



Therapeutic Targets for the Treatment of Chronic Cough

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Abstract

Purpose of Review Chronic cough, defined in adults as one lasting longer than 8 weeks, is among the commonest clinical problems encountered by doctors both in general practice and in hospital. It can exist as a distinct clinical problem or as a prominent and troublesome symptom for patients with common pulmonary conditions including asthma, chronic obstructive pulmonary disease and idiopathic pulmonary fibrosis.

Recent Findings Chronic cough impacts considerably on patients' daily-life activities and many patients are left frustrated by what they see as a complete lack of awareness among their doctors as how to treat their condition. Some of this arises from limited levels of physician knowledge about managing cough as a clinical problem but also because there are no very effective treatments that specifically target cough.

Summary In this article, we review the current clinical thinking regarding cough and the treatments that are currently used and those undergoing clinical development.

Keywords Cough · Cough receptor · Pharmacological targets · Novel · Ion channel

Introduction

Under normal physiological circumstances, coughing occurs with the primary purpose of protecting the lung from inhaled irritants and clearing unwanted airway secretions. Clinically, this defensive neural reflex can become dysregulated leading to episodes of cough evoked by relatively innocuous stimuli. This causes significant problems for those affected and is one of the most common reasons why people seek medical attention [1].

For clinical purposes, cough is defined as acute, sub-acute or chronic depending on its duration. An acute cough typically lasts no more than 3 weeks and is generally the result of a viral upper respiratory tract infection (URTI) associated with the common cold. Sub-acute cough lasts between 3 and 8 weeks

and is likely due to a slowly resolving post-viral cough. In adult patients, a cough persisting for more than 8 weeks is termed 'chronic' and can occur as an isolated clinical problem or associated with common respiratory and non-respiratory conditions such as chronic obstructive pulmonary disease (COPD), asthma, idiopathic pulmonary fibrosis (IPF), gastro-oesophageal reflux disease (GORD) and post-nasal drip or rhinosinusitis. Sometimes, despite extensive investigation and trials of empirical therapy, the cause for cough remains unexplained and in such instances is termed an idiopathic, or unexplained chronic cough [2, 3].

The prevalence of chronic cough may affect up to 13% of the general population and is associated with significant physical and psychological morbidity [4–8]. In a recent survey of almost 2000 people living with chronic cough, respondents reported their cough had impacted adversely on daily-life activities leading to many feeling fed-up and depressed [9]. Crucially, the majority reported undergoing numerous consultations with different physicians and receiving many unsuccessful therapeutic trials.

The increasing recognition of chronic cough as an important clinical problem has led to the development of guidelines to improve clinical care [10–12]. The underlying principle recommended in these guidelines is first to consider common respiratory diseases associated with obvious chest radiology and lung function abnormalities and then systematically

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evaluate the patient for other common causes such as asthma, reflux disease and upper airway disorders. While this approach is helpful and treatments such as inhaled steroids and anti-reflux therapy do help to control cough, a significant proportion of patients remain troubled [3]. In response to increasing numbers of patients referred for evaluation of cough, a number of specialist clinics have been set up within the UK, throughout Europe, the USA and Australasia. With a more detailed and expert assessment, there has been a fundamental change in understanding of the complex clinical aspects of cough. Central to this has been the notion that cough develops as a result of inflammation causing damage to the nerve cells (neuroinflammation) that conduct the cough signal from the lungs to the brain resulting in the clinical state now termed, cough hypersensitivity syndrome (CHS). Understanding the mechanisms responsible for this nerve damage (neuropathy) will be key to developing better treatments [13].

Cough Hypersensitivity Syndrome

The term ‘Cough Hypersensitivity Syndrome’ was first proposed by a European Respiratory Society Task Force and is defined as a ‘clinical syndrome characterised by troublesome coughing triggered by low levels of thermal, mechanical or chemical exposure’ [14]. The underlying concept is that the basal protective cough reflex transforms to a heightened state such that it is now triggered by low-level stimuli that would not usually cause cough. This state reflects precisely the clinical features reported by patients with CHS in so far as changes in ambient temperature, taking a deep breath, or talking, exposure to aerosol sprays or eating crumbly dry food are all sufficient to trigger bouts of coughing. The mechanisms responsible for this clinical state have yet to be determined but may parallel other disease concepts such as neuropathic pain. Following this line of thinking, cough triggered by innocuous stimuli might be similar to allodynia and excessive coughing in response to a noxious exposure (hypertussia) could be considered equivalent to hyperalgesia.

