



Cite this: *Polym. Chem.*, 2022, **13**, 69

## A versatile and straightforward process to turn plastics into antibacterial materials†

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We demonstrate that antibacterial activity can be conferred to common plastic materials using amphiphilic cationic methacrylate-based block copolymers, specifically quaternized poly(butyl methacrylate)-*b*-poly(*N,N*-dimethyl-aminoethyl methacrylate) (PBMA-*b*-PDMAEMA) with 64 mol% of DMAEMA and  $M_n$  close to 20 000 g mol<sup>-1</sup>. With 0.5–2 wt% of these copolymers simply dispersed in the given matrix by extrusion, the modified materials prove to be active against *E. coli*, *S. aureus*, *Listeria monocytogenes* and enterohemorrhagic *E. coli* without toxicity against two cell lines, HaCaT and L929 fibroblasts, while keeping the mechanical properties of the materials intact. In addition, the study of the mechanism of action shows that the antibacterial materials target the bacterial membrane, which is expected to avoid antibacterial resistance. Our protocol is a cost-effective solution to generate antibacterial materials for several applications, including packaging or medical devices, without modification of the production process.

Received 7th October 2021,  
Accepted 12th November 2021

DOI: 10.1039/d1py01344k

rsc.li/polymers

## Introduction

Infection remains the third leading cause of death in developed countries and the second worldwide. Indeed, the continuing decline of effectiveness of existing antibiotic options due to the ability of microbes to develop and disseminate mechanisms of resistance against multiple classes of traditional antimicrobials raise international healthcare concerns as they are associated with increased morbidity and mortality and also higher hospital costs.<sup>1–3</sup> Antimicrobial surfaces and antimicrobial materials are thus becoming a very important class of products that could contribute to mitigate the problem. Among them, antimicrobial plastics have received

considerable attention for numerous applications, including appliances, filters, packaging, nonwoven fabrics and textiles, and sanitary materials.<sup>4</sup> To confer antibacterial functionality to plastics, surface modification based on an array of methodologies has been exploited, such as surface casting, immersion, surface finishing, plasma treatment and deposition.<sup>5,6</sup> Although these methods afford plastic products with effective antibacterial properties, the processing methods are complex and are associated with high costs. Moreover, the antibacterial properties will be weakened if the antibacterial coating is damaged.

To obtain antibacterial functions, blending is a simpler method. Many antibacterial compounds can be employed and include either inorganic species (such as copper nanoparticles, silver nanoparticles and silver-doped glass) or organic entities (for example quaternary ammonium salts, *N*-haloamines, natural biocides such as thymol or carvacrol and classic antibiotics).<sup>4,7,8</sup> However, the sometimes low antimicrobial efficiency and leaching of antimicrobial components along with the poor durability of their antimicrobial properties limit their application.<sup>9</sup> To prepare long-lasting or non-leaching antibacterial composites, polymer-based compounds were used as additives to impart antibacterial properties to pristine materials.<sup>10–13</sup> Such an approach is usually carried out either with a small library of known antibacterial polymers such as poly(hexamethylene guanidine hydrochloride) (PHMG), chitosan or poly(*tert*-butylamino methacrylate).<sup>14,15</sup> In these cases,

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† Electronic supplementary information (ESI) available: Copolymer synthesis, mechanical properties, membrane permeation assay, surface characterization and ToF-SIMS analyses. See DOI: 10.1039/d1py01344k



because of the incompatibility of mixing the polymers with each other, a significant amount of the additive has to be inserted into the matrix, potentially altering the material properties (mechanical properties, chemical and thermal resistance, *etc.*) of the plastics.

More complex polymers that feature a backbone similar to the matrix and are decorated with antibacterial moieties or antibacterial polymers<sup>18–24</sup> have been developed to enhance the compatibilization of the additive to the plastics. Such approaches are efficient, yet specific macromolecular antibacterial architectures have to be designed for each matrix. Recently, we developed cationic amphiphilic poly(butyl methacrylate)-*b*-poly(*N,N*-dimethyl-aminoethyl methacrylate) (PBMA-*b*-PDMAEMA) diblock copolymers based on a methacrylate backbone that are potent antibacterial copolymers in solution as well as promising additives to specific polymer matrices (polystyrene (PS) and poly(methyl methacrylate) (PMMA)).<sup>16,17</sup> *N,N*-Dimethyl-aminoethyl methacrylate (DMAEMA) was chosen as the hydrophilic monomer and butyl methacrylate (BMA) as the hydrophobic monomer. A structure–activity analysis was subsequently performed, and we selected diblock copolymers with a molar composition of 64 mol% DMAEMA and obtained permanent cationic charges after quaternization of the amine groups by methyl iodide (MeI) to obtain the highest antibacterial activity (*i.e.*, low MIC values) along with the lowest toxicity. In addition, a number-average molar mass ( $M_n$  as measured by SEC/DMF) of close to 20 000 g mol<sup>-1</sup> was targeted. Such large

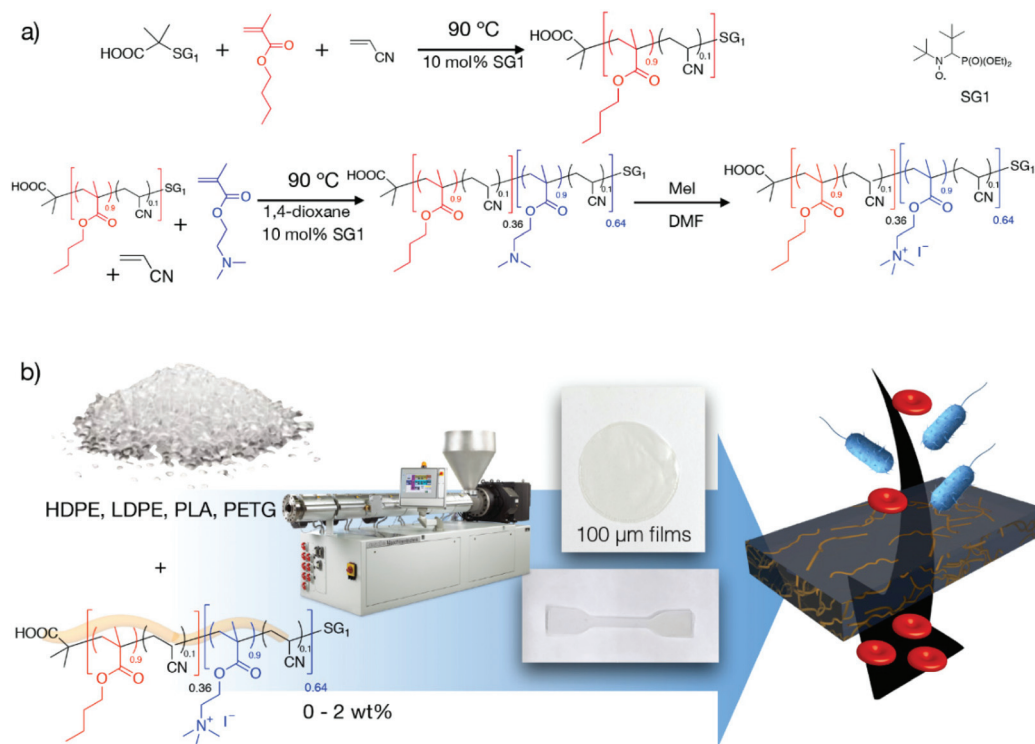
number average molecular weights,  $M_n$ , compared to those typically employed in antibacterial copolymers, were expected to ensure a good dispersion of the copolymer in common industrial organic polymer matrices, while avoiding any leaching of the active copolymer and thus a loss of activity over time.

