


Cite this: *RSC Adv.*, 2020, **10**, 8967

## Synthesis and characterization of carboxylated PBAMO copolymers as promising prepolymers for energetic binders

Minkyung Lim,<sup>†</sup><sup>a</sup> Meilan Bu,<sup>†</sup><sup>a</sup> Yoorim Jang,<sup>a</sup> Jongoh Jeong,<sup>b</sup> Sitae Noh<sup>\*c</sup> and Hakjune Rhee  <sup>\*ad</sup>

The carboxylated poly[3,3-bis(3-azidomethyl)oxetane] (PBAMO) copolymers (poly(BAMO-carboxylate)) were synthesized by substitution of poly[3,3-bis(3-chloromethyl)oxetane] (PBCMO) with potassium carboxylate and sodium azide in DMSO. The synthesized compounds were characterized using various analytical techniques, such as Fourier-transform infrared (FT-IR) spectroscopy, inverse-gated decoupling <sup>13</sup>C-nuclear magnetic resonance (<sup>13</sup>C NMR) spectroscopy, gel permeation chromatography (GPC), thermogravimetric analysis (TGA), differential scanning calorimetry (DSC), calorimetry, friction, and impact sensitivity analysis. These poly(BAMO-carboxylate) compounds have better thermal properties, with lower glass transition temperatures (ranging from  $-43$  °C to  $-51$  °C) than PBAMO ( $-37$  °C) and higher thermal decomposition temperatures ( $233$ – $237$  °C) than PBAMO ( $211$  °C). Moreover, poly(BAMO<sub>0.80</sub>-octanoate<sub>0.20</sub>) and poly(BAMO<sub>0.78</sub>-decanoate<sub>0.22</sub>) have higher heats of combustion (5226 and 5665 kJ mol<sup>-1</sup>, respectively) and negative formation enthalpies ( $-0.17$  and  $-0.55$  kJ g<sup>-1</sup>, respectively), while PBAMO has lower heat of combustion (3125 kJ mol<sup>-1</sup>) and positive formation enthalpy (0.06 kJ g<sup>-1</sup>). The poly(BAMO-carboxylate) compounds have higher values (38–50 J) than that of PBAMO (14 J) in the impact sensitivities. This is a valuable study for improving the properties of PBAMO, which is a high energetic polymeric binder but difficult to handle because of its sensitivity. Therefore, poly(BAMO-carboxylate) could be a good candidate as a prepolymer for designing the energetic polymeric binder.

Received 26th November 2019  
Accepted 25th February 2020

DOI: 10.1039/c9ra09881j  
rsc.li/rsc-advances

### 1. Introduction

Solid rocket propellants are mixtures consisting of polymeric binders, high-energy additives, curing agents, oxidizers, metallic fuel additives, burning rate modifiers, *etc.*<sup>1–5</sup> Among these components, the polymeric binder plays an important role in determining structural integrity, which in turn affects the performance of the propellant.<sup>6–8</sup> The binders are typically cross-linked polymers, and hydroxyl-terminated poly(butadiene) (HTPB) is the most widely used prepolymer. The HTPB propellant has high specific impulse with the ammonium perchlorate (AP) oxidizer. However, the HTPB/AP binder releases HCl gas, which causes environmental pollution due to

the propellant plumes.<sup>4,9</sup> Recent propellant research has been aimed at developing high-energy binders that can provide better performance and improve the overall energy level to replace HTPB. Azide-based ( $-N_3$ ) polymers like glycidyl azide polymer (GAP), poly[3,3-bis(3-azidomethyl)oxetane] (PBAMO), and poly[(3-azidomethyl)methyloxetane] (PAMMO) are attracting attention, because they release high energy when decomposed (Scheme 1).<sup>10–12</sup>

Among these polymers, GAP has a high positive enthalpy of formation ( $+957$  kJ kg<sup>-1</sup>), a low glass transition temperature ( $T_g = -49$  °C), low sensitivity, and good compatibility with high-energy oxidizers like ammonium dinitramide (ADN) and hydrazinium nitroformate (HNF).<sup>1,13–16</sup> Furthermore, GAP

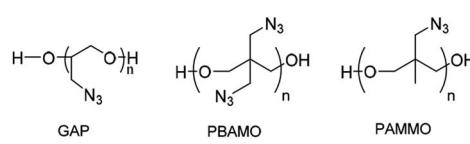
<sup>a</sup>Department of Bionanotechnology, Hanyang University, 55 Hanyangdaehak-ro, Sangnok-gu, Ansan, Gyeonggi-do, 15588, South Korea. E-mail: hrhee@hanyang.ac.kr

<sup>b</sup>NOROO Paint & Coatings Co., Ltd, 351, Bakdal-ro, Manan-gu, Anyang-si, Gyeonggi-do, 13977, South Korea

<sup>c</sup>Department of Materials Science and Chemical Engineering, Hanyang University, 55 Hanyangdaehak-ro, Sangnok-gu, Ansan, Gyeonggi-do, 15588, South Korea

<sup>†</sup>Department of Applied Chemistry, Hanyang University, 55 Hanyangdaehak-ro, Sangnok-gu, Ansan, Gyeonggi-do, 15588, South Korea

<sup>†</sup> These authors contributed equally.



