Anatomy and neuro-pathophysiology of the cough reflex arc

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Abstract
Coughing is an important defensive reflex that occurs through the stimulation of a complex reflex arc. It accounts for a significant number of consultations both at the level of general practitioner and of respiratory specialists. In this review we first analyze the cough reflex under normal conditions; then we analyze the anatomy and the neuro-pathophysiology of the cough reflex arc. The aim of this review is to provide the anatomic and pathophysiologic elements of evaluation of the complex and multiple etiologies of cough.

Keywords: Cough, C-Fibers, Rapidly adapting receptors, Reflex arc, Slowly adapting stretch receptors

Review

Cough is one of the most common symptoms for which outpatient care is asked [1], accounting for up to 40% of the practice activity [2]. Coughing is an important defensive reflex that enhances clearance of secretions and particulates from the airways and protects from aspiration of foreign materials occurring as a consequence of aspiration or inhalation of particulate matter, pathogens, accumulated secretions, postnasal drip, inflammation, and mediators associated with inflammation. Under normal conditions cough serves an important protective role in the airways and lungs, but in some conditions it may become excessive and nonproductive, and is troublesome and potentially harmful to the airway mucosa. These contrasting consequences of coughing can be attributed to the parallel afferent pathways regulating this important defensive reflex of the airways.

Each cough occurs through the stimulation of a complex reflex arc. This is initiated by the irritation of cough receptors which are found in the trachea, main carina, branching points of large airways, and more distal smaller airways; also, they are present in the pharynx. Laryngeal and tracheobronchial receptors respond to both mechanical and chemical stimuli. Chemical receptors sensitive to acid, heat, and capsaicin-like compounds trigger the cough reflex via activation of the type 1 vanilloid (capsaicin) receptor [3-5]. In addition, more airway receptors are in the external auditory canals, eardrums, paranasal sinuses, pharynx, diaphragm, pleura, pericardium, and stomach. These are probably mechanical receptors only, which can be stimulated by triggers such as touch or displacement.

Impulses from stimulated cough receptors traverse an afferent pathway via the vagus nerve to a ‘cough center’ in the medulla, which itself may be under some control by higher cortical centers. The cough center generates an efferent signal that travels down the vagus, phrenic, and spinal motor nerves to expiratory musculature to produce the cough.

Therefore, the cough reflex arc is constituted by:

1. **Afferent pathway**: Sensory nerve fibers (branches of the vagus nerve) located in the ciliated epithelium of the upper airways (pulmonary, auricular, pharyngeal, superior laryngeal, gastric) and cardiac and esophageal branches from the diaphragm. The afferent impulses go to the medulla diffusely.

2. **Central Pathway (cough center)**: a central coordinating region for coughing is located in the upper brain stem and pons.

3. **Efferent pathway**: Impulses from the cough center travel via the vagus, phrenic, and spinal motor nerves to diaphragm, abdominal wall and muscles. The nucleus retroambiguus, by phrenic and other spinal motor nerves, sends impulses to the inspiratory and expiratory muscles; and the nucleus
ambiguus, by the laryngeal branches of the vagus to the larynx.

The terminations of the vagal afferents are found in abundance in the airway mucosa and in the airway wall from the upper airways to the terminal bronchioles and lung parenchyma. Afferent neuronal subtypes can be identified based on their physicochemical sensitivity, adaptation to sustained lung inflation, neurochemistry, origin, myelination, conduction velocity (A-fiber, > 3 m/s; C-fiber, < 2 m/s), and sites of termination in the airways. These attributes can be used to identify at least three broad classes of airway afferent nerves:

1. Rapidly Adapting Receptors (RAR)
2. Slowly Adapting Stretch Receptors (SARs)
3. C-Fibers

Rapidly adapting receptors (RAR)
Functional studies of RARs suggest that they terminate within or beneath the epithelium of both intrapulmonary and extrapulmonary airways, but primarily the intrapulmonary airways. RARs are differentiated from other airway afferents by their rapid adaptation (in 1 - 2 seconds) to sustained lung inflations [6-19]. Other distinguishing properties of RARs include their sensitivity to lung collapse and/or lung deflation, their responsiveness to alterations in dynamic lung compliance (and thus their sensitivity to bronchospasm), and their conduction velocity (4 to 18 m/s), which is suggestive of myelinated axons. The sustained activation of RARs produced by dynamic lung inflation, bronchospasm, or lung collapse indicates that the adaptation of RARs is not attributable to an electrophysiologic adaptation. Perhaps RARs are thus better defined as dynamic receptors that respond to changes in airway mechanical properties (e.g. diameter, length, and interstitial pressures). RARs are sporadically active throughout the respiratory cycle, are activated by the dynamic mechanical forces accompanying lung inflation and deflation, and become more active as the rate and volume of lung inflation increase. RARs are activated by stimuli that evoke bronchospasm or obstruction resulting from mucus secretion or edema. Substances such as histamine, capsaicin, substance P, and bradykinin activate RARs in a way that can be markedly inhibited or abolished by preventing the local end-organ effects that these stimuli produce (e.g. bronchospasm and mucus secretion). RAR activation initiates reflex bronchospasm and mucus secretion through parasympathetic pathways. RARs can also respond to stimuli that evoke cough and fulfill many criteria for mediating cough. Further evidence for their role in coughing comes from studies of vagal cooling, which blocks cough at temperatures that selectively abolish activity in myelinated fibers (including RARs) while preserving C-fiber activity. RARs may act synergistically with other afferent nerve subtypes to induce coughing.

