Introduction

Chronic cough refers to a group of diseases in which cough symptoms last for more than 8 weeks. Common causes of chronic cough include upper airway cough syndrome, cough variant asthma, eosinophilic bronchitis, atopic cough, and gastroesophageal cough (1), and it can be effectively controlled in most patients by treatment of symptoms; however, there remain a small number of patients with unidentifiable etiology, or where treatment has poor efficacy once the etiology is identified, in whom the condition is referred to as unexplained chronic cough or refractory chronic cough (2-4). Patients with refractory chronic cough or unexplained chronic cough commonly have increased cough reflex sensitivity, which has been described as cough hypersensitivity syndrome. The adenosine triphosphate (ATP)-gated P2X3 receptor may be a key link in the activation of sensory neurons that regulate cough reflexes and has recently draw attention as a potential target for the treatment of refractory chronic cough, with a number of clinical studies validating the therapeutic effects of P2X3 receptor antagonists in patients with this condition. As the energy source for various cells in vivo, ATP localizes within cells under normal physiological conditions, and has physiological functions, including in metabolism; however, under some pathological circumstances, ATP can act as a neuromodulator and is released into the extracellular space in large quantities as a signal transduction molecule. In addition, ATP is involved in regulation of airway inflammation and the cough reflex. Here, we review the generation, release, and regulation of ATP during airway inflammation and its role in the etiology of cough hypersensitivity syndrome, including the potential underlying mechanism.

Keywords: Adenosine triphosphate (ATP); purinergic receptors; airway inflammation; cough hypersensitivity syndrome; TRPV1

The role of ATP in cough hypersensitivity syndrome: new targets for treatment

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Abstract: Clinically, chronic cough can be effectively controlled in most patients by etiological treatment; however, there remain a small number of patients whose cough has unidentifiable etiology or where treatment efficacy is poor following etiology identification, whose condition is described as unexplained chronic cough or refractory chronic cough. Patients with refractory chronic or unexplained chronic cough commonly have increased cough reflex sensitivity, which has been described as cough hypersensitivity syndrome. The adenosine triphosphate (ATP)-gated P2X3 receptor may be a key link in the activation of sensory neurons that regulate cough reflexes and has recently draw attention as a potential target for the treatment of refractory chronic cough, with a number of clinical studies validating the therapeutic effects of P2X3 receptor antagonists in patients with this condition. As the energy source for various cells in vivo, ATP localizes within cells under normal physiological conditions, and has physiological functions, including in metabolism; however, under some pathological circumstances, ATP can act as a neuromodulator and is released into the extracellular space in large quantities as a signal transduction molecule. In addition, ATP is involved in regulation of airway inflammation and the cough reflex. Here, we review the generation, release, and regulation of ATP during airway inflammation and its role in the etiology of cough hypersensitivity syndrome, including the potential underlying mechanism.
The antagonist of ATP P2X, receptor, AF-219/MK-7264 (gefaripant) (13), is useful for treatment of refractory chronic cough or unexplained chronic cough (14-16); however, there is no consensus on how ATP affects cough reflex sensitivity. In this study, we review the function of ATP in CHS etiology.

