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Understanding the development of the respiratory glands.

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17 Department of Craniofacial Development and Stem Cell Biology, Dental Institute,
18 King's College London, Floor 27 Guy's Tower, London Bridge, SE1 9RT London,
19 UK.
20

21
22 § author for Correspondence
23

24 abigail.tucker@kcl.ac.uk
25
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Abstract

The submucosal glands (SMGs) of the respiratory system are specialized structures essential for maintaining airway homeostasis. The significance of SMGs is highlighted by their involvement in respiratory diseases such as cystic fibrosis, asthma and chronic bronchitis, where their phenotype and function are severely altered. Uncovering the normal development of the airway SMGs is essential to elucidate their role in these disorders, however, very little is known about the cellular mechanisms and intracellular signals involved in their morphogenesis. This review describes in detail the embryonic developmental journey of the nasal SMGs and the postnatal development of the tracheal SMGs in the mouse. We also review the current knowledge of the genes involved in SMG organogenesis, in the hope of stimulating further research into the mechanisms required for successful SMG patterning and function.

Introduction

Airway mucus plays a vital role in maintaining respiratory homeostasis as it provides the first line of defence against airborne irritants in the nasal cavity, and is essential in the mucociliary process, ensuring no foreign particles reach the lungs. Not only does its thick consistency trap foreign particles; its protein constitution additionally contains bactericidal enzymes, reducing the risk of infection. In humans, 5% of the respiratory mucus is secreted by goblet cells, which are simple columnar cells found scattered within the respiratory epithelium (RE) (Reid, 1960). The nucleus of these cells is found basally, while the apex of the cell is rich in mucin secretory granules (Freeman, 1962). The remaining 95% of the mucus secretion is released by the nasal and tracheal submucosal glands (SMGs) that lie in the underlying submucosal layer beneath the RE (Reid, 1960).

The importance of understanding SMG development and function is increasingly emphasised as more and more evidence shows their involvement in an array of severe and life threatening respiratory diseases, with SMG hyperplasia and mucus hypersecretion common to most (Aikawa et al., 1992; Oppenheimer & Esterly, 1975; Reid, 1960). The most common hyper-secretory diseases in which SMGs play a role are cystic fibrosis (CF), asthma and chronic bronchitis. One of the earliest steps in CF pathophysiology is the occlusion and dilation of SMG ducts, evident in third trimester neonates with CF (Ornoy et al., 1987). Infants of CF then show early onset of hyperplasia and hypertrophy of SMGs (Oppenheimer & Esterly, 1975; Sturgess & Imrie, 1982). Cystic fibrosis Transmembrane conductance regulator (CFTR), the defected gene in patients suffering from cystic fibrosis (CF), is primarily expressed in the serous cells and some of the ductal cells of the SMGs (Engelhardt et al., 1992). While this suggests a functional defect in the glands of CF patients, defects in the developmental patterning of the SMGs in a CF mouse model have also been reported, suggesting an additional role during gland initiation (Borthwick et al., 1999).

Hyperplasia of the bronchial SMGs is a well defined feature of asthma airways with gland:bronchial wall area ratio being significantly increased in both chronic asthma sufferers and those that die of sudden attack, compared to control patients (Aikawa et al., 1992; Rogers, 2004). In sufferers of chronic bronchitis, the bronchial SMGs are hypertrophic, displaying mucus hyper-secretion and an increase in mucous cells compared to serous cell counterparts (Reid, 1960; Rogers, 2008).

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Considering the critical involvement of SMGs in the above respiratory conditions, little is known about the developmental and homeostatic mechanisms of the respiratory glands. To understand disease states, it is first important to have a good understanding of normal SMG development and the signalling factors, modulators and pathways involved in gland morphogenesis.

Temporal Localisation of the SMGs

The SMGs are hidden in the submucosal connective tissue running along the conductive airways of the respiratory system. Nasal SMGs are organised into various groups with names according to their location in the medial and lateral nasal cavity walls (Cuschieri & Bannister, 1974; Grüneberg, 1971), while in the trachea, SMGs lie in the submucosal layer between the cartilage rings and smooth muscle (Finkbeiner, 1999). In the mouse, SMGS are found in the upper trachea with the highest volume of tracheal glands found at the anterior boundary between the cricoid cartilage (CC) of the larynx and the first tracheal cartilage ring (C1) and extending no further than the sixth cartilaginous ring (C6) (Borthwick et al., 1999; Rawlins & Hogan, 2005). In humans these glands are also found at higher density in the anterior trachea, but are known to spread entirely through the tracheal tube all the way down to the submucosal layer of the bronchi (Borthwick et al., 1999; Sturgess & Imrie, 1982). In the mouse, nasal SMGs begin to develop on embryonic day (E) 12.5 with the appearance of the first lateral nasal gland (or Steno's gland) in the form of a single tube. The remaining subgroups of nasal SMGs then start to initiate, from different locations of the medial and lateral walls (Grüneberg, 1971). The tracheal SMGs appear postnatally as initial gland buds that then undergo morphogenesis with all tracheal SMGs initiated by postnatal day P21 (Borthwick et al., 1999; Rawlins & Hogan, 2005).

Cellular Composition of the SMGs

The SMGs boast a typical tree-like structure common to all branched organs such as the mammalian lungs, kidney, mammary glands, lacrimal glands and salivary glands. Research has shown that tracheobronchial SMGs in humans consist of three distinct domains in relation to the overlying RE: (A) the secretory region of the distal gland, (B) the medial collecting region and (C) the proximal ciliated duct (Meyrick et al., 1969; Wine & Joo, 2004) (Figure 1). The secretory region is made up of two main

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secretory cell types: serous cells and mucous cells. Glandular acini, “acinus” meaning “berry” in Latin, describes the cluster of serous secretory cells found at the distal gland that surrounds a central lumen (Meyrick & Reid, 1970) (Figure 1). In the trachea, the distal serous cells make up 61% of the overall secretory cell volume, forming a liquid rich in bactericidal enzymes, making it the primary defensive cell of the airway mucosa (Basbaum et al., 1990). Serous cells produce a watery solution primarily made up of proteoglycans and bactericidal proteins such as lactoferrin and lysozyme and some mucins (MUC2, MUC7) (Finkbeiner, 1999; Klockars & Reitamo, 1975; Masson et al., 1966). Serous tubules then lead into more proximal mucous tubules with mucous cells making up 39% of the remaining secretory gland volume (Basbaum et al., 1990) (Figure 1). Mucous cells produce a thick gel containing mucins MUC2 and MUC5 and antimicrobial peptide cathelicidin (Finkbeiner, 1999). The protein consistency of mucus provides ideal lubrication and acts as a chemical barrier to the exposed epithelium. The serum and mucus secreted by these specialized cells accumulates in a non-ciliated collecting duct and is then passed through a ciliated duct into the airway lumen (Figure 1). It is generally believed that mucus is expelled into the airway through the movement of the cilia within the ducts, however no concrete evidence indicates this is the case and it is suggested that the fluid is rather moved through a means of bulk flow (Ballard & Spadafora, 2007). Once secreted into the airway, mucus is swept away from the lungs to the pharynx by respiratory epithelial cilia movement where it is either swallowed and ingested, or is removed from the body by sneezing or coughing.

The Nasal SMGs and Their Secretions

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The nose is one of the initial entry points of inhaled air into the respiratory system. Therefore, the epithelial lining cells of the upper respiratory tract are exposed to the highest concentrations of inhaled toxins and debris. It is the anterior nasal SMGs, along with the goblet cells of the RE that provide the first line of defence within the airway. The function of these respiratory glands is to secrete mucus into the nasal cavity where it traps airborne pathogens within inspired air, preventing them from progressing to the lower airway tubes and lungs. These mucous secretions also play an important role in warming and humidifying air, preventing the drying out of the delicate respiratory tissue. The nose is the principal organ for sense of smell, and it is also believed that the mucus produced by the nasal SMGs dissolves odorant

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3 molecules to help them be picked up by olfactory receptors (Getchell & Getchell,
4 1992). Unlike humans, mice are obligate nasal breathers, therefore the nose and nasal
5 environment of these animals are critical in maintaining respiratory health and
6 homeostasis. For that reason, they provide an ideal model to study respiratory nasal
7 gland development and function.
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10 11 12 13 **The Nose**

14 In order to understand the developmental locations and progressions of the anterior
15 nasal glands, the structure of the nose must be understood. In humans and rodents, the
16 nose can be divided into three main areas known as the vestibule, the respiratory
17 region and the olfactory region (Figure 2). The nasal vestibule is the opening site of
18 the nose and is lined with a stratified squamous epithelium. Moving into the nasal
19 cavity a transition into a RE occurs (Figure 2). Gross et al. (1982) showed that in mice
20 the RE accounts for 47.5% and 45% of the total surface area of the nasal cavity in 7
21 week and 16 week old mice respectively. The periphery of the RE stops suddenly and
22 the epithelium becomes olfactory in nature. In the same study the olfactory epithelium
23 (OE) accounted for 45% and 47% of the nasal surface area in 7 week and 16 week old
24 mice respectively (Gross et al., 1982). The primary function of the RE is to moisten
25 and protect the cavity while the OE houses an enrichment of olfactory receptor
26 neurons, required for the sense of smell (Popp & Martin, 1984; Purves et al., 2004).
27 The RE and OE are supplied with mucosal secretions by goblet cells and the
28 respiratory SMGs, and the Bowman's glands respectively (Bojsen-Moller, 1964;
29 Frisch, 1967).
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44 To appreciate the morphology of the nasal cavity, the capsule can be divided into the
45 medial and lateral walls (Figure 2, B and C). The medial wall of the cavity is a simple
46 plane of tissue that covers either side of the cartilaginous nasal septum (Figure 2, C).
47 Shelves of bone known as conchae or turbinates curve the lateral wall of the cavity
48 (Figure 2, B and C). The most anterior shelf is referred to as the superior concha, the
49 middle shelf as the medial concha while the lower shelf is called the inferior concha
50 (Figure 2, B and C). By protruding from the lateral wall, these conchae create
51 channels within the nasal chamber over which air can flow. These channels are
52 known as meatuses and adopt the same name as their overlying concha (Figure 2, B
53 and C). Additionally, there are four pairs of connected air filled cavities found in the
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4 nose, known as the paranasal sinuses (Figure 2, C). The lining of these spaces is
5 continuous with the nasal RE and OE and secretions are drained directly or indirectly
6 into the nose (Figure 2, C, black dotted line). The maxillary sinus is the largest of the
7 sinuses and is located lateral to the main nasal chamber (Figure 2, C). The frontal
8 sinuses are the second largest of the group and are found above the eyes, and between
9 the eyes lie the ethmoidal sinuses (Figure 2, C). A fourth pair of sphenoid sinuses is
10 found behind the eyes and nasal cavity. The function of the paranasal sinuses is
11 disputable however it is believed they decrease the weight of the skull as well as
12 warming and humidifying inspired air with their mucosal secretions. In contrast to
13 humans, mice have been reported to have only anterior and posterior ethmoidal
14 sinuses, a true maxillary sinus, a secondary maxillary sinus, as well as observable
15 sphenoid sinuses (Jacob & Chole, 2006). SMGs are only found to drain into the true
16 maxillary sinus in mice (Jacob & Chole, 2006), and unlike humans, this cavity is not
17 completely closed off by the maxillary bones (Phillips et al., 2009). Despite these
18 anatomical differences however, the rodent maxillary sinus provides a useful tool in
19 investigating sinus SMG development and function.
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33 **Morphology and Development of the Anterior Nasal SMGs**

34 In the mid-twentieth century, researchers analysed and delineated anterior nasal SMG
35 topography and physiology (Bojsen-Moller, 1964; Brunner, 1942), however modern
36 investigation of these respiratory glands is severely lacking. It has been referenced in
37 the literature that Broman (1921) first described the topography of the anterior nasal
38 glands and created plastic reconstructions of the glands and ducts of rodents (Bojsen-
39 Moller, 1964; Grüneberg, 1971). According to the manuscripts published in the mid-
40 twentieth century, it was Broman (1921) who categorised and named the nasal glands
41 into different groups based on their location and timing of when they first arose
42 (Bojsen-Moller, 1964; Grüneberg, 1971). Although mentioned and reviewed in these
43 early anatomical studies, the work of Broman has been described as “lost in recent
44 literature” (Bojsen-Moller, 1964). Despite this, his primary names and descriptions of
45 the glands have continued throughout the years and his work undoubtedly provided
46 the backbone of future research on anterior respiratory gland development. While
47 previous studies have named and described the temporal location of the glands in
48 mice, and other mammals (Brunner 1942; Bojsen-Moller 1967; Moe & Bojsen-Moller
49 1971; Bojsen-Moller 1964), these manuscripts fail to give descriptive explanations
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3 and illustrative aids to help the researcher locate and differentiate all groups of glands.
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5 The aim of this review was to thoroughly investigate and describe the patterning of
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7 the anterior nasal glands and the tracheal SMGs in a mouse model, in the hope of
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9 clarifying the development and arrangement of the glands and encourage future
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11 research into their cellular and molecular behaviours.