Scheme 1 Examples of azide-based polymers for the energetic binder.



produces chlorine-free propellant plumes with ammonium nitrate (AN) instead of AP, thus curbing environmental pollution. However, the bulky and polar azide groups in GAP induce low flexibility of backbone due to hindrance of motion in molecular chain, leading to drawbacks such as poor low-temperature characteristics.<sup>17–19</sup> To improve the thermal properties of GAP, GAP copolymers such as fluorinated GAP copolymer,<sup>20</sup> GAP-THF copolymer,<sup>21</sup> GAP-PEG copolymer,<sup>22</sup> and GAP-PBAMO copolymer<sup>18,23,24</sup> have been synthesized. These GAP copolymers have better thermal properties than GAP.

Previously, we developed carboxylated GAP copolymers as prepolymers for designing high-energy binders.<sup>25</sup> We focused on improving the thermal properties of GAP copolymer by the introduction of long-chain carboxylates like butyrate, octanoate, 2-ethyl hexanoate, isonanoate, and decanoate.

The synthesized carboxylated GAP copolymers have better thermal properties owing to their lower glass transition temperatures (ranging from  $-48$  °C to  $-55$  °C) than GAP ( $-49$  °C) and thermal decomposition temperatures (228–230 °C) similar to that of GAP (227 °C). Moreover, carboxylated GAP copolymers have higher heats of combustion and negative formation enthalpies compared to GAP.

As an extension study of the synthesis of GAP copolymers, carboxylated PBAMO copolymers were synthesized and characterized herein. PBAMO containing two azide bonds has higher energy, but the glass transition temperature ( $T_g$ ) and crystallinity of PBAMO are relatively high due to its symmetric structure.<sup>26,27</sup> Because of these properties, practicality of PBAMO is limited;<sup>28</sup> therefore, the development of new PBAMO copolymers is of importance. Based on the carboxylated GAP copolymer study, PBAMO copolymer was substituted with 20% long-chain carboxylates by selecting butanoate, octanoate, and decanoate. The resultant carboxylated PBAMO copolymers have lower glass transition temperatures, higher thermal decomposition temperatures, and lower sensitivities than PBAMO. Considering its excellent performance, the carboxylated PBAMO

copolymer is a promising prepolymer candidate for energetic binders.

## 2. Results and discussion

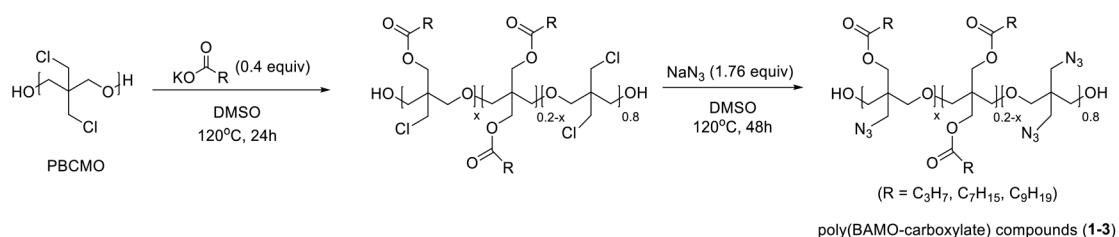
### 2.1 Synthesis and characterization of poly(BAMO-carboxylate) compounds

Some GAP copolymers substituted with long-chain carboxylates (butyrate, octanoate, 2-ethyl hexanoate, isonanoate, and decanoate) have been reported previously.<sup>25</sup> Poly(GA-carboxylate) compounds with three different azide : carboxylate molar ratios (90 : 10, 80 : 20, and 70 : 30) were synthesized and characterized. Most of these poly(GA-carboxylate) compounds have properties similar to each other and exhibited better properties than GAP. Therefore, we selected butanoate, octanoate, and decanoate to synthesize PBAMO copolymers substituted with 20 mol% carboxylate.