Pathogenesis of CHS—the association of TRPV1 with refractory chronic cough

CHS is heterogeneous, with the main underlying mechanisms involving excitatory changes in relevant pathways, including peripheral cough receptors, vagal afferent nerves, the higher cough center, efferent nerves, and effectors, of which activation of the transient receptor potential (TRP) pathway, airway inflammation, and cough center facilitation, may be involved in the development of increased cough reflex sensitivity (17). TRP ion channels distributed in the airways are mainly include TRP vanilloid 1 (TRPV1), TRP vanilloid 4 (TRPV4), TRP ankyrin 1 (TRPα1), and TRP cation channel subfamily M member 8 (TRPM8) (18). An altered phenotype of TRPV1 expression in airways and central sensory afferent nerves may underlie the development of cough hypersensitivity (19). TRPV1 is a ligand-gated non-selective cation channel protein, expressed in primary sensory neurons, pulmonary smooth muscle cells, bronchial and tracheal epithelial cells, and pulmonary dendritic cells (20), and can be activated by non-selective stimuli, such as capsaicin, acid (pH ≤5.9), and heat (>43 °C). Such activation opens calcium channels, causing calcium influx and producing an excitatory potential, which prompts vagal afferent C fibers to release neuropeptides, such as substance P (SP) and calcitonin gene-related peptide (CGRP), leading to local neurogenic inflammation or transmitting nerve impulses to the cough center, and increasing cough reflex sensitivity (18,21). TRPV1 was shown to mediate cough reflex in guinea pigs as early as 1995 (22). In guinea pig models with chronic cough, established by repeated aerosol inhalation of citric acid, increased capsaicin cough responses were also found to parallel tracheobronchial TRPV1 protein expression levels, suggesting that cough hypersensitivity is closely related to TRPV1 activation (23). Furthermore, TRPV1 expression is increased on sensory nerve fibers in the bronchial walls of patients with chronic cough (24). Thus, TRPV1 may be important in the development of CHS. Heightened cough responses to capsaicin inhalation have been described in patients with viral upper respiratory tract infections (25), asthma (26-28), chronic obstructive pulmonary disease (28,29), idiopathic fibrosis (30) and chronic cough (31,32). In addition, some clinical studies have found that leukotriene receptor antagonist (33), tiotropium (34) and bronchial thermoplasty (35) may ameliorate chronic asthmatic cough by altering the expression of TRPV1 to regulate capsaicin sensitivity; however, the clinical utility of TRPV1 antagonists is limited by their adverse effects, which include hyperthermia and impaired thermal sensation, as well as by their lack of stable pharmacokinetics and pharmacodynamics (36,37). In a double-blind randomized controlled trial conducted by Khalid et al., which was the first study to assess a TRPV1 receptor antagonist (SB-705498) as an antitussive agent in patients, SB-705498 failed to improve spontaneous cough frequency in patients with refractory chronic cough, but produced the reduction in capsaicin-evoked cough (38). Subsequently, Belvisi et al. validated that the potent TRPV1 inhibitor XEN-D051 showed superior efficacy and potency in preclinical and clinical capsaicin challenge studies, but was no associated with any improvements in objective cough frequency, further providing the evidence that TRPV1 is not a therapeutic effective target for refractory cough (39).

Role and potential mechanism of ATP in the etiology of CHS

ATP and P2 receptors

Under physiological conditions, cellular ATP functions in energy storage and energy supply, and the few ATP molecules released into the extracellular space are degraded to adenosine by extracellular nucleotidases; however, in inflamed airways, ATP is released into the extracellular space in large amounts, due to cell damage, stress, or hypoxia and, contributes to the development of disease and inflammation as a signaling molecule (9). Extracellular ATP can originate from nerve endings, red blood cells, platelets, skeletal muscle cells, ischemic and apoptotic cells, vascular smooth muscle cells, and endothelial or epithelial cells (8,40), and acts in an autocrine or paracrine manner on P2 purine receptors on the surface of target cells, thereby regulating cellular immune function.

In addition to cytolytic and extracellular secretion mechanisms, ATP release can even be induced by normal respiratory movements (41), when respiratory epithelium is subjected to stress; for example, by mechanical stimuli. Purinergic receptors include two major categories: P1 and
P2. The ligand for the P1 receptor is adenosine, which has four subtypes: A1, A2A, A2B, and A3. ATP-bound P2 receptors include two major families, P2XR and P2YR, the former being ligand-ATP-gated non-selective cation channels, with seven known receptor subtypes, P2X<sub>1</sub>-R, which allow the passage of cations, such as Ca<sup>2+</sup>, Na<sup>+</sup>, and K<sup>+</sup>; when activated. Of these subtypes, P2X<sub>R</sub> is widely distributed in the peripheral and central nervous systems, and P2X<sub>R</sub> is primarily found in sensory nerve cells, including the trigeminal nerve, dorsal root ganglia, and nodose ganglia, and is closely associated with cough (42). P2X<sub>R</sub> and P2X<sub>6</sub>/P2X<sub>R</sub> heteromers are moderately expressed in the lung, while P2Y<sub>R</sub> is a metabotropic G-protein coupled receptor expressed in almost all epithelial cells, and is responsible for fluid control and electrolyte transport, including P2Y<sub>R</sub>, P2Y<sub>1R</sub>, P2Y<sub>2R</sub>, P2Y<sub>4R</sub>, and P2Y<sub>11,14R8</sub> receptor subtypes (43). Among them, P2Y<sub>1R</sub>, P2Y<sub>2R</sub>, and P2Y<sub>4R</sub> are more strongly expressed in the Airways and lungs (43,44), while P2Y<sub>R</sub> is mainly related to transient platelet aggregation and deformation, and P2Y<sub>R</sub> and P2Y<sub>R</sub> are involved in regulation of the physiological functions of the respiratory system. (More specific distributions and functions of purinergic receptors are shown in Table S1).