12 13 *Lateral Nasal Gland 1 or the Steno's Gland*

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15 The lateral nasal gland 1, more commonly called the Steno's Gland, is the largest of
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17 the nasal glands (Grüneberg, 1971). The gland is located beneath the wall of the
18
19 maxillary sinus and releases its product into an extremely long excretory duct, which
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21 opens into the nasal cavity near the nasal vestibule. The Steno's gland, analogous to
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23 the anterior nasal gland in humans, is classified in some animals as solely serous in
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25 nature however in others it is seromucous (Moe & Bojsen-Moller, 1971; Bojsen-
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27 Moller, 1967). In dogs the Steno's gland is considered to play a principal role in
28
29 thermoregulation while in marine birds it is primarily a salt gland (Butler, 2002;
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31 Wells & Widdicombe, 1986). Studies in rodents have elucidated that its secretion
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33 contains bactericidal proteins indicating an essential role in airway protection (Moe &
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35 Bojsen-Moller, 1971).

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37 Between E12.0 and E12.5 the Steno's duct buds from the anterior respiratory
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39 epithelium (REA) on the septal side at the nasal vestibule (Figure 3, A-D). The
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41 budding of the duct occurs as the pseudostratified REA dips inwards and invaginates
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43 into the underlying mesenchyme (Figure 3, A-D). As the bud arises, an indentation in
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45 the REA is formed (Figure 3, C-D). At E13.5 duct elongation proceeds over the
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47 superior nasal meatus and distally extends through the mesenchyme of the middle
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49 concha (Figure 3, E-H). The REA indentation continues as the Steno's duct lumen,
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51 simultaneously with the growing duct. This was described by Grüneberg (1971)
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53 however no histological or illustrative description was provided. A day later at E14.5,
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55 the distal tip of the duct has reached the mesenchyme below the RE of the maxillary
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57 sinus cavity (Figure 3, I-L). At this location, duct elongation ceases and the Steno's
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59 gland begins to branch distally over the next 24 hours (Figure 3, J - red brackets). At
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61 E15.5, the extending gland branches and end buds are apparent (Figure 3, M-N).
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63 Branching of the Steno's gland continues over the next few days of embryonic
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65 development. The first sign of acinar cell differentiation is observed at E16.5 by
66
67 Alcian Blue stained mucus within acini. Continual branching proceeds at E17.5, when

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3 the gland expands (Figure 3. O-P). At this stage the majority of acinar cells are
4 producing mucus as indicated by Alcian Blue staining in the gland (Figure 3, O-P).
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8 *The Lateral Nasal Glands*

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10 The lateral nasal glands (LNGs) are found throughout the lateral wall of the nasal
11 chamber. Similar to the Steno's gland they have long ducts that open into the airway
12 lumen close to the nasal vestibule, however, these glands are found within the
13 submucosa of the lateral wall at more caudal locations. Up to 40 lateral nasal glands
14 have been described in rabbits while approximately 20 have been reported in adult
15 mice (Bojsen-Moller, 1964; Broman, 1921), however, it is unclear in this report how
16 the glands were counted. From our studies, we observe five LNGs (Steno's gland and
17 LNG2-5), which bud and elongate during embryonic murine development. This
18 suggests that either additional LNGs initiate postnatally, or that the large number of
19 glands previously described in adults represent branches associated with the same
20 gland. All of these LNGs bud from the posterior respiratory epithelium (REP) that
21 covers a lip of tissue that protrudes from the rostral end of the middle concha (Figure
22 4). LNG2 is the first to undergo bud initiation between E13.5 and E14.0 from the REP
23 at the nasal vestibule closest to the nares (Figure 4 and Figure 5, B-D). Unlike the
24 Steno's gland duct, which dips into the underlying mesenchyme forming a dent in the
25 REA, the budding of LNG2, and all the other LNGs, involves the invagination of a
26 solid swelling of cells from the REP into the mesenchyme. At E14.5, LNG2 caudally
27 extends as a solid cord of cells through the mesenchyme of the middle concha (Figure
28 5, F-H). At this stage LNG3 is observed to be at the late bud stage, having extended
29 slightly from the REP at a location anterior to the distal tip of the extending LNG2
30 duct, and lies just beneath the opening of the Steno's duct into the airway lumen
31 through the REA (Figure 4 and Figure 5, F-H). Additionally at E14.5, LNG4 and
32 LNG5 are observed budding from the REP covering the most rostral side of the
33 middle concha lip, and caudal to the distal end of the LNG2 duct respectively (Figure
34 5, F-J).
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55 By E15.5, the duct of LNG2 reaches its final destination beneath the maxillary sinus,
56 above the branching Steno's gland (Figure 5, K-M). The duct of LNG3 elongates
57 parallel to that of LNG2, with its distal tip found rostral to the end of the arrested
58 LNG2 duct (Figure 5, K-M). LNG4 elongates posteriorly through the middle concha
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4 and its distal tip emerges in the mesenchyme below the enclosing cartilage of the
5 nasal capsule (Figure 5, N-O). This suggests that LNG4, although structurally similar
6 to the other LNGs, could be following a different molecular signal that attracts its
7 elongating gland duct in a posterior direction, compared to a caudal attractant guiding
8 the other LNGs. Regardless of this difference in spatial-temporal development, all the
9 LNGs appeared to follow the same pattern of cavitation and branching
10 morphogenesis. The LNG5 duct elongates caudally through the middle concha and
11 arrests rostral to the ends of LNG2 and LNG3 (Figure 5, K).
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19 By E16.5, LNG2 has branched distally above the Steno's gland, showing the
20 formation of end buds (Figure 6, A-B). The beginning of lumen formation within the
21 acini of LNG2 is apparent at this stage (Figure 6, B – white arrows). No mucus
22 production is evident in acinar cells of LNG2 at this stage. The distal tip of LNG3
23 duct has reached the caudal end of the middle concha, where it was found lying
24 rostral to the branching LNG2 (Figure 6, A-B). LNG3 has begun branching
25 morphogenesis as the first signs of end buds are observed (Figure 6, B – green
26 arrows). LNG4 is at a similar stage of development with the formation of end buds
27 noted at its distal end below the nasal cartilage capsule (Figure 6, E-F). The duct of
28 LNG5 has elongated through the middle concha beneath the RE, parallel to the ducts
29 of LNG2 and LNG3, where it begins branching over the next 24 hours (Figure 6, I-J).
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40 By E17.5, LNG2 has undergone extensive branching and cells have arranged into
41 distinct acini (Figure 6, C-D). The distal gland of LNG3 has also undergone similar
42 branching and clear lumens are identified within acini (Figure 6, C-D). No mucus
43 staining by Alcian Blue is evident at E17.5 in both the LNG2 and LNG3. LNG4 has
44 formed numerous branches and terminal end buds and cavitation has begun within
45 acini (Figure 6, G-H). LNG5 has begun to form distal epithelial gland buds and lumen
46 formation has progressed from the duct to acini (Figure 6, K-L)
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53 *The Medial Nasal Glands*

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55 The medial nasal glands (MNGs) are found within the submucosa of the medial wall
56 on either side of the septum. Their long ducts open at rostral locations while their
57 glands progress caudally and are seen anterior to the vomeronasal organ in rodents.
58 These glands have been previously described in the rat, which can have four to five
59 MNGs that project into the septal mesenchyme (Bojsen-Moller, 1964). The MNGs
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3 have been reported as having serous acini by light microscopy (Kerjaschki, 1974).
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5 The rat MNGs do not stain with Periodic-Acid Schiff (PAS) or Alcian Blue staining
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7 (Bojsen-Moller, 1964). Analysis of the four MNGs of the mouse have also been
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9 carried out (Broman, 1921; Kerjaschki, 1974). These glands have previously been
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11 reported to be exclusively made up of serous cells (Bojsen-Moller, 1964; Kerjaschki,
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13 1974) and in our studies, no Alcian Blue staining is seen within their acini during
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15 embryonic stages, suggesting these glands are exclusively serous in nature. While all
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17 the MNGs look similar and undergo comparable developmental strategies,
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19 investigation into their ultrastructure has delineated distinguishable differences
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21 between these glands (Kerjaschki, 1974). It has been shown that MNG4, shows
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23 homologous characteristics to serous salivary glands, with its gland cells having
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25 common characteristics as serous cells, however MNG1-3 stand alone as their own
26
27 distinct gland type (Kerjaschki, 1974). What differentiates MNG1-3 from the fourth is
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29 that they possess “light” and “dark” endpiece cells within their glandular acini when
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31 analysed by electron microscopy (Kerjaschki, 1974). These “light” and “dark” cells
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33 have been described in the Bowman’s glands of the olfactory epithelium, where
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35 “light” cells contain electron-dense serous secreting vesicles while the “dark” cells
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37 contain electron-lucent mucous secreting vesicles (Getchell & Getchell, 1992). These
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39 compositional differences of these glands may also explain why in this study we see
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41 MNG1-3 elongate and branch in the same plane and at similar time points, when
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43 MNG4 arises in the more rostral septum and develops slightly later. While all are
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45 considered ‘Medial Nasal Glands’, for future studies it is advised to be aware of their
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47 differences.

