High-Throughput Preparation of Antibacterial Polymers from Natural Product Derivatives via the Hantzsch Reaction

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\textbf{Summary:} The Hantzsch and free-radical polymerization reactions were combined in a one-pot high-throughput (HTP) system to simultaneously prepare 30 unique polymers in parallel. Six aldehydes derived from natural products were used as the starting materials to rapidly prepare the library of 30 poly(1,4-dihydropyridines). From this library, HTP evaluation methods lead to the identification of a new antibacterial polymer. Mechanistic studies revealed the dihydropyridine group in the polymer side-chain structure plays an important role in resisting bacterial attachment to the polymer surface, thus leading to the antibacterial function of this polymer. This research demonstrates the value of multicomponent reactions (MCRs) in interdisciplinary fields by discovering new functional polymers for possible practical applications. It also provides insights to further developing functional polymers using MCRs and HTP methods with important implications in organic chemistry, polymer chemistry, and materials science.

\textbf{INTRODUCTION}

High-throughput (HTP) synthesis methods enable the rapid production of hundreds to even millions of samples for screening and identifying products with the desired properties/functions.(Briceno et al., 1995; Danielson et al., 1997; Hanak, 1970; Lehn and Eliseev, 2001; Rademann and Jung, 2000; Xiang et al., 1995) This effectively accelerates the pace of research, significantly increasing productivity and
resulting in remarkable benefits. In polymer chemistry, HTP strategies have been successfully applied to the discovery of new functional polymers. Many polymers have been synthesized in parallel and characterized via HTP methods for detailed studies of the subtle relationships between polymer structures and properties. As a result, many new functional polymers have been explored and used in interdisciplinary fields as sensors, gene/drug carriers, antibacterial surfaces, tissue engineering scaffolds, (stem)cell factories, etc. (Akinc et al., 2003; Algahtani et al., 2014; Anderson et al., 2004; Anderson et al., 2005; Anderson et al., 2006; Bosman et al., 2001; Celiz et al., 2014; Chapman et al., 2016; Goldberg et al., 2008; Gupta et al., 2010; Hook et al., 2012; Khan et al., 2010; Lynn et al., 2001; Mao et al., 2018; Mei et al., 2010; Mei et al., 2004a; Mei et al., 2004b; Neumann et al., 2017; Ting et al., 2015; Ting et al., 2016) This has opened new research directions in polymer science and expanded the application scope of polymers beyond traditional plastics and rubbers. Thus, the development of new HTP methods to synthesize, screen, and isolate new polymers with application values is important from both academic and industrial points of view.

Multicomponent reactions (MCRs) are defined as reactions where three or more reactants combine to effectively generate a single product. Recently, MCRs have aroused widespread attention in polymer chemistry because Meier and coworkers reported the synthesis of polycondensates via the tricomponent Passerini reaction. (Kreye et al., 2011) Since that report, more and more MCRs have been exploited for the preparation of elegant polymers containing multicomponent mainchains and side chains. Such MCRs include the Passerini, Ugi, Mannich, Biginelli, Hantzsch, and Kabachnik–Fields reactions, as well as metal-catalyzed and thiolactone-based MCRs. (Blasco et al., 2017; Deng et al., 2012; Espeel et al., 2011; Kakuchi and Theato, 2014; Kreye et al., 2011; Lee et al., 2013; Liu et al., 2014; Llevot et al., 2017; Sehlinger et al., 2013; Siamaki et al., 2011; Theato, 2015; Wu et al., 2017a; Zhang et al., 2016; Zhang et al., 2014; Zhao et al., 2016)

MCRs have been applied to the construction of polymer libraries in a HTP manner. (Mao et al., 2018; Wu et al., 2017b; Xue et al., 2016) MCRs quickly increase the number and diversity of polymer samples,
due to their multicomponent nature, effectively improving the synthesis efficiency for constructing polymer libraries. Selected robust MCRs have been carried out during controlled radical polymerization (CRP) to successfully prepare well-defined polymers with multicomponent pendant/chain end groups in a one-pot manner. (Yang et al., 2015; Zhang et al., 2015; Zhang et al., 2014; Zhu et al., 2013) This greatly simplifies the preparation of new polymers by avoiding the tedious separation and purification steps during a new monomer/initiator synthesis. HTP methods, MCRs, and a one-pot strategy straightforwardly improve the efficiency of polymer preparation; thus, the combination of HTP methods, MCRs, and a one-pot strategy might lead to a new methodology for facilely constructing polymer libraries. Herein, we report a one-pot HTP method that can quickly establish a library of poly(1,4-dihydropyridine)s (poly(1,4-DHP)s) via the Hantzsch reaction (a typical MCR) and free-radical polymerization (FRP) (Scheme 1).

Scheme 1. One-Pot HTP Method for Preparing a Library of Poly(1,4-DHP)s

The Hantzsch reaction, reported by Arthur R. Hantzsch in 1881,(Hantzsch, 1881) consists of four common components (aldehyde, ß-ketoester, 1,3-diketone, and NH₄OAc) and effectively produces 1,4-dihydropyridines (1,4-DHPs), which are candidate drugs for treating cardiovascular diseases.(Loev et al., 1974; Stout and Meyers, 1982) The Hantzsch reaction has been broadly studied in organic chemistry and
pharmaceutical chemistry; only recently has this venerable reaction been explored in polymer chemistry. Well-defined polymers with 1,4-DHP side groups have been efficiently prepared by the combination of the Hantzsch reaction and reversible addition–fragmentation chain transfer (RAFT) polymerization in a one-pot fashion. (Zhang et al., 2015) Here, we chose FRP instead of CRP to further simplify the polymerization process. Aldehydes derived from natural products were reacted with a commercially available monomer, 2-(acetoacetoxy) ethyl methacrylate (AEMA), via the Hantzsch reaction during the FRP process. The polymers were synthesized in parallel using one-pot reactions containing different combinations of six aldehydes ($A(X)$) and five 1,3-cyclohexanedione compounds ($B(Y)$) to simultaneously create $6 \times 5 = 30$ new polymers ($P(X)(Y)$).

The resulting polymers were screened via HTP measurement techniques to ultimately identify the polymer with the best antibacterial ability. This polymer was then blended with commodity polymers (polyethylene (PE), poly(propene) (PP), poly(ethylene terephthalate) (PET), poly(methyl methacrylate) (PMMA), and polyamide (PA66)) to dramatically improve the antibacterial ability of these general polymers without impairing the mechanical strength of the overall polymer system. This demonstrates the utility of new synthesis strategies in polymer chemistry to explore new functional polymers for practical applications.

RESULTS AND DISCUSSION

One-Pot HTP Preparation of Polymers via the Hantzsch Reaction and FRP.

In a representative procedure, $A(X)$, $B(Y)$, AEMA, and NH$_4$OAc were added to centrifuge tubes; the molar ratio of each component was $A(X)$: $B(Y)$: AEMA: NH$_4$OAc = 1: 1: 1: 1.5. Solvents (acetonitrile and 1,4-dioxane (1/1, v/v)) were then added followed by glycine (10 mol% with respect to aldehydes) and 2,2’-azobisisoheptonitrile (ABVN, 2 mol% to AEMA) as the catalyst for the Hantzsch reaction and the FRP initiator, respectively. The 30 charged tubes were purged with bubbling nitrogen for 20 min and then immersed in a 75 °C oil bath for 12 h. After quenching the polymerization in an ice–water bath, aliquots...
(20–50 µL) were taken for 1H NMR and gel permeation chromatography (GPC) analyses. For every tube, AEMA was polymerized in high conversions (95%–99%), and the β-ketoester groups of AEMA nearly completely disappeared (92–99%) (Figure S1 (P(X)(1) as a typical example), Table S1). All poly(1,4-DHP)s had high molecular weights (Mn(GPC): 16000–28000 g mol⁻¹, Figure S2 (P(X)(1) as a typical example), Table S1).

The P(X)(Y) were purified through simple precipitation in water followed by re-precipitation from tetrahydrofuran by diethyl ether. As a typical example, the 1H NMR spectra of P(X)(1) are presented in Figure 1.

**Figure 1.** a) Reaction conditions: acetonitrile/dioxane (1/1, v/v), glycine (10% to aldehydes), ABVN (2% to AEMA), 75 °C for 12 h. A(X): B(1): AEMA: NH₄OAc = 1:1:1:1.5. b) 1H NMR spectra (CDCl₃, 400M) of P(X)(1).
The specific peaks of the 1,4-DHP moieties (R1-CH) can be found at 3.95–4.55 ppm, whereas the peaks of the AEMA vinyl group (6.07, 5.55 ppm) have completely disappeared. The unreacted β-ketoesters (~3.48 ppm) in P(X)(1) were calculated as ~1–6% (Table S1). Other P(X)(Y) polymers showed similar results (Figure S3-S6, Table S1). Thus, the selected aldehydes and 1,3-cyclohexanedione derivatives were suitable components for the Hantzsch reaction and highly compatible with FRP.

To compare the effectiveness of the one-pot HTP method with the traditional monomer polymerization approach, P(X)(1) with 100% 1,4-DHP side groups were prepared in a two-step process (Figure S7-12). As a typical example, cinnamaldehyde (A(1)) was reacted with AEMA and dimedone (B(1)) via the Hantzsch reaction to obtain the target monomer M(1)(1) in 73.6% yield after column chromatography purification. M(1)(1) was polymerized through FRP to obtain the desired “pure” polymer P(1)(1). 1H NMR spectra indicated this polymer was nearly identical to that obtained by the one-pot HTP method (Figure S7). Similar results were observed when other aldehydes were used to prepare pure monomers and polymers (Figure S8-12). This suggests the feasibility of this one-pot HTP strategy to quickly build a library of target polymers in a time-saving and labor-saving manner.

**HTP Analyses of the Antibacterial Capabilities of P(X)(Y)**

1,4-DHP derivatives have been reported as potential antimicrobial agents, antioxidants, and anti-inflammatory agents. (Vijesh et al., 2011) The selected aldehydes in this study have been widely employed in the cosmetic, food, and pharmaceutical industries as flavor agents, antibacterial agents, and antioxidants. (Aziz and Karboune, 2018; Gyawali and Ibrahim, 2014) Thus, this library of poly(1,4-DHP)s might contain new functional polymers with bioactive properties. To test this hypothesis, the antibacterial capability of the resulting polymers was evaluated via HTP measurement techniques.

