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Novel 1,2,3-triazolium-functionalized inulin derivatives: synthesis, free radical-scavenging activity, and antifungal activity

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A new class of inulin derivatives possessing 1,2,3-triazolium charged units by associating “click reaction” with efficient 1,2,3-triazole quaternization were designed and synthesized. After the chemical modification, all the inulin derivatives showed good solubility in water, and exhibited excellent bioactivity compared to inulin. In the 1,1-diphenyl-2-picrylhydrazyl (DPPH) radical-scavenging assay, the scavenging activity of 1,2,3-triazolium inulin derivatives (**5a–5d**) (IC_{50} 0.21–0.29 mg mL^{−1}) was found to be higher than that of inulin derivatives (**4a–4d**) (IC_{50} 0.58–0.95 mg mL^{−1}) and inulin. **5a–5d** (IC_{50} < 0.01 mg mL^{−1}) also exhibited stronger inhibitory activity against superoxide anion formation than **4a–4d** (IC_{50} 0.02–0.09 mg mL^{−1}) and Vitamin C (IC_{50} 0.02 mg mL^{−1}). In the antifungal assays, **5a–5d** also inhibited the growth of tested phytopathogens more effectively with inhibitory indices of 43.10–82.56% at 1.0 mg mL^{−1}, compared with **4a–4d** and inulin. Based on these data, it is reasonable to propose that the introduction of 1,2,3-triazolium group through chemical modification is significant to enhance the free radical-scavenging activity and antifungal activity of inulin.

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1. Introduction

Inulin is a natural, biodegradable, and plant-derived storage polysaccharide, and has attracted broad attention due to its beneficial nutritional attributes for human health. Inulin also has advantages of non-toxicity, biodegradability, and biocompatibility, facilitating its applications in the food, feed, biofuel, water purification, and pharmaceutical industries. However, application of inulin has been limited due to its low bioactivity. Because the biological activity of a polysaccharide is related with its molecular structure, increasing attention has been paid to chemical modification of polysaccharide structure to improve its biological activity.^{1–7} To promote the antifungal activity of inulin, many inulin derivatives have been prepared such as *O*-(aminoethyl)inulin and 6-amino-6-deoxyinulin by chemical modification. Our previous publications also reported the improved antioxidant activities and antifungal activities of inulin derivatives after chemical modification.^{8–10}

Triazole derivatives represent an interesting class of heterocyclic compounds; they possess many biological activities such as antimicrobial, anti-tubercular, anti-inflammatory and

anticancer activities.^{11–15} These observations inspired us to modify inulin with thiazolyl as a substituent *via* “click reaction” to improve its bioactivity. The Huisgen 1,3-dipolar [3 + 2] cycloaddition of azides and alkynes has emerged as the front-runner of “click reaction”, and this synthetic approach is highly efficient, convenient and the reaction conditions are mild. For example, Ren *et al.* prepared a novel amphiphilic aminated inulin derivative *via* “click reaction”, and this inulin derivative owned stronger antibacterial property against *S. aureus* than inulin.⁸ Furthermore, 1,2,3-triazole is capable of either losing a proton (acting as a weak acid) or accepting a proton through the nitrogen lone pair of electrons (acting as a weak Brønsted base),^{16,17} which might be conducive to scavenging free radicals.

1,2,3-Triazolium cations have recently been developed by quaternization of 1,2,3-triazoles with halogenide. The quaternization of 1,2,3-triazoles could help improve solubility. Meanwhile, the positively charged moiety of the cationic molecules could react with the negatively charged components of fungi and bacteria, which is favorable for the biological activity of the inulin compound.^{17,18} However, there are very few reports on synthesis of inulin derivatives bearing 1,2,3-triazolium cations, and the effect of quaternization of 1,2,3-triazole moieties on the bioactivity of inulin derivative is still unknown.

On the basis of these observations, it was thought of synthesizing a new class of inulin derivatives, wherein 1,2,3-triazolium was linked to inulin as substituent. Application of “click reaction” allows us to prepare various inulin based materials *via* a one-step reaction from azido-inulin and

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terminal alkynes with different substituent groups. And the inulin derivative bearing 1,2,3-triazolium will be obtained by alkylation of 1,2,3-triazolyl inulin derivative. Through the above chemical modifications of inulin, the aimed products would have hydrophilic group (triazole or/and quaternary ammonium salt) and lipophilic group (acetyl), which enabled the target product to have a good surface activity. At the same time triazole is a biologically active group, which could be conducive to improving inulin biological activity. We therefore speculated that the aimed products might have improved biological activity, such as better antioxidant activity and stronger antifungal activity. The free radical-scavenging and antifungal activity of inulin and the synthesized inulin derivatives will be investigated to verify our design of structures.

2. Experimental

2.1. Materials

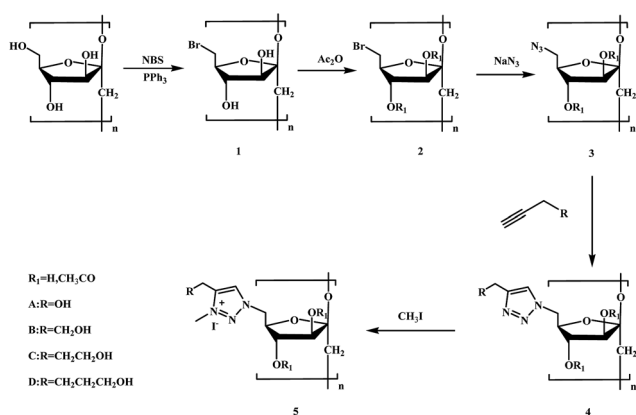
Inulin was purchased from E. Merck (Darmstadt, Germany). Its average degree of polymerization is around 20 fructosyl fructose. 3-Pyridinecarboxaldehyde was purchased from Aladdin Chemical Corp (Shanghai, China). Terminal alkynes (propargyl alcohol, 3-butyn-1-ol, 4-pentyn-1-ol, and 5-hexyn-1-ol) were purchased from Sigma-Aldrich (Shanghai, China) with a minimum purity of 98%. The other reagents such as iodomethane, sodium iodide, sodium hydroxide, cuprous iodide, potassium iodide, and solvents are analytical grade and were supplied by Sinopharm Chemical Reagent Co., Ltd. (Shanghai, China).

