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Synthesis and biological evaluation of novel sinomenine derivatives as anti-inflammatory and analgesic agent†

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Sinomenine (SIN) has long been known as an anti-inflammatory drug, while poor efficiency and large-dose treatment had limited its further application. A series of novel SIN derivatives **1–26** were designed and synthesized to improve its anti-inflammatory activity. The anti-inflammatory activity evaluation showed most of the derivatives exhibited enhanced anti-inflammatory activity *in vitro* compared to SIN. Compound **17** significantly inhibited LPS-induced secretion of pro-inflammatory factors NO (IC₅₀ = 30.28 ± 1.70 μM), and suppressed the expression of iNOS, IL-6 and TNF-α in RAW264.7 cells. Moreover, compound **17** showed excellent anti-inflammatory in mouse paw edema. Immunohistochemistry results revealed that compound **17** exerted anti-inflammatory activity by inhibiting the pro-inflammatory cytokine TNF-α. Furthermore, compound **17** exhibited an analgesic effect *in vivo*. The results attained in this study indicated that compound **17** had the potential to be developed into an anti-inflammation and analgesic agent.

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Introduction

Inflammation is a common pathological process, which occurs as a result of exposure of tissues and organs to harmful stimuli such as microbial pathogens, allergens or toxic cellular components.¹ The primary physical manifestations of inflammation are redness, swelling, heat, pain, and loss of function to the affected area.² Although inflammation is a beneficial automatic defence mechanism, uncontrolled inflammation is the main cause of many serious diseases such as rheumatoid arthritis, systemic lupus erythematosus, atherosclerosis, *etc.*^{3–5}

The commonly used anti-inflammatory drugs in clinics are mainly divided into glucocorticoids and nonsteroidal anti-inflammatory drug (NSAIDs).⁶ Glucocorticoids such as prednisolone, dexamethasone and hydrocortisone are one of the common clinical drugs, and exhibit good anti-inflammatory effects on inflammation caused by various causes. However, long-term use of these drugs can cause edema, hypokalemia, abnormal increase of blood sugar, digestive tract ulcers and other side effects.

NSAIDs have a great efficacy on inflammatory and pain by inhibiting cyclooxygenase (COX) activity. Currently, NSAIDs were among the most frequently used over-the-counter drugs in the world, accounting for 5% of all prescription drugs.⁷

Although the use of NSAIDs was clearly unavoidable, but the disadvantages of NSAIDs that may lead to an increased risk of cardiovascular disease and organ toxicity such as liver and kidney cannot be ignored.^{7,8} Yet, the development of more effective and less toxic drugs from natural products to treat acute inflammation as well as chronic inflammatory diseases remains a challenge in medicinal chemistry.^{9–14}

Sinomenine (SIN), an alkaloid extracted from the anti-rheumatism Chinese medicine *Sinomenium acutum* (Thunb.) Rehd. et Wils., has long been known as an anti-inflammatory drug.^{15–19} SIN has been reported to inhibit the production of proinflammatory mediators, such as TNF-α, nitric oxide (NO) and IL-6, both *in vitro* and *in vivo*.^{20–22} Therefore, SIN was regarded as lead compounds in the drug research, especially for anti-inflammatory medicines, while low activity had limited its clinical application.^{23–27} To improve activity, structural modification of SIN has attracted enormous interest in recent years.^{28,29} Studies have shown that the phenolic hydroxyl group of C-4 was easy to oxidize and decompose and may be a factor leading to allergic reactions *in vivo*. Structure–activity relationships (SARs) showed that the modification at C-1 and C-4 could enhance the activity of SIN. Several SIN derivatives have been reported and exhibit strong anti-inflammatory activity (Fig. 1).^{30–32} Introducing amino acid group into the drug molecule could enhance its anti-tumor and anti-inflammatory activity and increase its solubility.^{33,34} Meanwhile, nitrogen-based heterocycles in the drug design are almost 60% of unique small-molecule drugs. A large number of nitrogen-based heterocyclic derivatives including pyrazine, pyrimidine and