Although the causes of chronic cough vary, cough reflex hypersensitivity is likely to be a shared clinical and mechanistic characteristic. Therefore, while CHS may be present in a wide variety of conditions, common mechanisms likely underpin its development, offering great potential for the development of effective therapies.

Treatment of Chronic Cough

The recognition that chronic cough is characterised by hypersensitivity of the peripheral and central neural pathways involved in cough has expanded the range of potential therapeutic targets currently under evaluation. Here, we provide an

update on these therapies with a focus on proposed site of action (peripheral or central) and preclinical and clinical findings to date. To do this, we undertook a detailed appraisal of published peer-reviewed research over the past 5 years (January 2013–December 2018) using PubMed for articles in English. The search terms included ‘cough’ plus ‘receptor’, ‘pharmacological targets’, ‘novel’, ‘ion channel’. We also referred to publications accumulated because of our involvement in chronic cough research and treatment over the past 15 years. Articles resulting from these searches and relevant references cited in those articles were also reviewed. We also reviewed *ClinicalTrials.gov* over the last 5 years, and detail from this has been summarised in an accompanying table.

Treatments with a Peripheral Site of Action

Attenuating the heightened cough sensitivity while retaining the protective physiological capacity of the cough reflex is critical in the development of effective treatments. Sensory nerves of the airways express a variety of receptors and ion channels that when activated trigger cough. It seems logical that these channels/receptors may represent therapeutic targets for cough (Fig. 1). Here, we provide brief overview of experience with a number of these putative targets.

TRP Channels

Transient receptor potential (TRP) channels were first discovered and described in *Drosophila* in the 1960s but have since become important targets in chronic cough [15]. Manipulation of TRP channels on nociceptive neurons is proposed as an attractive strategy in the treatment of pain, and several drug developments targeting pain are focusing on these receptors. TRP channels are present in abundance in the airways and are expressed in many cell types of the airway including primary sensory afferent nerves, epithelial cells and smooth muscle cells [16].

TRP channels are cation channels comprising six transmembrane spanning domains with a pore between the fifth and sixth domains which allows the entry of calcium into a cell [17]. TRP channels are polymodal and directly activated by changes in temperature, chemicals, mechanical stimulation, pH and osmolality, which may evoke nociceptive responses including cough. The TRP channel superfamily encompasses 28 members that can be divided into 6 subfamilies based on sequence homology [17]. Of particular interest in relation to cough are members of the vanilloid (TRPV1, TRPV4), anykrin (TRPA1) and melastatin (TRPM8) families.

TRPV1 TRPV1 was the first channel to be considered as a key regulator of cough. TRPV1 is a polymodal channel activated by a variety of irritants including high temperature (>43 °C),