2.2. Analytical methods

FT-IR spectra were measured on a Jasco-4100 Fourier Transform Infrared Spectroscopy (JASCO Co., Ltd., Shanghai, China) with KBr disks. ^1H Nuclear Magnetic Resonance (^1H NMR) was measured with a Bruker AVIII-850 Spectroscopy with TCI Cryo-Probe (Bruker Tech. and Serv. Co., Ltd., Beijing, China).

2.3. The synthesis of inulin derivatives

The synthetic routes for the preparation of inulin derivatives are shown in Scheme 1.



Scheme 1 Synthetic routes for the preparation of inulin derivatives.

Hydrophobic 6-bromo-6-deoxyinulin (2) and hydrophobic 6-azido-6-deoxyinulin (3) were prepared according to the methods reported by Ren.⁸

Hydrophobic 6-bromo-6-deoxyinulin (2). ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 2.02 ppm ($\text{CH}_3\text{-C=O}$), δ 3.59–5.33 ppm (pyranose rings), 3.95 ppm (CH_2Br); FT-IR (thin film): ν 1743 (C=O of acetyl).

Hydrophobic 6-azido-6-deoxyinulin (3). Yields: 68.8%; ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 2.01 ppm ($\text{CH}_3\text{-C=O}$), δ 3.34–5.26 ppm (pyranose rings), 4.0 ppm (CH_2N_3); FT-IR (thin film): ν 2105 (C-6-azido), ν 1743 (C=O of acetyl).

Preparation of compound 4a–4d. A mixture of compound 3 (1.0 mmol), 12 mg cuprous iodide, terminal alkynes (2.0 mmol), and 3 mL triethylamine in 20 mL DMSO was stirred at 75°C for 48 h under argon. The mixture was filtered and the filtrate was collected and precipitated into ethanol. The precipitate was washed with saturated solution of potassium iodide and filtered. The unreacted alkyne was extracted in a Soxhlet apparatus with ethanol for two days. The products were dried at 60°C for 24 h.

4a. Yield: 62.1%; ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 1.99 ppm ($\text{CH}_3\text{-C=O}$), δ 7.83 ppm (triazole-5-H); FT-IR (thin film): ν 3409 (stretching vibration of OH), ν 1735 (C=O of acetyl), ν 786 (C-H of triazole).

4b. Yield: 82.5%; ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 1.99 ppm ($\text{CH}_3\text{-C=O}$), δ 2.77 ppm ($\text{OH-CH}_2\text{-CH}_2$), δ 7.77 ppm (triazole-5-H); FT-IR (thin film): ν 3424 (stretching vibration of OH), ν 1743 (C=O of acetyl), ν 790 (C-H of triazole).

4c. Yield: 85.0%; ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 1.72 ppm ($\text{OH-CH}_2\text{-CH}_2\text{-CH}_2$), δ 1.99 ppm ($\text{CH}_3\text{-C=O}$), δ 2.63 ppm ($\text{OH-CH}_2\text{-CH}_2$), 7.77 ppm (triazole-5-H); FT-IR (thin film): ν 3421 (stretching vibration of OH), ν 2942 (C-H of methylene), ν 1743 (C=O of acetyl), ν 794 (C-H of triazole).

4d. Yield: 80.3%; ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 1.42 ppm ($\text{OH-CH}_2\text{-CH}_2\text{-CH}_2\text{-CH}_2$), δ 1.61 ppm ($\text{OH-CH}_2\text{-CH}_2\text{-CH}_2$), δ 1.99 ppm ($\text{CH}_3\text{-C=O}$), δ 2.35 ppm ($\text{OH-CH}_2\text{-CH}_2$), δ 7.72 ppm (triazole-5-H); FT-IR (thin film): ν 3424 (stretching vibration of OH), ν 1743 (C=O of acetyl), ν 798 (C-H of triazole).

Inulin derivative bearing 1,2,3-triazolium (5) were prepared according to the methods reported by Tan.¹⁹ A solution of 1,2,3-triazole-functionalized inulin derivative (3) (1 mmol) and iodomethane (0.187 mL, 3 mmol) in 15 mL of DMSO was stirred at 60°C for 24 h. Afterwards, the remaining iodomethane was evaporated, and the reaction mixture was precipitated into 100 mL of acetone. The solid product was filtered, washed with acetone three times. After being dialyzed against deionized water for 48 h, the inulin derivative (5) was obtained by lyophilization of the aqueous solution.

5a. Yield: 98.5%; ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 2.0 ppm ($\text{CH}_3\text{-C=O}$), δ 4.35 ppm (N^+CH_3), 8.31 ppm (triazole-5-H); FT-IR (thin film): ν 3417 (stretching vibration of OH), ν 2956 and 2881 (stretching vibration of C-H in methyl), ν 1739 (C=O of acetyl), ν 790 (C-H of triazole).

5b. Yield: 94.1%; ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 2.0 ppm ($\text{CH}_3\text{-C=O}$), δ 2.76 ppm ($\text{OH-CH}_2\text{-CH}_2$), δ 4.35 ppm (N^+CH_3); δ 8.27 ppm (triazole-5-H); FT-IR (thin film): ν 3417 (stretching



vibration of OH), ν 2962 and 2877 (stretching vibration of C–H in methyl), ν 1743 (C=O of acetyl), ν 798 (C–H of triazole).

5c. Yield: 73.5%; ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 1.84 ppm (OH–CH₂–CH₂–CH₂), δ 2.04 ppm (CH₃–C=O), δ 2.96 ppm (OH–CH₂–CH₂), δ 4.23 ppm (N⁺CH₃), δ 8.29 ppm (triazole-5-H); FT-IR (thin film): ν 3409 (stretching vibration of OH), ν 2962 and 2881 (stretching vibration of C–H in methyl), ν 1743 (C=O of acetyl), ν 790 (C–H of triazole).