**Preliminary Screening** The resulting polymers were mixed with poly(ethylene) (PE) to prepare mixed hot-pressed samples. The weight ratio of P(X)(Y) in P(X)(Y)-PE was 33%. *Escherichia coli* (*E. coli*) and
*Staphylococcus aureus* (*S. aureus*) were used as representative Gram-negative and Gram-positive bacteria, respectively. Both bacteria were transfected with red fluorescent protein (RFP) for easy observation.

**Figure 2.** HTP analysis of the antibacterial capabilities of P(X)(Y). a) Schematic of the polymer arrays and the test procedure. b) LSCM images and b’) intensity map of *E. coli* on sample disks after 72-h incubation. c) LSCM images and c’) intensity map of *S. aureus* on sample disks after 72-h incubation. PE served as the control; each image is 200 × 200 μm. Three replicate samples were tested.

Briefly, 30 disk samples of P(X)(Y)-PE (diameter: 5 mm, thickness: 0.5 mm) were attached to three pieces of glass to form three mini-arrays (Figure 2a, 10 samples/piece). Pure PE disks served as the control. These polymer arrays were sterilized by a 75% ethanol aqueous solution and UV light irradiation (254
nm, 40 w, 30 min), then put into a Luria–Bertani (LB) broth (200 mL) followed by the addition of a suspension of planktonic *E. coli* or *S. aureus* (10 μL). The optical density of this suspension at UV ~ 600 nm (OD600) was approximately 1.0. The polymer arrays were incubated with bacteria for 72 h prior to observation by laser scanning confocal microscopy (LSCM) (Figure 2a). The fluorescence intensity at each sample disk reflected the number of attached bacteria.

From the LSCM images (Figure 2b, 2c), the P(X)(Y)-PE samples contained far fewer bacteria than the pure PE sample. This was interpreted as the antibacterial capability of these 1,4-DHP-containing polymers. The quantitative fluorescence intensity maps of the samples indicated almost no *E. coli* on **P(4)(1)**, **P(4)(3)**, **P(4)(4)**, and **P(5)(5)** (Figure 2b’); and almost no *S. aureus* on **P(1)(4)**, **P(4)(1)**, **P(4)(3)**, **P(4)(4)**, and **P(4)(5)** (Figure 2c’). The antibacterial efficiency of all poly(1,4-DHP)s was calculated by comparing the fluorescence intensity on P(X)(Y)-PE and PE (Table S2). **P(4)(1)**, **P(4)(3)**, and **P(4)(4)** demonstrated better antibacterial efficiency to both *E. coli* and *S. aureus* (>95%) than the other poly(1,4-DHP)s; thus, these three polymers were selected as the top three broad-spectrum antibacterial polymers for the next round of screening.

**Secondary screening** Commodity polymers are indispensable to everyday life. If items composed from these polymers can be imbued with antibacterial properties, the value of the items produced with such polymers would be increased greatly. Thus, the three best polymers from preliminary testing (Figure 3a) were blended into five commonly found commodity polymers (PE, PP, PET, PMMA, and PA66) in different ratios (33%, 20%, 10%, and 5%).
Figure 3. a) The three poly(1,4-DHP)s selected for further study after preliminary tests. b, c) Intensity map of the bacteria on sample disks (commodity polymers + P(X)(Y)), 72-h culture. Three replicate samples were tested, and pure commodity polymer disks served as the controls.

The antibacterial capabilities of these P(4)(1,3,4)-polymer samples were subjected to a similar test as before. Pure commodity polymer disks were used as the respective controls.

The fluorescence intensity maps (Figure 3b, 3c; Table S3) were obtained according to the LSCM images (Figure S13). The greater the P(4)(1,3,4) content, the fewer the bacteria found on the P(4)(1,3,4)-polymer blends, indicative of a concentration-dependent antibacterial property. When the poly(1,4-DHP) to commodity polymer blend was 10 wt.%, P(4)(4) offered the best antibacterial properties. Thus, P(4)(4) was selected for further research.

Development of a Model Antibacterial Utensil
For items made from these commodity polymer-antibacterial polymer blends to remain practical, the basic mechanical properties of the base commodity polymer must not be meaningfully compromised. Thus, the mechanical strength of the \( \text{P(4)(4)} \)-polymer samples (33 wt.% \( \text{P(4)(4)} \)) was tested using PE, PP, PET, PMMA, and PA66 (Figure S14). All five 33 wt.% \( \text{P(4)(4)} \)-polymer samples yielded the similar mechanical strength results as the pure-polymer controls. This suggests that commodity polymers can be easily blended with antibacterial \( \text{P(4)(4)} \) without a loss of mechanical strength.

Next, we applied the injection molding technique to form antibacterial plastic products from a blended \( \text{P(4)(4)} \) and PE mixture. PE was chosen as a representative sample owing to its easy machining and broad applicability as a packaging material. \( \text{P(4)(4)} \)-PE (33 wt.% of \( \text{P(4)(4)} \)) and PE were injection molded into plastic bowls (Figure 4a, diameter: 44 mm, height: 14 mm). These bowls were sterilized by a 75% ethanol aqueous solution and air-dried on a laboratory bench for 24 h. Then, various commonly consumed beverages (water, orange juice, milk, red wine, coffee, and black tea) (5 mL) were placed in the PE and \( \text{P(4)(4)} \)-PE bowls (Figure 4b). Glass covers were placed on the bowls, and the beverages were incubated at 37 °C. Aliquots were obtained at specified time points.

A plate-streaking experiment was performed to test for the presence of viable bacteria. After 24 h, beverage aliquots (100 µL) from the different bowls were taken and evenly coated on the surface of LB agar-coated petri dishes (diameter: 90 mm, thickness: 6 mm, LB: 1.5 g/L). These petri dishes were incubated at 37 °C prior to observation, and the original beverages were used as the controls (Figure 4c).
Figure 4. a) Bowls made from PE and P(4)(4)-PE. b) Commercially available beverages (5 mL) for testing and the experimental setup. c) Photographic images of the plate-streaking experiment: 100 μL sample on LB agar (15 g/L), 24-h culture.

Commercially available beverages are typically sterile; thus, almost no bacterial colonies were observed (Figure 4c, control). However, aliquots of water, orange juice, milk, and coffee from the pure PE bowls clearly contained bacteria, as demonstrated by the visible colonies grown on the LB agar petri
dishes (Figure 4c, PE). This suggests that the PE bowls were contaminated with germs when stored in the open air, leading to the contamination of these beverages inside. Notably, red wine and black tea samples from the pure PE bowls yielded similar results to the control during the LB agar growth test. This was attributed to the antibacterial compounds contained in the parent beverages, such as tannin, ethanol, and tea polyphenols (Figure 4c, PE). On the other hand, only milk in the P(4)(4)-PE bowl resulted in a small bacterial colony on the LB agar surface (Figure 4c, P(4)(4)-PE). This suggests that only milk contained sufficient nutrients to encourage growth of the small amount of bacteria present after 24 h of open-air storage and overcome the inherent antibacterial properties of the P(4)(4)-PE bowls.

Additional samples were obtained after 48 h and 72 h to repeat the plate-streaking experiments (Figure S15). Denser and denser bacterial colonies were observed in the pure PE group over time, indicating propagation of bacteria in beverages incubated in the pure PE bowls. The inherent antibacterial defenses of tea and red wine expired after 48 h and 72 h, respectively. Nevertheless, all beverages, except milk, in the P(4)(4)-PE bowls remained bacteria-free after 48 h of incubation (Figure S15a). Red wine and tea in P(4)(4)-PE bowls remained sterile even after 72-h incubation (Figure S15b). These results agreed well with the quantitative OD600 analyses (Figure S16), demonstrating great potential for the P(4)(4)-PE blended polymer as a new antibacterial material for utensils and food applications.

**Mechanism Study**

The possible mechanisms for antibacterial action of P(4)(4) were studied. Several polymers were used according to the different evaluation protocols.

**Bactericidal ability** Killing bacteria is a general strategy in developing antibacterial materials. Some 1,4-DHPs have been reported to effectively inhibit bacterial growth. Thus, the bactericidal ability of P(4)(4) was evaluated. Polyetherimide (PEI), a well-known antibacterial polymer that is lethal to bacteria,(Gibney et al., 2012) was used as the control. P(4)(4)-PE (33 wt.% P(4)(4)) and PEI-PE (33 wt.% PEI) disks were prepared as abovementioned. These two samples were attached to the bottom of a multi-well
plate and covered with LB broth (200 µL). Then, a suspension of planktonic *E. coli* or *S. aureus* (10 µL, OD600~1.0) was added. The plate was incubated at 37 °C for 24 h. The OD600 values were tested at different time intervals, and a pure PE disk was used as the blank control.

From the OD600 curves over time (Figure 5a), the rate of bacterial growth was slower with PEI-PE than with P(4)(4)-PE and PE, confirming the ability of PEI to kill bacteria. There was no observable difference in bacterial growth rates between P(4)(4)-PE and PE. This suggests P(4)(4) has negligible bactericidal effects. Thus, P(4)(4) achieves its antibacterial function by preventing bacterial adhesion instead of killing the bacteria outright.

**Figure 5.** a) OD600 values versus time in the presence of different polymers, 24-h culture. Data are represented as mean ± SD, n = 5. b) Intensity data of bacteria on sample disks after 72-h incubation. Data are represented as mean ± SD, n = 3; ** p < 0.01, compared with PE.

Then, the antibacterial abilities of samples were evaluated through the polymer-array method. After 72-h incubation, PEI effectively killed bacteria, resulting in much less bacteria on PEI-PE surface than on PE (Figure 5b, p < 0.01 compared with PE; Figure S17). However, almost no bacteria was detected on
the $\text{P}(4)(4)$-PE surface despite the poor sterilization ability of $\text{P}(4)(4)$ (Figure 5b, Figure S17). This suggests that resisting bacterial adhesion is a promising strategy to exploit new antibacterial materials.

**Molecular structures** Polymer functions are determined by their molecular structures. Thus, the relationship between the structure of $\text{P}(4)(4)$ and its antibacterial ability was studied.

![Molecular structures](image)

**Figure 6.** a) $\text{P}(4)(4)$, $\text{P}(\text{AEMA})$, P1 (polymer containing an aromatic group), P2 (polymer containing a long-branched olefin), and M1. b) Intensity data of bacteria on sample disks after a 72-h culture. Data are represented as mean ± SD, n = 3; * $p < 0.05$, compared with PE.

