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Synthesis and biological activities of local anesthetics

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Local anesthetics are mainly used in stomatology, ophthalmology, gynecology and surgery to temporarily relieve pain. Local anesthetics act on nerve endings or around nerve trunks, and are combined with specific sodium ion (Na⁺) channel sites on the nerve membrane. They can affect the membrane potential by reducing Na⁺ passage through sodium ion channels, thus blocking the generation and conduction of nerve impulses, reversibly blocking the generation and conduction of sensory nerve impulses. Local anesthetics are used for convenience in local surgical operations and treatments. Herein, we mainly review the research progress on local anesthetics and discuss the important aspects of design, synthesis and biological activity of various new compounds.

1. Introduction

Anesthetics are a type of drug that act on the nervous system, causing it to be inhibited and resulting in the loss of the sense of pain.^{1–4} According to the different scopes of action of drugs, the department of anesthetics is divided into general anesthetics and local anesthetics.^{5–7} General anesthetics act on the central nervous system, causing it to be reversibly inhibited, thereby resulting in a loss of consciousness and sensation. Especially pain sensation and reflexes disappear and skeletal muscle relaxes, thus achieving the effect of general anesthesia,

which is mainly used before surgery.^{8–11} In contrast, local anesthetics act on nerve endings or around nerve trunks, reversibly blocking the generation and conduction of sensory nerve impulses, and temporarily eliminating local sensation (mainly pain sensation) under the condition of consciousness. Moreover, it is convenient for local surgical operations and treatments.^{12–14} Local anesthetics are mainly used in stomatology, ophthalmology, gynecology and surgery to relieve pain temporarily.¹⁵ Local anesthetics combine with specific sodium ion (Na⁺) channel sites on the nerve membrane, and they can affect the membrane potential by reducing Na⁺ passage through the sodium ion channels, thus blocking the generation and conduction of nerve impulses.¹⁶ Local anesthetics decrease the excitability of nerve membranes, but have no effect on their resting potential.^{17–20} The application methods of local anesthesia can be divided into topical anesthesia, invasive anesthesia, conduction anesthesia, subarachnoid anesthesia and epidural anesthesia.^{21–24}

The first local anesthetic used was an active ingredient from coca leaves in South America (*Erythroxylum coca* Lam.). The Peruvians were known for chewing coca leaves to relieve pain as early as 1532. In 1860, Niemann extracted an alkaloid crystal from coca leaves and named it cocaine (1). Koller, an Austrian ophthalmologist, first formally used cocaine in clinical surgery in 1884. Also, he discovered a new local anesthesia technology, which led to the rapid development of local anesthetics.^{25–30} Cocaine has a local anesthesia effect and vasoconstriction effect, and it can reduce the bleeding of surgical wounds.^{31–34} Additionally, cocaine can be used in the nose, nasopharynx, mouth, throat and ear as surface anesthesia.^{35–39} However, cocaine is highly addictive, and easy to cause tissue irritation, allergic reactions and other toxic side effects. Also, its aqueous solution is not stable, and thus easily hydrolyzed during high-pressure disinfection; therefore, the clinical application of

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include the medicinal chemistry (drug design and synthesis, natural products semi-synthesis and full synthesis).



cocaine is greatly restricted, and it has long since been replaced by other local anesthetics.^{40–43} Because of the shortcomings of cocaine, researchers began to study and modify the structure of cocaine to find better local anesthetics. None of the three products from the hydrolysis of cocaine, ecgonine (2), benzoic acid and methanol, has the effect of local anesthesia.⁴⁴ Substitution of benzoic acid with other carboxylic acids can form esters with ecgonine, resulting in reduced or complete loss of anesthetic effects, proving that benzoic acid esters are an important structure of cocaine with local anesthetic effects. The alkaloid tropacocaine (3) isolated from the leaves of coca in Java had only the structure of benzoate ester and no carboxylate methyl ester group in its molecular structure, which had the general effect of local anesthesia. Therefore, the local anesthesia effect of benzoate ester was further confirmed.^{45–47}

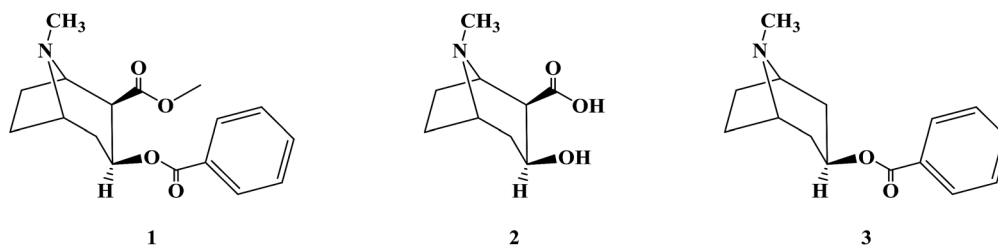
After determining the local anesthesia effect of benzoate ester, the structure of the parent nucleus of cocaine (ecgonine) was modified and simplified. When the *N*-methyl of ecgonine was removed, there was no significant effect on physiological activities, but the toxicity increased.⁴⁸ If the nitrogen atom was quaternized, the activity was lost. For ecgonine, Corning further simplified its structure, and designed and synthesized two piperidol benzoate derivatives, α -eucaine (4) and β -eucaine (5). Both of them had a local anesthetic effect similar to cocaine, but their aqueous solution property was relatively stable, and their toxicity was also low, which indicate that the ecgonine double ring structure is not necessary for the local anesthesia effect.^{49–53} Simultaneously, this further proves the importance of the benzoate ester part from the side. Due to the significant research work on benzoate ester compounds, benzocaine (6) was first synthesized in 1890 and proven to have a local anesthesia effect. Subsequently, orthoform (7) and new orthoform (8) were successfully synthesized. Both of them exhibited strong local anesthesia effects, but they were not water-soluble and could not be injected.^{54–56} They could only be used for local anesthesia, while their salt was too acidic to be used. Because of their shortcomings, further structural transformation led to the successful development of the local anesthetic procaine (9) in 1904, which overcame these shortcomings.

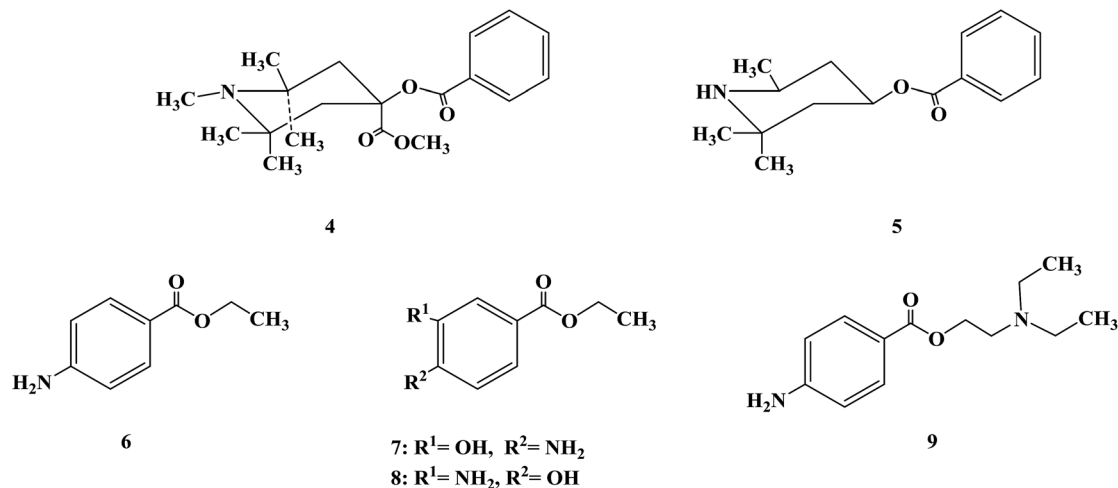
The discovery of procaine made researchers realize the important role of the *p*-aminobenzoate structure in local anesthetics.^{57–59} Upon further structural modification of procaine, aromatic acid esters were obtained as local anesthetics. Based

on the structure of aromatic acid esters, amide, aminoketone, aminoether and amino carbamate types of local anesthetics were developed (Table 1). In terms of drugs structure–activity relationship (SAR), the chemical structure of local anesthetics usually consists of three basic skeleton parts, a lipophilic aromatic ring, intermediate connecting functional group and hydrophilic amine group. In recent years, the local anesthetics used in clinical practice are mainly used to reduce toxicity and prolong the action time by changing the new dosage forms of existing local anesthetics.^{60,61} Herein, we mainly review the research progress of local anesthetics (2009–2019), including the design, synthesis and biological activity of various new compounds.

2. Synthesis and biological activities

Local anesthesia is a compound that blocks the conduction of nerve fibers when applied locally to nerve tissue at appropriate concentrations. It blocks the voltage-sensitive sodium channels of each nerve fiber, eliminating their action potential. The greatest practical advantage of local anesthetics is that their effects can be reversed when they were removed from nerve tissue. In this case, nerve function and action potential are completely restored, with no evidence of damage to nerve fibers or cellular structures. Compounds with amino groups linked to heterocyclic nuclei, such as lipophilic molecules, were shown to be more active and less toxic than benzene analogues. It has been reported that 2-aminobenzothiazole, 2-aminothiazole derivatives and 3-aminobenzothiazole-*d*-isothiazole derivatives have local anesthetic activity. In 2009, Carla F. *et al.*⁶² reported the design and synthesis of different thiazole and isothiazole derivatives for the study of local anesthesia. In the process of structural design of target compounds 1–32, they used the thiazole structure as the parent structure of the target compounds, and simultaneously retained the alkyl-acetamido or alkyl-propionamide groups that local anesthetic drugs have. In terms of structure–activity relationship (SAR), these target molecules were structurally similar to lidocaine, and thus they should be active in infiltration anesthesia and surface anesthesia. In the process of the synthesis of the target compounds, they used Scheme 1 as the synthetic route, and they chose the substituted thiazolamide as the starting material and dimethylformamide (DMF) as the reaction solvent to obtain





intermediate compounds 35–42. After further reaction with imine at high temperature in anhydrous benzene, the crude products of the target compounds were obtained and then crystallized with EtOH/Et₂O. All target compounds obtained good yields. The synthetic route had a few reaction steps, simple operation and easy reagents and raw materials. However, the synthetic route used benzene as the solvent, which will bring certain difficulties in the future industrial production. Thus, in the optimization of process conditions in the future, it is necessary to strengthen the in-depth study of solvents and try to choose low-toxic or non-toxic solvents. In the evaluation of local anesthetic activity, they chose the isolated sciatic nerve model to evaluate the local anesthetic effect of the target compounds. The results of local anesthesia screening *in vitro* showed that these compounds have a good local anesthesia effect, and the local anesthesia effect of some target compounds is better than that of the control product of lidocaine. To find the most promising target compounds, they also used the computer programs PASS and DEREK to predict the activity and toxicity of local anesthesia. The results of the QSAR study showed that the polarizability, polarity, hydrogen bond and molecular shape of the molecule had a positive and important effect on the local anesthetic activity of the target compounds. In addition, aromatic benzyl and single bond nitrogen also had a certain contribution to the local anesthetic activity of the target compounds. The data results of the local anesthesia model of frog and mouse showed that the calculated Pa of the local anesthesia effect of target compounds 5, 17, 27, 10 and 24 was in the range of 23–47% (the probability of passing was less than 50%), and thus these target compounds may become new local anesthetics, which need further research and development. Since these compounds are still in the process of *in vitro* screening, a follow-up study on their local anesthetic activity *in vivo* still needs to be carried out.