5d. Yield: 75.8%; ^1H NMR (500 MHz, $\text{DMSO}-d_6$): δ 1.55 ppm (OH–CH₂–CH₂–CH₂–CH₂), δ 1.73 ppm (OH–CH₂–CH₂–CH₂), δ 2.0 ppm (CH₃–C=O), δ 2.88 ppm (OH–CH₂–CH₂), δ 4.23 ppm (N⁺CH₃), δ 8.26 ppm (triazole-5-H); FT-IR (thin film): ν 3424 (stretching vibration of OH), ν 2958 and 2881 (stretching vibration of C–H in methyl), ν 1743 (C=O of acetyl), ν 802 (C–H of triazole).

2.4. Radical scavenging ability assay

DPPH-radical scavenging ability assay. The DPPH scavenging properties of the products were evaluated by the following method:¹ the test compound and 2 mL DPPH solution (180 $\mu\text{mol L}^{-1}$) in ethanol was incubated in a water bath at 25 °C for 30 min. Then, the absorbance of the remained DPPH radical was measured at 517 nm against a blank. Three replicates for each sample were tested and the scavenging effect was calculated using the following equation:

$$\text{Scavenging effect (\%)} = \left[1 - \frac{A_{\text{sample 517 nm}} - A_{\text{control 517 nm}}}{A_{\text{blank 517 nm}}} \right] \times 100\%$$

where $A_{\text{control 517 nm}}$ is the absorbance of the control (ethanol instead of DPPH for each concentration) and $A_{\text{blank 517 nm}}$ is the absorbance of the blank (distilled water instead of the sample). The antioxidant activity of test compound was expressed as IC_{50} , which was defined as the concentrations of test compound required for inhibition of the formation of DPPH radical by 50%. Vitamin C was used as a positive control.

Superoxide-radical scavenging ability assay. The superoxide radical scavenging ability was assessed following Xing's method with minor modification.⁹ Involving the test compound, 30 M phenazine mothosulfate (PMS), 338 M nicotinamide adenine dinucleotide reduced (NADH), and 72 M nitro blue tetrazolium (NBT) in Tris–HCl buffer (16 mM, pH 8.0), the reaction mixture was incubated at 25 °C for 5 min. The absorbance was read at 560 nm against blank. Three replicates for each sample were tested and the capability of scavenging superoxide radical was calculated using the following equation:

$$\text{Scavenging effect (\%)} = \left[1 - \frac{A_{\text{sample 560 nm}} - A_{\text{control 560 nm}}}{A_{\text{blank 560 nm}}} \right] \times 100\%$$

where $A_{\text{control 560 nm}}$ is the absorbance of the negative control (distilled water instead of NADH for each concentration) and $A_{\text{blank 560 nm}}$ is the absorbance of the blank (distilled water instead of the samples). The superoxide radical-scavenging activity was expressed as the IC_{50} value. Vitamin C was used as a positive control.

Hydroxyl-radical scavenging ability assay. The test of hydroxyl-radical scavenging ability was carried out according to Liu's method with minor modification.²⁰ The reaction mixture, a total volume of 4.5 mL, containing the samples of inulin, **4a–4d** or **5a–5d** (0.045, 0.09, 0.18, 0.36, and 0.72 mL), were incubated with EDTA–Fe²⁺ (220 μM), safranine O (0.23 μM), and H₂O₂ (60 μM) in potassium phosphate buffer (150 mM, pH 7.4) for 30 min at 37 °C. The absorbance of the mixture was measured at 520 nm. In the blank, samples were substituted with distilled water. Meanwhile, in the negative control, H₂O₂ was substituted with potassium phosphate buffer. Three replicates for each sample were tested. The capability of scavenging hydroxyl radicals was computed using the following equation:

$$\text{Scavenging effect (\%)} = \frac{A_{\text{sample 520 nm}} - A_{\text{blank 520 nm}}}{A_{\text{control 520 nm}} - A_{\text{blank 520 nm}}} \times 100\%$$

where $A_{\text{blank 520 nm}}$ is the absorbance of the blank at 520 nm; $A_{\text{sample 520 nm}}$ is the absorbance of the sample at 520 nm; $A_{\text{control 520 nm}}$ is the absorbance of the control at 520 nm. The antioxidant activity of test compounds was expressed as IC_{50} . Vitamin C was used as a positive control.

2.5. Antifungal assays

Antifungal assays were performed by following the plate growth rate method described by Guo *et al.*²⁸ Briefly, the compounds (inulin, **4a–4d**, and **5a–5d**) were dispersed in distilled water at 5.0 mg mL^{−1}. Then, each solution was added to sterilized potato dextrose agar to give a final concentration of 0.1, 0.5, and 1.0 mg mL^{−1}. After the mixture was cooled in the plate (6.0 cm diameter), 5.0 mm diameter of fungi mycelium was transferred to the test plate and incubated at 27 °C for 2–3 days. When fungi mycelium reached the edges of control plate (without the presence of samples), the antifungal index was calculated as follows:

$$\text{Antifungal index (\%)} = (1 - D_a/D_b) \times 100$$

where D_a is the diameter of growth zone in test plate and D_b is the diameter of the growth zone in control plate. Each experiment was performed in three replicates, and the data were shown as mean \pm SD.²⁹

3. Results and discussion

3.1. Chemical syntheses and characterization

Each step of synthesis was followed by FT-IR and ^1H NMR spectroscopy measurements. The FT-IR and ^1H NMR spectra of intermediate products, compound **4a–4d**, and **5a–5d** are shown in Fig. 1 and 2 respectively.

