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Synthesis, characterization, and functional evaluation of branched dodecyl phenol polyoxyethylene ethers: a novel class of surfactants with excellent wetting properties†

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A series of branched dodecyl phenol polyoxyethylene ethers (b-DPEO_n) were successfully synthesized via alkylation and ethylene oxide addition reactions. The alkylation reaction was conducted by using a branched internal olefin as the raw material. Furthermore, the conversion rate of the branched dodecene was measured to be 98.1% and the selectivity towards branched dodecyl phenol (b-DP) was 95.9%. Moreover, b-DPEO_n (b-DPEO₇, b-DPEO₁₀, b-DPEO₁₂) were obtained via the reaction of ethylene oxide with b-DP. Notably, b-DPEO₁₀ can efficiently reduce the surface tension of water below 31.55 mN m⁻¹ at the critical micelle concentration (cmc) and the cmc value in water was approximately 1.3×10^{-2} g L⁻¹ at 25 °C. The preferable wetting ability of b-DPEO₁₀ was superior to that of commercialized dodecyl phenol polyoxyethylene ether (c-DPEO_n), so it will be promoted and used in the textile and pesticide industries.

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Introduction

Surface tension is caused by the imbalance of the forces on the molecules at the interface of the two phases. Surfactant molecules usually feature a hydrophilic group at one end and a lipophilic group at the other, which can form an arrangement on the surface of the solution to improve the unbalanced state of the surface molecules, thereby reducing the surface tension of the solution.¹⁻⁴ Non-ionic surfactants represent an important subgroup of surfactants, whose hydrophobic groups are generally composed of alkyl-phenols, fatty alcohols, *etc.* On the other hand, their hydrophilic groups consist of polyoxyethylene (EO) or polyoxypropylene (PO) chains.⁵⁻⁸ The properties of non-ionic surfactants can be adjusted by changing the lengths of hydrophobic or hydrophilic groups and by modifying the structures of the hydrophobic moiety.^{9,10}

At present, commercialized alkylphenol polyoxyethylene ethers are obtained from alpha olefins (primarily the linear ones), while only a few branched alkylphenol polyoxyethylene ethers have been reported. However, studies have revealed that the hydrophobic base with a branched structure show better wetting and penetrating properties than those of linear ones, even when their types and molecular sizes are comparable.¹¹⁻¹⁵ Sha and coworkers successfully synthesized a novel branched fluorinated anionic surfactant 4-(3,3,4,4,5,5,5-heptafluoro-2,2-

bis-trifluoromethyl-pentyl)-benzene lithium phosphonate via a four-step route by using perfluoro-2-methyl-2-pentene as a starting material.¹⁶ This branched fluorinated anionic surfactant exhibits excellent surface activity, which reduces the surface tension of water to be 19.2 mN m⁻¹. Zhang *et al.* successfully obtained novel anionic surfactants alcohol polyoxyethylene ether carboxylates with three branched chains.¹⁷ One of these novel branched surfactants features allows for the reduction of surface tension of water to around 27 mN m⁻¹. Compared with linear surfactants, branched surfactants exhibit preferable wetting ability toward potential application, which was mainly caused by the difference in diffusion rate.¹⁸⁻²⁰

Although high-branched alkyl tails may lead to poor biodegradability, branched surfactants with limited short side-chains have been testified to have negligible effect on their biodegradability.²¹ With respect to the linear isomer, a branched alkyl amine oxide surfactant (C₁₀DAO-branched) showed the ability to form concentrated solutions without generating lyotropic liquid crystals²² or to tune supramolecular aggregation upon pH stimuli.²³ Moreover, C₁₀DAO-branched can keep low viscosity even in mixtures with other linear surfactants, thereby the rational design of branched alkyl structures has become an efficient strategy to increase the concentration of surfactants in detergent formulations.²⁴

In this research, we present a series of branched dodecyl phenol polyoxyethylene ether (b-DPEO_n), which were obtained through alkylation and ethylene oxide addition reaction in high yields, as shown in Scheme 1. It is noteworthy that the raw material non-terminal dodecene is a by-product of the butene

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Test of wetting ability

The coarse canvas is used in the test, and the specification is C/C 21/2 × 8/2. The canvas is cut into round cloth pieces with a diameter of 30 mm, with a mass range of 0.44 to 0.45 g. The 0.5 g L⁻¹ solution of surfactant in water is prepared, and the temperature is kept at 25 °C after dissolution. The round cloth pieces were put into the solution horizontally. The time from entering the solution to the beginning of sinking was denoted as the wetting time, and three parallel tests were carried out for each surfactant.

