



Azidated Ether-Butadiene-Ether Block Copolymers as Binders for Solid Propellants

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Azidated Ether-Butadiene-Ether Block Copolymers as Binders for Solid PropellantsShort title: **Azidated Ether-Butadiene Energetic Copolymers***Miriam Cappello, Pietro Lamia, Claudio Mura, Giovanni Polacco*, Sara Filippi**Department of Civil and Industrial Engineering**University of Pisa**Largo Lucio Lazzarino, 1 – 56122 Pisa, ITALY*

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Abstract

Polymeric binders for solid propellants are usually based on hydroxyl-terminated polybutadiene (HTPB), which does not contribute to the overall energy output. Azidic polyethers represent an interesting alternative, but may have poorer mechanical properties. Polybutadiene-polyether copolymers may join the advantages of both. Four different ether-butadiene-ether tri-block copolymers were prepared and azidated starting from halogenated and/or tosylated monomers and using HTPB as initiator. The presence of the butadiene block complicates the azidation step and reduces the storage stability of the azidic polymer. Nevertheless, the procedure allows modifying the binder properties by varying the type and the lengths of the energetic blocks.

1. Introduction

In the most common propellant formulations, the solid ingredients are mixed with a polymer, which is subsequently cast cured to give an elastomeric network. The polymer binder acts by wetting the solid thus providing a void-free matrix that allows the formulation to be cast into large and irregular cases [1]. The polymer provides mechanical integrity to the final product and, together with a plasticiser, ensures safety during handling, since it absorbs and dissipates energy from hazardous stimuli that may happen during storage and transportation [2]. The most commonly used binder is hydroxy-terminated polybutadiene (HTPB) cross-linked with isocyanates to give a polyurethanic network. Alternatively, carboxy-terminated polybutadiene and hydroxyl-terminated polyethers have been also suggested [2]. HTPB is used worldwide and its success from the mechanical and safety point of view is well known and recognized. However, from the propellant point of view, due to its inert character, there are many cases when it is just a dead weight, which does not contribute to the overall energy output and limits the performance of the composition unless there is a high solids loading. On the other hand, high solids loadings induce processing and vulnerability problems. Therefore, the main goal of binder research is to find a formulation that reduces vulnerability, shock and impact insensitivity without lowering performance. A possible approach is to substitute HTPB with alternative materials, such as the so-called energetic-polymers, that act as binder and at the same time contribute to the output energy. The simplest way to achieve this result is the inclusion of functional groups (such as nitro (NO_2), nitrate (NO_3) or azide (N_3)), which may increase the internal enthalpy of formation of the formulation and/or improve the overall oxygen balance. Of course, this has to be done by preserving all the main requirements of a binder, which should [3]:

- (a) be a liquid with a good processability at the mixing temperatures (30-60 °C);

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3 (b) be curable, with minimum evolution of heat at 40-80 °C and give an elastomeric
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5 network with good mechanical properties;
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7 (c) have a glass transition temperature possibly below -40 °C;
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10 (d) be compatible with all the other ingredients of the formulation.

11 During the last decades, several energetic binders have been synthesized and here we will
12 mention only the most promising ones, but the interested reader may find detailed information
13 and a much longer list in the reviews by Agrawal [4], Provatas [2], Badgujar et al. [5], Sikder
14 and Sikder [6] and Gaur et al. [7].

15 The first idea that comes to mind is the introduction of a limited quantity of energetic groups
16 directly into HTPB. The early studies to produce nitrated HTPB (NHTPB) suggested a
17 nitromercuration-demercuration route, [8], while later on Colclough et al. [1] started from
18 epoxidation of HTPB and then reacted N_2O_5 with epoxide groups to form dinitrate esters. Of
19 course, the level of nitration affects also thermal stability and mechanical properties and the
20 authors suggested that a nitration corresponding to 10% of double bonds gives a good
21 compromise between energy output, mechanical properties and miscibility with energetic
22 plasticizers.
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38 Alternatively, the energetic binders may have a chemical structure completely different from
39 HTPB and the most studied one are those containing nitro or azidic groups. In the first case,
40 popular polymers derives from cationic polymerization of 3,3-(nitratomethyl) methyl oxetane
41 3-nitratomethyl-3-methyl oxetane (NIMMO) and glycidyl nitrate (GLYN). Again, the OH-
42 terminated chains are able to crosslink into polyurethane rubbers. By comparing these nitrated
43 polymers, Agrawal [4] suggested that NHTPB has the lower production cost, while the other
44 two have better performances. In the case of azidic functionalities, there is a slightly longer
45 list of candidates, but the most studied one is glycidyl azide polymer (GAP). GAP has been
46 employed also as a plasticizer [4] and several publications may be found concerning its
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3 synthesis as well as its thermal behavior and explosive properties [9-25]. The GAP synthesis
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5 was first described in a patent filed in 1972 by Vandenburg [26] who did azidation of
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7 polyepichlorohydrin (PECH) by using sodium azide in dimethylformamide. Twenty years
8
9 later, Frankel et al. described a semi-industrial production in USA, sponsored by the Air
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11 Force Astronautic Laboratory [27]. The PECH was obtained by polymerization of
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13 epichlorohydrin (ECH) by using glycerol as the initiator to give a triol polymer easily cross-
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15 linkable with isocyanates. In the following decades, the synthetic route has not changed
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17 significantly and sodium azide remains the preferred reagent. The azidation can be performed
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19 in different organic solvents and in water [27, 28]. In the first case, the reaction is much faster
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21 and in dimethyl sulphoxide (DMSO) it is reported to occur at 90-95°C within 12- 18 h. In
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23 water, the use of a phase transfer catalyst is necessary and the reaction may take several days
24
25 before completion at comparable temperatures. Nevertheless, the two processes give GAP of
26
27 comparable quality. PECH may be in linear, star or branched form depending on the catalyst,
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29 initiator and operating conditions. Consequently, the number of hydroxyl functionalities per
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31 chain can be modulated in a wide range of values. Moreover, PECH is now commercially
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33 available as well as GAP. Starting from GAP, several alternative organic azides have been
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35 developed in the last years and the use of the azido group is gaining more and more attention
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37 since it has also other advantages like i.e. reducing the flame and smoke in exhaust gases, thus
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39 making the propellant formulations more eco-friendly [5]. Some examples of suggested
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41 polymeric azides are poly(3,3 bis(azidomethyl)oxetane-co- ϵ -caprolactone) [29], 3,3
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43 bis(azidomethyl)oxetane-tetrahydrofuran [30, 31] and polyglycidylazide-*b*-poly(azidoethyl
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45 methacrylate) [32]. Among the proposed ones, the polyoxetanes, first synthesized by Manser
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47 [33-35], gained an important role in the field of energetic binders. Manser started from
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49 monomers such as 3-nitratomethyl-3-methyl oxetane (NIMMO), 3,3-bis-
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51 (azidomethyl)oxetane (BAMO) and its analog monofunctional 3-azidomethyl-3-methyl
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3 oxetane (AMMO). Later on, he also described the synthesis of many different oxetanes with
4 cubyl and carboranyl groups, but the difficult syntheses involved have so far precluded their
5 evaluation in large-scale formulations [1]. In the case of BAMO and AMMO, a chlorinated or
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oxetane (AMMO). Later on, he also described the synthesis of many different oxetanes with cubyl and carboranyl groups, but the difficult syntheses involved have so far precluded their evaluation in large-scale formulations [1]. In the case of BAMO and AMMO, a chlorinated or tosylated monomeric precursor was azidated and then subjected to cationic ring-opening polymerization (CROP) by using a diol and a Lewis acid as catalyst. The classical Active Chain End (ACE) mechanism proceeds by donation of a proton from initiator to the oxetane, which then propagates with more oxetane monomers until the chain is terminated either with water or alcohol to give the hydroxy-terminated polymer. The number of hydroxyl functionalities per chain should theoretically coincide with those of the alcohol used as initiator.

The molecular weight can be adjusted by changing monomer feed rates and the ratio of diol to Lewis acid, but of course a post-polymerization chain elongation by using di-isocyanates is always possible. Unfortunately, the ACE mechanism has some disadvantages e.g. a lack of molecular weight control and product reproducibility, poor initiator incorporation and formation of cyclic oligomers [36]. For these reasons, an Activated Monomer Mechanism (AMM), which involves the concept of living polymerization, where the OH-terminated polymer reacts with an “active” monomer, may be preferable since it avoids the formation of unstable and highly active cationic propagating species and favors the molecular weight control and reproducibility of the results. Side reactions, including cyclization, are strongly reduced in AMM and well-defined linear products can be obtained [7]. However, it must be emphasized that it is not easy to drive the reaction toward the desired mechanism and often both mechanisms may be present at the same time [37, 38].