Slowly adapting stretch receptors (SARs)
SARs are highly sensitive to the mechanical forces that are put on the lung during breathing. SAR activity increases during inspiration and peaks just prior to the initiation of expiration [13]. SARs are thus thought to be the afferent fibers involved in the Hering-Breuer reflex, which terminates inspiration and initiates expiration when the lungs are adequately inflated. SARs can be differentiated from RARs in some species based on action potential conduction velocity, and in most species by their lack of adaptation to sustained lung inflations. SARs may also be differentially distributed throughout the airways: they appear to terminate primarily in the intrapulmonary airways. SARs also differ from RARs with respect to the reflexes they precipitate. SAR activation results in the central inhibition of respiration and the inhibition of the cholinergic drive to the airways, leading to decreased phrenic nerve activity and decreased airway smooth muscle tone (due to a withdrawal of cholinergic nerve activity) [14]. The sensory terminals of SARs assume a complex and varying position within the airway wall: most of these SARs are found in the peripheral airways (associated with alveoli or bronchioles). Occasionally, SAR dendritic arbors are associated with the bronchiolar smooth muscle. SARs may facilitate coughing by a central cough network via activation of brainstem second-order neurons of the SAR reflex pathway.

C-fibers
The majority of afferent nerves innervating the airways and lungs are unmyelinated C-fibers. They are similar in many ways to the unmyelinated somatic sensory nerves innervating the skin, skeletal muscle, joints, and bones that respond to noxious chemical and mechanical stimuli (called nociceptors). In addition to their conduction velocity (< 2 m/s), airway vagal afferent C-fibers are distinguished from RARs and SARs by their relative insensitivity to mechanical stimulation and lung inflation. C-fibers are further distinguished from RARs by the observation that they are directly activated by bradykinin and capsaicin, not indirectly through effects on smooth muscle or the airway vasculature. Moreover, prostaglandin E2, adrenaline, and adenosine, which by bronchodilating the airways might inhibit RAR activation by bradykinin and capsaicin, actually sensitize C-fibers to capsaicin and bradykinin through direct effects on their peripheral nerve terminals [15-17]. Morphologic studies in rats and in guinea pigs have revealed that afferent C-fibers innervate the airway epithelium as well as other
effectors structures within the airway wall. C-fibers may 
synthesize neuropeptides that are subsequently trans-
ported to their central and peripheral nerve terminals. 
C-fibers are generally quiescent throughout the respira-
tory cycle but are activated by chemical stimuli such as 
capsaicin, bradykinin, citric acid, hypertonic saline solu-
tion, and sulfur dioxide (SO₂). Reflex responses evoked 
by C-fiber activation include increased airway parasym-
pathetic nerve activity, and the chemoreflex, character-
ized by apnea (followed by rapid shallow breathing), 
bradycardia, and hypotension. Stimulants of C-fibers 
such as capsaicin, bradykinin, SO₂, and citric acid evoke 
cough in conscious animals and in humans, and capsai-
icin desensitization abolishes citric acid-induced cough-
ing in guinea pigs.

Sex-related differences in cough reflex sensitivity ex-
plain the observation that women are more likely than 
men to develop chronic cough [18-20].

The mechanical events of a cough can be divided into 
three phases [21]:

1. Inspiratory phase: Inhalation, which generates the 
   volume necessary for an effective cough.
2. Compression phase: Closure of the larynx combined 
   with contraction of muscles of chest wall, 
diaphragm, and abdominal wall result in a rapid rise 
in intrathoracic pressure.
3. Expiratory phase: The glottis opens, resulting in high 
   expiratory airflow and the coughing sound. Large 
   airway compression occurs. The high flows dislodge 
mucus from the airways and allow removal from the 
   tracheobronchial tree.

The specific pattern of the cough depends on the site 
and type of stimulation. Mechanical laryngeal stimulation 
results in immediate expiratory stimulation (some-
times termed the expiratory reflex), probably to protect 
the airway from aspiration; stimulation distal to the lar-
ynx causes a more prominent inspiratory phase, presum-
able to generate the airflow necessary to remove the 
stimulus.