Herein, we demonstrate that quaternized PBMA-*b*-PDMAEMA copolymers can be efficiently used as additives to confer antibacterial properties to four model industrial organic polymer matrices: poly(ethylene terephthalate glycol) (PETG), polylactide (PLA) and both high density and low density polyethylenes (HDPE and LDPE respectively). Such matrices were chosen since they have a large range of applications, especially for packaging, and are produced on a very large scale. The dispersion of the copolymer is performed by simply co-extruding a low amount of the antibacterial copolymer ( $\leq 2$  wt%) within the given matrix without impacting the innocuity of the material and its inherent mechanical properties (Fig. 1). Such an approach constitutes a simple, efficient and versatile solution to limit bacterial infections.

## Experimental section

### Materials

Commercial organic matrices, *i.e.* PLA (Natureworks), PETG (Skygreen S2008), HDPE (Dow Chemical KT 10000) and LDPE



**Fig. 1** (a) Synthesis of the antibacterial diblock copolymer B(A10%)Me as previously described<sup>16,17</sup> (PBMA-*b*-PDMAEMA,  $F_{\text{DMAEMA}} = 0.64$ ,  $M_n = 18\,400$  g mol<sup>-1</sup>, quaternization with MeI). (b) Preparation of non-leaching antibacterial materials *via* the blending of maximum 2 wt% of the B(A10%)Me antibacterial diblock copolymer with various organic matrices by extrusion.



(Total LA 0710), were used. All the reagents and solvents were used without further purification. The synthesis of the B(A10%)Me copolymer is described in the ESI† following our previous study. Human normal skin cells HaCaT were obtained from Creative Bioarray (Shirley, NY 11967, USA) and mouse fibroblastic lung cells—L929 (C3H/An)—were purchased from Sigma.

### Elaboration of antibacterial organic materials

The extrusion step was performed with a Thermo Scientific HAAKE MiniLab II mini-extruder. The pellets were dried at 90 °C under vacuum for 4 h before extrusion. Once dry, 5 g of pellets were manually loaded into the extruder. For mechanical testing, tensile specimens (type 5A) were prepared by injection moulding using a Thermo Scientific HAAKE Minijet II injector. The polymer melt at the exit of the extruder was collected in the nozzle heated to 200 °C of the injector and then injected with a pressure of 500 bar in a test specimen mold (tensile bar ISO527-2-5A) heated at 60 °C. Bioassays were performed on circular shaped films of 3 cm diameter and 100 µm thickness. Upon exiting the extruder, the recovered rushes were cooled under air and then cut into small pellets, and finally shaped using a Specac hydraulic press equipped with heating platens and a thin film kit. The pellets were pressed between two aluminium sheets heated at 170 °C and under 2 tons of pressure for 10 s.

### Antibacterial tests

The bactericidal activity of matrices was evaluated as previously described.<sup>16</sup> The bacteria used in the study were either Gram positive (*Staphylococcus aureus*, ATCC CRM-6538P) or Gram negative (*Escherichia coli*, ATCC 8739) and were routinely grown on lysogeny broth (LB) agar plates. The bactericidal activity of the different matrices was evaluated according to an adapted ISO 22196 procedure. Briefly, pieces of matrices (squares of 4 × 4 mm) were sterilely prepared from extruded films and were placed at the bottom of the wells of sterile polypropylene 96-well microplates (Greiner BioOne). Overnight bacterial suspensions (3 ml of LB broth) of *E. coli* or *S. aureus* were prepared from single colonies isolated on LB agar plates. Tubes were incubated overnight at 37 °C under agitation (200 rpm). The next day, bacterial suspensions (optical density OD > 1.0) were diluted 1/100 in 3 ml of fresh culture media and incubated at 37 °C and 200 rpm until the bacteria reached log phase growth (OD around 0.6). The bacteria were then diluted in sterile phosphate buffered saline (PBS) to reach a cell density of around 10<sup>5</sup> bacteria per ml. 10 µl of these suspensions (corresponding to ~1000 bacteria) were added onto the surface of matrices. After 60 min of incubation at 37 °C, the bacteria were collected by adding 90 µl of sterile PBS into the wells containing the matrices and repeatedly pipetted up and down. The bacteria were then serially diluted (1 in 10 dilution) in sterile PBS before plating 10 µl of each bacterial dilution onto LB agar plates. After overnight incubation at 37 °C, the plates were observed, and colonies were counted allowing the determination of the bacterial killing. Experiments were con-

ducted in independent triplicate ( $n = 3$ ). A similar protocol was used to evaluate the bactericidal activity of the matrices against food-borne bacterial pathogens, using *Listeria monocytogenes* (ATCC 15313) and enterohemorrhagic *E. coli* EHEC (ATCC K88). Finally, the bactericidal activity of the matrices was evaluated using the brine of commercial mozzarella naturally containing mixed bacterial community. Commercial mozzarella was bought from a food supermarket (Mozzarella Galbani, from Carrefour supermarket). The brine was collected sterilely and 10 µl of brine was used to evaluate the bactericidal effect of the matrices as explained above.

### Extractables and leachables

To identify and quantify the extractables, we used a dissolution/precipitation process. 6 g of the material were dissolved by refluxing in 50 mL of hot toluene under stirring. After complete dissolution, the polymer was precipitated by slowly pouring 100 mL of methanol under stirring. The precipitated polymers were removed from the additive solution by filtration on filter paper. The solution that contained the extractables was evaporated using a rotavapor at 60 °C under vacuum. To identify and quantify the leachables, 6 g of the material were immersed in 15 mL of ethanol for 72 h at 37 °C in glass vials. The material was then removed and ethanol that contained the leachables for these conditions of storage was evaporated using a rotavapor at 60 °C under vacuum.

Before injection in the HPLC system, the dry residue containing the extractables or leachables was dissolved in a mixture of 0.5 mL of tetrahydrofuran (HPLC grade Prolabo) and 0.5 mL of acetonitrile (HPLC grade Prolabo) and the solution was filtered on a 0.2 µm filter. 20 µL of the solution were injected on a Lichrospher 100 RP 18 (5 microns) using a 100% acetonitrile phase; the apparatus was a HPLC Ultimate 3000 series (Dionex) equipped with an automatic injector, a gradient pump and a UV detector with a diode array (190–400 nm). Characterization and possible identification of the compounds were performed by using their retention time and their UV spectrum and by comparing them to those of references.

## Results and discussion

### Antibacterial activity

Synthetic antimicrobial copolymers are inspired by antimicrobial peptides, *i.e.* synthetic amphiphilic cationic copolymers.<sup>25</sup> Since the pioneering work of DeGrado and colleagues,<sup>11</sup> studies covering synthetic antimicrobial copolymers experienced an exponential growth with more than 1500 publications per year. These constructs have many advantages such as simple manufacturing on a large scale. Another advantage of these synthetic copolymers compared to peptides is that a large library of monomers can be used, as well as diverse architectures (generally random or block copolymers), lengths ( $M_n$  between 1000 to 80 000 g mol<sup>-1</sup>) and compositions (various hydrophilic/hydrophobic balances).<sup>26–29</sup> This chemical variety allows identifying optimum antimicrobial systems that feature



the best compromise between high antimicrobial activity and low toxicity.<sup>25</sup> Several families of polymers have been reported in the literature, including polyacrylate and polymethacrylate derivatives, poly(oxa)norbornene, polymaleimides, polyionenes and polycarbonates.<sup>25,30</sup> Herein we prepare an antibacterial copolymer constituted of a methacrylate backbone<sup>16</sup> due to the exothermic mixing enthalpy of PMMA derivatives with a wide range of commercially relevant plastics (*e.g.*, poly(vinyl chloride) PVC, poly(styrene-*co*-acrylonitrile) PSAN, poly(vinylidene fluoride) PVDF, epoxy matrices, *etc.*), allowing for their facile incorporation into different organic matrices.<sup>31</sup> The diblock copolymer was synthesized *via* the NMP technique<sup>32</sup> using acrylonitrile ACN as the comonomer<sup>33</sup> to ensure good control of the polymerization reaction (PBMA-*b*-PDMAEMA,  $F_{\text{DMAEMA}} = 0.64$ ,  $M_n = 18\,400\text{ g mol}^{-1}$ ). ACN was used to lower the average main equilibrium constant,  $K$ , between the macroradical and macroalkoxyamine. This equilibrium is shifted towards active species in the case of methacrylate derivatives, impeding the obtainment of good control/livingness of the polymerization. The addition of only 10% of some comonomers restored the correct equilibrium, enabling living and controlled polymerization.<sup>33</sup>