Two-thirds of the world's best-selling drugs are chiral compounds, and chiral drugs play an important role in drug discovery and drug therapy. Chiral drugs (enantiomers) differ in pharmacology, pharmacokinetics and toxicity, while single enantiomers have obvious advantages in the treatment of diseases. Therefore, the development of efficient methods for the synthesis of chiral compounds with high yield and enantiomer purity has become one of the important topics in the field of chemistry. Xanthone (9*H*-xanthone-9-one) derivatives are natural compounds with extensive biological and pharmacological activities, which have been proven important in the synthesis of new compounds. There are few reports on the synthesis of chiral derivatives despite their structural diversity and biological activity. Some chiral xanthone derivatives (CXDs) show good biological activity in the central nervous system (CNS). These compounds can both stimulate nerve activity and act as sedatives. Xanthone derivatives also have antiarrhythmic, anticonvulsive, antitumor, local anesthesia, antibacterial and antifungal effects. Some of these compounds have been shown to induce antihypertensive and vascular relaxant effects by inhibiting Ca²⁺ influx through receptor manipulation and/or voltage-sensitive channels. In 2012, Athina G. *et al.*⁶³ designed and synthesized three chiral xanthone derivatives. In the process of the synthesis of target compounds 7a–7c, they chose an efficient and convenient synthesis route, as shown in Scheme 2. Under the catalysis of *N,N*-dimethyl-glycine, CuI and Cs₂CO₃, the Ullmann condensation reaction between intermediate compound 2 and *m*-methoxyphenol resulted in the formation of diaromatic ether compound 3. The coupling reaction was carried out at 90 °C for 14 h, and the yield of compound 3 was good (54%). Compound 4 was obtained by hydrolyzing methyl ester in a mixture of methanol/tetrahydrofuran and sodium hydroxide solution. Compound 5 was obtained by the molecular acylation of diaryl ether compound 4 with phosphorus pentoxide and methanesulfonic



Table 1 The common types of local anesthetics

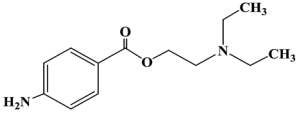
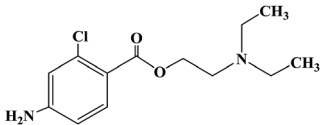
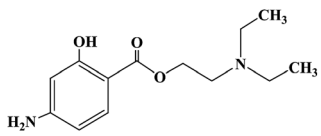
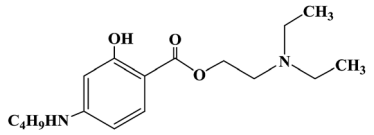
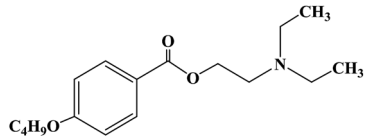
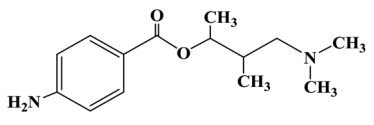
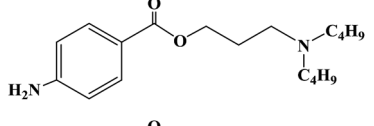
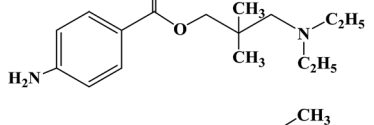
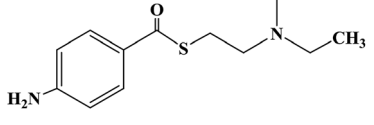
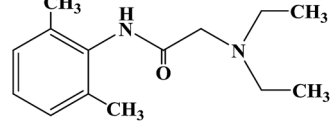
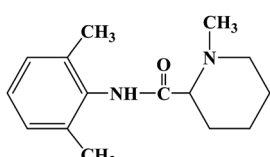
Compound	Structure	Time to market	Application methods
Procaine		1904	Infiltration anesthesia, conduction anesthesia, subarachnoid anesthesia and epidural anesthesia
Chloroprocaine		1952	Infiltration anesthesia, epidural anesthesia and conduction anesthesia
Hydroxyprocaine		1960	Infiltration anesthesia
Tetracaine		1988	Conduction anesthesia, subarachnoid anesthesia and epidural anesthesia
Oxybuprocaine		1975	Topical anesthesia
Tutocaine		1976	Topical anesthesia and infiltration anesthesia
Butacaine		1976	Topical anesthesia and infiltration anesthesia
Dimethocaine		1938	Topical anesthesia and infiltration anesthesia
Thiocaine		Halt sales	Topical anesthesia and infiltration anesthesia
Lidocaine		1948	Conduction anesthesia and epidural anesthesia
Mepivacaine		1986	Infiltration anesthesia, conduction anesthesia, epidural anesthesia and topical anesthesia



Table 1 (Contd.)

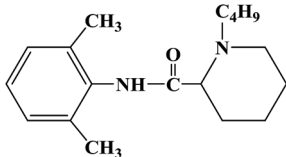
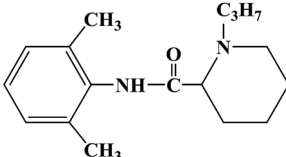
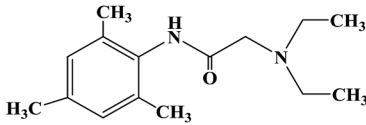
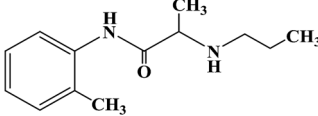
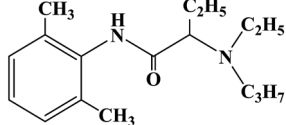
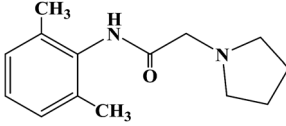
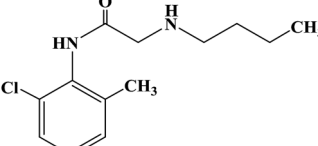
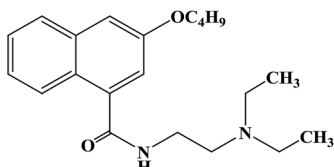
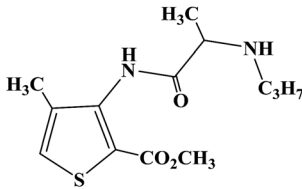
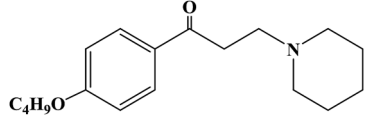
Compound	Structure	Time to market	Application methods
Bupivacaine		2000	Infiltration anesthesia, conduction anesthesia and epidural anesthesia
Ropivacaine		1996	Infiltration anesthesia, conduction anesthesia and epidural anesthesia
Trimecaine		1965	Infiltration anesthesia, surface anesthesia and epidural anesthesia
Prilocaine		1993	Infiltration anesthesia, topical anesthesia and epidural anesthesia
Etidocaine		1976	Epidural anesthesia
Pyrocaine		1964	Conduction anesthesia and epidural anesthesia
Butanilcaine		1982	Infiltration anesthesia and conduction anesthesia
Cinchocaine		1985	Topical anesthesia, subarachnoid anesthesia and epidural anesthesia
Articaine		2002	Infiltration anesthesia and subarachnoid anesthesia
Dyclonine		1956	Topical anesthesia

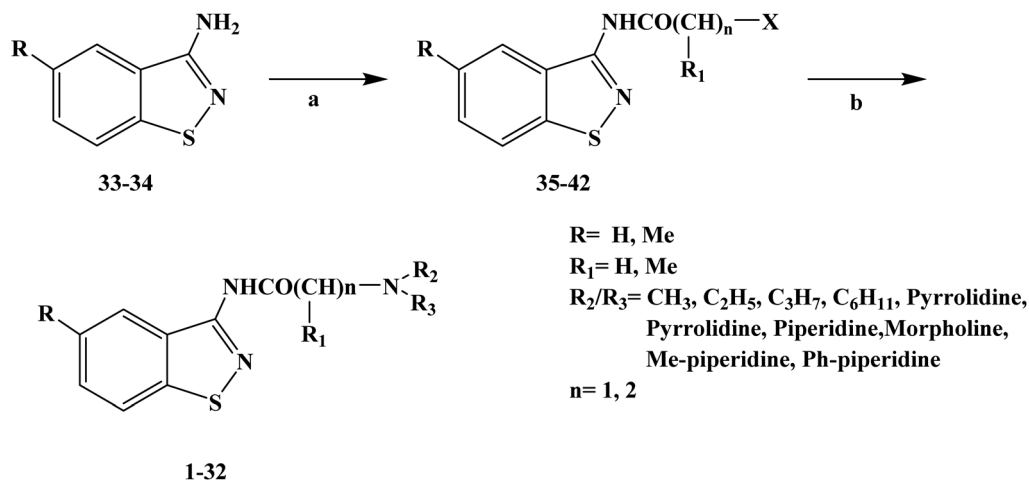


Table 1 (Contd.)