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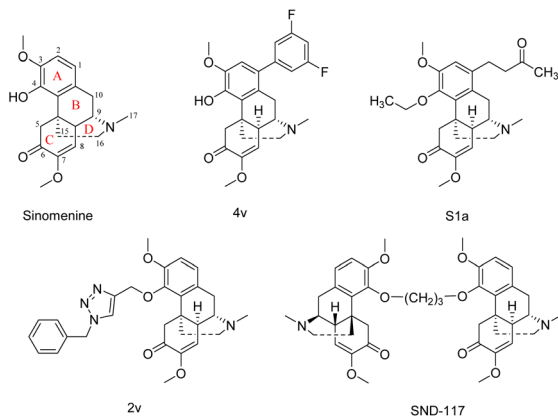


Fig. 1 Structures of sinomenine and previously synthesized compounds.

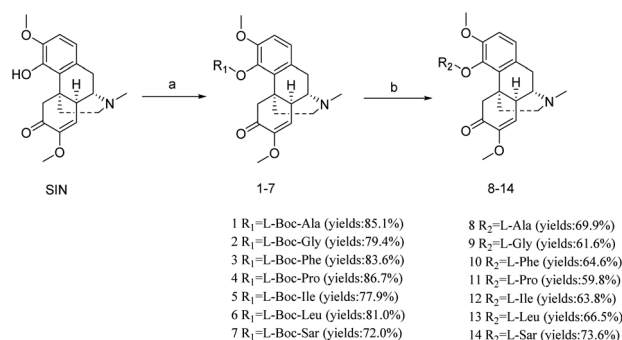
pyridazine have been discovered and described for their anti-inflammatory properties.³⁵

In this study, to improve the anti-inflammatory activity of SIN, a variety of amino acid and nitrogen-based heterocyclic biologically active small fragments were introduced to SIN to synthesize compounds 1–26. The anti-inflammatory activity and cytotoxicity of all the novel derivatives were evaluated by LPS-induced RAW264.7 inflammatory cell model. Then the preliminary anti-inflammatory mechanism of compound 17 was evaluated by qPCR experiment. Furtherly, the λ -carrageenan-induced paw oedema and acetic acid-induced writhing mouse model was used to determine the anti-inflammatory and analgesic effects of compound 17. This study lays the foundation for the subsequent development of new anti-inflammatory drugs of SIN and has practical significance for the expansion of clinical applications of SIN.

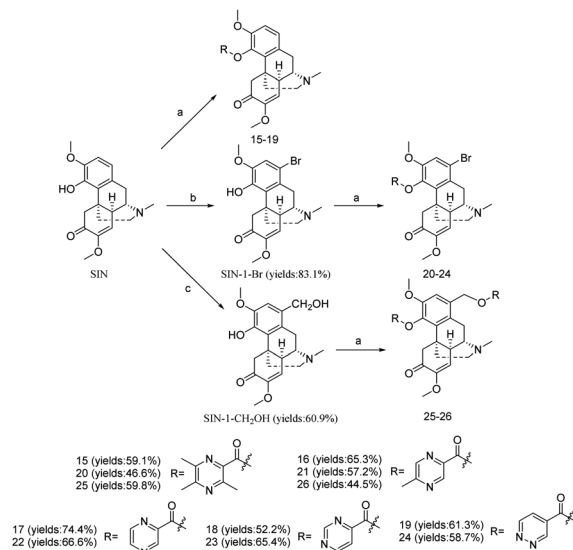
Results and discussion

Chemical synthesis

Amino acids are the basic substances necessary for human life activities and have many important physiological functions.^{36–38} Compounds 1–14 were prepared following the procedure Scheme 1. SIN was reacted with Boc-L-amino acid in dry



Scheme 1 Synthesis of compounds 1–14. Reagents and conditions: (a) Boc-L-amino acid, EDCI, DMAP, rt, 12 h; (b) TFA, rt, 30 min.



