Understanding the aetiology and resolution of chronic otitis media from animal and human studies

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ABSTRACT

Inflammation of the middle ear, known clinically as chronic otitis media, presents in different forms, such as chronic otitis media with effusion (COME; glue ear) and chronic suppurative otitis media (CSOM). These are highly prevalent diseases, especially in childhood, and lead to significant morbidity worldwide. However, much remains unclear about this disease, including its aetiology, initiation and perpetuation, and the relative roles of mucosal and leukocyte biology, pathogens, and Eustachian tube function. Chronic otitis media is commonly modelled in mice but most existing models only partially mimic human disease and many are syndromic. Nevertheless, these models have provided insights into potential disease mechanisms, and have implicated altered immune signalling, mucociliary function and Eustachian tube function as potential predisposing mechanisms. Clinical studies of chronic otitis media have yet to implicate a particular molecular pathway or mechanism, and current human genetic studies are underpowered. We also do not fully understand how existing interventions, such as tympanic membrane repair, work, nor how chronic otitis media spontaneously resolves. This Clinical Puzzle article describes our current knowledge of chronic otitis media and the existing research models for this condition. It also identifies unanswered questions about its pathogenesis and treatment, with the goal of advancing our understanding of this disease to aid the development of novel therapeutic interventions.

KEY WORDS: Chronic otitis media, Genetics, Animal models, Inflammation

Introduction

Otitis media (OM; see Box 1 for a glossary of terms) describes an inflammatory disease of the middle ear that consists of a set of inter-related clinical phenotypes. Chronic otitis media with effusion (COME, or ‘glue ear’; Box 1) affects 5-6% of children in high-income countries in their second year of life (Bhutta, 2014), becomes less prevalent in older children (Suarez Nieto et al., 1983) and is rare in adults. COME is characterised by mucosal hyperplasia, including the proliferation of mucus-secreting goblet cells in the epithelial lining of the antero-inferior middle ear cleft. These changes lead to serous or mucoid middle ear effusion (Fig. 1, Box 2), which impairs the transmission of airborne sound. COME is the most common cause of hearing loss in childhood (Robb and Williamson, 2016).

Chronic suppurative otitis media (CSOM; Box 1) is characterised by a persistent perforation of the tympanic membrane (Box 1, Fig. 2) with intermittent or constant discharge of pus through this perforation (a condition known as otorrhoea; Box 1 and see the accompanying case study in Box 2). CSOM is rare (<1%) in high-income countries but relatively common (>2%) in many low- and middle-income countries, and is highly prevalent (>4%) in some indigenous groups, such as in Australian Aboriginal, Pacific Islander, Native American and Inuit populations (Bhutta, 2015). CSOM is estimated to affect 65-million–330-million people worldwide (WHO, 2004). Epidemiological studies indicate that the highest incidence of CSOM occurs in childhood (Monasta et al., 2012), but others have suggested that prevalence continues to rise into adulthood (Shaheen et al., 2012; Chung et al., 2016).

The risk of developing either COME or CSOM involves a complex interplay between host immunity and microbial pathogenicity, which, in turn, is affected by host and microbe genetics, as well as by environmental factors (particularly those that affect risk of exposure to bacteria) and by therapeutic interventions (Fig. 3). Most cases of COME are preceded by a bacterial or viral infection of the middle ear, which causes acute otitis media (AOM; Boxes 1, 2). After an episode of AOM, the middle ear effusion becomes non-purulent [otitis media with effusion (OME); Box 1] and then usually resolves within days; however, in an estimated 8% of affected children it persists for more than 3 months and becomes chronic (COME) (Bhutta, 2014). What distinguishes the minority of children who develop chronic inflammation from those that do not is a key research question. Recurrent AOM (rAOM; Box 1) is also a risk factor for developing COME (Alho et al., 1995), but most children with COME do not suffer from rAOM.

Relatively little is known about the aetiology of CSOM, but it is thought to occur as a consequence of recurrent or persistent middle ear inflammation, which leads to a non-healing perforation of the tympanic membrane. Many affected individuals present without a history of preceding symptoms, but there is evidence that early or recurrent AOM (Fliss et al., 1991; Lasisi et al., 2007; van der Veen et al., 2006), or COME (Youngs, 1998), increases an individual’s risk of developing CSOM. Chronic otorrhoea can also develop in children after the insertion of grommets (Box 1).

In this Clinical Puzzle, we discuss our current knowledge of chronic OM and the existing research models that have been generated to investigate the factors that contribute to this disease. We also identify unanswered questions about the pathogenesis and treatment of these chronic ear conditions, and highlight the need for us to advance our understanding of the aetiology and biology of this
Box 1. Glossary of clinical terms

Acute otitis media (AOM): an ear infection, usually accompanied by symptoms of fever and pain in the ear.

Audigram: a test to measure hearing, with hearing thresholds measured in decibels.

Bulla: the middle ear cavity in animals.

Chronic otitis media with effusion (COME): otitis media with effusion present for at least 3 months.

Chronic suppurrative otitis media (CSOM): chronic otitis media with a perforated tympanic membrane and intermittent or continuous otorrhoea. Duration of otorrhoea to define disease is debated: some suggest 2 weeks, others 6 weeks, others 3 months.

Grommet: a small, hollow (also called a ventilation) tube inserted into the tympanic membrane to treat symptomatic COME and resolve effusion.

Myringoplasty: an operation to repair the tympanic membrane.

Myringotomy: an operation to make an incision in the tympanic membrane.

Otitis media (OM): inflammation of the middle ear.

Otitis media with effusion (OME): serous or mucoid effusion in the middle ear, without signs or symptoms of infection. Usually the effusion is tenacious; hence, OME is colloquially called ‘glue ear’.

Otorrhoea: discharge from the ear.

Otoscope: an instrument to examine the ear.

Recurrent AOM (rAOM): usually defined as more than three episodes of AOM in 6 months, or more than four episodes in a year.

Tympanic membrane: the eardrum, a membrane between the middle and outer ear that vibrates in response to sound.

Tympanometry: a clinical test to evaluate the pressure of the middle ear or the presence of fluid by observing the mobility of the eardrum in response to induced variations in the air pressure in the ear canal.

Box 2. Case study

A 5-year-old boy is taken to the doctor by his parents because both they and his schoolteachers have noticed that for several months his hearing seems poor. He is also behind his peer group in his spoken and written language, and there are concerns about poor behaviour. There is no preceding history of ear infections and the boy is otherwise well. The boy’s father also had hearing problems in childhood. Examination with an otoscope reveals evidence of effusion behind the tympanic membrane (see Fig. 1), giving a diagnosis of otitis media with effusion (OME; or ‘glue ear’). This is confirmed by tympanometry. An audigram reveals a 40-decibel hearing loss. After a discussion of the options, the parents decide to proceed with bilateral grommet insertion under general anaesthesia, which normalises hearing and leads to improvements in language development and behaviour.

After a year, the grommets fall out, but the left tympanic membrane has a residual perforation. Subsequently, the child suffers recurrent left-sided mucopurulent otorrhoea, which occurs every few months. The child is repeatedly treated with topical antibiotic drops but is unable to continue with his swimming lessons. At the age of 8 years, he undergoes a left myringoplasty, which successfully repairs the tympanic membrane. The child has no further problems. (See Box 1 for a glossary of clinical terms.)

disease in order to develop novel therapeutic interventions for this prevalent and chronic condition.