The Hantzsch ring in $\text{P}(4)(4)$ is composed from a long-branched olefin and aromatic groups. There are still some $\beta$-ketoester moieties (~2%) left in $\text{P}(4)(4)$. Thus, two polymers containing only the long-branched olefin and aromatic groups were prepared (Figure 6a, P1, P2; Figure S18, S19). A homopolymer of AEMA was also prepared (Figure 6a, P(AEMA), Figure S20). Meanwhile, a small molecule with similar Hantzsch ring as the side group of $\text{P}(4)(4)$ was prepared (Figure 6a, M1). These polymers (P1, P2, P(AEMA)) and M1 were mixed with PE to prepare samples, respectively. The antibacterial capabilities of these samples were studied via the polymer-array method. The sample composed of pure PE served as the blank.
The fluorescence signals of *E. coli* and *S. aureus* on the polymer samples were recorded (Figure S21). Quantitative data from the fluorescence intensity were calculated, and the number of bacteria present on the samples followed the order $\text{P(4)(4)} \ll \text{PAEMA} \approx \text{P2} < \text{P1} < \text{M1} \approx \text{PE}$ (Figure 6b). M1-PE had almost no antibacterial ability ($p = 0.118$ (*E. coli*), 0.137 (*S. aureus*), compared with PE), suggesting direct addition of free 1,4-DHP molecules to bulk commodity polymers would not be effective. $\text{P1}$, $\text{P2}$, and P(AEMA) demonstrated poor antibacterial capability ($p < 0.05$, compared with PE), whereas $\text{P(4)(4)}$ had the best antibacterial ability among all poly(1,4-DHP)s. These results suggest the polymer chain and 1,4-DHP pendant group both play key roles for the antibacterial function of $\text{P(4)(4)}$; the long-branched olefin and aromatic groups enhance the antibacterial ability of $\text{P(4)(4)}$ via a possible synergistic effect.

**Limitation of this study and possible Extension**

Here, a simple polymer (random structure with a broad polydispersity index (PDI)) with potential antibacterial applications was developed via a very simple HTP method (small sample pool and semi-quantitative measurements). This highlights the validity of HTP methods and MCRs in exploring new functional polymers. However, in addition to side-chain groups, the core polymer structure can also be important for polymer functions. Recent studies have reported that the antibacterial capability of polymers can be remarkably improved by tuning the polymer structures (i.e., monomer sequences and topology structures). (Judzewitsch et al., 2018; Kuroki et al., 2017; Lam et al., 2016; Namivandi-Zangeneh et al., 2018) Currently, well-defined polymers can be rapidly prepared via modern technologies in CRP, including single electron transfer-atom transfer radical polymerization (SET-ATRP), sulfur-free RAFT emulsion polymerization, photoinduced ATRP, and photoinduced electron/energy transfer-RAFT (PET-RAFT). (Anastasaki et al., 2016; Anastasaki et al., 2014; Boyer et al., 2016; Carmean et al., 2017; Chen et al., 2016; Engelis et al., 2017; Fors and Hawker, 2012; Gormley et al., 2018; Rosen and Percec, 2009; Xu et al., 2016; Zhang et al., 2013) The future combination of these modern CRP techniques with the method used here might offer new polymer libraries with different side groups, molecular weights, monomer
sequences, topology structures, etc. This will be helpful for developing new polymers with improved antibacterial capabilities and other bioactive functions.

CONCLUSIONS

In summary, we have developed a new one-pot HTP method by simultaneously carrying out the Hantzsch reaction and FRP in a high-throughput manner. Using this method, six aldehydes derived from natural products were used to rapidly synthesize 30 poly(1,4-DHP)s in one step. This highlighted the advantage of MCRs in amplifying molecular diversity and the superiority of a one-pot strategy in improving synthesis efficiency. The resulting poly(1,4-DHP)s were screened via HTP measurement methods to ultimately identify a polymer that could effectively resist bacterial attachment due to its unique 1,4-DHP side group. This opens the door to new opportunities for developing antibacterial compounds via the Hantzsch reaction. The developed one-pot method highlighted the value of HTP methods, natural product derivatives, and MCRs for exploring new functional polymers with potential applications in the real world. These results might prompt a broad study of MCRs and HTP methods in polymer science for developing other new functional polymers for interdisciplinary applications.

METHODS

All methods can be found in the accompanying Transparent Methods supplemental file.

Author contributions:
L. T. developed the concept and conceived the experiments. G. L., Y. L., H. W., and Y. Z. performed the laboratory experiments. L.T., G. L., Q. Z., and X. W. contributed to the experimental analyses. L. T. and G. L. wrote the manuscript. L. T. and Y. W. provide financial support to the research.

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Supplementary information:
Supplementary information (preparation of small molecules and polymers, antibacterial test, etc.) is available for this paper. Correspondence and requests for materials should be addressed to L. T.

Declaration of interests:

The authors declare no competing financial interests.

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High-Throughput Preparation of Antibacterial Polymers from Natural Product Derivatives via the Hantzsch Reaction

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Highlights

- A one-pot high-throughput (HTP) system was developed to prepare a polymer library
- A new antibacterial polymer was selected from this library via HTP measurements
- This antibacterial polymer was mixed in polyethylene to get antibacterial utensils
- Mechanism study revealed the Hantzsch reaction leads to new antibacterial polymers
High-Throughput Preparation of Antibacterial Polymers from Natural Product Derivatives \emph{via} the Hantzsch Reaction

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EXPERIMENTAL SECTION

1. Materials

All chemicals and solvents were purchased from commercial sources and used without further purification. 2-(Acetoacetoxy)ethyl methacrylate (AEMA, Aladdin, 95%), cinnamic aldehyde (MREDA, 99%), citronellal (MERYER, 99%), hyacinthin (J&K, 97.5%), myrac aldehyde (Energy, 97%), cyclamen aldehyde (MAKLIN, 92%), phellandral (MAKLIN, 97%), 5,5-dimethyl-1,3-cyclohexanedione (MAKLIN, 99%), 1,3-cyclohexanedione (Shaoyuan, 97%), 5-methyl-1,3-cyclohexanedione (NineDing, 98%), 5-phenyl-1,3-cyclohexanedione (Energy, 97%), 5-n-propyl-1,3-cyclohexanedione (Energy, 97%), ammonium acetate (MAKLIN, 98%), glycine (Ouhe, 98%), 2,2'-azobisisoheptonitrile (ABVN, Energy, 98%), polyethylene (PE, 500 mesh, ZhongLian Plastic), polypropylene (PP, 500 mesh, ZhongLian Plastic), polyethylene terephthalate (PET, 500 mesh, ZhongLian Plastic), poly(methyl methacrylate) (PMMA, 500 mesh,
ZhongLian Plastic), polyamide 66 (PA66, 500 mesh, ZhongLian Plastic), poly(ethylene imine) (PEI, $M_w \sim 10000$, 50 wt/% H$_2$O), 3-bromo-1-propanol (Ark-Pharm, 97%), methacryloyl chloride (HEOWNS, 99%), sodium borohydride (NaBH$_4$, LanGe, 98%).

Escherichia coli (E. coli), BL21 (competent cell of E. coli, Merck Millipore)), Staphylococcus aureus strain (S. aureus, CMCC), pCMV-N-mCherry (Beyotime), Luria-Bertani (LB) broth (Gibco), LB agar (Gibco), tryptic soy broth (TSB, Gibco), tryptic soy agar (TSA, Gibco), kanamycin sulfate (inalco), iso-propyl β-D-thiogalactoside (IPTG, inalco), phosphatic buffer solution (PBS, Gibco) were used as purchased.

2. Instruments

Gel permeation chromatography (GPC) analyses of polymers were performed using N,N-dimethyl formamide (DMF) containing 50 mM LiBr as the eluent. The GPC system is a Shimadzu LC-20AD pump system consisting of an auto injector, a MZ-Gel SD plus 10.0 μm guard column (50 × 8.0 mm, 10$^2$ Å) followed by two PLgel 5 μm MIXED-D columns (300 × 7.5 mm), and a Shimadzu RID-10A refractive index detector. The system was calibrated with narrow molecular weight distribution polystyrene standards ranging from 200 to 10$^6$ g mol$^{-1}$.

NMR spectra were obtained using a JEOL JNM-ECA400 spectrometer for all samples. The ESI-MS data were collected using a Micro TOF-QII Bruker. The FT-IR spectra were recorded in a transmission mode on a Perkin-Elmer Spectrum 100 spectrometer (Waltham, MA, USA).

Differential scanning calorimetry (DSC) was performed using a TA instrument Q2000 operated at a scanning rate of 10°C/min. Thermal gravimetric analysis (TGA) was conducted on a TA instrument Q50 with a heating rate of 20°C/min. The fluorescent images of bacteria were recorded by a Laser scanning Confocal Microscopy (Zeiss LSM 780). Bio-Rad Electroporator was used for the electrotransformation of E. coli and S. aureus. The tensile shear strength was measured using an MTS SYSTEMS (CHINA) Co.
Ltd. SANS CMT6503 electromechanically universal testing machine. Injection molding was performed by a micro injection molding machine (WZS10D, Shanghai Xinshuo precision instrument co., LTD).

3. Methods

3.1 2-(Methacryloyloxy)ethyl (E)-2,7,7-trimethyl-5-oxo-4-styryl-1,4,5,6,7,8-hexahydroquinoline-3-carboxylate (M(1)(1))

The monomers M(X)(1) were prepared via the Hantzsch reaction by different combinations of aldehydes (A(X)) and 5,5-dimethyl-1,3-cyclohexanedione (B(1)). Typically, A(1) (cinnamaldehyde, 660 mg, 5.0 mmol), AEMA (1.07 g, 5.0 mmol), B(1) (700 mg, 5.0 mmol), and ammonium acetate (578 mg, 7.5 mmol) were put in a 15-mL centrifuge tube. Then, glycine (38 mg, 0.5 mmol) and acetonitrile (6.0 mL) were added. The mixture was kept in an oil bath (75 °C) for 4 h. After removing volatiles under vacuum, the crude was purified by a column chromatography (petroleum ether/ethyl acetate: 5/1) to get monomer M(1)(1) as a yellow powder (1.65 g, yield: 73.6%).

All other monomers were prepared through the same procedure.

\(^1\)H NMR (400 MHz, CDCl\(_3\), \(\delta/\text{ppm}\)): 7.06-7.29 (m, 5H, ph), 6.20 (s, 1H, CH\(\text{CH=CH}\)), 6.19 (s, 1H, CH\(\text{CH=CH}\)), 6.12 (s, 1H, CH\(\text{=C}\)), 6.06 (s, 1H, NH), 5.57 (s, 1H, CH\(\text{=C}\)), 4.54 (s, 1H, CCHC\(_3\)), 4.25-4.44 (m, 4H, COOC\(\text{H}_2\)CH\(_2\)), 2.36 (s, 2H, CH\(\text{=C}=\text{O}\)), 2.09 (s, 3H, NHCH\(\text{CH}_3\)), 2.00 (s, 3H, CH\(_3\)C=CH\(_2\)), 1.90 (s, 2H, NHCH\(\text{CH}_3\)), 1.08 (s, 6H, CH\(_3\)CH\(_3\)).
$^{13}$C NMR (100 MHz, CDCl$_3$, δ/ppm): 196.15, 167.70, 167.29, 149.84, 147.27, 145.23, 135.96, 128.02, 127.98, 127.97, 127.96, 127.95, 126.96, 126.95, 126.19, 111.79, 105.18, 62.80, 61.51, 50.93, 40.58, 36.70, 32.70, 29.59, 27.13, 19.29, 18.38.