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49 Development of the MNGs is relatively simple to follow as their ducts elongate and
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51 branch through the clear plane of mesenchymal tissue on either side of the nasal
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53 septum. All MNGs bud from the RE covering either side of the nasal septum of the
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55 medial wall and elongate and branch within the mesenchyme of the septum (Figure 7,
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57 A-L). MNG1 and MNG2 are the first to bud between E14.0 and E14.5. At E14.5
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59 ducts of these two medial glands have elongated (Figure 7, A-D). By E15.5 these
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61 ducts have elongated into the central area of the septum while MNG3 has budded and
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63 elongated to the midline of MNG1 and MNG2 (Figure 7, E-H). Through another
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65 plane of mesenchyme it is clear that branching of MNG1 begins by E15.5 (Figure 7,
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67 inset image H). Branching progresses by E16.5 in MNG1 and end bud formation is

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3 also underway in MNG2 (Figure 7, I-J). MNG3 elongates adjacent to the ducts of
4 MNG1 and MNG2 at E16.5 (Figure 7, I-J) and begins branching over the next 24
5 hours. At E17.5, MNG1, 2 and 3 are all branching, with branches and terminal end
6 buds all crossing over one another (Figure 7, K-L). MNG4 has also elongated and
7 begun its branching at E17.5, indicated by the presence of end buds (Figure 7, K-L).
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12 *The Maxillary Sinus Glands*

14 The maxillary sinus glands (MSGs) bud and branch from the RE at both the anterior
15 and posterior walls of the maxillary sinus. They do not have long gland ducts like the
16 other anterior nasal glands but rather branch within the mesenchyme close to the
17 epithelium. Research on the MSG in rat found it to be an exclusively serous gland
18 (Vidić & Greditzer, 1971) and begins its development at E16 (Vidić, 1971). In the
19 mouse, we show that the MSGs undergo a similar method of development as in the
20 rat, however, is first seen to bud from the epithelium of the maxillary sinus at
21 approximately E15.0. The anterior and posterior sinus glands undergo the same
22 developmental patterning so for expediency the development of only the posterior
23 MSG will be explained. At E14.5, no evidence of a MSG is observed (Figure 8, A-B)
24 however by E15.5 branching of the MSG is apparent close to the RE (Figure 8, C-D).
25 At this stage the MSG is observed to invaginate into the mesenchyme with preformed
26 lumens within the branching epithelium. Branching of the MSG epithelium continues
27 over the next two days of embryonic development with lumens constantly present
28 within the end buds (Figure 8, E-H).
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42 **Signalling During Nasal SMG Development**

43 The Wnt signalling pathway is an evolutionary conserved pathway, critical for
44 controlling the development of an organism. Many developmental processes such as
45 cell proliferation, migration, polarity and fate determination rely on the regulation of
46 Wnt signalling (Komiya & Habas, 2008). Two categories of Wnt transduction
47 pathways are known, the canonical Wnt/ β -catenin cascade, and the non-canonical
48 Wnt/ β -catenin-independent cascades, which include the Wnt/Planar Cell Polarity
49 pathway and the Wnt/Calcium pathway. The former of these, the canonical Wnt
50 signalling pathway, is the most studied and involves the binding of extracellular Wnt
51 ligands to transmembrane bound Frizzled receptors and a co-receptor, low-density
52 lipoprotein receptor related protein 5 (LRP5) or LRP6. In the absence of Wnt ligand
53 binding, cytoplasmic β -catenin is constantly broken down by the Axin complex,
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4 consisting of Axin, glycogen synthase kinase 3 (GSK3), casein kinase 1 (CK1) and
5 the *adenomatous polyposis coli* gene product (APC). Together, GSK3 and CK1
6 phosphorylate β -catenin, rendering it recognisable to the E3 ubiquitin ligase subunit
7 β -Trcp (Davidson et al., 2005; Liu et al., 2002; Zeng et al., 2005). Subsequent
8 ubiquitin of β -catenin occurs, followed by its proteosomal degradation (He et al.,
9 2004). Upon Wnt ligand-receptor binding, the Axin complex is recruited to the cell
10 membrane bound receptor complex, preventing the β -catenin inhibitory effects of
11 these proteins, and thus β -catenin degradation (He et al., 2004). This permits β -
12 catenin translocation into the nucleus where it can activate Wnt-responsive genes. The
13 most studied nuclear mediators targeted by Wnt signalling belong to the lymphoid
14 enhancer factor 1/T-cell factor (LEF-1/TCF) family (Arce et al., 2006; Eastman &
15 Grosschedl, 1999).

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27 Wnt signalling negatively regulates branching morphogenesis in other branching
28 organs (Dean et al., 2005; Patel et al., 2011). *Ex vivo* culture of lung, lacrimal and the
29 submandibular salivary gland (SMDG) with WNT3A, leads to an inhibition of
30 clefting and a reduction in the number of epithelial branches (Dean et al., 2005; Patel
31 et al., 2011). Using the *Axin2^{lacZ}* reporter, it has been shown that Wnt/ β -catenin
32 signalling is restricted to the ductal structures of the developing SMDG, with its
33 localisation seen at E14.5, during the late *Pseudoglandular stage*, in the main gland
34 duct and by E15.5 spreading into the smaller ducts (Patel et al., 2011). Wnt activity
35 may be maintained in the ductal epithelial cells to ensure their long-range elongation
36 before the branching of a distal gland occurs. Wnt activity is maintained in the
37 intercalated ducts of the SMDG during postnatal development, however, reduces as
38 mice age (Hai et al., 2010). Enhanced Wnt activity is observed in ductal cells
39 following SMDG duct ligation, indicating a role of this pathway in SMDG
40 regeneration (Hai et al., 2010). Using the *Lef1* *-/-* mouse to study the biological role of
41 this gene, it was shown that knockout mice failed to develop organs such as teeth, hair
42 follicles and mammary glands (van Genderen et al., 1994).

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57 Using the β -catenin/Tcf- β gal reporter (TOPGAL) mouse, it has been shown that the
58 canonical Wnt pathway is active during early nasal gland bud formation (Driskell et
59 al., 2007). Additionally, the transcription factor *Lef1*, has been shown to be highly
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3 expressed in progenitor cells of early SMG buds in the mouse nose and trachea, and
4 the ferret trachea, and in *Lefl* knockout mice, amorphous aggregates of cells develop
5 where nasal SMGs are seen in wild types (Driskell et al., 2004; Duan et al., 1999,
6 1998). *Lefl* promoter expression has been shown to be regulated by Wnt-3a during
7 SMG morphogenesis, and in the *Wnt3-a* knockout mouse, SMG *Lefl* expression was
8 lost and cell proliferation of epithelial gland cells was reduced in SMG buds (Driskell
9 et al., 2007). While these studies have given insight into the necessity of Wnt
10 signalling for early SMG bud formation, the involvement of Wnt molecules in later
11 stages of respiratory gland development has not been investigated.
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21 The role of the Ectodysplasin A (EDA) pathway has been investigated in the
22 development of a number of ectodermal derived organs (Mikkola, 2009). Mutations in
23 the *Eda* pathway genes in both humans and mice leads to hypohidrotic ectodermal
24 dysplasia (HED) which gives rise to congenital defects in salivary glands, hair, teeth
25 and sweat glands (Clarke et al., 1987). Signalling through this pathway involves the
26 binding of EDA A1, a member of the tumor necrosis factor (TNF) family, to a death
27 domain transmembrane receptor, EDAR (Headon & Overbeek, 1999; Srivastava et
28 al., 1997). EDA ligand binding to EDAR leads to the recruitment of the intracellular
29 adaptor protein EDAR Associated Death Domain (EDARADD). Recruitment of
30 EDARADD to the receptor leads to the formation of a complex containing the
31 cytoplasmic proteins TNF Receptor Associated Factor 6 (TRAF6), TGF- β Activated
32 Kinase 1 (TAK1) and TAK Binding Protein 2 (TAB2) (Morlon et al., 2005). This
33 leads to the activation of the I κ B kinase complex (IKK), which is a compound of two
34 kinase sub-units, IKK1 and IKK2, as well as the regulatory component NF-kappaB
35 essential modulator (NEMO) (Döffinger et al., 2001; Morlon et al., 2005). Activation
36 of the IKK complex triggers recruitment and the phosphorylation of inhibitor kappa-B
37 (I κ B), a nuclear factor-kappaB (NF κ B) inhibitor. Phosphorylation of I κ B leads to its
38 degradation, and NF κ B translocates into the nucleus where it acts as a transcription
39 factor, regulating target genes involved in cell proliferation, differentiation and
40 survival (Gilmore, 2006; Oeckinghaus & Ghosh, 2009).
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58 A number of reports using the mouse model have shown the importance of *Eda*
59 signalling in both major and minor salivary gland morphogenesis. Analysis of the
60 *Tabby* mouse, a model of a naturally occurring mutation in the gene required for EDA

(Srivastava et al., 1997), gave rise to hypoplastic SMDGs with reduced numbers of convoluted tubules and a reduction in mucin immunolocalization (Blecher et al., 1983; Jaskoll et al., 2003). In contrast to this, the knockout mouse of the *Edar* receptor gene, known as *downless*, developed dysplastic SMDGs, showing a severe lack of ducts and acini, and virtually no detection of mucus (Jaskoll et al., 2003). Furthermore, addition of exogenous EDA protein addition to developing SMDGs in culture, as well as elevated expression of EDAR signalling in adult mice, gave rise to an increase in SMDG epithelial branches (Chang et al., 2009; Jaskoll et al., 2003). Addition of EDA protein to cultured *Eda* hemizygous (*Tabby*) SMDGs *in vitro* can also rescue branching in these mutant glands (Wells et al., 2010). Investigation of the minor salivary glands additionally emphasised the role of *Eda* signalling in gland branching morphogenesis as these glands of the tongue do not develop in *Tabby* and *downless* mice (Wells et al., 2011).

Analysis of the *Tabby* mouse in a classical study revealed that specific groups of the anterior nasal SMGs were completely absent while others developed normally (Grüneberg, 1971). It was reported that the Steno's gland and MSG develop in the *Tabby*, while some, but not all of the LNGs were completely absent (Grüneberg, 1971). Furthermore, all of the MNGs were reported missing (Grüneberg, 1971). Of the LNGs that did develop in the *Tabby*, these glands were described as rudimentary from the beginning, and their distal branched glands were also defective (Grüneberg, 1971). Defects in nasal glands have also been reported in HED patients, with patients suffering from dryness of the nasal mucosa, severe nasal crusting and foul smelling nasal discharge (Al-Jassim & Swift, 1996; Dietz et al., 2013).

Morphology and Development of the Tracheal SMGs

Developmental of the tracheal SMGs observed from animals models can be described in four distinct stages (Keswani et al., 2011; Thurlbeck et al., 1961). Stage 1 of development is the initial bud stage where an epithelium swelling is seen to immerge from the RE (**Error! Reference source not found.**, A). Stage 2 is characterised by the epithelial bud elongating into the underlying mesenchyme and cavitation occurs within this stalk to form a lumen (**Error! Reference source not found.**, B). This is followed by Stage 3 where canalized stalks begin to branch within the underlying mesenchyme (**Error! Reference source not found.**, C), while Stage 4 is classified as

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4 the cellular differentiation stage, indicated by mucus production within the gland
5 (**Error! Reference source not found.**, D). Unlike the nasal glands that develop for
6 approximately two days as a long gland duct, followed by the branching of a distal
7 gland the tracheal SMGs bud, elongate with a short stem, cavitate and begin
8 branching. This developmental mechanism is similar to that seen in the branching of
9 both the major and minor salivary glands, however in these structures branching
10 occurs before lumen formation (Melnick & Jaskoll, 2000; Teshima et al., 2011).