PBAMO is generally synthesized by using sodium azide with poly[3,3-bis(3-chloromethyl)oxetane] (PBCMO). First, we synthesized poly(BAMO-carboxylate) compounds by sequential addition of sodium azide and potassium carboxylate to PBCMO. However, due to the steric congestion in PBAMO containing two azide groups on one carbon, the substitution of the residual C–Cl bond with carboxylate was not completed at 120 °C. Thus, poly(BAMO-carboxylate) compounds were synthesized by the first substitution of PBCMO with relatively bulky carboxylate, followed by addition of azide group. Substitution of all C–Cl bonds was completed after 48 h using 1.76 equivalents of sodium azide at 120 °C. The synthesized poly(BAMO-carboxylate) compounds were analyzed by FT-IR spectroscopy, inverse-gated decoupling  $^{13}\text{C}$ -NMR spectroscopy, and GPC to confirm successful synthesis (Table 1).

The completion of the reaction was confirmed by FT-IR spectroscopy (Fig. 1). Compared with the starting material PBCMO, the azide N=N=N stretching band at  $2091\text{ cm}^{-1}$  and ester C=O stretching band at  $1735\text{ cm}^{-1}$  increased in intensity,

Table 1 Results of substitution reaction using PBCMO, potassium carboxylate, and sodium azide



Entry	Composition <sup>a</sup>	[Potassium carboxylate] : [sodium azide] mol%	R	Yield (%)	$M_w^b$ (g mol $^{-1}$ )	$M_n^b$ (g mol $^{-1}$ )	PDI $^b$	$T_g^c$ (°C)	$T_{dec}^d$ (°C)
1	PBCMO	—	—	—	2250	1850	1.22	—	—
2	PBAMO	—	—	—	2460	2030	1.21	-37	211
3	Poly(BAMO <sub>0.79</sub> -butyrate <sub>0.21</sub> ) (1)	80 : 20	$\text{C}_3\text{H}_7$	92	2720	2080	1.30	-43	233
4	Poly(BAMO <sub>0.80</sub> -octanoate <sub>0.20</sub> ) (2)	80 : 20	$\text{C}_7\text{H}_{15}$	90	2910	2390	1.22	-47	237
5	Poly(BAMO <sub>0.78</sub> -decanoate <sub>0.22</sub> ) (3)	80 : 20	$\text{C}_9\text{H}_{19}$	91	2980	2440	1.22	-51	237

<sup>a</sup> Calculated by inverse-gated decoupling  $^{13}\text{C}$  NMR spectroscopy. <sup>b</sup> Measured by GPC. <sup>c</sup> Measured by DSC. <sup>d</sup> Measured by TGA (reported 1st  $T_{dec}$ ).



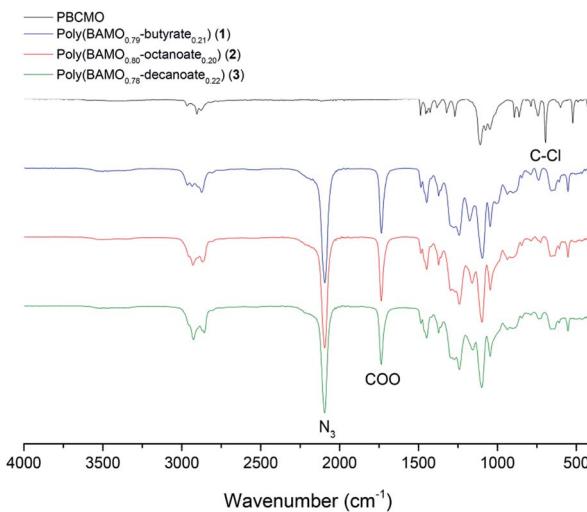


Fig. 1 IR spectra of PBCMO and poly(BAMO-carboxylate) compounds (1–3).

while the C-Cl stretching band at  $700\text{ cm}^{-1}$  disappeared as the reactions proceeded to completion.

The composition of the polymer was analyzed by a quantitative  $^{13}\text{C}$  NMR spectroscopic study based on inverse-gated decoupling.<sup>25,29,30</sup> The structures and successful synthesis of poly(BAMO-carboxylate) (1–3) were confirmed from the  $^{13}\text{C}$  NMR spectra.