Compound	Structure	Time to market	Application methods
Fallicaine		1957	Topical anesthesia
Quinisocaine		1957	Topical anesthesia
Pramocaine		1977	Topical anesthesia
Diperodon		1980	Topical anesthesia
Heptacaine		1984	Infiltration anesthesia

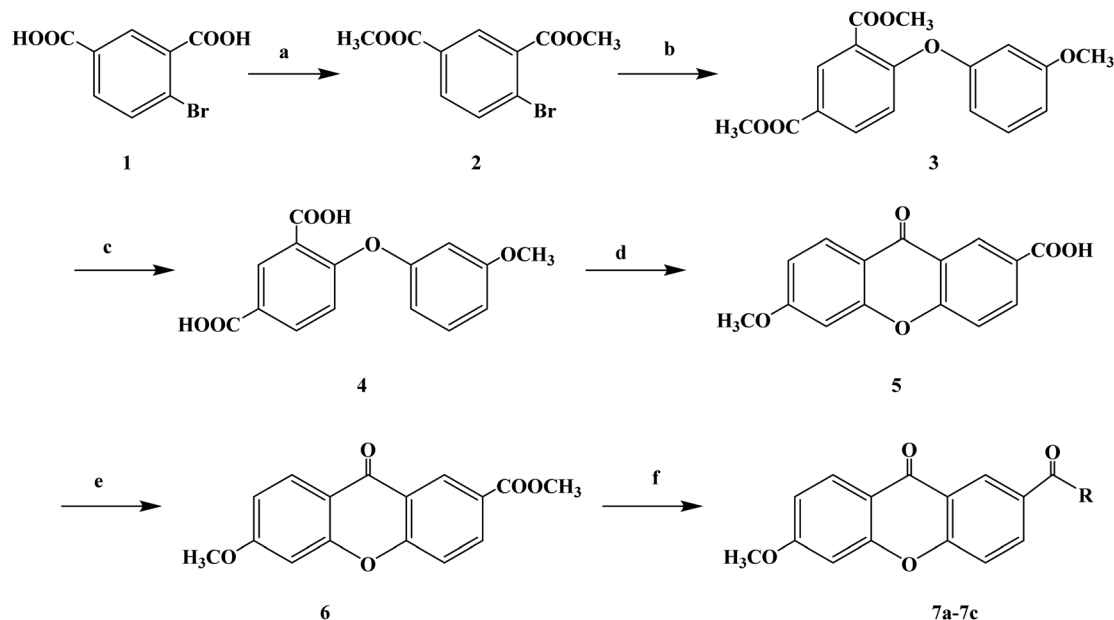
acid at room temperature. Finally, using TBTU and triethylamine as the catalyst, target compounds 7a–7c were obtained by reaction of tetrahydrofuran. The use of TBTU synthetic derivatives could be done at room temperature, resulting in high yield

(97%), in a short reaction time (30 min to 2 h), with the absence of racemization (enantiomeric excess was more than 99%). This synthetic route used cheap raw materials to synthesize the target compounds, which has the characteristics of mild



Scheme 1 Reagents and conditions: (a) ClCO(CHR)_nX, DMF and (b) NN(R), benzene.





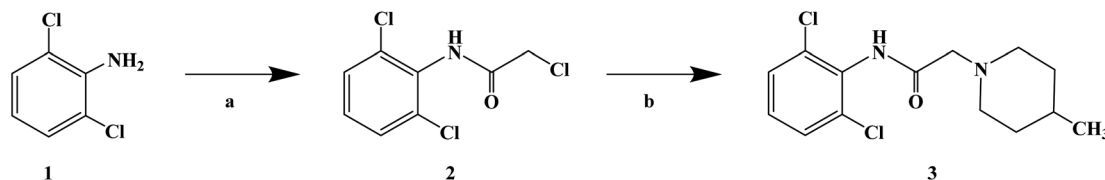
Scheme 2 Reagents and conditions: (a) methanol, H_2SO_4 , reflux, 17 h; (b) *m*-methoxyphenol, CuI, Cs_2CO_3 , *N,N*-dimethyl glycine, dioxane, N_2 , 90°C , 14 h; (c) methanol/tetrahydrofuran, NaOH, r.t., 18 h; (d) P_2O_5 , $\text{CH}_3\text{SO}_3\text{H}$, r.t., 22 h; (e) methanol, H_2SO_4 , reflux, 19 h; and (f) substitute amine, TBTU, triethylamine, dry tetrahydrofuran, r.t., 30 min to 2 h.

reaction temperature, relatively short reaction time and high overall yield. However, there are also many reaction steps, which bring some difficulties to the future process and need to be improved in the subsequent process conditions. The data from the local anesthesia activity study showed that when the concentration of CXDs (compounds **7a-7c**) was high, they could block the sciatic nerve conduction in rats, and had no obvious protective effect on hypotonic hemolysis. It could be seen that the nerve conduction block induced by CXDs was mainly caused by the sodium ion (Na^+) current. This effect could be isolated from their ability to stabilize the membrane, and became apparent only when the CXD (compounds **7a-7c**) concentration increased to the high micromolar range. The chiral xanthone derivatives compounds **7a-7c** are structurally similar to many local anesthetics, such as dibucaine, mepivacaine, lidocaine and bupivacaine, which may have the same molecular spatial structure and have the same pharmacophilia. The mechanism of nerve block induced by local anesthesia can be distinguished by its structure and by the way it crosses the cell membrane. Hydrophilic compounds diffuse to the axoplasm through open Na^+ channel pores, while lipid-soluble drugs can easily enter the Na^+ channel through the nerve membrane even if the (hydrophobic channel) is closed, achieving the effect of local anesthesia. The ability of compounds **7a-7c** to prevent hypotonic hemolysis was also evaluated, and the results showed that there were no excited erythrocytes and no voltage-gated sodium ion channels. In general, these chiral derivatives show good activity in local anesthesia, which need further research and development.

Sensory-neuron voltage-gated sodium (Na^+) channels play an important role in chronic painful neuropathy caused by peripheral nerve injury. Local anesthetics can block sodium

(Na^+) channels and effectively treat neuropathic pain and diabetic neuropathy. Lidocaine, a local anesthetic and antiarrhythmic agent, reduces pain by blocking voltage-gated sodium (Na^+) channels in sensory neurons around and at the central level of the spinal cord. Lidocaine is widely used clinically, but there is also a risk of nerve damage, particularly in patients with diabetic peripheral neuropathy, where neurotoxicity is more common. In 2013, Lilianna G. *et al.*⁶⁴ using lidocaine as the lead compound, designed and synthesized the lidocaine analogue *N*-(2,6-dichlorophenyl)-2-(4-methyl-1-piperidine) acetamide, which has less side effects, and evaluated its effect as local anesthesia. In the structural design of the target compound, lidocaine was modified through the bioisosteric principle, and the methyl group on the benzene ring in lidocaine was replaced by chlorine. This structural design improved the half-life and lipid solubility, and reduced the oxidation metabolism of the drug. The use of the cyclic analogue 4-methyl-piperidine instead of alkyl diethylamine generally did not change the pharmacokinetic effects such as lipophilicity and metabolism of the drug. The synthetic process shown in Scheme 3 was used to synthesize target compound **3**. Initially, sodium bicarbonate (NaHCO_3) was applied under the condition of acid and acetone as the solvent, and 2,6-dichloro (compound **1**) and 2-chloro aniline acetyl chloride were used as the raw materials, which were reacted at 40°C for 6 h to give the intermediate α -chlorine acetamide (compound **2**). Finally, compound **2** and 4-methyl piperidine were reacted for 4 h at room temperature to synthesize target compound **3**. The synthetic route was characterized by high yield (86.6%), simple operation and cheap reagents, thus reducing the synthesis cost. In the activity study, the formalin test was used to determine the safe activity of the compounds in the body of laboratory mice, which were used to





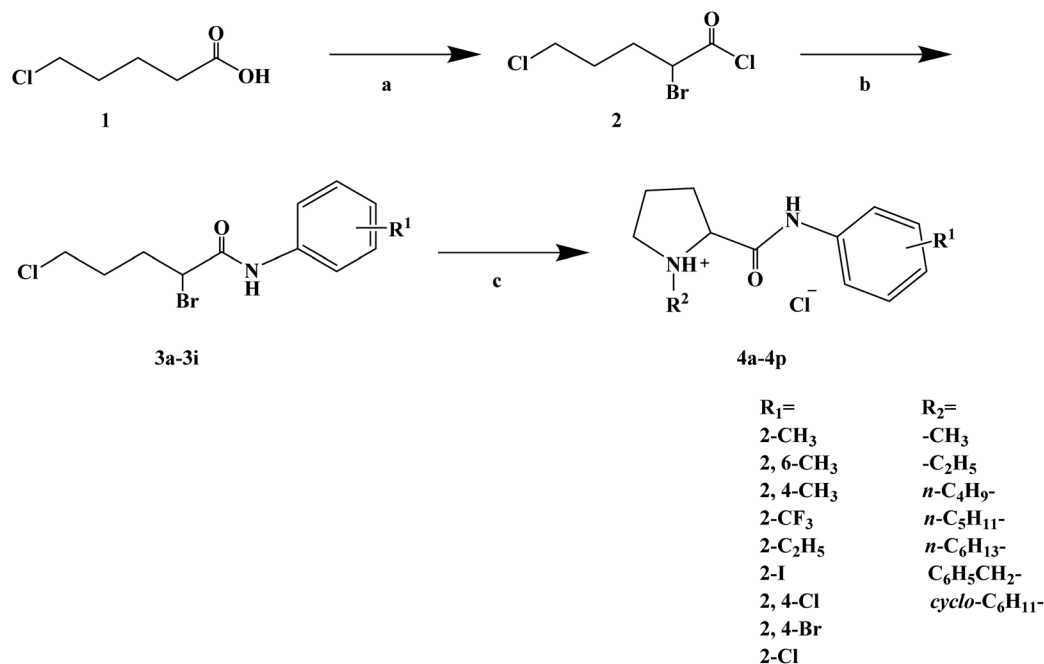
Scheme 3 Reagents and conditions: (a) 2-chloroacetyl chloride, NaHCO₃, acetone, 40 °C, 6 h; and (b) 4-methylpiperidine, r.t., stirring, 4 h.

evaluate the effect of local anesthesia. Two stages were used in the evaluation process. The first stage was acute pain response, and the second stage was the increase of primary afferent fiber input after tissue injury caused by chemical stimulation and local inflammatory reaction. The first stage of acute pain response began immediately after formalin management and rejection for more than 10 min. The second stage began about 15 min after formalin administration and lasted for about 1 h. A increased pain response in diabetic rats was observed with a lower dose of formalin than in the non-diabetic rats. Although 1% formalin had a higher harmful reaction to the non-diabetic rats, 0.5% formalin had no significant difference to the non-diabetic rats. This finding suggests that hyperalgesia in diabetic rats is caused by a sensitization process that changes both their peripheral and central levels. These results were consistent with previous observations using the same formalin concentration in the same model. The results showed that compound 3 has an anti-pain hypersensitivity effect because it could reduce the pain hypersensitivity behavior of diabetic rats. Although the mechanism of action of this compound is unknown, its mechanism may be similar to lidocaine due to its similar structure.