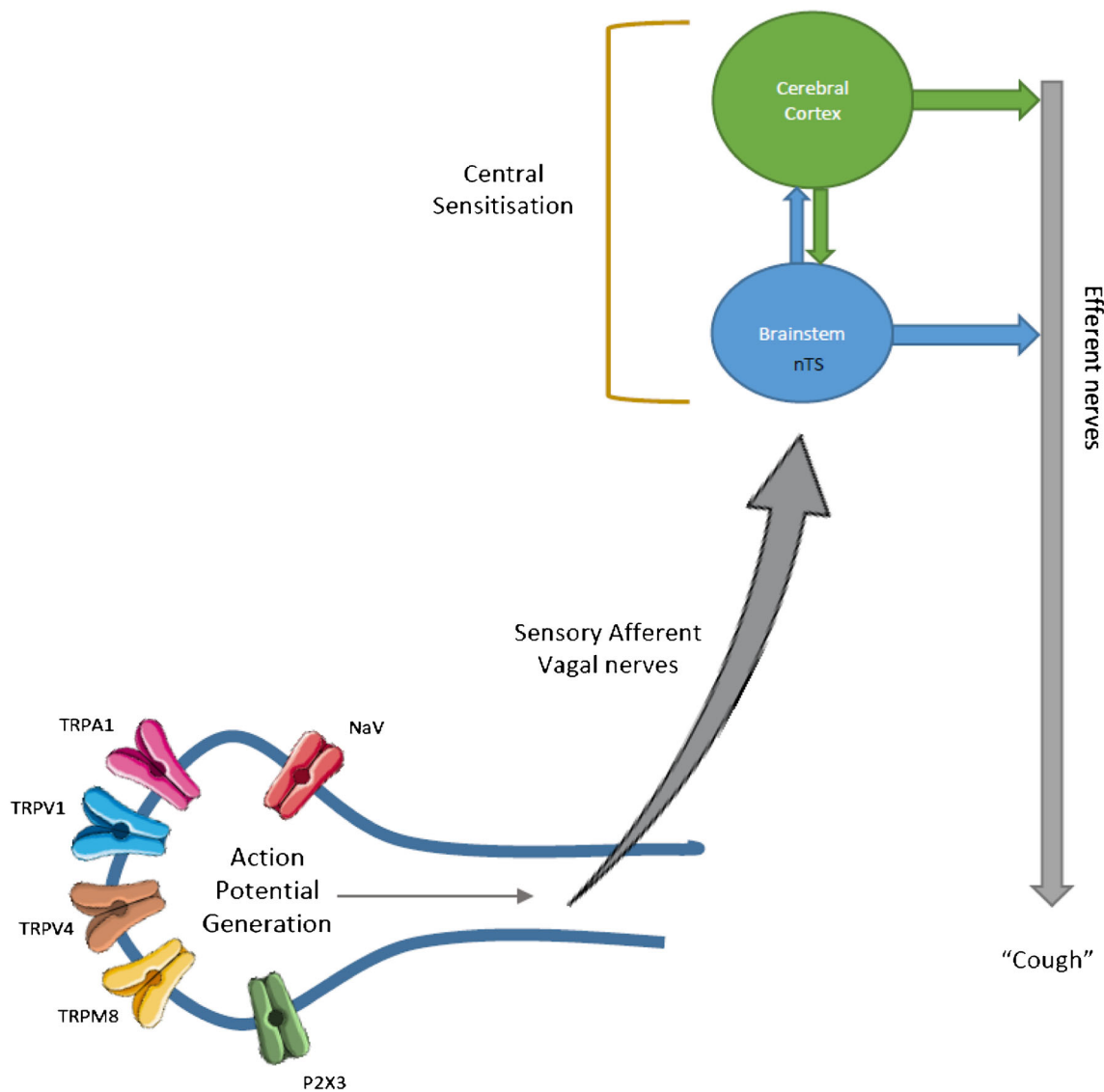


Fig. 1 Illustration of central and peripheral targets for antitussive agents. Current therapies in development that target the peripheral component of cough are transient receptor potential (TRP) channels (TRPV1, TRPV2, TRPV4 and TRPM8), purinergic receptors (P2X3) and voltage-gated sodium channels (NaV). Peripheral sensory neurons project to regions of the brainstem including the nucleus tractus solitarius (nTS).

Currently available centrally acting drugs target predominately μ -opioid receptors and N-methyl-D-aspartate (NMDA) receptors. Neurokinin-1 receptor (NK-1R) antagonists and agonists of the alpha7 (α 7) subtype of nicotinic acetylcholine receptors (nAChRs) are in early stages of evaluation. Figure adapted from Smith JA et al. *N Engl J Med* 2016; 375: 1544–51

inflammatory mediators (e.g. prostaglandins, bradykinin), low pH (< 5.9) and capsaicin [17, 18]. Capsaicin, the pungent ingredient of hot chilli peppers, readily induces cough on inhalation. Recent randomised clinical trials of the TRPV1 antagonists, SB-705498 and XEN-D0501, have shown marked reduction in capsaicin-induced cough reflex sensitivity but failed to reduce cough frequency in chronic cough patients [19, 20]. Based on evidence to date, it is unlikely that TRPV1 will be an effective therapeutic target for chronic cough.

