

Synthesis of budesonide conjugates and their anti-inflammatory effects: a preliminary study

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Purpose: Budesonide (Bud) is a nonhalogenated glucocorticoid with high anti-inflammatory potency and low systemic side effects. However, the poor water solubility of Bud affects its dissolution and release behavior, thus influencing its anti-inflammatory effect. This study was aimed at synthesizing and evaluating novel conjugates of Bud, hoping to increase the anti-inflammatory activity of Bud by improving its water solubility.

Materials and methods: Seven novel Bud conjugates (**3a–3g**) were designed and synthesized in this study. Besides, the equilibrium solubility, cell viability, in vitro and in vivo anti-inflammatory activity, and the hydrolysis behavior of the conjugates in different pH solutions, rat and human plasma, and rat lung homogenate were studied in detail.

Results: As compared to Bud, the equilibrium solubility of **3a**, **3c**, and **3e** was significantly increased; **3a**, **3b**, and **3c** significantly inhibited the interleukin-6 production in lipopolysaccharide-induced A549 cells; **3a** and **3e** could significantly decrease the xylene-induced ear edema; and **3a** and **3c** were gradually and slowly hydrolyzed into Bud in the alveolar fluid and lung homogenate and broken down quickly in plasma.

Conclusion: The amino acid ester compounds budesonide-21-glycine ester (**3a**) and budesonide-21-alanine ester (**3c**) were selected as potential conjugates of Bud. This study would provide a theoretical and an experimental basis for the in vivo process of glucocorticoids and the treatment of inflammatory diseases.

Keywords: budesonide, glucocorticoid, anti-inflammatory effect, equilibrium solubility, hydrolysis behavior

Introduction

Glucocorticoid has received much attention and has been widely used for the clinical treatment of allergy, asthma, autoimmune disease, and sepsis due to its powerful anti-inflammatory and immunosuppressive effects.^{1,2} Budesonide (Bud) is a kind of nonhalogenated glucocorticoid and characterized by a strong first-pass effect, a short half-time, less absorption and metabolism, and a high affinity for glucocorticoid receptors.³ In clinic, Bud is mainly used for the treatment of asthma, allergic rhinitis, and ulcerative colitis by inhalation and enteroclysis. Bud, which is a continuously inhaled corticosteroid, is commonly administered for the treatment of human asthma.^{4,5} Compared with other glucocorticoids in this class, Bud has stronger local anti-inflammatory and pharmaceutical effects, a longer action time, and a less systemic action; therefore, it could significantly reduce the adverse reactions when used in local treatment.^{6,7} However, the poor water solubility of Bud affects its dissolution and release behavior, thus influencing the drug concentration at the inflammatory site and its local bioavailability.^{8,9}

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In order to discover new compounds with a higher activity and explore the relationship between the structure and activity of glucocorticoid receptor modulators, a recent study selected 77 prednisolone C-21 heteroaryl thioethers as target molecules^{10–12} and investigated the effect of molecular force field on their drug activity by means of molecular docking and establishing their three-dimensional quantitative structure–activity relationship (3D-QSAR) models. The result indicated that the introduction of hydrophilic groups into the C₂₁ substituent position of glucocorticoid could enhance the drug activity.¹³ Therefore, we hypothesized that the anti-inflammatory activity of Bud would be increased by conjugating small hydrophilic groups to its 21-hydroxy group with an increase in water solubility.

In the present study, seven novel Bud conjugates were designed and synthesized by introducing small hydrophilic molecules into the 21-hydroxy group of Bud, that is, budesonide-21-glycine ester (**3a**), budesonide-21-phenylalanine ester (**3b**), budesonide-21-alanine ester (**3c**), budesonide-21-diethylamine acetate (**3d**), budesonide-21-dimethylamine acetate (**3e**), budesonide-21-N-methylpiperazine acetate (**3f**), and budesonide-21-morpholine acetate (**3g**). Furthermore, their equilibrium solubility, cell viability, in vitro and in vivo anti-inflammatory effect, and hydrolysis behavior in different pH solutions, rat and human plasma, and rat lung homogenate were studied in detail. Hopefully, the synthesized conjugates could improve the water solubility and thus increase the anti-inflammatory activity of Bud.

Materials and methods

Materials

Bud, N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide hydrochloride (EDCI), 4-dimethylaminopyridine (DMAP), Boc-glycine, Boc-phenylalanine, and Boc-alanine were purchased from Duodian Chemistry Co. Ltd. (Nanjing, China). Sodium iodide, diethylamine, dimethylamine hydrochloride, N-methylpiperazine, morpholine, lipopolysaccharide from *E. coli* O111:B4 (LPS-EB), and MTT were purchased from Aladdin Bio-Chem Technology Co. Ltd. (Shanghai, China). Human interleukin (IL)-6 ELISA kit was purchased from Dakewe Bioengineering Co. Ltd. (Shenzhen, China). The human alveolar adenocarcinoma cell line (A549) was kindly provided by Stem Cell Bank, Chinese Academy of Sciences (Shanghai, China). The blank plasma of healthy volunteers was given by the First Affiliated Hospital of Xi'an Jiaotong University (Xi'an, China).

Male Sprague Dawley rats (220–250 g) and male Kunming mice (20–25 g) were obtained from Laboratory

Animal Center of Xi'an Jiaotong University, SCXK 2007-001, Shaanxi, China. The animals were housed under a 12-hour light–dark cycle at a constant ambient temperature (22°C–25°C), with normal chow and water ad libitum. They were allowed to acclimatize for 1 week before the experiments were started. All animal care and experimental protocols were in strict accordance with the Guidelines of the Laboratory Animal Center of Xi'an Jiaotong University and approved by the Institutional Animal Care and Use Committee of Xi'an Jiaotong University (No XJTULAC2016-004).

Synthesis of amino acid ester conjugates of Bud

Bud (1.5 g, 3.5 mmol), EDCI (1.35 g, 7.0 mmol), and DMAP (0.24 g, 2.0 mmol) were dissolved in dichloromethane (10 mL). Boc-glycine (1.02 g, 5.8 mmol) dissolved in dichloromethane was added dropwise into the above solution, and the mixture was continuously stirred at 30°C for 2 hours. The reaction was monitored by a thin-layer chromatography (TLC) method, with a developer of petroleum ether/ethyl acetate (2:1). The dichloromethane was removed by rotary evaporation, and the residue was dissolved in ethyl acetate and washed twice with hydrochloric acid (2 mol/L) and saturated sodium bicarbonate solution, respectively. The product was dried with anhydrous magnesium sulfate, and the evaporation of solvent in vacuum afforded a crystallized yellow solid. The solid residue was placed onto a silica gel column and eluted with petroleum ether/ethyl acetate (4:1); then, a crystallized white solid was obtained (**2a**, 1.8 g, 88.6%).

Compound **2a** (0.3 g, 0.51 mmol) and trifluoroacetic acid (3.4 mL) were dissolved in dichloromethane (10 mL), and the mixture was cooled to 0°C and stirred for 4 hours. The reaction was monitored by a TLC method with a developer of petroleum ether/ethyl acetate (1:1). The mixture was adjusted to alkaline with a saturated sodium carbonate solution, washed twice with water, and dried with anhydrous magnesium sulfate. The product was then dried by rotary evaporation and washed twice with cold petroleum ether. Finally, a crystallized yellow solid was obtained (**3a**, budesonide-21-glycine ester; 0.28 g, 96.1%).

Budesonide-21-phenylalanine ester (**3b**) and budesonide-21-alanine ester (**3c**) were obtained similarly with the above methods. Boc-phenylalanine and Boc-alanine were used instead of Boc-glycine, respectively (Figure 1A).