It is important to introduce another aspect, related to safety during the synthetic process. In contrast to the above-mentioned polymers, the low molecular weight azide monomers are highly dangerous and may explode without apparent reasons during handling. In addition, the

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3 polymerization of energetic monomers requires careful control of reaction conditions since
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5 initiators may not be compatible with the energetic groups. Consequently, this polymerization
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7 is seen as an advanced technology and high-risk approach. Alternative to azidation-
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9 polymerization is the polymerization-azidation route, where the azidation step follows that of
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11 polymerization of the chlorinated-tosylated monomers. This is a low- tech., low-risk
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13 approach, but gives less opportunity to tailor the final properties of the polymer and suffers
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15 from the usual complications of modifying a macromolecule [36]. Taking into account the
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17 risks associated to the scale up for an industrial production, researchers are nowadays aiming
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19 toward the polymerization followed by azidation. With regard to the three candidate repeating
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21 units, BAMO is the one with the higher N content (50%w) followed by GA (42 w %) and
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23 AMMO (33%w). However, BAMO is a symmetric monomer and gives a crystalline
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25 homopolymer that cannot be used as energetic binder, but is suitable to provide the hard block
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27 of an energetic thermoplastic elastomer [7, 39]. In contrast, GA and AMMO are non-
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29 symmetric and provide amorphous character to the polymer. Therefore, the best solution to
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31 combine the amorphous character and the high energetic content seems to be the random
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33 copolymerization of BAMO with either GA or AMMO introduced in the minimum quantity
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35 necessary to break up the crystallization ability of the polymer. Barbieri et al described the
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37 synthesis of polyAMMO (PAMMO) and polyBAMO (PBAMO) homopolymers starting from
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39 the chlorinated and brominated precursors and followed by azidation with sodium azide in
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41 DMF [40]. In the same paper, poly(GA-*r*-BAMO) copolymers were also synthesized by using
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43 different molar ratios between the two repeating units. Later on, the GA/BAMO = 75/25
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45 molar ratio was considered as the optimal one and the effects of different operating conditions
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47 on copolymer characteristics were investigated [37]. Even though the operating conditions
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49 were set to favor a living character of the polymerization, the final product resulted in
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51 combined AMM and ACE mechanism. In particular, the latter is responsible for the formation
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3 of cyclic oligomers, which of course do not contribute to the formation of the binder network,
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5 but may be tolerated in the formulation since they behave as plasticizers perfectly miscible
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7 and compatible with the binder. The polymers were also subjected to preliminary
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9 characterizations in lab-scale propellant formulations [41, 42]. The same research group
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11 studied also the use of the tosylated precursor of AMMO to prepare PAMMO [43] and
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13 p(AMMO-*r*-BAMO) copolymers [38].

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15 Pisharat et al. choose to produce a thermoplastic elastomer and prepared a poly(BAMO-*b*-
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17 GA-*b*-BAMO) block copolymer by using hydroxyl terminated PECH as initiator for the
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19 polymerization of the chlorinated precursor of BAMO. The azidation was then performed by
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21 using NaN₃ in DMF [39].

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24 Another possible strategy is the joining of HTPB and energetic binders, in order to keep the
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26 advantages of both components. However, as it is easily predictable, due to their different
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28 composition and polarity, HTPB and the azidic polyethers are not compatible and the mixture
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30 is destined to phase separation [44]. Nonetheless, Manu et al. prepared blends of HTPB and
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32 GAP and determined their glass transition after cross-linking with isocyanates [44]. The glass
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34 transition temperatures were evaluated through differential scanning calorimetry (DSC) and,
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36 even though the reported spectra are not easy to interpret, the authors claim that the blends
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38 show a “dominant single glass transition” that indicates a micro-heterogeneous morphology
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40 of the interpenetrating network. This result is somehow in agreement with those reported by
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42 Mathew et al. who did a mechanical and thermal characterization of cross-linked GAP/HTPB
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44 networks [45]. The glass transition temperatures were evaluated by dynamic mechanical
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46 analysis and it was found that blends prepared with GAP content up to 30% showed a single
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48 transition in the loss tangent trace. Very recently, Ding et al. used a triazole curing system as
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50 alternative to the traditional isocyanates, starting from GAP and a propargyl-terminated
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52 polybutadiene, under the catalysis of cuprous chloride at ambient temperature [46].
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3 Nonetheless, it is not advisable to use HTPB and GAP or another azidic polymer in a simple
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5 blend that may separate before the curing process, thus giving the risk of inhomogeneity in
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7 the final product. A possible solution could be the use of a compatibilizer, which may be, e.g.
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9 a diblock copolymer made with the two components, but the best way to limit the problem of
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11 phase separation is the covalent bonding of the two polymers to form block copolymers. In
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13 block copolymers, usually the incompatibility between the blocks leads to a biphasic structure
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15 where the dispersed phase is organized in nanoscale blocks that may arrange in several
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17 configurations that have been widely studied both from the experimental and theoretical point
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19 of view [47]. Zhou et al. made a simulation study of the morphologies of GA-*b*-PB and GA-*b*-
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21 PB-*b*-GA di- and tri-block copolymers and concluded that there is a narrow range of
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23 compositions where a “bicontinuous” phase may form and improve the overall mechanical
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25 properties [48, 49]. Since after copolymerization, the polybutadiene block does not contain
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27 the OH terminal groups, in the above sentence and in what follows it is indicated as PB, while
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29 HTPB refers to the unreacted OH terminated homopolymer.

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33 The GAP/HTPB block copolymers have been synthesized and characterized by a few research
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35 groups. Eroğlu et al. first described the grafting of GAP onto HTPB, via free radical
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37 mechanism, by using a GAP “macroinitiator” obtained by reacting the azidic polymer with
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39 4,4' azo bis(4-cyanopentanoyl chloride) [50]. Later on, Murali and Raju used the same
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41 procedure and made a detailed characterization of the graft copolymer, which exhibited two
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43 distinct glass transition temperatures at -74 and -36 °C, corresponding to the PB and GA
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45 blocks respectively [51].

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49 Alternatively, the hydroxyl functionalities of HTPB may start the polymerization of a suitable
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51 monomer, thus directly giving a tri-block copolymer with PB as the central block. This was
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53 done for ECH almost contemporarily by Vasudevan and Sundararajan [52] and by
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55 Subramanian [53]. The main difference in the two procedures is the azidation step, done by
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3 using NaN_3 , in DMSO or in a mixture of dimethylacetamide and toluene. Similarly, Reddy et
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5 al. used HTPB as the central block and polymerized BAMO on it [54]. It is worth nothing that
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7 in this case the azidation followed by polymerization was chosen.

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9 In this work, the same idea of using HTPB as the polyalcohol that starts the ionic
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11 polymerization was followed to prepare block copolymers to be subsequently azidated. As
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13 already pointed out, the azidation of ether precursors can be performed by using many
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15 different operating conditions and solvents. However, in this case we found that the
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17 unsaturated polybutadiene middle block may lead to an undesired cross-linking that impedes
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19 the subsequent use as binder. Therefore, it has been necessary to test different precursors,
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21 solvents and reaction temperatures to obtain appropriate operating conditions to solve this
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23 problem. A mixture of dimethylacetamide (DMAc) and toluene, as suggested by Subramanian
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25 [53], showed to be suitable to limit the cross-linking problems during azidation. Four block
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27 copolymers were then prepared by using the same procedure: 1) GAP-PB-GAP copolymer
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29 starting from a tosylated glycidol, 2) GA/BAMO-PB-GA/BAMO copolymer from ECH and
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31 3,3-bis-(Brome methyl)oxetane (BBrMO, the brominated precursor of BAMO), 3)
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33 GA/BAMO-PB-GA/BAMO copolymer from tosylated glycidol (GT) and BBrMO and 4)
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35 AMMO-PB-AMMO copolymer starting from 3-tosyloxymethyl-3-methyl oxetane (TMMO).
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37 Therefore, polymers 2 and 3 are equal, with the difference just in the precursors. The first
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39 polymer (GAP-PB-GAP) was already synthesized by Vasudevan and Sundararajan [52] and
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41 by Subramanian [53], but starting from different precursors, while, to our knowledge, the
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43 other two polymers have not been described yet in the scientific literature. The synthetic
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45 pathways are described in Figure 1 (preparation of the monomers starting from the
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47 commercial products) and Figure 2 (synthesis of the polymeric precursors). In all cases, a
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49 polymeric non-energetic precursor was prepared and subsequently azidated.
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3 All the monomers and polymers were characterized by Fourier Transfer Infra Red (FTIR)
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5 analysis and Nuclear Magnetic Resonance (NMR). The thermal behavior of the polymers was
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7 evaluated by Differential Scanning Calorimetry (DSC) and Thermogravimetric Analysis
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9 (TGA).
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11 12 13 14 15 16 **2. Materials and methods**

17 18 *2.1. Materials*

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20 All the chemicals, unless differently stated, were purchased at Sigma Aldrich. N,N
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22 dimethylacetamide (99%), toluene (99,8%), dimethyl sulfoxide (DMSO) (99.7%),
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24 dimethylformamide (DMF) (99.8%), methanol (99.9%), ethanol (99.9%), sodium azide
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26 (>99.5%), boron trifluoride tetrahydrofuranate (BTF THF), triethylamine (TEA) ($\geq 99\%$), 3-
27
28 hydroxy-methyl-3-methyloxetane (HMMO) (98%), glycidol (G) (96%), toluene-4-sulfonyl
29
30 chloride (TsCl) ($\geq 99\%$), sodium chloride (>99%), sodium hydroxide (>98%), sodium
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32 carbonate anhydrous, sodium bicarbonate (>99.7%), magnesium sulfate (>99.5%),
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34 phosphorus pentoxide, hydrochloric acid, potassium hydroxide were used as received.
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36 Diethyl-ether, 3-bromo-2,2-bis(bromomethyl)propanol (BrBBrMP) from Chemos GmbH, was
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38 used as received. Dichloromethane (DCM) was dried with P_2O_5 and distilled at 40°C and 1
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40 atm, ECH and butanediol were distilled under reduced pressure. HTPB was Poly bd[®] R-
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42 45HTLO by Cray Valley, with the following main properties: viscosity = 8000 mPa.s at 23
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44 °C, hydroxyl value = 0.84 meq/g, hydroxyl functionality = 2.4-2.6 OH groups/chain, glass
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46 transition temperature = -75 °C. 2,2'-Methylenebis(6-tert-butyl-4-methylphenol) used as anti-
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48 oxidant was VulKanox BKF by Lanxess.
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2.2. Synthesis of monomers

2.2.1. Glycidyl tosylate GT

The synthesis was performed according to a procedure described by Nakabayashi et al [55]. A 250 mL flask, was fed with 50 mL of anhydrous toluene and 4.2 mL of TEA. In a beaker, a solution of 5.42 g TsCl in 12 mL of toluene was also prepared. The flask and the beaker were then conditioned at -15°C for 1.5 h. Then, the flask was fed with 2 mL of distilled G and dropwise with the TsCl solution. At the end of the feeding, the system was maintained at -15 °C for 24 h, filtered and distilled. The remaining liquid was dropped in cold petroleum-ether thus forming a white suspension that crystallizes at -15 °C. The GT was obtained with a 98.6% yield, as white anhydrous crystals.