TRPA1 The TRPA1 channel is activated by a range of physical and chemical factors including cold temperatures (<

17 °C), mechanical stimulation, inflammatory mediators (e.g. prostaglandins, bradykinin) and acrolein (a component of cigarette smoke) [21, 22]. In both animal models and healthy humans, the TRPA1 agonist cinnamaldehyde induces a dose-dependent cough response [23]. Animal studies have shown effectiveness of TRPA1 antagonists in reducing cough in response to tussive challenges. The TRPA1 antagonist HC-030031 reduces cough responses to both exogenous and endogenous activators of TRPA1 in guinea pigs [24, 25]. Glenmark have developed a TRPA1 inhibitor, GRC 17536, which showed inhibition of citric acid-induced cough in guinea pigs [26]. However, in a double-blinded placebo-controlled study, there was no reduction in

cough frequency over 24 h or no reduction in citric acid-induced cough [16].

TRPV4 TRPV4 is recognised as an osmosensor and responds to diverse stimuli including non-noxious temperatures (27–35 °C), shear stress and mechanical stimulation. TRPV4 activation has been shown to be associated with the release of adenosine 5'-triphosphate (ATP). The TRPV4 agonist, GSK1016790a, induced depolarisation of vagus nerves in both animal and human tissue and induced cough in conscious guinea pigs which was blocked by the TRPV4 antagonist HC067047 [27, 28]. The mechanism underlying TRPV4-induced ATP release has yet to be elucidated but a role for P2X3 receptors has been proposed [27]. However, a clinical trial with the TRPV4 antagonist, GSK2798745, was terminated early, presumably due to lack of efficacy.

TRPM8

TRPM8 is activated by cooling compounds such as menthol, icilin and eucalyptol. Although thought to play a role in cough induced by the inhalation of cold air, the TRPM8 agonist menthol also has protective and soothing effects and is a component of many over-the-counter cough remedies [29, 30]. The evidence for an antitussive role of TRPM8 is inconclusive and modulation of TRPM8 remains to be investigated as a potential target for chronic cough.

To date, pharmacological modulation of TRP channels for the treatment of cough has been disappointing. It remains to be seen if newer agents or strategies represent realistic options for the field of cough.

Purinergic Receptors

Purinergic receptors include subtypes of P2X ion channel receptors which are activated by ATP and have received considerable attention recently as potential targets for the treatment of pain and cough. ATP is released as a result of cellular stress due to injury or infection and in lung conditions such as asthma or COPD, elevated airway levels have been reported [31, 32]. P2X3 receptors expressed on airway sensory afferent nerves are considered to have a role in development of mechanical allodynia and hyperalgesia in animal models of pain [33]. Gefapixant (MK-7264), previously known as AF-219, is a P2X3 receptor antagonist recently shown to reduce cough frequency in a small placebo-controlled, crossover study of patients with unexplained chronic cough. High doses (600 mg bd) of the compound were used in this study, and all participants reported taste disturbances (hypogeusia or dysgeusia) [34]. A phase 2b study of gefapixant at lower doses has reported efficacy with less taste disturbance [35]. Currently, this compound represents the most promising therapeutic target in development.

Voltage-Gated Sodium Channels

Voltage-gated sodium channels (NaV) mediate the initiation and propagation of action potentials in afferent sensory nerves and represent a potential therapeutic target for cough. Lidocaine, a non-selective NaV channel blocker, attenuates the cough induced chemically by citric acid and capsaicin in awake guinea pigs [36, 37]. It has also been used clinically to alleviate cough and has been reported to be safe [38]. However, there are concerns regarding serious side effects of lidocaine including cardiac arrhythmias [39, 40]. A more targeted approach to NaV blockade seems sensible and of nine subtypes, NaV1.7, NaV1.8 and NaV1.9 are known to be expressed on vagal sensory neurons that innervate the lung and are likely to be involved in initiating cough [41, 42]. Selective inhibition of the NaV1.7 channel reduces excitability and sensory nerve activity and inhibits mechanically induced cough in guinea pigs [43, 44]. Recently, however, the results of a clinical trial of a novel blocker targeting NaV1.7 (GSK2339345) suggested no antitussive effect [45].

Treatments with a Central Site of Action

Sensory input from the upper and lower airways to the brainstem is relayed to higher brain regions where they modulate the varying sensory and motor responses that accompany coughing. The mechanisms underpinning the development of CHS may be akin to those described in pain where disordered peripheral and central neurophysiology are thought to be responsible for a variety of chronic pain syndromes. Chronic cough patients appear to have altered brain activity with evidence of both central sensitisation and dysfunctional inhibitory control [46]. A greater understanding of this disordered central processing is required and more selective targeting of pathways will be key to improved therapeutic control of troublesome coughing. Currently, the drugs used to treat or control cough that exert their therapeutic effect centrally do so at the expense of undesirable effects including sedation, respiratory depression and addiction. Here, we provide a brief overview of some of the common drugs with presumed central action used in treatment of chronic cough and highlight some more specific agents in early phase development.