Synthesis of acetates conjugates of Bud

Bud (2.0 g, 3.96 mmol) and chloroacetyl chloride (3.5 mL) were dissolved in N,N-dimethylformamide (10 mL), the

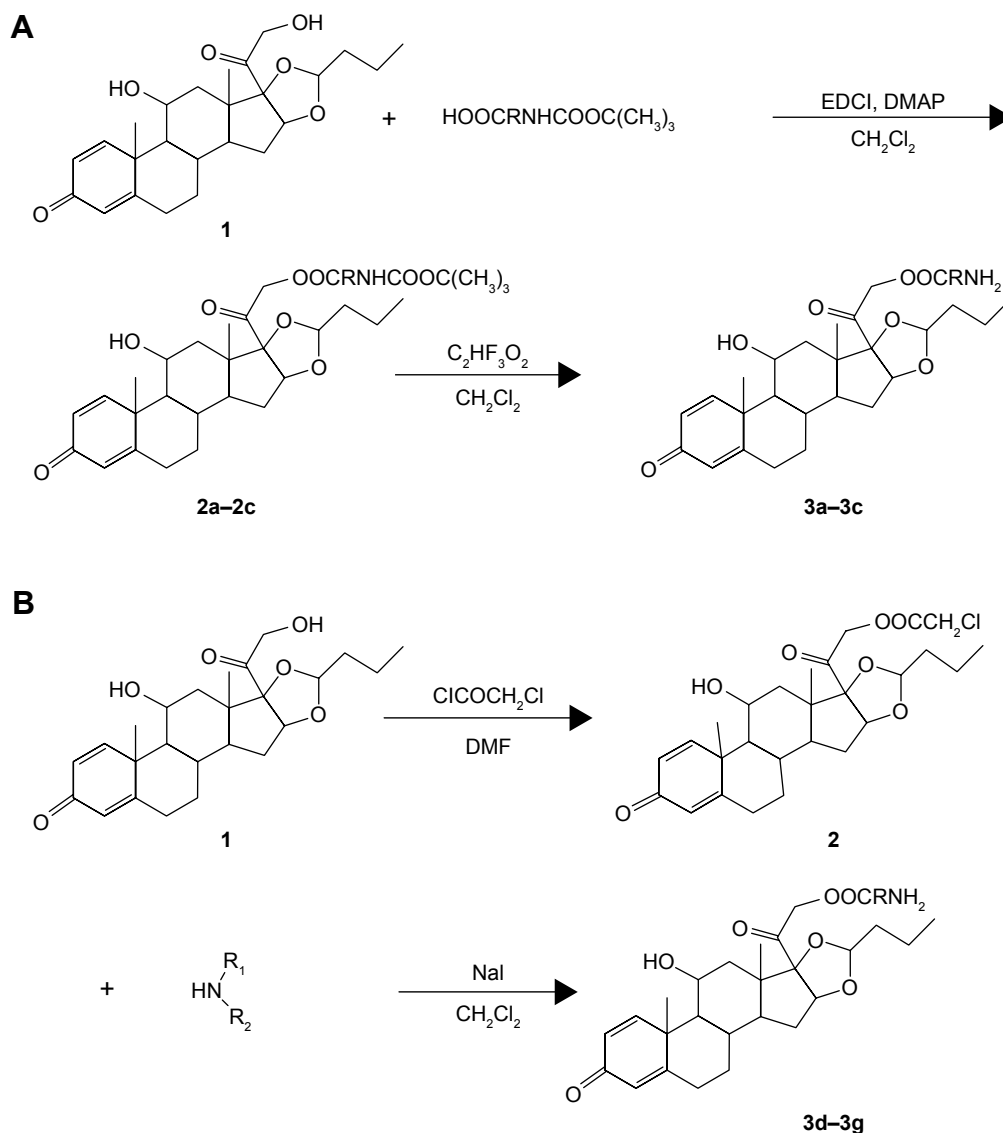


Figure 1 (A) Synthesis of amino acid ester conjugates of budesonide and (B) synthesis of acetates conjugates of budesonide.

Notes: **1**, budesonide; **2**, budesonide-21-chloroacetate; **2a–2c**, N-Boc-amino acid ester conjugates of budesonide; **3a**, budesonide-21-glycine ester; **3b**, budesonide-21-phenylalanine ester; **3c**, budesonide-21-alanine ester; **3d**, budesonide-21-diethylamine acetate; **3e**, budesonide-21-dimethylamine acetate; **3f**, budesonide-21-N-methylpiperazine acetate; **3g**, budesonide-21-morpholine acetate.

Abbreviations: DMAP, 4-dimethylaminopyridine; DMF, N,N-dimethylformamide; EDCI, N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide hydrochloride.

mixture was stirred at room temperature for 30 minutes, and the reaction was monitored by a TLC method with a developer of petroleum ether/ethyl acetate (3:1). The mixture was dispersed in water (100 mL) and filtered, and the residue was washed twice with water. After drying, a pale yellow solid was obtained (4.5 g). The solid residue was placed onto a silica gel column and eluted with petroleum ether/ethyl acetate (6:1), and then, a crystallized white solid was obtained (**2**, budesonide-21-chloroacetate; 2.84 g, 98.5%).

Compound **2** (0.3 g, 0.6 mmol), sodium iodide (0.09 g, 0.6 mmol), and diethylamine (0.72 mL) were dissolved in dichloromethane (10 mL), the mixture was refluxed for 30 minutes, and the reaction was monitored by a TLC method

with a developer of petroleum ether/ethyl acetate (1:1). The dichloromethane was removed by rotary evaporation, and the residue was washed twice with water and saturated sodium chloride solution, respectively. The product was dried with anhydrous sodium sulfate, and the evaporation of solvent in vacuum afforded a crystallized yellow solid (**3d**, budesonide-21-diethylamine acetate; 0.16 g, 74.5%).

Budesonide-21-dimethylamine acetate (**3e**, 77.9%), budesonide-21-N-methylpiperazine acetate (**3f**, 62.4%), and budesonide-21-morpholine acetate (**3g**, 67.1%) were obtained similarly with the above methods. Dimethylamine hydrochloride, N-methylpiperazine, and morpholine were used instead of diethylamine, respectively (Figure 1B).

Characterization of the synthesized Bud conjugates

The structures of the compounds were confirmed by differential scanning calorimetry (DSC), proton nuclear magnetic resonance (^1H NMR), and mass spectrometry (MS). The thermal behavior of each prodrug was analyzed by a differential scanning calorimeter (DSC822e; Mettler Toledo, Greifensee, Switzerland). Samples (5 mg) were placed in an aluminum crucible and heated at a scanning rate of $10^\circ\text{C}/\text{min}$ from 25°C to 400°C under nitrogen atmosphere. The ^1H NMR spectrum (in CDCl_3) of each sample was recorded on a Varian 400 spectrometer (Varian, Palo Alto, CA, USA) at 400 MHz using tetramethylsilane as an internal standard. The MS analysis was performed by a TSQ EnduraTM Triple Quadrupole Mass Spectrometer (Thermo Fisher Scientific, Waltham, MA, USA) equipped with an EASY-Max NG ion source system.

Equilibrium solubility of the synthesized Bud conjugates

Each conjugate and Bud powder (0.5 g) were separately dispersed in distilled water (50 mL) and continuously stirred at 25°C for 72 hours. At 0, 12, 24, 36, 48, 60, and 72 hours, 1 mL of the supernatant was sampled and filtered with $0.45\text{-}\mu\text{m}$ microporous membrane.¹⁴ The drug concentration was measured by high-performance liquid chromatography (HPLC) method to determine the equilibrium solubility of a parent drug as well as the conjugates.

The contents of Bud and the synthesized conjugates in experimental samples were measured by HPLC (Shimadzu, Kyoto, Japan) using a Diamonsil[®] ODS-2 C18 column (Dikma Technologies Co. Ltd., Beijing, China).¹⁵ The detecting conditions were as follows: (1) Bud: the mobile phase was 70% of methanol solution containing 0.1% acetic acid; (2) **3a** and **3b**: the mobile phase was a mixture of methanol/phosphate buffer (pH 8.0, 70:30, v/v); (3) **3c**: the mobile phase was a mixture of acetonitrile/phosphate buffer (pH 7.4, 70:30, v/v); and (4) **3d**, **3e**, **3f**, and **3g**: the mobile phase was a mixture of methanol/phosphate buffer (pH 8.0, 80:20, v/v). Samples were detected at 30°C , the wavelength was 242 nm, and the flow rate was 1.0 mL/min.

Cell viability of the synthesized Bud conjugates

The A549 cells were cultured in Roswell Park Memorial Institute-1640 medium, supplemented with 10% fetal bovine serum, 100 U/mL penicillin, and 100 $\mu\text{g}/\text{mL}$ streptomycin and incubated at 37°C in a humidified atmosphere with 5% CO_2 .¹⁶ Throughout the experiment, the cells were used

in the logarithmic phase of growth. They were plated into a 96-well plate at a density of 5×10^3 cells/well. The following day, the cells were treated with or without 200 μL of different concentration of drug solution for 72 hours. The medium was removed and replaced with a medium containing 5 mg/mL MTT and incubated for 4 hours. Then, the medium was removed, and the product was solubilized using dimethyl sulfoxide (DMSO). Finally, the absorbance was detected at 490 nm using a microplate reader (Bio-Tek Instruments Inc., Winooski, VT, USA) according to the manufacturer's instructions.¹⁷ The inhibitory rate of cell growth was calculated using the following equation, and the 5% inhibiting concentration (IC_{50}) was calculated from the dose–response curves.

$$\text{Inhibitory rate (\%)} = \frac{\text{OD}_{\text{test}} - \text{OD}_{\text{blank}}}{\text{OD}_{\text{control}} - \text{OD}_{\text{blank}}} \times 100\%$$

Ability of Bud conjugates to inhibit IL-6 production

The A549 cells were treated with test compounds (10^{-11} – 10^{-6} M) in 100 μL of medium and incubated in a 48-well plate for 1 hour. Then, LPS-EB solution (5 μL , 10 $\mu\text{g}/\text{mL}$) was added in each well and incubated for 24 hours. Finally, supernatants were collected and assayed for the IL-6 level by ELISA.¹⁸ The absorbance was detected at 450 nm by a microplate reader. The inhibitory rate of IL-6 production was calculated, and the half maximal inhibitory concentration of different compounds was calculated from the dose–response curves.