IR (ν/cm$^{-1}$): 3287, 2959, 1716, 1692, 1603, 1488, 1377, 1279, 1214, 1149, 1113, 1082, 1045, 941, 884, 814, 783, 695, 658, 566.

ESI-MS: observed (expected): 450.2286 (450.2280) [M + H$^+$].

3.2 2-(Methacryloyloxy)ethyl 4-(2,6-dimethylhept-5-en-1-yl)-2,7,7-trimethyl-5-oxo-1,4,5,6,7,8-hexahydroquinoline-3-carboxylate (M(2)(1))

$^1$H NMR (400 MHz, CDCl$_3$, δ/ppm): 6.12 (s, 1H, CH$_2$=C), 5.83 (s, 1H, NH), 5.57 (s, 1H, CH$_2$=C), 5.05 (m, 1H, CH=C(CH$_3$)$_2$), 4.25-4.44 (m, 4H, COOCH$_2$CH$_2$), 4.05 (s, 1H, CCHC), 2.36 (s, 2H, CH$_2$C=O), 2.09 (s, 3H, NHCCCH$_3$), 2.00 (s, 3H, CH$_3$C=CH$_2$), 1.90 (s, 2H, NHCCCH$_2$), 1.60 (s, 3H, CH=CCH$_3$), 1.55 (s, 3H, CH=CCH$_3$), 1.25 (m, 7H, CH$_2$CHCH$_3$CH$_2$CH$_2$), 1.08 (s, 6H, CH$_2$C(CH$_3$)$_2$), 0.86 (d, 3H, CHCH$_3$, J = 4.2 Hz).

$^{13}$C NMR (100 MHz, CDCl$_3$, δ/ppm): 196.28, 167.70, 167.27, 148.98, 145.00, 136.03, 130.73, 128.93, 126.21, 112.53, 105.74, 62.85, 61.53, 50.98, 44.86, 41.34, 38.32, 37.49, 32.42, 29.85, 27.70, 27.43, 25.81, 25.62, 20.24, 19.51, 18.40, 17.68.
IR (ν/cm⁻¹): 3675, 3285, 2971, 2901, 1719, 1694, 1637, 1604, 1480, 1451, 1405, 1393, 1319, 1241, 1214, 1164, 1075, 1056, 938, 891, 813, 776, 661.

ESI-MS: observed (expected): 472.3061 (472.3063) [M + H⁺].

3.3 2-(Methacryloyloxy)ethyl 4-benzyl-2,7,7-trimethyl-5-oxo-1,4,5,6,7,8-hexahydroquinoline-3-carboxylate (M(3)(1))

\[ \text{IR (ν/cm}^{-1}\text{): 3675, 3285, 3062, 2959, 1719, 1693, 1603, 1486, 1452, 1383, 1320, 1296, 1279, 1220, 1165, 1106, 1079, 1048, 1004, 941, 814, 753, 699, 595.} \]

ESI-MS: observed (expected): 438.2287 (438.2280) [M + H⁺].

3.4 2-(Methacryloyloxy)ethyl 2,7,7-trimethyl-4-(4-(4-methylpent-3-en-1-yl) cyclohex-3-en-1-yl)-5-oxo-1,4,5,6,7,8-hexahydroquinoline-3-carboxylate (M(4)(1))

\[ 1^1 \text{H NMR (400 MHz, CDCl}_3, \delta/\text{ppm): 6.95-7.25 (m, 5H, ph), 6.12 (s, 1H, CH}_2=\text{C), 5.76 (s, 1H, NH), 5.57 (s, 1H, CH}_2=\text{C), 4.31 (s, 1H, CCHC), 4.18-4.30 (m, 4H, COOCH}_2\text{CH}_2, 2.62 (m, 2H, phCH}_2, 2.36 (s, 2H, CH}_2=\text{O), 2.09 (s, 3H, NHCCCH}_3, 2.00 (s, 3H, CH}_3\text{C=CH}_2, 1.90 (s, 2H, NHCCCH}_2, 1.08 (s, 6H, CH}_3\text{CCH}_3.} \]

\[ 1^3 \text{C NMR (100 MHz, CDCl}_3, \delta/\text{ppm): 196.08, 167.99, 167.21, 149.74, 145.27, 137.36, 135.94, 131.24, 126.10, 124.50, 120.43, 110.02, 103.37, 62.74, 61.74, 50.94, 41.99, 41.31, 37.53, 34.11, 32.39, 29.85, 27.63, 26.48, 19.45, 17.67.} \]

IR (ν/cm⁻¹): 3675, 3286, 3062, 2959, 1719, 1693, 1603, 1486, 1452, 1383, 1320, 1296, 1279, 1220, 1165, 1106, 1079, 1048, 1004, 941, 814, 753, 699, 595.

ESI-MS: observed (expected): 438.2287 (438.2280) [M + H⁺].
\[ \begin{align*}
\text{H NMR} (400 \text{ MHz}, \text{CDCl}_3, \delta/\text{ppm}): & \quad 6.12 \text{ (s, 1H, CH\textsubscript{2}=C)}, 5.95 \text{ (s, 1H, NH)}, 5.57 \text{ (s, 1H, CH\textsubscript{2}=C)}, 5.29 \text{ (m, 1H, CHCH\textsubscript{2}CH=CH\textsubscript{2})}, 5.05 \text{ (d, 1H, COOCH\textsubscript{2}CH\textsubscript{2})}, 4.12 \text{ (s, 1H, CCH\textsubscript{2}CH\textsubscript{2}CH=CH\textsubscript{2})}, 2.36 \text{ (s, 2H, CH\textsubscript{2}CH=CH\textsubscript{2})}, 2.09 \text{ (s, 3H, NHCCCH\textsubscript{3})}, 2.00 \text{ (s, 3H, CH\textsubscript{3}C=CH\textsubscript{2})}, 1.95-2.06 \text{ (m, 4H, CH\textsubscript{2}CH\textsubscript{2}CH=CH\textsubscript{2})}, 1.70-2.10 \text{ (m, 7H, CH\textsubscript{2}CHCH\textsubscript{2}CH\textsubscript{2})}, 1.90 \text{ (s, 2H, NHCCCH\textsubscript{2})}, 1.81 \text{ (s, 3H, CHCCCH\textsubscript{3})}, 1.76 \text{ (s, 3H, CHCCCH\textsubscript{3})}, 1.08 \text{ (s, 6H, CH\textsubscript{3}CH\textsubscript{3})}.
\end{align*} 
\]

\[ \begin{align*}
\text{C NMR} (100 \text{ MHz}, \text{CDCl}_3, \delta/\text{ppm}): & \quad 196.08, 167.80, 167.21, 149.70, 145.27, 137.36, 135.97, 131.24, 126.10, 124.50, 120.43, 110.14, 103.37, 62.74, 61.47, 50.97, 41.99, 41.31, 37.53, 34.11, 32.39, 29.85, 29.23, 27.63, 27.34, 26.48, 25.70, 19.48, 18.30, 18.30, 17.67.
\end{align*} 
\]

IR (\(\nu/\text{cm}^{-1}\)): 3675, 3281, 2968, 2901, 1720, 1694, 1606, 1483, 1383, 1280, 1214, 1153, 1105, 1076, 1045, 997, 938, 883, 788, 754, 610, 588, 565.

ESI-MS: observed (expected): 510.3224 (510.3219) [M + H\textsuperscript{+}].

3.5 2-(Methacryloyloxy)ethyl 4-(1-(4-isopropylphenyl)propan-2-yl)-2,7,7-trimethyl-5-oxo-1,4,5,6,7,8-hexahydroquinoline-3-carboxylate (M(5)(1))
$^1$H NMR (400 MHz, CDCl$_3$, δ/ppm): 6.95-7.09 (m, 4H, ph), 6.12 (s, 1H, CH$_2$=C), 5.89 (s, 1H, NH), 5.57 (s, 1H, CH$_2$=C), 4.32-4.44 (m, 4H, COOCH$_2$CH$_2$), 4.19 (s, 1H, CCHC), 2.92 (m, 2H, CH$_2$ph), 2.30 (m, 1H, phCH), 2.36 (s, 2H, CH$_2$C=O), 2.09 (s, 3H, NHCH$_3$), 2.00 (s, 3H, CH$_3$C=CH$_2$), 1.90 (s, 2H, NHCH$_2$), 1.55-1.65 (m, 1H, CH$_2$CHCH$_3$), 1.10 (d, 3H, CH$_2$CHCH$_3$, J = 6.5 Hz), 1.08 (s, 6H, CH$_3$CCH$_3$), 0.67 (d, 6H, CH$_3$CHCH$_3$, J = 6.5 Hz).

$^{13}$C NMR (100 MHz, CDCl$_3$, δ/ppm): 197.13, 167.23, 164.74, 146.93, 139.16, 137.36, 135.94, 135.41, 129.02, 126.51, 126.07, 125.03, 124.04, 110.04, 103.57, 68.15, 62.36, 51.98, 46.55, 40.24, 33.78, 32.99, 32.40, 30.03, 28.36, 27.19, 25.31, 24.16, 18.37, 17.87, 15.22.

IR (ν/cm$^{-1}$): 3285, 2958, 1720, 1697, 1604, 1483, 1382, 1275, 1217, 1150, 1113, 1076, 1048, 998, 938, 886, 856, 812, 783, 654, 590, 566, 554.

ESI-MS: observed (expected): 508.3069 (508.3063) [M + H$^+$].

3.6 2-(Methacryloyloxy)ethyl 4-hexyl-2,7,7-trimethyl-5-oxo-1,4,5,6,7,8-hexahydroquinoline-3-carboxylate (M(6)(1))
$^1$H NMR (400 MHz, CDCl$_3$, δ/ppm): 6.12 (s, 1H, CH$_2$=C), 6.05 (s, 1H, NH), 5.57 (s, 1H, CH$_2$=C), 4.25-4.44 (m, 4H, COOCH$_2$CH$_2$), 4.00 (s, 1H, CCHC), 2.36 (s, 2H, CH$_2$C=O), 2.09 (s, 3H, NHCCCH$_3$), 2.00 (s, 3H, CH$_3$C=CH$_2$), 1.90 (s, 2H, NHCCH$_2$), 1.09-1.50 (m, 10H, CH(CH$_2$)$_2$CH$_3$), 1.08 (s, 6H, CH$_3$CCH$_3$), 0.86 (s, 3H, CH$_2$CH$_3$).