Opiates (Morphine Sulphate/Codeine/Tramadol)

Opiates have been used to control cough for over 200 years. The notion that opiates are an effective cough suppressant has arisen from old studies in humans and animal studies [47, 48]. One commonly used drug is codeine, regarded as a weak opioid and found naturally in extracts of the poppy. The

evidence for its efficacy is conflicting and given its unfavourable side effect profile, codeine is not an ideal antitussive [49–51].

The only randomised controlled study of opiates conducted in patients with chronic intractable cough was undertaken by Morice and colleagues which showed that 5 mg of morphine sulphate twice daily for 2 months was an effective antitussive [52]. More recently, the antitussive properties of tramadol, an opioid medication with a similar structure to codeine and morphine, have been investigated in a pilot study and shown to be effective [53].

Dextromethorphan, which is an active ingredient of many popular brands of ‘over-the-counter’ cough preparations, is considered to have opiate properties. However, as its primary use is to control acute cough associated with the common cold, it is outside the scope of this review. However, a detailed review of this compound and other centrally acting drugs used for acute cough is available elsewhere [54].

GABA-Related Compounds (Gabapentin, Pregabalin)

Gabapentin and pregabalin are centrally acting neuromodulators primarily used for the treatment of neuropathic pain and epilepsy [55]. Both are structurally related to the inhibitory neurotransmitter, γ -aminobutyric acid (GABA), although their precise mode of action is unknown. They are known to modulate the release of neurotransmitters including substance P and may exert antitussive effects through attenuation of post-synaptic excitability [56]. One randomised placebo-controlled trial has shown that treatment of refractory chronic cough with gabapentin is both effective and well tolerated [57]. These efficacy claims have been supported in a recent systematic literature review [58].

Pregabalin has also been shown to improve cough in patients with what was termed sensory laryngeal neuropathy, and more recently, the combination of pregabalin and speech pathology treatment (SPT) was shown to be more effective in improving cough quality of life than SPT alone [59, 60]. Based on the existing evidence, gabapentin has recently received recommendation in the CHEST Guideline and Expert Panel Report as a treatment option for chronic cough [3].

N-Methyl-D-Aspartate (NMDA) Receptors

N-methyl-D-aspartate (NMDA) receptors are expressed within the peripheral and central nervous system and are believed to play a role in the synaptic transmission of cough-associated vagal afferents which project from the lungs to the brainstem [61]. Dextromethorphan is thought to exert its effect at least in part through activation of NMDA receptors [62]. Preclinical studies have also suggested the antitussive effects of memantine, an NMDA receptor channel blocker used in the treatment of Alzheimer’s

disease [63, 64]. Despite this promising data, a recent small open-label pilot study of a novel NMDA receptor antagonist, V3381, showed only marginal efficacy and concerns regarding tolerability [65].

Amitriptyline

Amitriptyline is used for the treatment of depression and lower doses are also prescribed for the treatment of chronic neuropathic pain. This drug is an inhibitor of serotonin re-uptake but may also have effects on the muscarinic, adrenergic and histaminergic systems [66, 67]. Treatment of chronic cough with amitriptyline is reported to reduce cough, and in an open-label study of treatment for 10 days, improvements in cough frequency and severity when compared with codeine were reported [68]. However, in another study conducted over a longer duration (2 months), the efficacy of amitriptyline in patients with chronic cough was short-lived, and the majority of subjects discontinued treatment [69].

Tachykinin Receptor Antagonists

The tachykinins, substance P, neurokinin A and neurokinin B are released both from the peripheral endings of afferent nerves (predominately C-fibres) and from central neural structures. The tachykinin receptor, neurokinin 1 receptor has gained attention as a target for chronic cough treatment. The NK1R antagonist, aprepitant, has been shown to reduce cough in patients with lung cancer [70]. A pilot study of serlopitant, a centrally acting NK1R antagonist, resulted in significant improvements in objective cough frequency and sustained reductions in daytime cough frequency [71]. Currently, serlopitant is undergoing further clinical evaluation (Table 1). Orvepitant, another NK1 receptor antagonist, is also in phase 2 clinical trials (Table 1).