In vivo anti-inflammatory study

To evaluate the anti-inflammatory effect of test compounds, the xylene-induced ear edema test was conducted.¹⁹ Mice in test groups received different amounts of conjugates at an equivalent dose of 200 $\mu\text{g}/\text{kg}$ Bud, and mice in the control group received a solvent instead of a drug solution. Sixty mice were randomly divided into six groups ($n=10$): control group (2% DMSO), Bud group (200 $\mu\text{g}/\text{kg}$), **3a** group (226 $\mu\text{g}/\text{kg}$), **3c** group (233 $\mu\text{g}/\text{kg}$), **3d** group (252 $\mu\text{g}/\text{kg}$), and **3e** group (239 $\mu\text{g}/\text{kg}$). Mice in each group was administrated by intraperitoneal injection (0.1 mL/10 g of body weight). After 1 hour, each mouse received 20 μL of xylene on the right ear, and the left ear remained untreated. The mice were sacrificed 1 hour after xylene application, and 6-mm circular sections of the ears were taken. The sections of both sides of ears were weighed, and the ear edema was considered as the weight difference between the right ear and the left ear.²⁰

Biological stability of the synthesized Bud conjugates

Analytical methods of Bud conjugates

The contents of **3a**, **3c**, **3d**, and **3e** in different samples were determined by HPLC using the methods described above. The linearity of concentrations for each compound was assessed by a calibration that obtained by plotting the peak area of each sample vs its concentration (0.2–500 µg/mL), respectively. The absolute recoveries of each compound from different samples were calculated by analyzing spiked samples at 5.0, 50.0, and 500.0 µg/mL, respectively, and each concentration level was tested three times. For precision evaluation, the samples (5.0, 50.0, and 500.0 µg/mL) were analyzed three times within 1 day (intraday) and within 5 days (interday) for calculating the relative standard deviation (RSD), respectively.

Stability of Bud conjugates in different pH solutions

Then, **3a**, **3c**, **3d**, and **3e** (1 mg) were incubated in different media including pH 1.2, 6.8, and 7.4 (containing 2% DMSO) with a final concentration of 200 µg/mL.²¹ The samples were constantly shaken at 37°C for 48 hours. At 0, 1, 2, 4, 8, 12, 24, and 48 hours, 200 µL of the mixture was collected, filtered with 0.45-µm microporous membrane, and analyzed by HPLC. The residual contents of each conjugate were calculated, and the hydrolysis curves were drawn.

Stability of Bud conjugates in rat plasma

Fresh rat plasma was diluted 5.7 times (v/v), and **3a**, **3c**, **3d**, and **3e** (1 mg) were added in the diluent (containing 2% DMSO) with a final concentration of 200 µg/mL and incubated at 37°C for 24 hours. At 0, 0.5, 1, 2, 4, 8, 12, and 24 hours, 200 µL of the mixture was collected, filtered with 0.22-µm microporous membrane, and analyzed by HPLC. The residual contents of each conjugate were calculated, and the hydrolysis curves were drawn.

Stability of Bud conjugates in human plasma

Blank plasma was collected from healthy volunteers. **3a**, **3c**, **3d**, and **3e** (1 mg) were added in human plasma (containing 2% DMSO) with a final concentration of 200 µg/mL. The stability study was conducted as above.

Stability of Bud conjugates in rat lung homogenate

Rat lung was collected, weighed, and then homogenized with three times of normal saline (m/v). Then, **3a**, **3c**, **3d**, and **3e** (1 mg) were added in rat lung homogenate (containing

2% DMSO) with a final concentration of 200 µg/mL. The stability study was conducted as described above.

Statistical analyses

Statistical analyses were performed with SPSS Version 19.0 (IBM Corporation, Armonk, NY, USA). The Shapiro–Wilk test was used to test distribution. The One-way analysis of variance followed by the Fisher's least significant difference test was used to test normally distributed data. The results were presented as mean ± SD. $P < 0.05$ was considered to be statistically significant.

Results

Preparation and characterization of Bud conjugates

Figure 1A shows the synthesis of amino acid ester conjugates of Bud. The esterification was reacted between the 21-hydroxyl group of Bud (**1**) and the carboxyl group of Boc-amino acids (Boc-glycine, Boc-alanine, or Boc-phenylalanine) under the catalysis of EDCI/DMAP to produce the N-Boc-amino acid ester conjugates of Bud (**2a–2c**). Finally, **2a–2c** were deprotected with trifluoroacetic acid, and the final products (**3a–3c**) were obtained. Figure 1B shows the synthesis of acetates conjugates of Bud. The esterification was reacted between the 21-hydroxyl group of Bud (**1**) and chloroacetyl chloride to produce budesonide-21-chloroacetate (**2**). Then, **2** was reacted with the amino group of diethylamine, dimethylamine, N-methylpiperazine, and morpholine, respectively, under the catalysis of sodium iodide to obtain the final products (**3d–3g**). The synthetic routes used in this study are simple with mild reaction conditions, less side reactions, and high yields. The products were characterized by ¹H NMR, DSC, and MS. As shown in Figure 2, **3a–3g** were successfully synthesized.

Bud: ¹H NMR (CDCl₃): δ 7.30 (dd, 1H, C_{1-H}), 6.28 (dd, 1H, C_{2-H}), 6.03 (s, 1H, C_{4-H}), 4.58–4.45 (m, 3H, C_{21-H}, C_{16-H}), 4.27–4.15 (m, 2H, C_{24-H}, C_{11-H}), 2.36 (d, 2H, C_{23-H}), 2.37–1.96 (m, 2H, C_{6-H}), 1.95–1.42 (m, 3H, C_{8-H}, C_{12-H}, C_{7-H}), 1.72 (m, 1H, C_{15-H}), 1.54–1.42 (m, 5H, C_{12-H}, C_{15-H}, C_{14-H}, C_{25-H}), 1.54–1.18 (m, 3H, C_{19-H}), 1.32 (m, 2H, C_{26-H}), 0.95–0.81 (m, 5H, C_{7-H}, C_{9-H}, C_{18-H}), 0.95 (m, 3H, C_{27-H}); **3a**: Yield: 0.24 g, 96.1%. mp: 106°C–107°C. ¹H NMR (CDCl₃): δ 7.29 (dd, 1H, C_{1-H}), 6.25 (ddd, 1H, C_{2-H}), 5.99 (d, 1H, C_{4-H}), 4.72–4.95 (m, 3H, C_{21-H}, C_{16-H}), 4.46–4.59 (m, 2H, C_{24-H}, C_{11-H}), 3.54 (dt, 2H, C_{23-H}), 2.55–2.31 (m, 2H, C_{6-H}), 1.97–2.13 (m, 3H, C_{8-H}, C_{12-H}, C_{7-H}), 1.79 (m, 1H, C_{15-H}), 1.51–1.68 (m, 5H, C_{12-H}, C_{15-H}, C_{14-H}, C_{25-H}), 1.44 (s, 3H, C_{19-H}), 1.37 (m, 2H, C_{26-H}), 0.95–1.17 (m, 5H, C_{7-H}, C_{9-H}, C_{18-H}),