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16 In CD1 mice, the first signs of tracheal bud initiation can be identified adjacent to the
17 cricoid cartilage in the ventral trachea at late E18.5. At P0 some ventral glands were
18 observed at Stage 1 of budding, while others had extended and branched and had
19 already begun Stage 3 of lumen formation and branching (Figure 10, C). At birth,
20 some gland buds were also found in more dorsolateral planes, with a wave of
21 initiation starting in the ventral regions and spreading more dorsally (Figure 10, D).
22 Initiation before birth is earlier than previously described for the tracheal SMGs
23 (Rawlins % Hogan, 2005) and may represent differences in genetic background
24 (Borthwick et al., 1999; Innes & Dorin, 2001). Different inbred strains of mice show
25 different posterior extensions of the SMGs between cartilage rings, while CFTR
26 knockout mice have an increase in posterior extension with mutants showing glands
27 as far as the eighth cartilage ring compared to the fourth ring in WT littermates
28 (Borthwick et al., 1999; Innes & Dorin, 2001)

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40 By P1, branching continues in both the ventral and dorsal glands, as well as continued
41 induction of new buds throughout the dorsolateral walls. By P2, more newly forming
42 SMGs arose in the anterior tracheal mesenchyme in dorsolateral positions (Figure 10,
43 F), while ventral glands progressed to the branching and mucus producing (Figure 10,
44 E). By P4, the ventral and dorsolateral glands continued to undergo extensive
45 branching and cellular differentiation (Figure 10, G and H). The first signs of SMG
46 formation in a posterior direction between the first and second cartilage rings was
47 observed in the ventral trachea at approximately P4 (Figure 10, G). By P8, established
48 anterior glands were continuing their branching as well as new glands appearing
49 between C2 and C3 and progressively at P12 glands were found between C3 and C4
50 in ventral cross-sections. At P15 glands had developed between C4 and C5 and were
51 producing mucus. During postnatal development, glands arose in a gradient from the
52 ventral to dorsal areas of the trachea, in conjunction with their posterior development.
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3 By P22, glands were well established and observed as posterior as C6 and initiation of
4 new glands appeared to cease (Borthwick et al., 1999; Rawlins & Hogan, 2005).
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9 **Signalling During Tracheal SMG Development**

10 To date, no investigation into the role of the canonical Wnt signalling pathway has
11 been carried out on the tracheal SMGs. Similar to the nasal glands however, tracheal
12 SMGs have been reported to be completely absent in the adult *Tabby* mouse, as well
13 as in the early *crinkled* mice, the model of *Edaradd* mutation, at P7, indicating that
14 EDA signalling is required for early SMG initiation and budding (Rawlins & Hogan,
15 2005). Complimentary to these findings, human patients with HED have been
16 reported to have reduced numbers of seromucous respiratory glands, asthma like
17 symptoms and respiratory tract infections (Callea et al. 2013). Furthermore, allergic
18 response is increased in those with HED (Vanselow et al. 1970).
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28 The bone morphogenetic proteins (BMPs) belong to the transforming growth factor β
29 (TGF β) superfamily (Miyazono et al., 2010). The BMPs play a multitude of roles
30 during early embryonic patterning and morphogenesis such as the regulation of cell
31 proliferation, differentiation, apoptosis and cell-fate determinations (Hogan, 1996;
32 Kishigami & Mishina, 2005). Twenty BMPs have been identified, all having
33 heterogeneous functions during developmental processes such as neurogenesis, bone
34 and cartilage formation, and organogenesis (Hogan, 1996; Zhao, 2003).
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43 In general, the BMP pathway involves the extracellular BMP ligands binding to the
44 cell membrane bound BMP receptors (BMPR). These receptors are serine-threonine
45 kinase receptors that belong to two groups, Type I and Type II (Massagué et al.,
46 1994). BMPs bind to the Type I BMPR with higher affinity to that of Type II (Kirsch
47 et al., 2000; Wrana et al., 1994). In order for ligand binding to stimulate an
48 intracellular signal, heterodimerization of one Type I and one Type II BMPR is
49 required. Upon BMP ligand binding to the Type I receptor, the Type II receptor is
50 recruited (Carreira et al., 2014; Kirsch et al., 2000). The Type II BMPR then catalyses
51 the phosphorylation of the intracellular serine-threonine domain of the Type 1 BMPR,
52 recruiting a cytoplasmic receptor-activated Smad protein (either Smad1, 5 or 8).
53 (Wrana et al., 1994). Phosphorylation of one of these Smads triggers the recruitment
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3 of another cytoplasmic Smad, Smad4 (Whitman, 1998). This dimeric Smad complex
4 can now translocate into the nucleus where BMP target genes can be activated. A
5 number of extracellular antagonists also function in regulating the pathway. These
6 include such soluble proteins as Noggin, Gremlin and Chordin, which form
7 complexes with BMP ligands, preventing them from binding to their cell receptors
8 (Walsh et al., 2010; Yanagita, 2005).
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12 It has been previously shown that BMP signalling is critical for the development and
13 patterning of the trachea and lungs (Eblaghie et al., 2006; Li et al., 2008; Weaver et
14 al., 1999). During SMDG development, *Bmp4* is mesenchymally expressed and
15 addition of BMP4 protein to developing SMDGs *in vitro* SMDG inhibits gland
16 budding (Hoffman et al., 2002), indicating an antagonizing role in the control of the
17 number of gland branches. Furthermore, when SMDGs are cultured with BMP
18 antagonist sFRP1, branching of SMDGs is induced with a 26% increase observed in
19 the amount of gland buds in treated glands compared to controls (Patel et al., 2011).
20 In contrast BMP7 seems to enhance gland branching. In the *Bmp7* knockout mouse, a
21 reduction in epithelial branches of the SMDG as well as disorganized mesenchyme
22 surrounding the gland was observed (Jaskoll et al., 2002).
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26 The only investigation into the roles of BMP signalling in respiratory gland
27 development have been shown during that of the tracheal SMGs (Rawlins & Hogan,
28 2005). Using the *Bmp4^{lacZ}* reporter mouse, it was reported that at P2 when tracheal
29 buds were initiating from the RE, *Bmp4* was expressed throughout the underlying
30 mesenchyme. During later development at P7, *Bmp4* detection was reduced and found
31 only in the mesenchyme close to the newly formed gland branches (Rawlins &
32 Hogan, 2005). By P28, *Bmp4* expression had reduced even further, and was only
33 detected in a few mesenchymal cells (Rawlins & Hogan, 2005). The exact role of
34 BMP4, or other BMPs, during respiratory gland development has not been elucidated.
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38 Fibroblast growth factors (FGFs) make up one of the largest family of polypeptide
39 proteins critical for tissue patterning and development of branching organs. Today, 22
40 members (FGF1-FGF22) of the FGF family are known within the human genome
41 (Ornitz & Itoh, 2001). Four cell membrane-bound FGF receptors (FGFRs) have been
42 identified in vertebrates, each having different affinities for different FGF ligands.
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The four receptors, FGFR1-FGFR4, are composed of the common protein structure of most tyrosine kinase (TK) receptors (Johnson et al., 1990). The initiation of an intracellular signalling transduction pathway relies on extracellular binding of FGF ligands its target receptor, leading to receptor dimerization and intracellular phosphorylation of the TK domains of each receptor (Jiang & Hunter, 1999; Schlessinger, 1988). Following TK phosphorylation, a number of intracellular signalling cascades are triggered that are required for a multitude of developmental processes. These include the phospholipase C gamma (PLC- γ) pathway which initiates calcium release and activates protein kinase C to influence cell motility (Burgess et al., 1990; Mohammadi et al., 1991), and the phosphatidylinositol-3 kinase (PI3K) which activates Akt/protein kinase B needed for the mediation of cell survival (Ong et al., 2001). The most common route of FGF signalling during developmental processes is through the activation of the rat sarcoma homologue (RAS)/mitogen-activated protein kinase (MAPK) cascade. This pathway functions in activating target genes involved in cellular differentiation and proliferation.

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The roles of a number FGFs have been shown to be critical for gland morphogenesis. *Fgf10* is expressed in the mesenchyme underlying the presumptive salivary gland epithelium from E11.5 (Wells et al., 2013). Jaskoll et al. (2005) showed in *Fgf10* null mice, as well as the *Fgfr2b* knockouts, that the SMDG initiated to the *Initial Bud* stage at E12.5 however it was unable to develop further, and at E13.5 any epithelial evidence of the SMDG was absent. Culture of SMDGs with FGF7 gave rise to moderate stalk extension and induced epithelial bud enlargement (Koyama et al., 2008). Inhibition of salivary gland branching by FGFR2b antisense oligonucleotides was also rescued by the addition of exogenous FGF7 which stimulated cell proliferation and end bud formation (Steinberg et al., 2005). Using *Fgf8* hypomorphic embryos, as well as *Fgf8* conditional knockout mutants, it has also been described that SMDG bud initiation is independent of *Fgf8*, however expression is required for continued SMDG branching morphogenesis and survival (Jaskoll et al., 2004a). At approximately E13.5 when the lacrimal gland bud arises on the temporal side of the eye, *Fgf10* expression is seen in the periocular mesenchyme adjacent to the budding gland and distal to the tip of the gland as it is elongating at E14.5 (Makarenkova et al., 2000). Investigation of *Fgf10* $-/-$ mice at E18.5 also showed the complete absence of the lacrimal gland (Makarenkova et al., 2000). While *Fgf7* expression was not as

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evident as that of *Fgf10* in the periocular mesenchyme during normal *in vivo* lacrimal gland development, application of an FGF7 bead to lacrimal gland explant cultures induced ectopic gland bud formation, similarly to FGF10 protein application, however not at as high a rate (Makarenkova et al., 2000). The respiratory gland phenotype has not been investigated in any *Fgf* homozygous knockout mice, however, Rawlins and Hogan (2005) established that in *Fgf10* heterozygous mice at P20 fewer tracheal SMGs developed. Of the glands that did develop, they were only located anteriorly by the CC and C1 and had not undergone full branching while no glands were seen between the more posterior cartilaginous rings (Rawlins & Hogan, 2005). The role of FGF signalling during nasal and tracheal SMG development has not been investigated. It has also been shown that tracheal cartilage ring defects occur in *Fgf10* homozygous mice, which may also have a secondary affect on tracheal SMG development (Sala et al., 2011). It is therefore possible that in addition to a direct defect of FGFs on gland elongation and branching, a tracheal ring defect might compromise the arrangement of the tracheal SMG development.

33 34 35 36 37 38 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 55 56 57 58 59 60 **Future Directions**

In this review we describe a detailed account of anterior nasal and tracheal SMG development. We highlight that the SMGs show distinct morphological stages of development, beginning with bud formation, followed by duct elongation, distal gland branching and acinar differentiation (Table 1). Now that the morphology of these glands has been elucidated, further research into the mechanisms involved in their budding and arborisation is encouraged. As we report here, studies have looked at early bud induction of the nasal glands, and the roles played by the canonical Wnt signalling pathway (Driskell et al., 2007; Duan et al., 1999) (Figure 11). The involvement of Wnt activity during later duct elongation and distal gland branching stages of the nasal SMGs is however unknown. In addition, the role of Wnt signalling during tracheal SMG development has not been analysed (Figure 11). Requirement for Ectodysplasin A signalling has been emphasised by the absence of some nasal and all tracheal SMGs in the *Tabby* mutant (Grüneberg, 1971; Rawlins & Hogan, 2005) (Figure 11). In keeping with this mouse data, human patients of HED also develop airway obstruction and pulmonary infection (Dietz et al., 2013). The nasal gland phenotype observed in the *Tabby* mouse indicates that different subsets of nasal SMGs utilise different cues, with the Steno's gland and the MSG developing normally

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in the absence of Eda signalling, while the medial nasal glands are completely absent (Grüneberg 1971). Thus glands with similar morphology may develop using different signalling pathways. This is also emphasised by the difference in lumen formation used to create a final hollow duct. The Steno's gland develops with a preformed lumen, similar to the lung, while the other LNGs and MNGs develop as solid cords of cells that are cavitated at later stages to create a lumen, similar to salivary glands (Figure 11). Thus there are different mechanisms at play to form anatomically similar structures. Further down the respiratory tree, participation of *Fgf10* during trachea SMG development has been shown to be essential for successful gland branching morphogenesis (Rawlins & Hogan, 2005) (Figure 11). Analysis of the role of *Fgf10* during nasal SMG development, or its requirement for early tracheal SMG bud initiation has not been explained however. Furthermore, while previous studies have hinted at roles played by the signalling factors mentioned above, other developmentally important pathways during SMG organogenesis have not been touched upon. For example, Sonic hedgehog (*Shh*) expression is localised to ducts and terminal end buds of the developing SMDG (Jaskoll et al. 2004b). In the *Shh* $-/-$ mouse at E18.5, an undifferentiated dysplastic gland forms, suggesting that during SMDG development *Shh* is required for gland growth and cellular differentiation (Jaskoll et al. 2004b). Interestingly, mammary gland branching morphogenesis is independent of SHH signalling, as glands in the *Shh* $-/-$ mouse are indistinguishable of those observed in wild type littermates (Michno et al., 2003). This data collectively poses interesting questions for the role and regulatory functions of *Shh* in SMG development. Additionally, epidermal growth factors (EGFs) have been described to be essential for successful development of both SMDGs and mammary glands (Kashimata et al., 2000; Luetke et al., 1999), however this pathway has not been studied in any of the respiratory SMGs. To this end, continual investigation is imperative to delineate the cellular mechanisms and intracellular cues that are required for successful SMG morphogenesis. This will provide insight into how development of SMGs influences function, and thus help clarify their role in pulmonary diseases.