During vigorous coughing, intrathoracic pressures may 
reach 300 mm Hg and expiratory velocities approach 
800 kilometers per hour [22]. While these pressures and 
velocities are responsible for the beneficial effects of 
cough on mucus clearance, they are also responsible for 
many of the complications of cough, including exhaus-
tion, self-consciousness, insomnia, headache, dizziness, 
musculoskeletal pain, hoarseness, excessive perspiration, 
urinary incontinence [23]. Cough-induced rib fractures 
are another painful and potentially serious complication 
of chronic cough. Fractures often involve multiple ribs, 
particularly ribs five through seven. Women with 
decreased bone density are at the greatest risk of this 
complication; however, fractures can occur in patients 
with normal bone density as well [24].

A nonproductive cough is a well-recognized complica-
tion of treatment with angiotensin converting enzyme 
(ACE) inhibitors, occurring in up to 15% of patients trea-
ted with these agents [25]. Although the pathogenesis of 
the cough is not known with certainty, it has commonly 
been hypothesized that accumulation of bradykinin, 
which is normally degraded in part by ACE, may stimu-
late affecter C-fibers in the airway [26].

The important observation that cough does not appear 
to occur with increased frequency in patients treated with 
amphotericin II receptor antagonists (which do not increase 
kinin levels) is consistent with the kinin hypothesis.

Lesions that compress the upper airway, including ar-
teriovenous malformations and retrotracheal masses, 
may present with chronic cough [27-29]. Cough can also 
be a symptom of tracheobronchomalacia, which results 
from loss of rigid support of the large airways and in-
spiratory collapse, and is usually seen in conjunction 
with obstructive lung disease in patients with a history of 
cigarette smoking [30].

Laryngeal sensory neuropathy has been identified as 
the cause of chronic cough in 18 of 26 patients with 
acute onset of cough that was often associated with lar-
yngospasm or throat clearing [31]. Chronic tonsillar en-
largement has been proposed as a cause of chronic 
cough, but clinical evidence of this association is limited. 
One series of 236 patients referred for evaluation in a 
specialized clinic noted tonsillar enlargement in the ab-
ence of other known causes of chronic cough in 8 
(3.4%) individuals [32]. Following tonsillectomy, these 
patients had decreased cough sensitivity and significantly 
improved symptom control. At this moment, these 
observations need further investigations.

Irritation of the external auditory canal by impacted 
foreign bodies or cerumen is another unusual cause of 
chronic dry cough [33]. The etiology of the ‘ear-cough’ 
(or oto-respiratory) reflex is related to stimulation of the 
auricular branch of the vagus nerve (Arnold’s nerve) 
[34,35]. Another rare cause of chronic cough is Holmes-
Adie syndrome due to autonomic dysfunction affecting 
the vago nerve [36] patients present anisocoria, abnor-
mal deep tendon reflexes, and patchy areas of hyperhi-
drosis or anhidrosis.

In adults, habit (also known as ‘psychogenic’) cough 
may rarely be the cause of a chronic cough that remains 
troublesome despite a thorough evaluation, including 
ruling out tic disorders.

Differences among several sites from which cough 
stimuli can originate may result in variations in the 
sounds and patterns of coughing.

Laryngeal stimulation produces a choking type of 
cough without a preceding inspiration.
Inadequate mucociliary clearance mechanisms (as in bronchiectasis or cystic fibrosis) may produce a pattern of coughing with less violent acceleration of air and a sequence of interrupted expirations without any intervening inspiration.

Awareness of cough varies considerably: it can be distressing when it appears suddenly, especially if associated with discomfort due to chest pain, dyspnea, or copious secretions, while a cough that develops over decades (e.g. in a smoker with chronic bronchitis) may be hardly noticeable or may be considered normal by the patient.

Since cough is an important defensive reflex, required to maintain the health of the lungs, people who do not cough effectively are at risk of atelectasis, recurrent pneumonia, and chronic airways disease from aspiration and retention of secretions. Many disorders can impair the ability to cough effectively, which may result in persistent cough. The elderly, newborns, lung transplant recipients, and patients with paralysis or neuromuscular disorders have a poorly developed and/or compromised cough reflex, and are rendered highly susceptible to lung infections and aspiration pneumonia. Patients with paralysis or neuromuscular disorders (including rib fractures and chest wall deformities) may not generate the expected coughing with less violent acceleration of air and secretion movement necessary for effective coughing [38,39].

Conclusions
Based on this complex mechanism, the treatment often requires an early symptomatic approach in order to prevent the vicious cycle of cough perpetuating cough and a chronic disorder which has been reported to be the fifth most common complaint seen by primary care physicians in the world and the third in Italy [41].

Competing interests
The authors declare that they have no competing interests.

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