

ANO1 in Cystic Fibrosis

Subjects: **Biochemistry & Molecular Biology**

Contributor: Tabary Olivier , Christie MITRI

Cystic fibrosis (CF) is the most common of rare hereditary diseases in Caucasians, and it is estimated to affect 75,000 patients globally. CF is a complex disease due to the multiplicity of mutations found in the CF transmembrane conductance regulator (CFTR) gene causing the CFTR protein to become dysfunctional. Although CFTR is the main chloride channel in the lungs, others could, e.g., anoctamin-1 (ANO1 or TMEM16A), compensate for the deficiency of CFTR.

cystic fibrosis

anoctamin-1

calcium-activated chloride channel

CFTR-independent therapy

1. Introduction

Cystic fibrosis (CF), an autosomal recessive genetic multiorgan disease, is caused by an absent or dysfunctional CF transmembrane conductance regulator (CFTR) channel that mainly mediates chloride anion transport across the apical membrane of epithelial cells. On the pulmonary level, CF leads to persistent pulmonary infections, chronic inflammation, and mucus plugging in the airways, causing irreversible lung damage. To date, more than 2000 different mutations in the CFTR gene have been identified [1]. A classification system groups mutations into six classes according to the functional consequences they generate on the CFTR protein: (1) class I: no functional CFTR protein; (2) class II: CFTR trafficking defects; (3) class III: defective channel gating; (4) class IV: decreased channel conductance; (5) class V: reduced synthesis of CFTR; (6) class VI: decreased CFTR stability at plasma membrane. Today, many symptomatic therapeutics (antibiotics, mucus thinners, bronchodilators, supplements to prevent malnutrition, etc.) are available to treat patients with CF, which has lengthened their life expectancy from 5 years in 1960 to over 50 years. New curative treatments aimed at rescuing CFTR functionality have emerged. There are four FDA-approved CFTR modulators developed by Vertex Pharmaceuticals (Kalydeco®, Orkambi®, Symdeko®, and the latest Trikafta®) [2], although the proven efficacy of these correctors and potentiators is limited only to particular mutations. There are still 15% of patients without any CFTR-directed therapeutics. Hence, there is an interest in finding an alternative strategy to treat patients with CF independently of CFTR mutations. Alternative ion channels have been suggested to bypass CFTR dysfunction [3][4], such as ENaC [5], the solute carrier 26A9 (SLC26A9) [6], and calcium-activated chloride channel (CaCCs), including anoctamin-1 (ANO1 or TMEM16A) [4]. Such approaches might be efficient therapies for all patients, regardless of their CF mutations.

2. Anoctamin-1

CaCCs were described for the first time in 1981 in *Rana pipiens* eggs [7] and then in *Xenopus laevis* oocytes [8]. Today, CaCCs have been identified in many cellular types (neurons, epithelial cells, smooth muscle cells,

pancreatic cells, etc.) and have been shown to play essential roles in cellular functions [9][10][11]. Among CaCCs is the anoctamin family (ANO for anion channel and OCTA for their eight predicted transmembrane domains or TMEM16), consisting of 10 proteins (ANO1–10). ANOs might play an essential role in development due to their temporal and spatial differential expression in many developing tissues. We can separate ANOs into two groups: ANO1-2 and the rest. However, ANO2 has different biophysical characteristics to ANO1 and its expression is limited to the vomeronasal epithelium [12][13]. In 2008, ANO1 was identified as a CaCC by three independent research groups [14][15][16].

3. ANO1 in Cystic Fibrosis

3.1. Overview

A study showed that the activity of CaCCs, in general, was increased in the nasal epithelium *in vivo* of patients with CF [17]. In 1986, another team demonstrated the presence of chloride current at the apical membrane of epithelial cells of CF in the presence of ionized calcium [18]. Moreover, another study showed a decrease in ATP-induced chloride efflux in the primary bronchial epithelial cell line [19]. The identification of ANO1 as a CaCC later on and its involvement in many deregulated processes in patients with CF made it a real therapeutic target. In the last few years, more research has been dedicated to ANO1 in CF.

