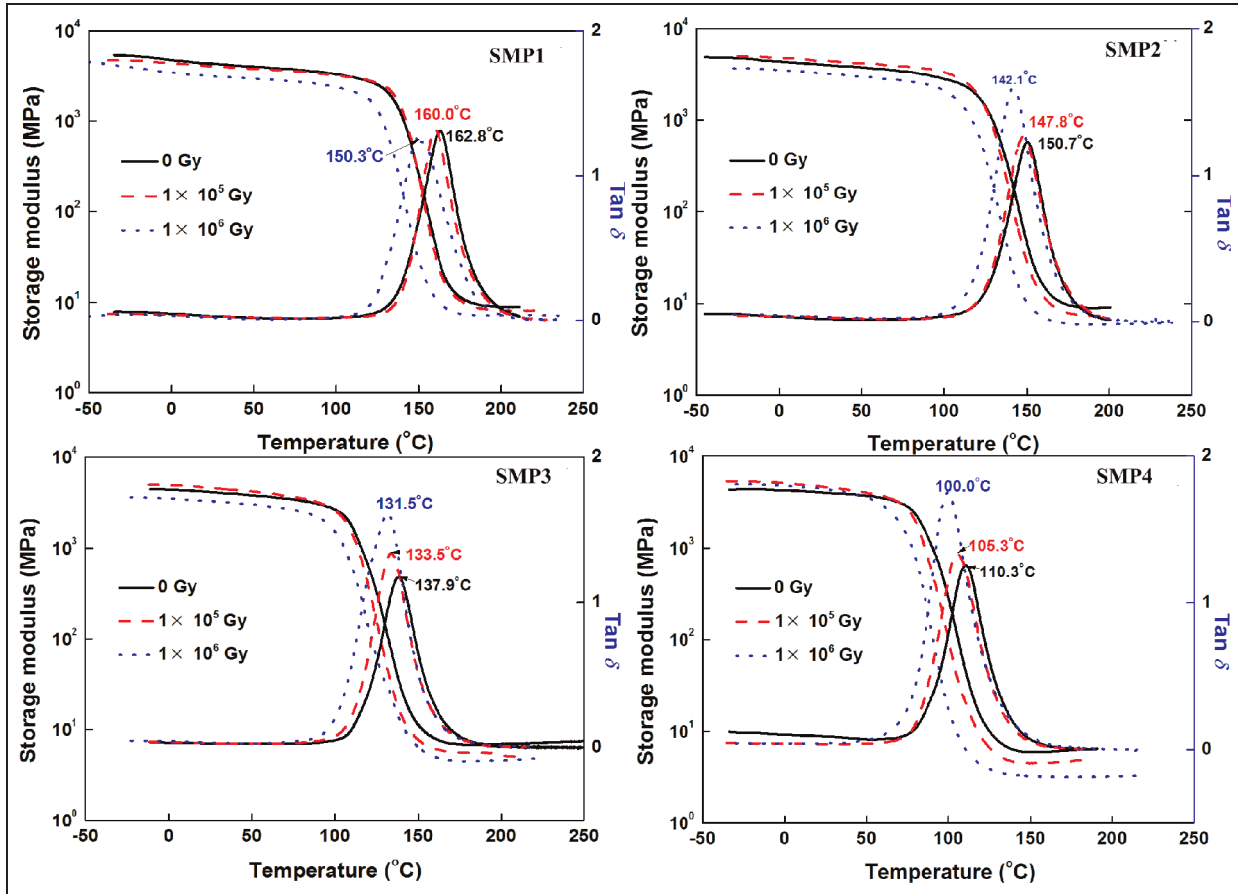


Figure 2. DMA curves of the epoxy-based SMPs before and after γ -radiation. DMA: dynamic mechanical analysis; SMP: shape memory polymer.