Nicotinic Acetylcholine Receptors

Nicotinic acetylcholine receptors, or nAChRs, respond to the neurotransmitter acetylcholine and to nicotine and are found in the central and peripheral nervous system. In a recent study of healthy non-smokers, marked antitussive effects of nicotine-containing vapours produced by e-cigarettes were reported [72]. It is now believed that the $\alpha 7$ ($\alpha 7$) subtype of the nAChRs, activating GABAergic interneurons in the brainstem, is responsible for the antitussive effect of nicotine [73]. Bradanicline, a selective agonist of the $\alpha 7$ subunit is currently being tested as a potential treatment for chronic cough in a phase 2 randomised, double-blind, dose-escalation and crossover study.

Table 1 Compounds currently undergoing evaluation as treatments for chronic cough

Compound	Number	Placebo arm	Dose regime	Study type	Status	Primary outcome measure(s) (time frame)	Main results	Completion date
<i>TRPV1 antagonists</i>								
XEN-D0501	27	Matched placebo	4 mg, BID, Day 1–13, 4 mg OD Day 14	Phase 2, double-blind, crossover assignment, RCT	Completed	Change from baseline at the end of each treatment period in objective daytime cough frequency in COPD patients on XEN-D0501 compared to control	Failed to show improvements in spontaneous cough frequency	Completed: February, 2015
	18	Matched placebo	4 mg BID Day 1–13, 4 mg OD Day 14	Phase 2, double-blind, crossover assignment, RCT	Completed	Change from baseline at the end of each treatment period in objective daytime cough frequency on XEN-D0501 compared to control	Failed to show improvements in spontaneous cough frequency	Completed: May 2015
<i>TRPV4 antagonists</i>								
GSK2798745	17	Matched placebo	Oral, 4.8 mg day 1, 2.4 mg OD for 6 days	Phase 2, double-blind, crossover assignment, RCT	Active, Not recruiting	Number of coughs during daytime hours following 7 days of dosing	Terminated due to lack of efficacy	Expected: April, 2019
<i>P2X3 antagonists</i>								
Gefapixant (MK-7264) previously known as AF-219	30	Matched placebo	50 mg, BID, 16 days	Phase 2, double-blind, crossover RCT, dose-escalation study	Completed	Reduction in cough frequency (2 weeks)	Cough frequency was reduced by 75% when patients were allocated to AF-219 compared when allocated to placebo ($p=0.0003$)	Completed: February, 2016
	24	Matched placebo	Two 7.5 mg, four 7.5 mg or one 50 mg tablet, BID, 8 weeks	Phase 2, double-blind, parallel assignment, RCT	Completed	Reduction in cough frequency (8 weeks)	Results not posted	Completed: May, 2016
	50	Matched placebo	50 mg	Phase 2, double-blind, crossover RCT	Completed	Cough reflex sensitivity to capsaicin as measured by cough counts (4 h)	Results not posted	Completed: May, 2016
	23	Matched placebo	45 mg, BID	Phase 2, double-blind, parallel assignment, RCT	Completed	AE (up to 6 weeks)	Results not posted	Completed: June 2018
	36	Matched placebo	100 mg	Phase 2, double-blind, crossover RCT	Completed	Treatment discontinuations (up to 4 weeks) 1. Cough reflex sensitivity to capsaicin in participants who received gefapixant 100 mg and placebo (period 1 and period 2 combined) (avg: 1, 3 and 5 h post-dose)	Results posted on ClinicalTrials.gov	Completed: October 2016

Table 1 (continued)