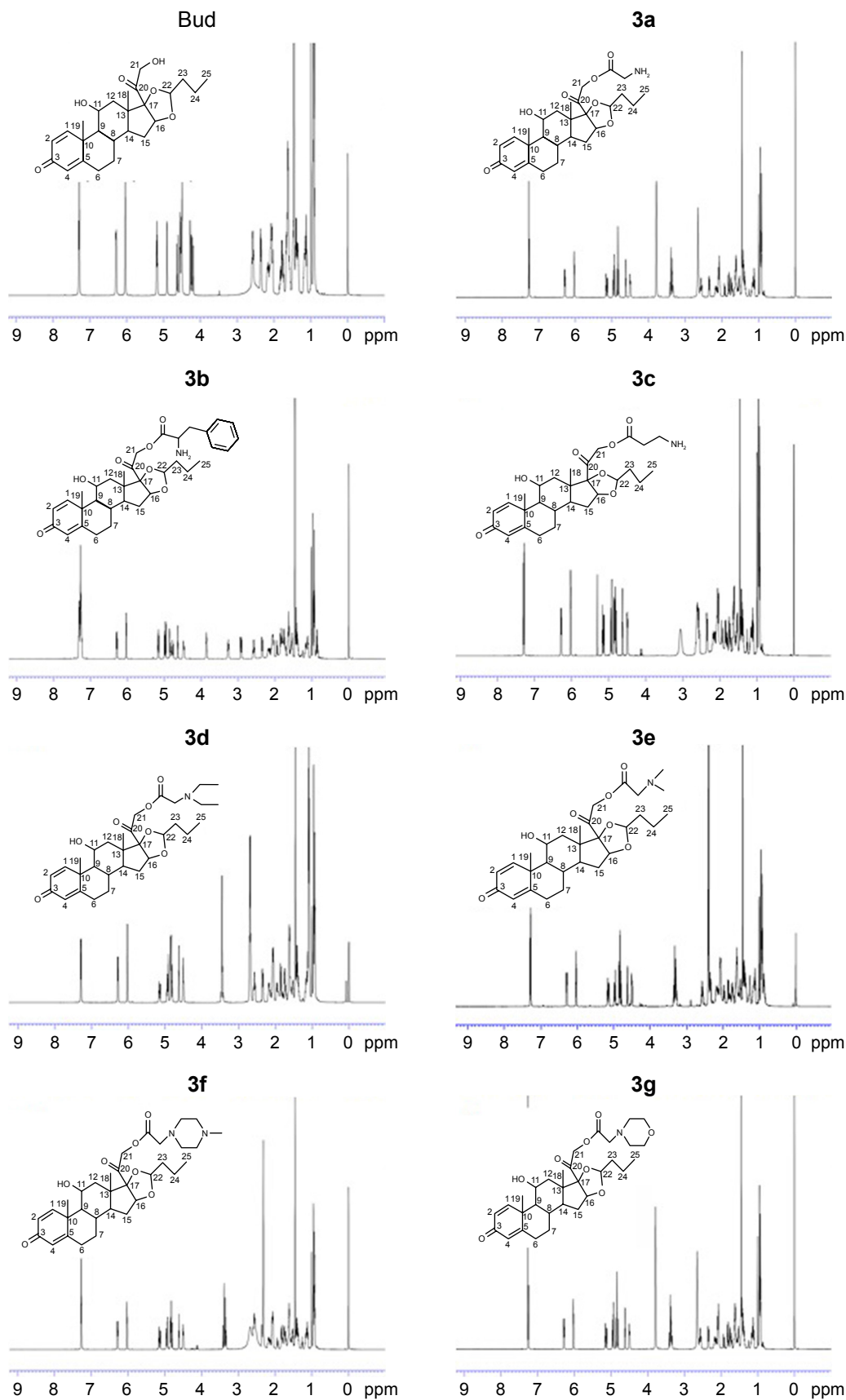


Figure 2 ^1H NMR spectra (in CDCl_3) of Bud conjugates.

Notes: **3a**, budesonide-21-glycine ester; **3b**, budesonide-21-phenylalanine ester; **3c**, budesonide-21-alanine ester; **3d**, budesonide-21-diethylamine acetate; **3e**, budesonide-21-dimethylamine acetate; **3f**, budesonide-21-N-methylpiperazine acetate; **3g**, budesonide-21-morpholine acetate.

Abbreviations: Bud, budesonide; CDCl_3 , chloroform-d; ^1H NMR, proton nuclear magnetic resonance.

0.90 (td, 3H, C_{27-H}); **3b**: Yield: 0.19 g, 68.7%. mp: 95°C–96°C. ¹H NMR (CDCl₃): δ 7.27 (m, 6H, C_{1-H}, Ph-5H), 6.03–6.28 (ddd, 1H, C_{2-H}, C_{4-H}), 5.16 (dd, 1H, Ph-H), 4.79–4.97 (m, 3H, C_{16-H}, C_{21-H}), 4.63 (t, 1H, C_{25-H}), 4.47 (dd, 1H, C_{11-H}), 3.85 (dt, 1H, C_{23-H}), 3.27 (ddd, 1H, C_{24-H}), 2.92 (dd, 1H, C_{24-H}), 2.58 (td, 1H, C_{6-H}), 2.35 (dd, 1H, C_{6-H}), 2.05–2.15 (m, 3H, C_{7-H}, C_{8-H}, C_{12-H}), 1.55–1.84 (m, 6H, C_{12-H}, C_{14-H}, C_{15-H}, C_{26-H}), 1.46 (d, 3H, C_{19-H}), 1.39 (m, 2H, C_{27-H}), 1.11–1.17 (m, 2H, C_{7-H}, C_{9-H}), 0.98 (d, 3H, C_{18-H}), 0.94 (td, 3H, C_{28-H}). MS: m/z 578.27 [M+H]⁺; **3c**: Yield: 0.23 g, 86.5%. mp: 84°C–85°C. ¹H NMR (CDCl₃): δ 7.29 (m, 1H, C_{1-H}), 6.28 (ddd, 1H, C_{2-H}), 6.03 (d, 1H, C_{4-H}), 4.49–4.93 (m, 5H, C_{11-H}, C_{16-H}, C_{21-H}, C_{25-H}), 3.06 (m, 2H, C_{23-H}), 2.59 (m, 3H, C_{6-H}, C_{24-H}), 2.34 (dd, 1H, C_{6-H}), 2.04–2.16 (m, 3H, C_{7-H}, C_{8-H}, C_{12-H}), 1.52–1.83 (m, 6H, C_{12-H}, C_{14-H}, C_{15-H}, C_{26-H}), 1.46 (s, 3H, C_{19-H}), 1.41 (td, 2H, C_{27-H}), 1.11–1.17 (m, 2H, C_{7-H}, C_{9-H}), 0.98 (d, 3H, C_{18-H}), 0.93 (td, 3H, C_{28-H}). MS: m/z 502.26 [M+H]⁺.

3d: Yield: 0.16 g, 74.5%. mp: 147°C–148°C. ¹H NMR (CDCl₃): δ 7.29 (m, 1H, C_{1-H}), 6.28 (d, 1H, C_{2-H}), 6.03 (s, 1H, C_{4-H}), 4.82–4.93 (dd, 3H, C_{16-H}, C_{21-H}), 4.63 (t, 1H, C_{24-H}), 4.50 (m, 1H, C_{11-H}), 3.44 (m, 2H, C_{23-H}), 2.57–2.69 (dd, 5H, C_{6-H}, C_{1-H}), 2.34 (d, 1H, C_{6-H}), 2.08–2.16 (m, 3H, C_{7-H}, C_{8-H}, C_{12-H}), 1.54–1.82 (m, 6H, C_{12-H}, C_{14-H}, C_{15-H}, C_{25-H}), 1.46 (s, 3H, C_{19-H}), 1.40 (td, 2H, C_{26-H}), 1.12–1.16 (m, 2H, C_{7-H}, C_{9-H}), 1.09 (t, 6H, C_{2-H}), 0.98 (d, 3H, C_{18-H}), 0.92 (m, 3H, C_{27-H}). MS: m/z 544.39 [M+H]⁺; **3e**: Yield: 0.17 g, 77.9%. mp: 135°C–136°C. ¹H NMR (CDCl₃): δ 7.28 (m, 1H, C_{1-H}), 6.29 (ddd, 1H, C_{2-H}), 6.03 (d, 1H, C_{4-H}), 4.81–4.97 (m, 3H, C_{16-H}, C_{21-H}), 4.62 (t, 1H, C_{24-H}), 4.51 (m, 1H, C_{11-H}), 3.30 (m, 2H, C_{23-H}), 2.57 (td, 1H, C_{6-H}), 2.35–2.40 (m, 7H, C_{6-H}, C_{1-H}), 2.07–2.15 (m, 3H, C_{7-H}, C_{8-H}, C_{12-H}), 1.62–1.83 (m, 4H, C_{12-H}, C_{14-H}, C_{15-H}), 1.53 (m, 2H, C_{25-H}), 1.46 (s, 3H, C_{19-H}), 1.39 (td, 2H, C_{26-H}), 1.15 (m, 2H, C_{7-H}, C_{9-H}), 0.99 (d, 3H, C_{18-H}), 0.93 (m, 3H, C_{27-H}). MS: m/z 516.27 [M+H]⁺; **3f**: Yield: 0.13 g, 62.4%. mp: 174°C–175°C. ¹H NMR (CDCl₃): δ 7.27 (m, 1H, C_{1-H}), 6.29 (ddd, 1H, C_{2-H}), 6.03 (d, 1H, C_{4-H}), 4.81–4.94 (m, 3H, C_{16-H}, C_{21-H}), 4.62 (t, 1H, C_{24-H}), 4.50 (m, 1H, C_{11-H}), 3.36 (m, 2H, C_{23-H}), 2.64 (m, 4H, C_{1-H}), 2.54 (m, 5H, C_{6-H}, C_{2-H}), 2.32–2.35 (m, 4H, C_{6-H}, C_{3-H}), 2.06–2.16 (m, 3H, C_{8-H}, C_{7-H}, C_{12-H}), 1.60–1.81 (m, 4H, C_{12-H}, C_{14-H}, C_{15-H}), 1.53 (m, 2H, C_{25-H}), 1.46 (s, 3H, C_{19-H}), 1.38 (m, 2H, C_{26-H}), 1.11–1.16 (m, 2H, C_{7-H}, C_{9-H}), 0.98 (d, 3H, C_{18-H}), 0.93 (m, 3H, C_{27-H}). MS: m/z 571.26 [M+H]⁺; **3g**: Yield: 0.14 g, 67.1%. mp: 161°C–162°C. ¹H NMR (CDCl₃): δ 7.26 (m, 1H, C_{1-H}), 6.28 (ddd, 1H, C_{2-H}), 6.03 (d, 1H, C_{4-H}), 4.82–4.95 (m, 3H, C_{16-H}, C_{21-H}), 4.62 (t, 1H, C_{24-H}), 4.50 (m, 1H, C_{11-H}), 3.79 (s, 4H, C_{2-H}), 3.36 (m, 2H, C_{23-H}), 2.65 (s, 4H, C_{1-H}), 2.57 (td, 1H, C_{6-H}), 2.35 (dd, 1H, C_{6-H}), 2.08–2.16 (m, 3H, C_{8-H}, C_{7-H}, C_{12-H}), 1.61–1.82 (m, 4H, C_{12-H}, C_{14-H}, C_{15-H}), 1.54 (m, 2H, C_{25-H}), 1.45 (s, 3H, C_{19-H}), 1.37 (m, 2H, C_{26-H}),