DMAEMA was selected as the hydrophilic monomer and BMA as the hydrophobic monomer. The copolymer was quaternized using methyl iodide prior to use (Fig. 1a).<sup>16</sup> To be consistent with our previous work,<sup>16,17</sup> this copolymer will be denoted B(A10%)Me herein. Specifically, “B”, “A” and “Me” refer respectively to the block copolymer, acrylonitrile and methyl iodide. To prepare the antibacterial materials, pellets of commercial grades of various polymer matrices (LDPE, HDPE, PETG, PLA) and the antibacterial diblock copolymer B(A10%)Me are blended during the co-extrusion using a micro-extruder that allows working on a gram scale (Fig. 1b). The experimental conditions of the extrusions are given in the ESI (ESI – Table S1†) and are very close to the recommendations of the matrix suppliers. The strips that were obtained directly from the micro-extruders were hot pressed to obtain films of 3 cm diameter and 100 micron thickness (refer to section I.4 of the ESI†). Also, it was possible to directly employ – after micro-extrusion – a mini-injection moulding machine to prepare samples for mechanical testing (tensile bar ISO527-2-5A). We used a ratio of 0, 0.5 and 2 wt% B(A10%)Me copolymer in the organic matrices to ensure that the antibacterial agent is only an additive, expecting that the mechanical properties of the matrices will not be altered.

The antibacterial activity of the prepared solid materials was subsequently investigated using *S. aureus* and *E. coli* as model Gram-positive and Gram-negative bacteria respectively, following an ISO-22196 adapted procedure.<sup>16</sup> Indeed, the size of the sample is supposed to be  $5 \times 5\text{ cm}$  in this procedure and thus requiring a high amount of the additive and film to be prepared. To screen the various matrices and concentration of the additives more efficiently, the surface that was used in our study was decreased to  $4 \times 4\text{ mm}$ , enabling the use of 96-well plates. The tests were performed in triplicate with an exposure time of 1 h (Fig. 2). It has been reported that syn-

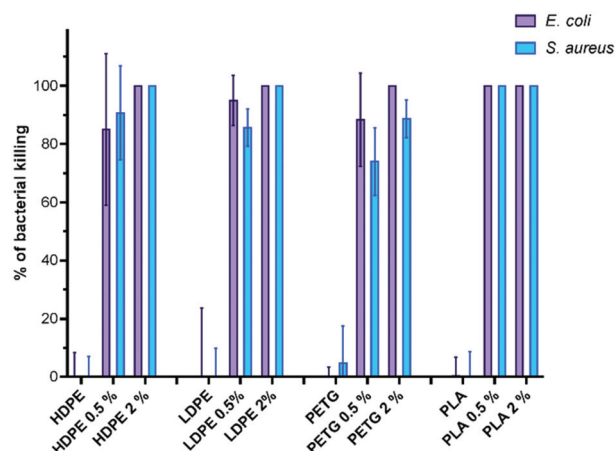


Fig. 2 Bactericidal activity of the matrices on *E. coli* and *S. aureus* after 1 h of contact. Matrices containing 0, 0.5 or 2 wt% of B(A10%)Me were exposed to *E. coli* or *S. aureus* bacteria for 1 h at 37 °C before plating and counting of live bacteria. The results correspond to means  $\pm$  S.D. ( $n = 3$ ).

thetic antibacterial copolymers are very efficient, displaying biocide action after a few minutes.<sup>34</sup> The results shown in Fig. 2 correspond to the percentage of bacteria that was killed compared to the negative control.

In all cases, the pristine matrices did not show any antibacterial activity. The incorporation of 2 wt% of B(A10%)Me proved to be sufficient to confer high antibacterial activity to commercial PETG, LDPE, HDPE and PLA matrices. The bacterial killing ranged from 88% (for PETG) to 100% (for HDPE, LDPE and PLA) on *S. aureus* and was 100% with all matrices for *E. coli*.

At 0.5 wt% of incorporation of B(A10%)Me, bactericidal activities were lower for HDPE, LDPE and PETG matrices (bacterial killing ranging from 85 to 95% for *E. coli* and from 74 to 90% for *S. aureus*). Importantly, among the various matrices, only the additivated PLA matrix remained fully active with no viable bacteria (either *S. aureus* or *E. coli*) observed after one hour of contact with only 0.5 wt% of incorporation of the B(A10%)Me copolymer. These tests confirm the antibacterial activity of these films, but more studies are required to investigate their antibiofilm behaviour as well as their effectiveness against aging.

### Mechanism of action

The mechanism of action of such antibacterial methacrylate-based copolymers has been studied in the literature and is hypothesized to be mainly a membrane destabilization mechanism leading to the lysis of the bacterial membrane.<sup>35</sup> Thus we were interested in investigating the mechanism of action of the herein presented antibacterial copolymer embedded into a polymer matrix. A propidium iodide assay was used to establish if the bactericidal activity of the matrices is related to membranolytic effects.<sup>36</sup> Indeed, such compounds are known to enter cells and form a fluorescent complex with DNA only

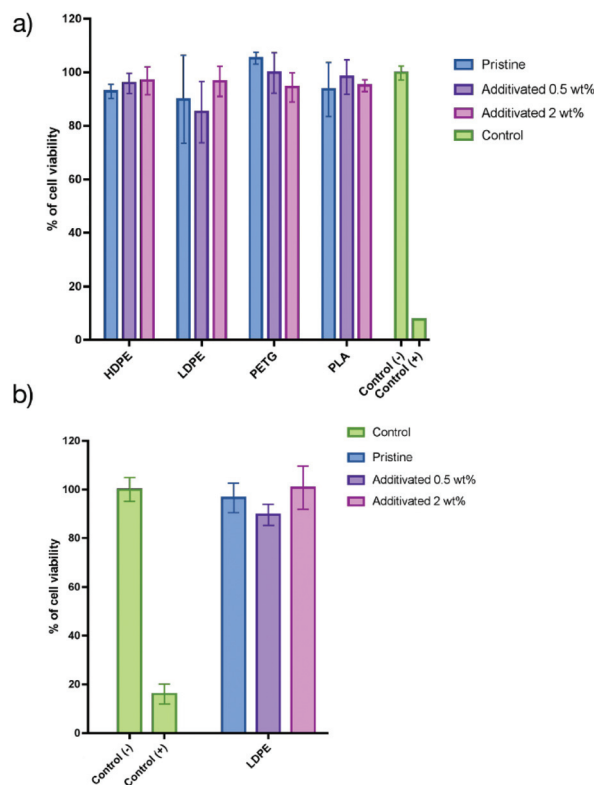


after membrane damage and are therefore good probes to monitor membrane integrity.<sup>36–38</sup> Incubation of *E. coli* as model bacteria and propidium iodide onto PLA – that displayed very good antibacterial properties – was monitored by fluorescence (the details of the test are described in the ESI†).