C-6-Br inulin derivative was synthesized firstly by the reaction between the primary hydroxyls with NBS (Scheme 1), as NBS and Ph₃P could selectively replace primary hydroxyl groups of polysaccharide with bromine.⁸ Then the secondary hydroxyls of the C-6-Br inulin derivative were protected by acetic anhydride to bring hydrophobic groups into inulin backbone (**2**). The azidation of compound **2** could be conveniently achieved through a nucleophilic substitution at C-6 with sodium azide to get C-6-azido inulin derivative (**3**). The introduction of hydrophobic groups



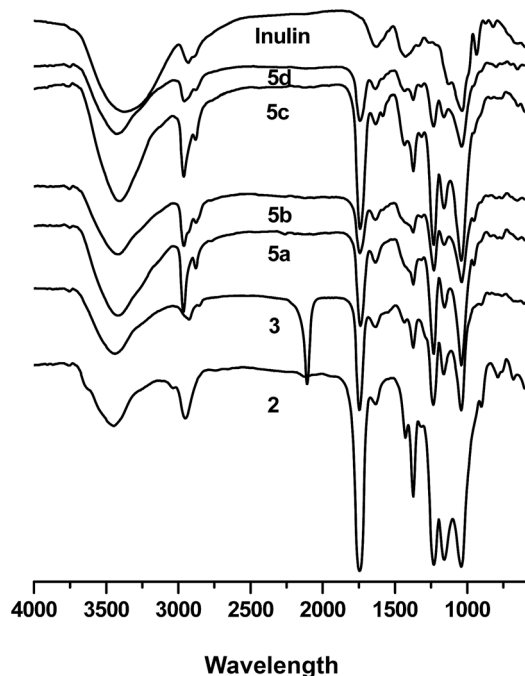


Fig. 1 FT-IR spectra of intermediate products and inulin derivatives 5a–5d.

could effectively reduce the water solubility of inulin derivative (2 and 3), which facilitated the removal of inorganic salt and acetic anhydride by simply washing the products with water. In the spectrum of compound 3, characteristic peak of C-6-azido is observed at 2105 cm^{-1} , and a strong peak at 1743 cm^{-1} was assigned to the vibration of the $\text{C}=\text{O}$ bonds of the acetyl ester (Fig. 1). In the ^1H NMR spectrum of compounds 2 and 3 (Fig. 2), the signals at 3.34–5.33 ppm were assigned to the protons of inulin. New signals, which appear at 2.01 ppm, could be assigned to the protons of $\text{CH}_3-\text{C}=\text{O}$ (Fig. 2).

As long as we got C-6-azido inulin derivative (3), the ‘click chemistry’ could be performed in an elegant way with terminal alkynes to synthesize inulin derivative bearing 1,2,3-triazole. The peak at 2105 cm^{-1} in compound 3 spectrum disappears when the C-6-azido in compound 3 was transformed to 1,2,3-triazoles and a new absorption band at about $794\text{--}789\text{ cm}^{-1}$ appears (Fig. 1). The appearance of the proton in 1,2,3-triazole at 7.72–7.83 ppm further proved the successful ‘click reaction’. Besides, the new chemical shift at 2.77 ppm (in 4b), 1.72 and 2.63 ppm (in 4c), 1.42, 1.61, and 2.35 ppm (in 4d), were assigned to the rest of methylenes.

Subsequently, alkylation of 1,2,3-triazolyl inulin derivative (4) was conducted by reacting with iodomethane in DMSO. The ^1H NMR spectrum of 5 corroborated the quaternization reaction through the disappearance of the peak, corresponding to the proton of the 1,2,3-triazole group (initially present at 7.72–7.83 ppm in $\text{DMSO}-d_6$) and the quantitative appearance of new signals for the 1,2,3-triazolium proton and the N-3 methyl protons of the quaternizing group at 8.26–8.31 ppm and 4.23–4.35 ppm, respectively.²¹

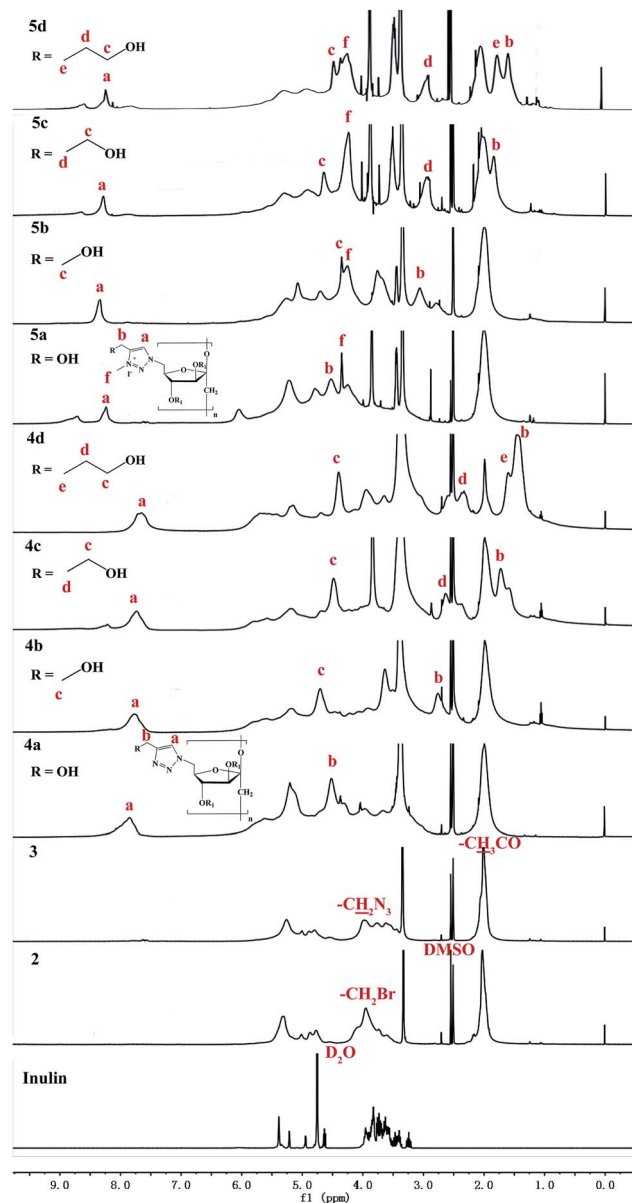


Fig. 2 ^1H NMR spectra of intermediate products and inulin derivatives.