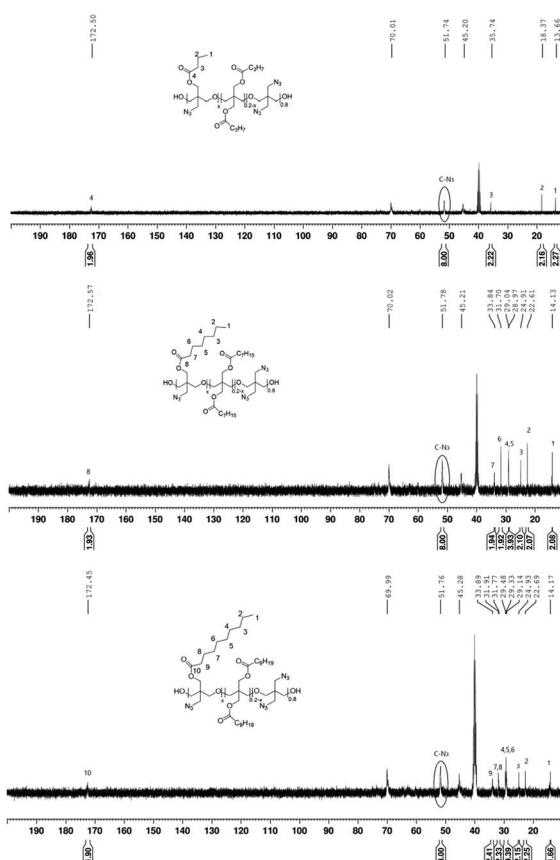


Fig. 2 Inverse-gated decoupling  $^{13}\text{C}$  NMR spectra of poly(BAMO-carboxylate) compounds.

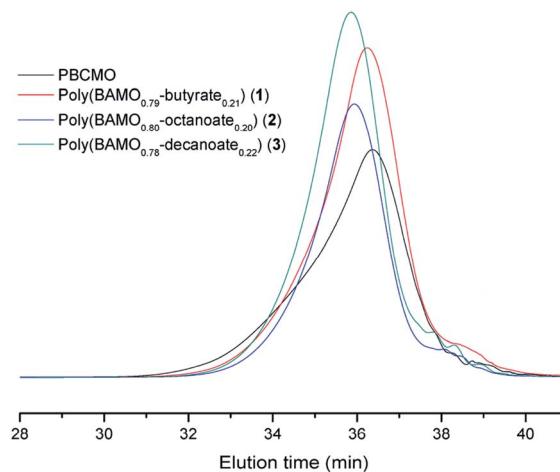


Fig. 3 GPC curves of PBCMO and the poly(BAMO-carboxylate) compounds (1–3).

NMR spectra. The compositions of poly(BAMO<sub>0.79</sub>-butylate<sub>0.21</sub>) (1), poly(BAMO<sub>0.80</sub>-octanoate<sub>0.20</sub>) (2), and poly(BAMO<sub>0.78</sub>-decanoate<sub>0.22</sub>) (3) were calculated by averaging the integral values of each peak in the inverse-gated decoupling  $^{13}\text{C}$  NMR spectra (Fig. 2). The substitution ratio of azide and carboxylate in each compound was almost similar to the target value of 80 : 20 (azide : carboxylate).

The dispersity values and the average molecular weights of poly(BAMO-carboxylate) compounds (1–3) were analyzed by GPC. The GPC chromatograms are shown in Fig. 3, and the detailed data are listed in Table 1. The substitution reaction gave high yields (90–92%), with average molecular weights ranging from  $2080\text{ g mol}^{-1}$  to  $2440\text{ g mol}^{-1}$ , which represented an increase from the average molecular weight of PBCMO ( $1850\text{ g mol}^{-1}$ ) and polydispersity index (PDI =  $M_w/M_n$ ) values that ranged from 1.22 to 1.30. The PDI values of all the poly(BAMO-carboxylate) compounds (1–3) are close to 1.00, indicating that the copolymers were successfully synthesized.

## 2.2 Thermal properties

Thermal stability and low glass transition temperature are critical factors for increasing propellant insensitivity. DSC and

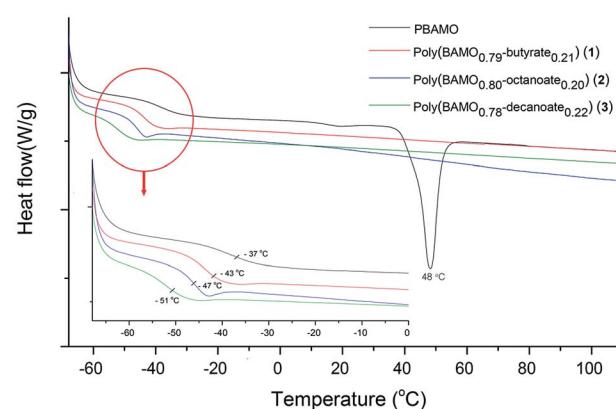


Fig. 4 DSC curves of PBAMO and the poly(BAMO-carboxylate) compounds (1–3).