**ATP may be involved in the etiology of CHS**

Airway inflammation and remodeling are common pathological changes in chronic cough caused by various intrapulmonary and extrapulmonary diseases; however, airway injury caused by persistent cough also leads to airway inflammation (45,46), which presents as pathological changes, such as mucosal vasodilatation, plasma exudation, tissue edema, epithelial injury, and inflammatory cell infiltration (46). Numerous studies have indicated the presence of extracellular ATP during the development of airway inflammation. Esther et al. found that ATP levels in bronchoalveolar lavage are inversely correlated with lung function in children with pulmonary cystic fibrosis, while extracellular purines, including ATP, can be used as biomarkers of neutrophilic airway inflammation (47). Basoglu et al. found that aerosol inhalation of ATP can produce cough in 70% of healthy subjects and 90% of those with mild intermittent asthma; however, ATP had no bronchoconstriction effect on healthy subjects. Extracellular ATP is involved in neuropathic bronchoconstriction, inflammation, and cough caused by P2 receptor sensitization in airway sensory nerves, which may contribute to the pathogenesis of asthma and obstructive airway diseases (10). Fowles et al. conducted a randomized controlled trial that recruited 20 healthy volunteers and 20 patients with chronic cough, with the same sex ratio in both groups. Each group was subjected to cough provocation tests with ATP and AMP; the primary endpoints were the differences between ATP and AMP, as well as the substance concentration provoking cough after 2 and 5 cough exposures (C2 and C5), between healthy volunteers and patients with chronic cough. They found that aerosol inhalation of ATP could produce cough in both healthy subjects and patients with chronic cough, and that C2 and C5 values were significantly lower in patients with chronic cough than those in healthy volunteers, while the response to ATP was more pronounced (12). Under pathological conditions, binding of extracellular ATP to P2 receptors can produce different physiological effects. ATP activates P2X<sub>2,3R</sub>, located on C and Aδ fibers, to stimulate vagal afferent nerve endings in the lungs, leading to the release of local neurogenic inflammatory mediators, bronchoconstriction, and cough (48,49). By activating P2Y<sub>4R</sub>, ATP causes the release of inflammatory factors, including IL-6 and IL-8, which amplifies inflammatory effects on the airway (50,51). IL-6 and IL-8 also contribute to the release of thromboxane from the airway epithelium via P2Y<sub>i</sub> and P2Y<sub>α</sub>, causing allergic bronchospasm (52). Exonuclease E-NPP1 expression is decreased in blood leukocytes from patients with severe acute bronchial asthma episodes, and P2X<sub>7</sub> and P2Y<sub>12</sub> receptors mediate increased production of endogenous ATP, exacerbating asthma (53). Pelleg et al. found that ATP enhances IgE-mediated histamine release from mast cells (54). In addition, histamine also induces ATP release from subcutaneous and smooth muscle fibroblasts (55,56). Kamei et al. suggested that ATP did not cause cough directly, but may enhance cough reflex sensitivity in guinea pigs by becoming rapidly adapted to cough receptors, rather than the C-afferent fiber pathway, with P2X receptors an important link in this process (11,56).