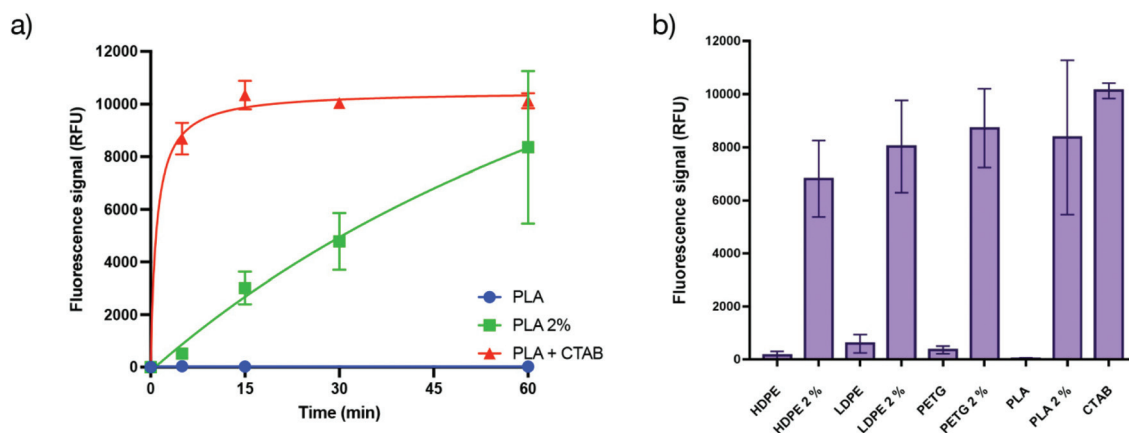
Cetyltrimethylammonium bromide (CTAB) was used as a positive control (added in a solution on a non-additivated PLA surface) since this compound is known to lyse efficiently all the bacteria present in the medium and would represent the upper limit for the fluorescence value. The experiments performed using the additivated PLA present an increase in the fluorescence value with time, reaching the upper limit within the error margin of the positive control (Fig. 3a), confirming that the antibacterial activity is based on the permeabilization of the bacterial membrane. The comparison of the increase in the fluorescence intensity between the positive control and the additivated PLA shows that the process of permeabilization of the copolymer embedded into the matrix is slower than the liquid biocide reference, yet remains rapid ( $\sim 90\%$  permeabilization in less than 1 h) compared to the activity of leachable biocides that generally requires hours to be efficient. The other matrices (HDPE, LDPE and PETG) were also tested and showed similar behaviours within the error margin (Fig. 3b).

### Cytotoxicity study

Once the antibacterial activity had been demonstrated, we proceeded to assess the *in vitro* innocuity of such materials using human normal keratinocytes as a model (HaCaT cells). Such kinds of organic matrices could be utilized for applications in which the main contact is with the skin. The cells were incubated for 48 h on the films and the cell viability was determined using the resazurin assay (the details of the tests are described in the ESI†). The results are presented in Fig. 4a. The cell viability on all pristine polymer organic matrices was close to 100% compared to the positive control.



**Fig. 4** (a) Innocuity testing of the matrices using human normal keratinocytes (HaCaT cells) as a model. HaCaT cells grown in a 96-well plate were exposed to matrices for 48 h before measurement of the cell viability using the resazurin assay. CTAB (300  $\mu\text{M}$ ) was used as the positive control of cell toxicity. The results correspond to means  $\pm$  S.D. ( $n = 3$ ). (b) Fibroblast viability referring to the MTT test results after 48 h of contact between the cells and the material; the control (+) corresponds to phenol. The results correspond to means  $\pm$  S.D. ( $n = 6$ ).



**Fig. 3** (a) Time dependent increase in the fluorescence intensity monitored *via* the propidium iodide assay using *E. coli* in the case of the PLA matrix (blue), PLA + 2 wt% B(A10%)Me (green), and PLA + cetyltrimethylammonium bromide (CTAB) at 300  $\mu\text{M}$  that was used as a membranolytic molecule (red). (b) Maximum of the fluorescence intensity observed after 60 min of contact *via* the propidium iodide assay depending on the organic matrix (poly(ethylene terephthalate glycol) (PETG), polylactide (PLA), high density polyethylene (HDPE) or low density polyethylene (LDPE) with 0 or 2 wt% of the B(A10%)Me antibacterial copolymer. The results correspond to means  $\pm$  S.D. ( $n = 3$ ).

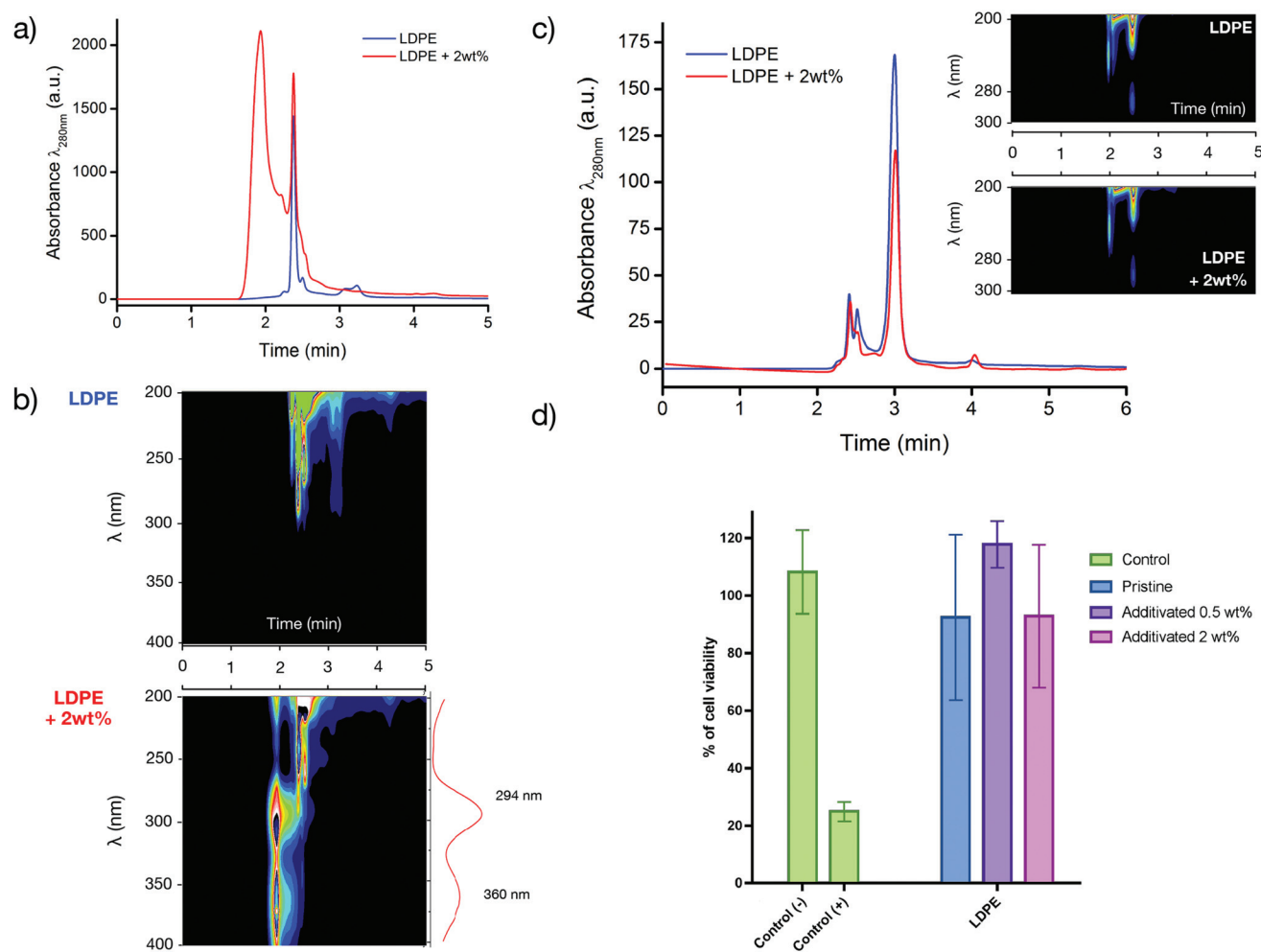


When 0.5 wt% of the additive was embedded into the matrices, the cell viability was identical to that of the pristine matrices, demonstrating that there is no impact on the additive at such low concentrations. When the amount of the additive was increased from 0.5 to 2 wt%, the HaCaT cell viability (Fig. 4a) was still close to 100%, showing that the additive does not impact the innocuity of all the materials. In order to assess if these results are dependent on the cell line, similar experiments were performed using L929 fibroblasts (Fig. 4b) using LDPE films. In that case, the cell viability is also not dependent on the concentration of the additive, showing that this approach confers antibacterial activity without altering the cell viability for all the matrices.