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Figure Legends

Figure 1. Schematic illustration outlining the cellular composition of the treacheobronchial SMGs.

The tracheal glands are composed of sacs of serous cells, which form acini. Serous cells secrete a watery like secretion rich in bactericidal enzymes. Acini run into tubules of mucous cells, which secrete a thicker gel like substance rich in glycosylated proteins. These two types of secretions are then combined in a collecting duct, whose wall is composed of non-ciliated columnar epithelial cells. The collecting duct runs into a ciliated duct made of more respiratory epithelial-like cells that possess beating cilia on their apical surfaces. Mucus is secreted through this ciliated duct into the airway lumen where it covers the surface epithelium.

Figure 2. Anatomy of the nose and paranasal sinuses in humans.

(A) Sagittal view of the three distinct areas of the nose; vestibule, respiratory and olfactory regions. (B) Sagittal view of the lateral wall showing the curved concha and the inter-conchal meatuses. (C) Frontal section through the nasal chamber showing the position of the medial walls (yellow) on either side of the septum (S). Image shows the location of the superior concha (SC), the medial concha (MC) and the inferior concha (IC) of the lateral wall (orange) and their underlying meatuses. The frontal image also delineates the location the paranasal sinuses. SM=superior meatus; MM=middle meatus; IM=inferior meatus; FS=frontal sinus; ES=ethmoidal sinus; MS=maxillary sinus. Dotted black lines indicate opening of sinuses into nasal cavity found at a more caudal plane.

Figure 3. Steno's duct elongation and branching morphogenesis.

(A-D) The epithelial budding of the Steno's duct is seen between E12.0 and E12.5 from the REA. (E-H) By E13.5, the Steno's duct elongates over the nasal cavity and caudally extends through the mesenchyme of the medial concha. (I-L) By E14.5 the distal tip of the Steno's duct has reached the mesenchyme beneath the maxillary sinus to the location of where the Steno's gland will branch (J, red bracket). (M-N) The Steno's gland begins to branch from the Steno's duct between E15.0 and E15.5. The first signs of end buds are seen at E15.5. (O-P) By E17.5 the Steno's gland is continually branching and acinus differentiation is well established with most acinar

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cells producing mucus (blue arrows). Maxillary sinus cavity (MS); Steno's gland (yellow arrows and dotted line); Division of RE to OE shown by pink arrows. C,G,K scale bar= 300 μ m, D,H,L scale bar = 500 μ m, M & O scale bar = 500 μ m; N & P scale bar = 250 μ m.

Figure 4. Budding locations of the LNGs from the protruding lip of the middle concha.

The LNG2 bud is first seen at E13.5 arising from the REP at the most rostral area of the conchal lip. It will extend caudally through the middle concha as development continues. At E14.5, LNG3 arises from the anterior area of the lip, underneath the opening of the Steno's duct through the REA, and will extend caudally adjacent to the duct of LNG2. At E15.5, LNG4 initiates below the origin of the LNG2 and extends posteriorly through the mesenchyme of the concha where it will eventually branch beneath the nasal cartilage capsule, anterior to the palatal shelf. LNG5 also initiates at E15.5, where it will elongate adjacent to the ducts of LNG2 and LNG3. Gland buds (2-5) shown in green, direction of future extending duct shown by red arrows. Schematic is taken at a sagittal view. Axis describes anterior (A), posterior (P), rostral (R) and caudal (C).

Figure 5. LNG duct elongation.

(A) Sagittal (S) and horizontal (H) planes through an E13.5 mouse head showing the location of the LNG2 bud. (B) Horizontal representation of LNG2 budding from the middle conchal lip. (C) Sagittal histological section showing the location of the middle conchal lip and the LNG2 bud in the nasal region. (D) The epithelial bud of LNG2 at E13.5. (E-F) Schematics showing location and appearance of the elongated LNG2 and the budding LNG3-5 at E14.5. (G-H) At E14.5 the LNG2 has elongated caudally as a solid stem. LNG3 has budded and begun to elongate and LNG4 is seen at a slightly earlier budding stage. (I-J) LNG5 is at its initial bud formation stage at E14.5. (K) Schematic showing the progression of the LNGs by E15.5. (L-M) A sagittal section shows the distal tips of LNG2 and LNG3 ending within the mesenchyme above the branching Steno's gland (yellow). The distal tip of LNG5 is found at a more rostral location adjacent to these ducts (shown in K). (N-O) LNG4 has elongated posteriorly with its distal tip emerging below the nasal cartilage

capsule. Left column of sagittal sections scale bar = 500 μ m; right column scale bar = 100 μ m.

Figure 6. Branching morphogenesis of the LNGs.

At E16.5 the LNGs begin branching of their distal glands. **(A-B)** LNG2 is the first to undergo branching between E15.5 and E16.5. At E16.5 many end buds are apparent and cavitation has begun within the gland acini (**B** - white arrows). LNG3 has also begun to branch at E16.5 as seen by the appearance of end buds (**B** - green arrows). **(C-D)** At E17.5, LNG2 and LNG3 have branched and formed distinctive acini with lumens. **(E-F)** At E16.5 the onset of branching of the LNG4 beneath the nasal capsule is evident (**F** - green arrows). **(G-H)** By E17.5, LNG4 has formed branches and end buds, and lumen formation has begun within acini (**H** - white arrows). **(I-J)** LNG5 is seen close to the ducts of LNG2 and LNG3, with possible lumen formation beginning. A clear lumen is seen in the ducts of LNG2 and LNG3 at this stage (**J** - white arrows). **(K-L)** By E17.5, LNG5 has begun to branch with a clear lumen seen within its duct and lumen formation at its initial stages within acini. Left column of each age scale bar = 500 μ m; right column scale bar = 250 μ m.

Figure 7. MNG development.

(A-D) The development of the MNGs begins between E14.0-E14.5 with the budding and elongation of the ducts of MNG1 and MNG2. **(E-H)** The MNG1 and MNG2 ducts elongate to the central septal area by E15.5 and branching has initiated at the distal tip of MNG1 (inset image **H**). MNG3 has also budded and elongated, with its distal end adjacent to the midline of the MNG1 and MNG2 ducts (**E-F**). **(I-J)** Branching of MNG1 has progressed by E16.5 with an array of branches and terminal end buds apparent at this stage. MNG2 has also begun branching. The distal tip of MNG3 has extended close the branching ends of the other MNGs. **(K-L)** By E17.5, branching continues in all the MNGs in the central region of the septum. MNG4 has emerged by E17.5 as an elongated duct and the onset of branching has begun. Image A scale bar = 500 μ m; image B scale bar = 250 μ m.

Figure 8. MSG development.

The maxillary sinus gland initiates between E14.5 and E15.5. **(A-B)** No MSG primordium is evident at E14.5. **(C-D)** At E15.5 the MSG is seen to have invaginated

and branched into the mesenchyme close to its opening in the RE. The MSG continues to branch and grow at E16.5 (E-F) and E17.5 (G-H). MSG = red, Steno's gland = yellow, LNG2 = green. MS = maxillary sinus cavity. Image A scale bar = 500 μ m; Image B scale bar = 250 μ m.

Figure 9. Stages of tracheal SMG development

(A) Stage 1: Tracheal SMG buds are first seen to invaginate from the RE into the underlying mesenchyme. (B) Stage 2: Elongation of the bud and cavitation of the epithelial stalk occurs. (C) Stage 3: Epithelial stalk undergoes clefting and branching (D) Stage 4: Cellular differentiation is observed indicated by mucus staining within the glands by Alcian Blue. AL= airway lumen, L= lumen, M= mucus. Scale bar = 25 μ m.

Figure 10. Normal postnatal tracheal SMG development.

(A-B) Schematic representations of the ventral (A) and more dorsal (B) tissue sections cut through postnatal tracheal tissue. (C) Glands are seen to have reached the lumen formation stage in the ventral trachea by P0, while more dorsally they are seen at the branching stage (D). (E) At P2 ventral glands have continued to branch and cellular differentiation is indicated by mucus staining within gland lumens. Continued budding of new glands is seen in more anterior dorsal positions (F). (G-H) By P4 glands have branched extensively in the ventral mesenchyme as well as in the dorsal trachea. Glands are also seen at the branching stage between C1 and C2 in the ventral trachea (G). CC=cricoid cartilage, C1=first cartilage ring, C2=second cartilage ring etc. Black arrow = developing gland, green arrow = lumen formation, blue arrow = mucus production. Scale bar=100 μ m.

Figure 11. Possible paracrine factors required for early respiratory gland development.

(A) Steno's gland bud induction does not require EDA signalling, however the roles of other developmental molecules such as FGFs, BMPs, and WNTs have not been investigated during Steno's gland development. (B) The medial nasal glands have been reported to be absent in the *Tabby* mouse, indicating EDA signalling is imperative for their development. *Lef-1* expression has also been described in early nasal gland buds, indicating the involvement of WNT signalling. Requirement for

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3 FGF or BMP signalling has not been investigated in the medial nasal glands. (C)
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5 Tracheal SMG investigation has shown the expression of *Edar* in the tracheal RE, and
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7 the absence of tracheal SMGs in the *Tabby* mouse, emphasising the role of EDA
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9 signalling during SMG bud induction. Branching morphogenesis of tracheal SMGs is
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11 also reduced in *Fgf10* +/- adult mouse, indicating its need for successful
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13 development. While *Bmp4* is expressed surrounding development tracheal SMGs, the
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15 exact role of this molecule or the BMP pathway has not been investigated. Finally,
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17 WNT signalling has not been analysed during tracheal SMG development.
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Tables

Table 1. Approximate stages of the distinct morphological events of anterior nasal gland development.

