Mechanisms of respiratory innervation during embryonic development

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Abbreviations: ASM, airway smooth muscle; BDNF, brain derived neurotrophic factor; GDNF, glial cell-derived neurotrophic factor; NEBs, neuroendocrine bodies; NGF, nerve growth factor; NT, neurotrophin

During embryogenesis, the development of the respiratory tract is closely associated with the formation of an extensive neuronal network. While the topic of respiratory innervation is not new, and similar articles were published previously, recent studies using animal models and genetic tools are breathing new life into the field. In this review, we focus on signaling mechanisms that underlie innervation of the embryonic respiratory tract.

Introduction

The mammalian respiratory tract consists of the trachea and lung. It arises from ventral foregut endoderm.1 After progenitor specification, the lung primordia bifurcate ventrally-laterally to form two primary lung buds. These buds continue to invade the surrounding mesenchyme, elongate and branch to ultimately form a tree-like structure of epithelium tubules and alveoli. Along with lung bud formation, the trachea forms ventrally and separates from the primitive esophagus that is formed in the dorsal side of the foregut. Meanwhile, the mesenchyme forms from lateral plate mesoderm and gives rise to other cell types in the respiratory tract, such as airway smooth muscle (ASM), trachea cartilage, lymphatics, and blood vessels.1 In mice, the trachea and lung form around E9, and the respiratory tract continues to develop after birth before reaching maturity around 3–4 weeks postnatally.1,3

During respiratory tree development, a complex neuronal network forms.4 This network includes axons from extrinsic neurons, whose cell bodies are located outside of the respiratory tract, and intrinsic neurons whose cell bodies reside in the trachea and major bronchi and cluster to form ganglia (Fig. 1).5,6 Previous studies identified the location of extrinsic neuronal cell bodies by using a combination of retrograde/anterograde labeling techniques, immunohistology, and microscopy.7,12 Extrinsic neurons within the dorsal and ventral respiratory nuclei in the medulla oblongata and within the jugular and nodose ganglia supply parasympathetic efferents and most of the sensory afferents, respectively (Fig. 1C).6,13-15 These axons travel along the vagus nerve to innervate ASM and neuroendocrine bodies (NEBs) in the lung epithelium (Fig. 1C).14,16 Some efferents also connect to intrinsic neurons that provide post-ganglionic parasympathetic input to the trachea and bronchi (Fig. 1B).6 In addition, sensory afferents from the dorsal root ganglia connects with thoracic ganglia to supply sympathetic innervation to the blood vessels and submucosal gland (Fig. 1A).6 This complex neural network functions to control breathing, smooth muscle tone, and mucous secretion, and to trigger reflexes such as cough.13-15

Although lung innervation has been described by previous studies in humans, primates, rodents, and several other animals, signals that regulate respiratory neurogenesis are not fully understood.17-20 Here, we review current knowledge on mechanisms of respiratory neurogenesis during embryonic development. Most studies on neurogenic signals were performed in mice, where ASM and NEBs are the major targets of innervation. This review is designed to highlight key findings in the development of respiratory tract innervation rather than a comprehensive overview of every study published in this field. We apologize to authors whose contribution is not acknowledged.

Intrinsic Neurogenesis Within the Respiratory Tract

Earlier observations in the airways of humans and other species show that intrinsic neurons express neural crest cell markers, suggesting their neural crest origin.21,22 Follow-up studies using engrafment of avian neural tissues and lineage labeling in mouse embryos definitively prove that intrinsic neurons in the respiratory tract are exclusively derived from vagal neural crest cells.23,24 These neural crest cells generate both neurons and glial cells that cluster to form ganglia, mostly found in the dorsal trachea and upper respiratory tract (Fig. 1B).23,24 As the size of the airway tapers off along the proximal-distal axis of the respiratory tree, there are fewer intrinsic neurons.23,24 In mice, a small number of intrinsic neurons are located in the secondary and tertiary bronchi with little to none in the distal lung (Fig. 1B).