**P2 receptor antagonists can be used to treat unexplained refractory chronic cough**

Selective P2X<sub>7</sub>R antagonists can be used to treat unexplained refractory chronic cough. P2X<sub>7</sub>R is an important vector for cough hypersensitivity that is mainly expressed on C-vagal afferent nerve fibers (57). A recent study using the whole-cell patch clamp technique in 1321N1 cells expressing human P2X<sub>R</sub> and P2X<sub>2,7</sub>R receptors...
found that the reversible selective P2X,R and P2X\gamma,R antagonist, MK-7264 (gefapixant), exerted its antagonistic effects through negative allosteric modulation. In addition, in vivo experiments demonstrated that MK-7264 had high oral bioavailability in a rat model of neuropathic sensitization, could treat neuralgia, and had comparable efficacy to gabapentin, supporting its clinical application for diseases involving chronic hypersensitivity (13). In a randomized placebo-controlled clinical trial, Morice et al. demonstrated that oral administration of 100 mg of the P2X\beta receptor antagonist, AF-219/MK-7264, in patients with chronic cough significantly reduced cough reflex sensitivity induced by inhaled ATP challenge 5 hours later; however, it had no inhibitory effect on cough hyperresponsiveness induced by capsaicin and citric acid challenge, indicating that AF-219/MK-7264 only inhibits P2X\beta receptors on airway cough receptors, and is not associated with the inhibition of receptors such as TRPV1 (16). Further, a single-center randomized, double-blind, placebo-controlled phase II clinical study, conducted by Rayid Abdulqawi et al. (14), showed that oral administration of the P2X\alpha receptor antagonist, AF-219/MK-7264 (600 mg twice daily for 2 weeks), could significantly reduce cough frequency and severity, and improve the quality of life in patients with refractory chronic cough. Despite good antitussive effects, where cough frequency can be reduced by 75% in patients administered AF-219 compared with those receiving placebo, AF-219/MK-7264 was associated with frequent dysgeusia as a side-effect; all patients with refractory chronic cough taking AF-219 experienced this symptom, and six discontinued the drugs due to intolerance, which usually appeared within a few hours after administration, fully recovering by 24 hours after discontinuation. Subsequent dose-finding studies showed that reducing the dose of medication significantly diminished the incidence and extent of dysgeusia, while maintaining high efficacy, indicating that AF-219/MK-7264 has great promise for final inclusion in clinical treatment. To explore efficacy and tolerability of gefapixant (AF-219/MK-7264), the P2X\beta receptor antagonist, for the treatment of chronic cough, Smith et al. designed two randomized, double-blind, placebo-controlled, crossover, dose-escalation studies (58). Both studies were composed of two 16-day treatment periods with either 3–7 days (Study 1) or 14–21 days (Study 2) washout periods. Study 1 was the high-dose cohort, in which gefapixant was received at four BID dose levels (50, 100, 150 and 200 mg); study 2 investigated a lower range of four BID dose levels (7.5, 15, 30 and 50 mg); both doses escalated every 4 days. This work suggested that compared with placebo, gefapixant dose ≥30 mg ameliorated cough mostly and taste disturbances were closely related to the doses, most obvious at dose ≥150 mg. Gefapixant could maintain the maximum efficacy at a lower dose of 50 mg than previous research twice daily, well tolerated. Recent work about gefapixant in the treatment of refractory or unexplained chronic cough, a randomized, double-blind, controlled, parallel-group, phase 2b trial, was reported in Lancet Respir Med by Smith et al. (59), first to show the anti-tussive efficacy of P2X\beta antagonist maintained over 12 weeks. Dosing regimen used in this study was set at 7.5 to 50 mg twice daily as a dose response plateau in the former dose-escalation research evaluating doses of gefapixant from 7.5 to 200 mg (58). 253 patients enrolled were randomly assigned to receive placebo or one of three doses (7.5, 20, or 50 mg) of oral gefapixant twice daily for 84 days and were found that inhibiting P2X\beta receptor with gefapixant at a dose of 50 mg twice daily had good safety and tolerability, significantly reducing cough frequency in patients with refractory and unexplained chronic cough after 12-week treatment. This large-scale, multicenter study validated and complemented previous preliminary proof-of-concept studies on the anti-tussive efficacy of P2X\beta antagonist and investigations on the balance between efficacy and tolerability (Table S2). At present, an ongoing global multicenter phase III clinical trial of low-dose gefapixant for treatment of refractory chronic cough has enrolled 2,000 subjects, and is expected to obtain positive efficacy results, representing an optimistic development for patients with refractory chronic cough (15). Moreover, studies by Smith et al. also found that gefapixant did not eliminate excessive cough completely (59) and failed to show a good response to patients with lower daily cough frequencies (14), which can be explained by the heterogeneity among patients with chronic refractory cough (60,61). There may exist other processes related to the cough hypersensitivity in refractory chronic cough patients rather than ATP-P2X pathways. Coughing of those subjects with poor efficacy may be mainly produced by hypersensitivity and continuous exposure to inhaled stimuli, rather than by endogenous ATP-dependent mechanisms (62).