Scheme 2 Synthesis of the compounds 15–26. Reagents and conditions: (a) EDCI, DMAP, rt, 12 h; (b) NBS, rt, 10 h; (c) (CH₂O)_n, HCl, 60 °C, 2 h.

dichloromethane (DCM) in the presence of 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide hydrochloride (EDCI) and 4-dimethylaminopyridine (DMAP) to yield compounds 1–7. Subsequently, deprotection was performed with trifluoroacetic acid (TFA) in dry DCM at room temperature to afford compounds 8–14.

The 3,5,6-trimethylpyrazine-2-chloride (TMP-Cl) was synthesized according to our previous study.³⁹ In Scheme 2, SIN was reacted with N-heterocycles in dry DCM in the presence of EDCI and DMAP to yield compounds 16–19. As the H-1 of SIN was easily halogenated by N-bromosuccinimide (NBS), the SIN-1-Br was prepared. Then it was reacted with nitrogen-based heterocyclic to yield compounds 20–24. In the synthesis of compound 25–26, the SIN was firstly reacted with para-formaldehyde to afford the intermediate SIN-1-CH₂OH, then it was reacted with N-heterocycles to yield compounds 25–26.

Bioassay for anti-inflammatory activity *in vitro*

In this section, the LPS-induced RAW264.7 inflammatory cell model was used to evaluate the anti-inflammatory effects of compounds *in vitro*.⁴⁰ NO is an important endogenous free radical involved in a variety of physiological functions and pathophysiological states. Griess assay was used to evaluate the inhibitory effect of compounds 1–26 on the overproduction of NO. If the amount of NO decreases after treatment with compounds, the inflammatory response is considered to be suppressed and the corresponding compound is an effective anti-inflammatory agent.⁴¹

Before the anti-inflammatory activity assay, we screened the cytotoxicity of the tested compound to avoid false-positive results.⁴² The cytotoxicity of compounds 1–26 on RAW264.7 cells was evaluated by MTT assay. The IC₅₀ values of these compounds were summarized in Table 1. As shown, the



Table 1 The cytotoxicity of SIN and compounds in RAW264.7 cells

Compound	Cytotoxicity IC ₅₀ (μM)	Compound	Cytotoxicity IC ₅₀ (μM)
SIN	>100	13	>100
Indo	>100	14	>100
1	19.11 ± 2.43	15	17.88 ± 2.24
2	>100	16	>100
3	29.62 ± 7.48	17	>100
4	>100	18	>100
5	>100	19	>100
6	>100	20	21.88 ± 3.51
7	>100	21	>100
8	>100	22	52.69 ± 3.19
9	>100	23	>100
10	>100	24	>100
11	>100	25	54.87 ± 7.77
12	>100	26	>100

cytotoxicity IC₅₀ of SIN was >100 μM. Among them, the toxicity of compounds **1** (IC₅₀ = 19.11 ± 2.43 μM), **3** (IC₅₀ = 29.62 ± 7.48 μM), **15** (IC₅₀ = 17.88 ± 2.24 μM), **20** (IC₅₀ = 21.88 ± 3.51 μM), **22** (IC₅₀ = 52.69 ± 3.19 μM) and **25** (IC₅₀ = 54.87 ± 7.77 μM) was significantly increased compared to SIN. SARs shown that introduction of tetramethylpyrazine at C-4 could significantly improve the toxicity of SIN, but the modification of C-1 reduced the increase of toxicity. In addition, the cytotoxicity IC₅₀ of most of the compounds did not increase significantly and all exceeded 100 μM.

Then, LPS-induced RAW 264.7 cells were used to evaluate the inhibitory activity of all compounds **1–26** on NO production, and calculate the IC₅₀. As depicted in Table 2, the IC₅₀ of SIN was 70.86 ± 1.00 μM, indicating that the anti-inflammatory activity of SIN was relatively weak. The anti-inflammatory activity of most of the compounds increased after modification of amino acids and nitrogen-based heterocyclic ring compared to SIN. The elementary SARs of NO inhibitory activity indicated that introduction of protected amino acid (Boc group) at C-4 could enhance anti-inflammatory activities, and deprotect the Boc group could reduce anti-inflammatory efficacy. Among the nitrogen-containing heterocyclic compounds, the compound with pyrazine group showed stronger activity compared to these compounds with pyridazine and pyrimidine. While the effect of substituted halogen atoms at the C-1 position on the anti-inflammatory activity of the compounds was less. Among all the derivatives, compound **17** showed strong anti-inflammatory activity (IC₅₀ = 30.28 ± 1.70 μM), which was twice as strong as SIN, and it was worth to further investigate its mechanism of action and anti-inflammatory activity *in vivo*.