3.2. Solubility of the inulin derivatives

Inulin has good water solubility. As the introduction of acetyl greatly reduced the water solubility of the inulin derivatives, hydrophobic 6-azido-6-deoxyinulin (3) exhibited worse water solubility. After ‘click reaction’ between azido and terminal alkynes, 1,2,3-triazole was formed in inulin backbone. Because of the hydrophilic moiety at the end of the molecular chains, triazole enabled inulin derivative (4) with better water solubility. After the quaternization, the water solubility of inulin derivative (5) was further improved. Therefore, compounds 4 and 5 showed favourable water solubility, and their aqueous solutions could be prepared at $0.1\text{--}1.0\text{ mg mL}^{-1}$ at room temperature.



3.3. Radical scavenging activity

As metabolic products of human body, free radical damage is of great threat to the health.²² Free radical, including hydroxyl radicals and superoxide anion, are normally generated in human body and scavenged by antioxidant defences system when free radical remains at physiological concentrations.²³ Oxidative stress resulting from the toxic effects of free radicals on tissue plays an important role in the pathogenesis of various neurodegenerative diseases such as Alzheimer's disease, Parkinson's disease, and ischemic-reperfusion injury.²⁴ Free radical can also cause pathological damages like cancer disease, diabetes, atherosclerosis, coronary heart disease, and many other diseases associated with aging to the organism, and lead to harmful alterations in foods and pharmaceutical industry.^{20,25} It is of great interest among researchers to develop antioxidant supplements to help the human body reduce oxidative scratch. Moreover, restrictions over the use of synthetic antioxidants butyl hydroxyl anisid (BHA) and butylated hydroxytoluene (BHT) in food further strengthen the concept of using naturally occurring compounds as antioxidants.²⁶ In order to study the effect of those chemical modifications mentioned above on the biological activity of inulin, it is essential to evaluate whether these compounds are able to scavenge the free radicals using different assay systems such as the DPPH radical-scavenging, superoxide radical-scavenging, and hydroxyl radical-scavenging assay.

The DPPH radical scavenging activity of inulin and its related derivatives was tested by their ability to bleach the stable radical DPPH (Fig. 3). This assay provided information on the reactivity of the compounds with a stable free radical. Because of the odd electron, DPPH shows a strong absorption band at 517 nm in visible spectroscopy (deep violet color). As this electron becomes paired off in the presence of a free radical scavenger, the absorption vanishes, and the resulting decolorization is stoichiometric with respect to the number of electrons taken up.²⁴ Vitamin E, a reference drug, exhibited scavenging activity with IC_{50} value of 24.5 μ M (IC_{50} 0.01 mg mL⁻¹).²⁴ As a positive control, Vitamin C was tested with IC_{50} < 0.1 mg mL⁻¹. Inulin showed relatively weak scavenging activity against DPPH radical, and the scavenging index was 19.03% at 1.6 mg mL⁻¹. In this test the synthesized inulin derivatives (4a–4d and 5a–5d)

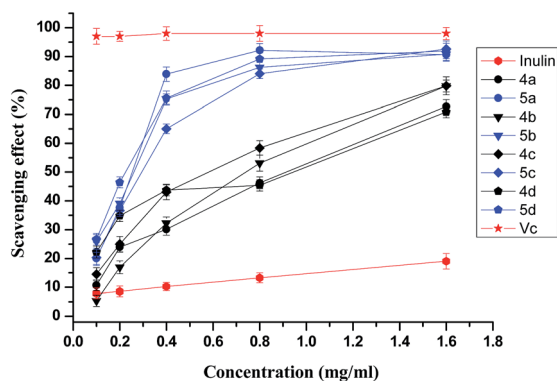


Fig. 3 DPPH-radical scavenging ability of inulin and inulin derivatives.

gave much stronger DPPH-radical scavenging ability compared with inulin. Meanwhile, compounds 5a–5d (IC_{50} 0.21–0.29 mg mL⁻¹) was more efficient than compound 4a–4d (IC_{50} 0.58–0.95 mg mL⁻¹).

The superoxide radical-scavenging activity of inulin and derivatives synthesized was also evaluated based on their ability to quench the superoxide radical generated from the PMS/NADH reaction (Fig. 4). This assay provides information on the reactivity of test compounds with superoxide free radicals, independently of any enzymatic activity. The generation of superoxide anions was markedly inhibited by Vitamin C with an IC_{50} value of 0.02 mg mL⁻¹, and superoxide dismutase (SOD) with an IC_{50} value of 3.5 units per mL.²⁴ Inulin, 4a–4d, and 5a–5d inhibited superoxide anion formation in a concentration-dependent manner, but 5a–5d showed more potent scavenging activity (IC_{50} < 0.01 mg mL⁻¹), followed by 4a–4d (IC_{50} of 4a–4c 0.02–0.09 mg mL⁻¹, IC_{50} of 4d < 0.01 mg mL⁻¹), and inulin (IC_{50} 1.34 mg mL⁻¹), respectively.