Recent data reported the absence of calcium-induced chloride currents in epithelial cells of ANO1 KO mice [20]. A subsequent study showed that CFTR expression requires ANO1 in plasma membranes, indicating a close relationship between the two chloride channels [21]. Moreover, ANO1 expression and chloride activity were decreased in CF [22]. According to several observations ((1) ANO1's absence decreases airway secretion, (2) ANO1 is ubiquitously expressed in all the tissue affected by CF, including airway epithelial cells where ANO1 is a secondary chloride channel, (3) ANO1 provides a chloride pathway that is CFTR-independent, and (4) ANO1 is involved in HCO_3^- secretion, which is highly important for fluid secretion and mucus hydration [23]), increasing ANO1 activity can probably compensate for CFTR deficiency (**Figure 1**). Different approaches targeting ANO1 have been developed to bypass CFTR dysfunction.

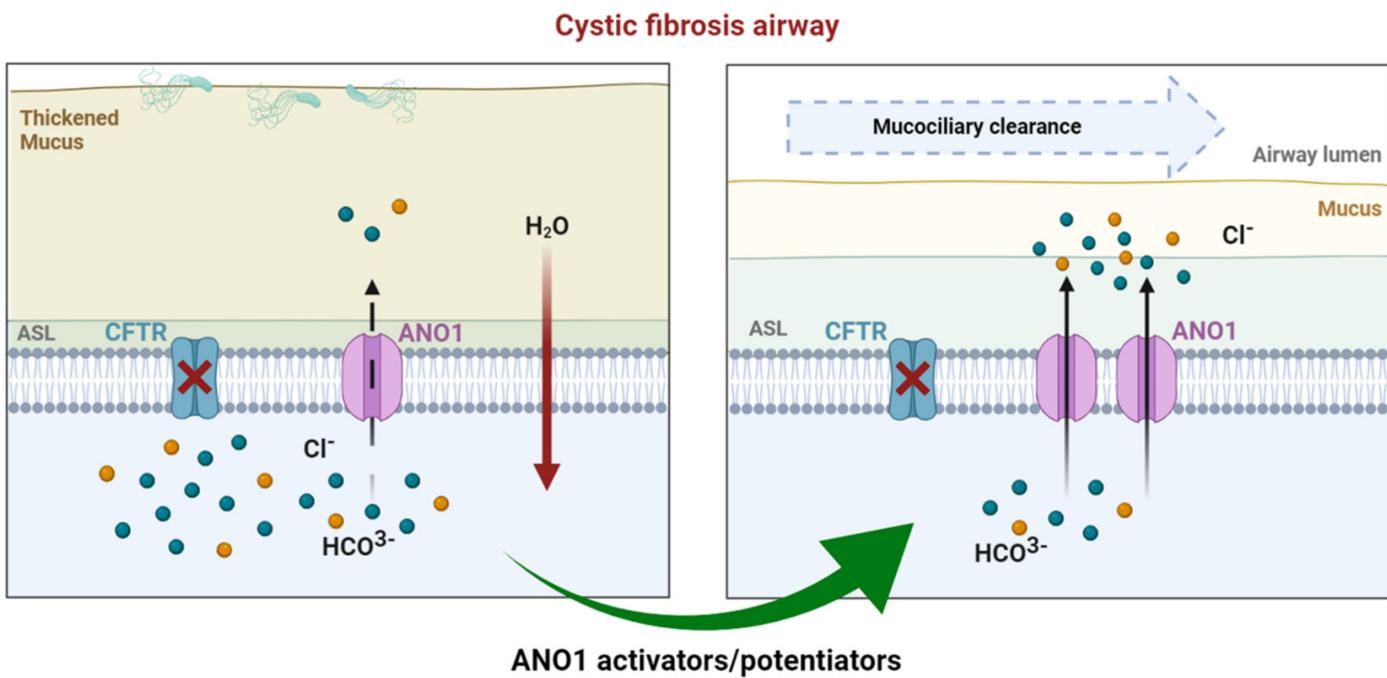


Figure 1. In CF airways, dysfunctional CFTR leads to compromised chloride efflux. Sodium entry is upregulated, leading to a dehydrated air surface liquid (ASL) and impaired mucociliary clearance favoring mucostasis, causing chronic inflammation and infection. In healthy airways, ANO1 is colocalized with CFTR within the apical membrane of epithelial cells, contributing to ion and water homeostasis. In CF ciliated cells, the expression of ANO1 is also diminished. Modulating ANO1, as an alternative CF therapy, could compensate for defective CFTR and, thus, enhance fluid secretion by ciliated epithelial cells, regulating ASL height and pH.

3.2. Drug Approaches Targeting ANO1 in Cystic Fibrosis

Many molecules have been used to modulate ANO1's activity, but few were shown to be efficient and, most importantly, specific to ANO1 (**Table 1**).

Table 1. Summary of ANO1 inhibitors and activators used in CF.

Inhibitors	Specificity	Assay	References
ANI9	Not specific	In vitro	[24][25]
CCinh-A01	Not specific	In vitro, in vivo	[26]

Inhibitors	Specificity	Assay	References
DIDS	Not specific	In vitro	[27]
Diphenylamine-2-carboxylate (DPC), 5-nitro-2-(3-phenylpropylamino) benzoic acid	Not specific	In vitro	[28]
Flufenamic acid	Not specific	In vitro	[29] [30]
Niclosamide	Not specific	In vitro, in vivo	[31]