Test of polyethylene glycol content

Test instrument. Agilent Technologies 1260 Infinity LC.

Agilent LC column: ZORBAX SB-C18, 4.6 × 150 mm × 5 μm.

The content of polyethylene glycol was analyzed by LC, and the content of polyethylene glycol was obtained from the standard curve. The function of the standard curve was $\lg A = 1.5025 \lg x + 2.1948$, where A was the peak area and x was the concentration of polyethylene glycol. The test method of the sample was as follows: weighed m_1 g of the sample in a 10 mL volumetric flask, dissolved it in a methanol aqueous solution. The mass concentration of the aqueous methanol solution was 80%. The formula for the content of polyethylene glycol was as follows:

$$X = \frac{x \cdot 10}{m_1} \times 100\%$$

Results and discussion

Synthesis of b-DP

Excess phenol was typically added in the alkylation reactions with the presence of olefin and phenol as raw materials to

reduce the content of multi-substituted products and increase the yield of mono-substituted ones.²⁵ In this paper, the conversion rate of the reaction was calculated based on the remaining content of dodecene in the products, showing that the main by-products in the reaction were multi-substituted alkylphenol. The alkylation reaction also yielded other organics including decyl phenol and cetyl phenol. However, their contents were less than 0.5% wt and therefore they were not discussed in this study. As shown in Fig. 1, lowering the reaction temperature, increasing the molar ratio of phenol to olefin, and reducing the reaction time exclusively resulted in the reduction of the selectivity of multi-substituted products. As a consequence, conversion rate of olefins decreased and the difficulty of the separation and purification increased. To maintain the high conversion rate of olefins and high selectivity of mono-substituted products, the reaction condition was optimized. The amount of catalyst added was 5%, the molar ratio of phenol to olefin was 4 : 1, the reaction temperature was 90 °C, and the reaction time was 5 h. Under this condition, the conversion rate of branched non-terminal dodecene and the selectivity of b-DP was determined to be 98.1% and 95.9%, respectively.

Synthesis of b-DPEO_n

Characterization of b-DPEO_n. After separating and purifying, b-DPs further reacted with ethylene oxide to yield three kinds of b-DPEO_n (b-DPEO₇, b-DPEO₁₀, and b-DPEO₁₂). The structures were analyzed by FTIR and ¹H NMR, and the results were shown in Fig. 2 and S1,† respectively.

The main characteristic absorption peaks of the DPEO_n can be found in the FTIR spectra. The peak of 3450 cm⁻¹ corresponds to the stretching vibration of the O-H group in the phenolic hydroxyl group, and the peaks of 1640, 1600, 1510, and 1460 cm⁻¹ can be assigned to the stretching vibration of the

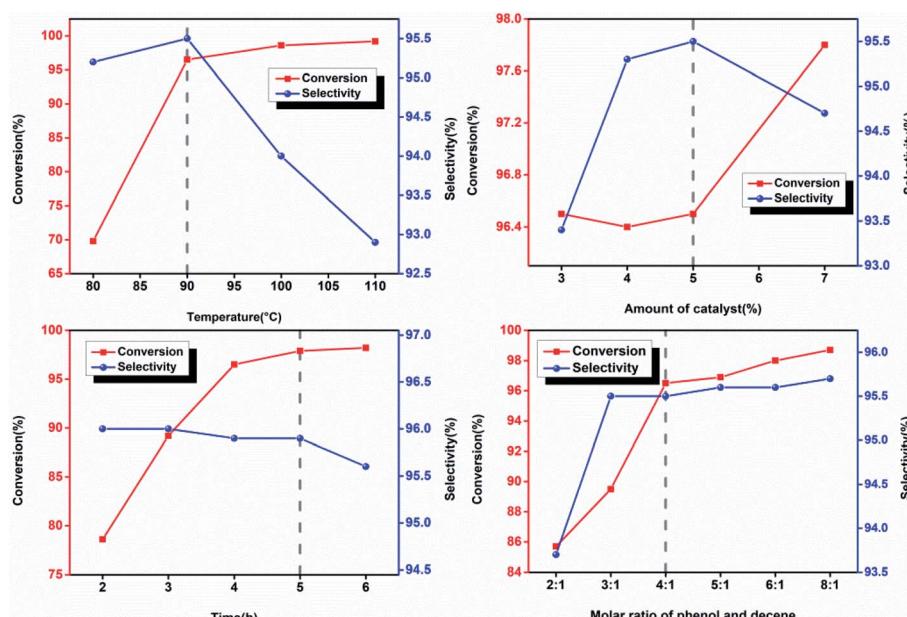


Fig. 1 Effects on the synthesis of b-DP.