γ -radiation had no obvious effect on the elastic modulus ratio in this study. After γ -radiation of 1×10^5 Gy, the storage modulus and $\tan \delta$ curves slightly move to the lower temperature. As the storage modulus reveals the stiffness of the material, it could be noted that the stiffness of the polymer is stable after a low dose of γ -radiation. After γ -radiation of 1×10^6 Gy, the transition region of the storage modulus curves and the $\tan \delta$ peaks moved to a lower temperature for about 5°C – 12°C , less than 10%, also predicting the decrease of T_g . At the same time, the $\tan \delta$ peaks in Figure 2 became higher and broader with the incremental radiation. The amplitude of $\tan \delta$ peaks (Table 1) indicates the damp property of the polymer, and the width of the curve reveals the homogeneity of the segments in the network. In the case of SMP3, the $\tan \delta$ peak value increased from 1.26 (0 Gy) to 1.75 (1×10^6 Gy). A higher $\tan \delta$ in general indicates that materials are more resistant to impact (or cracking) through better dissipation of mechanical energy (Lu et al., 2011). This phenomenon was gradually more obvious with the increasing linear monomer from SMP1 to SMP4. Then, it is obvious that after γ -radiation of 1×10^6 Gy, the homogeneity of the segments in the network was

weakened. It is noteworthy that T_g values obtained from the DMA ($\tan \delta$) were 30°C – 40°C higher than those from the DSC test. The method of determining T_g by DMA can cause disagreements, as at least five ways are in current use. Depending on the industry standards or background of the operator, the peak or onset of the $\tan \delta$ curve, the onset of the storage modulus decrease, or the onset or peak of the loss modulus curve may be used. The values obtained from these methods can differ by up to 25°C from each other for the same run. In addition, a 10°C – 20°C difference from the DSC is also seen in many materials (Menard, 2003). Since the value obtained from the peak of $\tan \delta$ curve is generally the highest one among the above methods, the difference in T_g values obtained from the DMA ($\tan \delta$) and DSC test is reasonable.

Effect of γ -radiation on quasi-static mechanical properties

To investigate the effect of γ -radiation on mechanical behaviors, tensile tests for the epoxy-based SMP samples before and after irradiation were all carried out at 20°C (Figure 3). Compared with non-irradiation

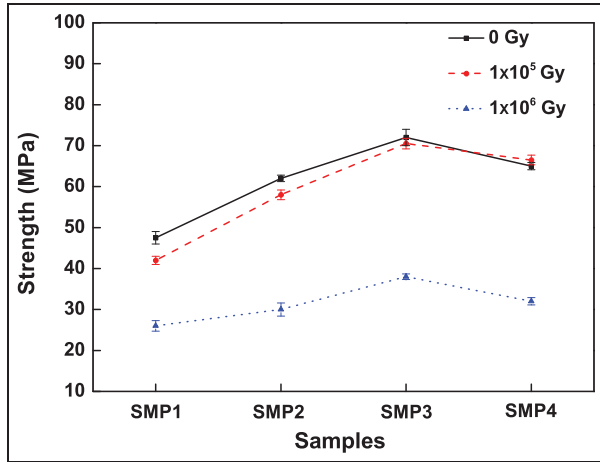


Figure 3. Tensile strength of epoxy-based SMPs before and after γ -radiation.

SMP: shape memory polymer.

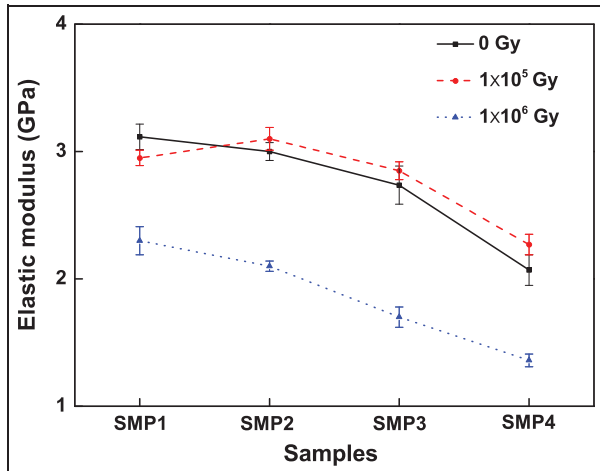


Figure 4. Elastic modulus of epoxy-based SMPs before and after γ -radiation.

SMP: shape memory polymer.

samples, the tensile strength changed little after irradiation of 1×10^5 Gy. Tensile strength of SMP1, SMP2, and SMP3 decreased slightly with the rates from 2.2% to 11%, while that of SMP4 increased slightly with a rate of 2.3%. However, after γ -radiation of 1×10^6 Gy, the tensile strength of all samples reduced significantly, with the rates of 11%–51%.