1.11–1.16 (m, 2H, C_{7-H}, C_{9-H}), 0.98 (d, 3H, C_{18-H}), 0.92 (m, 3H, C_{27-H}). MS: m/z 558.26 [M+H]⁺.

Equilibrium solubility of Bud conjugates

Figure 3 presents the results of solubility study of conjugates. The data indicated an improvement in the solubility of the drug after binding to hydrophilic carriers. The equilibrium solubility values of **3a**, **3b**, **3c**, **3d**, **3e**, **3f**, and **3g** in 72 hours were 0.4015, 0.0035, 0.3301, 0.0126, 0.0345, 0.0189, and 0.0104 g/100 g water, respectively. Particularly, the solubility values of **3a**, **3c**, and **3e** were significantly increased 180-fold, 149-fold, and 16-fold ($P < 0.01$) as compared to Bud (0.0022 g/100 g water).

Cytotoxicity of Bud conjugates

In order to select the experimental concentrations of Bud conjugates for this study, the cytotoxicity of conjugates to A549 cells was determined. The IC₅₀ of Bud conjugates on A549 cell lines was regarded as the maximum safe concentration. As shown in Figure 4, Bud and the synthesized conjugates all showed certain cytotoxicity. However, the IC₅₀ values of **3f** and **3g** (13.35±0.11 and 43.30±0.42 nM, respectively) were significantly lower than those of Bud and five other conjugates, which indicated its higher cytotoxicity. Combined with the result of solubility study, **3f** and **3g** were excluded from the following studies.

Inhibition of IL-6 in LPS-induced A549 cells

To determine the in vitro anti-inflammatory effect of Bud conjugates, the inhibition of IL-6 production in LPS-induced A549 cells was investigated. As shown in Figure 5, Bud

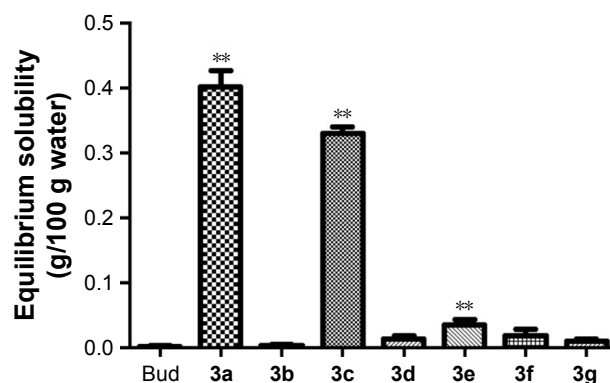


Figure 3 Equilibrium solubility of Bud conjugates at room temperature in distilled water for 72 hours.

Notes: $**P < 0.01$ compared with Bud group. Data are expressed as mean ± SD, n=3. One-way ANOVA followed by the Fisher's LSD test was used for data analysis. **3a**, budesonide-21-glycine ester; **3b**, budesonide-21-phenylalanine ester; **3c**, budesonide-21-alanine ester; **3d**, budesonide-21-diethylamine acetate; **3e**, budesonide-21-dimethylamine acetate; **3f**, budesonide-21-N-methylpiperazine acetate; **3g**, budesonide-21-morpholine acetate.

Abbreviations: ANOVA, analysis of variance; Bud, budesonide; LSD, least significant difference.

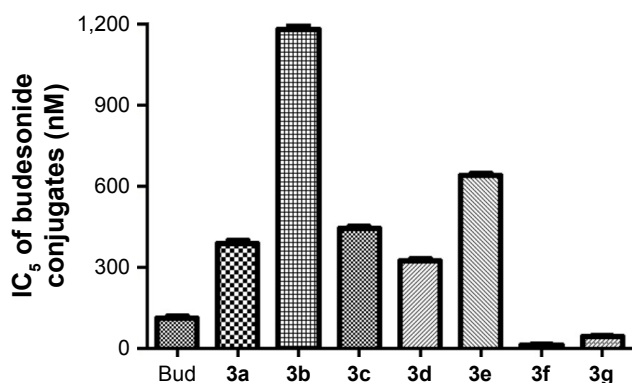


Figure 4 IC_{50} of Bud conjugates on A549 cell lines.

Notes: The cell viability was analyzed by MTT assay. Data are expressed as mean \pm SD, $n=8$. **3a**, budesonide-21-glycine ester; **3b**, budesonide-21-phenylalanine ester; **3c**, budesonide-21-alanine ester; **3d**, budesonide-21-diethylamine acetate; **3e**, budesonide-21-dimethylamine acetate; **3f**, budesonide-21-N-methylpiperazine acetate; **3g**, budesonide-21-morpholine acetate.

Abbreviations: Bud, budesonide; IC_{50} , 5% inhibiting concentration.

and the five conjugates (1 nM) resulted in an effective anti-inflammatory effect, and their inhibitory rates on IL-6 were $>50\%$. Compared with Bud, the conjugates significantly inhibited the IL-6 production in LPS-induced A549 cells ($P<0.01$); particularly, the inhibitory rates of **3a**, **3b**, and **3c** were almost twice as much as that of Bud.

In vivo anti-inflammatory study

To evaluate the in vivo anti-inflammatory activity of Bud conjugates, a xylene-induced mouse ear edema model was used; **3b** was also excluded from this study because of its poor

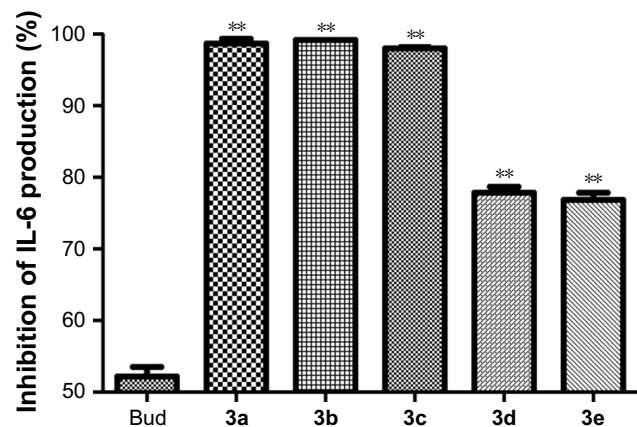


Figure 5 Bud conjugates (1 nM) inhibit the production of IL-6 on LPS-induced A549 cell lines.