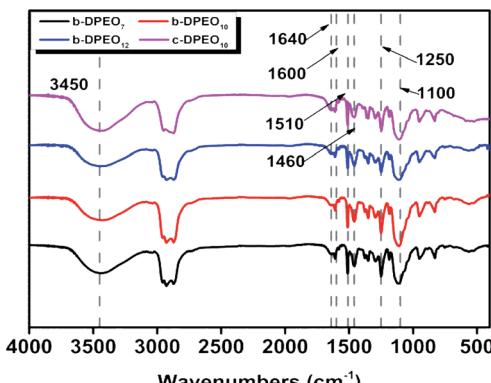


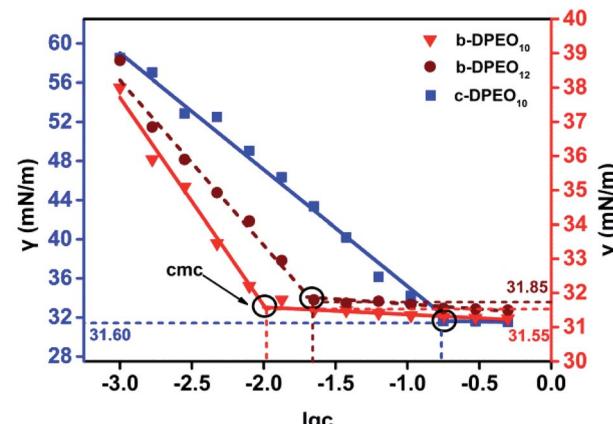
Fig. 2 FTIR spectra.

benzene ring. The anti-symmetric stretching vibration peak of C—O—C group and the symmetric stretching vibration peak locate at 1250 cm^{-1} and 1100 cm^{-1} , respectively. Combined with the analysis of the ^1H NMR, it can be seen that b-DPEO₁₀ and c-DPEO₁₀ have similar chemical shifts regions of phenyl H, methyl H and methylene H, which further confirm their structural relevance. However, the branched chain in b-DPEO₁₀ leads to more complicated chemical environments for the H on the hydrocarbon chain, as exemplified by the larger integration in the range of 0–2 ppm due to the presence of more methyl H in the branched one.

Properties of b-DPEO_n. The hydroxyl value can be used to define the average molecular weight of the product. Furthermore, the contents of polyethylene glycol may affect the surface activity of the product. These two parameters are summarized in Table 1.

It can be seen from Table 1 that all the hydroxyl values of the synthesized samples are close to the theoretical values, while the hydroxyl value of b-DPEO₁₂ is higher than the calculated value. This result combined with the polyethylene glycol content suggests that the hydroxyl value of b-DPEO₁₂ is higher because the reaction time was longer and more polyethylene glycol was formed. The hydroxyl value of c-DPEO₁₀ is higher than the theoretical value, which is mainly caused by the naming convention in the industry. According to the hydroxyl value calculation, its actual degree of polymerization should be around 9. The molecular weights of the b-DPEO_n were in a reasonable range based on the theoretical hydroxyl values, which allows for the measurement of surface tension and test of performance.

Surface tension of b-DPEO. The surface tension (γ) was taken as the ordinate, and the logarithm of the concentration ($\lg c$)

Fig. 3 Plots of γ – $\lg c$ of the DPEO_n at 25 °C.

was taken as the abscissa to get Fig. 3. Because the solubility of b-DPEO₇ is modest, the surface tension research cannot be carried out under the same experimental condition as the other samples. The data in Table 2 can be calculated by formulas (1)–(4). Compared with the commercialized samples, the cmc value of the products are significantly lower, which was mainly due to the influence of the branched hydrophobic chains. The minimum surface area A_{\min} of the molecules with branched chains is larger, which reduces the saturated adsorption amount (Γ_{\max}) at the interface. Therefore, micelles can be easily formed in the solution at low concentrations, and the correspondingly the cmc value is lower. Although the adsorption capacity of b-DPEO_n at the interface is not as large as that of the commercialized sample, the hydrocarbon chain of the later one is prone to bend due to its linear nature. This phenomenon results in the lower $-\text{CH}_3$ group number in the outermost layer of the commercialized sample than that of the branched ones. Since the surface energy of $-\text{CH}_3$ group was higher than $-\text{CH}_2$ group, the γ_{cmc} of b-DPEO_n can be comparable to that of the commercialized sample.