Elastic modulus of epoxy-based SMPs before and after γ -radiation is shown in Figure 4. After irradiation of 1×10^5 Gy, only SMP1 saw a 5.3% decrease in elastic modulus, while the others increased by 2%–9.6%, and they all changed slightly. After irradiation of 1×10^6 Gy, the elastic modulus decreased significantly with the rates of 16%–47%. These results suggest that elastic modulus first increased slightly and then reduced significantly with an increase in γ -radiation.

Currently, there are no specific or uniform evaluation indicators for SMPs under radiation. In practice, the main method is to compare the mechanical properties of structural materials for aerospace before and after irradiation. In this article, the epoxy-based SMPs showed good resistance to irradiation of 1×10^5 Gy, and after being irradiated by 1×10^6 Gy radiation, the tensile strength and elastic modulus were above 26 MPa and 1.36 GPa, respectively, showing great potential in aerospace structural materials.

Effect of γ -radiation on shape memory properties

The shape recovery time under different actuating temperatures (T_g , $T_g + 10^\circ\text{C}$, $T_g + 20^\circ\text{C}$, $T_g + 30^\circ\text{C}$, and $T_g + 40^\circ\text{C}$) of the epoxy-based SMPs before and after γ -radiation is shown in Figure 5. It is worth noting that below 1×10^5 Gy γ -radiation, the shape recovery speed above the respective glass transition temperatures was faster than that without irradiation. As the shape switching temperature was set to match with T_g , when T_g decreased after irradiation, the actuating temperature was correspondingly high. However, the recovery time became a little longer for the samples exposed to a dose below 1×10^6 Gy γ -radiation probably because the recovery force became much smaller due to the damage of large amount irradiation. But generally speaking, the recovery time after irradiation fluctuated slightly around the time before irradiation, and when the actuating temperature was high enough, for example, $T_g + 40^\circ\text{C}$, the recovery time was quite similar.

The shape recovery process of SMP4 at 100°C is shown in Figure 6, as an example. According to the experimental results, the shape recovery ratio of all the samples were greater than 95%, which means that the samples recovered nearly the full initial shape and γ -radiation has little effect on shape recovery ratio. Therefore, after γ -irradiation doses of 1×10^5 Gy and 1×10^6 Gy, the shape memory properties of epoxy-based SMPs are basically stable.

Conclusion

In this article, the thermomechanical properties and shape memory behaviors of the epoxy-based SMPs before and after γ -radiation were investigated systematically. The radiation source was ^{60}Co , and the radiation dosage was 1×10^6 Gy for the first time. The influence of γ -radiation on thermal and mechanical properties of the SMPs was characterized by means of DSC, DMA, and tensile test. The shape memory behaviors of the SMPs before and after irradiation were demonstrated through a “U”-shape bending test. Although this work is only a baseline assessment of the durability of epoxy-based SMPs under γ -radiation, the results are quite instructive in selecting and evaluating material for aerospace structures. We may draw the