Notes: $**P<0.01$ compared with Bud group. Data are expressed as mean \pm SD, $n=3$. One-way ANOVA followed by the Fisher's LSD test was used for data analysis. **3a**, budesonide-21-glycine ester; **3b**, budesonide-21-phenylalanine ester; **3c**, budesonide-21-alanine ester; **3d**, budesonide-21-diethylamine acetate; **3e**, budesonide-21-dimethylamine acetate; ANOVA, analysis of variance; Bud, budesonide; IL-6, interleukin-6.

Abbreviations: ANOVA, analysis of variance; Bud, budesonide; IL-6, interleukin-6; LPS, lipopolysaccharide; LSD, least significant difference.

solubility. It was observed that there were significant differences in ear edema between the control group and test groups. Besides, as shown in Figure 6, Bud conjugates significantly decreased xylene-induced ear edema when compared with Bud. Especially, **3a** and **3e** inhibited ear edema by 30.67% and 46.67%, respectively, while Bud presented a relatively low inhibition of 17.19%.

Biological stability of the synthesized Bud conjugates

A linear relationship was established between the concentration (0.2–500 $\mu\text{g/mL}$) and the peak area of each sample. Table 1 lists the regression equations of **3a**, **3b**, **3c**, and **3e** in different tissue samples for HPLC analysis. The absolute recoveries of the above four conjugates at each concentration (5.0, 50.0, and 500.0 $\mu\text{g/mL}$) were $>95\%$. The recoveries had no significant difference between the three concentrations. For precision assay, both the intraday and interday RSDs in all tissue samples were $<5\%$. These data validated that the HPLC method was acceptable and reproducible (Table 2).

Bud conjugates have an ester bond that is susceptible to acidic and basic conditions. Figure 7 shows the drug content that was determined following the hydrolysis of the conjugates in different pH solutions. The amount of Bud released in hydrochloric acid solution (pH 1.2) and phosphate buffer solutions (pH 6.8 and 7.4) were calculated to evaluate the hydrolysis behavior of conjugates over time; **3a**, **3d**, and **3e**

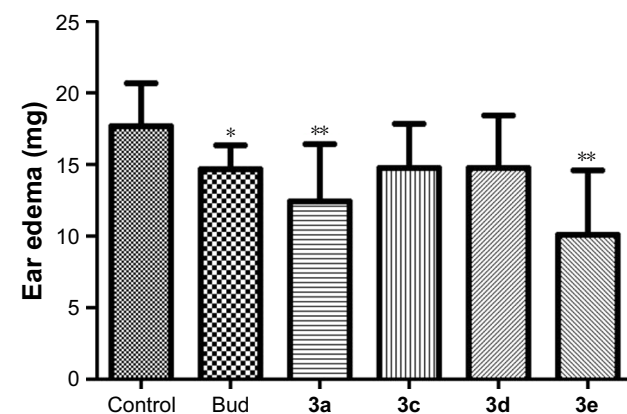


Figure 6 Effects of Bud conjugates on xylene-induced ear edema.

Notes: Ear edema was considered as the weight difference between the right ear and the left ear. Mice received 2% DMSO solution (control), 200 $\mu\text{g/kg}$ of Bud, and different amounts of conjugates (equivalent to 200 $\mu\text{g/kg}$ Bud), respectively. $*P<0.05$ vs control group; $**P<0.01$ vs control group. Data are expressed as mean \pm SD, $n=10$. One-way ANOVA followed by the Fisher's LSD test was used for data analysis. **3a**, budesonide-21-glycine ester; **3c**, budesonide-21-alanine ester; **3d**, budesonide-21-diethylamine acetate; **3e**, budesonide-21-dimethylamine acetate. **Abbreviations:** ANOVA, analysis of variance; Bud, budesonide; DMSO, dimethyl sulfoxide; LSD, least significant difference.

Table 1 Regression equations of budesonide conjugates in different tissue samples for HPLC analysis

Samples	Equations	R ²	LOQ (µg/mL)
3a			
Phosphate buffer	y=25,812x-138,166	0.9995	0.4
Rat plasma	y=7,485.4x-43,139	0.9974	0.4
Human plasma	y=1,709x+18,631	0.9992	0.4
Rat lung homogenate	y=902.3x+26,761	0.9919	0.4
3c			
Phosphate buffer	y=33,600x+38,829	0.9998	0.4
Rat plasma	y=4,654.1x-45,180	0.9920	0.4
Human plasma	y=1,750.2x-9,513	0.9990	0.4
Rat lung homogenate	y=953.4x+7,973	0.9915	0.4
3d			
Phosphate buffer	y=29,258x+99,124	0.9999	0.4
Rat plasma	y=3,162.2x-6,084	0.9924	0.4
Human plasma	y=4,399x+18,835	0.9980	0.4
Rat lung homogenate	y=3,306.7x-3,568.4	0.9953	0.4
3e			
Phosphate buffer	y=24,838x+141,746	0.9997	0.4
Rat plasma	y=5,033.7x+31,322	0.9939	0.4
Human plasma	y=4,503.2x+143,087	0.9952	0.4
Rat lung homogenate	y=5,291.8x+30,999	0.9910	0.4

Notes: **3a**, budesonide-21-glycine ester; **3c**, budesonide-21-alanine ester; **3d**, budesonide-21-diethylamine acetate; **3e**, budesonide-21-dimethylamine acetate. **Abbreviations:** HPLC, high-performance liquid chromatography; LOQ, limit of quantitation.

were barely hydrolyzed in pH 1.2 solution with the remaining 76.15%, 93.35%, and 88.53% of conjugates in 48 hours, respectively. However, conjugates hydrolyzed more quickly under neutral and basic conditions, especially in pH 7.4; **3a**, **3d**, and **3e** were almost completely hydrolyzed in 24 hours.

Figure 8 shows the residual content of conjugates after incubation with rat plasma, human plasma, and rat lung homogenate for 24 hours. As shown in Figure 8A, the four conjugates were gradually hydrolyzed in rat plasma by the action of carboxylesterase; **3a** and **3e** were rapidly hydrolyzed in 6 hours, <6% of **3c** was left in 12 hours, and **3d** was completely hydrolyzed in 24 hours. Figure 8B shows the hydrolysis behavior of conjugates in human plasma. Similar to the above result, **3a** was rapidly hydrolyzed in 6 hours, <2% of **3e** was left in 8 hours, and **3c** and **3d** were completely hydrolyzed in 8 and 24 hours, respectively.

When conjugates were incubated in the rat lung homogenate, a rapid increase was observed in the hydrolysis profile. As shown in Figure 8C, **3d** and **3e** were rapidly and completely hydrolyzed in 2 and 6 hours, respectively; 84.04% of **3a** was hydrolyzed in 6 hours and then almost completely hydrolyzed in 12 hours; 53.86% and 87.75% of **3c** were hydrolyzed after 12 and 24 hours of incubation, respectively. Hydrolysis behaviors of conjugates were

different in three conditions and were more significant in rat lung homogenate.

Discussion

Bud is a kind of nonhalogenated glucocorticoid developed in recent years, with the local anti-inflammatory activity superior to the systemic effect. Clinically, it is mainly used in the treatment of asthma, allergic rhinitis, and ulcerative colitis by inhalation and enteroclysis. Based on the study of quantitative structure–activity relationship of glucocorticoid receptor modulators, the hydrophilic groups at the C₂₁ substituent position of glucocorticoids could enhance the drug activity. Besides, the poor water solubility of drugs affects its dissolution and release behavior and thus influences the drug concentration at the inflammatory site and its local bioavailability.

In this study, seven novel Bud conjugates were synthesized (**3a–3g**); moreover, the equilibrium solubility, cell viability, in vitro and in vivo anti-inflammatory effect, and the hydrolysis behavior of the synthesized conjugates were investigated in detail. The analytical methods of Bud conjugates by HPLC in different pH solutions and biological samples were established for detecting the content of compounds in different medium. As shown in Figure 3, the equilibrium solubility study suggested that the solubility of **3a**, **3c**, and **3e** was significantly increased than that of Bud (180-fold, 149-fold, and 16-fold, respectively; $P < 0.01$), which has achieved the goal of enhancing the water solubility of Bud. Besides, the results also showed that the solubilization effect of amino acid esters was more effective than that of acetates.