$^{13}$C NMR (100 MHz, CDCl$_3$, δ/ppm): 196.14, 167.62, 167.29, 149.90, 145.38, 136.08, 126.15, 111.33, 104.67, 62.81, 61.47, 51.02, 41.08, 36.34, 32.59, 32.01, 29.84, 29.75, 27.09, 25.01, 22.76, 19.43, 18.36, 14.16, 14.15.

IR (ν/cm$^{-1}$): 3281, 2926, 1722, 1693, 1605, 1483, 1386, 1278, 1216, 1163, 1094, 1048, 995, 954, 884, 813, 780, 747, 641, 618, 587, 575, 558.

ESI-MS: observed (expected): 432.2759 (432.2750) [M + H$^+$].

3.7 Polymerization of M(X)(1) to get P(X)(1)
The ‘pure’ polymers P(X)(1) were prepared by radical polymerization of M(X)(1). For example, monomer M(1)(1) (450 mg, 1.0 mmol), 2,2’-azobis-(2,4-dimethylvaleronitrile) (ABVN, 5 mg, 0.02 mmol) were dissolved in dimethyl formamide (DMF, 1.0 mL). The mixture was purged by nitrogen flow for 20 min, then kept in an oil bath (75 °C) for 12 h. The polymerization was quenched in an ice–water bath. After precipitation in diethyl ether three times, P(1)(1) was obtained as a yellow powder (400 mg, yield: 88.9%).

All other ‘pure’ polymers were prepared through the same procedure by using different monomers.

### 3.8 HTP-One pot preparation of polymers (P(X)(Y))

Polymers P(X)(Y) were prepared via a one-pot HTP method. For a representative synthesis, A(1) (cinnamaldehyde, 660 mg, 5.0 mmol), AEMA (1.07 g, 5.0 mmol), B(1) (5,5-dimethyl-1,3-cyclohexanedi-one, 700 mg, 5.0 mmol), and ammonium acetate (578 mg, 7.5 mmol) were charged to a 15-mL centrifuge tube. Then, glycine (38 mg, 0.5 mmol), ABVN (25 mg, 0.1 mmol), and acetonitrile/1,4-dioxane (3.0/3.0 mL) were added. The mixture was purged by bubbling nitrogen for 20 min, then sealed and placed in an oil bath (75 °C) for 12 h. The polymerization was quenched in an ice–water bath. A 20–50 µL aliquot was taken for ^1^H NMR and GPC analyses. The polymer was purified by precipitation in water followed by re-precipitation from THF with diethyl ether to obtain a yellow powder (P(1)(1) (1.85 g, yield: 82.3%).

All other polymers were simultaneously prepared using the same procedure with the respective A and B starting materials.

### 3.9 Red-fluorescent-protein (RFP) transferred bacteria

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RFP transferred bacteria were prepared according to previous literatures. BL21 is a competent cell line of *E. coli*. BL21 cells were transferred to a chilled electroporation cuvette with a precooled plasmid solution (2 μL, pCMV-N-mCherry in PBS, 100 μg/mL, kanamycin resistance). This cuvette was put in the electroporation instrument for electric shock with an exponential decay pulse (2.5 kV, 200-Ω resistance, 25-μF capacitance, 4.5 ms). Then, LB medium (1 mL) was added to the electroporation cuvette, and the contents were transferred to a 1.5-mL tube for incubation (1.5 h, 180 r/min, 37 °C). The mixture was plated on a LB agar plate containing antibiotic (kanamycin sulfate: 50 μg/mL) and incubated at 37 °C for 12 h. One small colony was isolated and incubated in 200 mL of LB medium for 2 h. Then, IPTG (200 mM, 2 mL) was added to induce gene expression (28 °C, 4 h) to obtain the RFP-*E. coli*.

RFP-*S. aureus* was similarly prepared.

### 3.10 Poly(acetoacetoxy ethyl methacrylate) (P(AEMA))

AEMA (2.14 g, 10 mmol) and ABVN (50 mg, 0.2 mmol) were dissolved in DMF (20 mL) and put in a 50 mL polymerization tube. This tube was sealed with a rubber septum and purged by nitrogen flow for 20 min, and kept in an oil bath (75 °C) for 12 h. The polymerization was quenched in an ice–water bath, and the polymer (P(AEMA)) was purified by precipitation in diethyl ether three times as a white powder (1.82 g, yield: 85.0%).

### 3.11 Poly((4-(4-methylpent-3-en-1-yl)cyclohex-3-en-1-yl)methyl methacrylate) (P1)

**A4-OH**: Myrac aldehyde (A4, 9.62 g, 50 mmol) and sodium borohydride (NaBH₄, 2.27 g, 60 mmol) were dissolved in methanol (50 mL), and put in an ice bath for 4 h. After removing volatiles under vacuum,
A4-OH was purified by a column chromatography (petroleum ether/ethyl acetate = 10/1) as a colorless oil (8.59 g, 88.4% yield).

\(^1\)H NMR (400 MHz, CDCl\(_3\), \(\delta/\text{ppm}\)): 5.45 (d, 1H, CH\(_2\)CH\(_2\)C=CH, \(J = 6.8\) Hz), 5.05 (t, 1H, CH\(_2\)CH\(_2\)CH=CH, \(J = 6.9\) Hz), 3.58 (m, 2H, CH\(_2\)O), 2.20-2.10 (m, 7H, CH\(_2\)CH\(_2\)CH=CH), 2.10-2.00 (m, 4H, CH\(_2\)CH\(_2\)-), 1.85 (s, 3H, CH=CC\(_3\)CH), 1.82 (s, 3H, CH=CC\(_3\)CH).

\(^13\)C NMR (100 MHz, CDCl\(_3\), \(\delta/\text{ppm}\)): 137.89, 131.47, 124.44, 119.54, 67.91, 37.79, 36.44, 28.30, 27.88, 26.55, 25.79, 25.75, 17.77.

IR (\(\nu/\text{cm}^{-1}\)): 3324, 2965, 2913, 1670, 1437, 1376, 1261, 1232, 1139, 1087, 1049, 943, 892, 825, 673, 600, 581, 563.

ESI-MS: observed (expected): 195.1757 (195.1749) [M + H\(^+\)].

\(M(A4)\): A4-OH (1.94 g, 10 mmol) and trimethylamine (2.0 g, 20 mmol) was solved in CH\(_2\)Cl\(_2\) (20 mL) followed by adding methacryloyl chloride (1.3 g, 12 mmol) dropwise slowly. The mixture was kept at 20°C for 6 h. M(A4) was purified by a column chromatography (petroleum ether/ethyl acetate = 10/1) as a colorless oil (1.95 g, 74.4% yield).

\(^1\)H NMR (400 MHz, CDCl\(_3\), \(\delta/\text{ppm}\)): 6.12 (s, 1H, CH\(_2\)CH=CH\(_3\)), 5.67 (s, 1H, CH\(_2\)CH=CH\(_3\)), 5.45 (d, 1H, CH\(_2\)CH=CH=CH, \(J = 6.8\) Hz), 5.05 (d, 1H, CH\(_2\)CH\(_2\)CH=CH, \(J = 6.9\) Hz), 3.58 (m, 2H, CH\(_2\)O), 2.10-2.20 (m, 7H, CH\(_2\)CH\(_2\)CH=CH), 2.00-2.10 (m, 4H, CH\(_2\)CH\(_2\)CH=CH), 2.00 (s, 3H, CH\(_3\)C=CH\(_2\)), 1.85 (s, 3H, CH=CCH\(_3\)), 1.82 (s, 3H, CH=CCH\(_3\)).

\(^13\)C NMR (100 MHz, CDCl\(_3\), \(\delta/\text{ppm}\)): 167.63, 137.82, 136.62, 131.52, 125.32, 124.38, 119.25, 69.01, 37.76, 33.21, 28.45, 27.73, 26.53, 25.87, 25.79, 18.44, 17.77.
IR (v/cm⁻¹): 3676, 2967, 2914, 1718, 1638, 1452, 1403, 1376, 1320, 1296, 1104, 1066, 1049, 1013, 984, 937, 813, 760, 726, 683, 580.

ESI-MS: observed (expected): 285.1839 (285.1831) [M + Na⁺].

**P1:** M(A4) (2.62 g, 10 mmol) and ABVN (50 mg, 0.1 mmol) were dissolved in DMF (20 mL). The mixture was purged by nitrogen flow for 20 min, and kept in an oil bath (75°C) for 12 h. The polymerization was quenched in an ice-water bath, and P1 was precipitated in diethyl ether three times as a white powder (2.08 g, 79.4% yield).

### 3.12 Poly(3-(2,6-dioxo-4-phenylcyclohexyl)propyl methacrylate) (P2)

**B4-OH:** 5-Phenyl-1,3-cyclohexanedione (B4, 9.40 g, 50 mmol), potassium iodide (KI, 10 g, 60 mmol), potassium carbonate (K₂CO₃, 13.8 g, 50 mmol) and 3-bromo-1-propanol (6.9 g, 50 mmol) were dissolved in acetone (50 mL). This solution was refluxed for 12 h. After removing acetone under vacuum, B4-OH was purified by a column chromatography (petroleum ether/ethyl acetate = 5/1) as a colorless oil (8.86 g, 72.1% yield).

¹H NMR (400 MHz, CDCl₃, δ/ppm): 7.10-7.45 (m, 5H, ph), 5.48 (s, 1H, CH₂OH), 4.06 (m, 2H, CH₂OH), 3.92 (m, 1H, CCHC), 3.41 (t, 1H, CH₂CHCH₂, J = 5.2 Hz), 2.62 (m, 4H, CH₂CHCH₂), 2.60 (m, 2H, CHCH₂CH₂), 2.00 (m, 2H, CH₃CH₂OH).

¹³C NMR (100 MHz, CDCl₃, δ/ppm): 199.18, 199.18, 142.70, 128.90, 128.90, 127.17, 126.77, 126.77, 67.75, 62.27, 43.91, 43.91, 39.43, 36.71, 31.50.
IR (ν/cm⁻¹): 3403, 3029, 2944, 1717, 1598, 1497, 1453, 1403, 1374, 1352, 1318, 1289, 1257, 1245, 1231, 1207, 1161, 1044, 1007, 975, 940, 910, 869, 816, 760, 733, 699, 657, 615, 600.