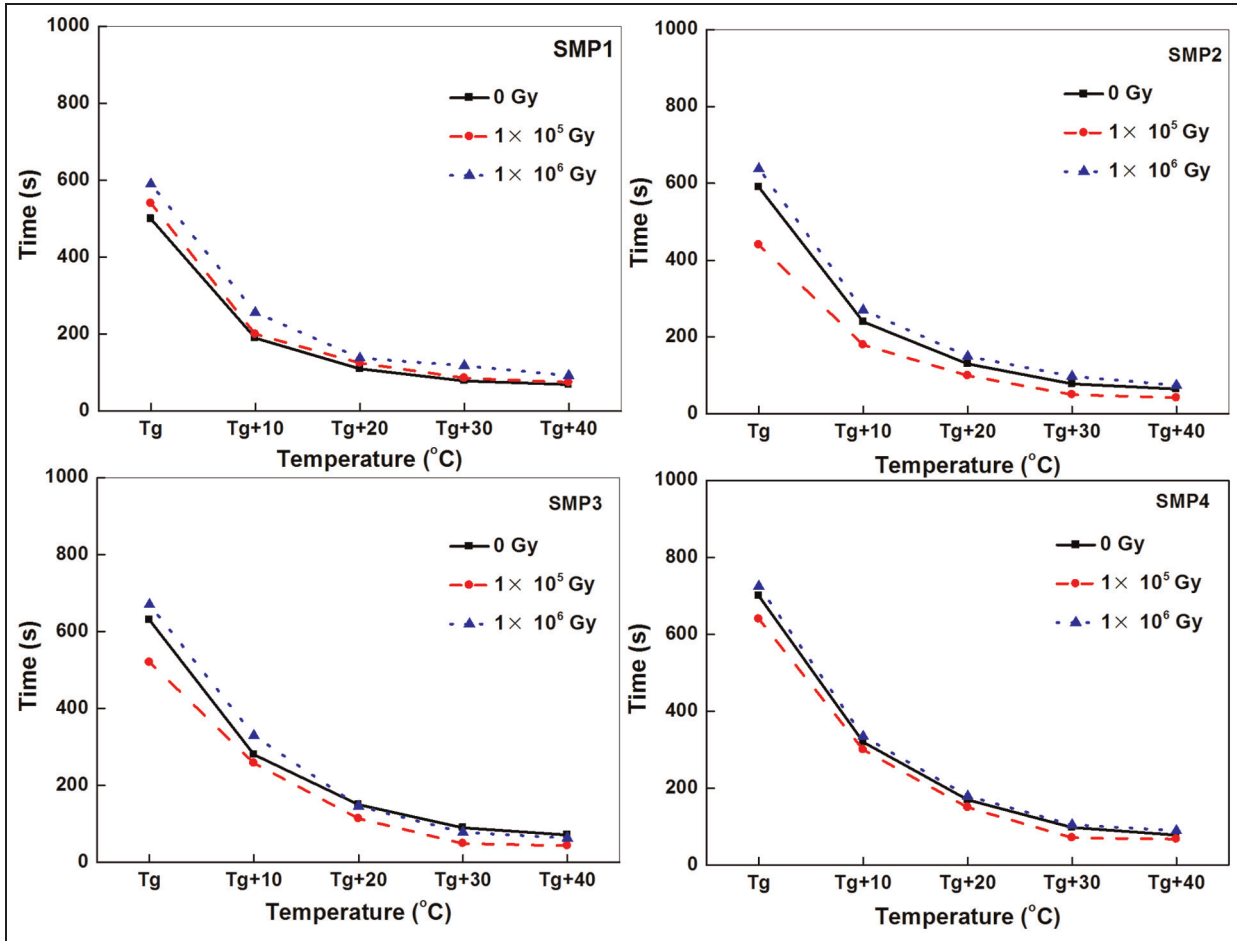


Figure 5. Time for recovery of the epoxy-based SMPs before and after γ -radiation. SMP: shape memory polymer.

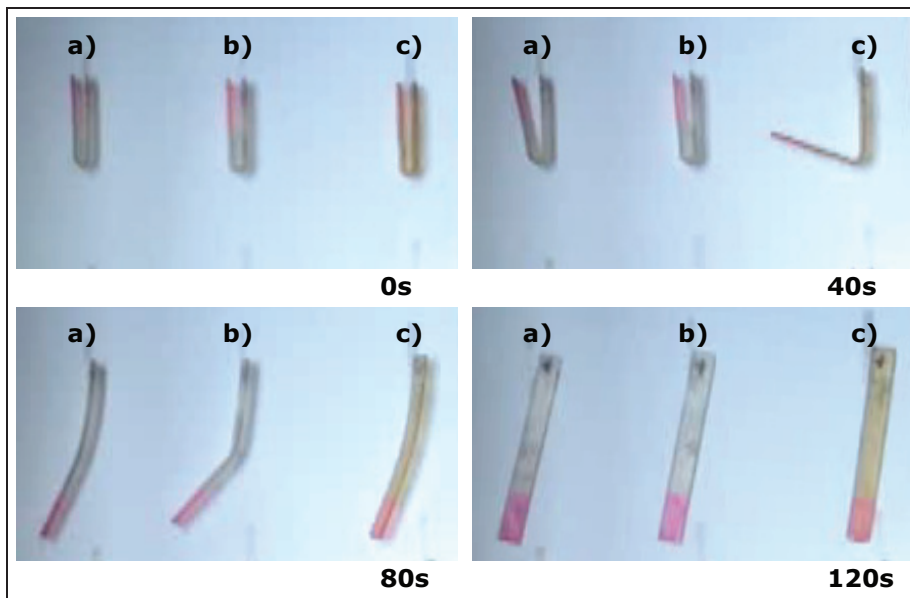


Figure 6. Shape recovery process of SMP4 at 100°C. (a) Before radiation, (b) after 1×10^5 Gy, and (c) after 1×10^6 Gy. SMP: shape memory polymer.