The main advantage of blending the antibacterial copolymer with the matrix is related to the non-leaching of the active

antibacterial additive. The co-extrusion ensures a long-lasting efficiency compared to leaching approaches (either antibiotics/biocides or inorganic salts), avoiding the increase of antibiotic resistance due to the dispersion of antibiotics in the environment. Besides the action mechanism of the antibacterial copolymer involving alteration of the bacterial membrane, the eventual leaching of such materials in the environment will not induce bacterial resistance. The non-leaching feature was assessed by using LDPE as the model polymer and ethanol as the solvent.

First, we determined the extractables, *i.e.*, all the molecules that will be able to diffuse and migrate outside the polymer matrix and that will be extracted using aggressive solvent conditions. To identify and quantify the extractables, a dissolution/precipitation process was conducted using toluene



**Fig. 5** (a) Comparison of the HPLC chromatograms of the extractables of the LDPE with and without 2 wt% of the antibacterial additive. (b) The corresponding 3D diagram (retention time and absorbance wavelength) of the HPLC chromatogram presented in (a); the peak between 2 and 3 min is attributed to a phenolic antioxidant (BHT); the peak at short time for the LDPE with the antibacterial additive (1.8–2.0 min) was attributed to an extractable of the antibacterial copolymer, featuring characteristic absorbance bands at 294 and 360 nm. (c) HPLC chromatograms of the leachables from LDPE with (red curve) and without (blue curve) the antibacterial additive; the main leachable was BHT; the extractable of the antibacterial copolymer was not detected (no compound at short retention time having characteristic UV absorption at 360 and 294 nm). (d) Fibroblast viability referring to MTT test results for the extract solutions (DMEM solution in contact with the material for 48 h). Cells were placed in contact with the extract solution for 48 h. The control (+) corresponds to phenol. The results correspond to means  $\pm$  S.D. ( $n = 7$ ).



under reflux. After complete dissolution, the matrix was precipitated by slowly pouring 100 mL of methanol under stirring. The filtrate was subsequently analysed by HPLC (Fig. 5a). A peak at 1.5–2 min was observed that was in good agreement with the B(A10%)Me copolymer (refer to Fig. S3 in the ESI†). Unlike extractables that require the dissolution of the matrix, leachables refer to the compounds that are effectively leached inside a media under conditions that are generally milder and closer to normal conditions of use. In our case, the films were immersed in 15 mL of ethanol for 72 h at 37 °C in glass vials.

The material was subsequently removed and the supernatant ethanol that contained the leachables (for these conditions of storage) was analysed (Fig. 5c).

Analysis of the leachables in ethanol by HPLC showed similar chromatograms for the LDPE with and without the antibacterial additive (Fig. 5c): the main leachable in this case was BHT, a phenolic antioxidant that was originally present in the pristine matrix. To confirm that the antibacterial copolymer and/or degradation side-products do not leach out and do not impact the biocompatibility of the matrix, the strips was also immersed in a culture medium for 48 h at 37 °C and this medium was then later used for incubating L929 fibroblasts. The results presented in Fig. 5d showed no decrease in cell viability, invariant to the amount of the antibacterial copolymer

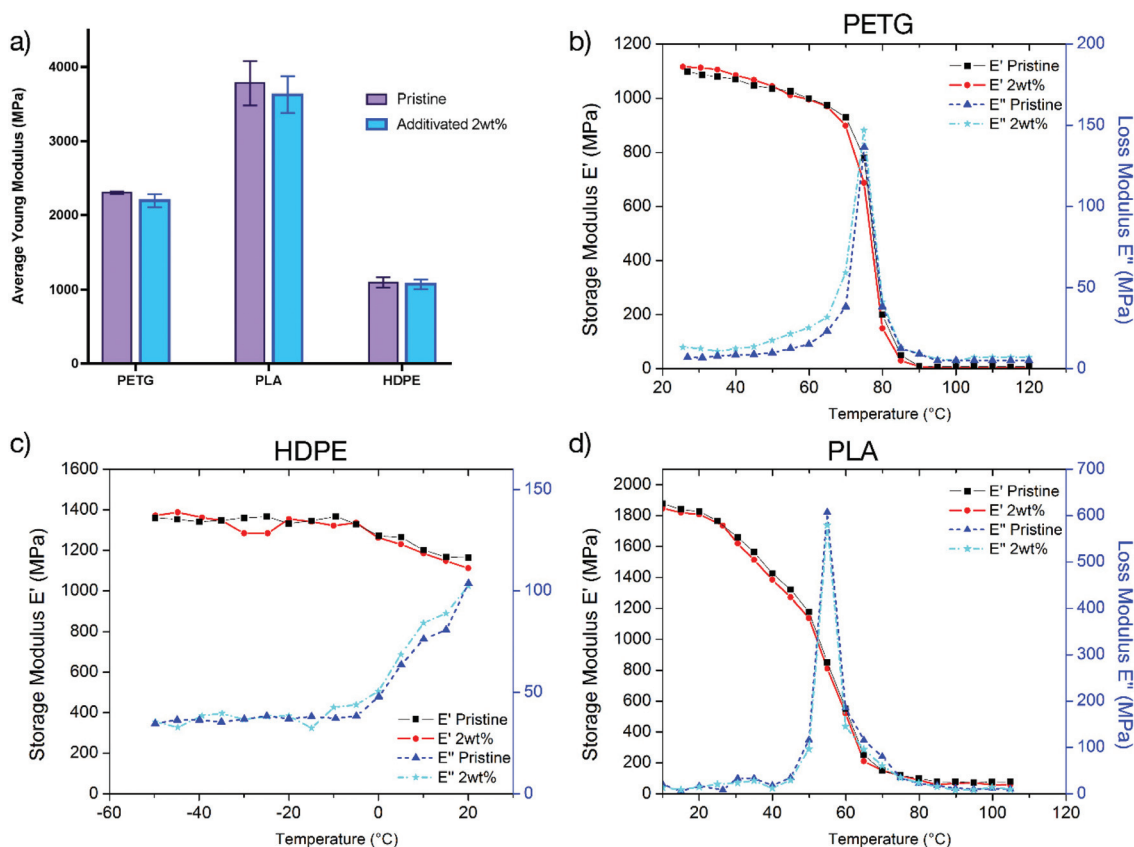
used as an additive (Fig. 5d). The absence of cytotoxicity of the additivated matrix is of prime importance to envision a biomedical application of the prepared antibacterial materials.