Compound	Number	Placebo arm	Dose regime	Study type	Status	Primary outcome measure(s) (time frame)	Main results	Completion date
						2. Cough reflex sensitivity to citric acid in participants who received gefapixant 100 mg and placebo (period 1 and period 2 combined) [time frame: average of 1, 3 and 5 h post-dose] 3. Cough reflex sensitivity to ATP in participants who received gefapixant 100 mg and placebo (period 1 and period 2 combined) [time frame: average of 1, 3 and 5 h post-dose] 4. Cough reflex sensitivity to distilled water in participants who received gefapixant 100 mg and placebo (period 1 and period 2 combined) [time frame: average of 1, 3 and 5 h post-dose]		
	253	Matched placebo	7.5, 20 or 50 mg BID depending on randomisation	Phase 2, double-blind, Parallel RCT	Completed	1. Change from baseline in awake objective cough frequency after 12 weeks of treatment (day 84) (baseline visit (day -1), Day 84)	Results posted on ClinicalTrials.gov	Completed: November, 2016
	188	Matched placebo	45 mg, BID, 7 days	Phase 2a, double-blind, sequential assignment, RCT	Recruiting	1. Awake cough per hour on day 3		Expected: June, 2019
	720	Matched placebo	15 or 45 mg, BID	Phase 3, double-blind, parallel assignment, RCT	Recruiting	1. 24-h coughs per hour at week 12 (12 weeks)		Expected: July, 2020
	160	Matched placebo	15 or 45 mg (BID)	Phase 3, double-blind, parallel assignment RCT	Not yet recruiting	2. Percentage of participants experiencing at least one AE during treatment and follow-up (54 weeks) 3. Percentage of participants who discontinued treatment due to an AE (52 weeks)		Expected: October, 2020
	1290	Matched placebo	15 or 45 mg (BID)	Phase 3, double-blind, parallel, RCT	Recruiting	1. Number of participants experiencing at least one AE 2. Number of participants discontinuing study drug due to an adverse event 1. 24-h coughs per hour at week 24 (24 weeks)		Expected: January, 2021

Table 1 (continued)

Compound	Number	Placebo arm	Dose regime	Study type	Status	Primary outcome measure(s) (time frame)	Main results	Completion date
BAY1817080	108	Matched placebo	4 different doses	Phase 1/Phase 1, two-part, double-blind, RCT	Recruiting	<p>2. Percentage of participants experiencing at least one AE during treatment and follow-up (up to 54 weeks)</p> <p>3. Percentage of participants who have a study drug discontinued due to an AE (up to 52 weeks)</p> <p>1. Frequency of treatment-emergent adverse events in study part 1 [time frame: up to 5 weeks]</p> <p>2. Severity of treatment-emergent adverse events in study 1 [time frame: up to 5 weeks]</p> <p>3. Frequency of treatment-emergent adverse events in study part 2 [time frame: up to 12 weeks]</p> <p>4. Severity of treatment-emergent adverse events in study part 2 [time frame: up to 12 weeks]</p> <p>5. 24-h cough counts [time frame: at week 1 in period A]</p> <p>6. 24 h cough counts [time frame: at week 2 in period A]</p> <p>7. 24 h cough counts [time frame: at week 3 in period A]</p> <p>8. 24 h cough counts [time frame: at week 1 in period B]</p> <p>9. 24 h cough counts [time frame: at week 2 in period B]</p> <p>10. 24 h cough counts [time frame: at week 3 in period B]</p>		
BAY1902607	88	Matching placebo or Midazolam	4 different doses	Phase 1/phase 2, two-part, double-blind, RCT	Recruiting	<p>1. Frequency of treatment-emergent AE (approx. 5 weeks, part 1)</p> <p>2. Severity of treatment-emergent AE (approx. 5 weeks, part 1)</p> <p>3. Frequency of treatment-emergent AE (approx. 10 weeks, part 2)</p> <p>4. Severity of treatment-emergent AE (approx. 10 weeks, part 2)</p> <p>5. AUC of midazolam without BAY 1902607 (at day 1, part 1)</p> <p>6. AUC of midazolam in combination with BAY 1902607 (at day 13, part 1)</p>	Expected: July 2019	

Table 1 (continued)