Bud is a first-line drug for the treatment of corticosteroid dependent and independent asthma, which exerts its potent anti-inflammatory activity primarily by reducing the number of inflammatory cells and inhibiting the expression of inflammatory factors, such as IL-6, IL-8, and tumor necrosis factor- α .²² A549 cell is one of the main cell models used in the study of airway epithelial cell, which is first suffered damage when the respiratory tract is exposed to hazardous substance.²³ In this study, the in vitro inflammation model was established by using LPS-treated A549 cell lines. LPS was used as a stimulating factor, which is existed in the outer membrane of Gram-negative bacteria and elicits strong immune responses in animals. The inhibition of IL-6 by Bud and its conjugates at different concentrations in LPS-induced A549 cells was evaluated to reflect their anti-inflammatory effects. The data showed that the inhibitory rates of **3a**, **3b**, **3c**, **3d**, **3e**, and Bud at 1 nM were all above 50%. Besides, the inhibitory effect of the conjugates

Table 2 Recovery and precision for the detection of budesonide conjugates in different tissue samples by HPLC

Samples	Concentration ($\mu\text{g/mL}$)	Recovery (%)	Intraday RSD (%)	Interday RSD (%)
3a				
Rat plasma	5.0	100.97 \pm 1.27	0.48	1.28
	50.0	100.11 \pm 2.18	0.83	1.47
	500.0	100.05 \pm 0.73	1.26	2.15
Human plasma	5.0	101.94 \pm 1.12	1.23	1.36
	50.0	99.50 \pm 1.01	1.04	1.55
	500.0	99.08 \pm 1.68	1.48	2.29
Rat lung homogenate	5.0	99.54 \pm 1.55	0.78	2.42
	50.0	101.98 \pm 0.69	1.65	2.73
	500.0	100.59 \pm 0.35	1.35	2.51
3c				
Rat plasma	5.0	101.95 \pm 1.32	0.62	1.28
	50.0	100.91 \pm 0.58	0.29	1.33
	500.0	100.12 \pm 0.37	1.24	2.44
Human plasma	5.0	100.22 \pm 1.17	2.15	1.36
	50.0	100.90 \pm 1.33	1.18	2.05
	500.0	99.92 \pm 1.32	1.73	2.93
Rat lung homogenate	5.0	101.03 \pm 1.85	2.25	2.35
	50.0	99.67 \pm 0.71	1.71	2.74
	500.0	101.75 \pm 0.93	1.91	2.87
3d				
Rat plasma	5.0	99.98 \pm 1.76	1.25	1.25
	50.0	103.23 \pm 0.45	0.36	1.72
	500.0	101.09 \pm 1.85	1.45	2.58
Human plasma	5.0	100.67 \pm 1.33	2.27	1.30
	50.0	101.27 \pm 1.81	0.57	1.18
	500.0	100.40 \pm 0.44	0.23	1.63
Rat lung homogenate	5.0	101.70 \pm 1.16	1.02	2.09
	50.0	100.37 \pm 0.48	1.78	2.17
	500.0	101.30 \pm 1.22	1.72	2.37
3e				
Rat plasma	5.0	101.75 \pm 1.22	0.55	1.28
	50.0	100.02 \pm 0.80	0.24	2.32
	500.0	101.66 \pm 1.58	1.12	2.30
Human plasma	5.0	100.79 \pm 2.37	2.67	1.31
	50.0	100.62 \pm 0.73	1.33	2.13
	500.0	100.23 \pm 0.56	1.94	2.55
Rat lung homogenate	5.0	99.63 \pm 1.63	0.74	1.42
	50.0	99.61 \pm 1.59	1.51	2.35
	500.0	100.45 \pm 1.26	2.25	2.73

Notes: Data are expressed as means \pm SD, n=3. **3a**, budesonide-21-glycine ester; **3c**, budesonide-21-alanine ester; **3d**, budesonide-21-diethylamine acetate; **3e**, budesonide-21-dimethylamine acetate.

Abbreviations: HPLC, high-performance liquid chromatography; RSD, relative standard deviation.

was significantly higher than that of Bud ($P < 0.01$). Therefore, the synthesized conjugates possessed a desirable in vitro anti-inflammatory activity at a relatively low concentration. This result is in accordance with those reported in the previous papers. Wang et al established 3D-QSAR models for a series of prednisolone conjugates by using the comparative molecular field analysis and comparative molecular similar indices analysis.¹³ The 3D-QSAR contour maps suggested that hydrophilic substituents at C₂₁ could enhance the anti-inflammatory activity of prednisolone.

Based on the results of the equilibrium solubility and the in vitro inflammation study, **3a**, **3c**, **3d**, and **3e** were screened for further studies. The in vivo study on the anti-inflammatory activity of Bud conjugates was evaluated by the xylene-induced ear edema test. Xylene-induced ear edema is an acute and highly reproducible inflammatory response and characterized by releasing inflammatory mediators, which stimulate ear edema by increasing vascular permeability and promoting vasodilation. Therefore, the xylene-induced ear edema test presented fluid accumulation

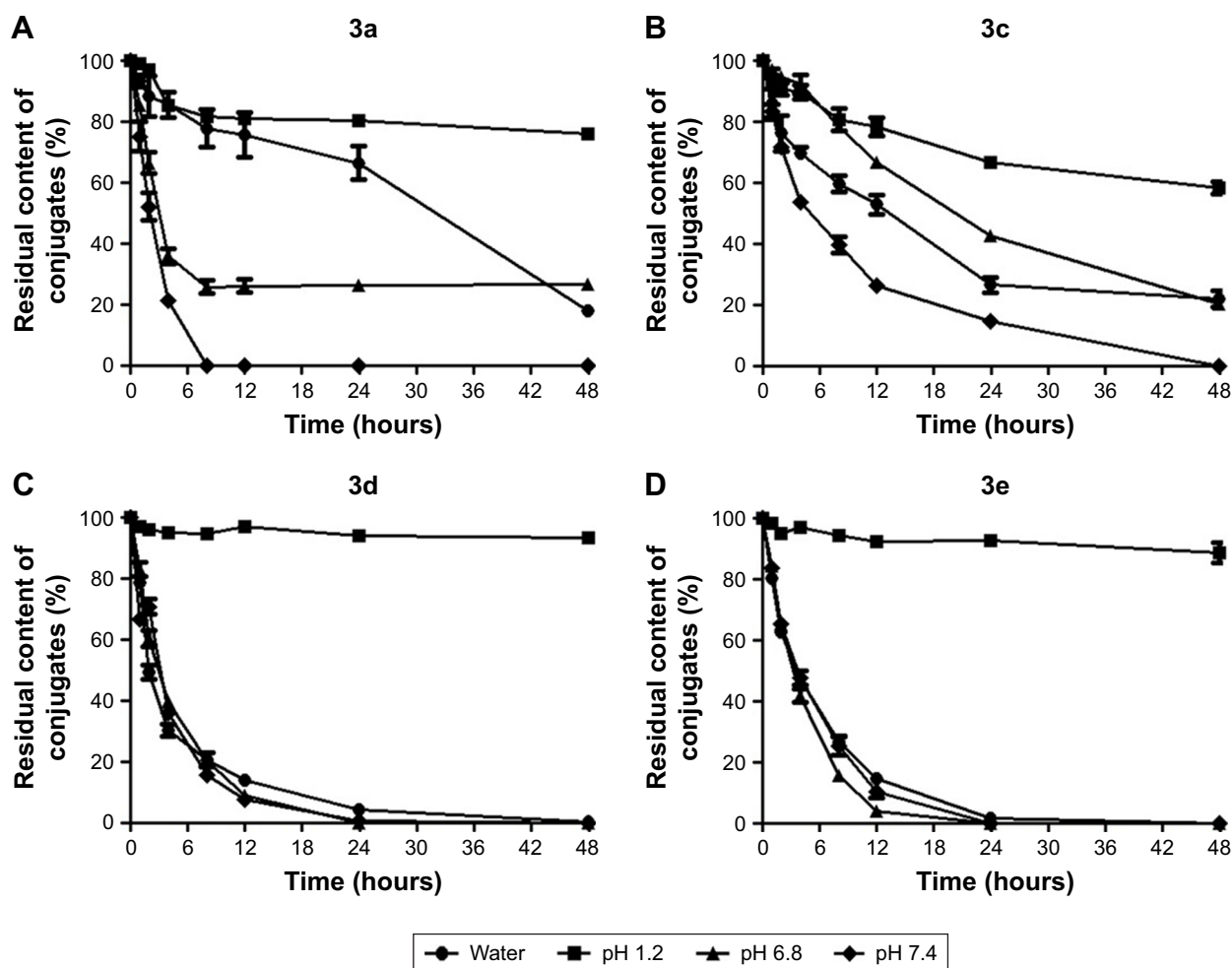


Figure 7 Hydrolysis behavior of budesonide conjugates in different pH solutions.

Notes: The residual content of conjugates that decreased with time in 48 hours was measured by HPLC. Data are expressed as mean \pm SD, $n=3$. **3a**, budesonide-21-glycine ester; **3c**, budesonide-21-alanine ester; **3d**, budesonide-21-diethylamine acetate; **3e**, budesonide-21-dimethylamine acetate.