### Mechanical properties

With applications ranging from biomaterials to packaging, the mechanical properties of the materials are of paramount importance. The modification of the materials to confer antibacterial activity should ideally not affect their mechanical properties. In our approach, the use of low amounts (0.5–2 wt%) of the antibacterial copolymer as an additive is not expected to alter the bulk properties of the materials. For confirmation, tensile tests were performed on PETG, PLA and HDPE samples to determine Young's modulus. The samples of both pristine and additivated matrices (2 wt%) were prepared by injection moulding.

We assessed the materials with the higher amount of the antibacterial copolymer to determine the maximum modification of the mechanical properties. The comparison of these values (Fig. 6a, details in the ESI†) demonstrates that the incorporation of the copolymer in the matrix does not change the mechanical properties of the pristine material.

To confirm the conservation of the mechanical properties, dynamic mechanical analysis (DMA) tests were performed on



**Fig. 6** (a) Young's modulus of the pristine and additivated matrices containing 2 wt% of the B(A10%)Me copolymer as obtained by the tensile test. (b–d) Comparison between the dynamic properties storage ( $E'$ ) and loss moduli ( $E''$ ) curves as a function of temperature for PETG (b), HDPE (c) and PLA (d) without the copolymer (pristine) or with 2 wt% of the B(A10%)Me copolymer.



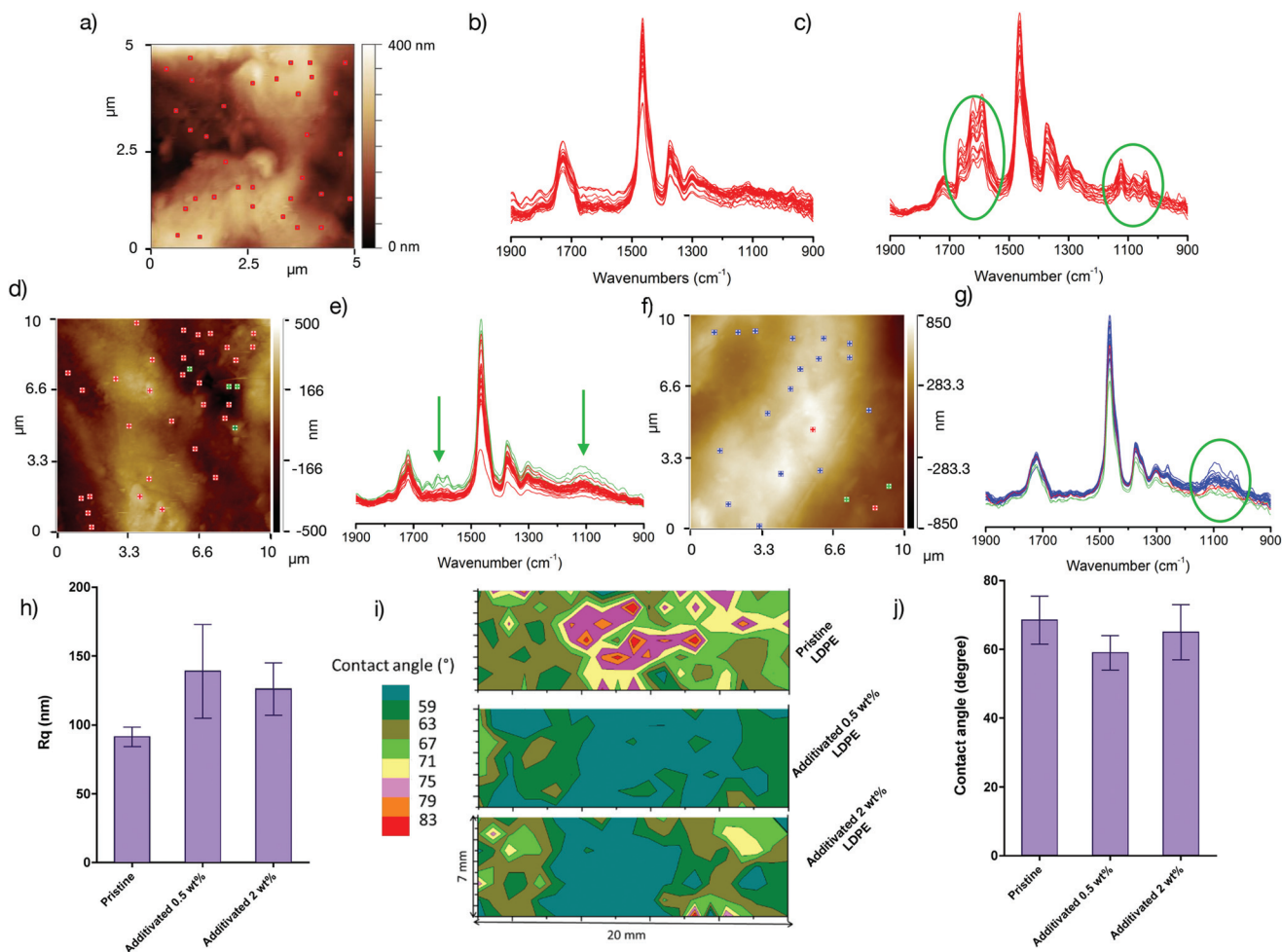
PETG, PLA and HDPE samples to determine the viscoelastic properties over a range of temperatures.

The comparison of storage ( $E'$ ) and loss moduli ( $E''$ ) curves as a function of temperature shown in Fig. 6b–d demonstrates that the materials have the same viscoelastic properties over a large range of temperature. In conclusion, the addition of the copolymer into the polymer matrix does not change the mechanical properties of the materials over a large range of temperatures and strain levels.

### Surface characterization

According to the antibacterial studies, the use of a macromolecular antibacterial additive that is not leached out in the medium confers good activity by membrane permeabilization even with a small amount of the copolymer. The localization of the copolymer in the materials is thus an intriguing question. Due to the blending in the micro-extruder, one would

expect a homogeneous dispersion of the amphiphilic copolymer into the polymer matrix, but the partial or complete migration of the copolymer to the surface cannot be excluded. To answer this question, we performed time-of-flight secondary ion mass spectrometry (ToF-SIMS) analyses on PETG, LDPE and PLA materials containing 2 wt% of copolymers (see the ESI† – section VI for details). Using ToF-SIMS, the specific mass response of the copolymer was detected in the background of the matrix. Even if all the copolymer migrated towards the surface of the materials, the low amount of the additive was still difficult to detect. As shown in Fig. S2,† there is no increase in the intensity in the ToF-SIMS spectrum of the organic matrix with the copolymer at  $m/z$  values corresponding to peaks in the copolymer spectrum, for any of the three materials. These results showed that it was not possible to detect the antibacterial additive on the surface samples even when using the sensitive ToF-SIMS analysis. These indirect



**Fig. 7** (a) AFM height image ( $5 \times 5 \mu\text{m}$ ) of the LDPE film surface without the antibacterial copolymer; (b) FTIR spectra corresponding to the red points in figure a; (c) FTIR spectra corresponding to another area of the film surface without the antibacterial copolymer; (d) AFM height image ( $10 \times 10 \mu\text{m}$ ) of the LDPE film surface with 0.5% of the antibacterial copolymer; (e) FTIR spectra corresponding to the red and green points in figure d; (f) AFM height image ( $10 \times 10 \mu\text{m}$ ) of the LDPE film surface with 2 wt% of the antibacterial copolymer; (g) FTIR spectra corresponding to the blue, red and green points in figure f; (h) root-mean-square roughness  $R_q$  for the different LDPE films; (i) water contact angle mapping of the LDPE film surface for LDPE without the antibacterial copolymer and containing 0.5 and 2 wt% of the copolymer. A  $7 \times 20 \text{ mm}$  surface was investigated with 300 pL water droplets; (j) water mean contact angle for each surface (for the 168 droplets deposited onto the surface).



results prove that whatever the matrix is, the copolymer does not migrate to the surface and likely stays well dispersed in the material.