**Chronic otitis media can resolve**

What is unusual about both COME and CSOM is that these conditions can resolve spontaneously. Many other chronic inflammatory disorders, such as rheumatoid arthritis or multiple sclerosis, demonstrate a clinical course of persistent or recurrent inflammation, and rarely result in permanent resolution (Nathan and Ding, 2010).

For patients with COME, effusion can be resolved through the insertion of grommets (ventilation tubes), although the mechanism underlying this effect is unknown. Around one fifth of children treated with grommets in infancy will have a recurrence of effusion once the grommets extrude from the tympanic membrane, yet, by the age of 6 years, hearing will have normalised in almost all affected children, irrespective of treatment (Johnston et al., 2004; Khodaverdi et al., 2013).

Resolution can also occur in CSOM, and healing of the tympanic membrane has been noted in some patients after treatment with topical antibiotics (Gupta et al., 2014; Smith et al., 1996). However, spontaneous resolution occurs less commonly than in cases of COME. In a study of 549 children in Greenland, 9% were found to have CSOM and, when a subset of this cohort was followed up after 15 years, the tympanic membrane had healed in only one third (Jensen et al., 2012).

**Existing model systems for chronic otitis media**

The mouse has become the preferred animal model for OM research owing to the availability of suitable reagents, low husbandry costs, genetic tractability, a well-characterised immune response and well-defined microbiological status (Fig. 4) (Bhutta, 2012). The long-tailed chinchilla has also been used for OM research because its large middle ear and Eustachian tube more closely resemble the anatomy of humans, and make it easier to recover effusion for microbiological or immunological analysis (Doyle, 1985; Jurcisek et al., 2003). For these reasons, the chinchilla has proven to be especially useful in vaccine studies.
and, because the chinchilla is outbred, it also better models the heterogeneity of immunological response compared to inbred animals (Green et al., 1994; Novotny et al., 2013). Other rodents used to study OM include rats, guinea pigs and gerbils (Bhutta, 2012; Sabirov and Metzger, 2008). Non-rodent animal models are rarely used to study this condition.

In vitro cell-culture models to study OM have also been developed using immortalised human middle ear epithelial cells (Chun et al., 2002), rat mucosal explants (Hill et al., 1992) and murine primary middle ear epithelial cells (Mulay et al., 2016; Tsuchiya et al., 2005) (Fig. 4D). Although these cell-culture models enable host-pathogen interactions to be assessed at the epithelial surface (Samuel et al., 2002), bacterial exposure and persistence, and cytokine production, they do not fully replicate the complexity of the in vivo microbial and immune responses in vivo. In this schematic, white broken arrows denote postulated links between OM-related conditions and unbroken arrows represent known links. Risk factors and therapies associated with pathogen exposure (left) and with host characteristics and inflammatory response (right) for acute and chronic forms of OM are shown. There are two forms of chronic mucosal OM: COME (chronic OM with effusion) and CSOM (chronic suppurative OM). Note that OME and COME can occur without antecedent AOM. The causes of CSOM are not well understood. AOM, acute otitis media; rAOM, recurrent AOM.
2008; Palacios et al., 2004; Val et al., 2016a), they cannot fully recapitulate the complexity of host-pathogen interaction in vivo, which is needed to understand the pathophysiology of chronic OM.

There are currently no experimental models that fully replicate the development or progression of chronic OM in humans. AOM is induced in animal models either by injection of bacteria directly into the bulla (Box 1; Oishi et al., 2013) or by intranasal inoculation coupled with viral co-infection (Langeries et al., 2012) or nasal pressurisation to facilitate ascension of bacteria from the nose to the middle ear (Chaney et al., 2011; Fig. 4A). The injection of less-virulent bacteria or of a reduced bacterial load leads to histological changes that resemble OME rather than AOM (Hermansson et al., 1988). In such models, inflammation is short-lived, the infection resolves within days and transition from AOM to chronic OM is not observed. Chronic OM (without preceding AOM) can be induced in mice or rats either through surgical obstruction of the Eustachian tube (Varsak and Santa Maria, 2016; Santa Maria et al., 2015; Hirano et al., 2016) or through genetic mutation (Zheng et al., 2006) (Fig. 4B,C).

Current genetic mutant mouse models of chronic OM are listed in Table S1 and reveal that a wide range of biological mechanisms can result in chronic OM in mice. OM penetrance ranges from ∼30% in the BpifA1-null mouse mutant (BpifA1 encodes the innate immune response protein BPI fold-containing family A member 1) (Bartlett et al., 2015) to ∼80% in a mouse carrying a point mutation at the immunomodulatory Mecom (MDS1 and EVI1 complex) locus (Parkinson et al., 2006). OM mouse models recapitulate many of the features of human chronic OM. The effusion that accumulates in the middle ear (bulla) varies from serous in mice that carry a point features of human chronic OM. The effusion that accumulates in the middle ear (bulla) varies from serous in mice that carry a point mutation in the gene encoding the G-protein-coupled receptor oxoglutarate receptor 1), and in mice carrying a point mutation in the pattern-recognition receptor Tlr4 (Toll-like receptor 4) (Kerschner et al., 2013; MacArthur et al., 2006). Cholesterol granulomas are seen in mice with a semi-dominant point mutation in the gene encoding ribosomal protein L38 (RpL38) and in mice with a semi-dominant mutation in the gene encoding the nuclear scaffold lamin A/C proteins (Lmna) (Noben-Trauth and Latoche, 2011; Zhang et al., 2012). Foreign-body granulomas have also been found in mice null for ectodysplasin A (Eda) and its receptor (Edar), which are involved in ectodermal morphogenesis (Azar et al., 2016).

It is important to note that the pathology of mouse models also differs from certain aspects of human COME and CSOM. No mouse mutant develops the tenacious fluid that typifies the effusion of the human mucoid form of COME, although there is evidence of a modest increase in mucus production. Mucin genes are upregulated in the mucosa of the Osrgr1 mouse mutant, and there is goblet cell hyperplasia in Lmna, Eda and Edar mutants. Goblet cell hyperplasia is also seen with OM in mice carrying a null mutation in the following genes: the chromatin-remodelling gene Chd7 (chromodomain-helicase-DNA-binding protein 7), the transcriptional co-activator Ey4 (EYA transcriptional coactivator and phosphatase 4), the immunomodulatory gene Tgf (TGFB induced factor homeobox 1) and the structural protein Sh3pxd2b (SH3 and PX domains 2B) (Tian et al., 2012; Depreux et al., 2008; Tateossian et al., 2013; Yang et al., 2011). Goblet cell hyperplasia also occurs as a result of chromosomal microdeletion in the Df1 mouse model of the human 22q11 deletion syndrome (Fuchs et al., 2013). Importantly, COME in children may resolve either spontaneously or after successful grommet treatment, whereas spontaneous remission of chronic OM is not documented in animal models.

The hallmark features of human CSOM, namely tympanic membrane perforation and purulent otorrhoea, are uncommon sequelae in genetic mouse models. In the Mecom mutant, otorrhoea occurs in conventionally housed low-health-status mice over 6 months of age but not in high-health-status specific-pathogen-free (SPF) conditions (Parkinson et al., 2006). Many other mouse OM models have not been assessed at this age and so it is possible that they could also develop otorrhoea if allowed to age. The findings in the Mecom mouse support the argument that laboratory mice should experience more normal environmental exposure to natural pathogens in order to better model human microbial exposure (Beura et al., 2016).