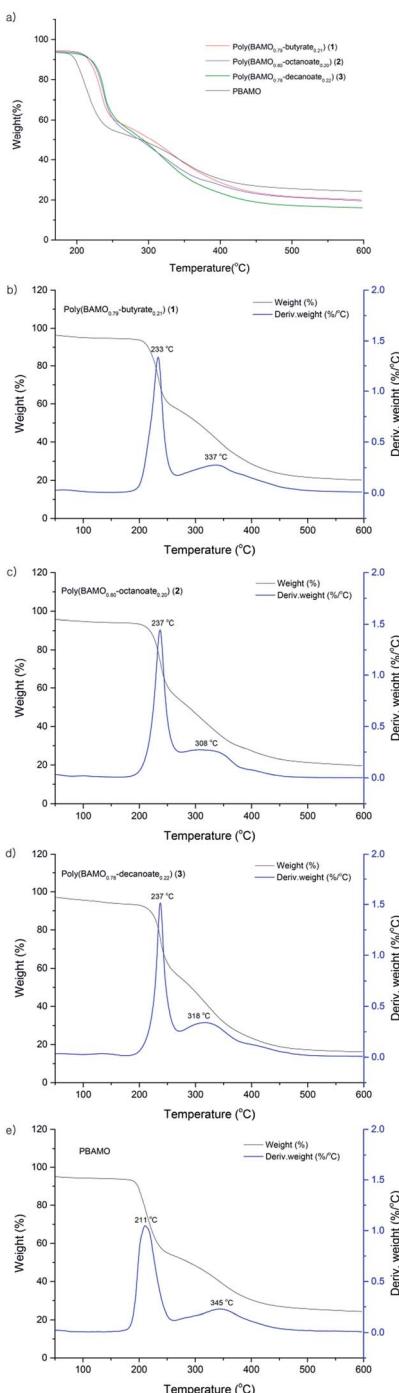


Fig. 5 TGA and DTG curves of the poly(BAMO-carboxylate) compounds (1–3) and PBAMO.

TGA experiments were performed to determine the thermal properties. To confirm the effect of carboxylate substitution, the DSC and TGA experiments of pure PBAMO were also performed. The obtained values are summarized in Table 1. The three poly(BAMO-carboxylate) (1–3) have similar  $T_g$  values ranging from  $-43$  °C to  $-51$  °C (Table 1 and Fig. 4). In Fig. 4, a melting temperature (48 °C) was observed only in PBAMO.<sup>18</sup>

The  $T_g$  values of the three poly(BAMO-carboxylate) were lower than that of PBAMO, and the longer the alkyl chain is, the lower glass transition temperature is. Partial introduction of long-chain carboxylates into PBAMO reduces the intramolecular and intermolecular forces in the skeleton, thereby lowering the  $T_g$ .<sup>31</sup> In addition, it is expected that  $T_g$  is lowered because the symmetric structure of PBAMO is broken and the crystallinity is lowered due to random substitution of carboxylate. This property facilitates processability in the preparation of the propellant binder.<sup>6,32</sup>

The thermal decomposition temperatures ( $T_{dec}$ ) of compounds 1–3 were measured by TGA to determine their thermal stabilities. As shown in Fig. 5, there are two main exothermic peaks in compounds 1–3, ranging from 233 to 237 °C and from 308 to 337 °C, attributed to the energy release due to azido-decomposition forming  $N_2$  gas and the decomposition of polymer backbone. The compounds 1–3 have similar thermal decomposition temperatures that are about 20 °C higher than that of PBAMO (211 °C), *i.e.* 1–3 have better thermal stabilities than PBAMO.

### 2.3 Energetic properties

Besides thermal stability, sensitivity against stimuli and energetic properties is also important for the application of polymeric binders. The friction and impact sensitivities of synthesized copolymers were measured by NATO STANAG 4487 and NATO STANAG 4489, respectively.

As shown in Table 2, the friction sensitivities of all the compounds was above the detection limit, and thus could not be compared. Compounds 1–3 have lower impact sensitivities than PBAMO, because the azide contents of compounds 1–3 were lower than that of PBAMO. The impact sensitivity of 1 was 38 J, which is higher than that of PBAMO (14 J). Notably, the impact sensitivities of 2 and 3, having longer alkyl chains, are above the detection limit of 50 J. Therefore, the properties of PBAMO, which is difficult to handle because of its sensitivity, must be improved.

To check the energetic properties and to improve sensitivity, the energies of combustion ( $\Delta U_{comb}$ ) of PBAMO, poly(BAMO<sub>0.80</sub>-octanoate<sub>0.20</sub>) (2), and poly(BAMO<sub>0.78</sub>-decanoate<sub>0.22</sub>) (3) were measured by a Parr Bomb Calorimeter 6200. Then, the

Table 2 Friction and impact sensitivity of PBAMO and poly(BAMO-carboxylate) compounds (1–3)

	Poly(BAMO <sub>0.79</sub> -butylate <sub>0.21</sub> ) (1)	Poly(BAMO <sub>0.80</sub> -octanoate <sub>0.20</sub> ) (2)	Poly(BAMO <sub>0.78</sub> -decanoate <sub>0.22</sub> ) (3)	PBAMO
Friction sensitivity (N)	>360	>360	>360	>360
Impact sensitivity (J)	38	>50	>50	14