Fig. 5 shows the hydroxyl radical-scavenging ability of inulin and synthesized derivatives at various concentrations. Mannitol, a classical hydroxyl radical scavenger, can significantly inhibited deoxyribose degradation with an IC_{50} value of 4.6 mM.²⁴ The hydroxyl-radical scavenging ability of inulin and Vitamin C is weak, and the maximum inhibitory activity in the test concentration range (0.1–1.6 mg mL⁻¹) is 48% and 30% respectively. The hydroxyl-radical scavenging ability of inulin is weak, and the maximum inhibitory activity in the test concentration range (0.1–1.6 mg mL⁻¹) is 48%. The synthesized inulin derivatives (4a–4d and 5a–5d) also show much stronger hydroxyl-radical scavenging ability compared with inulin, and

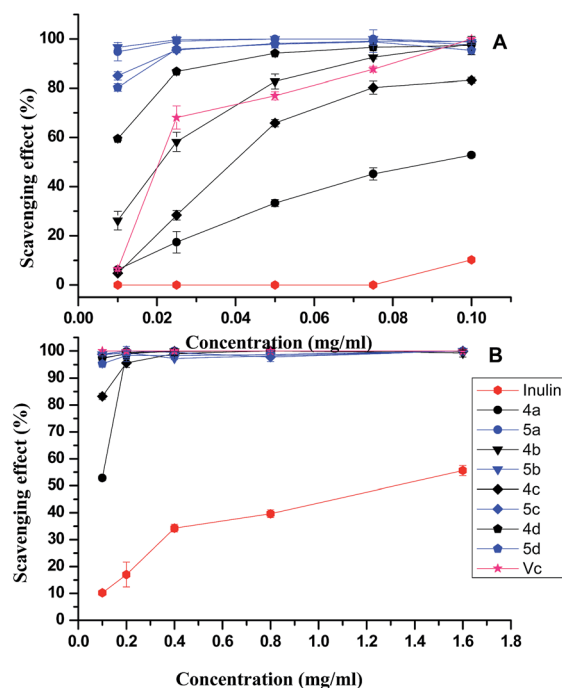


Fig. 4 Superoxide-radical scavenging ability of inulin and inulin derivatives at different concentrations: (A) 0.01–0.1 mg mL⁻¹; (B) 0.1–1.6 mg mL⁻¹.



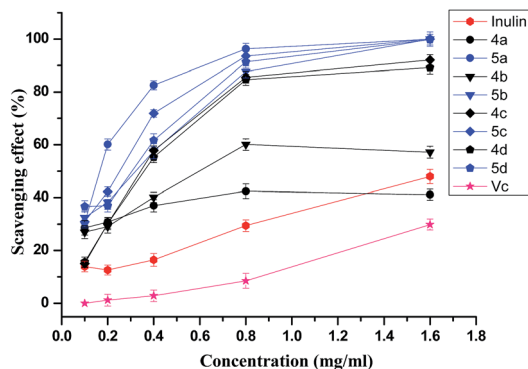


Fig. 5 Hydroxyl-radical scavenging ability of inulin and inulin derivatives.

suppressed hydroxyl-radical in a concentration-dependent manner. The inhibitory activity was observed in the following order: **5a–5d** (IC_{50} 0.16–0.32 mg mL⁻¹) > **4a–4d** (IC_{50} 0.34–0.59 mg mL⁻¹) > inulin.

On the basis of IC_{50} , it is obvious that inulin derivatives with triazole or triazolium show much better antioxidant activity due to the introduction of functional groups—triazole groups. Free radical chain reactions may be inhibited by adding preventive antioxidants that retard the formation of free radicals or stabilize free radicals.^{27,28} Owing to the slightly polarized nature of the C(5)–H bond, 1,2,3-triazoles have gained recognition as excellent hydrogen donors,²⁹ which can form stable free radicals. Meanwhile, the conjugated double bonds allow electron delocalisation across the molecule thus stabilize the radical.²⁶ Furthermore, it is apparent that the inulin derivatives with triazolium group (**5a–5d**) own better hydroxyl-radical scavenging ability. Since the C(5)–H \cdots A⁻ binding ability is strongly enhanced by converting the triazole unit into a triazolium cation, the latter is expected to be a more efficient anion captor,²⁹ which may help stabilize the free radicals form. On the other hand, other factors such as the surface activity affected by the structure of polymers and the different mechanisms involved in the radical antioxidant reactions in different systems may also affect the capacity of test compounds to react with and quench different radicals. The scavenging effect on DPPH radicals and superoxide radicals represents direct radical-scavenging activity. However, in the hydroxyl radical scavenging assay, hydroxyl radicals are generated by the Fenton reaction and the inhibition could be attributed to the inhibition of radicals or the Fe²⁺ chelating effect of the test compounds.²⁴

Many polysaccharides themselves have relatively weak antioxidant activity. The hydroxyl radical scavenging index of starch and hyaluronan was 22% and 40% at 1.6 mg mL⁻¹ respectively,³⁰ which is weaker than that of inulin. Chitosan, one of the most important polysaccharide, has stronger scavenging activity on hydroxyl radicals with IC_{50} 0.48 mg mL⁻¹.³¹ Recently, chemical modification of polysaccharides is increasingly reported for its potential of improving the biological activity of polysaccharides. Fan reported antioxidant activity of silk peptides grafted carboxymethyl chitosan, and the highest scavenging activity of

DPPH was 24.86%, 91% of hydroxyl radical and 36.8% of H₂O₂ at the concentration of 0.5–2.5 mg mL⁻¹.³² Double quaternized chitosan derivatives showed better scavenging ability than chitosan, with more than 90% scavenging indices against hydroxyl radicals and DPPH radicals at 1.6 mg mL⁻¹.³³ Tan synthesized aminopyridinium-grafted starch derivative and found antioxidant property of synthesized starch derivatives was affected by the different positions and the number of amino groups on pyridinium rings.³⁴ The experimental data above and related literature demonstrated that the chemical modification of polysaccharides was conducive to improving the free radical scavenging activity of them.

3.4. Antifungal activity

Plant pathogenic fungi negatively affect a large number of important fruits and vegetables, and limit crop production worldwide, especially in developing countries. For example, the premature plant death caused by *C. lagenarium* can result in a yield loss of 6–48%.³⁵ Control of these plants-threatening fungi could benefit the production of related vegetables, fruits, and crops. In this paper, two common harmful *Gibberella zeae* and *C. lagenarium* were selected to estimate the antifungal activity of inulin and the synthesized inulin derivatives (**4a–4d** and **5a–5d**). The results are shown in Fig. 6 and 7.