Several authors have attempted to induce CSOM in rodents through surgical means. In wild-type mice, surgical tympanic membrane perforation followed by the introduction of infection does not lead to chronic otorrhoea, and the tympanic membrane usually heals (Wang et al., 2014). Tympanic perforation in mutant Mecom mice also heals within 5 days, despite the presence of a pre-existing chronic purulent effusion (Bhutta et al., 2014). CSOM can be reliably induced in mice and rats by a combination of tympanic membrane perforation, blockade of the Eustachian tube, prevention of tympanic membrane healing (through grommet insertion or through the application of the matrix metalloprotease inhibitor KB-R7785) and by infection with Pseudomonas aeruginosa or Streptococcus pneumoniae (Santa Maria et al., 2015; Silva et al., 2012). It is still uncertain whether these are good models of CSOM because, in human disease, the Eustachian tube is usually normal or only partially obstructed (Bhat et al., 2009). Further studies of disease pathogenesis with these CSOM models are warranted.

Chronic OM is a feature of human syndromic conditions, such as Down syndrome, hypohidrotic ectodermal dysplasia (HED), primary ciliary dyskinesia (PCD), mucopolysaccharidosis and 22q11.2 deletion syndrome, and is also observed in the mouse strains that bear the comparable genetic lesions (Table S1). Other current mouse models of chronic OM have syndromic features, and so the pathology and mechanisms described in these models must be translated cautiously to non-syndromic disease in humans. In addition, some syndromic mouse models, such as perinatal lethal neurofibromatosis type 2, have OM (Giovannini et al., 2000), but this is not a feature of the equivalent human condition.

The role of pathogens in chronic otitis media
COME is often preceded by AOM, and so it is likely that bacterial and/or viral pathogens initiate inflammation in such cases (Fig. 5). It is generally accepted that infection with an upper respiratory virus often precedes bacterial AOM. The main bacteria that cause AOM are S. pneumoniae (pneumococcus), non-typeable Haemophilus influenzae (NTHi) and Moraxella catarrhalis (Ngo et al., 2016). These bacteria are also found in the middle ear effusion of children with chronic OM, but microbiome studies reveal that a multitude of other bacterial species are also present, in both COME (Chan et al., 2016; Jervis-Bardy et al., 2015) and CSOM (Neeff et al., 2016).

The role of bacteria in contributing to the persistence of inflammation in COME is not clear. There is some evidence that children who have recurrent episodes of AOM are more likely to have middle ear effusion (Alho et al., 1995) but it is uncertain
whether this represents repeated episodes of acute resolving effusion or continuous non-resolving effusion. Different studies using culture or molecular identification have detected bacterial DNA in 14-73% of effusions from children with COME (this wide range may reflect differing sensitivity of detection methods), and NTHi is more likely to be present than is pneumococcus (Ngo et al., 2016). Live bacteria have been found in a biofilm matrix on mucosal surfaces and/or in middle ear effusion (Thornton et al., 2013; Hall-Stoodley et al., 2006), intracellularly within mucosal cells and planktonically within the effusion (Thornton et al., 2011) (Fig. 5B, C). There is evidence that the effusion in COME is more likely to resolve in children who are given antibiotics (Venekamp et al., 2016), although that effect is relatively small.

In CSOM, bacteria play a more definite role in disease perpetuation. The prevalence of CSOM correlates with socioeconomic deprivation and malnutrition, both between and within countries (Bhutta, 2014; Lasisi et al., 2007; Chadha et al., 2006; Shaheen et al., 2012; Ooledge and Nwawolo, 2003). These factors likely have an impact on immune responses, on the risk of pathogen exposure and on pathogen load. Microbiological culture of middle ear effusion from children with CSOM yields a mix of aerobic and anaerobic bacteria (Verhoeff et al., 2006), usually with a predominance of Staphylococcus aureus and P. aeruginosa (Mittal et al., 2015). Bacteria in CSOM also exist in a biofilm (Lee et al., 2009; Saunders et al., 2011; Gu et al., 2014; Homae et al., 2009). Topical or oral antibiotics may stop otorrhoea in patients with CSOM, but treatment will frequently fail and it is also
not known how often antibiotic treatment enables the long-term resolution of disease, including healing of the tympanic membrane (Macfadyen et al., 2005, 2006).

All chronic OM mouse mutants develop disease spontaneously without the need for bacterial challenge. Wild-type SPF mice have a diverse nasal microbiome (Krone et al., 2014) and nasal commensals are a potential source for bulla infection. Nasal commensals have been cultured from the bullae of various mouse mutants that carry null mutations, including in genes involved in ectodermal morphogenesis (Eda, Edar and Mcph, which encodes microcephalin) (Azar et al., 2016; Chen et al., 2013), in chromatin remodelling (Chd7) (Tian et al., 2012), in transcription (Eya4, Dfi1 and Isl1, which encodes ISL LIM homeobox 1) (Depreux et al., 2008; Hilton et al., 2011; Fuchs et al., 2013), in protein synthesis (Rpl38) (Noben-Truth and Latoche, 2011) and in immune signalling (Mecom and Nfkbia, which encodes NFKB inhibitor alpha) (Schmidt-Ullrich et al., 2001; Parkinson et al., 2006). However, not all bulla fluids are culture positive (Table S1). The human otopathogens NTHi, pneumococcus and Moraxella catarrhalis have not been detected by PCR in Chd7 or Ovx1 mouse mutants, but M. catarrhalis was detected in the Spag6 (sperm-associated antigen 6)-null mutant, which has defects thought to result from disruption of the ciliary cytoskeleton (Li et al., 2014). Anaerobic culture and microbiome studies have yet to have performed in these OM mutants. Antimicrobial (azithromycin) treatment does not suppress OM in Eya4 mutants (Depreux et al., 2008). In Mecom mutants, OM initiates later, but occurs with the same frequency under germ-free and SPF conditions, suggesting that respiratory irritants, such as ammonia and dust from the cage environment, can act as inflammatory stimuli to initiate OM. Intranasal challenge of Mecom mutants with NTHi results in high rates of bulla infection lasting up to 56 days, and shows the potential of this model for treatment and prevention studies for chronic OM. There is no evidence of NTHi intraepithelial infection or biofilm formation in Mecom mice (Hood et al., 2016).

**The role of host factors in chronic otitis media**

The relative importance of host factors in the initiation and perpetuation of chronic OM can be gauged from estimates of the heritability of disease. Twin studies in young children with COME have suggested that heritability of the duration of middle ear effusion is high at 0.73 (Casselbrant et al., 1999). Studies of the Inuit population in Greenland report that parental history is an important predictor of CSOM in children, independent of socioeconomic status (Jensen et al., 2011; Koch et al., 2011).