Table 3 Energy values of PBAMO and the poly(BAMO-carboxylate) compounds (2, 3)

	PBAMO	Poly(BAMO <sub>0.80</sub> -octanoate <sub>0.20</sub> ) (2)	Poly(BAMO <sub>0.78</sub> -decanoate <sub>0.22</sub> ) (3)
Formula (repeating unit)	C <sub>5</sub> H <sub>8</sub> N <sub>6</sub> O	C <sub>8.2</sub> H <sub>14.2</sub> N <sub>4.8</sub> O <sub>1.8</sub>	C <sub>9</sub> H <sub>15.6</sub> N <sub>4.8</sub> O <sub>1.8</sub>
FW (repeating unit) [g mol <sup>-1</sup> ]	168.16	208.84	219.86
-ΔU <sub>comb</sub> [J g <sup>-1</sup> ]	18 590	25 023	25 716
-ΔH <sub>comb</sub> [kJ mol <sup>-1</sup> ]	3125	5226	5665
Δ <sub>f</sub> H <sup>o</sup> <sub>m</sub> [kJ mol <sup>-1</sup> ]	10.78	-35.05	-121.33
Δ <sub>f</sub> H <sup>o</sup> [kJ g <sup>-1</sup> ]	0.06	-0.17	-0.55

combustion enthalpies ( $\Delta H_{\text{comb}}$ ) were calculated at 25 °C based on the combustion energies ( $\Delta H_{\text{comb}} = \Delta U_{\text{comb}} + \Delta nRT$ ;  $\Delta n = \Delta n_i(\text{product, g}) - \Delta n_i(\text{reactant, g})$ ). The enthalpies of formation ( $\Delta_f H^o$ ) can be obtained by using the following equation:  $\Delta_f H^o(\text{copolymer}) = a\Delta_f H^o(\text{CO}_2) + 0.5b\Delta_f H^o(\text{H}_2\text{O}) - \Delta_c H^o(\text{copolymer})$  (for the composition  $C_aH_bN_cO_d$ ). It is based on the Hess thermochemical cycle at 25 °C with the combustion reactions of the repeating unit, as shown in Scheme 1. Here, the heats of formation for H<sub>2</sub>O (l) and CO<sub>2</sub> (g) are -286 and -394 kJ mol<sup>-1</sup>, respectively.<sup>31</sup>

As shown in Table 3, the combustion enthalpies are always negative because combustion is an exothermic reaction. The combustion enthalpies of compounds 2 and 3 have more negative values in than that of PBAMO, because they have larger repeating units (greater molar mass) and  $|\Delta U_{\text{comb}}|$  than the latter. It suggests that compounds 2 and 3 release higher energies during the combustion process than PBAMO. Additionally, the enthalpies of formation of 2 and 3 are negative, while that of PBAMO is positive (0.06 kJ g<sup>-1</sup>), because they have higher carbons and hydrogen contents in the repeating unit. In this case, contributions of the CO<sub>2</sub> and H<sub>2</sub>O terms in the equation for the enthalpies of formation increase beyond the  $\Delta_c H^o$  term.<sup>33</sup> Therefore, compounds 2 and 3 have negative formation enthalpies, which indicate the product is more stable than the constituent elements. In contrast, the positive formation enthalpy of PBAMO suggests the opposite.

### 3. Experimental

#### 3.1 Materials and instruments

PBCMO was supplied by J. CHEM. Inc. Butanoic acid, decanoic acid, and octanoic acid were supplied by PMC KOREA CO., LTD. The other chemicals were purchased from commercial sources (Sigma Aldrich and TCI), and were used without further purification, unless specifically mentioned. For the quantitative analysis of the product, the inverse-gated decoupling <sup>13</sup>C NMR spectra were obtained in DMSO with a Bruker NMR and zpig30 pulse program. The FT-IR spectra were collected on an Alpha FT-IR spectrometer from Bruker using a diamond ATR accessory in the wavenumber range 4000 to 400 cm<sup>-1</sup>. The friction and impact sensitivities were determined using NATO STANAG 4487 and NATO STANAG 4489, respectively. DSC was performed on a TA instruments Q 1000 using aluminum pans at the heating rate of 10 °C min<sup>-1</sup> under nitrogen flow, in the temperature range from -70 °C to 110 °C. TGA was performed

on a PERKIN ELMER TGA Q500 V6.2 Build 187 under nitrogen atmosphere at a heating rate of 5 °C min<sup>-1</sup>, from 25 °C to 600 °C. The molecular weights of the materials were determined by WATERS 515, using THF as the flow solvent at the flow rate of 1.0 mL min<sup>-1</sup>. The heat of combustion was determined by the Parr Bomb Calorimeter 6200.