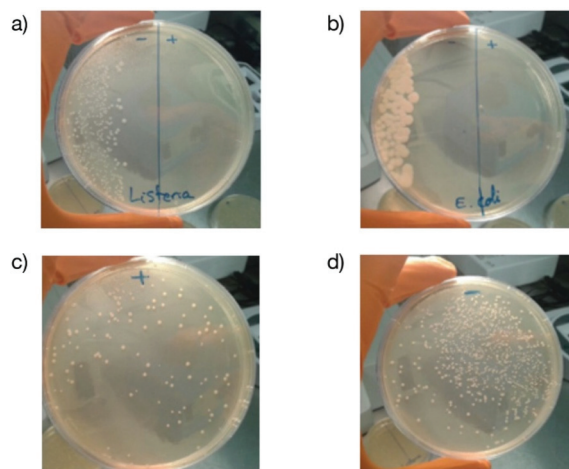
To obtain further insights, we focused on the matrix that differs the most from the copolymer in terms of the structure (LDPE with an apolar totally carbon-based backbone) to maximize the chance to detect the copolymer. The surface of the LDPE films was characterized by atomic force microscopy (AFM) in the tapping mode and nanoscale infrared spectroscopy (nano-IR), allowing the acquisition of local Fourier transform infrared spectroscopy (FTIR) spectra of the sample with a high lateral resolution (all the techniques are described in detail in the ESI†). The pristine LDPE film has a heterogeneous and rough (Fig. 7a) surface as shown by the local FTIR spectra (Fig. 7b) and the hydrophobicity mapping (Fig. 7i). The non-homogeneous surface polarity and low contact angles observed for such an apolar polymer (Fig. 7j) may be the result of surface oxidation (C=O band at  $1724\text{ cm}^{-1}$  in Fig. 7b) and/or of the presence of lubricant-rich areas (see amide bands between  $1700$  and  $1590$  in Fig. 7c). The addition of the antibacterial copolymer results in a slight increase in the roughness (Fig. 7h). The hydrophobicity is nevertheless not significantly modified and rich antibacterial copolymer areas on the film surface cannot be certainly attributed: the oxidation band and amide lubricant bands (green arrow in Fig. 7e) are still observed, yet no other bands appear significantly, even if, for the film containing 2 wt% of the copolymer, small bands at around  $1100\text{ cm}^{-1}$  (ether stretching band in Fig. 7g) are locally detected and may be the result of the presence of the copolymer but cannot be ascribed with certainty to the additive.

The combination of all these techniques showed that the surface is not sufficiently modified to detect the copolymer. This is a striking difference with the materials whose antibacterial properties are conferred by various surface modification techniques.<sup>39</sup>

### Antibacterial packaging application

Finally, to demonstrate that our approach could be realistically envisioned as a solution to prevent bacterial infection in a real-world scenario, we prepared antibacterial LDPE films containing 2 wt% of the copolymer as an additive and test them as packaging for mozzarella as already reported.<sup>40</sup>

The idea was to show whether such a plastic that is commonly used to prepare films used in food packaging applications could be useful to (i) avoid the development of pathogenic bacteria such as *Listeria monocytogenes* and enterohemorrhagic *E. coli* that are responsible every year for contamination widely reported in the media<sup>41,42</sup> and (ii) preserve the bacteria within the cheese that are necessary to impart gustative properties.<sup>43</sup> Fig. 8a and b clearly prove that both *Listeria monocytogenes* and enterohemorrhagic *E. coli* were completely killed after contact with the additive containing film. In the case of bacteria already present in cheese, we observe that a large fraction of them survived (Fig. 8c and d) although with a lower amount than the pristine film. This result could be inter-



**Fig. 8** (a and b) Development of pathogenic bacteria (*Listeria monocytogenes* (a) and enterohemorrhagic *E. coli* (b)) in contact with LDPE with 2 wt% of the B(A10%)Me copolymer (noted +) compared to the pristine LDPE as a reference (noted -). (c and d) Survival of cheese bacteria from mozzarella brine after contact with LDPE with 2 wt% of the B(A10%)Me copolymer (noted +) (c) compared to the pristine LDPE as a reference (noted -) (d).

esting for increasing the shelf-life of food and allows us to envision the possibility of using B(A10%)Me copolymer containing plastics as food packaging materials.

## Conclusions

We demonstrate that simply adding an amphiphilic synthetic cationic diblock copolymer based on methacrylic monomers into common organic matrices (such as HDPE, LDPE, PLA and PETG) *via* an extrusion process confers antibacterial properties to the pristine materials after 1 h of exposure. The effect of the concentration for such additives has been studied and we demonstrate that even 0.5 wt% is sufficient to impart good antibacterial efficiency (>80% of bacterial killing) for both *E. coli* and *S. aureus*. The increase to 2 wt% led to complete inhibition of both bacteria on HDPE, LDPE and PLA and more than 90% of bacterial killing on PETG. The mechanism of action was also investigated and the permeabilization of the bacterial membrane was monitored *via* the propidium iodide assay, suggesting that the killing of bacteria occurred by contact and membrane destabilization. This action mechanism is expected to limit the development of bacterial resistance. In addition, the study on HaCaT and L929 fibroblast cell lines reveals no toxicity of the materials. Further analysis of the materials demonstrated that the mechanical properties over a wide range of temperatures are not altered. The stable mechanical properties are of prime importance for the utilization of these materials in their specific applications.

For a better understanding of the mode of action of such materials, complete characterization of the surface has been carried out. ToF-SIMS, nano-IR spectroscopy and AFM did not



reveal any specific modification of the surface, suggesting that there is a homogeneous incorporation of the additive in the materials without complete migration towards the surface. This is a critical advantage compared to antibacterial materials whose properties are given by a surface modification (antibacterial coatings). Lastly, to demonstrate its application in food packaging, a LDPE film containing 2 wt% of the additive was shown to inhibit the growth of *Listeria monocytogenes* and enterohemorrhagic *E. coli*. At the same time, the additive containing film slows down the development of cheese bacteria in mozzarella brine, which could increase the shelf-life of the food contained in the LDPE film.

Our methodology therefore constitutes a straightforward and cost-effective solution to generate intrinsically and long-lasting antibacterial materials for a plethora of applications, including packaging, disposable materials and medical devices.

## Author contributions

Y. G. D. G. and C. L. conceived the original idea of the project. S. H. conducted the copolymer synthesis and material preparations. G. M. and A. M.-P. performed the measurements of material properties. S. F., J. S., A. D.-B. and N. Y. conducted the material characterization. S. H. and M. M. performed the antibacterial and innocuity tests. L. M. and C. B.-K. conducted the surface characterization of the materials by ToF-SIMS. C. B.-K., Y. G. and C. L. wrote the original manuscript. All authors contributed to the scientific planning and discussions.

## Conflicts of interest

D. G., M. M., Y. G. and C. L. are named inventors on patent applications (French Patent Application FR1756390 (2017) and PCT WO2019/008176 (2018)) filed by the CNRS, Ecole Centrale Marseille, and Aix-Marseille University on the antibacterial materials described in the current study.

## Acknowledgements

The authors thank the CNRS, SATT Sud-Est and Aix-Marseille University for support. C. B.-K. acknowledges the Australian Research Council (ARC) for funding in the context of a Laureate Fellowship and the Queensland University of Technology (QUT) for continued support including *via* its Centre for Materials Science.