2.2.2. 3,3-bis(bromomethyl)oxetane (BBrMO)

A 1L three-necked round bottom flask, fitted with a reflux condenser, a nitrogen inlet and a mechanical stirrer set at 180 rpm was fed with 50 g di BrBBrMP and 100 mL of ethanol. The flask was then immersed in a water bath set at 6 ± 1 °C and 100 mL of a 1.77 M solution of NaOH in ethanol were added drop by drop in about 0.5 h. Once all the NaOH solution was fed to the reactor, the bath temperature raised to 70 °C and the solution was kept under stirring for further 1 h. Then, the solution was cooled to room temperature, filtered under reduced pressure (6500 Pa) and mixed with 50 mL of DCM. The organic phase was washed several times with distilled water and BBrMO was finally obtained after distillation at 30,000 Pa and 61 °C, with a 93.0 % yield.

2.2.3. 3-tosyloxymethyl-3-methyl oxetane (TMMO)

The synthesis was performed in the solid state, according to a procedure described by Kazemi et al [56]. A mortar was fed with 31.8 g of anhydrous Na₂CO₃, 10.2 g of HMMO

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3 (Na₂CO₃/HMMO=3/1 molar ratio) and manually milled with a pestle for about 7 min. Then
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5 28.7 g of TsCl (TsCl/HMMO=1.5/1 molar ratio) were added and milled for another 23 min.
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7 Finally, 28.05 g of KOH were added (KOH/HMMO=5/1 molar ratio) and milled for about 50
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9 min to remove the unreacted TsCl. The whole reaction was conducted in a glove box under
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11 nitrogen atmosphere. The obtained solid was added to 200 mL of diethyl-ether and filtered
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13 several times. The obtained clear liquid phase was finally subjected to distillation at 48°C and
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15 8500 Pa thus inducing the crystallization of TMMO that was obtained with a 40 % yield.
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21 The structure and purity of all monomers was verified by Fourier transform infrared
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23 spectroscopy (FTIR) and by nuclear magnetic resonance (¹H-NMR).
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27 *2.3. Polymerizations*

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29 As stated above, the block copolymers formed through the growing of the energetic segments
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31 on each end of preformed HTPB, act as initiator. The catalyst and operating conditions were
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33 set in order to favor the cationic ring opening polymerization by AMM. In all the
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35 polymerizations, the quantities of BTF THF and HTPB were such that the molar ratio
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37 between BTF THF and the OH functionalities of the polymer was equal to 2/1, while the
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39 molar ratio between monomers and OH functionalities was equal to 50/1. Table 1 reports the
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41 quantities used for a typical polymerization of each block copolymers, being in the last
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43 column indicated the theoretical N₃ content after azidation, calculated from the hypothesis of
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45 100 % yield in both the polymerization and azidation reactions. The polymerization procedure
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47 was as follows: A 1L three-necked round bottom flask, fitted with a reflux condenser, a
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49 nitrogen inlet and a mechanical stirrer set at 120 rpm was fed with 80 ml of DCM, 0.72 mL of
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51 BTF THF, about 4 g of HTPB and maintained 2 h under stirring at room temperature. Then,
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53 the reactor was covered with aluminum foil, immersed in a water bath at 20±0.5 °C and the
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3 monomers, previously dissolved in 30 mL of DCM, were added drop-wise to the reactor
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5 through a dropping funnel. From the end of the monomer feeding, the reaction mixture was
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7 kept in the same conditions for 20 h (120 h when using tosylated monomers) and then
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9 hydrolyzed with 400 mL of an aqueous solution of NaCl (10% w). The mixture was
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11 maintained under vigorous stirring for further 2 h and then the organic and aqueous phases
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13 were separated. The aqueous phase was washed with DCM, subsequently recovered and
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15 added to the organic phase. The organic phase was washed once with a water/methanol =
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17 50/50 v/v solution to remove unreacted monomers and catalyst, then several times with the
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19 aqueous solution of NaCl, dried with MgSO₄, filtered and distilled at 45 °C, under vacuum
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21 (6500 Pa) in order to remove all the DCM. All the polymers were obtained with a 95-97 %
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23 yield.
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29 30 *2.4 Azidation of the polymeric precursors*

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32 The azidation was conducted with the well-known second-order reaction with SN₂ type
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34 mechanism, by using sodium azide and a polar solvent. This technique of azidation is almost
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36 “universal” and in the last decades many researchers, which tested several solvents and
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38 operating conditions as well as many different leaving groups, described it. In our research
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40 group, it was previously used to substitute chlorine, bromine and tosyl groups, in either DMF
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42 or DMSO, with temperatures ranging from 90 up to 150 °C [37, 38, 40, 43]. Pisharat and Ang
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44 azidated PECH in DMF, at 120 °C for 12h [39], while Manser suggested DMSO at 80 °C also
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46 for the azidation of the oxetanic monomers [33]. Many other examples of azidic compounds
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48 can be found in the literature, with DMF and DMSO the most used solvents. However, we
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50 found that neither DMF nor DMSO were suitable for the azidation step, because before
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52 obtaining the complete azidation of the polymers, an incipient cross-linking reaction was
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54 observed, thus leading to partially soluble materials. This happened in several attempts made
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3 at testing temperatures ranging from 100 to 130 °C. The addition of small quantities of
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5 Vulkanox BKF as antioxidant only reduced the phenomenon. Since this procedure is well
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7 established and was already tested several times in our laboratory for GAP, BAMO and
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9 PAMMO polymers and copolymers, which never caused such inconvenience, the cross-
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11 linking must be correlated with the presence of the unsaturations in the polybutadiene middle
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13 block. However, Vasudevan and Sundararajan, which used DMSO at 105 °C for 10 h, did not
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15 describe the cross-linking problem in their work [52]. In our case, the use of a toluene/DMAc
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17 mixture, as suggested by Subramanian [53], in combination with small quantities of Vulkanox
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19 BKF, gave good results and was adopted for all azidations.
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23 A typical azidation procedure was as follows: About 2 g of polymeric precursor were
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25 dissolved in 100 ml of DMAc/toluene (50/50 v/v) solution and fed in a 250 mL three-necked
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27 round bottom flask, fitted with a reflux condenser, a nitrogen inlet and a magnetic stirrer.
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29 After adding a small quantity of Vulkanox BKF (about 1% by weight with respect to the
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31 polymer), the flask was immersed in an oil bath set at 90 ± 1 °C and conditioned for 30
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33 minutes. Then, NaN_3 was added in 10% molar excess with respect to the stoichiometric
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35 quantity and the temperature of the bath raised to 95 °C while maintaining the system under
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37 constant stirring. Periodically, samples were taken from the reactor and subjected to FTIR
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39 analysis to evaluate the degree of azidation. Once the IR spectra reached a stationary “state”,
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41 the reaction medium was filtered under vacuum in order to remove the formed sodium salt’s
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43 and the unreacted sodium azide. The reaction time strongly varied depending on the
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45 monomeric precursors (Table 2).
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50 The solution was then washed several times with an aqueous solution of NaCl (10 % w) in
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52 order to complete the salts removal. The aqueous phases resulting from the washings were
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54 mixed together and washed with DCM to remove possible traces of polymer, while the
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56 organic phase was dried with MgSO_4 , filtered again and distilled at 45 °C, under vacuum
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(1000 Pa) in order to remove all the organic solvents. A small quantity (about 1.5 % by weight) of Vulkanox BKF was finally added to the polymer before storage. All the azidated copolymers were obtained with a yield higher than 95%.

The structure and purity of all synthesized polymers (before and after azidation) was checked by FTIR and NMR, while their thermal properties were determined by DSC and TGA.

2.5. Characterization of the monomers and polymers

Fourier transform infrared spectroscopy was performed on a Bruker Tensor 27 and nuclear magnetic resonance ($^1\text{H-NMR}$, $^{13}\text{C-NMR}$) on a VXR300 and INOVA600 instruments.

Chemical analysis and FTIR Chemical Imaging were carried out by Spectrum Spotlight FTIR Imaging System from Perkin Elmer. Chemical imaging analysis, in transmittance mode, allowed combining optical microscopy and infrared analysis of micro and macro areas of thin films prepared by solvent evaporation on a microscope slide. The morphological analysis was performed also by a LEICA DM LB fluorescence microscope. The molecular weight distributions of the polymers were measured from solution in CHCl_3 (4 mg/mL) by using a Gel Permeation Chromatography (GPC) apparatus Jasco PU-1580, equipped with PL Mesopore column, calibrated with low polydispersity polystyrene standards.