#### 3.2 General procedure for the synthesis of potassium carboxylate

For the synthesis of potassium carboxylate, methanol (50 mL), carboxylic acid (110 mmol), and potassium hydroxide (100 mmol) were added to a 100 mL round-bottom flask. The reaction mixture was stirred at room temperature for 2 h. After evaporating methanol, the potassium carboxylate was filtered off, washed with hexane, and dried in the oven at 40 °C for 24 h.

#### 3.3 General procedure for the synthesis of PBAMO copolymers based on carboxylate

The PBAMO copolymers were synthesized by the reaction of PBCMO, sodium azide, and potassium carboxylate. PBCMO (3.10 g, 20 mmol) was dissolved in DMSO (20 mL) in a 100 mL round-bottom flask at 60 °C. Potassium carboxylate (8 mmol) was added and stirred at 120 °C for 24 h. Then, sodium azide (2.29 g, 35.2 mmol) was added to the reaction mixture and stirred at 120 °C for 48 h. After the completion of the reaction, the mixture was cooled and 50 mL of ethyl acetate was added. The organic layer was washed with distilled water (50 mL × 4) to remove DMSO, dried over MgSO<sub>4</sub>, filtered through a pad of Celite, and concentrated under reduced pressure.

**Poly(BAMO<sub>0.79</sub>-butyrate<sub>0.18</sub>) (1).** <sup>13</sup>C NMR (100 MHz, DMSO-*d*<sub>6</sub>): 172.5, 70.0, 51.7, 45.2, 35.7, 18.4, 13.7; IR: 2873, 2093, 1735, 1448, 1270, 1096, 553 cm<sup>-1</sup>.

**Poly(BAMO<sub>0.80</sub>-octanoate<sub>0.20</sub>) (2).** <sup>13</sup>C NMR (100 MHz, DMSO-*d*<sub>6</sub>): 172.6, 70.0, 51.8, 45.2, 33.8, 31.7, 29.0, 29.0, 24.9, 22.6; IR: 2871, 2094, 1736, 1448, 1241, 1098, 553 cm<sup>-1</sup>.

**Poly(BAMO<sub>0.78</sub>-decanoate<sub>0.22</sub>) (3).** <sup>13</sup>C NMR (100 MHz, DMSO-*d*<sub>6</sub>): 172.5, 70.0, 51.8, 45.3, 33.9, 32.0, 31.8, 29.5, 29.3, 29.1, 24.9, 22.7, 14.2; IR: 2857, 2095, 1736, 1448, 1241, 1099, 553 cm<sup>-1</sup>.

### 4. Conclusions

Poly(BAMO-carboxylate) compounds were synthesized by the substitution of poly[3,3-bis(3-chloromethyl)oxetane] (PBCMO) with potassium carboxylate and sodium azide in DMSO, and



were characterized by FT-IR and  $^{13}\text{C}$  NMR spectroscopy. The molecular composition of poly(BAMO-carboxylate) compounds was confirmed by inverse-gated decoupling  $^{13}\text{C}$  NMR spectroscopy, and the average molecular weight was measured by GPC. The poly(BAMO-carboxylate) compounds were found to have molecular compositions similar to the theoretically calculated ones. The thermal properties were measured by DSC and TGA. The energy properties were determined in terms of the sensitivity and heats of combustion and formation. The poly(BAMO-carboxylate) compounds were found to have lower glass transition temperatures and higher thermal decomposition temperatures than PBAMO. Poly(BAMO<sub>0.80</sub>-octanoate<sub>0.20</sub>) and poly(BAMO<sub>0.78</sub>-decanoate<sub>0.22</sub>) have negative formation enthalpies and higher heats of combustion than PBAMO. Moreover, the impact sensitivities of poly(BAMO-carboxylate) compounds were lower than those of PBAMO. Consequently, poly(BAMO-carboxylate) compounds exhibited better thermal and energetic properties than PBAMO as a polymeric binder for solid propellants. These properties facilitate processability in the preparation of the propellant binder. Therefore, poly(BAMO-carboxylate) is a good candidate as a prepolymer for designing energetic polymeric binders.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This research was supported by the Defense Acquisition Program Administration (DAPA 912575201) and the Korean Ministry of Education through the BK21-Plus project.