Gland Name	Approx. age of budding	Approx. age of initial lumen formation	Approx. age of onset of branching
Steno's Gland	E12.0	-	E15.5
LNG2	E13.5	E14.5-E15.0	E16.5
LNG3	E14.0-14.5	E15.0	E16.5
LNG4	E14.5	E15.5	E16.5
LNG5	E14.5	E15.5	E17.0
MNG1	E14.0	E14.5	E15.5
MNG2	E14.5	E15.5	E16.5
MNG3	E15.5	E16.5	E17.0
MNG4	E16.0	E17.5	E17.5
MSG	E14.5-E15.0	-	E15.0

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Figures

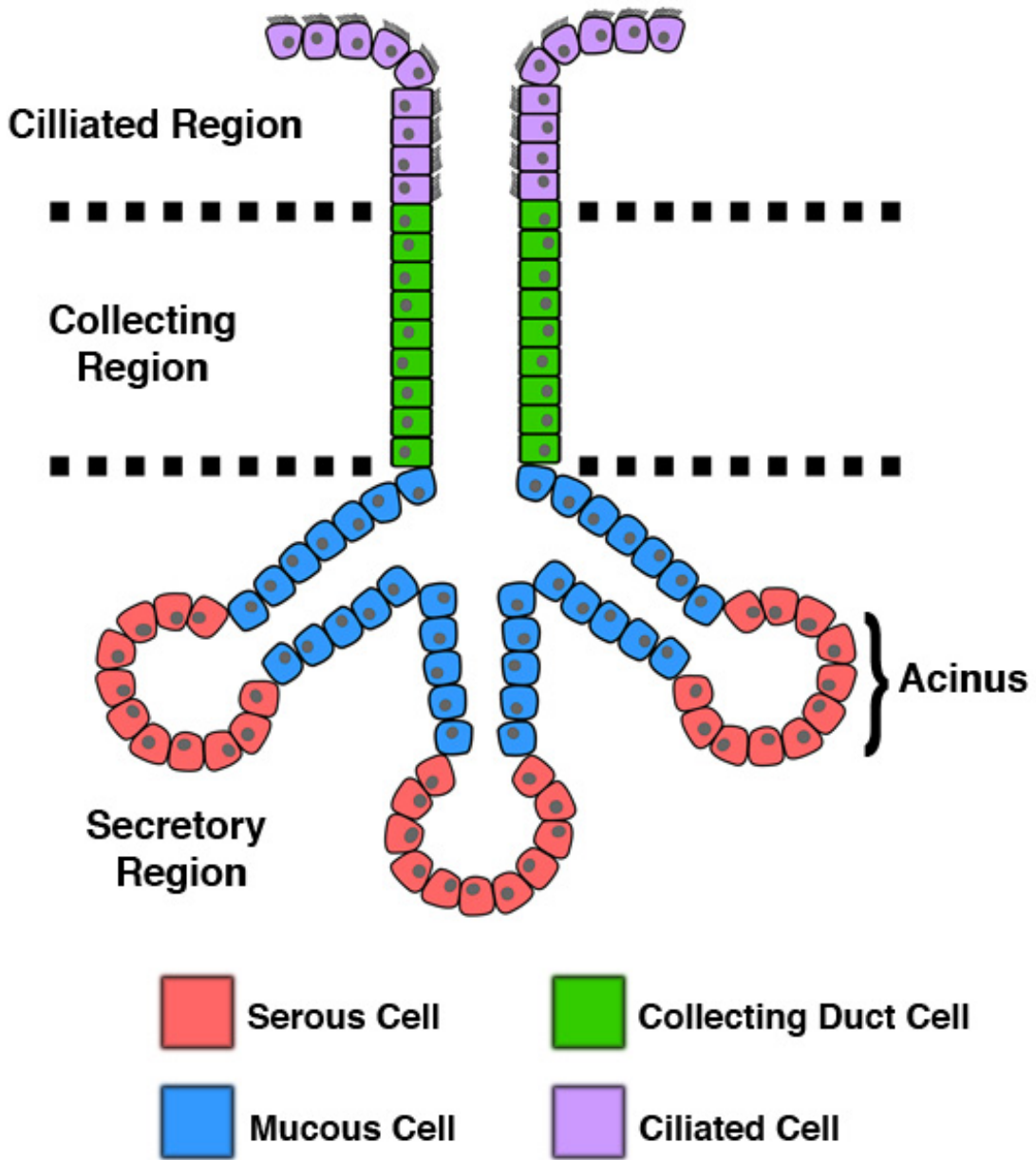


Figure 1.

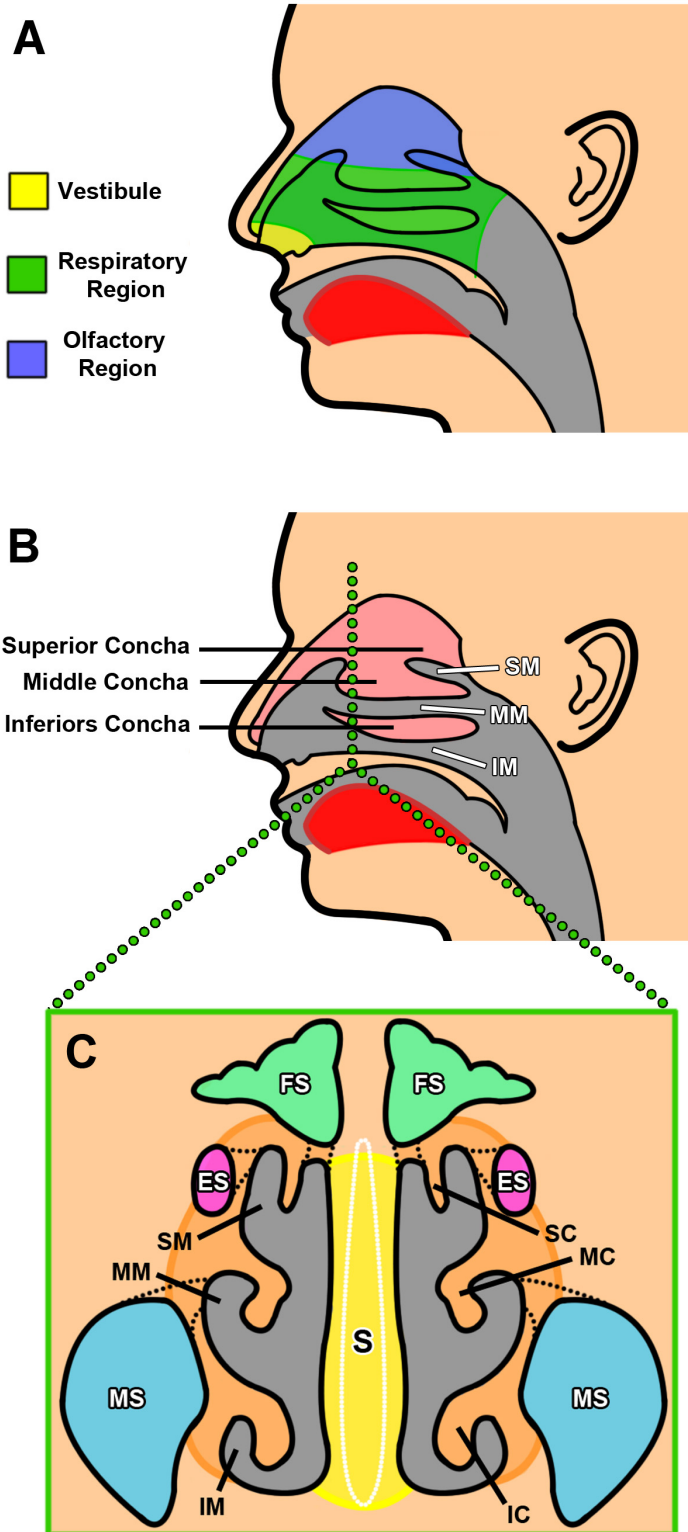


Figure 2.

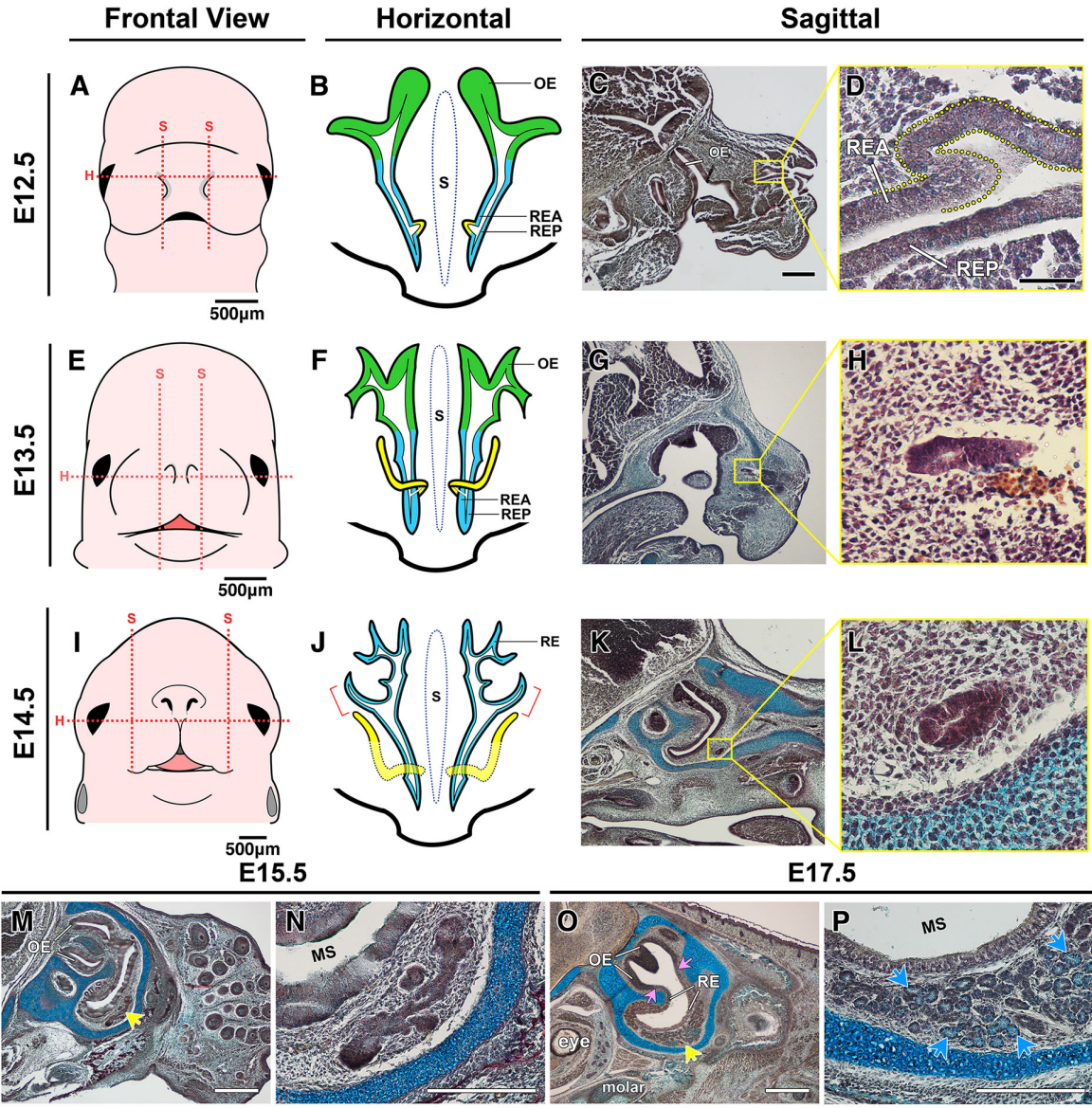


Figure 3.