Thermogravimetric analysis (TGA) was done by using a TA Q500 apparatus, under nitrogen atmosphere, with a heating rate of $10^\circ\text{C}/\text{min}$ until 600°C and using samples of about 5 mg.

Thermal properties were studied by differential scanning calorimetry (DSC) performed with a Pyris 1 scanning calorimeter from Perkin Elmer, by using aluminum pans. The sample mass was in the range of 3-5 mg and spectra were collected from 50°C at a heating rate of $10^\circ\text{C}/\text{min}$, until 350°C .

3. Results and discussion

3.1. Chemical characterization (IR – NMR)

IR and NMR spectra were recorded for all monomers, precursors and polymers synthesized in this work. In order to limit the number of figures, we report only two H-NMR spectra, showing the comparison between GT/BBrMO-PB-GT/BBrMO and GA/BAMO-PB-GA/BAMO in the range corresponding to the tosyl group signals (Figure 3).

With regard to the IR spectra, they are reported in Figures 4 and 5, showing the main peaks involved in the azidation process. In all cases, but one, azidation appeared to be quantitative. The only polymer without complete substitution of the leaving groups is AMMO-PB-AMMO, where the peaks relative to the tosyl group are clearly visible even after 140 h of azidation reaction. A quantitative evaluation of the residual tosyl groups indicates a degree of azidation close to 50 %. The peaks highlighted in the figures are: 554 cm^{-1} = N_3 bending, 670 cm^{-1} = C-Br stretching, 744 cm^{-1} = C-Cl stretching, 1190 cm^{-1} = SO_2 symmetric stretching, 1280 cm^{-1} = N_3 symmetric stretching, 1363 cm^{-1} = SO_2 asymmetric stretching, 1600 cm^{-1} = aromatic C=C vibration, 2100 cm^{-1} = N_3 asymmetric stretching.

From the reaction times reported in Table 2, it can be concluded that the azidation of the tosyl group is the bottleneck of the reported synthetic strategies. This is somehow unexpected, because the tosyl is a good leaving group. However, it should be considered that the group is attached to a polymeric chain, so there are several aromatic rings close one to each other and this may determine a high steric hindrance as well as a low mobility of the chains even when in solution. In the case of the GT/BBrMO copolymer, probably the alternation of GT and BBrMO groups allows sufficient mobility of the chain, so that a complete azidation was obtained in a short time. Therefore, even if potentially better than halogen atoms, the tosyl group may determine a slowing down of the azidation kinetics and also limits the maximum degree of azidation obtainable within the reaction times adopted in this work.

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3 An interesting point concerning FTIR is that a comparison of the GA-PB-GA spectrum
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5 (labeled as N. 6 in Figure 5) with the corresponding one reported in the paper by Subramanian
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7 [53] clearly shows that in the latter the N_3 peak is significantly smaller. This means that the
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9 synthetic procedure adopted in this work enables formation of copolymers with a significantly
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11 higher azide content.
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14 With regard to the characterization of the copolymers, during the revision process it was
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16 asked if the alcohol moieties in the HTPB chain were really used as the primary initiator site
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18 rather than adventitious moisture. In the latter case, we would have obtained the polyether
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20 instead of a block copolymer. Therefore, depending on the primary initiator, the final product
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22 may contain block copolymer, polyether and HTPB. Block copolymer alone is the desired
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24 product, with all HTPB used as initiator and no moisture involved. A mechanical mixture of
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26 block copolymer and polyether will form if both HTPB and moisture act as initiator. A
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28 mechanical mixture of block copolymer and HTPB is obtained if HTPB is only partially used
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30 as initiator. A mechanical mixture of HTPB and polyether is the worst case, with no HTPB
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32 used as initiator. And so on, with other combinations possible. To clarify this point, we can
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34 start from the above-described incompatibility between the ether and butadiene blocks [44],
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36 which leads to a macroscopic phase separation when blended. For this reason, the
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38 morphologies of the polymeric precursors were compared with those of the mechanical blends
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40 prepared from the polymers constituting the single blocks. For this purpose, the ether blocks
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42 were appositely synthesized, by using the same synthetic strategy described for the block
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44 copolymers, simply substituting HTPB with butanediol. Particular care was taken in order to
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46 assure the use of identical operating conditions during polymerizations (reagent purity,
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48 volumes, solvent quantities and so on). The mechanical blends (MB) were prepared by
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50 solvent evaporation from a DCM solution where HTPB and polyether were added in
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52 quantities corresponding to the composition calculated from the yields of the copolymer
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3 synthesis. Moreover, MB and copolymers were subjected to fluorescence microscopy and
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5 Spectrum Spotlight FTIR Imaging. As an example of the obtained results (qualitatively
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7 similar for all the synthesis and corresponding MB), Figure 6 shows the images and spectra
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9 recorded for PGT and HTPB. The MB is on the left hand side of the Figure and the product of
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11 the synthesis (named product in what follows) is on the right hand side. The fluorescence
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13 microscopies of MB (Figure 6a) and the product (Figure 6b) are completely different. The
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15 latter is quite homogeneous and shows some area with slightly different fluorescence
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17 intensity, but no visible phase separation. In contrast, the MB is markedly biphasic, with dark
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19 roundish drops dispersed in a bright matrix. A comparison of these images with those (not
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21 reported) of the homopolymers suggests that the dark phase corresponds to HTPB. However,
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23 the dark phase and brightness inhomogeneity may also derive from voids and/or variations of
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25 the sample thickness. Therefore, the samples were analyzed by Spectrum Spotlight FTIR
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27 Imaging, which uses optical microscopy (Figures 6c and 6d) and allows building a chemical
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29 maps (Figures 6e and 6f). The maps represent the normalized peak area between 2980 and
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31 2700 cm^{-1} (stretching of aliphatic C-H bonds). This peak is present in both polyether and
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33 HTPB, but its intensity is significantly higher in the latter. Therefore, the grey level in the
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35 map reflects the composition of the sample: a high HTPB content corresponds to white color
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37 in the map. Figure 6e shows areas with markedly different grey level and confirms that the
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39 round domains of MB are mainly composed of HTPB. In contrast, Figure 6f does not show
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41 any significant variations in grey level. Moreover, the arrows indicate the spectra recorded in
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43 correspondence of the “+” symbols reported in Figures 6e and 6f. The two spectra on the left
44
45 hand side differ one from each other and result similar to those of the polyether and HTPB
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47 homopolymers (not reported). In contrast, the right hand side spectra are very similar, even if
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49 recorded in areas corresponding to the maximum possible difference in grey scale. Based on
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51 the IR spectra and maps, we can conclude that the shadows and apparent inhomogeneity
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3 observable in Figures 6b, 6d and 6f do not correspond to significant variation in chemical
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5 composition. Therefore, we can exclude that the reaction product is a mechanical mixture of
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7 HTPB and polyether.
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10 Moreover, a phase separation is observable also while adding very small quantities of HTPB
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12 to the product. This excludes also that the product could be a mechanical mixture of block
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14 copolymer and HTPB. The last proof that all HTPB was used to build the polyether blocks
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16 comes from an extraction with n-hexane. The mechanical blend was immersed in n-hexane,
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18 and stirred vigorously at room temperature for a few minutes, after which a consistent part of
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20 the blend showed to be immiscible with the solvent. Then, the hexane-phase was recovered
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22 and the solvent evaporated. The resulting polymeric phase was HTPB alone. The same
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24 procedure was followed for the copolymer and, in that case, no HTPB was recovered after
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26 hexane evaporation.
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29 The presented results demonstrate that HTPB worked as initiator for the polyether, but leave
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31 the doubt that the ether could polymerize starting from the above-mentioned water moieties.
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33 This possibility is validated from the fact that the addition of small quantities of polyether to
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35 the synthesis product did not give phase separation (contrary to what happened while adding
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37 HTPB). Unfortunately, this hypothesis could not be verified with solvent extraction, because
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39 we did not find a solvent selective toward the ether phase. Several polar solvents were tried,
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41 but all of them partially dissolved the HTPB, probably due to its OH functionalities. A more
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43 enhanced extraction procedure was necessary and the GPC column showed to be useful.
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47 As an example, Figure 7 shows the GPC analysis of HTPB and GT-PB-GT copolymer. The
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49 weight molecular weight distribution of the copolymer resulted bimodal, and the curve was
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51 subjected to deconvolution (dashed lines). The obtained results are summarized in Table 3
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53 and show that the right hand peak of the copolymer has a molecular weight slightly higher
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55 than HTPB. In contrast, the left hand peak has a very low molecular weight, thus
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3 remembering the values usually associated with the formation of cyclic oligomers composed
4 of 3-5 repeating units [40]. Therefore, the correct interpretation of the GPC is that the high-
5 molecular weight peak is the block copolymer, while the low-molecular weight peak is due to
6 the presence of polyether not linked to HTPB and grown from OH functionalities of different
7 origin. The synthesis product is a mechanical mixture of block copolymer and polyether, thus
8 both HTPB and some adventitious moisture worked as initiator. The polyether content was
9 calculated from the area of the peaks and the results are reported in the last column of Table 3.
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11 In all cases, the oligomer content is around 20 % by weight of the total amount.
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13 We can thus affirm that all the HTPB chains acted as initiator for the polymerization, but
14 some of the OH moieties present in the reaction medium did the same and led to the
15 formation of polyether chains not covalently linked to the polybutadiene. The product thus
16 contains polyether oligomers that are difficult to separate from the copolymer. Nevertheless,
17 if these oligomers are linear, they should be OH functionalized and participate to the curing
18 with isocyanates while forming the polymeric network that constitutes the final binder.
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20 Otherwise, in case they are cyclic oligomer, they will act as plasticizers in the binder
21 formulation.
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43 *3.2 Thermal characterization (DSC – TGA)*