Novel P2X,R antagonists are also being explored. BLU-5937 is a non-competitive antagonist with high selectivity for P2X,R homotrimers. In guinea pig cough models, BLU-5937 significantly reduced the increase in cough episodes induced by histamine or ATP in a dose-
dependent manner. Further, in rat models of behavioral taste response, BLU-5937 had no significant effect on taste, compared with control animals (63). BLU-5937 has excellent pharmaceutical properties, including high oral bioavailability, stable metabolism, no blood-brain barrier permeability, and good tolerability, and is currently in clinical phase I development. Besides, S-600918, a new antagonist, highly selective for P2X3 homomer compared with P2X2/3, heteromer, is expected to reduce cough frequency with minimal taste-related side effects in patients with refractory chronic cough (64). Molecules exhibiting greater selectivity for P2X3 compared with P2X2/3 are also under investigation, such as BAY 1817080 and BAY1902607 (42). The results of the above studies indicate that ATP may be involved in the formation of CHS, and show that P2X3R antagonists have a clear therapeutic effects on refractory chronic cough; however, in view of the high incidence of dysgeusia, further dose response studies are needed to achieve cough improvement at well-tolerated doses. Further, BLU-5937, as a non-competitive antagonist with high potency, selectivity for the P2X3 receptor, and excellent pharmaceutical properties for the treatment of chronic cough, strongly warrants additional investigation in clinical practice. Gefapixant, the P2X3 receptor antagonist, is expected to be the first therapy approved for the treatment of refractory chronic cough. In addition, refractory chronic cough can be treated by therapies targeting other channels, such as neuromodulators and speech therapy, indicating that the pathogenesis of excess cough are unlikely occur independently, and coughing in different patients may be dominated by different mechanisms (42). Consequently, for the management of different patients with refractory chronic cough, their clinical characteristics must be analyzed to achieve accurate phenotyping in line with various preclinical and clinical researches to predict their potential underlying mechanisms and choose the optimal therapy.

Possible mechanism of the involvement of ATP in the etiology of CHS

ATP and TRPV1

TRPV1 is mainly expressed in primary afferent sensory neurons, smooth muscle cells of the lung, bronchial and small airway epithelial cells, and dendritic cells of the lung, and is involved in the activation of a variety of neuroinflammatory pathways (20,65), usually mediated by capsaicin. In addition, almost all major neurogenic inflammatory signaling pathways converge on TRPV1, to increase the excitability of C fibers during airway inflammation (21), and produce neurohypersensitivity, causing patients to respond to minimal stimulation, thus forming a vicious cycle and leading to therapeutic difficulties. Clinically, airway wall injury, due to mechanical shear stress generated during prolonged and repeated coughing in patients with refractory chronic cough, may increase ATP release into the airways (66,67), recruit inflammatory cells, and initiate or aggravate airway inflammation and remodeling (68). Extracellular ATP itself can directly depolarize vagal nerve fibers in animals and humans (57), with similar effects to capsaicin, producing transient but unsustained receptor potential pathways that sensitize vagal sensory fiber terminals in the lung, resulting in increased bronchoconstriction and cough reflex sensitivity, an effect that may be associated with activation of TRPV1 (8). In addition, Ma et al. found that ATP is involved in HCl-induced activation of TRPV1 receptors in the esophageal mucosa (69) and that TRPV1 can also mediate acid-stimulated ATP release from porcine bladder mucosa (70). ATP can interact with TRPV1, as demonstrated by studies in TRPV1-deficient mice, as it induces thermal hyperalgesia in wild-type, but not TRPV1-deficient, mice (71). TRPV1 expressed by C afferent vagal nerve fiber terminals is closely related to cough reflex sensitivity. During airway inflammation, a variety of inflammatory mediators can activate TRPV1, so that C afferent vagal nerve fiber neurolemma is rapidly depolarized, leading to excitement and enhancing cough reflex sensitivity. However, combined with the finding of Morice et al. that oral administration of the P2X3 receptor antagonist, AF-219/MK-7264, did not affect cough reflex sensitivity to either capsaicin or citric acid in patients with chronic cough (16), whether ATP-mediated changes in cough reflex sensitivity are related to TRPV1 is uncertain and further studies are warranted.

TRPV4 and ATP

TRPV4 is a non-selective calcium channel, widely expressed in the respiratory tract, including epithelium (human), macrophages (human and rat), and airway smooth muscles (human and guinea pig). It is also expressed in the trigeminal and dorsal root ganglia. A variety of physical and chemical stimuli, including mechanical stimulation, mild thermal stimulation above 24 °C, and arachidonic acid metabolites (72), can activate TRPV4. Further, TRPV4 can act as an osmotic sensor, to participate in cough development. Buday et al. also confirmed that TRPV4 mediates the cough response
induced by citric acid and hypotonic solution (72-75). Activation of TRPV4 causes depolarization of vagal fibers and activation of Aδ, rather than C, fibers and the transmission of impulses to the nodose ganglia causes an increase in intracytoplasmic calcium concentration, which exacerbates cough, or causes airway smooth muscle contraction, in humans and animals; the jugular ganglion does not contribute to this process, and TRPV4 antagonists can reduce cough (75,76). Studies on human primary alveolar macrophages and bronchial epithelial cells exposed to cigarettes have shown that ATP release can be reduced using specific antagonists of TRPV1 or TRPV4, and that TRPV1 or TRPV4 agonists can directly induce ATP release (74). TRPV4 also mediates ATP release from human lung fibroblasts and subsequently acts on the purinergic receptor, P2YR, in an autocrine or paracrine manner (77).