## Notes and references

- 1 M. Baym, L. K. Stone and R. Kishony, *Science*, 2016, **351**, 5.
- 2 M. F. Chellat, L. Raguz and R. Riedl, *Angew. Chem., Int. Ed.*, 2016, **55**, 6600–6626.
- 3 M. I. Hutchings, A. W. Truman and B. Wilkinson, *Curr. Opin. Microbiol.*, 2019, **51**, 72–80.
- 4 A. Munoz-Bonilla and M. Fernandez-Garcia, *Prog. Polym. Sci.*, 2012, **37**, 281–339.
- 5 M. M. Konai, B. Bhattacharjee, S. Ghosh and J. Haldar, *Biomacromolecules*, 2018, **19**, 1888–1917.
- 6 K. G. Neoh, M. Li, E. T. Kang, E. Chiong and P. A. Tambyah, *J. Mater. Chem. B*, 2017, **5**, 2045–2067.
- 7 L. Karam, R. Roustom, M. G. Abiad, T. Ei-Obeid and I. N. Savvaidis, *Int. J. Food Microbiol.*, 2019, **291**, 42–47.
- 8 A. Orsuwan, S. Kwon, N. Bumbudsanpharoke and S. Ko, *Food Control*, 2019, **100**, 176–182.
- 9 J. Bruenke, I. Roschke, S. Agarwal, T. Riemann and A. Greiner, *Macromol. Biosci.*, 2016, **16**, 647–654.
- 10 A. Kyzioł, W. Khan, V. Sebastian and K. Kyzioł, *Chem. Eng. J.*, 2020, **385**, 123888.
- 11 G. N. Tew, D. H. Liu, B. Chen, R. J. Doerksen, J. Kaplan, P. J. Carroll, M. L. Klein and W. F. DeGrado, *Proc. Natl. Acad. Sci. U. S. A.*, 2002, **99**, 5110–5114.
- 12 G. N. Tew, R. W. Scott, M. L. Klein and W. F. DeGrado, *Acc. Chem. Res.*, 2010, **43**, 30–39.
- 13 D. Campoccia, L. Montanaro and C. R. Arciola, *Biomaterials*, 2013, **34**, 8533–8554.
- 14 S. Ye, D. Wei, X. Xu, Y. Guan and A. Zheng, *Polym. Adv. Technol.*, 2020, **31**, 1847–1856.
- 15 H. Zuo, D. Wu and R. Fu, *J. Appl. Polym. Sci.*, 2012, **125**, 3537–3544.
- 16 B. T. Benkhaled, S. Hadiouch, H. Olleik, J. Perrier, C. Ysacco, Y. Guillaneuf, D. Gimes, M. Maresca and C. Lefay, *Polym. Chem.*, 2018, **9**, 3127–3141.
- 17 D. Gimes, Y. Guillaneuf, C. Guillaneuf, M. Maresca and C. Ysacco, *Solid organic antibacterial material*, 2018, WO2019008176A1.
- 18 G. Liu, Q. Zhang, Y. Li, X. Wang, H. Wu, Y. Wei, Y. Zeng and L. Tao, *iScience*, 2020, **23**, 100754.
- 19 W. Cao, D. Wei, A. Zheng and Y. Guan, *Eur. Polym. J.*, 2019, **118**, 231–238.
- 20 M. R. Badrossamay and G. Sun, *React. Funct. Polym.*, 2008, **68**, 1636–1645.
- 21 F. Chen, X. Ding, Y. Jiang, Y. Guan, D. Wei, A. Zheng and X. Xu, *ACS Omega*, 2020, **5**, 10481–10488.
- 22 D. Wei, Y. Ding, T. Wang, J. Yang, Y. Guan and A. Zheng, *J. Appl. Polym. Sci.*, 2017, **134**, 44190.
- 23 R. Cuervo-Rodriguez, F. Lopez-Fabal, J. L. Gomez-Garces, A. Munoz-Bonilla and M. Fernandez-Garcia, *Macromol. Biosci.*, 2017, **17**, 1700258.
- 24 J.-M. Thomassin, S. Lenoir, J. Riga, R. Jérôme and C. Detrembleur, *Biomacromolecules*, 2007, **8**, 1171–1177.
- 25 C. Ergene, K. Yasuhara and E. F. Palermo, *Polym. Chem.*, 2018, **9**, 2407–2427.
- 26 A. C. Engler, J. P. K. Tan, Z. Y. Ong, D. J. Coady, V. W. L. Ng, Y. Y. Yang and J. L. Hedrick, *Biomacromolecules*, 2013, **14**, 4331–4339.
- 27 P. R. Judzewitsch, T. K. Nguyen, S. Shanmugam, E. H. H. Wong and C. Boyer, *Angew. Chem., Int. Ed.*, 2018, **57**, 4559–4564.



- 28 A. Kuroki, P. Sangwan, Y. Qu, R. Peltier, C. Sanchez-Cano, J. Moat, C. G. Dowson, E. G. L. Williams, K. E. S. Locock, M. Hartlieb and S. Perrier, *ACS Appl. Mater. Interfaces*, 2017, **9**, 40117–40126.
- 29 A. Kuroki, A. K. Tchoupa, M. Hartlieb, R. Peltier, K. E. S. Locock, M. Unnikrishnan and S. Perrier, *Biomaterials*, 2019, **217**, 119249.
- 30 J. Tan, J. Tay, J. Hedrick and Y. Y. Yang, *Biomaterials*, 2020, **252**, 120078.
- 31 A. V. Ruzette and L. Leibler, *Nat. Mater.*, 2005, **4**, 19–31.
- 32 J. Nicolas, Y. Guillaneuf, C. Lefay, D. Bertin, D. Gigmes and B. Charleux, *Prog. Polym. Sci.*, 2013, **38**, 63–235.
- 33 E. Guegain, Y. Guillaneuf and J. Nicolas, *Macromol. Rapid Commun.*, 2015, **36**, 1227–1247.
- 34 I. Sovadinova, E. F. Palermo, M. Urban, P. Mpiğa, G. A. Caputo and K. Kuroda, *Polymers*, 2011, **3**, 1512–1532.
- 35 H. Takahashi, G. A. Caputo, S. Vemparala and K. Kuroda, *Bioconjugate Chem.*, 2017, **28**, 1340–1350.
- 36 W. A. Dengler, J. Schulte, D. P. Berger, R. Mertelsmann and H. H. Fiebig, *Anti-Cancer Drugs*, 1995, **6**, 522–532.
- 37 L. B. Oyama, S. E. Girdwood, A. R. Cookson, N. Fernandez-Fuentes, F. Prive, H. E. Vallin, T. J. Wilkinson, P. N. Golyshin, O. V. Golyshina, R. Mikut, K. Hilpert, J. Richards, M. Wootton, J. E. Edwards, M. Maresca, J. Perrier, F. T. Lundy, Y. Luo, M. Zhou, M. Hess, H. C. Mantovani, C. J. Creevey and S. A. Huws, *npj Biofilms Microbiomes*, 2017, **3**, 9.
- 38 E. Di Pasquale, C. Salmi-Smail, J. M. Brunel, P. Sanchez, J. Fantini and M. Maresca, *Chem. Phys. Lipids*, 2010, **163**, 131–140.
- 39 J. Hasan, R. J. Crawford and E. P. Lvanova, *Trends Biotechnol.*, 2013, **31**, 31–40.
- 40 B. Agrillo, M. Balestrieri, M. Gogliettino, G. Palmieri, R. Moretta, Y. T. R. Proroga, I. Rea, A. Cornacchia, F. Capuano, G. Smaldone and L. De Stefano, *Int. J. Mol. Sci.*, 2019, **20**(3), 601.
- 41 A. Gerard, S. El-Hajjaji, E. Niyonzima, G. Daube and M. Sindic, *Int. J. Dairy Technol.*, 2018, **71**, 825–843.
- 42 M. Gouali and F. X. Weill, *Presse Med.*, 2013, **42**, 68–75.
- 43 C. Pelaez and T. Requena, *Int. Dairy J.*, 2005, **15**, 831–844.