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45 In the scientific literature there is a long list of papers that analyze the thermal decomposition
46 of polymers (especially GAP) containing the azidic functionalities, alone or after curing and
47 addition of plasticizers [17, 20, 21, 23, 25, 31, 57]. Usually, the easiest way to determine the
48 thermal stability of this kind of polymers is by TGA and DSC analysis. However, due to the
49 violent kinetics of decomposition, it may not be easy to obtain reliable and reproducible
50 results. Selim et al., in the case of GAP, used 2-3-mg samples, and kept the ramp rates in the
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3 range of 0.2-5°C/min since a greater amount of polymer or higher heating rate lead to
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5 explosion during the decomposition experiment [14]. In all cases, while subjected to a
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7 temperature sweep procedure, the azidic polymers degrade in a two-stage process. The first
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9 one, occurring at about 240-270 °C, is exothermic and corresponds to the abstraction of N₃
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11 groups from the polymeric chain. The second one, partially overlapped with the first one, but
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13 shifted to higher temperatures, generally occurs without significant heat release and
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15 corresponds to the progressive decomposition of the remaining polymeric chain. In one of the
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17 first and most cited work, Kubota and Sonobe suggested that the first process corresponds to
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19 the release of a nitrogen molecule from the N₃ group, with the formation of a nitrile C≡N
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21 bond [58]. The following studies confirmed that the decomposition starts with the initial
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23 rupture of N-N₂ bond, with elimination of molecular nitrogen. Then, rearrangement to a
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25 polymeric imine and/or acrylonitrile following elimination of hydrogen molecule has been
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27 proposed. In the final products of decomposition, gaseous (N₂, CO, HCN, NH₃, CH₂O, CH₄,
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29 C₂H₂, C₂H₄, CH₃CHO, CH₂CHCHNH, CH₃CHNH, H₂O etc.) and larger molecules as
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31 benzene, pyrrole and furan have been identified [18, 23, 24]. However, in all cases, the weight
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33 loss experimentally measured during the first stage is higher than the theoretical one based on
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35 sole N₂ release. This is due to the above-mentioned partial overlapping with the incipient
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37 second stage of decomposition, but also to the occurrence of other, parallel reactions during
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39 the first stage. Various mechanisms and possible involved reactions have been proposed by
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41 analyzing the residual solid and the evolved gases during decomposition. As an example,
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43 Eroğlu and Güven used FTIR and observed that the decrease of N₃ content was accompanied
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45 by the appearance of signals related to N-H bending, -C=N-C-H, -C=N-C-C, and N-H
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47 stretching. At the same time, the signal of the C-O-C ether bridge remained unchanged thus
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49 suggesting that the main chain was not yet involved in the decomposition process. Based on
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51 these observations, a mechanism based on the formation of intra and inter-molecular cross-
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3 linking was suggested for the first stage of degradation [13]. Reshmi et al. used DSC and
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5 TGA coupled with pyrolysis gas chromatography–mass spectrometric technique and showed
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7 that during its first stage of decomposition, 1,6-bis (azidoacetoxy) hexane preferentially
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9 forms the corresponding diimine by elimination of N_2 [59]. Then, at higher temperatures,
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11 there is formation of diimines by elimination of CO_2 , diols through elimination of CO and
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13 HCN and diene due to CO_2 and CH_2NH elimination. The photodegradation under ultraviolet
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15 (UV) irradiation was also investigated by Sahu et al. [15] and by Wang et al. [19]. The latter
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17 combined a tunable synchrotron vacuum ultraviolet photoionization and molecular-beam
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19 sampling mass spectrometry and observed that the UV radiation lowered the onset of the
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21 thermal decomposition of GAP and caused the appearance of a larger number of free radical
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23 species, compared to thermal decomposition alone.
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27 The DSC and TGA analysis are reported in Figures 8-10 and summarized in Tables 4 and 5. In
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29 both Tables, T_i and T_f indicate the initial and final temperatures of the degradation step. The
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31 produced heat and weight loss are reported after normalization with respect to both the total
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33 weight and the theoretical N_3 content of the polymer. In Table 5, Δw is the weight loss with
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35 respect to the total initial mass and $\Delta w N_3$ (%) is the weight loss compared to the theoretical
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37 one expected if all azidic groups release one N_2 molecule. Finally, W_r is the residual weight at
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39 the end of the test (600 °C). The DSC traces are reported only for the azidic polymers. It is
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41 useful to underline that such high temperatures have only scientific interest, not practical,
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43 because the behavior of the polymers in propellant will not be the same as in the individual
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45 state due to the oxidizer presence around.
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49 As already stated, the N_3 group decomposition is responsible for the exothermic peak
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51 observed in DSC and the first weight loss in the TGA thermograms. However, this
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53 decomposition may follow different pathways and be partially superposed to other
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55 degradation phenomena, which depend not only on the operating conditions, but also on the
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3 polymer structure. Therefore, the weight loss and energy release during this stage are
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5 unequivocally determined in the literature. As an example, in the case of GAP, the following
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7 values (here expressed as J/g of N₃) were reported: 4,352 [13], 4,835 [14] and 6,166 [21],
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9 representing the heat of formation of the azido group around 8500 J/g of N₃ [27]. In our case,
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11 the decomposition is probably “complicated” by the presence of the butadiene block that may
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13 interact with the formed species. The enthalpies reported in Table 4 are substantially in
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15 agreement with those reported in the literature. The value obtained for the AMMO-PB-
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17 AMMO copolymer confirms the estimation of about 50% substituted tosyl groups. Moreover,
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19 in the spectrum of this polymer there is the superposition with a second peak (centered at
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21 about 275 °C) that corresponds to the exothermic degradation of the tosyl group and well
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23 visible in the spectra of the tosylated precursor (not reported here).
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27 With regard to TGA (Figures 9 and 10), a first general observation is that all the azidated
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29 copolymers, but AMMO-PB-AMMO, have a significant residual weight at 600 °C (see last
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31 column of Table 5). In contrast, the precursors show a much lower final weight and HTPB
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33 alone has almost no residual weight at the same final temperature. This difference between
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35 azidated polymers and precursors suggests that during thermal decomposition some chemical
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37 interactions may occur between the azidic functionalities and the butadiene backbone.
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39 Probably, this is strictly connected with the above-mentioned tendency of the copolymers to
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41 cross-link during azidation and subsequent storage. It may be supposed that the first
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43 decomposition products interact with the butadiene main chain and form a highly cross-linked
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45 network that slows down the degradation process at higher temperatures. Of course, the
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47 explanation of different behavior of the AMMO-PB-AMMO copolymer can be based on its
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49 low degree of azidation. This difference among azidated copolymer and non-energetic
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51 precursors is coherent with the shape of the TGA curves. The precursors show a two-step
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53 degradation, well separated by a horizontal plateau. A comparison with the HTPB curve
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3 suggests that the first step is relative to the ether block, while the second one is the butadiene
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5 block. The azidic copolymers show again a two-step degradation, but differently shaped. The
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7 first step, already attributed to the N_3 functionality, is sharp, while the second one corresponds
8
9 to a gradual weight loss that extends in a wide temperature range and ends at about 500 °C.
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11 Again, AMMO-PB-AMMO copolymer is the only one with an intermediate behavior (curve 8
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13 in Figure 10 shows three distinct steps).

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16 The weight losses during the first decomposition step are higher than expected if each N_3
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18 functionality releases one nitrogen molecule (Table 5). This confirms that the main chain is
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20 already subjected to other degradation phenomena (mainly ascribable to the ether block) that
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22 superpose to the degradation of the azidic functionalities.
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25 A last point to be underlined is that polymers 2 and 3 in Figure 9 have very slight differences
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27 in the TGA and DSC curves, which is a further confirmation of their substantial equivalence.
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29 The AMMO-PB-AMMO copolymer is the one that shows the lower enthalpy release during
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31 decomposition of the azidic functionalities. This is in accordance with the already observed
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33 incomplete substitution of the tosyl functionalities with the azidic ones. From DSC and FTIR
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35 we already calculated that this corresponds to about one-half of the theoretical azidic
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37 functionalities. However, this value must be analyzed by taking into account also the TGA
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39 curves. The TMMO-PB-TMMO precursor shows two very distinct and sharp peaks
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41 corresponding to the degradations of the ether and butadiene blocks. However, the weight loss
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43 (16.6 %) associated to the latter is higher than the theoretical value (8.0%) corresponding to a
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45 100% conversion during polymerization, thus suggesting that the TMMO blocks may be
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47 shorter than expected. Therefore, with respect to the expected structure, the AMMO-PB-
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49 AMMO copolymer may have both shorter ether blocks and incomplete degree of tosyl
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51 functionalities. After N_3 decomposition, the weight loss of the azidic copolymer shows two
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53 partially superposed consecutive steps related to ether and HTPB blocks degradation.
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4. Conclusions

The synthesis of ether-butadiene-ether energetic block copolymers was done by a classical cationic homo and co-polymerization of oxetanes and oxiranes, using HTPB as diol-initiator and followed by the azidation process. Four copolymers were synthesized, with the external linear blocks constituted by azido homo and copolymers. Probably due to the presence of water moieties, a small amount of polyether oligomers not linked with HTPB were also obtained as undesired side products. With respect to the corresponding polymers prepared by using a low molecular weight, saturated diol, such as i.e. 1-4 butanediol, the use of HTPB complicates the synthetic procedure, since the azide could have a side reaction with the double bonds to cause some species, which could lead to cross-linking. This may lead to the formation of a cross-linked block copolymer, which, of course, is not suitable as polymeric binder for solid propellants. The cross-linking phenomenon may happen either during the polymerization, azidation or purification steps, or also during prolonged storage at room temperature. This restricts the range of operating conditions available for the synthetic process, limiting the suitable solvents, as well as the temperatures and duration of the reactions and purification procedures. Moreover, the addition of an anti-ageing compound to the final product and the storage in an inert and dark atmosphere is necessary. The tosyl group may slow down the kinetics of the azidation step and may also lead to incomplete substitution. In conclusion, the synthesis of these azidic copolymers with an internal unsaturated block is not simple and presents several drawbacks not easy to manage. Nevertheless, the appropriate choice of the operating conditions allows these drawbacks to be partially overcome and results in a versatile tool for the synthesis of polymeric binders, whose properties can be modulated by changing the type and length of the ether blocks.