As seen in Fig. 6, inulin does not inhibit the growth of *Gibberella zeae* at the tested concentrations. Compared with inulin, after the introduction of triazole groups, compound **4a–4d** have evident antifungal activities against *Gibberella zeae* and the inhibitory indices of compound **4a–4d** are 9.82%, 25.65%, 29.73%, and 20.21% at 1.0 mg mL⁻¹, respectively. Meanwhile, the inhibitory indices of all the samples ascend with the increasing concentration. It is obvious that inulin derivatives containing triazole (**4a–4d**) show much better antifungal activity due to the introduction of functional groups—thiazolyl groups. Furthermore, inulin derivatives containing triazolium (**5a–5d**) exhibit higher inhibitory indices than those of inulin derivatives containing triazoles (**4a–4d**), and the inhibitory indices of compound **4a–4d** are 43.10%, 61.65%, 67.57%, and 82.56% at 1.0 mg mL⁻¹, respectively. These data suggest that stronger antifungal activity of inulin derivative (**5a–5d**) may mainly benefit from the alkylation of 1,2,3-triazole. The cationic triazolium may interact with the anionic components on the cell

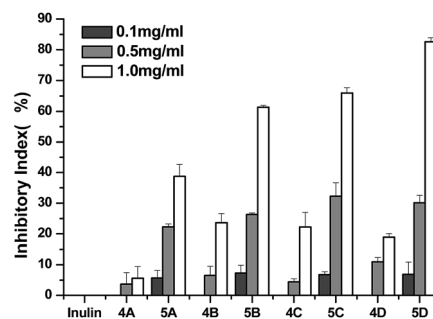


Fig. 6 The antifungal activity of inulin and inulin derivatives against *Gibberella zeae*.



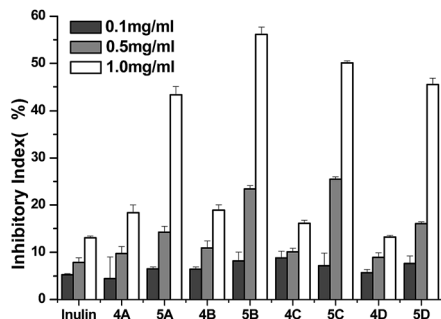


Fig. 7 The antifungal activity of inulin and inulin derivatives against *C. lagenarium*.

wall of fungi, such as glucan, mannan, proteins, and lipids; this leads to the formation of an impervious layer around the cell. The layer could prevent the transportation of essential nutrients, such as glucose, into the cell and may also disturb the cell wall and, thereby, cause severe leakage of the cell constituents and, ultimately, cell death.¹⁸

Fig. 7 shows the antifungal activity of inulin and all the derivatives against *C. lagenarium*. All the samples show antifungal activity against *C. lagenarium*, and the inhibitory indices of them mount up with increasing concentration. The antifungal activity of inulin is weak against *C. lagenarium* and the inhibitory index is 6.0% at 1.0 mg mL⁻¹. The inhibitory indices of inulin derivatives containing triazoles (**4a–4d**) are 20.0%, 20.2%, 16.6%, and 23.2% at 1.0 mg mL⁻¹, respectively. And the inhibitory indices of inulin derivatives containing triazolium (**5a–5d**) are 45.31%, 57.93%, 50.42%, and 46.93% at 1.0 mg mL⁻¹, respectively. Similar to the antifungal activity against *Gibberella zeae*, eight inulin derivatives give much stronger antifungal activity than unmodified inulin, and the inhibitory indices of inulin derivatives containing triazolium (**5a–5d**) are higher than those of inulin derivatives containing triazoles (**4a–4d**) at the same concentration. The results further confirmed that triazolium groups grafted into the synthesized inulin derivatives contributed a lot to the antifungal action and consequently increase the antifungal activity of them. Previous studies had proven that quaternized chitosan had a higher cationic charge than chitosan, which were more prone to interaction with anionic components of the cell wall of fungi. As a result, they would have had stronger antifungal activities.¹⁸

4. Conclusions

In this paper, we have proposed a straightforward synthetic route to a new class of inulin derivatives possessing 1,2,3-triazolium charged units by associating “click reaction” with efficient 1,2,3-triazole quaternization. The radical scavenging activity against three kinds of free radicals and antifungal activity against two kinds of phytopathogens were estimated. All the inulin derivatives showed good solubility in water, and exhibited higher radical scavenging activity and antifungal activity than inulin. Moreover, the synthesized inulin derivative bearing 1,2,3-triazolium showed more significant biological

activity than inulin derivative with 1,2,3-triazole and inulin. The mechanism of the obtained biological activity was also discussed. The 1,2,3-triazolium would be a more efficient anion captor than triazole after quaternization, which might help stabilize the free radicals form. Meanwhile, the 1,2,3-triazolium had a higher cationic charge, which were more prone to interaction with anionic components on fungi cell wall. Further study will be carried out to ascertain this hypothesis. The experiment data demonstrated that the chemical modification of inulin with 1,2,3-triazolium functional groups was conducive to improving the biological activity of inulin. These findings mentioned above bring further evidence that inulin derivatives are active and have the potential of becoming alternatives of free radical scavenger.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- 1 Y. Hu, J. Zhang, C. Yu, Q. Li, F. Dong, G. Wang and Z. Guo, *Int. J. Biol. Macromol.*, 2014, **70**, 44–49.
- 2 W. Tan, J. Zhang, F. Luan, L. Wei, Y. Chen, F. Dong, Q. Li and Z. Guo, *Int. J. Biol. Macromol.*, 2017, **102**, 704–711.
- 3 W. Tan, Q. Li, L. Wei, P. Wang, Z. Gao, Y. Chen, F. Dong and Z. Guo, *Mater. Sci. Eng., C*, 2017, **76**, 1048–1056.
- 4 Q. Li, C. Zhang, W. Tan, G. Gu and Z. Guo, *Molecules*, 2017, **22**, 156–165.
- 5 J. Ahmed, L. Thomas, A. Taher and A. Joseph, *Carbohydr. Polym.*, 2016, **152**, 639–647.
- 6 S. Adak and R. Banerjee, *Carbohydr. Polym.*, 2016, **150**, 359–368.
- 7 L. Fan, J. Yang, H. Wu, Z. Hu, J. Yi, J. Tong and X. Zhu, *Int. J. Biol. Macromol.*, 2015, **79**, 830–836.
- 8 F. Dong, J. Zhang, C. Yu, Q. Li, J. Ren, G. Wang, G. Gu and Z. Guo, *Bioorg. Med. Chem. Lett.*, 2014, **24**, 4590–4593.
- 9 J. Ren, J. Liu, F. Dong and Z. Guo, *Carbohydr. Polym.*, 2011, **83**, 1240–1244.
- 10 J. M. Ren, P. B. Wang, F. Dong, Y. Feng, D. J. Peng and Z. Y. Guo, *Carbohydr. Polym.*, 2012, **87**, 1744–1748.
- 11 X. Ouyang, X. Chen, E. L. Piatnitski, A. S. Kiselyov, H. Y. He, Y. Mao, V. Pattaropong, Y. Yu, K. H. Kim, J. Kincaid, L. Smith 2nd, W. C. Wong, S. P. Lee, D. L. Milligan, A. Malikzay, J. Fleming, J. Gerlak, D. Deevi, J. F. Doody, H. H. Chiang, S. N. Patel, Y. Wang, R. L. Rolser, P. Kussie, M. Labelle and M. C. Tuma, *Bioorg. Med. Chem. Lett.*, 2005, **15**, 5154–5159.