Unlike the capsaicin-mediated cough reflex, extracellular ATP-mediated enhancement of the cough reflex is not attenuated by desensitization of C-fibers, suggesting that the mechanism by which ATP enhances the cough reflex may also involve Aδ-fibers (8). In a study by Morice et al., the P2X,R antagonist, gefapixant, was found to increase the cough threshold for inhalation of distilled water, without having a significant effect on cough threshold on inhalation of capsaicin or citric acid, which also suggests that ATP-related changes in cough reflex sensitivity may not be related to C fibers (16). Further, Gu et al. injected the potent TRPV4 agonist, GSK1016790A, into the right atrium of anesthetized rats with spontaneous breathing, causing rapid shallow breathing observed in anesthetized rats, an effect that could be abolished by the selective TRPV4 antagonist, GSK2193874, or vagotomy. Moreover, patch clamp experiments showed that GSK1016790A does not directly activate bronchopulmonary sensory neurons, possibly acting indirectly on TRPV4-expressing cells in the lung and airways to regulate respiration (72). In addition, Bonvini et al. found that TRPV4 agonists and hypotonic solutions can induce depolarization of isolated vagal and Aδ fibers (non-C fibers) in mouse, guinea pig, and human tissues, using calcium imaging techniques, in vivo electrophysiology, and in vivo animal models of cough. Further, this effect was blocked by TRPV4 or P2X,R antagonists, which have similar effects, while the TRPV4 agonist, GSK1016790A, caused slower activation of Aδ fibers than capsaicin and citric acid, indicating that TRPV4 activation has an indirect effect on Aδ fiber activation, rather than a direct effect (75). Hence, it is speculated that TRPV4 may mediate ATP-activation-induced depolarization of vagal fibers and activation of Aδ fibers.

Overall, we propose that endogenous ATP release and P2XR activation are prerequisites for the TRPV4-mediated cough reflex, and that TRPV4-ATP-P2X signaling may be the main driver of hypotonicity-induced airway afferent nerve fiber activation. Nevertheless, Buday et al. also found that TRPV4 antagonists partially inhibited citric acid-induced cough in guinea pigs, while having no effect on cough induced by distilled water and hypertonic fluid. TRPV4-mediated activation of airway afferent nerve fibers does not fully explain osmolarity-related cough (73). The mechanism underlying TRPV4-induced ATP release remains unclear and requires further study. However, a clinical research by Ludbrook et al. suggested that the selective TRPV4 channel blocker GSK2798745 had no effect on reducing cough frequency (78). Therefore, whether TRPV4 antagonists can be used as an alternative therapy to P2X,R antagonists for patients with refractory chronic cough remains to be validated.

**Summary of mechanism of action**

In summary, ATP is released into the extracellular space during airway inflammation, where it binds to corresponding P2 receptors to mediate the cough reflex and participate in the formation of cough hypersensitivity. The underlying mechanism may be related to TRPV4-mediated activation of Aδ afferent vagal fibers. Periodic force, generated during activation of TRPV4 (for example by hypotonic solution, low pH, citric acid, or inflammatory metabolites) or cough, damages airway epithelium and causes it to release a large amount of ATP, thereby stimulating P2X receptors on activated cough receptors, producing signal transduction, and further activating Aδ afferent vagal fibers. Nerve impulse transmission to nodose ganglia causes increased intracytoplasmic calcium concentration. The conduction of peripheral stimuli to the cough center will result in increased cough reflex sensitivity, further inducing TRPV1 expression in airway sensory nerve fiber terminals. In addition, ATP may also promote inflammatory cells to release inflammatory factors or act directly on P2 receptors on a few nodose ganglion C fibers, producing neurogenic inflammation and activating TRPV1 in vagal nerve endings, thereby generating a vicious cycle of increased cough reflex sensitivity, rendering the cough difficult to cure.