Compound	Number	Placebo arm	Dose regime	Study type	Status	Primary outcome measure(s) (time frame)	Main results	Completion date
						7. Cmax of midazolam in combination with BAY 1902607 (at day 13, part 1) 8. Cmax of midazolam without BAY 1902607 (at day -1, part 1) 9. AUC (0-tlast) of midazolam in combination with BAY 1902607 (at day 13, part 1) 10. AUC (0-tlast) of midazolam without BAY 1902607 (at day -1, part 1) 11. Number of coughs experienced by patient (24 h, part 2)		
BLU-5937	90	Matched placebo	Oral tablet, single ascending doses (6 dose levels) or multiple ascending doses (3 dose levels)	Phase 1, double-blind, sequential assignment, RCT	Completed	Number and severity of treatment-emergent adverse events (TEAEs) (up to 48 h after the last dose)	Demonstrated good safety and tolerability profile. Antitussive effects observed with little alteration of taste perception	Completed: October, 2018
Voltage-gated sodium channels								
GSK2339345	16	Matched placebo	Oral inhalation, unit dose strength 1000 µg	Phase 2, double-blind, crossover, RCT	Completed	1. Total cough count over 8 h at visits 1, 2 and 3 (Part A) 2. Total cough count excluding transient coughs over 8 h at visits 1, 2 and 3	Failed to illustrate an antitussive response	Completed: October, 2014
NK-1 receptor antagonist								
Serlopitant (a.k.a. VPD-737)	185	Matched placebo	5 mg tablet	Phase 2, double-blind, RCT	Completed	1. Change from baseline in 24-h objective cough frequency		Completed: September, 2018
Orvepitant	244	Matched placebo	10–30 mg, OD, 12 weeks	Phase 2, double-blind, parallel assignment, RCT	Recruiting	1. Awake objective cough frequency		Expected: December, 2018
Nicotinic receptor agonist								
Bradanicline	48	Matched placebo	Oral, dose escalation, 3 different doses	Phase 2, double-blind, sequential assignment, RCT	Not yet recruiting	1. Awake coughs per hour at days 7, 14, 21, 43, 50, 57		Expected: October, 2019

Table 1 (continued)

Compound	Number	Placebo arm	Dose regime	Study type	Status	Primary outcome measure(s) (time frame)	Main results	Completion date
Other drugs in clinical trials for chronic cough treatment								
Azithromycin (Zithromax)	50	–	250 mg, 3 times/week	Phase 3, single group assignment, open label	Unknown	1. Change from baseline in cough on the Leicester Cough Questionnaire (LCQ) score at week 12 (12 weeks after treatment start)	Failed to show improvements in LCQ compared to placebo	Expected: December, 2016
PA-101 (RVT-1601)	52	Matched placebo	Inhalation, 40 mg, 3 times daily	Phase 2, double-blind, crossover RCT	Completed	2. Change of cough-specific quality of life 1. Cough Frequency	No treatment benefit for PA101 observed in chronic idiopathic cough patients	Completed: February, 2016

OD once daily, *BID* twice daily, *AE* adverse event, *RCT* randomised controlled trial

Other Medications for the Treatment of Chronic Cough

Azithromycin

A study by Hodgson et al. investigated the potential effects of azithromycin, a macrolide, on chronic cough [74]. Treatment with azithromycin for 8 weeks failed to improve health status of chronic cough patients when compared with placebo. This study however suggested there may be potential benefit of macrolide use in the treatment of cough associated with asthma.

PA-101 Cromolyn Sodium Formulation

PA-101 is a novel formulation of cromolyn sodium and thought to act as a mast cell stabiliser. It has undergone study in idiopathic chronic cough and in patients with pulmonary fibrosis, where cough is often a troublesome symptom. In a randomised placebo-controlled trial where PA-101 was delivered via a high-efficiency eFlow nebuliser, no improvement in chronic idiopathic cough patients was observed [75]. However, the study did report reduction in cough in patients with idiopathic pulmonary fibrosis. These findings suggest some degree of heterogeneity in mechanisms responsible for CHS.

Conclusion

Chronic cough is an extremely common clinical problem that can exist in isolation or as the dominant symptom for patients across a wide range of common chronic respiratory conditions. Chronic cough is often difficult to control, largely because of the lack of effective medicines. Over the last two or three decades, there has been enormous progress in terms of optimising clinical evaluation, and gaining consensus on how to best manage patients with troublesome cough. There have also been important developments in the mechanistic understanding of cough and elucidating the associated neurobiology, and with this have come a number of promising therapeutic targets. Some of these are at advanced stages of clinical development with drugs targeting the P2X3 receptor most promising. However, many challenges remain and partnerships are required, bringing clinicians and patients, scientists and industry together with government agencies to overcome these. This approach is likely to speed up drug discovery with the ultimate aim of providing better treatments for cough patients.

Compliance with Ethical Standards

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