## Notes and references

- 1 M. S. Eroğlu and O. Güven, *Polymer*, 1998, **39**, 1173–1176.
- 2 B. Gaur, B. Lochab, V. Choudhary and I. K. Varma, *J. Macromol. Sci., Polym. Rev.*, 2003, **43**, 505–545.
- 3 E. Landsem, T. L. Jensen, F. K. Hansen, E. Unneberg and T. E. Kristensen, *Propellants, Explos., Pyrotech.*, 2012, **37**, 581–591.
- 4 Y. M. Mohan, Y. Mani and K. M. Raju, *Des. Monomers Polym.*, 2012, **9**, 201–236.
- 5 J. Deng, G. Li, M. Xia, Y. Lan and Y. Luo, *J. Appl. Polym. Sci.*, 2016, **133**, 43840.
- 6 R. G. Sracer and D. M. Husband, *Propellants, Explos., Pyrotech.*, 1991, **16**, 167–176.
- 7 Y. Wu, Y. Luo and Z. Ge, *Propellants, Explos., Pyrotech.*, 2015, **40**, 67–73.
- 8 M. Cappello, S. Filippi, L. Mori and G. Polacco, *Propellants, Explos., Pyrotech.*, 2017, **42**, 974–981.
- 9 Y. Zhou, X.-P. Long and Q.-X. Zeng, *J. Appl. Polym. Sci.*, 2012, **125**, 1530–1537.
- 10 I. K. Varma, *Macromol. Symp.*, 2004, **210**, 121–129.
- 11 Y. M. Mohan, K. M. Raju and B. Sreedhar, *Int. J. Polym. Mater.*, 2006, **55**, 441–455.
- 12 M. Cappello, S. Filippi, L. Mori and G. Polacco, *Int. J. Polym. Mater.*, 2017, **42**, 974–981.
- 13 A. M. Kawamoto, J. A. S. Holanda, U. Barbieri, G. Polacco, T. Keicher, H. Krause and M. Kaiser, *Propellants, Explos., Pyrotech.*, 2008, **33**, 365–372.
- 14 Y. Zhang, J. Zhao, P. Yang, S. He and H. Huang, *Polym. Eng. Sci.*, 2012, **52**, 768–773.
- 15 J.-F. Pei, F.-Q. Zhao, X.-D. Song, X.-N. Ren, H.-X. Gao, T. An, J. An and R.-Z. Hu, *J. Anal. Appl. Pyrolysis*, 2015, **112**, 88–93.
- 16 G. Li, H. Dong, M. Liu, M. Xia, C. Chai and Y. Luo, *Polym. Int.*, 2017, **66**, 1037–1043.
- 17 B. S. Min and S. W. Ko, *Macromol. Res.*, 2007, **15**, 225–233.
- 18 S. Pisharath and H. G. Ang, *Polym. Degrad. Stab.*, 2007, **92**, 1365–1377.
- 19 B. Li, Y. Zhao, G. Liu, X. Li and Y. Luo, *J. Therm. Anal.*, 2016, **126**, 717–724.
- 20 M. Xu, Z. Ge, X. Lu, H. Mo, Y. Ji and H. Hu, *RSC Adv.*, 2017, **7**, 47271–47278.
- 21 Y. M. Mohan and K. M. Raju, *Int. J. Polym. Mater.*, 2006, **55**, 203–217.
- 22 Y. M. Mohan, M. P. Raju and K. M. Raju, *Int. J. Polym. Mater.*, 2005, **54**, 651–666.
- 23 U. Barbieri, G. Polacco, T. Keicher and R. Massimi, *Propellants, Explos., Pyrotech.*, 2009, **34**, 427–435.
- 24 J.-F. Pei, F.-Q. Zhao, H.-L. Lu, X.-D. Song, R. Zhou, Z.-F. Yuan, J. Zhang and J.-B. Chen, *J. Therm. Anal.*, 2016, **124**, 1301–1307.
- 25 H. Kim, Y. Jang, S. Noh, J. Jeong, D. Kim, B. Kang, T. Kang, H. Choi and H. Rhee, *RSC Adv.*, 2018, **8**, 20032–20038.
- 26 Y. Oyumi and T. B. Brill, *Combust. Flame*, 1986, **65**, 127–135.
- 27 C. Zhang, J. Li, Y.-J. Luo and B. Zhai, *J. Energ. Mater.*, 2015, **33**, 305–314.
- 28 K. E. Hardenstine, G. V. S. Henderson, L. H. Sperling, C. J. Murphy and G. E. Manser, *J. Polym. Sci., Part B: Polym. Phys.*, 1985, **23**, 1597–1609.
- 29 M. Cao, T. Li, J. Liang and G. Du, *Polymers*, 2017, **9**, 109.
- 30 M. Cao, T. Li, J. Liang, Z. Wu, X. Zhou and G. Du, *Polymers*, 2016, **8**, 391.
- 31 S. Hafner, T. Keicher and T. M. Klapötke, *Propellants, Explos., Pyrotech.*, 2017, **43**, 126–135.
- 32 J. Blackwell, M. R. Nagarajan and T. B. Hoitink, *Polymer*, 1982, **23**, 950–956.
- 33 E. Diaz, P. Brousseau, G. Ampleman and R. E. Prud'homme, *Propellants, Explos., Pyrotech.*, 2003, **28**, 101–106.