**Future prospects**

The treatment of refractory chronic cough remains a
common medical problem facing clinicians. Although a number of studies have verified the effectiveness of P2X3R for the treatment of refractory chronic cough, the high incidence of the adverse reaction of dysgeusia limits the sample size of clinical trials. Further, the optimal dose, treatment cycle, underlying mechanism of action, and safety of the long-term application of P2X3R for the treatment of refractory chronic cough are unclear. A global multicenter study is ongoing. New drugs, highly specific for P2X3R targets, need to be studied and developed. Simultaneously, basic research to further explore the specific mechanism underlying the involvement of ATP in the etiology of cough hypersensitivity is warranted, to provide theoretical support for the clinical research.

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### Table S1 Distributions and functions of purinergic receptors

| Purinergic receptors | Ligands | Types | Subtypes | Expression positions | Functions |
|----------------------|---------|-------|----------|----------------------|----------|
| **P1**               | Adenosine | AR    | A<sub>1</sub> | Highly expressed in the CNS and less in lungs | Bidirectional regulating the airway contraction and inflammation (79) |
|                      |         |       | A<sub>2A</sub> | Highly expressed in the brain striatum, immune cells of the spleen, lymphocytes, thymus and platelets and moderately in heart, lungs and blood vessels | Mediating inflammatory response (79) |
|                      |         |       | A<sub>2B</sub> | Widely distributed in heart, lungs, spleen, kidneys, colon, etc., especially in the vasculature | Bidirectional adjusting the function of CNS; Inhibiting myocardial injury and promoting angiogenesis; regulating intestinal motility and inflammation, etc. (80) |
|                      |         |       | A<sub>3</sub> | Eosinophils of lungs and liver | Inhibiting the aggregation and degranulation of neutrophils and eosinophils, thus anti-inflammatory (79) |
| **P2**               | ATP     | P2XR  | P2X<sub>1</sub> | Mainly expressed in smooth muscle cells | Involved in the regulation of smooth muscle contraction and relaxation, especially closely related to bladder functions (81) |
|                      |         |       | P2X<sub>2</sub> | Widely distributed in peripheral and central nervous system | Co-expressed with P2X<sub>3</sub> and involved in modulating sensory functions (43) |
|                      |         |       | P2X<sub>3</sub> | Sensory cells of trigeminal nerve, dorsal root ganglion of spinal nerve and nodose ganglion | Closely related to cough and important in pain regulation (42) |
|                      |         |       | P2X<sub>4</sub> | Widely distributed in brain, spinal cord, sympathetic nerve and sensory ganglion and moderately in lungs | Involved in the regulation of allergic airway inflammation, airway remodeling and physiological processes of the CNS, including pain and Parkinson, etc. (82,83) |
|                      |         |       | P2X<sub>5</sub> | Highly expressed in heart | Involved in regulating the physiological function of the cardiovascular system (81) |
|                      |         |       | P2X<sub>6</sub> | P2X<sub>4</sub>/P2X<sub>6</sub> heteromer is widely expressed in the CNS and moderately in lungs | Co-expressed with P2X<sub>3</sub>; exciting P2X<sub>6</sub> itself does not cause current (81,84) |
|                      |         |       | P2X<sub>7</sub> | Highly expressed in pancreas, liver, heart, thymus and brain and moderately in lungs | Involved in inflammation and pain regulation (85,86) |
| **P2YR**             | P2Y<sub>1</sub> |       | Widely distributed in heart, brain, lungs, placenta and co-expressed with P2Y<sub>10</sub> in platelets | Involved in the transient formation and aggregation of platelets; anti-inflammatory; dilating blood vessels; promoting the growth of neurons and nerve fibers (87) |
|                      | P2Y<sub>2</sub> |       | Epithelial cells in airways | Involved in secretion of mucin by glands and airway allergic reaction (40) |
|                      | P2Y<sub>4</sub> |       | Co-expressed with P2Y<sub>5</sub>, mainly in gastrointestinal | Mediating Cl<sup>-</sup> secretion in gastrointestinal tract; maintaining Na<sup>+</sup> balance in cochlea (88) |
|                      | P2Y<sub>6</sub> |       | Cardiovascular system dominated | Regulating inflammation, promoting smooth muscle contraction and cardiovascular disease (89) |
|                      | P2Y<sub>11</sub> |       | Brain and white blood cells | An important regulator of immune cell survival (90) |
|                      | P2Y<sub>12</sub> |       | Platelets | Related to TLR2-mediated random migration of microglial cells and transient platelet formation and aggregation (91) |
|                      | P2Y<sub>13</sub> |       | Spleen and brain | Involved in stem cell proliferation and differentiation (90) |
|                      | P2Y<sub>14</sub> |       | Universally expressed in vivo and significantly in various immune cells | Involved in modulating inflammation and immune response (90) |