In addition to intrinsic neurons in the respiratory tract, vagal neural crest cells also give rise to enteric neurons in the gastrointestinal tract.25,26 These two groups of neural crest cells likely

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migrate together initially. Upon separation of the trachea from the esophagus at E10.5 in mouse embryos, neural crest cells that migrate into the space between the esophagus and the trachea begin to take different paths.23,24

Innervation of the gastrointestinal tract by enteric neural crest cells is well characterized compared with the respiratory innervation by intrinsic neurons. Expressed in the enteric wall, glial cell derived neurotrophic factor (GDNF) is an essential chemoattractant for enteric neural crest cells.25-30 GDNF belongs to a family that also includes neurturin, artemin, and persephin. Each family member binds to unique GDNF family co-receptors 1, 2, 3 and 4 (GFRα1–4), respectively.31,32 When the GDNF family ligand binds to the GFRα co-receptor, the common tyrosine kinase receptor RET is recruited for downstream signaling.31,32 Enteric neural crest cells predominantly express GFRα1.23,30 Consistently, genetic disruption of GDNF or RET diminishes the migration of enteric neural crest cells and subsequent formation of the enteric nervous system.27-30,33

In contrast, GDNF is not expressed in the trachea, and GDNF deficiency has no effect on the respiratory intrinsic neurons.23,24 This suggests that respiratory neural crest cells depend on different chemo-attractants for migration. Further characterization of other GDNF family member and receptor expression shows that neurturin is expressed in the respiratory tract, and the respiratory neural crest cells express both GFRα1 and GFRα2.23 However, the neurturin knockout mouse embryos have normal airway intrinsic innervation.23 In addition, loss of RET function, which demolishes the signaling activity of all GDNF family members, does not affect the number of respiratory intrinsic neurons at E14.5.24 These findings indicate that the respiratory neural crest cells are independent of the GDNF family for migration into the respiratory tract. However, Ret deficiency leads to a 50% reduction in the size of the ganglia at E18.5.23 Thus, GDNF family signaling, through the Ret receptor, likely plays a role in the survival, proliferation and/or differentiation of these neural crest cells after they reach the respiratory tract. Signals for the migration of neural crest cells in the respiratory tract have yet to be identified.

In addition to the difference in essential migratory signals between the respiratory and gastrointestinal neural crest cells, cells that migrate into these two organs also differ in their progenitor status. Recently, using avian embryos, the Burns lab has shown that after graft into the vagal neural tube, neural crest cells that have already reached the gut can still migrate into the lung and the gut.24 However, neural crest cells collected from the lung fail to migrate after graft.24 These findings suggest that respiratory neural crest cells are committed once they reach their destination whereas gastrointestinal neural crest cells maintain their migratory potentials even after they migrate into the gut.

**ASM Innervation by Extrinsic Neurons**

While intrinsic neurons likely provide parasympathetic innervation to the trachea and main bronchi, a large body of evidence indicates that intrinsic neurons do not contribute significantly to lung innervation (Fig. 1C). First, most intrinsic neurons are located in the trachea and main bronchi.20,23,24 Only a small number of intrinsic neurons are found in the secondary and tertiary bronchi with little to none in the distal lung.23,24 Second, intrinsic neurons have short axons, suggesting that they function locally.35 Third, although RET mutant embryos have a reduction in the size of resident ganglia at E18.5, they have no defects in lung innervation.35 Finally, vagotomy results in an almost complete loss of innervation of airway targets, such as NEBs.36 Collectively, these findings indicate that the lung is innervated predominantly by extrinsic neurons. However, the role of intrinsic lung innervation cannot be fully elucidated until functional data are obtained from animal models selectively deficient in intrinsic neurons.

During embryogenesis, the outgrowth of axons into the distal lung is closely associated with the formation of ASM. Studies in the fetal porcine and human lung show that the developing tubules are covered in a layer of ASM and ensheathed in a newly formed neuronal network.19,20,37 Two large nerve trunks run the length of the bronchial tree. They give rise to a network of bundles, with fine fibers covering up to the growing tips of the airways.19,20 This close temporal and spatial relationship between ASM formation and axon outgrowth suggest an ASM-derived trophic mechanism for innervation.