During the development of new effective local anesthetics, the *N,N*-dialkyl derivatives of glycine gradually evolved into *N*-alkyl substituted isoproline and proline. These new local anesthetics have the biological activity of arrhythmia in addition to the effect of local anesthesia. Studies have shown that the local anesthesia and antiarrhythmic action mechanism is the same, mainly by blocking voltage-gated sodium channels, calcium channels and potassium channels, resulting in membrane stability to prevent the initiation of action potential. In the process of drug development, the development of local anesthetics and antiarrhythmic drugs is usually studied and developed simultaneously. In 2013, Kalinin D. V. *et al.*⁶⁵ reported the synthesis and biological activity of *N*-alkyl derivatives of proline aniline. In terms of synthesis, the designed synthetic route focused on two key biological activity substitutions of target compounds **4a–4p**, and the substitutions of the tetrahydropyrrole ring and benzene ring (Scheme 4). The bromination of 5-chloropentanoic acid (compound **1**) was accomplished by the action of bromine with phosphorus trichloride to produce 2-bromo-5-chloropentanoic acid. The chloropentanoic acid was then converted into the corresponding acyl chloride by SOCl₂. Compound **2** was treated with anilines with different substituents under the condition of chloroform as the solvent to synthesize 2-bromo-5-chloro-*N*-aryl amide (compound **3**). The *N*-alkyl proline aryl amines underwent an intramolecular

cyclization reaction after compound **3** was heated in toluene, which were further converted into the corresponding hydrochloride compounds **4a–4p** under the condition of HCl. The synthetic route has mild reaction conditions, relatively simple operation, and the reagents used are cheap and easy to obtain. Simultaneously, the synthetic route does not use precious metal catalysts, and the total yield varies from 37% to 75%, but the total yield of most of the target compounds is not high, which needs to be optimized *via* the reaction conditions to improve the total yield, reduce the cost of synthesis, and provide favorable conditions for the realization of industrialization. In terms of biological activity, all the synthetic target compounds (**4a–4p**) were evaluated as local anesthesia *in vitro* and *in vivo*. The primary selection of the local anesthetic activity of the target compounds was to evaluate the biological activity of rabbit cornea by local anesthetic effect of a common surface anesthesia model. Further studies were conducted to evaluate the compounds with the best local anesthetic activity using a rat infiltration anesthesia model. The results of the structural–activity relationship (SAR) analysis showed that the substituents on the aromatic local anesthetics usually included one *o*-methyl group, sometimes two *o*-methyl groups or even *p*-methyl group. Among the synthesized compounds, the best biological activity was observed when the substituents on the aromatic ring were *o*-methyl. When the number of carbon atoms in alkyl chain R¹ increased from methyl to ethyl, the surface anesthetic effect of the compounds was significantly reduced. Simultaneously, the number of substituents on the aromatic ring also affected the local anesthesia effect. With an increase in the number of methyl groups, the intensity and duration of the local anesthesia effect of the compound increased. The intensity and duration of the surface anesthetic effect of the compounds were significantly reduced by changing the substituent methyl group on the aromatic ring to trifluoromethyl or halogen. The *N*-alkyl substituents R² in the pyrrolidone ring also play an important role in the surface local anesthetic activity. Accordingly, –CH₃, –CH₂CH₃, *n*-C₄H₉, *n*-C₅H₁₁, *n*-C₆H₁₃, benzyl, and cyclohexyl were investigated. In the surface anesthesia experiments, when the substituents were methyl and ethyl, the compounds had no anesthetic effect. With an increase in the carbon chain length, the duration of the surface anesthesia was significantly prolonged, and the local anesthesia effect of the cyclohexyl substituted group was similar to that of the *N*-butyl substituted group. In local anesthesia tests *in vivo*, all the compounds worked quickly within 1–2 minutes of administration, blocking muscle contractions. The anesthesia time range of target compounds **4c**, **4d**, **4l** and **4o** was 35.0 ± 4.1 to 126.3 ± 24.3 min,





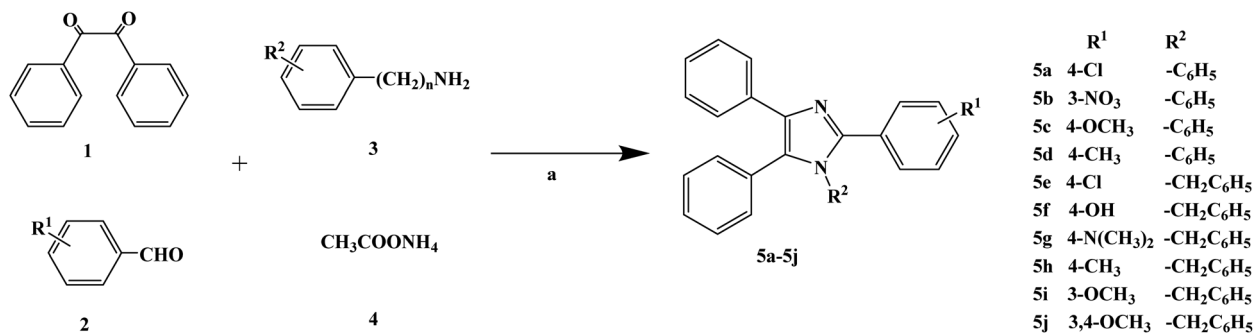
Scheme 4 Reagents and conditions: (a) (i) Br₂, PCl₃, 80 °C, 20 h; (ii) SOCl₂, reflux, 1.5 h; (b) substituted anilines, CHCl₃, Et₃N, reflux, 1 h; and (c) (i) RNH₂, KI, toluene, 6 h; (ii) HCl, CH₃CH₂OCH₂CH₃.

which was similar to the control drugs. However, the total anesthesia time of compound **4c** was shorter than that of the positive control substance (12.5 ± 2.9 min), and the complete prevention of stimulated muscle contraction was 102.5 ± 37.9 min and 58.8 ± 17.5 min, respectively. The local anesthesia time of compound **4d** was longer, where the maximum block time was 20.0 ± 13.5 min, and the recovery was slow after anesthesia. Compounds **4l** and **4o** were basically the same in intensity and duration of surface local anesthesia, but compounds **4d**, **4l** and **4o** showed obvious characteristics in the invasive anesthesia test. The duration of compound **4l** anesthesia was 115.0 ± 12.2 min, while the duration of infiltration anesthesia induced by compound **4o** was longer (126.3 ± 24.3 min). In general, compounds **4d**, **4l** and **4o** were more active in inducing surface anesthesia than the positive controls. Meanwhile, the toxicity of compound **4o** was significantly reduced, which make the compounds suitable for further research and development.

Heterocyclic compounds have the advantages of easy structural modification, structural diversity and good synthesis selectivity, and thus they are often used as preferred compounds in the research and development of new drugs. There are many types of heterocyclic compounds, among which those containing the imidazole structure are the most attractive to drug chemists. This is because compounds containing an imidazole structure have a wide range of medicinal value with different pharmacological effects, such as anti-malarial, anti-histamine, anti-bacterial, anti-inflammatory, anti-tuberculosis, analgesia, anti-insect drugs and anti-genic animal effects. In 2015, Yan R. *et al.*⁶⁶ reported the β -cyclodextrin-propyl sulfonic acid (β -CD-PSA) catalysed one-pot synthesis of 1,2,4,5-

tetrasubstituted imidazoles as local anesthetic drugs. They designed a total of 10 target compounds (compounds **5a–5j**). They used R¹ instead of a benzene ring and R² instead of the imidazoles to get 1,2,4,5-tetrasubstituted imidazoles (compounds **5a–5j**). Efficient synthetic methods for imidazole compounds have been widely studied. In the synthesis of 1,2,4,5-tetrasubstituted imidazoles compounds, usually with 1,2-diketone, hydroxy ketones or aldehydes ketonoxime, the primary amine and ammonium acetate group are divided into four synthetic raw materials, with the use catalysts of HY zeolite, silicone/NaHSO₄, BF₃-SiO₂, molecular iodine, HClO₄-SiO₂, InCl₃·3H₂O or potassium dodecatungstocobaltatetratehydrate (K₅-CoW₁₂O₄₀·3H₂O) and functionalized silica (SiO₂-Pr-SO₃H). 1,2,4,5-Tetranomeric imidazole compounds are synthesized *via* a one-pot method with or without solvent. However, currently, the methods for synthesizing 1,2,4,5-tetrasubstituted imidazoles compounds have disadvantages such as long reaction time, low product yield, harsh reaction conditions, cumbersome processes and the use of toxic reagents. Yan R. *et al.* provided a green and economic synthetic method to obtain 1,2,4,5-tetrasubstituted imidazoles compounds, which used β -CD-PSA as a catalyst, as shown in the synthetic route in Scheme 5. In synthetic process, firstly the catalyst β -CD-PSA required for the reaction was synthesized under laboratory conditions, and then the conditions for the synthesis of the target compounds were optimized. The final selection for the one-pot synthetic method involved benzyl (compound **1**), aniline or benzylamine (compound **2**), aromatic aldehyde (compound **3**) and ammonium acetate (compound **4**) as raw materials, β -CD-PSA as the catalyst, reaction temperature of 100 °C under solvent-free conditions, and stirring for 15 min to 4 h, with the total yield