ESI-MS: observed (expected): 247.1339 (247.1334) [M + H⁺].

**M(B4):** B4-OH (2.5 g, 10 mmol) and trimethylamine (2.0 g, 20 mmol) were solved in CH₂Cl₂ (25 mL). Methacryloyl chloride (1.3 g, 12 mmol) was added dropwise slowly. The mixture was kept at 25°C for 1 h. After removing the white solid by filtration and the volatiles under vacuum, M(B4) was purified by a column chromatography (petroleum ether/ethyl acetate = 5/1) as a colorless oil (2.67 g, 85.1% yield).

1H NMR (400 MHz, CDCl₃, δ/ppm): 7.10-7.45 (m, 5H, ph), 6.12 (s, 1H, CH₂=C), 5.57 (s, 1H, CH₂=C), 4.21 (t, 2H, CH₂O, J = 6.2 Hz), 4.02 (m, 1H, CCHC), 3.41 (t, 1H, CH₂C₃H₂, J = 6.7 Hz), 2.62 (m, 4H, C₃H₂CHC₂H₂), 2.60 (m, 2H, CHC₂H₂CH₂), 2.00 (m, 2H, CH₂C₃H₂CH₂), 1.98 (s, 3H, CH₂=CC₃H₃).

13C NMR (100 MHz, CDCl₃, δ/ppm): 198.93, 176.91, 167.34, 142.68, 136.19, 128.90, 128.90, 127.16, 126.77, 126.77, 125.91, 102.80, 65.38, 61.08, 43.91, 39.40, 36.60, 28.06, 18.40.

IR (ν/cm⁻¹): 3675, 2958, 2901, 1716, 1654, 1602, 1497, 1469, 1454, 1428, 1403, 1374, 1351, 1318, 1296, 1257, 1231, 1207, 1161, 1044, 1007, 975, 940, 910, 869, 816, 760, 733, 699, 657, 613, 588.

ESI-MS: observed (expected): 337.1422 (337.1416) [M + Na⁺].

**P2:** M(B4) (1.57 g, 10 mmol) and ABVN (50 mg, 0.2 mmol) were dissolved in DMF (20 mL). This solution was purged by nitrogen flow for 20 min, then kept in an oil bath (75°C) for 12 h. The polymerization was quenched in an ice–water bath. P2 was purified by precipitation in diethyl ether three times as a white powder (1.37 g, 87.3% yield).

### 3.13 Ethyl 2-methyl-4-(4-(4-methylpent-3-en-1-yl)cyclohex-3-en-1-yl)-5-oxo-7-phenyl-1,4,5,6,7,8-hexahydroquinoline-3-carboxylate (M1)
Myrac aldehyde (A(4), 1.92 g, 10 mmol), ethyl acetoacetate (1.30 g, 10 mmol), 5-phenyl-1,3-cyclohexanedione (B(4), 1.89 g, 10 mmol), and ammonium acetate (1.16 g, 15 mmol) were put in a 15-mL centrifuge tube. Then, glycine (76 mg, 1 mmol) and acetonitrile (9.0 mL) were added. The mixture was kept in an oil bath (75°C) for 4 h. Acetonitrile was removed under vacuum, and the crude was purified by a column chromatography (petroleum ether/ethyl acetate: 5/1) to get M1 as a yellow oil (3.86 g, yield: 81.6%).

$^1$H NMR (400 MHz, CDCl$_3$, δ/ppm): 7.10-7.33 (m, 5H, ph), 6.08 (s, 1H, NH), 5.29 (s, 1H, CHCH$_2$CH=CH=C), 5.09 (s, 1H, CH$_2$CH$_2$CH=CH=C), 4.13 (m, 2H, CH$_2$O), 4.12 (s, 1H, CCHC), 3.51 (m, 1H, phCH), 2.36 (m, 2H, CH$_2$C=O), 2.09 (s, 3H, NHCCH$_3$), 1.90 (s, 2H, NHCC$_2$H), 1.95-2.06 (m, 4H, CCH$_2$CH$_2$CH), 1.70-2.10 (m, 7H, CH$_2$CHCH$_2$CH$_2$), 1.81 (s, 3H, CHCCH$_3$), 1.76 (s, 3H, CHCH$_3$), 1.17 (m, 3H, CH$_3$CH$_2$O).

$^{13}$C NMR (100 MHz, CDCl$_3$, δ/ppm): 195.62, 168.44, 150.57, 144.00, 142.64, 137.34, 131.33, 128.79, 128.79, 127.05, 126.84, 126.84, 124.61, 120.49, 111.40, 104.00, 59.88, 43.78, 41.97, 41.57, 39.58, 37.64, 34.90, 34.55, 29.32, 27.68, 26.63, 25.80, 19.36, 17.78, 14.49.

IR (v/cm$^{-1}$): 3288, 2980, 2908, 1738, 1696, 1641, 1605, 1485, 1449, 1372, 1236, 1215, 1157, 1105, 1070, 1045, 937, 915, 846, 786, 762, 699, 634, 607, 560.

ESI-MS: observed (expected): 474.3015 (474.3008) [M + H$^+$].

3.14 HTP measurements of the antibacterial of polymers
Preliminary screening: As a typically example, P(1)(1) (1.5 g) and PE powder (3.0 g) were mixed and grinded evenly in a mortar. The mixture was manufactured to a square membrane (40 × 40 mm, thickness: 0.5 mm) by the hot-press technology under 150°C. Then, P(1)(1)-PE discs (diameter: 5 mm) were prepared by using a puncher. Other P(X)(Y)-PE samples were similarly prepared.

All P(X)(Y)-PE samples were attached on glass slides (25 mm × 75 mm × 1 mm) to get mini-arrays (10 samples/piece). These mini-arrays were used to test the antibacterial ability of P(X)(Y) with PE as the control. Briefly, these polymer mini-arrays were sterilized by a 75% ethanol aqueous solution and UV light (254 nm, 40 w, 30 min), then put into a Luria-Bertani (LB) broth (200 mL) followed by addition of a suspension of planktonic E. coli or S. aureus (10 μL). The optical density of this suspension at UV~600 nm (OD600) is approximately 1.0. The polymer arrays were incubated with bacteria for 72 h followed by washing twice with water and fixation with paraformaldehyde (4%) prior to observation by laser scanning confocal microscopy (LSCM, excitation wavelength: 543 nm; emission wavelength: 566–719 nm).

Secondary screening: P(X)(Y)-polymer samples were prepared by abovementioned method. These samples were attached on glass slides for antibacterial test as abovementioned. Commodity polymers (PP, PET, PMMA, PA-66) were used as the controls.

Antibacterial capabilities of all other polymers and small molecules (PEI, P(AEMA), P1, P2, and M1) were similarly tested through the polymer-array method.

3.15 Mechanical properties of P(4)(4)-polymer samples

The tensile shear strength of P(4)(4)-polymer samples were tested according to the national standard method (GBT 1040.3-2006). As a typical example, P(4)(4) (5 g) was mixed with PE (10 g) to prepare dumbbell-shaped splines by hot pressing. The dumbbell-shaped splines were in agreement with the national standard (length: 120 mm, breadth: 25 mm, thickness: 0.5 mm; length of narrow part: 25 mm, breadth of narrow part: 6 mm, thickness of narrow part: 0.5 mm). These splines were tested by an MTS
SYSTEMS (CHINA) Co. Ltd. SANS CMT6503 electromechanically universal testing machine (tensile speed: 5 mm/min; 25°C).

All other P(4)(4)-polymer splines were prepared and tested through the same procedure. General polymers were used as the controls. The data are presented as mean ± SD (n = 6).

3.16 Injection molding for plastic bowls

The plastic bowls were produced by injection molding technology. Typically, the uniform mixture of P(4)(4)-PE (33 wt.% of P(4)(4), 3 g) was added to the machine and heated (150 °C, 2 minutes) to melt-down. Then, the molten polymer was injected into a mold for bowl (0.6 Mpa, 1 minute) followed by keeping at 25 °C (normal atmosphere, 10 minutes) to get a plastic P(4)(4)-PE bowl.

The PE bowls were prepared through the same procedure.

3.17 Statistical Analyses

Results were analyzed with SPSS 25.0 and MedCalc 18.1 and are presented as mean ± SD as indicated. Comparisons were performed between two groups using a two-tailed Student’s t-test or ANOVA test when comparing more than two conditions. For all analyses, *p < 0.05 and **p < 0.01 were considered statistically significant. Each experiment was carried out at least three times independently. The sample size is pre-estimated to ensure statistical analysis and no sample was optionally excluded from analysis. No blinding was done in the analyses and quantifications.
Supporting Data

Figure S1. $^1$H NMR spectra (CDCl$_3$, 400M) of crude $P(X)(1)$ after HTP polymerization for conversion calculation.
Figure S2. GPC traces of P(X)(1).
Figure S3. a) Reaction conditions: acetonitrile/dioxane (1/1, v/v) as the solvent, glycine (10% to aldehyde) as the Hantzsch reaction catalyst, ABVN (2% to AEMA) as the initiator for FRP, 75 °C, and 12 h. A(X): B(2): AEMA: NH₄OAc = 1:1:1:1.5. b) ¹H NMR spectra (CDCl₃, 400M) of P(X)(2).
Figure S4. a) Reaction conditions: acetonitrile/dioxane (1/1, v/v) as the solvent, glycine (10% to aldehydes) as the Hantzsch reaction catalyst, ABVN (2% to AEMA) as the initiator for FRP, 75 °C, and 12 h. A(X): B(3): AEMA: NH₄OAc = 1:1:1:1.5. b) ¹H NMR spectra (CDCl₃, 400M) of P(X)(3).
**Figure S5.** a) Reaction conditions: acetonitrile/dioxane (1/1, v/v) as the solvent, glycine (10% to aldehyde) as the Hantzsch reaction catalyst, ABVN (2% to AEMA) as the initiator for FRP, 75 °C, and 12 h. \( A(X) \):

\[ \text{B(4)}: \text{AEMA: NH}_4\text{OAc} = 1:1:1:1.5. \]

b) \( ^1\text{H} \) NMR spectra (CDCl\(_3\), 400M) of \( P(X)(4) \).
Figure S6. a) Reaction conditions: acetonitrile/dioxane (1/1, v/v) as the solvent, glycine (10% to aldehyde) as the Hantzsch reaction catalyst, ABVN (2% to AEMA) as the initiator for FRP, 75 °C, and 12 h. A(X): B(5): AEMA: NH₄OAc = 1:1:1.5. b) ¹H NMR spectra (CDCl₃, 400M) of P(X)(5).
Figure S7. a) Preparations of P(1)(1) from the traditional monomer–polymer method and one-pot strategy.