CNS, central nervous system; TLR2, toll-like receptor 2.
| Study | Study type | Recruited patients | Drug dose and course of treatment | Primary endpoint measure | Main findings | Side effects | Conclusion |
|-------|------------|--------------------|----------------------------------|--------------------------|---------------|-------------|------------|
| Abdulqawi R et al. 2015 (14) | Randomized, double-blind, placebo-controlled Phase II study | 24 RCC patients | Oral AF-219 600 mg or placebo for 2 weeks, bid, then crossover for 2 weeks | Cough frequency | 75% reduction in cough frequency vs. placebo control, accompanied by significant improvements in CQLQ scores and VAS scores | All patients experienced dysgeusia, which disappeared after drug withdrawal | P2X3 antagonists significantly improve cough symptoms in RCC patients, but further dose-response studies are needed |
| Morice AH et al. 2019 (16) | Randomized, double-blind, placebo-controlled Phase II study | 24 patients with chronic cough vs. 12 healthy subjects | Cough sensitivity to ATP, citric acid, capsaicin, and distilled water was measured at 1, 3, and 5 h after oral administration of gefapixant (100 mg) or placebo | Change in C2 and C5 cough thresholds after drug administration vs. before drug administration | Gefapixant inhibited cough sensitivity to ATP or distilled water, but had no significant effect on cough responses elicited by capsaicin or citric acid | Dysgeusia occurred in 75% of healthy subjects and 25% of patients with chronic cough | Gefapixant may improve cough symptoms in patients with CC, and cough hypersensitivity in RCC may be mediated by the TRPV4/ATP pathway |
| Smith JA et al. 2020 (58) | Two randomized, double-blind, placebo-controlled, crossover, dose-escalation studies | In total 59 patients with refractory chronic cough were randomized; 29 in study 1, 30 in study 2 with 18 subjects participating in both studies | Both studies were composed of two 16-day treatment periods with either 3–7 day (Study 1) or 14–21 day (Study 2) washout periods. In study 1 gefapixant was received at four BID dose levels (50, 100, 150 and 200 mg); study 2 investigated a lower range of four BID dose levels (7.5, 15, 30 and 50 mg); both doses escalated every 4 days | Awake cough frequency assessed with 24 h ambulatory cough monitor at baseline and on day 4 of each dose | Gefapixant doses ≥30 mg produced maximal improvements in coughing compared with placebo; taste disturbances were closely related to doses, apparently maximal at dose ≥150 mg | Four subjects terminated study drug early due to AEs, 3 in study 1 and 1 in study 2; only one termination was due to taste disturbance | P2X3 antagonism with gefapixant demonstrates antitussive efficacy and improved tolerability at lower doses than previously investigated. Longer duration studies are warranted |
| Smith JA et al. 2020 (59) | Randomised, double-blind, controlled, parallel-group, phase 2b trial | 253 patients with refractory or unexplained chronic cough aged 18–80 years | Patients enrolled were randomly assigned to gefapixant 7.5 mg (n=64), 20 mg (n=63), or 50 mg (n=65), or a matching placebo (n=63) twice daily for 12 weeks | Placebo-adjusted change from baseline in awake cough frequency after 12 weeks | Inhibiting P2X3 receptor with gefapixant at a dose of 50 mg twice daily had good safety and tolerability, significantly reducing cough frequency in patients with refractory and unexplained chronic cough after 12-week treatment | Taste-related adverse events, oral paraesthesia, and oral hypoesthesia increased in frequency in a dose-dependent manner. Dysgeusia and hypogeusia were the most common adverse events in this study | Gefapixant shows promise as a novel therapy for chronic cough, and further studies examining longer-term antitussive benefit are warranted |

CC, chronic cough; RCC, refractory chronic cough; CQLQ, Cough Quality of Life Questionnaire; VAS, visual analog score; C2, minimum challenge inhalation concentration required to induce ≥ two coughs; C5, minimum challenge inhalation concentration required to induce ≥ five coughs; AEs, adverse events.
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