Effect of compound **17** on iNOS, IL-6 and TNF-α mRNA expression in LPS-induced RAW264.7 cells

The inflammatory response is generally regulated by a complex network of cytokines and chemokines; clinical interventions have clearly demonstrated that IL-6, TNF-α are essential in this process. IL-6 and TNF-α contributed to the exacerbation of the inflammatory response by activating endothelial cells and attracting immune cells to accumulate at the site of

Table 2 The inhibitory effects on NO production of SIN and compound **1–26** in LPS-induced RAW264.7 cells^a

Compound	Activity IC ₅₀ (μM)	Compound	Activity IC ₅₀ (μM)
SIN	70.86 ± 1.00	13	69.19 ± 9.02
Indo	44.67 ± 2.29	14	41.69 ± 3.66
1	NS	15	NS
2	42.27 ± 2.26	16	50.40 ± 4.62
3	NS	17	30.28 ± 1.70
4	51.38 ± 3.89	18	63.99 ± 1.32
5	44.11 ± 2.24	19	77.81 ± 6.04
6	59.50 ± 3.56	20	NS
7	59.28 ± 4.64	21	48.69 ± 2.06
8	68.87 ± 3.70	22	NS
9	53.36 ± 2.51	23	58.84 ± 5.59
10	72.27 ± 1.64	24	53.36 ± 4.64
11	52.77 ± 4.67	25	NS
12	60.40 ± 3.32	26	69.10 ± 3.76

^a NS: data not shown because activity IC₅₀ is higher than cytotoxicity IC₅₀.

inflammation. In addition, the excessive production of NO could affect the NF-κB signaling pathway, which in turn affects the expression of cytokines and chemokines including IL-6 and TNF-α.⁴³ In this section, qPCR was performed to qualification of genes including iNOS, IL-6, and TNF-α.

The effect of compound **17** with different concentrations on iNOS, IL-6 and TNF-α expression are shown in Fig. 2. Compared with the control group, the expression of iNOS mRNA was significantly increased in the model group after LPS induction, indicating that the inflammation model was successfully established. Compared with the model group, the compound **17** with high dose group (60 μM) could significantly inhibit the expression of iNOS mRNA. Compared with the model group, compound **17** with low dose group (15 μM), medium dose group (30 μM) and high dose group (60 μM) could significantly inhibit the expression of IL-6 mRNA. The treatment group inhibited TNF-α mRNA expression in a dose-dependent manner. Compared with the model group, compound **17** with the low dose group (15 μM) had no significant effect on the expression of TNF-α mRNA; the medium dose group (30 μM) could significantly reduce the expression of TNF-α mRNA; while the high dose group (60 μM) could inhibit the expression of TNF-α mRNA extremely significantly.

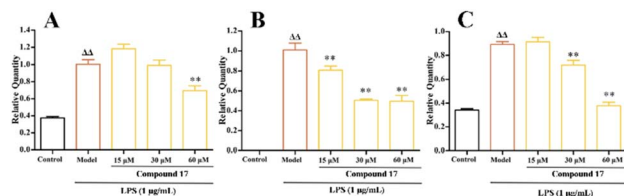


Fig. 2 Effect of compound **17** on LPS-induced iNOS, IL-6 and TNF-α mRNA expression in RAW264.7 cells ($\bar{x} \pm s$, $n = 3$). (A) qPCR results of iNOS mRNA levels. (B) qPCR results of IL-6 mRNA levels. (C) qPCR results of TNF-α mRNA levels. Compared with the control group, $\Delta\Delta P < 0.01$; compared with the model group, $**P < 0.01$.