$^1$H NMR spectra (CDCl$_3$, 400M) of b) M(1)(1), c) “pure” P(1)(1), and d) P(1)(1) via the one-pot Hantzsch–FRP method.
Figure S8. a) Preparations of P(2)(1) from the traditional monomer–polymer method and one-pot strategy.

$^1$H NMR spectra (CDCl$_3$, 400M) of b) M(2)(1), c) “pure” P(2)(1), and d) P(2)(1) via the one-pot Hantzsch–FRP method.
Figure S9. a) Preparations of P(3)(1) from the traditional monomer–polymer method and one-pot strategy.

$^1$H NMR spectra (CDCl$_3$, 400M) of b) M(3)(1), c) “pure” P(3)(1), and d) P(3)(1) via the one-pot Hantzsch–FRP method.
**Figure S10.** a) Preparations of P(4)(1) from the traditional monomer–polymer method and one-pot strategy. $^1$H NMR spectra (CDCl$_3$, 400M) of b) M(4)(1), c) “pure” P(4)(1), and d) P(4)(1) via the one-pot Hantzsch–FRP method.
**Figure S11.** a) Preparations of P(5)(1) from the traditional monomer–polymer method and one-pot strategy. $^1$H NMR spectra (CDCl₃, 400M) of b) M(5)(1), c) “pure” P(5)(1), and d) P(5)(1) via the one-pot Hantzsch–FRP method.
Figure S12. a) Preparations of P(6)(1) from the traditional monomer–polymer method and the one-pot strategy. $^1$H NMR spectra of (CDCl$_3$, 400M) of b) M(6)(1), c) “pure” P(6)(1), and d) P(6)(1) via a one-pot Hantzsch–FRP system.
Figure S13. LSCM images of the bacteria on sample disks, three replicate samples were tested, commodity polymers served as the controls. a) Commodity polymers + P(4)(1). b) Commodity polymers + P(4)(3). c) Commodity polymers + P(4)(4).
Figure S14. a) Typical samples of P(4)(4)-polymers. b1–5) Stress–strain curves of commodity polymers (black) and P(4)(4)-polymer blends (red). c) Tensile strength of commodity polymers and P(4)(4)-polymer blends. Data are represented as mean ± SD, n = 6.
Figure S15. Images of the plate-streaking experiment: 100 μL sample on LB agar (diameter: 90 mm, thickness: 6 mm, LB: 1.5 g/L), a) 48-h culture, and b) 72-h culture.

Figure S16. OD600 values versus time in the presence of different commercially available beverages. Data are represented as mean ± SD, n = 5.
Figure S17. LSCM images of *E. coli* and *S. aureus* on sample disks after 72-h incubation. PE served as the control. Each image is 200 × 200 μm.

Figure S18. $^1$H NMR spectrum (CDCl$_3$, 400M) of P1.
Figure S19. $^1$H NMR spectrum (CDCl$_3$, 400M) of P2.

Figure S20. $^1$H NMR spectrum (DMSO-$d_6$, 400M) of P(AEMA).
Figure S21. LSCM images of *E. coli* and *S. aureus* on sample disks after 72 h incubation. PE served as the control; each image is $200 \times 200 \ \mu m$. 

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**Table S1.** The Hantzsch-type polymers (P(X)(Y)).

| Polymer | Conv. (vinyl)<sup>a</sup> | Conv. (Hantzsch)<sup>a</sup> | M<sub>n</sub><sup>b</sup> | PDI<sup>b</sup> | T<sub>g</sub><sup>c</sup>(°C) | T<sub>d</sub><sup>d</sup>(°C) |
|---------|--------------------------|-------------------------------|----------------|----------------|-----------------|-----------------|
| P(1)(1) | 95.0%                    | 97.0%                         | 16000          | 1.64           | 115.48          | 207             |
| P(2)(1) | 98.0%                    | 94.0%                         | 23000          | 2.93           | 88.21           | 252             |
| P(3)(1) | 98.0%                    | 98.0%                         | 22000          | 1.83           | 102.56          | 232             |
| P(4)(1) | 98.0%                    | 98.0%                         | 27000          | 3.20           | 106.82          | 221             |
| P(5)(1) | 99.0%                    | 99.0%                         | 28000          | 6.28           | 111.35          | 227             |
| P(6)(1) | 99.0%                    | 98.0%                         | 21000          | 1.69           | 121.34          | 308             |
| P(1)(2) | 97.0%                    | 95.0%                         | 16000          | 1.39           | 90.65           | 228             |
| P(2)(2) | 97.0%                    | 92.0%                         | 19000          | 1.85           | 88.35           | 263             |
| P(3)(2) | 97.0%                    | 93.0%                         | 23000          | 1.96           | 98.72           | 251             |
| P(4)(2) | 98.0%                    | 94.0%                         | 28000          | 1.35           | 102.86          | 216             |
| P(5)(2) | 98.0%                    | 95.0%                         | 22000          | 3.42           | 105.68          | 226             |
| P(6)(2) | 99.0%                    | 98.0%                         | 19000          | 1.48           | 85.23           | 283             |
| P(1)(3) | 97.0%                    | 93.0%                         | 20000          | 1.48           | 117.24          | 208             |
| P(2)(3) | 96.0%                    | 92.0%                         | 24000          | 1.77           | 93.61           | 256             |
| P(3)(3) | 98.0%                    | 93.0%                         | 23000          | 2.14           | 105.63          | 236             |
| P(4)(3) | 99.0%                    | 94.0%                         | 30000          | 1.37           | 99.97           | 224             |
| P(5)(3) | 99.0%                    | 95.0%                         | 20000          | 3.35           | 108.86          | 267             |
| P(6)(3) | 98.0%                    | 98.0%                         | 22000          | 1.48           | 103.52          | 282             |
| P(1)(4) | 98.0%                    | 93.0%                         | 24000          | 1.26           | 112.65          | 237             |
| P(2)(4) | 98.0%                    | 98.0%                         | 25000          | 1.71           | 73.91           | 292             |
| P(3)(4) | 98.0%                    | 98.0%                         | 26000          | 1.94           | 97.89           | 219             |
| P(4)(4) | 97.0%                    | 98.0%                         | 23000          | 1.83           | 97.12           | 208             |
| P(5)(4) | 99.0%                    | 99.0%                         | 20000          | 2.30           | 103.65          | 268             |
| P(6)(4) | 99.0%                    | 99.0%                         | 26000          | 1.64           | 101.13          | 283             |
| P(1)(5) | 95.0%                    | 97.0%                         | 23000          | 1.47           | 56.57           | 246             |
| P(2)(5) | 96.0%                    | 97.0%                         | 21000          | 1.79           | 84.16           | 250             |
| P(3)(5) | 97.0%                    | 96.0%                         | 24000          | 2.38           | 112.72          | 238             |
| P(4)(5) | 98.0%                    | 99.0%                         | 19000          | 2.07           | 109.56          | 256             |
| P(5)(5) | 98.0%                    | 98.0%                         | 25000          | 3.33           | 99.77           | 259             |
| P(6)(5) | 99.0%                    | 99.0%                         | 21000          | 1.55           | 96.40           | 307             |

a. Calculated by <sup>1</sup>H NMR (CDCl<sub>3</sub>, 400 MHz).
b. Measured by GPC using DMF as the eluent (1 mL/min).
c. Measured by DSC at a scanning rate of 10 °C/min.
d. Temperature at weight-loss ratio of 5%, measured by TGA with a heating rate of 20 °C/min.

Table S2. The fluorescence intensity of *E. coli* and *S. aureus* on the sample disks.

|      | *E. coli* | A.E. (%)<sup>b</sup> (E. coli) | *S. aureus* | A.E. (%)<sup>b</sup> (S. aureus) |
|------|-----------|-------------------------------|-------------|----------------------------------|
| PE   | 3.828±0.653 | 0                             | 4.386±0.583 | 0                                |
| A1   |           |                               |             |                                  |
| B1   | 1.528±0.207 | 60.06                         | 1.034±0.150 | 76.42                            |
| B2   | 2.338±0.344 | 38.92                         | 0.460±0.157 | 89.51                            |
| B3   | 2.207±0.281 | 42.35                         | 2.116±0.271 | 51.76                            |
| B4   | 1.042±0.142 | 72.78                         | 0.004±0.001 | 99.91                            |
| B5   | 0.805±0.103 | 78.97                         | 1.962±0.240 | 55.27                            |
| A2   |           |                               |             |                                  |
| B1   | 0.790±0.111 | 79.36                         | 1.703±0.257 | 61.17                            |
| B2   | 1.397±0.158 | 63.51                         | 3.806±0.519 | 13.22                            |
| B3   | 0.511±0.086 | 86.65                         | 1.721±0.251 | 60.78                            |
| B4   | 0.244±0.036 | 93.63                         | 0.816±0.114 | 80.37                            |
| B5   | 1.211±0.172 | 68.36                         | 1.827±0.238 | 58.34                            |
| A3   |           |                               |             |                                  |
| B1   | 0.248±0.039 | 93.52                         | 1.896±0.250 | 56.77                            |
| B2   | 1.076±0.120 | 71.89                         | 2.283±0.307 | 47.95                            |
| B3   | 1.032±0.130 | 73.04                         | 2.112±0.271 | 51.85                            |
| B4   | 0.532±0.071 | 86.10                         | 2.968±0.392 | 32.33                            |
| B5   | 2.276±0.316 | 40.54                         | 1.421±0.186 | 67.60                            |
| A4   |           |                               |             |                                  |
| B1   | 0.004±0.001 | 99.90                         | 0.004±0.001 | 99.91                            |
| B2   | 0.244±0.036 | 93.63                         | 0.816±0.114 | 80.37                            |
| B3   | 0.097±0.019 | 97.47                         | 0.237±0.032 | 95.60                            |
| B4   | 0.002±0.000 | 99.95                         | 0.004±0.001 | 99.91                            |
| B5   | 1.164±0.162 | 69.59                         | 0.053±0.010 | 98.79                            |
| A5   |           |                               |             |                                  |
| B1   | 0.786±0.119 | 79.47                         | 1.311±0.178 | 70.11                            |
| B2   | 1.818±0.237 | 52.51                         | 1.601±0.227 | 63.50                            |
| B3   | 1.456±0.207 | 61.96                         | 2.407±0.312 | 45.12                            |
| B4   | 0.142±0.024 | 96.29                         | 0.427±0.041 | 90.26                            |
|       | B1        | B2        | B3        | B4        | B5        |
|-------|-----------|-----------|-----------|-----------|-----------|
|       | 1.157±0.158 | 0.664±0.097 | 0.708±0.109 | 0.711±0.103 | 1.728±0.206 |
| A6    | 69.78     | 82.65     | 81.50     | 81.43     | 54.86     |
|       | 1.182±0.157 | 2.968±0.407 | 2.431±0.327 | 2.270±0.307 | 2.622±0.339 |
|       | 73.12     | 32.33     | 44.57     | 48.24     | 40.22     |
|       | 0.004±0.001 | 0.304±0.029 | 2.323±0.347 | 2.130±0.309 | 3.828±0.653 |
|       | 99.90     | 92.06     | 39.32     | 44.36     | 0         |
|       | 0.004±0.001 | 0.693±0.101 | 0.650±0.089 | 1.418±0.517 | 4.386±0.583 |
|       | 99.91     | 84.20     | 85.18     | 67.67     | 0         |
|       | 0.010±0.001 | 0.340±0.059 | 1.417±0.199 | 1.555±0.187 | 4.008±0.527 |
|       | 99.75     | 91.52     | 64.65     | 61.20     | 0         |
|       | 0.003±0.001 | 0.406±0.051 | 1.256±0.189 | 2.080±0.331 | 3.116±0.402 |
|       | 99.90     | 86.97     | 59.69     | 33.25     | 0         |
|       | 0.008±0.001 | 0.599±0.087 | 0.780±0.116 | 2.080±0.287 | 3.469±0.716 |
|       | 99.77     | 82.73     | 77.52     | 40.04     | 0         |
|       | 0.007±0.001 | 0.232±0.039 | 0.662±0.102 | 1.962±0.379 | 3.176±0.394 |
|       | 99.78     | 92.70     | 79.16     | 38.22     | 0         |
|       | 0.001±0.002 | 0.056±0.011 | 0.571±0.071 | 1.481±0.249 | 99.72     |
|       | 99.72     | 98.59     | 85.66     | 62.82     | 0         |
|       | 0.009±0.001 | 0.090±0.010 | 0.424±0.057 | 1.681±0.210 | 47.30     |