Scheme 5 Reagents and conditions: (a) solvent free, catalysed by β -CD-PSA, 100 °C, 15 min to 4 h.

of the final target compounds of 68–96%. The synthetic route has the advantages of simple operation, high total yield, no solvent and no pollution gas product, and it is environmentally friendly. Simultaneously, the catalyst β -CD-PSA used in the reaction process can be recycled and reused, and the recovery operation is simple and the recovery rate is high. In the study of the local anesthetic activity, they measured the local anesthetic activity of the target compounds by corneal anesthesia and mouse tail anesthesia. Firstly, two types of anesthesia models, surface anesthesia and infiltration anesthesia, were used to evaluate the local anesthesia activity of the target compounds *in vitro*. The results of local anesthetic activity *in vitro* showed that compounds **5g** and **5i** had a good local anesthetic effect (Table 2), while the other compounds had no significant local anesthetic effect. Among them, the local anesthetic effect of compound **5g** was better than the positive control product lidocaine. From the analysis of the drug structural activity relationship (SAR), the presence of the benzylamine group has a significant effect on the local anesthetic effect of the target compounds, and aniline replaces benzylamine to eliminate the local anesthetic activity of the target compounds. Thus, the duration of the local anesthetic activity of compounds **5g**, **5i** and **5h** was further determined by rat sciatic nerve block. The test results showed that the local anesthesia duration of compound **5g** was similar to that of the positive control product lidocaine. In the study of local anesthetic activity *in vivo*, the acute toxicity and mouse therapeutic index test of compound **5g** were

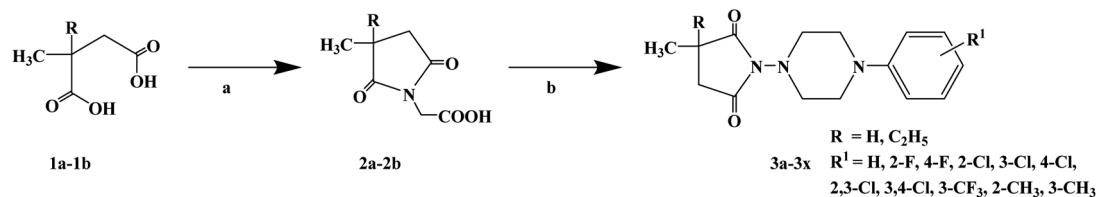
investigated. The results showed that the acute toxicity of the compound **5g** was less than that of the positive control product lidocaine. In terms of therapeutic index, that of compound **5g** was significantly higher than that of the positive control product lidocaine. The local anesthetic activities of the target compounds were screened *via* various methods and the results showed that these compounds are potential the local anesthetics. Combined with its effective synthetic method and obvious the local anesthetic activity, compound **5g** may be a potential lead compound for further drug development.

To determine the chemical structure characteristics of anti-convulsants, after decades of continuous research and numerous attempts, the basic chemical structure of anticonvulsants and nitrogen heterocycles containing an imine structure was finally determined, and it was found that there are usually carbonyl and phenyl or alkyl substituents in the structure of nitrogen heterocycles. In 2016, Jolanta O. *et al.*⁶⁷ reported the design and synthesis of a series of 3-methyl-3-methyl-2,5-dioxo-pyrrolidin-1-yl-acetic acids and 3-ethyl-3-methyl-2,5-dioxo-pyrrolidin-1-yl-acetic acids as anticonvulsants. In the structural design, they introduced a methyl or ethyl and methyl substituent at the imine ring 3 site to evaluate the effect of the substituent on activity. These compounds were originally used as anticonvulsants, but some convulsants have analgesic effects in clinical practice. Therefore, some compounds have local anesthetic effects in subsequent studies. The synthetic process for the target compounds is shown in Scheme 6. Starting with 3-

Table 2 The local anesthetic activity of target compounds 5a–5j

Compound	Corneal anesthesia (%)	Mouse tail anesthesia IC ₅₀ (mol L ⁻¹)
5a	20.3 ± 8.9	(11.3 ± 0.11) × 10 ⁻²
5b	12.4 ± 9.3	(5.7 ± 0.42) × 10 ⁻²
5c	15.5 ± 12.3	(6.5 ± 0.23) × 10 ⁻²
5d	10.32 ± 0.12	(4.3 ± 0.54) × 10 ⁻²
5e	21.65 ± 4.7	(3.4 ± 0.32) × 10 ⁻²
5f	5.5 ± 3.5	(5.4 ± 0.14) × 10 ⁻²
5g	121.23 ± 12.3	(1.6 ± 0.26) × 10 ⁻²
5h	65.65 ± 5.8	(9.2 ± 0.16) × 10 ⁻²
5i	95.6 ± 13.2	(7.4 ± 0.18) × 10 ⁻²
5j	20.3 ± 10.3	(4.1 ± 0.45) × 10 ⁻²
Lidocaine	100	(2.1 ± 0.25) × 10 ⁻²





Scheme 6 Reagents and conditions: (a) glycine, 190–200 °C, 1.5 h; (b) arylpiperazine, CDI, DMF, r.t., 24 h.

methyl succinic acid (compound **1a**) and 3-ethyl-3-methyl succinic acid (compound **1b**), the intermediates 3-methyl-3-methylsuccinic acid (compound **2a**) and 3-ethyl-3-methylsuccinic acid (compound **2b**) were synthesized by condensation reaction with glycine at 190–200 °C for 1.5 h. Compound **2a** and compound **2b** obtained from the first step were again reacted with arylpiperazine. Target compounds **3a-3x** were obtained using dimethylformamide (DMF) as the solvent and *N,N*-carbonyldiimidazole (CDI) as a dehydrator, with stirring at room temperature, giving yields ranging from 50% to 80%. This synthetic route has only a few reaction steps, the suitable solvent is cheap and easy to obtain, and the total yield is relatively high. However, the reaction temperature of the first step is relatively high, and a large amount of energy is required under high temperature conditions. Moreover, the reactor requirements are also relatively high, and the risk factor is relatively high. Local anesthetics block nerve conduction by altering the function of voltage-gated sodium channels and are used as adjunctive analgesics in the treatment of neuropathic pain. Compounds **3g**, **3i**, **3j**, **3l**, **3r** and **3x** were selected to test the effect of local anesthesia by the tail immersion test (Table 3). During the test, compounds **3j** and **3r** at a concentration of 2% significantly prolonged the response time of the animals to thermal stimulation (124% and 61%), while that of the positive control substance was 84%. Meanwhile, the binding ability of compounds **3g**, **3i**, **3j**, **3l**, **3r** and **3x** to sodium channels and calcium channels (Na⁺ and Ca²⁺ channels) was evaluated *in vitro* (Table 4). The test results showed that compound **3x** had the best binding effect on Na⁺ and Ca²⁺ channels, with a binding capacity of 63.5 and 82.4 (positive control substances of 53.9 and 57.8), respectively. Thus, compound **3x** may be a new local anesthetic, which needs to be further studied.

Local anesthetics can cause a reversible loss of sensation in a specific area of the body and control acute and chronic pain

during surgery. However, the use of local anesthetics in medical practice also has serious side effects, leading to the need for safe and effective local anesthetics. The literature shows that benzimidazole, amide, benzotriazole, acetamide and other compounds can effectively inhibit sodium (Na⁺) channels and have obvious local anesthesia and antiarrhythmic activity. The chloroacetamido moiety also exists in many compounds and plays an important role in biological activities. In 2017, Ashish A. *et al.*⁶⁸ reported a series of 4-(2-chloroacetamido) benzoic acid derivatives, including 9 amide compounds, 1 imidazole compound and 8 ester compounds derivative (Scheme 7). A total of 19 target compounds were obtained (**4a-4i**, **5a**, and **6a-h**), and their local anesthetic activities were further studied. In the process of structurally designing the target compounds, they investigated relevant literature and found that many commercial local anesthetics contained the 4-aminobenzoic acid structure, in which the presence of an ester or amide bond is a key feature. Thus, they also chose 4-aminobenzoic acid as the parent structure. In the structural modification of the parent group, alkyl groups with strong ionization properties were replaced by amides, and carbonyl groups were placed next to the amides to enable them to enter the active pocket. A chloromethamphetamine was substituted on the carbonyl carbon of the amide. The presence of a chlorine atom will increase the ionization property of the amide through the induction effect, thus increasing its solubility in water. In the process of synthesizing the target compounds, the 4-aminobenzoic acid was synthesized by Fischer's esterification, Schotten-Baumann and substitution reactions. The 4-aminobenzoic acid (compound **1**) and chloroacetyl chloride were treated with 35% sodium hydroxide solution for 10 min at a reaction temperature of 4–10 °C. The hydrochloric acid was then acidified to obtain the first intermediate 4-(2-chloroacetamido) benzoic acid (compound **2**). Then compound **2** and thionyl chloride were

Table 3 The local anesthetic activity in the tail immersion test

Compound	Concentration (%)	Reaction time to painful stimulus (s)	Prolongation of the time reaction (%)
3g	2	5.41 ± 0.53	12
3i	2	5.66 ± 0.60	17
3j	2	10.80 ± 1.74	124
3l	2	5.53 ± 1.06	14
3r	2	7.79 ± 0.95	61
3x	2	7.06 ± 0.76	46
Phenytoin	2	8.90 ± 0.80	84