Identifying genetic loci that are associated with chronic OM is one way in which to elucidate potential disease mechanisms. A number of human genetic-association studies for OM have been reported (reviewed in Rye et al., 2012a), but most early studies comprised small cohorts (and so were underpowered or at high risk of false discovery) and, in many, the condition was poorly defined (Bhutta, 2013). More recent studies have featured larger cohorts and better phenotyping (Table 1) but, nevertheless, they remain too small to discover loci that are associated with modest relative risk. The only OM association to have been replicated is at the Fbxo11 locus. Fbxo11 was initially associated with OM in an Australian cohort (Rye et al., 2011), with nominal evidence of association, and then replicated in a UK cohort (Bhutta et al., 2017) and a US cohort (Segade et al., 2006), albeit at different polymorphisms at Fbxo11. Data from the Fbxo11 mouse model (see below) suggest that a mutation in Fbxo11 perturbs transforming growth factor (TGF)-β signalling in the middle ear.

The factors that contribute to host susceptibility to chronic OM have also not been elucidated. The onset of AOM is presumed to involve the detection of (bacterial) antigens by innate immune receptors (such as the Toll-like receptors) and by other pattern-recognition molecules on cells within the middle ear mucosa, which will recruit neutrophils and lymphocytes to the middle ear. However, we do not know whether it is signalling and regulation by the mucosa, leukocytes, or both, that perpetuates inflammation in the chronic condition, and whether the inflammatory mechanisms involved are organ-specific or mirror mechanisms of chronic inflammation found in other tissues (Buckley et al., 2013; Val et al., 2016b).

Much of the older literature suggests that poor aeration of the middle ear due to ventilatory dysfunction of the Eustachian tube is a key factor in acute and chronic OM (Bluestone, 2005), yet there are no validated tests of Eustachian tube function to support this notion (Smith and Tysome, 2015). Moreover, anatomical modelling suggests that a narrowing of the Eustachian tube is unlikely to significantly impede gas exchange (Sadé et al., 2004). Additionally, no consistent or significant differences in Eustachian tube parameters have been demonstrated in individuals affected by OM.

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**Table 1. Human genetic linkage or association studies for susceptibility to OM**

<table>
<thead>
<tr>
<th>Study type</th>
<th>Reference</th>
<th>Cohort</th>
<th>Findings</th>
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<tbody>
<tr>
<td>Genome-wide association study</td>
<td>Allen et al., 2013</td>
<td>373 cases with rAOM or COME treated with grommets</td>
<td>Association with rs1110060 in Klf7 intron. Not replicated in an independent cohort</td>
</tr>
<tr>
<td>Genome-wide association study</td>
<td>Rye et al., 2012b</td>
<td>416 cases of OM, 1075 controls. Identified by parental questionnaire of symptoms and/or clinical findings</td>
<td>Top hits at CAPN14 (OR 1.86) and GALNT14 (OR 1.6). Not replicated in two independent cohorts (Allen et al., 2014)</td>
</tr>
<tr>
<td>Genome-wide association study</td>
<td>Einarsdottir et al., 2016</td>
<td>803 OM cases with rAOM or COME, 2073 controls</td>
<td>Association with 80 kb region on chromosome 19. Replication in UK cohort in opposite direction for risk</td>
</tr>
<tr>
<td>Linkage study</td>
<td>Chen et al., 2011</td>
<td>139 families with rAOM or COME treated with grommets</td>
<td>Linkage at 2.2 Mb region at 19q</td>
</tr>
<tr>
<td>Linkage study</td>
<td>Rye et al., 2014</td>
<td>468 families with rAOM or COME treated with grommets</td>
<td>Linkage at 4.2 Mb region at 10q23.6</td>
</tr>
<tr>
<td>Linkage study</td>
<td>Casselbrant et al., 2009</td>
<td>403 families with children treated with grommets</td>
<td>Linkage at 17q12 and 10q22.3</td>
</tr>
<tr>
<td>Candidate gene study</td>
<td>Bhutta et al., 2017</td>
<td>1296 families with COME treated with grommets</td>
<td>Association with polymorphism at Fbxo11 and TGF1</td>
</tr>
<tr>
<td>Candidate gene study</td>
<td>Hafren et al., 2015</td>
<td>624 cases with rAOM or COME, 778 controls</td>
<td>Association with polymorphism at TLR4. Not replicated</td>
</tr>
<tr>
<td>Candidate gene study</td>
<td>Rye et al., 2011</td>
<td>434 families with rAOM or COME treated with grommets</td>
<td>Association with polymorphism at Fbxo11. Replicated</td>
</tr>
</tbody>
</table>

*Only selected studies with a reasonable sample size are shown, although cited studies are nevertheless underpowered by the standards applied to detect genetic associations to a common disease. rAOM, recurrent acute otitis media; CAPN14, calpain 14; COME, chronic otitis media with effusion; GALNT14, polypeptide N-acetylgalactosaminyltransferase 14; Mb, megabase; OM, otitis media; OR, odds ratio.
(Sadé and Luntz, 1989; Sadé et al., 1986). However, in Djf1 and Tbx1 (T-box 1)-null mouse mutants (which model human 22q11 deletion syndrome), developmental hypoplasia of the levator veli palatini muscle, which is involved in opening and closing the anterior pharyngeal portion of the Eustachian tube, impairs the experimental transit of dye from the bulla (Fuchs et al., 2013; Liao et al., 2004). In Fbxo11 (Hardisty et al., 2003) and Eya4 (Depreux et al., 2008) mutants, the Eustachian tube can be malpositioned, narrowed or misshapen and, in Eya4 mutants, its opening can be blocked by inflammatory polyps. An adrenergic signalling defect in Ddbh (dopamine beta-hydroxylase) mutants might also impair tubal function (Maison et al., 2010). In addition, mouse mutants with craniofacial abnormalities, such as domed heads and alterations in the cranial base, can have Eustachian tube anomalies; for example, the angle at which the tubes join the nasopharynx is more acute in Sh3pxd2b, Lmna and Chd7 mutants (Zhang et al., 2012; Yang et al., 2011; Tian et al., 2012). In Rpl38 and Edar mutants, the Eustachian tube undergoes pathological dilatation through loss of adjacent submucosal glands (Azar et al., 2016; Noben-Trauth and Latoche, 2011), and, in Eda and Edar mutants, the gating function of the Eustachian tube is reduced, permitting larger foreign-body particles to enter the bulla (Azar et al., 2016). There are anatomical defects in the middle ear of other mouse mutants that could also be relevant; for example, defective postnatal bulla cavitation in the Eya4 null mutant (Depreux et al., 2008), hypoplastic but proportionate skulls in the microcephaly-associated Mcph1-null mutant (Chen et al., 2013), shortened and distorted nasal bones with small bullae in compound mutants of the transcription factors Ets1 (ETS proto-oncogene 1, transcription factor) and Fli1 (Fli-1 proto-oncogene, ETS transcription factor) (Carpinelli et al., 2015), and perturbation of epithelial growth in the Fbxo11 and Tgft mutants (Hardisty et al., 2003; Tatoossian et al., 2013).