following conclusions from this: (1) T_g of the epoxy-based SMPs confirmed by the DSC curves decreased by 7°C–15°C with an increase in irradiation dose and the DMA test showed that the homogeneity of the segments in the network was weakened after exposure to the large amount of γ -radiation (1×10^6 Gy). (2) The stiffness of the polymer is stable under a low dose of γ -radiation (1×10^5 Gy), while the tensile strength and elastic modulus could, respectively, maintain above 26 MPa and 1.36 GPa under 1×10^6 Gy γ -radiation. (3) The shape recovery ratios of the SMPs were all above 95%. The recovery speeds of the irradiated specimens fluctuated slightly around that of non-irradiated ones, and the speed had little difference when the actuating temperature was high enough above T_g . The acquired good properties under irradiation successfully expanded the applications of epoxy-based SMPs in aerospace and other harsh environments.

Funding

This work is supported by the National Natural Science Foundation of China (Grant No.11225211, No. 11272106).

References

- Alvarado-Tenorio B, Romo-Urbe A and Mather PT (2011) Microstructure and phase behavior of POSS/PCL shape memory nanocomposites. *Macromolecules* 44(14): 5682–5692.
- Barrett R, Taylor R, Keller PN, et al. (2007) Design of a solar array to meet the standard bus specification for operation responsive space. In: *48th AIAA/ASME/ASCE/AHS/ASC structures, structural dynamics, and materials conference*, vol. 8, Honolulu, HI, 23–26 April. Waikiki, HI: *American Institute of Aeronautics and Astronautics Inc.*
- Behl M and Lendlein A (2007) Shape-memory polymers. *Materials Today* 10(4): 20–28.
- Ebewele RO (2000) *Polymer Science and Technology*. Boca Raton, FL: CRC Press.
- Fulcher JT, Lu YC, Tandon GP, et al. (2010) Thermomechanical characterization of environmentally conditioned shape memory polymer using nanoindentation. In: *Behavior and mechanics of multifunctional materials and composites*, vol. 7644, San Diego, CA, 8–11 March.
- Gall K, Yakacki CM, Liu YP, et al. (2005) Thermomechanics of the shape memory effect in polymers for biomedical applications. *Journal of Biomedical Materials Research Part A* 73(3): 339–348.
- Ge Q, Luo XF, Rodriguez ED, et al. (2012) Thermomechanical behavior of shape memory elastomeric composites. *Journal of the Mechanics and Physics of Solids* 60(1): 67–83.
- Hu JL, Dong ZE, Liu Y, et al. (2009) The investigation about the shape memory behavior of wool. In: Vincenzini P and Paradiso R (eds) *Smart Textiles*, vol. 60. Stafa-Zurich: Trans Tech Publications Ltd, pp. 1–10.
- Huang WM, Yang B, An L, et al. (2005) Water-driven programmable polyurethane shape memory polymer: demonstration and mechanism. *Applied Physics Letters* 86(11): 114105.
- Huang WM, Zhao Y, Wang CC, et al. (2012) Thermo/chemo-responsive shape memory effect in polymers: a sketch of working mechanisms, fundamentals and optimization. *Journal of Polymer Research* 19: 9952.
- Kunzelman J, Chung T, Mather PT, et al. (2008) Shape memory polymers with built-in threshold temperature sensors. *Journal of Materials Chemistry* 18(10): 1082–1086.
- Lan X, Liu YJ, Lv HB, et al. (2009b) Fiber reinforced shape-memory polymer composite and its application in a deployable hinge. *Smart Materials & Structures* 18(2): 024002.
- Lendlein A and Kelch S (2002) Shape-memory polymers. *Angewandte Chemie: International Edition* 41(12): 2034–2057.
- Lendlein A and Langer R (2002) Biodegradable, elastic shape-memory polymers for potential biomedical applications. *Science* 296(5573): 1673–1676.
- Leng JS, Lan X, Liu YJ, et al. (2009a) Electroactive thermoset shape memory polymer nanocomposite filled with nanocarbon powders. *Smart Materials & Structures* 18(7): 074003.
- Leng JS, Wu XL and Liu YJ (2009b) Effect of a linear monomer on the thermomechanical properties of epoxy shape-memory polymer. *Smart Materials & Structures* 18(9): 095031.
- Leng JS, Zhang DW, Liu YJ, et al. (2010) Study on the activation of styrene-based shape memory polymer by medium-infrared laser light. *Applied Physics Letters* 96: 111905.
- Lu YC, Fulcher JT, Tandon GP, et al. (2011) Microscale thermomechanical characterization of environmentally conditioned shape memory polymers. *Polymer Testing* 30(5): 563–570.
- Luo XF and Mather PT (2010) Conductive shape memory nanocomposites for high speed electrical actuation. *Soft Matter* 6(10): 2146–2149.
- Ma MM, Guo L, Anderson DG, et al. (2013) Bio-inspired polymer composite actuator and generator driven by water gradients. *Science* 339: 186–189.
- Menard KP (2003) Thermomechanical and Dynamic Mechanical Analysis. In: Lobo H and Bonilla JV (eds), *Handbook of Plastics Analysis*, Chapter 6. New York: Marcel Dekker, Inc.
- Mohr R, Kratz K, Weigel T, et al. (2006) Initiation of shape-memory effect by inductive heating of magnetic nanoparticles in thermoplastic polymers. *Proceedings of the National Academy of Sciences of the United States of America* 103(10): 3540–3545.
- Nardo LD, Bertoldi S, Tanzi MC, et al. (2011) Shape memory polymer cellular solid design for medical applications. *Smart Materials & Structures* 20(3): 035004.
- Paakinaho K, Heino H, Pelto M, et al. (2012) Programmed water-induced shape-memory of bioabsorbable poly(D,L-lactide): activation and properties in physiological temperature. *Journal of Materials Science: Materials in Medicine* 23(3): 613–621.
- Qi HJ, Nguyen TD, Castro F, et al. (2008) Finite deformation thermo-mechanical behavior of thermally induced shape memory polymers. *Journal of the Mechanics and Physics of Solids* 56(5): 1730–1751.
- Rousseau IA and Xie T (2010) Shape memory epoxy: composition, structure, properties and shape memory performances. *Journal of Materials Chemistry* 20(17): 3431–3441.