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Figure captions

- Figure 1 Synthesis of the monomers. G = Glycidol; GT = glycidol tosylate; HMMO = 3-hydroxy-methyl-3-methyloxetane; TMMO = 3-tosyloxymethyl-3-methyl oxetane; BrBBrMP = 3-bromo-2,2-bis(bromomethyl)propanol; BBrMO = 3,3-bis(bromomethyl)oxetane.
- Figure 2 Synthesis and azidation of the block copolymers.
- Figure 3 H-NMR spectra of: (1) GT/BBrMO-PB-GT/BBrMO and (2) GA/BAMO-PB-GA/BAMO from (1).
- Figure 4 FT-IR spectra of: 1) GT/BBrMO-PB-GT/BBrMO; 2) GA/BAMO-PB-GA/BAMO from 1; 3) GA/BAMO-PB-GA/BAMO from 4; 4) ECH/BBrMO-PB-ECH/BBrMO.
- Figure 5 FT-IR spectra of: 5) GT-PB-GT; 6) GA/PB/GA; 7) TMMO-PB-TMMO; 8) AMMO-PB-AMMO.
- Figure 6 GT-HTPB. Fluorescence microscopy of MB (a) and synthesis product (b); optical microscopy of MB (c) and synthesis product (d); chemical map of MB (e) and synthesis product (f). FTIR spectra corresponding to the positions indicated by the + symbols in (e) and (f).
- Figure 7 GPC analysis of HTPB and GT-PB-GT copolymer. The dashed lines represent the deconvolution peaks of the GT-PB-GT distribution.
- Figure 8 DSC traces of the first decomposition step for the azidated copolymers.
- Figure 9 TGA analysis of HTPB, GA/BAMO-PB-GA/BAMO copolymers and their respective precursors.
- Figure 10 TGA analysis of HTPB, GA-PB-GA, AMMO-PB-AMMO copolymers and their respective precursors.

Table 1 - Composition of the reaction mixtures for the polymerizations.

polymer	HTPB (g) (mole)	BBrMO (g) (mole)	ECH (g) (mole)	GT (g) (mole)	TMMO (g) (mole)	BTF THF (g) (mole)	DCM (mL)	N ₃ (w%)
GT-PB-GT	1.5 $5.3 \cdot 10^{-4}$	-	-	13.65 0.06	-	0.336 0.0024	110	35
GT/BBrMO-PB-GT/BBrMO	1.48 $5.3 \cdot 10^{-4}$	3.6 0.015	-	10 0.044	-	0.336 0.0024	110	38
ECH/BBrMO-PB-ECH/BBrMO	3.99	10.0	11.4	-	-	0.91 0.0065	110	38
TMMO-PB-TMMO	0.99 $3.5 \cdot 10^{-4}$	-	-	-	10.1 0.039	0.21 0.0015	110	28

Table 2 – Reaction time for the azidation of the copolymers.

monomeric precursor	GT	GT/BBrMO	ECH/BBrMO	TMMO
Azidation time (h)	140	22	16	140

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Table 3. GPC results. Mn= number average molecular weight; Mw= weight average molecular weight.

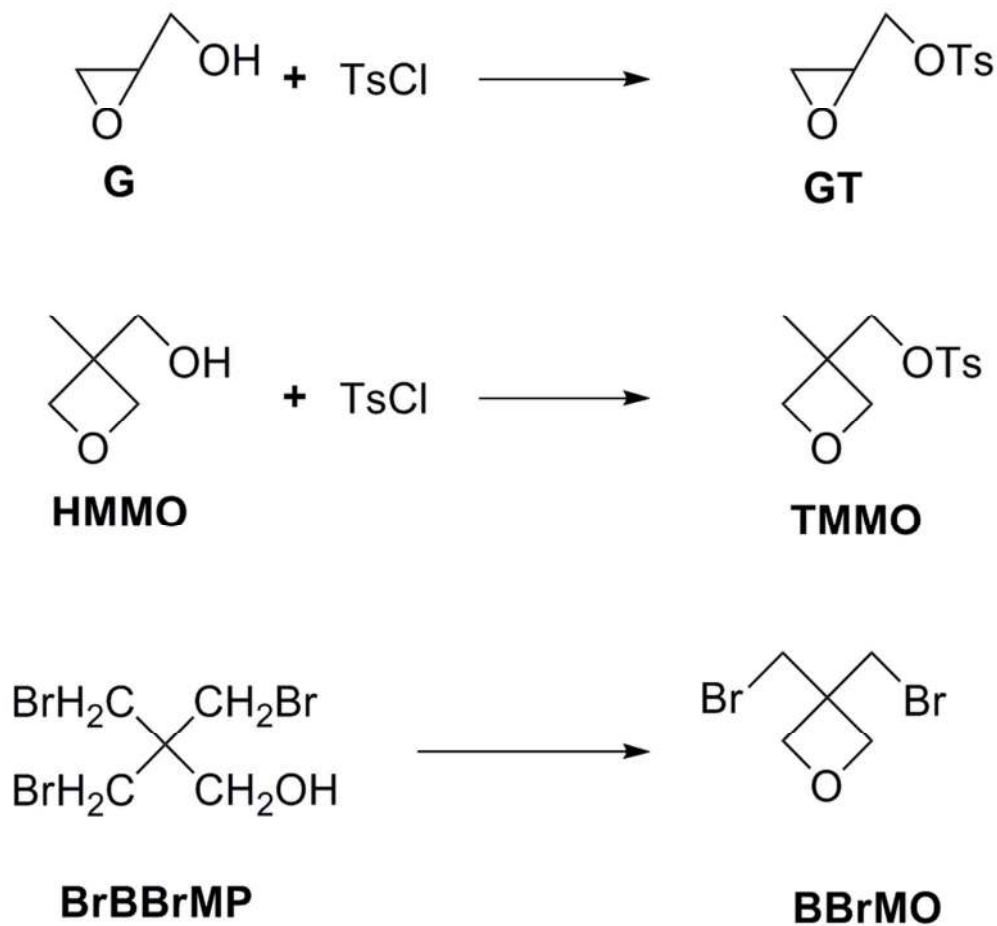
polymer	Mn (g/mole)	Mw (g/mole)	Mn 1 st peak (g/mole)	Mw 1 st peak (g/mole)	Mn 2 nd peak (g/mole)	Mw 2 nd peak (g/mole)	Polyether content (% wt)
HTPB	4700	11400	-	-	-	-	-
GT-PB-GT	1400	6230	1180	1340	6770	14850	21
ECH/BBrMO-PB-ECH/BBrMO	610	13420	550	810	6650	15200	22
GT/BBrMO-PB-GT/BBrMO	860	4990	750	1340	6930	14330	20
TMMO-PB-TMMO	1850	9430	900	1020	7060	16200	15

Table 4 – DSC data corresponding to the first decomposition step.

Polymer	T _i (°C)	T _f (°C)	ΔH (J/g)	ΔH (J/g N ₃)
GA-PB-GA	205	270	1,380	3,940
GA/BAMO-PB-GA/BAMO from ECH/BBrMO-PB-ECH/BBrMO	175	240	1,740	4,575
GA/BAMO-PB-GA/BAMO from GT/BBrMO-PB-GT/BBrMO	210	270	1,875	4,940
AMMO-PB-AMMO	175	255	630	2,250

Table 5 – TGA data for energetic polymers and corresponding precursors.