The importance of mucociliary defects in the perpetuation of middle ear inflammation is not known. In primary cilia dyskinesia (PCD) syndrome, patients have an increased incidence of middle ear effusion, and mouse mutants with null mutations in the following ciliary structural proteins have OM and rhinitis due to impaired mucociliary clearance: Dnah5 (dynein axonemal heavy chain 5), Cby1 (chibby family member 1, beta catenin antagonist), Spag6, Dnah11 (dynein axonemal heavy chain 11), Odf2 (outer dense fiber of sperm tails 2), Tll1 (tubulin tyrosine ligase-like 1), Ulk4 (unc-51-like kinase 4), Kif27 (kinesin family member 27), Dpdc (deleted in primary ciliary dyskinesia homolog) and Shk36 (serine/threonine kinase 36) (Ibanez-Tallon et al., 2002; Voronina et al., 2009; Li et al., 2014; Lucas et al., 2012; Kunimoto et al., 2012; Vogel et al., 2010, 2012). Mcph1 mutant mice are also suspected to have cilia defects, as well as the null mutant at Poren (porcupine O-acyltransferase), a gene involved in the processing of proteins by the endoplasmic reticulum (Chen et al., 2013; Biechele et al., 2013). In Lmna, Chd7, Eya4 and Phex (phosphate regulating endopeptidase homolog, X-linked) mutants, inflammation causes the loss of ciliated cells in the bulla epithelium, whereas perturbation of phosphorylation in the fibroblast growth factor (FGF)23/prostaglandin (PG)E2 pathways in the Phex hypomorph might also reduce the aqueous periocular layer and impair the clearance of overlying mucus (Han et al., 2012; Zhang et al., 2012; Tian et al., 2012; Depreux et al., 2008).

**Mechanisms leading to chronic inflammation and resolution**

What enables chronic OM to resolve is unclear, but this seems an important avenue of research. This is because, by better understanding the clearance mechanisms that lead to resolution, we might be able to clinically manipulate the disease to hasten resolution, and thereby reduce disability.

At a molecular level, the resolution of inflammation in tissues other than the middle ear is complex, involving the depletion of chemokines, the downregulation of pro-inflammatory cytokines, the upregulation of pro-resolution mediators, neutrophil apoptosis and the alternative activation of macrophages (Sugimoto et al., 2016). It seems likely that similar mechanisms will operate in mucosal cells and in leukocytes to resolve inflammation in the chronically inflamed middle ear, but which of these mechanisms is important, and how they are regulated, is not known.

Existing clinical treatments for chronic OM can be highly effective, but investigation into their mechanisms of action has been limited. In COME, symptoms result from effusion, and the insertion of grommets is effective in resolving that effusion (Browning et al., 2010). Grommets are often presumed to have a rheological effect but there is no evidence that grommets affect the ventilatory function of the Eustachian tube in the short (van der Avoort et al., 2009), medium (Takahashi et al., 1990; van Heerbeek et al., 2001; Streetemans et al., 2005), or long (Cayé-Thomasen et al., 2008) term. Myringotomy (Box 1) in a mouse model reduced tissue hypoxia and inflammatory effusion, suggesting that oxygen tension is important for middle ear homeostasis, and this might be an alternative explanation for the efficacy of grommets (Bhutta et al., 2014). Once grommets extrude, middle ear effusion can recur, leading to a repeat operation in one quarter of treated children (Browning et al., 2010). This suggests that grommets have little long-term effect on the goblet cell hyperplasia that generates middle ear effusion.

In CSOM, symptoms result from perforation of the tympanic membrane, which is associated with ongoing intermittent or chronic inflammatory and infected otorrhoea. Mechanisms underlying the healing or non-healing of the tympanic membrane are not well understood (Jung et al., 2013) but, where the tympanic membrane does not heal spontaneously or following medical treatment with antibiotics, surgical repair is often successful. Nevertheless, one in six attempts at surgical repair will fail, which may be due to a number of factors; however, evidence of continuing chronic inflammation in the contralateral ear (in the form of COME) has been shown to be predictive of failure (Hardman et al., 2015).

Molecular targeting may offer more reliable clinical resolution of chronic OM in the future. This notion is still some distance from the bedside, but mouse models have suggested pathways that could be targeted. For example, genome-wide transcriptional analysis of acute OM, induced in mice through transbullar injection, reveals an early response at 6 h that is dominated by immune and defence proteins, and a later response at 24-48 h that is dominated by immunoregulatory proteins (Hernandez et al., 2015). However, inflammation resolves within 5-7 days of inoculation, and so this model provides little insight into the molecular signatures that underlie the transition to chronic disease. In mouse mutants, deficits in innate immunity are known to contribute to persistent inflammation. In the Tlr4 mouse mutant, this occurs via altered responses to Gram-negative bacteria and, unusually for mouse mutants, the inflammatory changes extend into the inner ear (MacArthur et al., 2006). Other examples include the BpsiA1 mutant, in which the loss of the antimicrobial/surfactant protein product SPLUNC1 impairs auditory tube function (Bartlett et al., 2015), and the hypohidrotic ectodermal dysplasia (HED) mutants IkbαΔN, EdarαβΔJdJ, Edaαγ and Edaαβ, in which the nasal and nasopharyngeal glands and their products are absent (Schmidt-Ullrich et al., 2001; Azar et al., 2016). Impaired bulla mucosa...
secretion and response to Gram-negative bacteria is predicted in the Isl1 mutant owing to the known interaction of this gene with innate immune signalling pathways (Hilton et al., 2011). Impaired adrenergic signalling in Dhb mutants might also affect the systemic and mucosal adaptive immune system (Maison et al., 2010).

Other models have suggested that perturbation of the TGF-β, NF-κB or HIF (hypoxia-inducible factor) signalling pathways can drive chronic OM. In Fbxo11+/− and Tgff1−/− mutants, TGF-β signalling is disrupted (Tateossian et al., 2015, 2013). TGF-β regulates the differentiation, proliferation and activation of several immune cells and, in sites other than the middle ear, persistent TGF-β activation has been found to promote the transition from acute to chronic inflammation, including fibrosis (Yoshimura et al., 2010). TGF-β levels also correlate with duration of effusion in children with COME (Zhao et al., 2009). Both HIF and NF-κB signalling are activated in response to cellular stress, which is induced by pathogens but also by chemical or physical damage (via NF-κB signalling) or cellular hypoxia (via HIF signalling). There is considerable crosstalk between these pathways (D’Ignazio et al., 2016) and they induce inflammation as part of a strategy to promote cellular survival, but persistent activation can lead to non-resolving inflammation. MECOM acts as an inducible negative regulator of NF-κB, and loss of Mecom function in mice results in an elevated response to inflammatory stimuli, including to challenge with NTHi (Xu et al., 2012). In Mecom, Fbxo11 and Tgff1 mutant mice, leukocytes in the bulla fluid respond to inflammatory hypoxia by upregulating VEGF (vascular endothelial growth factor), a downstream effector in the HIF pathway. In Mecom mutants, tissue hypoxia extends to the mucosa, suggesting that hypoxia might be a common finding in the chronically inflamed middle ear (Cheeseman et al., 2011). HIF signalling was also upregulated in response to Eustachian tube blockage in a rat model of OM (Huang et al., 2012). VEGF signalling has been demonstrated in effusions of children with COME, and NF-κB in the mucosa of patients with CSOM (Sekiyma et al., 2011; Jesic et al., 2014).

Systemic administration of VEGF-receptor antagonists has been shown to ameliorate the progression of hearing loss in young Mecom mice before chronic OM is established (Cheeseman et al., 2011). Molecules to target NF-κB and TGF-β pathways have also been developed (Gilmore and Garbati, 2011; Fabregat et al., 2014) but have yet to be trialled for chronic OM.