Wetting ability of b-DPEO_n. As depicted in Fig. 4, the wetting ability of b-DPEO_n was tested under three different mass concentrations, showing that the performance of b-DPEO_n are better than c-DPEO₁₀. Specifically, b-DPEO_n shows good wetting ability at the concentration of 0.1 g L^{-1} and its performance can be optimized at a concentration of 0.5 g L^{-1} . The pc_{20} of b-DPEO_n is higher, implying that b-DPEO_n has a higher efficiency of reducing surface tension. At the same time, the sizes of b-DPEO_n in water are smaller than that of the linear surfactant c-DPEO₁₀. Branched surfactant can penetrate to the surface of the internal fibers more quickly to complete the directional

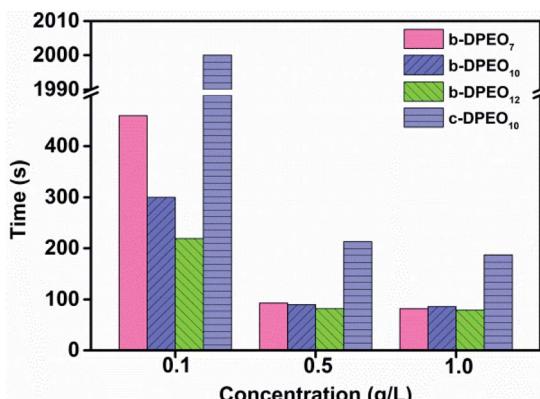
Table 1 Hydroxyl values and polyethylene glycol contents

Sample	Hydroxyl value/mg KOH g ⁻¹	Polyethylene glycol content/% wt	Theoretical hydroxyl value/mg KOH g ⁻¹
b-DPEO ₇	96.07	1.81	98.37
b-DPEO ₁₀	80.26	2.13	79.89
b-DPEO ₁₂	73.17	2.76	70.93
c-DPEO ₁₀	85.79	2.60	79.89



Table 2 Surface tension of surfactants

Sample	cmc/g L ⁻¹	$\gamma_{cmc}/\text{mN m}^{-1}$	$\Pi_{cmc}/\text{mN m}^{-1}$	pc ₂₀	$\Gamma_{max} \times 10^{10}/\text{mol cm}^{-2}$	A_{min}/nm^{-2}
b-DPEO ₁₀	0.013	31.55	40.45	5.48	0.99	1.68
b-DPEO ₁₂	0.020	31.85	40.15	5.72	0.85	1.95
c-DPEO ₁₀	0.200	31.60	40.40	2.41	2.09	0.80

Fig. 4 Wetting ability of b-DPEO_n.

arrangement, which give rise to a better wetting ability than the linear one. In contrast, a recent work showed that the cmc of a branched amine-oxide was higher than that of the linear analogue.²³ It appears that the effect of branches depends on the final architecture of the tail, which regulates the critical packing parameter of the surfactants bearing short side chains.

Conclusions

The b-DP was obtained through the reaction of branched non-terminal dodecenes and phenol. The optimal reaction condition was as follows: the amount of catalyst added is 5%, the molar ratio of phenol to branched dodecene is 4 : 1, the reaction temperature is 90 °C, and the reaction time is 5 h. Under this condition, the conversion rate of branched non-terminal dodecene is 98.1%, and the selectivity of b-DP is 95.9%. A series novel surfactants b-DPEO_n were synthesized by the ethylene oxide addition reaction of b-DP. The characterizations of FTIR and ¹H NMR confirmed that the structures of b-DPEO_n and the commercialized linear sample are similar. It was found that the b-DPEO₁₀ exhibits excellent surface activity to reduce the surface tension of water below to 31.55 mN m⁻¹ at the cmc and the cmc value in water is about 1.3×10^{-2} g L⁻¹ at 25 °C. Compared with c-DPEO_n, b-DPEO_n exhibited preferable wetting ability. This work can provide a feasible solution for the application of internal olefins, by-products of the olefin oligomerization unit.

Author contributions

Xing Wang: data curation, writing-original draft, formal analysis. Junfeng Qian: conceptualization, methodology, validation. Zhonghua Sun: investigation, methodology. Zhihui Zhang:

writing-review & editing. Mingyang He: project administration, conceptualization, funding acquisition.

Conflicts of interest

The authors declare no conflict of interest.

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