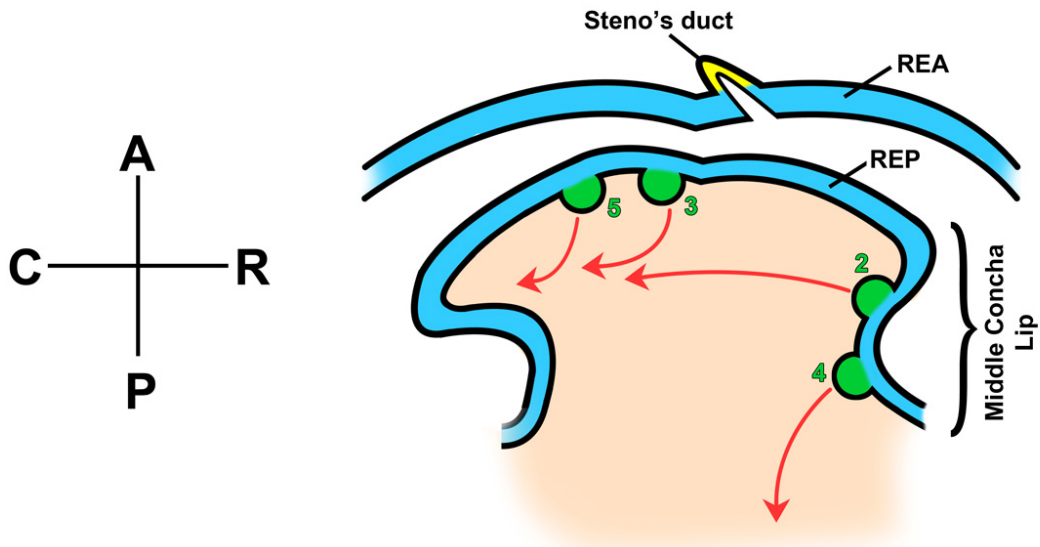


Figure 4.

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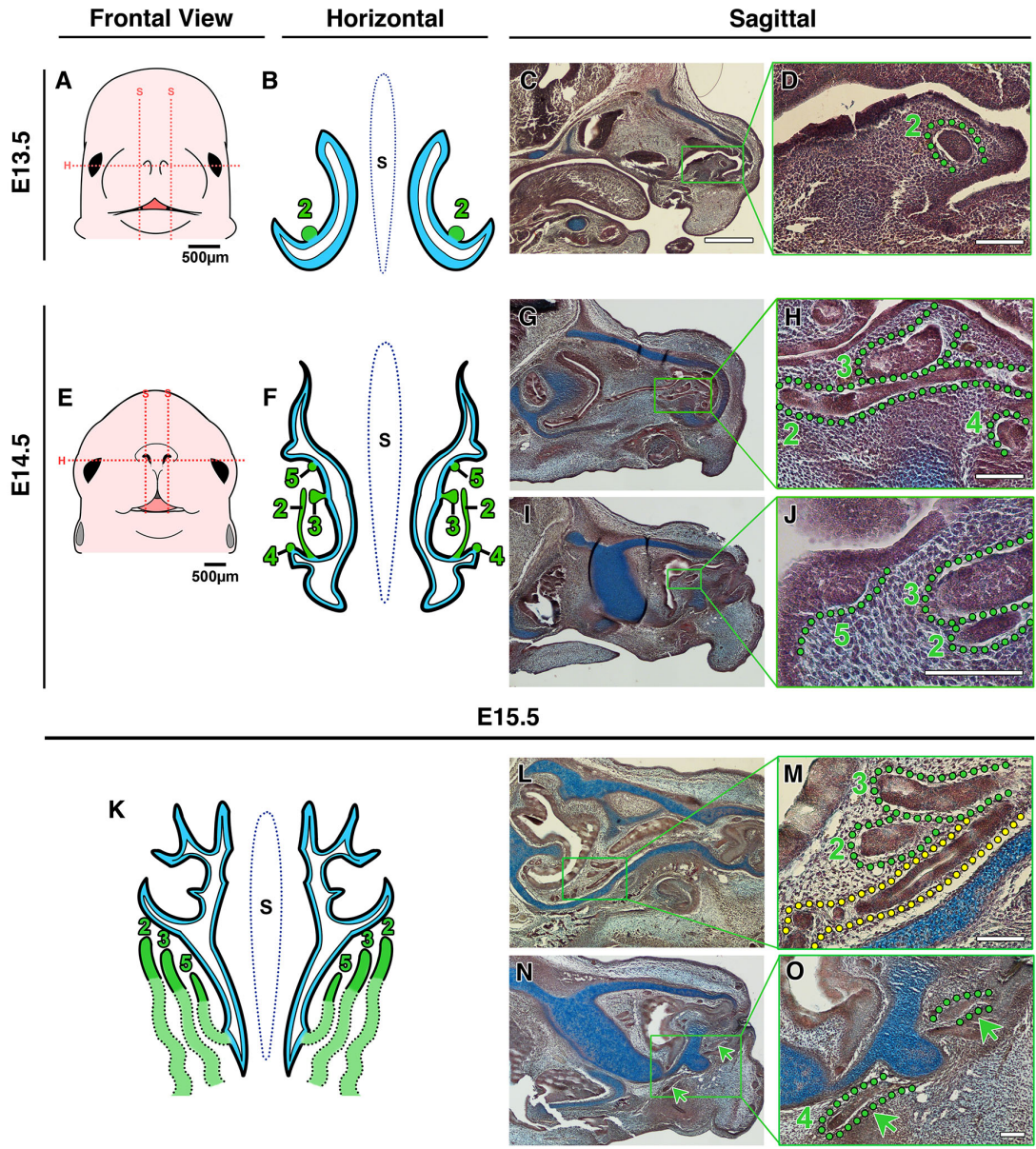


Figure 5.

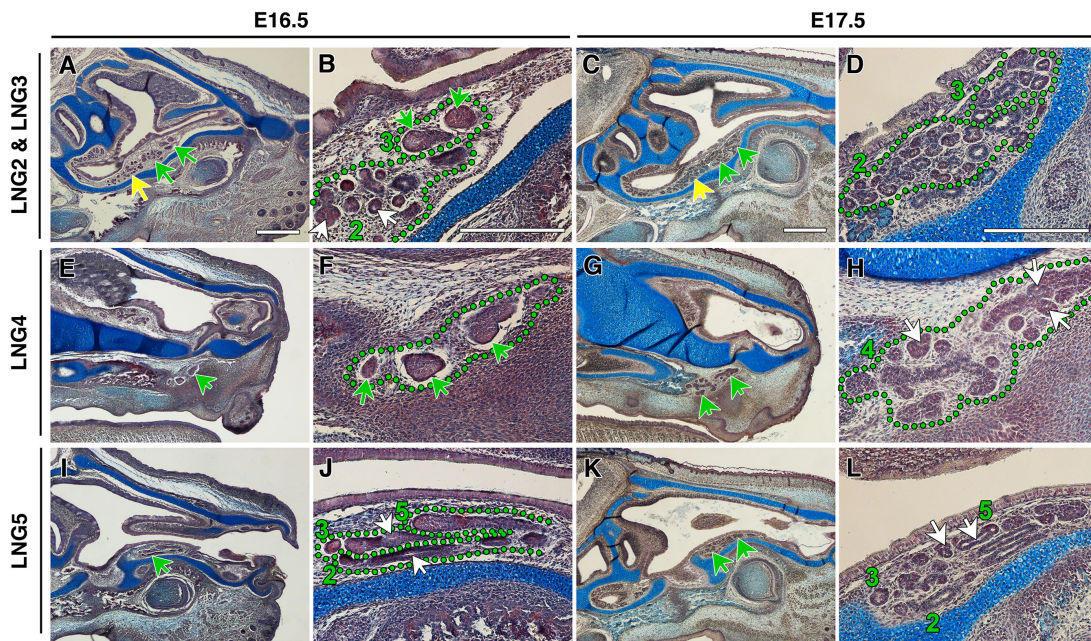


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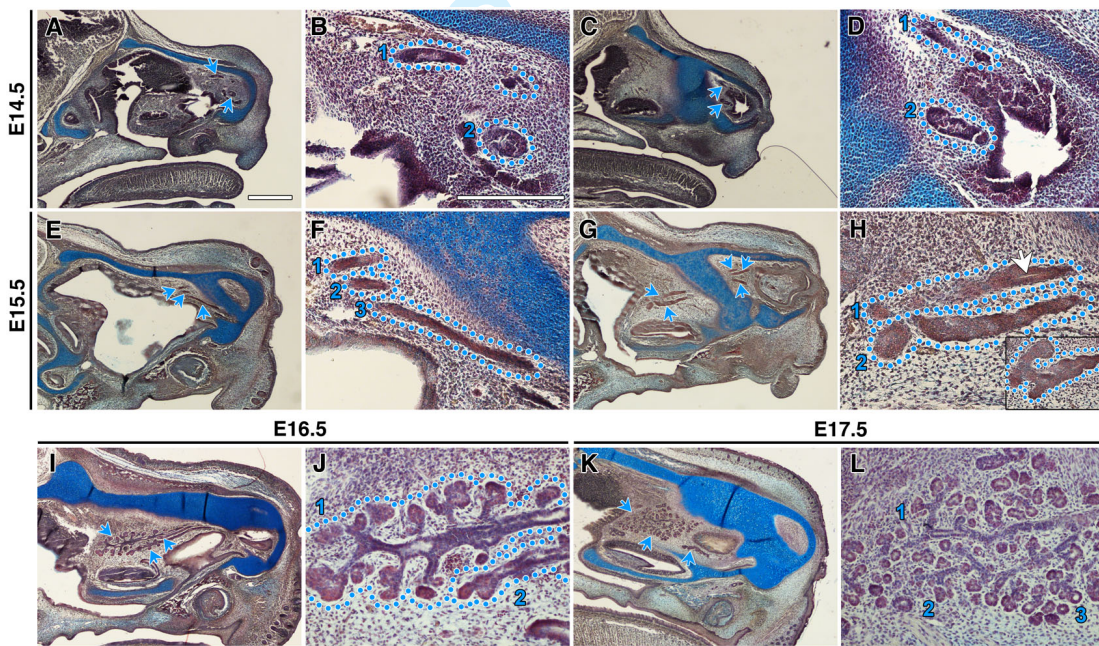


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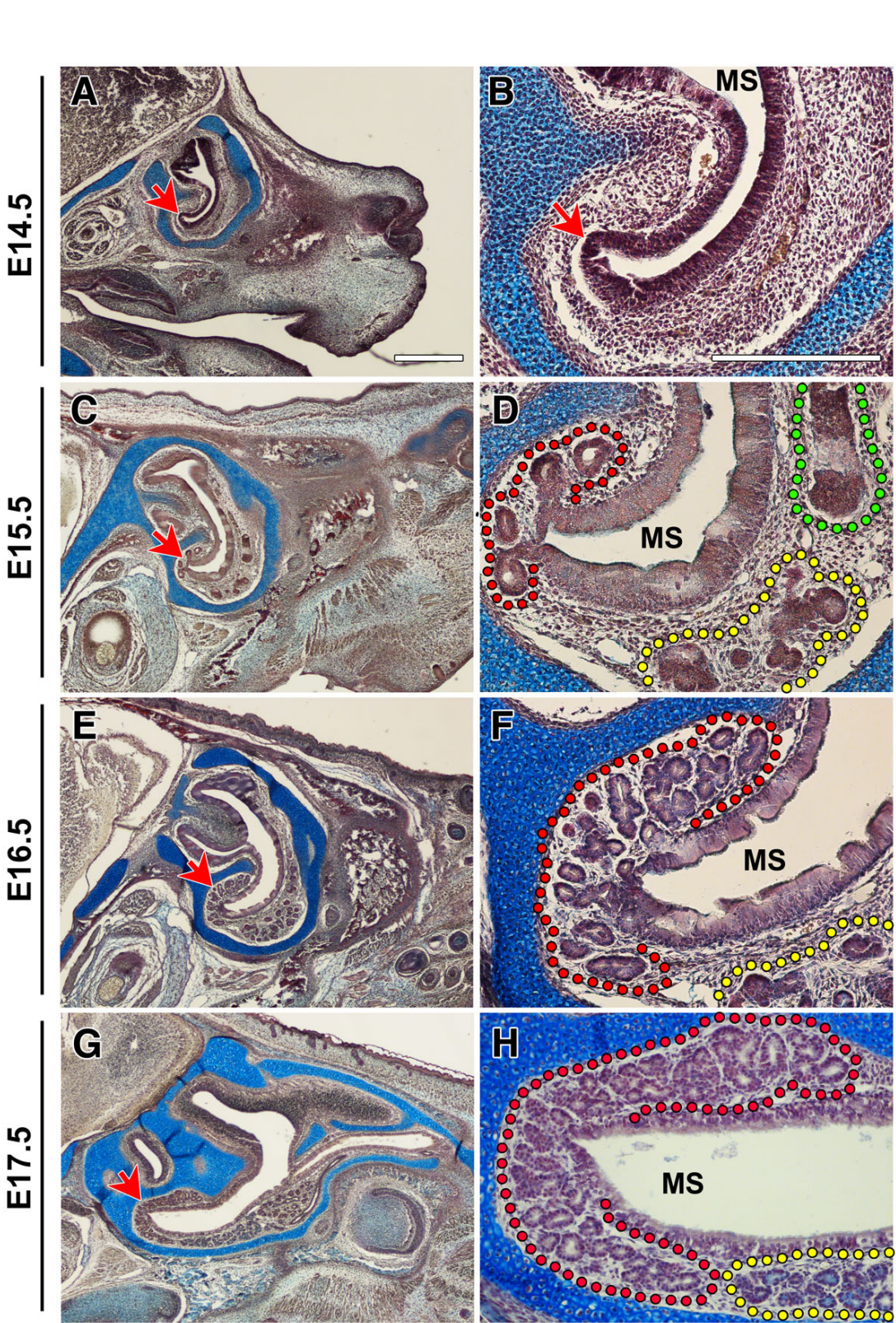


Figure 8.