**Conclusions and future outlook**

Despite the considerable prevalence of chronic OM worldwide, especially in childhood, many unanswered questions remain about its aetiology and about how to treat this disease (as summarised in Box 3). In terms of epidemiology, existing studies provide some understanding of the microbiological and host factors that predispose children to COME, but we do not understand what initiates the disease in those without a history of AOM, nor whether CSOM is a more severe variant of COME.

In terms of elucidating the pathobiology of chronic OM, we still need to understand the relative roles of mucosal versus leukocyte biology in the initiation and perpetuation of middle ear disease, and the role of pathogens and their interaction with host tissues. We do not know whether ventilatory dysfunction in the Eustachian tube is as important a mechanism in pathogenesis as was historically proposed (Bluestone, 2005). This question has become more pertinent with the recent development of balloons, which are used to surgically dilate the Eustachian tube with the aim of permanently improving ventilation of the middle ear in individuals with chronic middle ear disease (Miller and Elhassan, 2013; Norman et al., 2014). The efficacy of these balloons has not been established. Where COME is concerned, epidemiological studies tell us that host factors likely play a significant role. Yet, to date, human genetic-association studies have been underpowered, often poorly phenotyped and have an insufficient number of cohorts to enable the replication of their results.

In addition to human studies, animal models can be a powerful way to explore disease mechanisms. Mice have been used extensively to study chronic OM and its associated conditions, but none fully recapitulate the clinical features of COME or CSOM. The recent development of a mouse model of chronic otorrhea, using surgical perforation of the tympanic membrane and Eustachian tube obstruction, represents a significant advance (Varsak and Santa Maria, 2016), but the model still lacks some important characteristics of human disease. The existence of large-scale mouse mutagenesis programs and new gene-editing techniques, such as CRISPR/Cas9, should also be harnessed to identify relevant mutants (Horii and Hatada, 2017), with a focus on identifying those models that faithfully recapitulate human disease and rejecting those that do not. To identify such models, we need to better understand how individuals are genetically predisposed to chronic OM and how the disease progresses. Any potentially informative model thus identified needs to then undergo detailed molecular and biological study to identify the underlying pathogenic mechanisms and the means by which these can be therapeutically manipulated to resolve effusion and/or otorrhea.

We have little understanding of how current clinical treatments for chronic OM work at a molecular level. If we understood this better, perhaps we could offer more effective or reliable therapies. For example, antibiotics can sometimes stop otorrhea in CSOM but we do not know to what extent they enable its long-term resolution and the healing of the tympanic membrane. We do not understand why the integrity of the tympanic membrane has such a profound effect on middle ear immunology. Creating a perforation of the tympanic membrane using grommets in children with COME leads to resolution of effusion but, conversely, surgical repair of a perforated tympanic membrane in individuals with CSOM leads to resolution. The use of bulla explants from animal models might provide an avenue for furthering our understanding of the role of the tympanic membrane in the aetiology of this disease.

Laboratory mice are widely used in OM research and, although most genetic models of chronic OM are syndromic, they provide important insights into, and a means by which to explore, the homeostatic mechanisms of the middle ear cleft. In particular, the innate immunity of the middle ear is likely to be relevant to the

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**Box 3. Outstanding clinical and basic research questions**

- What leads to CSOM? Longitudinal epidemiological studies of OM are needed to address this question.
- What are the relative roles of mucosal biology, leukocyte biology, pathogens and Eustachian tube function in the perpetuation and resolution of COME and CSOM?
- How do we generate larger and well-phenotyped cohorts for genetic studies into human chronic OM?
- How do we create new and better animal models of chronic OM, and better evaluate their phenotypic relevance to human disease?
- How should we investigate the mechanisms that underlie current therapeutic interventions, including the long-term effects of antibiotic therapy for CSOM, and the immunobiological effects of perforation or repair of the tympanic membrane?
aclioogy of non-syndromic chronic OM in humans, and we need therefore to better understand its mechanisms.

We should also not overlook the opportunities to study the transition between disease and health by extending the time course of induced AOM models. Our current chronic OM mouse models might pass the point where changes such as mucosal fibrosis are reversible. The prospects for generating a genetic mouse mutant that is an exact model of non-syndromic COME or CSOM are doubtful given species differences in size and anatomy, such as the presence of adenoidal lymphoid tissue and bulla mastoid cells in humans (Bhutta, 2012). Nevertheless, there is scope to discover more about the pathobiology of chronic OM in mutant mice, and to use them as novel translational models to validate candidate genes, pathways and experimental tools. New techniques, such as designer-nuclease gene editors, make it possible to also engineer candidate OM genes in large animal species (Whitelaw et al., 2016), and to use them as realisation of the prevalence and morbidity associated with chronic inflammation of the middle ear.

Competition interests
The authors declare no competing or financial interests.

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References


Homoe, P., Bjarnsholt, T., Wessman, M., Sørensen, H. C. F. and Johansen, H. K.

Ibanez-Tallon, I., Gorokhova, S. and Heintz, N.

Jama 296, 948-954.

JAMA 75, 64-70.

JAMA Otolaryngol. 207, 1071-1079.

JAMA Otolaryngol. 134, 431-441.


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Supplementary Table S1
Current genetic mutant mouse models of chronic otitis media (OM). PCD = primary ciliary dyskinesia

<table>
<thead>
<tr>
<th>mouse gene</th>
<th>reference</th>
<th>human syndrome or condition</th>
<th>adult mouse phenotypes additional to otitis media</th>
<th>craniofacial abnormality</th>
<th>Eustachian tube</th>
<th>bulla bacteria</th>
<th>inner ear</th>
<th>comment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Down Syndrome</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>T&lt;sub&gt;s&lt;/sub&gt;65D&lt;sub&gt;n&lt;/sub&gt; Trisomy</td>
<td>(Han et al., 2009)</td>
<td>Down syndrome</td>
<td>behavioural and cognitive deficits, short narrow palate</td>
<td>short narrow palate</td>
<td></td>
<td></td>
<td></td>
<td>increase in goblet cells, effusion, fibrous proliferation Staph. coagulase-negative, Bordetella avium, Burkholderia cepacia labyrinth serous effusion; hair cells, stria vascularis &amp; spiral ganglion normal</td>
</tr>
<tr>
<td>Dp(16)1Yey Trisomy</td>
<td>(Bhutta et al., 2013)</td>
<td>Down syndrome</td>
<td></td>
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<td></td>
<td></td>
<td></td>
<td>thickened mucoperiosteum, polyps</td>
</tr>
</tbody>
</table>
### Primary Ciliary Dyskinesia (PCD)