The extrinsic neurons depend on the nerve growth factor (NGF) family for ASM innervation. The NGF family includes NGF, brain-derived neurotrophic factor (BDNF), neurotrophin 3 (NT3), and NT4.38,39 They signal through high-affinity tyrosine kinase Trk receptors with relative selectivity: TrkA for NGF, TrkB for BDNF and NT4, and TrkC for NT3. Both BDNF and NT4 are expressed by embryonic ASM.40 The BDNF knockout embryos have reduced axon branches and shortened axons targeting the ASM without any change in lung morphogenesis or ASM differentiation.40 Thus, BDNF serves as a target-derived neurotrophic factor for ASM innervation by extrinsic neurons during embryogenesis.40 These findings also provide further evidence that intrinsic neurons and extrinsic neurons require distinct neurogenic signals for innervation of the respiratory tract.

To ensure appropriate innervation, BDNF expression needs to be temporally coordinated with ASM differentiation. Notably, BDNF mRNA is expressed as early as E11.5 in the lung mesenchyme prior to its differentiation into ASM.40 To coordinate, a post-transcriptional regulation is at play to repress the translation of BDNF mRNA until ASM is formed. One of the mechanisms of post-transcriptional regulation of BDNF expression is through a microRNA, miR-206. MiR-206 is expressed in lung mesenchyme, and its expression is downregulated upon ASM differentiation. In addition, miR-206 targets BDNF mRNA for degradation.40 Furthermore, the miR-206 knockout mice exhibit premature airway innervation.40 Collectively, these findings support miR-206 as a post-transcriptional regulator for coordinated BDNF protein expression, ASM differentiation, and ASM innervation.

Additionally, NT4, which binds to the same TrkB receptor as BDNF, may play a redundant role in ASM innervation. Consistent with this hypothesis, previous studies showed that mice deficient in both BDNF and NT4 have a diminished number of neurons in the nodose-petrosal ganglion complex, one of
Innervation is involved in the pathogenesis of respiratory diseases. Identification of the signals required for disease-related neural plasticity will likely provide groundwork for identification of new therapeutic targets.

Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

NEBs Innervation

NEBs are specialized clusters of pulmonary neuroendocrine cells that originate from the airway epithelium in the mammalian lung. Whereas solitary pulmonary neuroendocrine cells are found in the trachea, bronchioles, and terminal airways, NEBs are localized only in the intrapulmonary airways. The cytoplasm of NEB contains secretory granules that are loaded with bioactive molecules, such as neuropeptide, monoamines, and purine transmitters. NEBs are usually found at or near the bifurcation sites of the airway and juxtapose invariant Clara cells, a cell population with stem-cell like properties. Due to these unique structural, chemical, and positional properties, NEBs have been speculated to function as mechanoreceptors and play a role in O2 sensing and regeneration of the distal pulmonary epithelium. However, precise roles of NEBs remain elusive.

NEBs in the lung epithelium are innervated by a mixture of sensory and cholinergic nerves. These sensory afferents originate from the nodose ganglia and dorsal root ganglia, and the cholinergic efferents come from the brain stem and intrinsic ganglia. NEB innervation by P2X3+ and P2X7+ axons is reduced in NT4−/− mice, indicating that NT4 is required for NEB purinergic innervation. Whether NEB innervation regulates the role of NEBs during homeostasis and regeneration of the lung epithelium is unknown.

Concluding Remarks

Despite recent progress on the mechanisms of respiratory innervation, future studies are required to fully understand these processes. For example, additional signals that control NEB innervation remain to be identified. In addition, mechanisms underlying ASM innervation in postnatal life need to be further investigated. Mounting evidence indicates that altered innervation is involved in the pathogenesis of respiratory diseases. Identification of the signals required for disease-related neural plasticity will likely provide groundwork for identification of new therapeutic targets.
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