In general, the RNA reduction of iNOS, IL-6 and TNF- α genes interprets the effects of compounds 17 on anti-inflammatory activity in RAW264.7 cells.

Bioassay for anti-inflammatory activity *in vivo*

Mouse paw edema method is widely used for the screening and evaluation of anti-inflammatory drugs due to its reliability and reproducibility.^{44,45} In this section, compound 17 with relative potent anti-inflammatory activity *in vitro* was selected and further evaluated with a λ -carrageenan-induced mouse paw edema test using the indomethacin as the positive drug. As shown in Fig. 3, after the injection of λ -carrageenan for 2 h in the model group mice, the paws showed obvious swelling, and the edema rate was around 60%.

The indomethacin group (10 mg kg⁻¹) significantly inhibited the development of edema at 2–6 h. Administration (i. g.) of compound 17 at both 25 and 50 mg kg⁻¹ doses significantly attenuated edema formation at 2–6 h. This suggested that compound 17 might be a potential anti-inflammatory agent with some *in vivo* anti-inflammatory activity.

The histological examination results were shown in Fig. 4. In the blank group, the articular cartilage was smooth and the subcutaneous tissue was dense without significant edema (Fig. 4A). After injection of λ -carrageenan, paw biopsies of control animals showed marked vasodilation and tissue edema. The subcutaneous tissue gap was enlarged, but articular cartilage was smooth and no lesions were observed (Fig. 4B). Paw biopsies of animals treated with indomethacin and compound 17 showed a reduction in the λ -carrageenan-induced inflammatory response. Among them, the compound 17 with high dose group (50 mg kg⁻¹) showed a reduction of edemas, and the subcutaneous tissue was dense (Fig. 4E).

The effects of compound 17 on the levels of TNF- α were shown in Fig. 5. No significant positive brownish-yellow staining was observed in the blank group, indicating less TNF- α expression (Fig. 5A). Compared with the blank group, the model group showed a large amount of brownish-yellow colouring at the articular cartilage, around the blood vessels and in

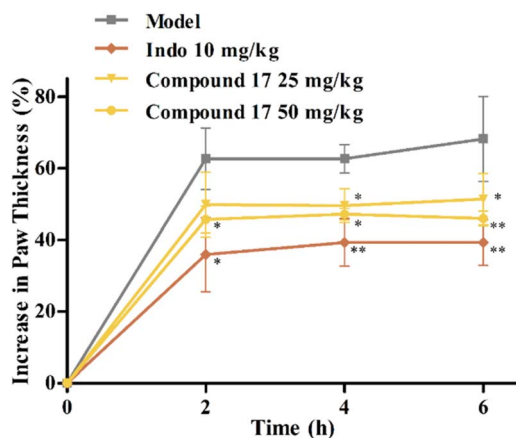


Fig. 3 *In vivo* anti-inflammatory activities of compound 17 ($\bar{x} \pm \text{SEM}$, $n = 8$). Compared with the model group, * $P < 0.05$, ** $P < 0.01$.

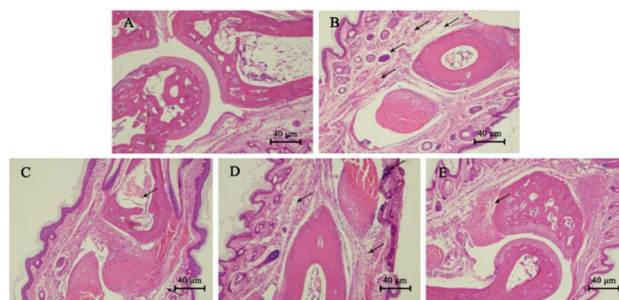


Fig. 4 Effects of compound 17 on morphological changes in paw edema tissue by hematoxylin–eosin staining. (A) Blank group. (B) Model group. (C) Indo group. (D) Compound low dose group (25 mg kg⁻¹). (E) Compound 17 high dose group (50 mg kg⁻¹).