<table>
<thead>
<tr>
<th>Mutant</th>
<th>Mutant Type</th>
<th>Phenotype</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Mdnah5</em> -/-</td>
<td>(Ibanez-Tallon et al., 2002)</td>
<td>PCD hydrocephaly, growth retardation, mortality 2-3 weeks, situs inversus, domed head</td>
<td></td>
</tr>
<tr>
<td><em>Dnahc11</em> iv/iv</td>
<td>(Lucas et al., 2012)</td>
<td>PCD situs inversus, rhinitis, sinusitis</td>
<td>immotile tracheal cilia</td>
</tr>
<tr>
<td><em>Cby</em> +/-</td>
<td>(Voronina et al., 2009)</td>
<td>PCD rhinitis, growth retardation, postnatal mortality by d30, anaemia, reduced subcutaneous fat, no situs inversus</td>
<td>cilia deficiency and motility deficits, experimental challenge with <em>P. aeruginosa</em></td>
</tr>
<tr>
<td><em>Odf2</em> ΔEx6,7/ΔEx6,7</td>
<td>(Kunimoto et al., 2012)</td>
<td>PCD rhinitis, stertor, growth retardation, infertility, hydrocephaly variable</td>
<td></td>
</tr>
<tr>
<td><em>Till1</em> +/-</td>
<td>(Vogel et al., 2012)</td>
<td>PCD not known with this mutation</td>
<td>PCD rhinosinusitis, male infertility, normal lungs, supplicative exudate</td>
</tr>
<tr>
<td>Mutant</td>
<td>Genotype</td>
<td>Phenotype</td>
<td>Pathological Findings</td>
</tr>
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<td>------------</td>
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<td>-----------------------------------------------</td>
<td>--------------------------------------------------------------------------------------</td>
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<tr>
<td>Spag6 -/-</td>
<td>(Li et al., 2014)</td>
<td>PCD</td>
<td>hydrocephalus, sperm motility defects, morbidity and mortality after 6 months</td>
</tr>
<tr>
<td>Ulk4 -/-</td>
<td>(Vogel et al., 2012)</td>
<td>PCD</td>
<td>PCD, rhinosinusitis, hydrocephalus, mortality &lt; 4 months</td>
</tr>
<tr>
<td>Kif27 -/-</td>
<td>(Vogel et al., 2012)</td>
<td>PCD</td>
<td>PCD, rhinosinusitis, hydrocephalus, mortality &lt; 8 weeks</td>
</tr>
<tr>
<td>Stk36 -/-</td>
<td>(Vogel et al., 2012)</td>
<td>PCD</td>
<td>PCD hydrocephalus, morbidity mortality &lt; 6 weeks</td>
</tr>
<tr>
<td>Dpcd -/-</td>
<td>(Vogel et al., 2012)</td>
<td>PCD</td>
<td>PCD, rhinosinusitis, hydrocephalus, male infertility</td>
</tr>
<tr>
<td>Mucopolysaccharidosis</td>
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<tr>
<td><strong>Ids KO</strong></td>
<td>(Hong et al., 2012)</td>
<td>mucopolysaccharidosis type II, Hunter syndrome X-linked</td>
<td>TM thickened, microCT opacity, bulla exudate</td>
</tr>
<tr>
<td><strong>Naglu -/-</strong></td>
<td>(Heldermon et al., 2007)</td>
<td>mucopolysaccharidosis type III, Sanfilippo syndrome</td>
<td>behavioural and sensory deficits, reduced lifespan</td>
</tr>
<tr>
<td><strong>Idua -/-</strong></td>
<td>(Schachern et al., 2007)</td>
<td>mucopolysaccharidosis type I</td>
<td>mucosa thickened and granulation tissue formation</td>
</tr>
<tr>
<td><strong>Gus mps2J/mps2J</strong></td>
<td>(Vogler et al., 2001)</td>
<td>mucopolysaccharidosis type VII</td>
<td>reduced survival, reduced lifespan, skeletal dysostosis, thickened tail</td>
</tr>
</tbody>
</table>
### 22q11.2 Deletion Syndrome (DiGeorge Syndrome or Velo-Cardio-Facial Syndrome)

<table>
<thead>
<tr>
<th>Genotype</th>
<th>Reference</th>
<th>Syndrome Overview</th>
<th>Cardiovacular Abnormalities</th>
<th>Levator Veli Palatini Hypoplasia</th>
<th>Thickened Mucosa, Vascularization</th>
<th>E. coli, Lactococcus lactis, Pantoea sp.</th>
<th>Hair Cell Loss</th>
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</thead>
<tbody>
<tr>
<td>$Df1^+/+$</td>
<td>(Fuchs et al., 2013)</td>
<td>22q11.2 Deletion Syndrome</td>
<td>none reported</td>
<td>levator veli palatini hypoplasia</td>
<td>thickened mucosa, vascularization, reduced numbers of ciliated cells</td>
<td></td>
<td>no hair cell loss detected</td>
</tr>
<tr>
<td>$Tbx^+/+$</td>
<td>Fuchs et al., 2015</td>
<td>22q11.2 Deletion Syndrome</td>
<td>cardiovascula r abnormalities</td>
<td>levator veli palatini hypoplasia</td>
<td>mucosa proliferation, vascularization, effusion, increased goblet cells</td>
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### Hypohidrotic ectodermal dysplasia

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<tbody>
<tr>
<td>$Eda Ta/Y, Eda Ta/Y, Eda dlJ/dlJ$</td>
<td>(Azar et al., 2016)</td>
<td>Hypohidrotic Ectodermal dysplasia (HED) OMIM 300451; EDAR 604095</td>
<td>rhinitis, nasopharyngitis, otitis media, HED, loss of nasal and nasopharyngeal submucosal glands</td>
<td>secondary dilation</td>
<td>mucosal thickening, polyps, foreign body granuloma</td>
<td>Gemella sp. Enterococcus sp. E. coli, Staphylococcus sp., Staph. aureus</td>
<td></td>
<td></td>
</tr>
<tr>
<td>$IKB \alpha AN$</td>
<td>(Schmidt-Ullrich et al., 2001)</td>
<td>HED, immune defects lymph nodes absent, leukocytosis, impaired macrophage function, growth retardation</td>
<td>domed head</td>
<td>mucosal thickening</td>
<td>Staphylococcus aureus</td>
<td>normal</td>
<td>Leishmania challenge</td>
<td></td>
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</table>
### Other syndromes