- 12 S. Eswaran, A. V. Adhikari and N. S. Shetty, *Eur. J. Med. Chem.*, 2009, **44**, 4637–4647.
- 13 H. Park, Y. J. Bahn and S. E. Ryu, *Bioorg. Med. Chem. Lett.*, 2009, **19**, 4330–4334.
- 14 Y. P. Hou, J. Sun, Z. H. Pang, P. C. Lv, D. D. Li, L. Yan, H. J. Zhang, E. X. Zheng, J. Zhao and H. L. Zhu, *Bioorg. Med. Chem.*, 2011, **19**, 5948–5954.
- 15 D. Abu-Rahma Gel, M. Abdel-Aziz, E. A. Beshr and T. F. Ali, *Eur. J. Med. Chem.*, 2014, **71**, 185–198.
- 16 L. Liu, S. He, S. Zhang, M. Zhang, M. D. Guiver and N. Li, *ACS Appl. Mater. Interfaces*, 2016, **8**, 4651–4660.
- 17 W. Tan, J. Zhang, F. Luan, L. Wei, Q. Li, F. Dong and Z. Guo, *Int. J. Biol. Macromol.*, 2017, **101**, 845–851.
- 18 J. Ren, J. Liu, R. Li, F. Dong and Z. Guo, *J. Appl. Polym. Sci.*, 2012, **124**, 2501–2507.
- 19 W. Tan, Q. Li, Z. Gao, S. Qiu, F. Dong and Z. Guo, *Carbohydr. Polym.*, 2017, **157**, 236–243.
- 20 J. Liu, H. Sun, F. Dong, Q. Xue, G. Wang, S. Qin and Z. Guo, *Carbohydr. Polym.*, 2009, **78**, 439–443.
- 21 R. Sood, M. M. Obadia, B. P. Mudraboyina, B. Zhang, A. Serghei, J. Bernard and E. Drockenmuller, *Polymer*, 2014, **55**, 3314–3319.
- 22 Z. Peng, M. Liu, Z. Fang and Q. Zhang, *Int. J. Biol. Macromol.*, 2012, **50**, 1254–1259.
- 23 G.-H. Wu, T. Hu, Z.-Y. Li, Z.-L. Huang and J.-G. Jiang, *Food Chem.*, 2014, **148**, 351–356.
- 24 P. Mahakunakorn, M. Tohda, Y. Murakami, K. Matsumoto and H. Watanabe, *Biol. Pharm. Bull.*, 2004, **27**, 38–46.
- 25 D. Wei, W. Cheng, Y. Wei and L. Zhang, *Glycoconjugate J.*, 2012, **29**, 167–172.
- 26 M. A. Soobrattee, V. S. Neergheen, A. Luximon-Ramma, O. I. Aruoma and T. Bahorun, *Mutat. Res., Fundam. Mol. Mech. Mutagen.*, 2005, **579**, 200–213.
- 27 P.-J. Park, W.-K. Jung, K.-S. Nam, F. Shahidi and S.-K. Kim, *J. Am. Oil Chem. Soc.*, 2001, **78**, 651–656.
- 28 E. S. H. El Ashry, E. El-Rafey, N. Rezki, H. H. Abou-Elnaga, W. M. A. Bakry and Y. M. Boghdadi, *J. Saudi Chem. Soc.*, 2014, **18**, 443–449.
- 29 J. M. Aizpurua, R. M. Fratila, Z. Monasterio, N. Perez-Esnaola, E. Andreieff, A. Irastorza and M. Sagartzazu-Aizpurua, *New J. Chem.*, 2014, **38**, 474–480.
- 30 S. Yang, Z. Guo, F. Miao, Q. Xue and S. Qin, *Carbohydr. Polym.*, 2010, **82**, 1043–1045.
- 31 Z. Guo, H. Liu, X. Chen, X. Ji and P. Li, *Bioorg. Med. Chem. Lett.*, 2006, **16**, 6348–6350.
- 32 Y. A. Skorik, A. S. Kritchenkov, Y. E. Moskalenko, A. A. Golyshev, S. V. Raik, A. K. Whaley, L. V. Vasina and D. L. Sonin, *Carbohydr. Polym.*, 2017, **166**, 166–172.
- 33 L. Wei, Q. Li, W. Tan, F. Dong, F. Luan and Z. Guo, *Molecules*, 2017, **22**, 501.
- 34 Z. G. W. Tan, Q. Chen, F. Dong and Q. Li, *Starch*, 2017, **69**, 1–8.
- 35 Y. Chen and G. Dai, *J. Sci. Food Agric.*, 2012, **92**, 1937–1943.