Polymer	N ₃ degradation			W _r
	T _i -T _f (°C)	Δw (%)	Δw N ₃ (%)	
HTPB				0.5
GT-PB-GT				4.7
GA-PB-GA	210-280	47	201	15
ECH/BBrMO-PB-ECH/BBrMO	-	-		2.2
GA/BAMO-PB-GA/BAMO from ECH/BBrMO-PB-ECH/BBrMO	175-275	41.2	164	24.1
GT/BBrMO-PB-GT/BBrMO				6.1
GA/BAMO-PB-GA/BAMO from GT/BBrMO-PB-GT/BBrMO	200-275	43.8	175	25.6
TMMO-PB-TMMO				6.3
AMMO-PB-AMMO	159-250	12.5	70	4.2



38 Figure 1 - Synthesis of the monomers. G = Glycidol; GT = glycidol tosylate; HMMO = 3-hydroxy-methyl-3-
39 methyloxetane; TMMO = 3-tosyloxymethyl-3-methyl oxetane; BrBBRMP = 3-bromo-2,2-
40 bis(bromomethyl)propanol; BBRMO = 3,3-bis(bromomethyl)oxetane.
41 81x76mm (300 x 300 DPI)

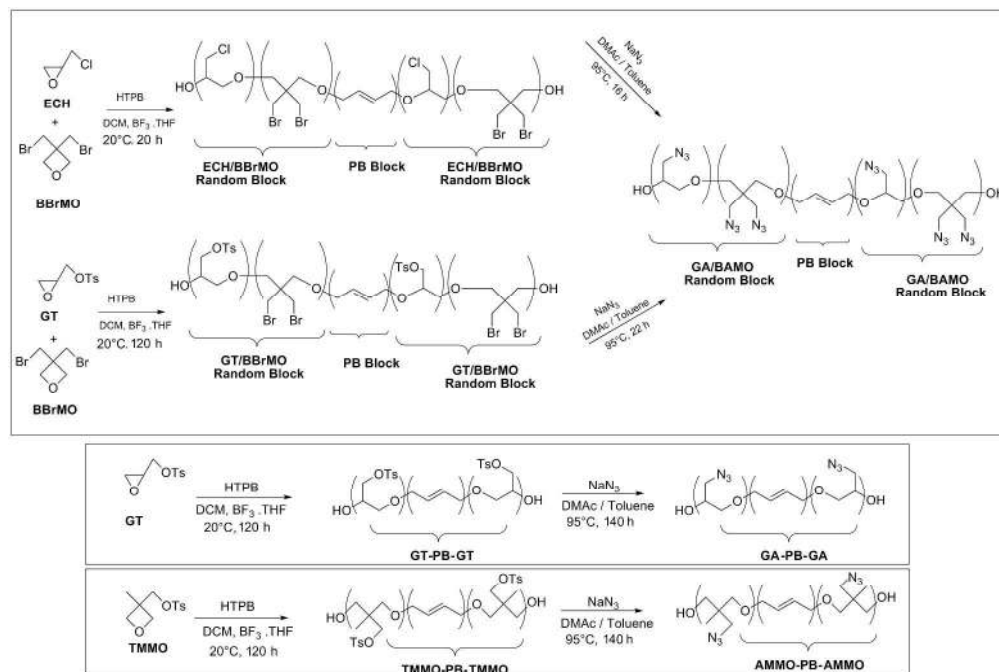


Figure 2 - Synthesis and azidation of the block copolymers.
272x183mm (300 x 300 DPI)

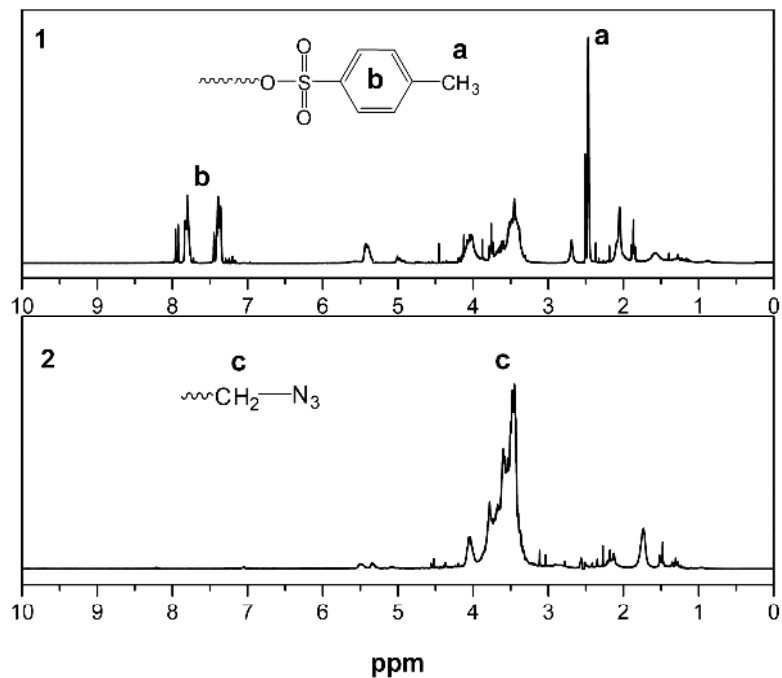


Figure 3 - $^1\text{H-NMR}$ spectra of: (1) GT/BBrMO-PB-GT/BBrMO and (2) GA/BAMO-PB-GA/BAMO from (1).
227x174mm (300 x 300 DPI)

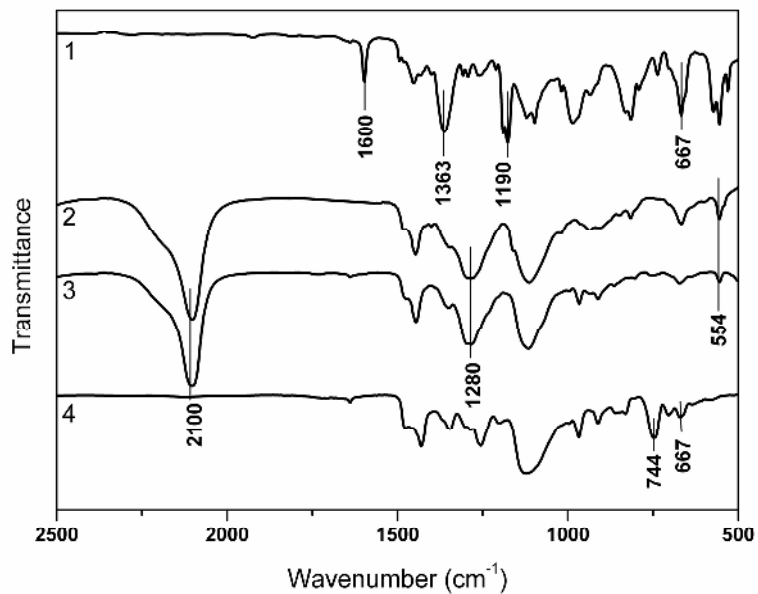


Figure 4 - FT-IR spectra of: 1) GT/BBrMO-PB-GT/BBrMO; 2) GA/BAMO-PB-GA/BAMO from 1; 3) GA/BAMO-PB-GA/BAMO from 4; 4) ECH/BBrMO-PB-ECH/BBrMO.
207x144mm (300 x 300 DPI)

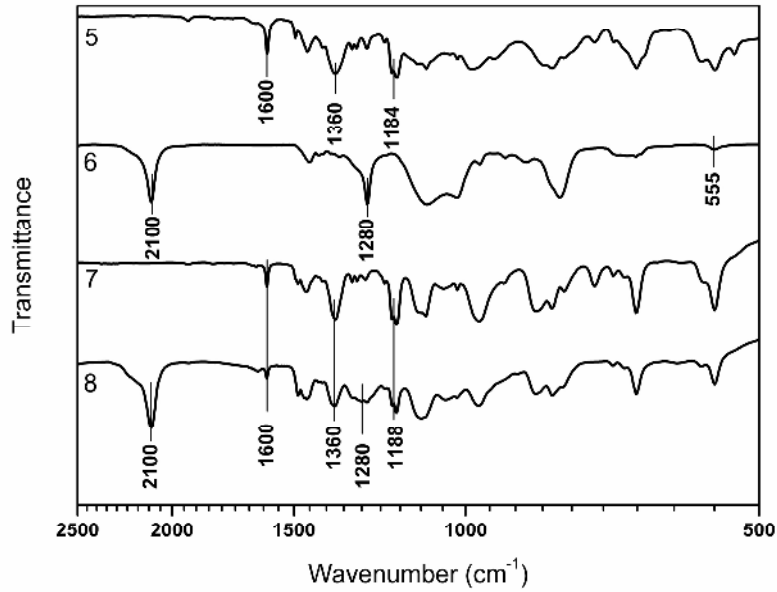


Figure 5 - FT-IR spectra of: 5) GT-PB-GT; 6) GA/PB/GA; 7) TMMO-PB-TMMO; 8) AMMO-PB-AMMO.
207x144mm (300 x 300 DPI)