<table>
<thead>
<tr>
<th>Syndrome</th>
<th>Description</th>
<th>Rhinitis</th>
<th>Polycystic Kidney Disease</th>
<th>Whole Genome</th>
<th>Characteristic of Control Cells</th>
<th>Spectroscopy</th>
<th>Normal Cilia</th>
<th>Functional Assay</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Porcn +/del</strong></td>
<td>(Biechele et al., 2013)</td>
<td>Focal Dermal Hypoplasia (Goltz Syndrome, OMIM#305600)</td>
<td>rhinitis, bronchopneumonia, mild hydrocephalus, dermal and skeletal defects</td>
<td>bulla exudate</td>
<td>cilia defect suspected</td>
<td></td>
<td>Streptococci present in mutants and wild-type. PCR for human H. influ, S. pneumo, M. catarrh</td>
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<td></td>
</tr>
<tr>
<td><strong>Chd7 Ome/+</strong></td>
<td>(Tian et al., 2012)</td>
<td>CHARGE syndrome abnormalities of eye, heart, choana, genitalia, middle and inner ear</td>
<td>keratitis sicca, growth retardation, circling behaviour</td>
<td>increased lateral angulation</td>
<td>increased goblet cells, reduced ciliated cells</td>
<td>normal hair cells and spiral ganglia</td>
<td>Muc5AC, Muc5B, Tgfβ1</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Lmna Dhe/+</strong></td>
<td>(Zhang et al., 2012)</td>
<td>LMNA laminopathies includes Hutchinson-Gilford progeria syndrome, pinna &amp; external ear canal anomaly, conductive hearing loss</td>
<td>epidermal dysplasia, craniofacial defects and reduced pinna, hyperphosphatemia and increased calcium x phosphate product in female mutants</td>
<td>increased lateral angulation, increased width of bony segment more horizontal</td>
<td>mucosal thickening increased goblet cells, haemorrhage, erythrophago cytosis, cholesterol granuloma, abscessation</td>
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</tr>
<tr>
<td>Mutant</td>
<td>Phenotype</td>
<td>Genotype</td>
<td>Mortality</td>
<td>Bone/Mucosa</td>
<td>Other</td>
<td>Azithromycin Treatment</td>
<td>Cilia Defect</td>
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<tr>
<td><em>Ets1 +/-</em> and <em>Fli1 +/-</em></td>
<td>Jacobsen syndrome, perinatal mortality, <em>Ets</em> and <em>Fli1</em> thrombocytopenia, short nasal bone, malformed septum</td>
<td>(Carpinelli et al., 2015)</td>
<td>none detected</td>
<td>inflamed mucosa, small bullae, deformed stapes</td>
<td>none detected</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>Mcph1 tm1a/tm1a</em></td>
<td>Microcephaly, mental retardation, homozygotes infertile, ocular abnormalities, increased B cells &amp; micronucleated normochromic erythrocytes, small skull</td>
<td>(Chen et al., 2013)</td>
<td>normal</td>
<td>thickened mucoperiosteum</td>
<td>Streptococcus sp.</td>
<td>cilia defect suspected</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>Eya4 -/-</em></td>
<td>Senoineural hearing loss, cardiomyopathy, none reported</td>
<td>(Depreux et al., 2008)</td>
<td>reduced size, malposition</td>
<td>reduced bulla size, mucosal hyperplasia, reduced cilia, increased goblet cells</td>
<td>azithromycin treatment has no effect on phenotype</td>
<td></td>
<td></td>
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<td></td>
</tr>
<tr>
<td><em>Phex Hyp-Duk/Y</em></td>
<td>X-linked hypophosphatemic rickets, small body size and short tail, slightly abnormal skull shape, hyperplasia of ciliated epithelium, increased goblet cells, reduced ciliated cells, overlaid by patches mucin-like material</td>
<td>(Han et al., 2012)</td>
<td></td>
<td></td>
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</tr>
</tbody>
</table>
### Anatomical defects

<table>
<thead>
<tr>
<th>Rpl38 Ts/+</th>
<th>(Noben-Trauth and Latoche, 2011)</th>
<th>skeletal abnormality, short misshapen tail, hyperphosphataemia</th>
<th>shortened nose</th>
<th>enlarged</th>
<th>polyps, cholesterol granuloma, mineralisation, ectopic calcification, osteogenesis</th>
<th>Staphylococcus spp, Enterococcus sp.</th>
<th>no abnormality</th>
<th>rescued by Rpl38 cDNA</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sh3pxd2b nee homozygote</td>
<td>(Yang et al., 2011)</td>
<td>glaucoma, growth retardation, adipose deficits, hearing impairment</td>
<td>domed head, rounded skull, short maxilla and nose</td>
<td>inflammatory cells in ET lumen, increased lateral with concurrent reduced horizontal angulation</td>
<td>thickened fibrous mucosa with increased goblet cells and decreased ciliated cells, TM thickened</td>
<td>inner ear inflammation minor</td>
<td>increased Tnf and Tlr2</td>
<td></td>
</tr>
</tbody>
</table>

### TGF-β signalling pathway

| Fbxo11 Jf/+ | (Hardisty et al., 2003, Hardisty-Hughes et al., 2006) | homozygote cleft palate, eyelids open at birth, heterozygotes growth retardation | short snout | New-born and d50 mice ET narrowed and misshapen | bulla reduced size, mucosa dilated capillaries & lymphatics, polyps, calcification in mice > 11 months | abnormal endocochlear potentials, possible impaired stria function |
|---|---|---|---|---|---|---|---|
| Tgif -/- | (Tateossian et al., 2013) | holoprosencephaly | homozygote placental defect and minor hydrocephalus | minor skull shortening and reduced size | normal acute angle to nasopharynx | thickened mucoperiosteum, increased goblet cells | none at 2 months | bulla fluids elevated MMP, VEGF, TNFa |
Innate immunity

<table>
<thead>
<tr>
<th>Model</th>
<th>Description</th>
<th>Genotype</th>
<th>Phenotype</th>
<th>Infection</th>
<th>Other Effects</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mecom (Evi1) Jbo/+</td>
<td>Extra digit, derepression of NF-kB signalling</td>
<td>(Azar et al., 2016, Bhutta et al., 2014, Cheeseman et al., 2011, Hood et al., 2016, Parkinson et al., 2006, Xu et al., 2012)</td>
<td>Mucosa thickened polyps, vascularization</td>
<td>Staph sp., Staph aureus, Proteus mirabilis, E. coli, Moraxella sp.</td>
<td>Bulla fluids elevated cytokines and VEGF. Experimental NTHi infection</td>
</tr>
<tr>
<td>Tlr4 Lps-d/Lps-d</td>
<td>Lumen filled by exudate</td>
<td>MacArthur et al., 2006</td>
<td>Mucosal fibrosis, granulation tissue, vascularization and bulla bone remodelling</td>
<td>Normal</td>
<td>Round window and organ of Corti inflammation</td>
</tr>
<tr>
<td>BpifAI (Splunc-1) +/-</td>
<td>Normal</td>
<td>(Bartlett et al., 2015)</td>
<td>Mucosal thickening and polyps, TM thickened, mucopurulent OM</td>
<td>Special stains for bacteria and fungi negative</td>
<td></td>
</tr>
<tr>
<td>Dbh +/-</td>
<td>Absence of measurable epinephrine / norepinephrine, altered metabolism, loss adrenergic modulation auditory tube function</td>
<td>(Maison et al., 2010)</td>
<td>Exudate or fibrous tissue in the vicinity of the round window, thickening</td>
<td>Normal</td>
<td>Physiological changes susceptibility to infection via compromised local and systemic</td>
</tr>
<tr>
<td>Isl1Drsh/+</td>
<td>(Hilton et al., 2011)</td>
<td>thermoregulation, cardiovascular tone, maternal behaviour. Deficits in motor function, learning &amp; memory</td>
<td>round window</td>
<td>mucosa thickened by granulation tissue, mucosa and tympanic membrane vascularised, fusion of malleus and incus</td>
<td>Proteus sp.</td>
</tr>
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</tbody>
</table>

**Miscellaneous**

<table>
<thead>
<tr>
<th>Nf2 KO</th>
<th>(Giovannini et al., 2000)</th>
<th>neurofibromatosis</th>
<th>osteosarcoma, molar eruption delayed or absent, growth retardation, reduced survival</th>
<th>otitis media present but not described in detail</th>
<th></th>
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</tr>
</thead>
</table>
Oxgr1 -/-

(Kerschner et al., 2013)

mucosal fibrosis, haemosiderin laden macrophages, mineralization of ossicles

Gram stains for bacteria & PCR for human otopathogens H. influe, S.pneumo, M. catarr are negative

upregulation of Muc5B, Muc 19

References for table S1