- Small W, Wilson TS, Benett WJ, et al. (2005) Laser-activated shape memory polymer intravascular thrombectomy device. *Optics Express* 13(20): 8204–8213.
- Tandon GP, Goecke K, Cable K, et al. (2009) Durability assessment of styrene- and epoxy-based shape-memory polymer resins. *Journal of Intelligent Material Systems and Structures* 20(17): 2127–2143.
- Tandon GP, Goecke K, Cable K, et al. (2010) Environmental durability of fabric-reinforced shape-memory polymer composites. *Journal of Intelligent Material Systems and Structures* 21(14): 1365–1381.
- Tmopking SS, Bowles DE, Slmep WS, et al. (1988) Response of composite materials to the space station orbit environment. In: *A collection of technical paper of a symposium of AIAA SDM issues of International Space Station*, Williamsburg, VA, 21–22 April.
- Tobushi H, Hayashi S and Kojima S (1992) Mechanical properties of shape memory polymer of polyurethane series—basic characteristics of stress-strain-temperature relationship. *JSME International Journal Series I: Solid Mechanics Strength of Materials* 35(3): 296–302.
- Xie T (2010) Tunable polymer multi-shape memory effect. *Nature* 464(7286): 267–270.
- Xu B, Fu YQ, Ahmad M, et al. (2010) Thermo-mechanical properties of polystyrene-based shape memory nanocomposites. *Journal of Materials Chemistry* 20(17): 3442–3448.
- Xu T, Li GQ and Pang SS (2011) Effects of ultraviolet radiation on morphology and thermo-mechanical properties of shape memory polymer based syntactic foam. *Composites Part A: Applied Science and Manufacturing* 42(10): 1525–1533.
- Yakacki CM, Lyons MB, Rech B, et al. (2008) Cytotoxicity and thermomechanical behavior of biomedical shape-memory polymer networks post-sterilization. *Biomedical Materials* 3(1): 015010.
- Youssef HA. (2009) Effect of electron beam irradiation on the properties of high styrene rubber/styrene butadiene rubber blends. *Journal of Macromolecular Science Part A: Pure and Applied Chemistry* 46(6): 636–642.
- Yu K, Xie T, Leng JS, et al. (2012) Mechanisms of multi-shape memory effects and associated energy release in shape memory polymers. *Soft Matter* 8(20): 5687–5695.
- Zhao Y, Wang CC, Huang WM, et al. (2011) Buckling of poly(methyl methacrylate) in stimulus-responsive shape recovery. *Applied Physics Letters* 99(13): 131911.