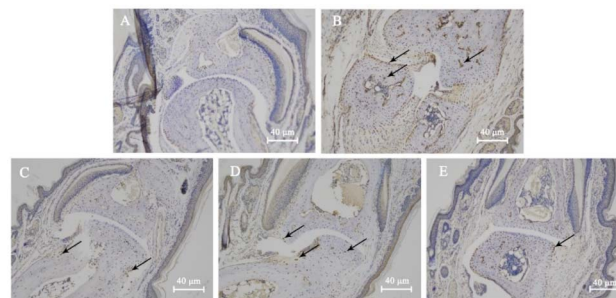


Fig. 5 Representative immunohistochemical staining for the levels of TNF- α . (A) Blank group. (B) Model group. (C) Indo group. (D) Compound low dose group (25 mg kg⁻¹). (E) Compound 17 high dose group (50 mg kg⁻¹).

the bone, indicating a large amount of TNF- α expression (Fig. 5B). The high dose group significantly inhibited TNF- α secretion around osteocytes compared with the model group, and only a small amount of positive coloration was seen at the articular cartilage (Fig. 5E), indicating that compound 17 significantly inhibited the expression of TNF- α inflammatory cytokines at the toe joint in mice. These results were also confirmed by the qPCR analysis (Fig. 2).

Bioassay for analgesia activity *in vivo*

The acetic acid-induced writhing test has been widely used for testing the analgesic effect.^{46,47} It is a non-selective model for analgesic studies, given that an intraperitoneal injection of acetic acid triggers the release of cytokines and chemokines. The effect of compound 17 on the writhing response in mice were shown in Fig. 6. The present result showed the positive control drug Voltaren (19.5 mg kg⁻¹) could reduce the number of writhes considerably. And the SIN at a dose of 200 mg kg⁻¹ and the compound 17 at a dose of 50 mg kg⁻¹ exhibited almost no analgesic effects. However, compound 17 at a dose of 100 and 200 mg kg⁻¹ could significantly decrease the number of writhes compared to that of the control group. Meanwhile, the analgesic activity of the compound 17 at a dose of 200 mg kg⁻¹ was stronger than that of Voltaren group. The results showed



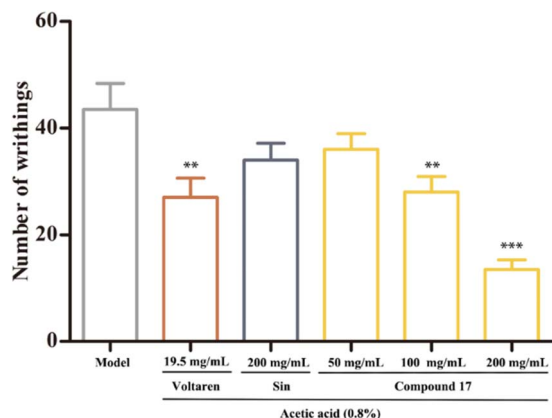


Fig. 6 Anti-nociceptive effects of compound 17 in mice. Compared with the model group, ** $P < 0.01$.

that compound 17 significantly reduced the number of writhes induced by acetic acid, suggesting that compound 17 has peripheral analgesic effects.

Molecular docking

NO is an important signal molecule in the body, which is produced by iNOS catalyzing *L*-arginine. iNOS is a carrier of gas-to-cell intercellular information transmission, which is activated in the inflammatory response, which increases iNOS expression and catalyzes NO production. NF- κ B is a classic inflammatory signaling pathway. Activation of NF- κ B stimulates the release of IL-6, TNF- α and NO when stimulated, which may worsen the inflammatory response.