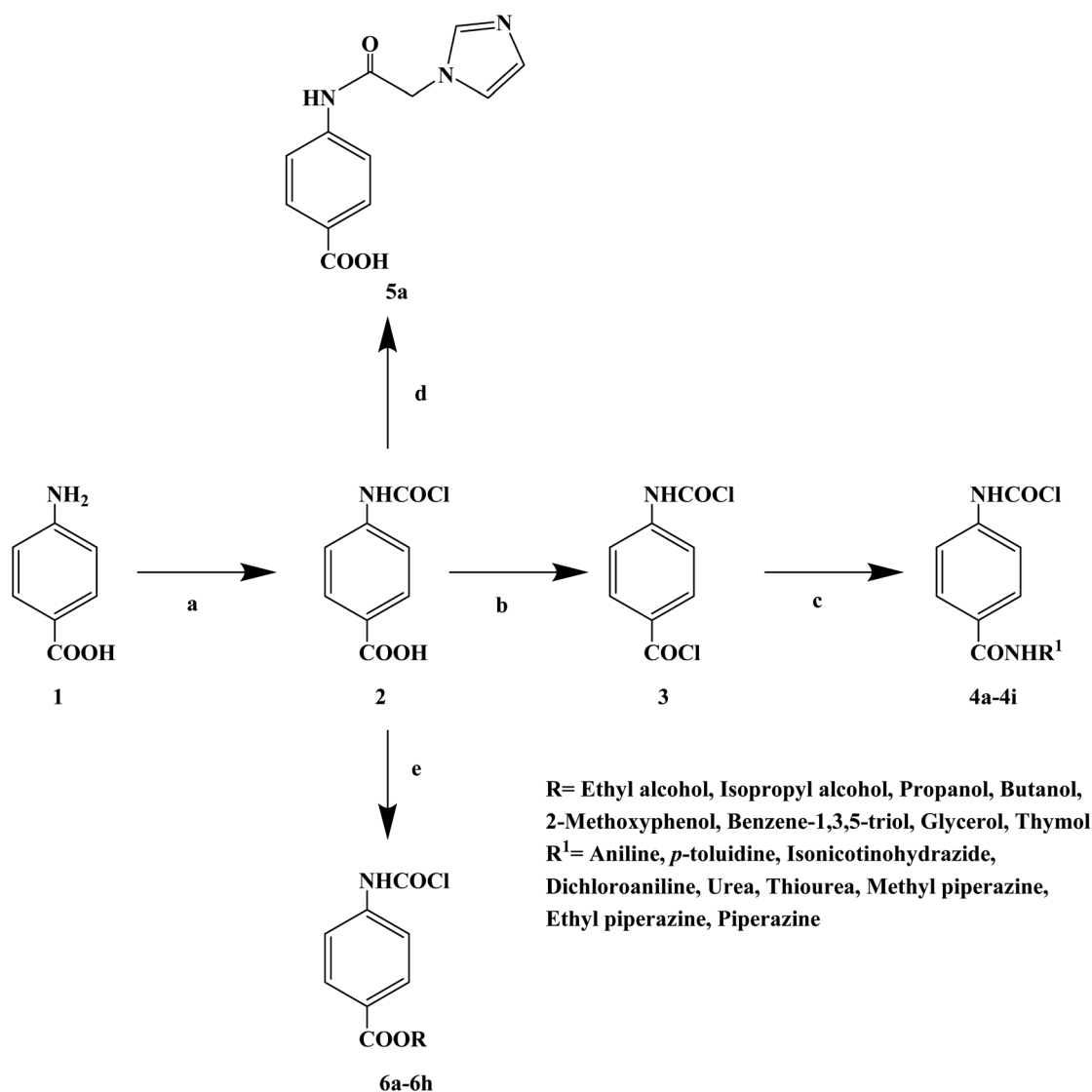


Table 4 *In vitro* binding assay results at a concentration 100 μ M

Compound	Inhibition of control specific binding (%)	
	Voltage-sensitive Na ⁺ channels	Voltage-sensitive L-type Ca ²⁺ channels
3g	36.6	13.5
3i	28.5	17.2
3j	20.9	4.2
3l	33.4	71.0
3r	32.5	30.8
3x	63.5	82.4
Phenytoin	53.9	57.8

continuously stirred in pyridine to form a second intermediate 4-(2-chloroacetamido)benzoyl chloride (compound 3). Then the corresponding amine dissolved in pyridine was added and vigorously mixed with compound 3, and after completion of the

reaction with sulfuric acid treatment, 4-(2-chloroacetamido)-*N*-substituted benzamide derivatives were obtained (compounds 4a–4i). Compound 2 and imidazole were treated in dimethyl sulfoxide (DMSO). The mixture was heated at 60 °C for 5 h and substituted for chloride ions in the presence of sodium carbonate to form imidazole derivatives (compound 5a). Compound 1 and the corresponding alcohols were heated in the presence of hydrochloric or sulfuric acid and refluxed for 4–6 h to obtain the corresponding esters (compounds 6a–6h). The synthetic route has the advantages of short reaction time, cheap reagents and mild reaction conditions. However, the total yield of some target compounds in this synthetic route was low (13%), which needs to be optimized to improve the total yield. The activity of the invasive local anesthesia was determined by Bianchi's method. The results of infiltration local anesthesia showed that compounds 4b, 5a, 6a, 6b and 6d showed good local anesthetic activity, which was close to the local anesthetic effect of lidocaine hydrochloride. Analysis of the structure–



Scheme 7 Reagents and conditions: (a) chloroacetyl chloride, HCl, 4–10 °C, stirring, 10 min; (b) thionyl chloride, pyridine, 0–4 °C, stirring; (c) corresponding amide derivatives, conc. H₂SO₄, reflux; (d) imidazole, AlCl₃, reflux; and (e) corresponding alcoholic derivative, conc. H₂SO₄, reflux.



Table 5 Infiltration local anesthetic activity ($n = 6$)

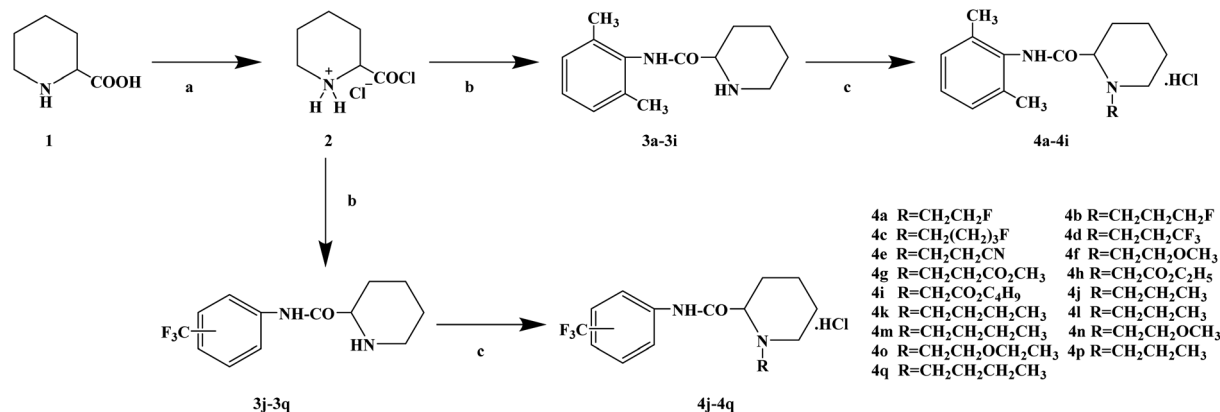
Compound	Enduring time at 15 min (s)	Enduring time at 30 min (s)	Enduring time at 45 min (s)	Mean enduring time (s)	%
4b	82.5 ± 1.66	75 ± 1.89	75 ± 2.08	77.5	61
5a	109 ± 1.41	102 ± 1.18	94.5 ± 1.26	101.8	81.44
6a	108.5 ± 0.04	100.5 ± 0.19	100.5 ± 0.17	103.2	81.87
6b	90 ± 0.56	71.5 ± 0.81	71.5 ± 0.77	77.7	61.76
6d	103 ± 1.47	96.5 ± 1.84	96.5 ± 1.66	98.7	78.45
Lignocaine	143.5 ± 2.11	117 ± 2.70	117 ± 2.91	125.8	100

activity relationship (SAR) results showed that compounds **4a–4i** were amide derivatives of aromatic and aliphatic amines such as aniline, *p*-toluidine, dichloroaniline, urea, thiourea, methyl piperazine, ethyl piperazine and piperazine. The amides of aniline compound **4a** and *p*-toluidine compound **4b** showed significant local anesthetic effects. The other compounds **4c–4i** exhibited a very low local anesthetic effect because both amide nitrogens were ionizable and easily dissolve in water, thus losing the local anesthetic effect. The imidazole derivative compound **5a** exhibited significant local anesthetic activity, which has all the basic chemical structures of local anesthetics and its activity was 81%. Compounds **6a–6d** were ester derivatives of ethyl, isopropyl, propyl and butanol, respectively, with a single ionizing group and their activity was between 60–80%. Compounds **6e–6h** are phenolic ester compounds composed of 2-methoxyphenol, phloroglucinol, glycerol and thymol, respectively, with relatively low local anesthetic activity. *In vivo*, the rat sciatic nerve block method was used to further study the infiltration local anesthesia activities of compounds **4b**, **5a**, **6a**, **6b** and **6d** (Table 5). The experimental results showed that compounds **4b**, **5a**, **6a**, **6b** and **6d** were effective, among which compound **5a** showed obvious infiltration local anesthetic activity. The effect of compound **5a** on blocking the sciatic nerve in rats was better than that of lidocaine (53 min), and that of lidocaine was 27 min. The activity of compound **4b** was lower than that of the control (11 min). In general, compound **5a** is a new local anesthetic, which has the potential for further research and development.