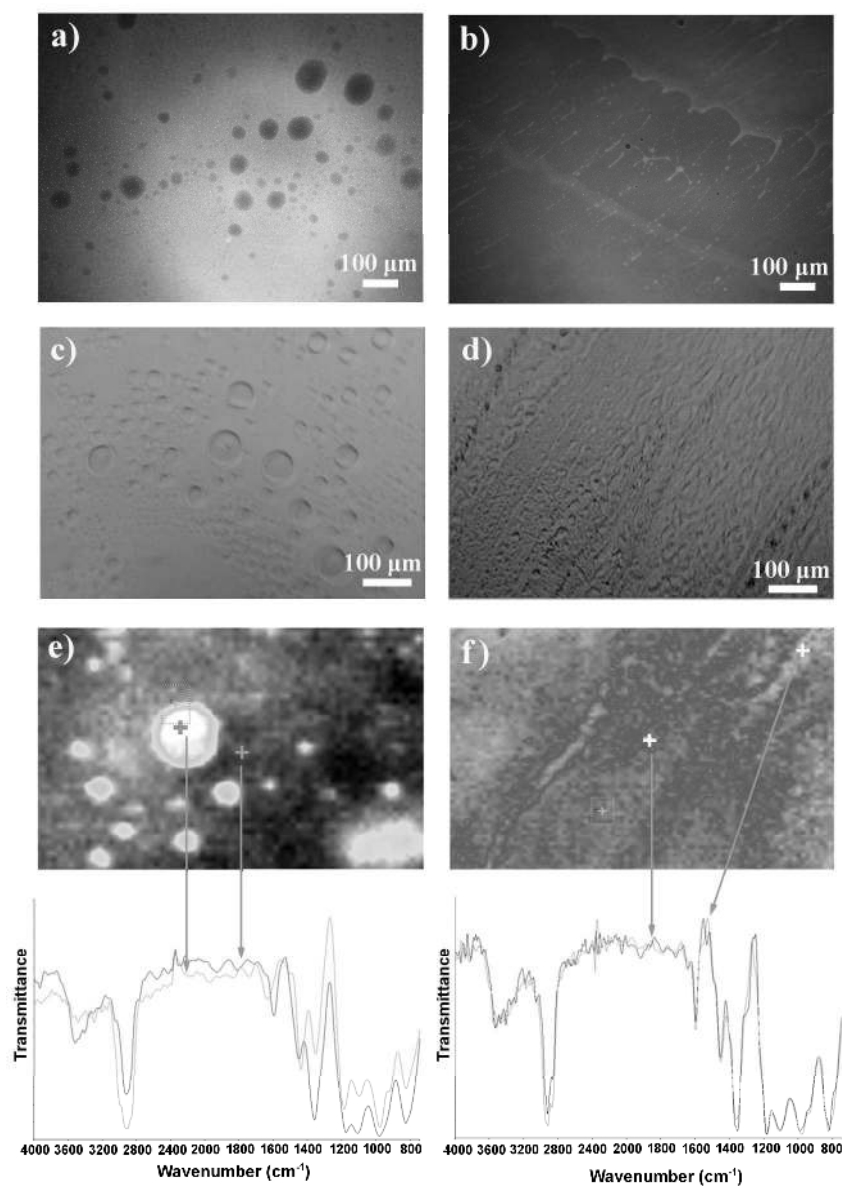


Figure 6 GT-HTPB. Fluorescence microscopy of MB (a) and synthesis product (b); optical microscopy of MB (c) and synthesis product (d); chemical map of MB (e) and synthesis product (f). FTIR spectra corresponding to the positions indicated by the + symbols in (e) and (f).
175x249mm (300 x 300 DPI)

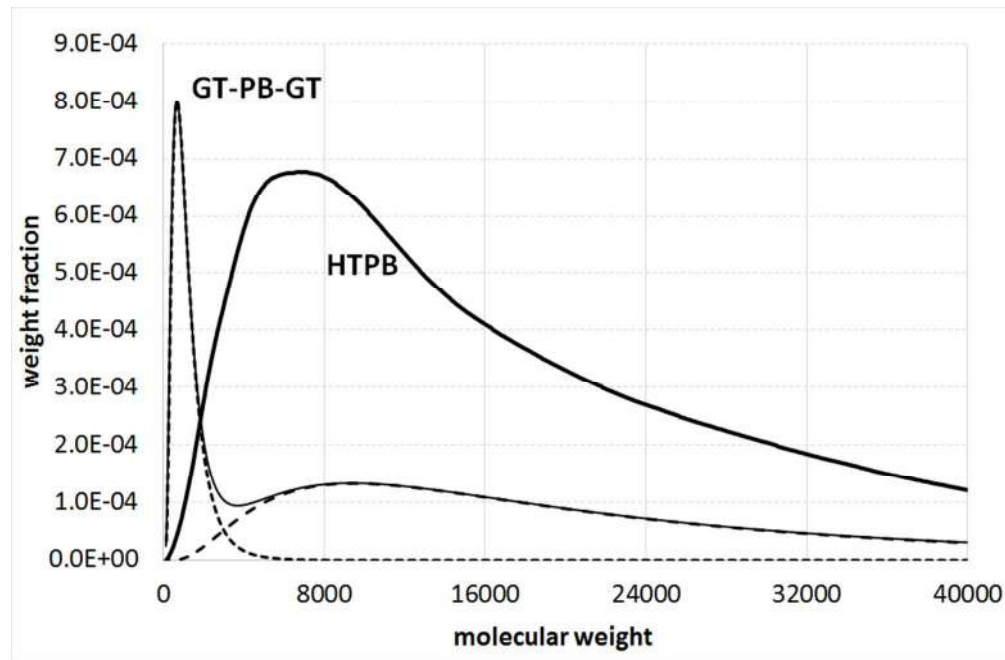


Figure 7 GPC analysis of HTPB and GT-PB-GT copolymer. The dashed lines represent the deconvolution peaks of the GT-PB-GT distribution.

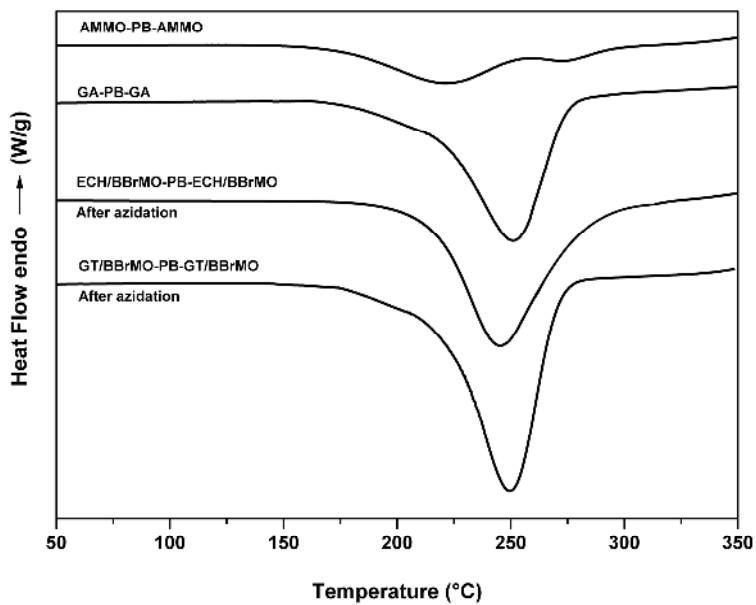


Figure 8 DSC traces of the first decomposition step for the azidated copolymers.
210x148mm (300 x 300 DPI)

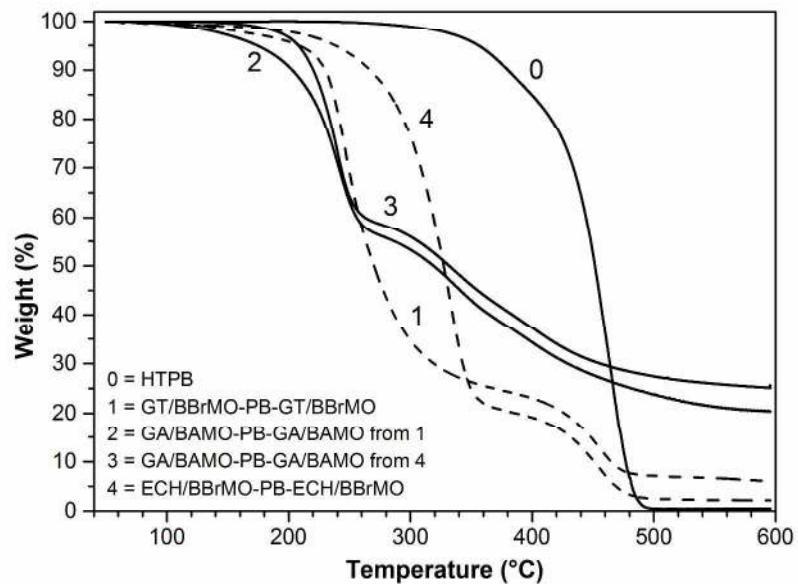


Figure 9 TGA analysis of HTPB, GA/BAMO-PB-GA/BAMO copolymers and their respective precursors.
286x199mm (300 x 300 DPI)

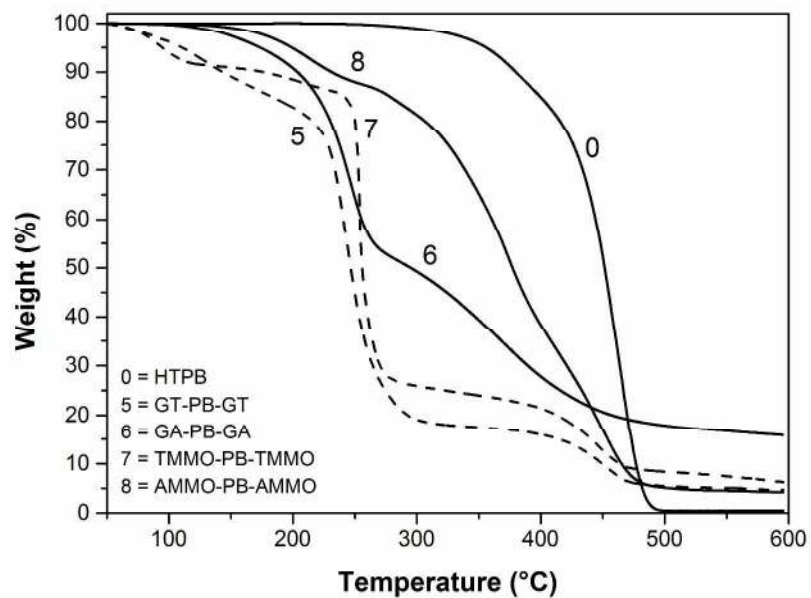


Figure 10 TGA analysis of HTPB, GA-PB-GA, AMMO-PB-AMMO copolymers and their respective precursors.
286x199mm (300 x 300 DPI)