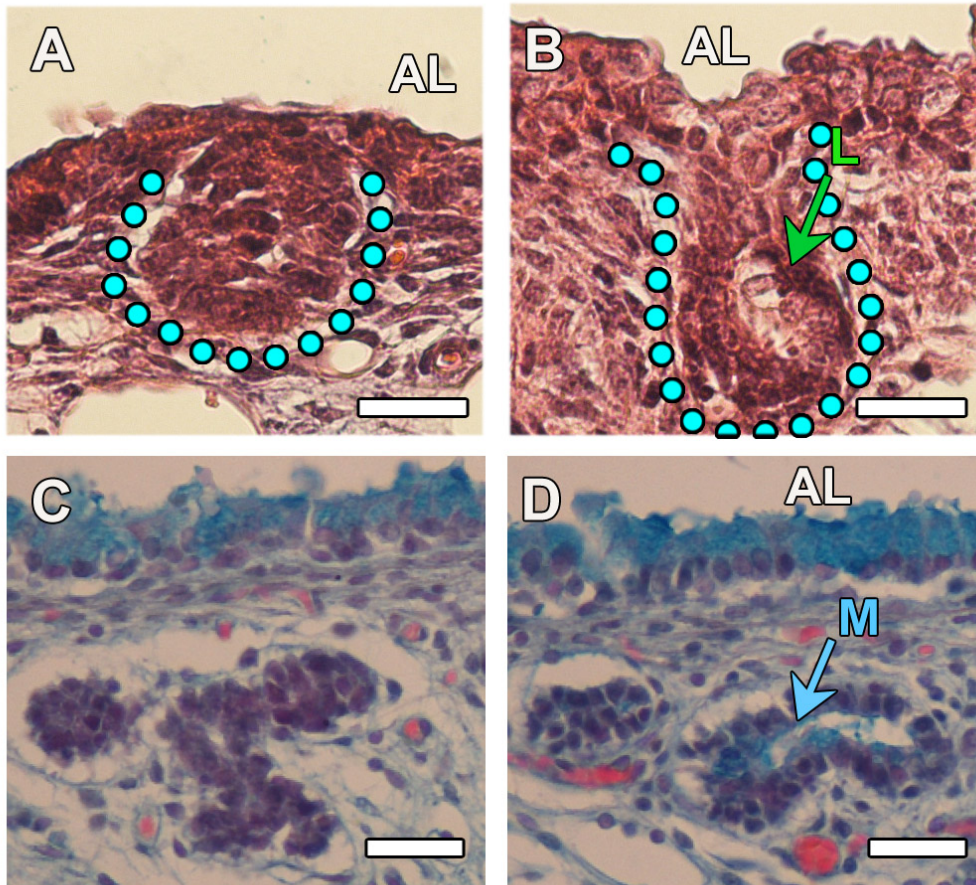


Figure 9.

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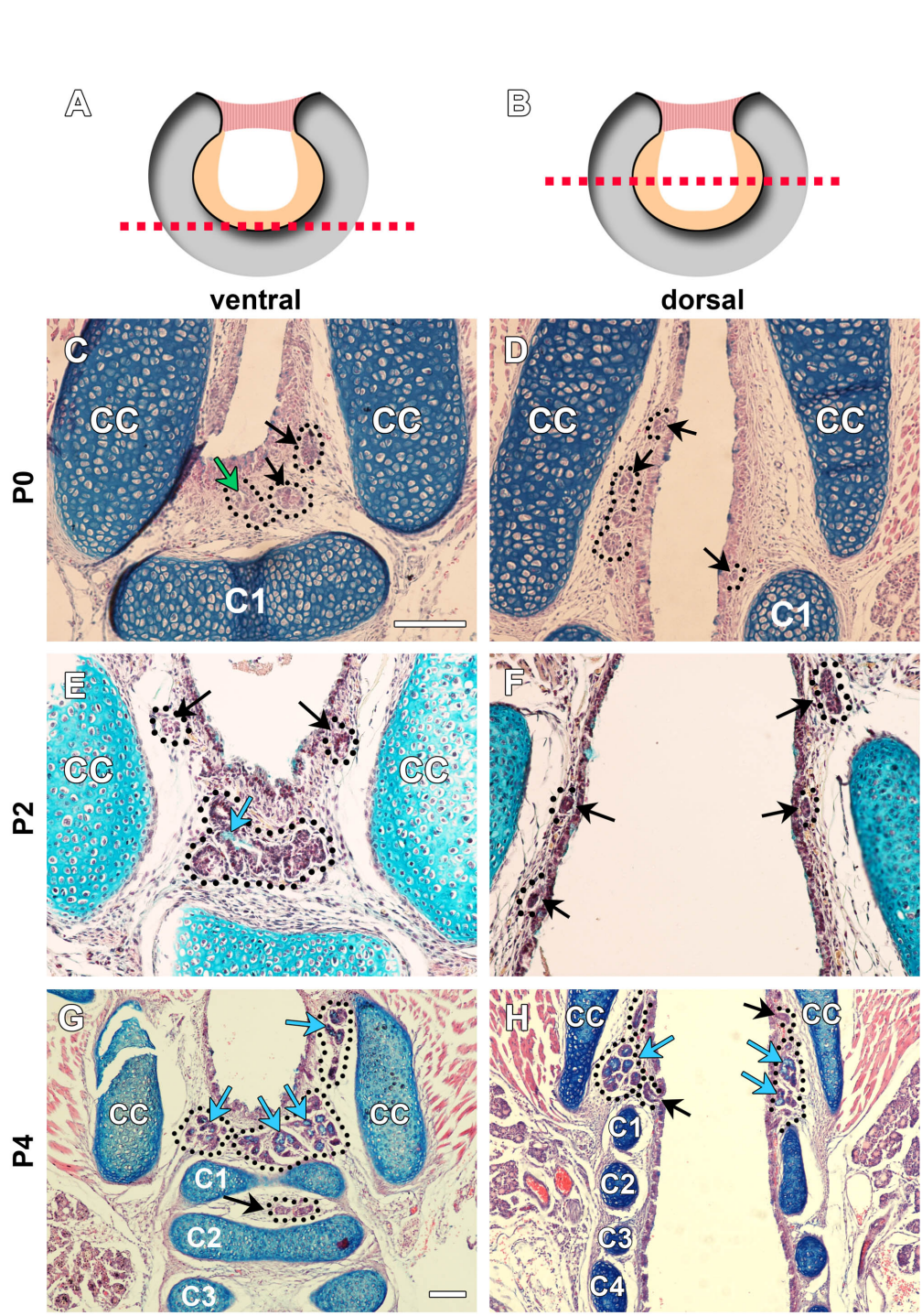


Figure 10.

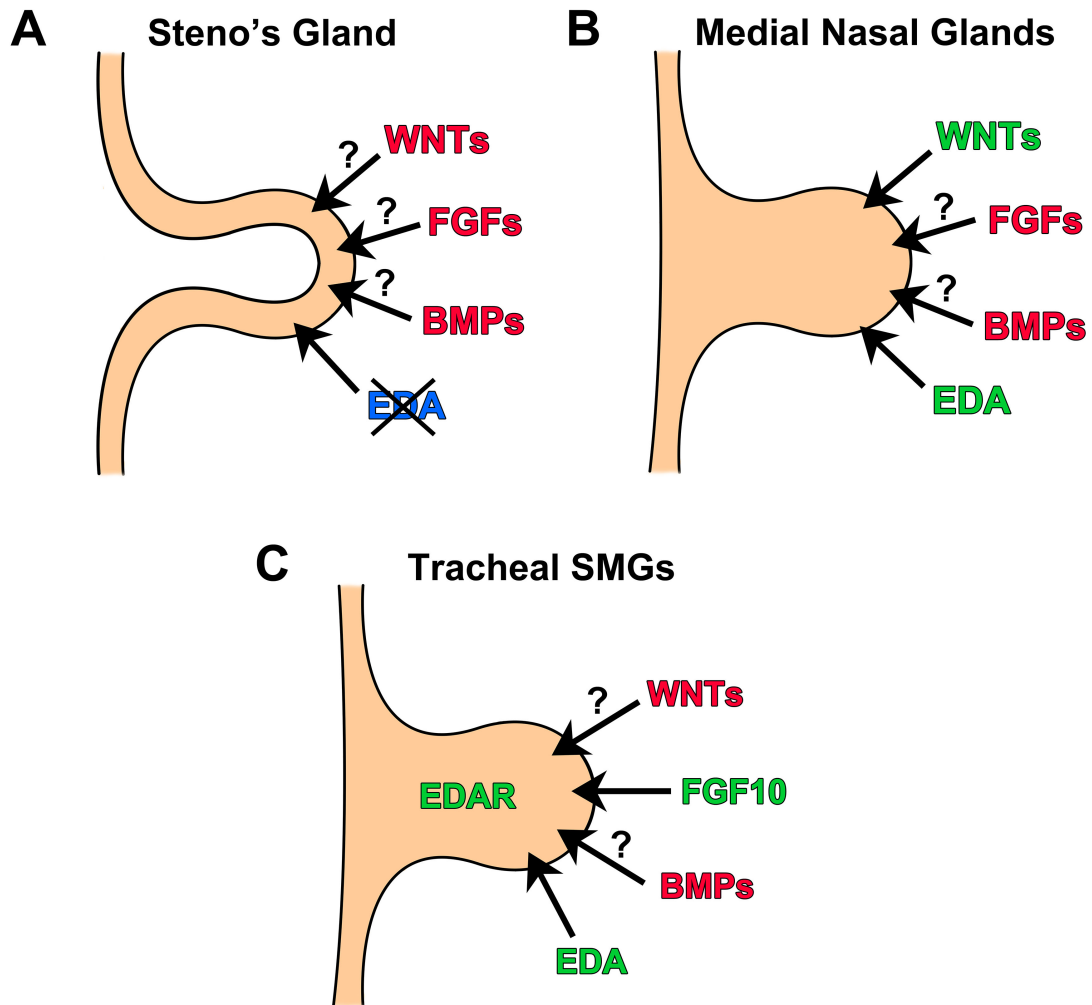


Figure 11.

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