Ropivacaine is the *S*-enantiomer of an *N*-alkyl pipercoloxylidene derivative, which is the first local anesthetic with chiral activity, and is widely used in clinical infiltration anesthesia, conduction anesthesia and epidural anesthesia. It has a long of local anesthesia and analgesic effect. However, ropivacaine also has serious safety risks in clinical practice. When the concentration of ropivacaine in human blood is too high, it may cause toxicity to the cardiovascular and central nervous system, and even cause allergic reactions in some patients. Thus far, the mechanism of the effect of ropivacaine on local anesthesia is not clear. Ropivacaine is a multitarget drug that acts on the gamma-aminobutyric acid a receptor (GABAA-R) and *N*-methyl-D-aspartate acid receptor (NDMA-R). Sodium (Na⁺) channels are a key target of local anesthetics and these two receptors regulate sodium channels. Previous studies on the structural modification of ropivacaine mainly focused on the substitution of –CH₃

on the phenyl group or the substitution of –CH₂CH₂CH₃ on piperidine with different alkyl groups. In 2017, Wen L. *et al.*⁶⁹ reported the design and synthesis of ropivacaine analogues for local anesthesia. In the process of structural design, they used ropivacaine as the lead compound to design two series of compounds, **4a–4q** (17 new substituted imines). In the first series of compounds, **4a–4i**, different substituents were selected to replace –CH₂CH₂CH₃ on piperidine. In the second series of compounds, **4j–4q**, the methyl groups were replaced by –CF₃ at the *o*-positions, *m*-positions and *p*-positions. Meanwhile, the –CH₂CH₂CH₃ on piperidine ring was also substituted and modified. The process for the synthesis of the target compounds is shown in Scheme 8. The synthetic route takes piperic acid (compound **1**) as the starting material, hydrochloric acid and sulfoxide chloride as additives, and toluene as the reaction solvent to convert compound **1** into acyl chloride salt (compound **2**). Compound **2** was then treated with substituted aniline and reacted at 58 °C for 5 h to form compounds **3a–3i** and **3j–3q**. Finally, bromoalkyl and hydrochloric acid were used to treat compounds **3a–3i** and **3j–3q**. Potassium carbonate (K₂CO₃) was used as an acid dressing agent and dimethylformamide (DMF) as the reaction solvent in *N*-alkylation reaction. The *N*-alkylation reaction lasted 10 h at 80 °C, and the salt reaction lasted 5 min at room temperature to obtain the final target compounds **4a–4q**. The total yield of the target compounds ranged from 17.5% to 87.7%. The synthetic route has the advantages of mild reaction conditions, cheap reagents and simple operation. However, using this synthetic route, the total yield of some products is too low, and too low yield will bring great problems to the synthesis cost, which needs to be further optimized in follow-up work. In the evaluation of the local anesthesia effect, sciatic nerve block activity, infiltration anesthesia activity, corneal anesthesia activity and spinal cord anesthesia activity were used as evaluation indexes. Ropivacaine was used as a positive control substance to test the local anesthesia activity *in vitro*. Firstly, the local anesthesia effect of all the target compounds **4a–4q** was screened by a sciatic nerve block test in toads *in vitro* (Table 6). The preliminary screening results *in vitro* showed that these compounds increased the blocking effect of the sciatic nerve on electrical stimulation, with ED₅₀ values ranging from 0.012 to 0.64 (positive control for ropivacaine was 0.013), with the highest activity shown by compound **4b**. In terms of latent period, that of target compounds **4a–4q** ranged from 27.7 to 59.4 min. Based on the





Scheme 8 Reagents and conditions: (a) (i) HCl, PhCH₃, r.t., 1 h; (ii) PhCH₃, SOCl₂, 55 °C, 1 h; (b) substituted anilines, 58 °C, 5 h; and (c) (i) RBr, K₂CO₃, DMF, 80 °C, 10 h; (ii) HCl, r.t., 5 min.

Table 6 The sciatic nerve block activities of compounds 4a–4q

Compound	ED ₅₀ (mg mL ⁻¹)	Latent period
4a	0.014	35.1
4b	0.012	28.9
4c	0.021	35.1
4d	0.021	34.6
4e	0.023	37.8
4f	0.021	37.5
4g	0.20	38.6
4h	0.64	39.3
4i	0.42	59.4
4j	0.026	32.8
4k	0.027	28.6
4l	0.021	33.7
4m	0.022	32.8
4n	0.026	29.0
4o	0.046	27.7
4p	0.32	28.9
4q	0.37	34.8
Ropivacaine	0.013	35.5

results of the preliminary *in vitro* screening, compounds 4a, 4b, 4c, 4j and 4l were selected to test the efficacy of invasive anesthesia in guinea pigs. The results of the infiltration anesthesia test showed that the local anesthetic effect of compounds 4c and 4l was similar to that of the positive control ropivacaine, and the local anesthetic activity of other compounds was lower than that of the positive control. Furthermore, compounds 4a, 4b, 4c, 4j and 4l were used to test the local surface anesthesia effect of these compounds (Table 7). The results of the surface anesthesia test showed that compound 4l had a similar local anesthetic effect as the positive control ropivacaine, while the effect of the other compounds was poor in comparison with the positive control. Finally, compounds 4a, 4b, 4c, 4j and 4l were tested for spinal anesthesia in order to further study their local surface anesthesia effect. The experimental results showed that the ED₅₀ produced by compounds 4l and 4b was 5.02 and 7.87, respectively, while the effects of compounds 4a, 4c and 4j were poor. The evaluation of local anesthesia *in vitro* found that compound 4l had the best activity, and thus molecular docking

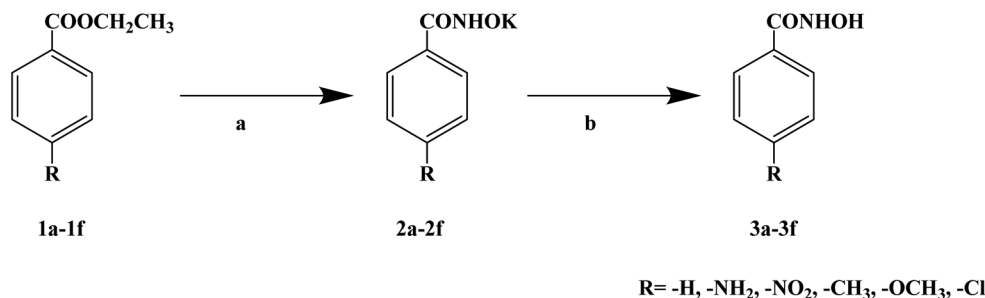
of compound 4l and ropivacaine was conducted to further study its local anesthesia mechanism. The molecular docking results showed that compound 4l interacts with receptor proteins of VGSC, GABAA-R and NDMA-R, which may help optimize and predict the activity of these ropivacaine analogues as potential local anesthetics.

Most ester anesthetics produce *p*-aminobenzoate (PABA) metabolically after use, which among the PABA and sulfa drugs have anaphylaxis and antagonistic effects. However, amide local anesthetics rarely cause anaphylaxis after use, but these drugs have a certain relationship with cardiac toxicity. Hydroxamic acid is a biologically active compound, which is a potential source in future drug development. In 2019, Ekhlas S. *et al.*⁷⁰ reported for the first time that benzohydroxamic acid with local anesthetic activity was converted into 6 benzohydroxamic acid analogues 3a–3f from *p*-substituted ethyl benzoate as a raw material (Scheme 9). The synthetic strategy was to react substituted benzoate with hydroxylamine hydrochloride in equimolar amount, and then treat intermediate products 2a–2f with potassium hydroxide. The intermediate products were in the form of potassium salt, and target compounds 3a–3f were finally acidified with acetic acid. The synthetic route only takes two steps to obtain the target compounds (3a–3f), which is simple and easy to operate, and the reagents used are cheap and easy to obtain, thus laying a solid foundation for future

Table 7 The infiltration anesthesia activities, corneal anesthesia activities and spinal anesthesia activities of selected compounds

Compound	ED ₅₀ (mg mL ⁻¹)		
	Infiltration anesthesia	Corneal anesthesia	Spinal anesthesia
4q	1.06	5.68	19.11
4b	3.02	2.64	7.87
4c	0.88	5.42	29.30
4j	4.02	4.25	34.56
4l	0.66	0.87	5.02
Ropivacaine	0.61	0.85	4.27





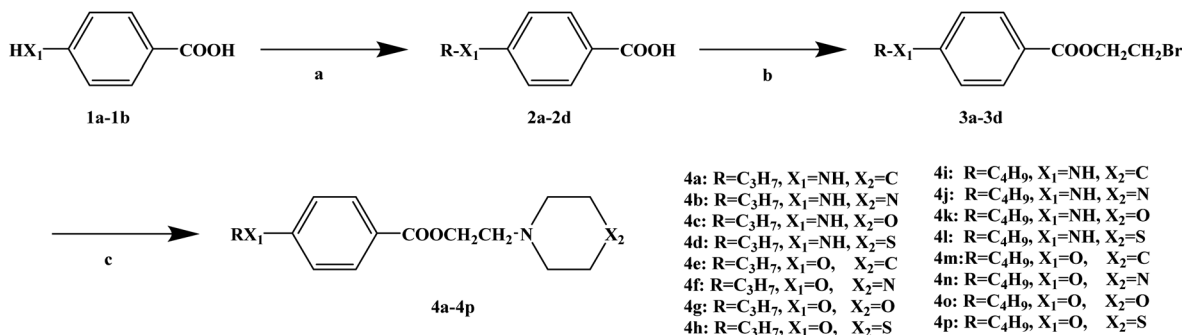
Scheme 9 Reagents and conditions: (a) $\text{NH}_2\text{OH}\cdot\text{HCl}$, KOH , CH_3OH , ice bath, 5 min; and (b) CH_3COOH , stirring.

industrialization. The total yield of all the analogues was medium, ranging from 41% to 75%, and the total yield was improved by optimizing the reaction conditions. In the evaluation of local anesthesia activity, the frog foot withdrawal reflex method was used to test the activity of the benzohydroxamic acid analogues, and benzocaine was used as the positive control. The local anesthetic activity tests were conducted with 5% dimethyl sulfoxide (DMSO) and 0.65% sodium hydroxide (NaOH) solutions, respectively. Three different concentrations of $40\ \mu\text{g mL}^{-1}$, $100\ \mu\text{g mL}^{-1}$ and $200\ \mu\text{g mL}^{-1}$ were selected for each test regimen. In 5% DMSO, most of the six benzohydroxamic acid analogues **3a–3f** showed a good local anesthesia effect. In particular, compounds **3d** and **3e** exhibited basically the same local anesthetic effect as the positive control product benzocaine at the same dose level. The drug structure–activity relationship (SAR) studies suggested that the better local anesthesia effect of compounds **3d** and **3e** was due to the fact that both methyl ($-\text{CH}_3$) and methoxy ($-\text{OCH}_3$) groups was located in the contra-apoplectic region of the strong electron donor. The presence of electrically active substituents in the adjacent or opposite position may increase the anesthetic effect of local anesthetics. When the local anesthetic activity was tested in 0.65% NaOH aqueous solution (sodium salt), compounds **3d** and **3e** also had a good local anesthetic effect when used as a sodium salt (0.65% NaOH). Among the six benzohydroxamic acid analogues **3a–3f**, compounds **3d** and **3e** showed a good local anesthesia effect, which was comparable with that of the positive control product benzocaine. These compounds have the advantages of good thermal stability, not easily hydrolyzed in water, no allergic phenomenon due to the non-release of PABA, and no antagonistic effect of clinical drugs. Thus, compounds **3d** and **3e** can be used as new local anesthetics for further research and development.