a. 37°C, 72-h culture.
b. The antibacterial efficiency (A.E.) of polymers, PE served as the control.

Table S3. The fluorescence intensity of the bacterial on sample disksa.

|       | %b | E.coli | A.E.(%)c (E. coli) | S.aureus | A.E.(%)c (S. aureus) |
|-------|----|--------|-------------------|----------|---------------------|
| PE    | 33 | 0.004±0.001 | 99.90          | 0.004±0.001 | 99.91               |
|       | 20 | 0.304±0.029 | 92.06          | 0.693±0.101 | 84.20               |
|       | 10 | 2.323±0.347 | 39.32          | 0.650±0.089 | 85.18               |
|       |  5 | 2.130±0.309 | 44.36          | 1.418±0.517 | 67.67               |
|       |  0 | 3.828±0.653 | 0              | 4.386±0.583 | 0                   |
| PP    | 33 | 0.001±0.001 | 99.75          | 0.003±0.001 | 99.90               |
|       | 20 | 0.340±0.059 | 91.52          | 0.406±0.051 | 86.97               |
|       | 10 | 1.417±0.199 | 64.65          | 1.256±0.189 | 59.69               |
|       |  5 | 1.555±0.187 | 61.20          | 2.080±0.331 | 33.25               |
|       |  0 | 4.008±0.527 | 0              | 3.116±0.402 | 0                   |
| P(4)(1)| 33 | 0.008±0.001 | 99.77          | 0.007±0.001 | 99.78               |
|       | 20 | 0.599±0.087 | 82.73          | 0.232±0.039 | 92.70               |
|       | 10 | 0.780±0.116 | 77.52          | 0.662±0.102 | 79.16               |
|       |  5 | 2.080±0.287 | 40.04          | 1.962±0.379 | 38.22               |
|       |  0 | 3.469±0.716 | 0              | 3.176±0.394 | 0                   |
| PET   | 33 | 0.011±0.002 | 99.72          | 0.009±0.001 | 99.72               |
|       | 20 | 0.056±0.011 | 98.59          | 0.090±0.010 | 97.18               |
|       | 10 | 0.571±0.071 | 85.66          | 0.424±0.057 | 86.71               |
|       |  5 | 1.481±0.249 | 62.82          | 1.681±0.210 | 47.30               |
|          | 0   | 3.983±0.497 | 0   | 3.190±0.422 | 0   |
|----------|-----|-------------|-----|-------------|-----|
| **PA-66**| 33  | 0.009±0.001 | 99.78| 0.010±0.002 | 99.75|
   | 20  | 0.840±0.146 | 79.43| 0.159±0.029 | 96.03|
   | 10  | 1.061±0.267 | 74.02| 0.746±0.175 | 81.37|
   | 5   | 1.654±0.257 | 59.50| 1.598±0.264 | 60.09|
   | 0   | 4.084±0.622 | 0   | 4.004±0.517 | 0   |
| **PE**  | 33  | 0.002±0.001 | 99.95| 0.004±0.001 | 99.91|
   | 20  | 0.050±0.008 | 98.69| 0.254±0.048 | 94.21|
   | 10  | 0.330±0.069 | 91.38| 0.616±0.089 | 85.96|
   | 5   | 1.601±0.248 | 58.18| 2.010±0.421 | 54.17|
   | 0   | 3.828±0.653 | 0   | 4.386±0.583 | 0   |
| **PP**  | 33  | 0.007±0.001 | 99.83| 0.003±0.001 | 99.90|
   | 20  | 0.316±0.041 | 92.12| 0.267±0.037 | 91.43|
   | 10  | 1.589±0.308 | 60.35| 1.425±0.245 | 54.27|
   | 5   | 1.973±0.307 | 50.77| 1.737±0.207 | 44.26|
   | 0   | 4.008±0.527 | 0   | 3.116±0.402 | 0   |
| **P(4)(3)**| 33 | 0.015±0.003 | 99.57| 0.007±0.002 | 99.78|
   | 20  | 0.221±0.037 | 93.63| 0.242±0.037 | 92.38|
   | 10  | 1.112±0.200 | 67.94| 1.110±0.145 | 65.05|
   | 5   | 1.549±0.194 | 55.35| 1.447±0.206 | 54.44|
   | 0   | 3.469±0.716 | 0   | 3.176±0.394 | 0   |
| **PET** | 33  | 0.007±0.002 | 99.82| 0.004±0.001 | 99.87|
   | 20  | 0.126±0.027 | 96.84| 0.254±0.038 | 92.04|
   | 10  | 0.362±0.042 | 90.91| 0.426±0.057 | 86.65|
   | 5   | 1.438±0.037 | 63.90| 1.419±0.029 | 55.52|
   | 0   | 3.983±0.497 | 0   | 3.190±0.422 | 0   |
| **PMMA**| 33  | 0.018±0.004 | 99.56| 0.014±0.003 | 99.65|
   | 20  | 0.183±0.028 | 95.52| 0.155±0.028 | 96.13|
   | 10  | 1.732±0.207 | 57.59| 0.767±0.107 | 80.84|
   | 5   | 1.708±0.216 | 58.18| 1.115±0.270 | 72.15|

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|    |    | P(4)(4) | PE | PP | PET | PMMA | PA-66 |
|----|----|---------|----|----|-----|------|-------|
| 0  | 4.084±0.622 | 0 | 4.004±0.517 | 0 |    |      |       |
| 33 | 0.004±0.001 | 99.90 | 0.013±0.005 | 99.70 |    |      |       |
| 20 | 0.340±0.049 | 91.12 | 0.211±0.049 | 95.19 |    |      |       |
| 10 | 0.491±0.097 | 87.17 | 0.695±0.109 | 84.15 |    |      |       |
| 5  | 0.553±0.105 | 85.55 | 1.291±0.208 | 70.57 |    |      |       |
| 0  | 3.828±0.653 | 0 | 4.386±0.583 | 0 |    |      |       |
| 33 | 0.018±0.004 | 99.55 | 0.028±0.009 | 99.10 |    |      |       |
| 20 | 0.157±0.022 | 96.08 | 0.216±0.047 | 93.07 |    |      |       |
| 10 | 0.244±0.039 | 93.91 | 0.308±0.044 | 90.12 |    |      |       |
| 5  | 0.829±0.100 | 79.32 | 1.122±0.112 | 63.99 |    |      |       |
| 0  | 4.008±0.527 | 0 | 3.116±0.402 | 0 |    |      |       |
| 33 | 0.019±0.002 | 99.45 | 0.005±0.002 | 99.84 |    |      |       |
| 20 | 0.083±0.014 | 97.61 | 0.233±0.057 | 92.66 |    |      |       |
| 10 | 0.303±0.049 | 91.27 | 0.671±0.094 | 78.87 |    |      |       |
| 5  | 1.481±0.027 | 57.31 | 1.462±0.028 | 53.97 |    |      |       |
| 0  | 3.469±0.716 | 0 | 3.176±0.394 | 0 |    |      |       |
| 33 | 0.007±0.001 | 99.82 | 0.012±0.003 | 99.62 |    |      |       |
| 20 | 0.004±0.001 | 99.90 | 0.077±0.022 | 97.59 |    |      |       |
| 10 | 0.193±0.038 | 95.15 | 0.194±0.029 | 93.92 |    |      |       |
| 5  | 0.694±0.103 | 82.58 | 0.390±0.062 | 87.77 |    |      |       |
| 0  | 3.983±0.497 | 0 | 3.190±0.422 | 0 |    |      |       |
| 33 | 0.009±0.001 | 99.78 | 0.007±0.001 | 99.83 |    |      |       |
| 20 | 0.095±0.017 | 97.67 | 0.299±0.042 | 92.53 |    |      |       |
| 10 | 0.313±0.041 | 92.34 | 0.542±0.071 | 86.46 |    |      |       |
| 5  | 1.250±0.203 | 69.39 | 1.055±0.151 | 73.65 |    |      |       |
| 0  | 4.084±0.622 | 0 | 4.004±0.517 | 0 |    |      |       |

a. 37°C, 72 h culture.

b. The ratio of selected polymers (P(4)(1,3,4)) added in several typical commodity polymers (PE, PP, PET, PMMA, PA66).

c. The antibacterial efficiency (A.E.) of polymers. Commodity polymers served as the controls.
Reference:

1) Nickoloff, J. A., *Electroporation protocols for microorganisms*. Humana Press: Totowa, N.J., 1995; p xix, 372 p.