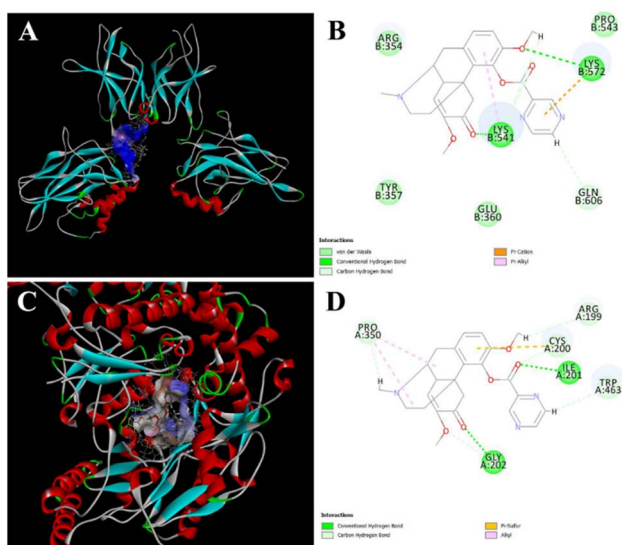


Fig. 7 Molecular docking. Predicted docking pose of compound 17 in complex with NF- κ B (PDB 1VKX) and iNOS (PDB 4NOS). (A) Overall structure of NF- κ B/compound 17 complex. The hydrophilic surface is coloured in blue. (B) Local view of residues interacting with NF- κ B and compound 17. (C) Overall structure of iNOS/compound 17 complex. The hydrophilic surface is coloured in blue. (D) Local view of residues interacting with iNOS and compound 17.

To better understand the interaction of compound 17 with NF- κ B and iNOS, CDOCKER was running by fitting compound 17 into the active site of NF- κ B and iNOS. As shown in Fig. 7A and B, compound 17 skeleton was placed in the active cavity of NF- κ B P65 formed by ARG 354, TYR 357, GLU 360, LYS 541, GLN 606, LYS 572, PRO 543 *etc.* van der Waals, hydrogen bonds, pi-cation and pi-alkyl interactions between compound 17 and NF- κ B protein further stabilized the interaction. As shown in Fig. 7C and D, compound 17 skeleton was placed in the active cavity of iNOS formed by PRO 350, GLY 202, TRP 463, ILE 201, CYS 200, ARG 199 *etc.* Hydrogen bonds, pi-sulfur and alkyl interactions between compound 17 and iNOS protein further stabilized the interaction.

Conclusions

Above all, a series of novel SIN derivatives were designed and synthesized, all of them were characterized by ^1H NMR, ^{13}C NMR and HR-MS. *In vitro* anti-inflammatory activity and cytotoxicity evaluation showed that most of the derivatives exhibited stronger anti-inflammatory activity compared to SIN, especially the compound 17 showed the most potent activity. Study of anti-inflammatory mechanism *in vitro* showed that compound 17 could inhibit the expression of iNOS mRNA, reduce the overproduction of NO and inhibit the expression of IL-6 and TNF- α mRNA. Experiments *in vivo* showed that compound 17 exhibited an ameliorating effect on λ -carrageenan induced paw oedema in mice. Pathological analysis and immunohistochemical results also indicated that it showed an ameliorating effect on subcutaneous edema of the mouse plantar and inhibited the expression of TNF- α inflammatory cytokines. The acetic acid-induced writhing test also demonstrated that compound 17 had good analgesic activity. Altogether, the results of the series of SIN derivatives supported the potential value of anti-inflammatory and had practical significance for the expansion of clinical applications of SIN.

Ethical approval

All animal procedures were performed in accordance with the Guidelines for Care and Use of Laboratory Animals of Beijing University of Chinese Medicine and experiments were approved by the Animal Ethics Committee of Beijing University of Chinese Medicine.

Author contributions

Feng Gao: conceptualization, methodology, validation, writing – original draft. Ziqi Dai: investigation, validation. Qianwen Wu: investigation. Desheng Cai: investigation. Mingjun Lu: data curation. Zijie Zhang: investigation. Qi Zeng: investigation. Tong Zhang: writing – reviewing and editing. Bingxian Shang: writing – reviewing and editing. Bing Xu: funding acquisition, supervision. Haimin Lei: resources, supervision, project administration, funding acquisition.



Conflicts of interest

There are no conflicts to declare.

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