The action mechanism of local anesthetics is that the drug binds to the receptors of sodium (Na^+) channels on the nerve membrane, preventing sodium ions from crossing the ion channels and blocking the conduction of nerve impulses to achieve the purpose of local anesthesia. Currently, there are many types of local anesthetic drugs used in clinic, such as tetracaine, dolicaine, dacrone and pramocaine. However, these local anesthetic drugs have many shortcomings, such as the toxicity of tetracaine is relatively high, and the drug dose is too prone to central nervous system and cardiovascular toxicity. However, procaine has some disadvantages such as short

duration of anesthesia. It is expected that local anesthetics have the characteristics of fast acting time, long lasting acting time and low toxic and side effects. Huiying *et al.*⁷¹ reported that a series of novel benzoate compounds with local anesthetic effects were designed and synthesized with tetracaine and pramocaine as lead compounds. In the structural design of the target compounds, tetracaine and pramocaine were combined into a new structure by using the principle of splitting, and the structure was further optimized based on this new structure. A series of target compounds were obtained by optimizing the molecular structure after splicing based on the principle of bioisostere formation and modification with alkyl groups. From the structure activity relationship (SAR) of the drug, the designed target compounds also have three basic components, including a lipophilic section, a link section and a hydrophilic section. The synthetic route for the target compounds is shown in Scheme 10. The first alkylation step used 4-aminobenzoic acid (**1a**) or 4-hydroxybenzoic acid (**1b**) as the starting material, and R-Cl in dimethylformamide (DMF) to arrange it. In alkylation step, sodium carbonate anhydrous (Na_2CO_3) was used as the deacid reagent and dimethylaminopyridine (DMAP) as the catalyst, and the reflux reaction was completed in 12 h to obtain compounds **2a–2d**. The esterification step used compounds **2a–2d** and 2-bromoethanol in tetrahydrofuran (THF), and compounds **3a–3d** were obtained. The esterification step used N,N' -carbonyldiimidazole (CDI) as the dehydrant, and the reflux reaction was completed in 8 h. Also, the second alkylation step used compounds **3a–3d** and six-membered heterocyclic nitrogen in toluene, and compounds **4a–4p** were obtained. In this step, potassium carbonate anhydrous (K_2CO_3) was used as the deacid reagent, and the reflux reaction was completed in 10 h. The synthetic route has a high total yield (76.3–90.4%), mild reaction conditions, and simple and safe operation. However, in the second step of this route, CDI was used as the dehydrating agent, and the price of CDI is relatively high, which increases the synthesis cost and is not conducive to the development of industrialization. Therefore, the selection of dehydrating agent needs to be improved due to the preferential treatment of subsequent process conditions. In the study of local anesthetic activity, rabbit corneal reflection was used to study the surface anesthetic effect of the target compound activity screening *in vitro*, and based on this, further mice abdomen acupuncture tests to study the target compounds infiltrating anesthesia effect, and with the rat tail nerve block





Scheme 10 Reagents and conditions: (a) R-Cl, Na₂CO₃, DMAP, reflux, 12 h; (b) 2-bromoethanol, CDI, THF, reflux, 8 h; and (c) six-membered heterocyclic nitrogen, K₂CO₃, toluene, reflux, 10 h.

test to study the effect of nerve block anesthesia using the target compounds. The studies on local anesthesia activity showed that these compounds had a good local anesthesia effect. From the analysis of the drug structure–activity relationship (SAR), it could be seen that substituents on benzene ring have a great influence on the local anesthesia effect. With an increase in substituent carbon chain length, the anesthetic effect time was prolonged. In the screening of the local anesthesia effect *in vitro*, the target compounds **4d**, **4g**, **4j**, **4k**, **4n** and **4o** showed a good surface anesthesia effect. In the further infiltration anesthesia test, the selected target compounds also showed good activity. The action time of compounds **4n** and **4o** was 4.5 h, which was better than the 2 h of the positive control substances tetracaine and pramocaine. When performing block anesthesia on compounds **4d**, **4g**, **4j**, **4k**, **4n** and **4o**, the experimental results showed that these compounds all showed the characteristics of quick analgesic effect and quick analgesic effect (Table 8). Among them, the analgesic initiation time of compound **4n** was 6.4 ± 0.2 min, and the duration analgesic duration was 293.5 ± 3.4 min, which were better than that of the positive control. In addition, acute toxicity was evaluated by orally administering compounds **4d**, **4g**, **4j**, **4k**, **4n** and **4o** to mice (Table 4). The results of the acute toxicity test showed that the LD₅₀ of compounds **4k**, **4n** and **4o** was higher, among which compound **4n** was 976.5 ± 4.5 mg kg⁻¹. Furthermore, in the study on the mechanism of action of local anesthetics of the target compounds, they found that the target compounds enter sodium channels and bind to the receptor, thereby affecting the passage of sodium ions to achieve the effect of local anesthetics.

In general, these newly designed local anesthetics include aromatic acid esters, amides, and imidazoles, which were mainly concentrated in amides (Table 9). Most of these target compounds were structurally modified based on commercial local anesthetics, which are traditional commercial local anesthetics such as lidocaine, ropivacaine and dyclonine. Imidazoles are novel local anesthetics used in local anesthesia. In terms of the synthesis of these compounds, cheap and easy starting materials are basically used, the conventional chemical synthetic method is used, and the ideal total yield is obtained, where the total yield of some target compounds can reach more than 90%. Simultaneously, the total yield of some target compounds is low, which needs to be optimized for the subsequent synthetic process to lay a solid foundation for the future industrialization. In terms of local anesthetic activity, most of these target compounds show good local anesthetic activity, and the local anesthetic effect of most of the target compounds is similar or better than the positive control substance. All of these compounds have potential for further research and development and are waiting to become the next generation of local anesthetics. However, there are still many follow-up studies, and most of the local anesthetic effects of these target compounds are still limited to *in vitro* screening or simple activity studies *in vivo*. Thus, there is still a long way to go before they become local anesthetics on the market. This field needs continuous efforts, and it is hoped that these compounds can become a new local anesthetics and serve human health.

Table 8 The target compounds of block anesthesia and acute toxicity test

Compound	Onset of analgesia (min)	Duration of analgesia (min)	LD ₅₀ (mg kg ⁻¹)
4d	8.8 ± 0.4	230.7 ± 3.2	543.7 ± 2.3
4g	8.3 ± 0.4	254.6 ± 3.6	854.6 ± 3.7
4j	12.5 ± 0.7	221.8 ± 3.0	735.1 ± 3.2
4k	7.2 ± 0.3	279.9 ± 3.2	907.2 ± 4.3
4n	6.4 ± 0.2	293.5 ± 3.4	976.5 ± 4.5
4o	6.6 ± 0.2	286.4 ± 3.3	928.9 ± 4.9
Tetracaine	12.3 ± 0.9	172.3 ± 2.6	435.4 ± 3.3
Pramocaine	10.4 ± 0.8	166.5 ± 2.3	472.1 ± 3.6



Table 9 Synthesis and bioactivity summary of the target compounds

Schemes	Structure type	Yields	Biological activity	Research phase	References
1	Amides	45–60%	Similar to lidocaine	<i>In vitro</i>	62
2	Aromatic acid esters	97–99%	Similar to lidocaine	<i>In vitro</i>	63
3	Amides	86.6%	Similar to lidocaine	<i>In vivo</i>	64
4	Aromatic acid esters	37–75%	The anesthesia time was 126.3 ± 24.3 min and infiltration anesthesia time was 126.3 ± 24.3 min	<i>In vitro</i> and <i>in vivo</i>	65
5	Imidazoles	68–96%	<i>In vitro</i> the mouse tail anesthesia IC ₅₀ was $(1.6 \pm 0.26) \times 10^{-2} \text{ mol L}^{-1}$	<i>In vitro</i> and <i>in vivo</i>	66
6	Amides	50–80%	Better than that of lidocaine	<i>In vitro</i>	67
7	Amides	13–79%	<i>In vitro</i> , the response time of animals to thermal stimulation 61% to 124%	<i>In vitro</i> and <i>in vivo</i>	68
8	Amides	17.5–87.7%	<i>In vitro</i> ED ₅₀ values ranging from 0.012 to 0.64 and <i>in vivo</i> ED ₅₀ was 5.02 and 7.87	<i>In vitro</i> and <i>in vivo</i>	69
9	Amides	41–75%	It's similar to benzocaine	<i>In vitro</i>	70
10	Aromatic acid esters	76.3–90.4%	The analgesic initiation time was 6.4 ± 0.2 min and the duration analgesic duration was 293.5 ± 3.4 min	<i>In vitro</i> and <i>in vivo</i>	71

3. Conclusions and perspective

Local anesthetics act on nerve endings or around nerve trunks, reversibly blocking the generation and conduction of sensory nerve impulses, and temporarily eliminating local sensation under the condition of consciousness, which are convenient for local surgical operation and treatment. Herein, we mainly reviewed the research progress of local anesthetics and discussed the important aspects of design, synthesis and biological activity of various new compounds. These target compounds are mainly concentrated in amides, which basically modify the structure of existing local anesthetics to obtain local anesthetics with better biological activity. However, imidazoles are novel compounds that have local anesthesia activity. In the synthetic process, these target compounds are generally synthesized by conventional reagents and conventional methods, and the final total yield is relatively ideal. In terms of local anesthetic activity of these target compounds, most of them show good activity *in vitro* or *in vivo*, and some of them have the same or better local anesthetic activity than the positive control products. However, these target compounds also have many shortcomings, which need to be continuously optimized in the follow-up work. In terms of the design of target compounds, the target compounds are mainly amide compounds, and there are few target compounds with new structures. Thus, more novel compounds with new structures should be designed in the future, and the diversity of compound structure should be the focus. In terms of the synthesis of target compounds, the total yield of some compounds is low, which will increase the synthesis cost and hinder the development of industrialization. In terms of the activity of local anesthesia, some compounds are still in the stage of *in vitro* screening, and there is still a lot of work in the follow-up study of the activity *in vivo*. Meanwhile, most of these compounds lack of toxicological research. Thus, the latest research progress of local anesthetics can provide novel compounds for local anesthesia.

Conflicts of interest

There are no conflicts to declare.

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