Abbreviation: HPLC, high-performance liquid chromatography.

at the treatment site, and the inhibition of this fluid accumulation was regarded as an anti-inflammatory effect.²⁴ Figure 6 shows the significant inhibition of ear edema by conjugates. Similar to the in vitro study's findings, Bud and its conjugates significantly decreased ear edema when compared with the control ($P<0.01$); particularly, **3a** and **3e** were significantly more effective than Bud. Hereby, **3a**, **3c**, **3d**, and **3e** have been demonstrated with a desirable anti-inflammatory effect by in vitro and in vivo models, which also provide further evidence that the hydrophilic groups at the C_{21} substituent position enhance the drug activity, and the hydrophilic conjugates of Bud possess a high research value.

Bud conjugates are susceptible and instable in aqueous solutions because the existing ester bond is to be easily hydrolyzed. In the present study, the hydrolysis behavior of the synthesized conjugates was evaluated in different pH solutions, rat and human plasma, and rat lung homogenate, respectively.

The pH value of the exhaled breath condensate in patients with bronchial asthma is about 6.5–7.0.²⁵ Therefore, **3a**, **3c**, **3d**, and **3e** were incubated in hydrochloric acid solution (pH 1.2) and phosphate buffer solutions (pH 6.8 and 7.4), respectively. At time intervals, drug solutions were sampled and analyzed by HPLC to calculate the residual content of conjugates. As a result, Bud conjugates were easier to be hydrolyzed under a weakly alkaline condition, they presented a pH-dependent hydrolysis manner, and their hydrolysis rate increased with the pH value of the medium. Besides, the acetates conjugates of Bud showed a higher rate than that of the amino acid ester conjugates. Previously, Varshosaz et al prepared three dextran–Bud conjugates with different molecular weights of dextran (molecular weight =10,000, 70,000, and 500,000) and different degrees of substitution by using budesonide-21-hemisuccinate as a starting material.^{26,27} Their chemical stability was studied in 0.1 N HCl and phosphate buffer solutions of pH 6.8 and 7.4. The data showed

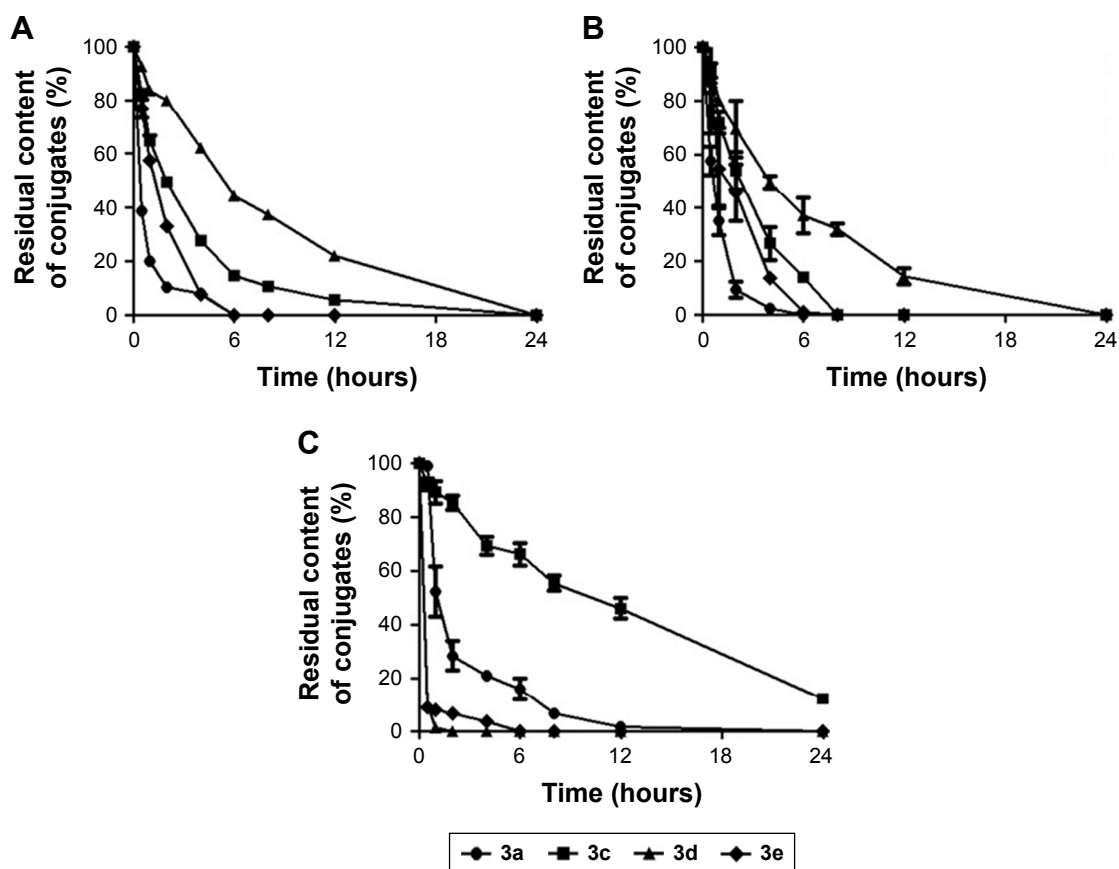


Figure 8 Hydrolysis behavior of budesonide conjugates in rat plasma (A), human plasma (B), and rat lung homogenate (C).

Notes: The residual content of conjugates that decreased with time in 24 hours was measured by HPLC. Data are expressed as mean \pm SD, $n=3$. **3a**, budesonide-21-glycine ester; **3c**, budesonide-21-alanine ester; **3d**, budesonide-21-diethylamine acetate; **3e**, budesonide-21-dimethylamine acetate.

Abbreviation: HPLC, high-performance liquid chromatography.

that the conjugates were easier to be hydrolyzed in alkaline media, which is similar to our findings. This indicated that the hydrolysis of conjugates was mediated by the cleavage of ester bond, and the main affecting factor on their hydrolysis behavior is the pH value of release media.

Second, Bud conjugates were incubated with rat and human plasma. The activity of carboxylesterase in rat plasma is about 5.7 times that in human plasma; therefore, fresh rat plasma was diluted to 5.7 times with normal saline and is then used in the biological stability study.^{28,29} As a result, the hydrolysis rate of the conjugates in plasma is $3a > 3e > 3c > 3d$; particularly, about 90.0% of **3a** was rapidly hydrolyzed in 2 hours, and 85.7% of **3d** was hydrolyzed in 12 hours in human plasma. This result suggested that most conjugates would be broken down quickly into Bud in the blood and then be metabolized in a few hours; therefore, they would not significantly affect the pharmacokinetic process of Bud and ensure the safety of the conjugates.

Finally, the hydrolysis character of conjugates was investigated in rat lung homogenate, and the hydrolysis rate of the

conjugates presented a trend as $3d > 3e > 3a > 3c$; particularly, **3d** was rapidly and completely hydrolyzed in 2 hours, and 53.8% of **3c** was hydrolyzed in 12 hours. Taken together, **3a** and **3c** would be gradually and slowly hydrolyzed into Bud in the pH environment of alveolar fluid and the lung homogenate; thus, they could have a therapeutic effect through both the form of Bud conjugates and the produced parent drug, as well as prolong the action time of Bud. While absorbed into the blood, **3a** and **3c** could be quickly broken down into Bud and thereby reduce the possible systemic adverse reactions of **3a** and **3c**.

Above all, the amino acid ester compounds **3a** and **3c** were screened as potential conjugates of Bud. Their in vitro and in vivo characteristics would provide a theoretical and an experimental basis for the intracorporal process of glucocorticoids and the treatment of inflammatory diseases. However, further study is required to confirm their pharmacokinetic and pharmacodynamic properties in detail. Currently, studies are made to evaluate the therapeutic effect of **3a** and **3c** on asthma guinea pig model induced by ovalbumin.

Conclusion

In this study, seven novel conjugates of Bud (**3a–3g**) were designed and synthesized. Based on their equilibrium solubility, cell viability, in vitro and in vivo anti-inflammatory effect, and their hydrolysis behavior in different media, two amino acid ester compounds (**3a** and **3c**) were screened as potential conjugates of Bud. As a result, the equilibrium solubility of **3a** and **3c** was significantly increased when compared to Bud, and they showed a significant anti-inflammatory effect in both in vitro and in vivo experiments. On this basis, further studies are made to evaluate their in vivo pharmacokinetics and anti-asthmatic effects in order to provide an experimental basis for further application.

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Disclosure

The authors report no conflicts of interest in this work.

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