Niflumic acid	Not specific	In vitro, in vivo	[32]
Plumbagin	Not specific	In vitro	[33]
Quercetin	Not specific	In vitro, in vivo, clinical trial (phase II)	[34] [35] [36]
Tannic acid	Not specific	In vitro	[37]
T16ainh-A01	Specific	In vitro	[26]
Activators			
Denufosal (INS37217)	Not specific	In vitro, in vivo, clinical trial (phase III failed)	[38] [39] [40] [41] [42] [43] [44]

Inhibitors	Specificity	Assay	References
Duramycine (MOLI1901)	Not specific	In vitro, in vivo, clinical trial (phase II failed)	[45]
Eact	Not specific	In vitro	[26]
ETD002 (or ETX001)	Specific	In vitro, in vivo, clinical trial (phase I)	[46]
Interleukin 4	Not specific	In vitro	[14]
[38][39][40][41]			ok place. therapy, s and led
Resveratrol	Not specific	In vitro, in vivo, clinical trial	[47][48][49]
TSB ANO1	Specific	In vitro, in vivo, preclinical	[50]

might have led to an increase in mucus in the airways. Therefore, CaCC activators need to target CaCCs directly without elevating cytoplasmic calcium for more targeted therapy and efficacy [42].

ANO1 identification paved the way for the development of specific activator molecules which would, without modifying the calcium signaling, obtain a more sustained activation over time, leading to better efficiency.

Another novel drug is an ANO1 potentiator (ETD002) developed by Enterprise Therapeutics (based in the University of Sussex Innovation Center, UK) and acquired by Roche (Genentech) in October 2020. This inhaled molecule demonstrated an upregulation of ANO1, which boosts epithelial fluid secretion and mucus clearance in primary CF bronchial epithelial cells and ovine models. Unlike denufosal, intracellular calcium measurements checked that ETD002 did not affect calcium mobilization, coherent with a direct effect on ANO1. A phase I study to test the safety of ETD002 in healthy participants is in progress [24]. The mechanism via which ETD002 potentiates ANO1 activity is still unclear.

Our team has also worked on an innovative alternative approach using a locked nucleic acid (LNA)-enhanced antisense oligonucleotide (ASO). A previous study demonstrated that microRNA (miR-9) contributes to the downregulation of ANO1 expression and activity by directly targeting its 3'UTR. ASO ANO1 binds to the 3'UTR target site of ANO1 mRNA, preventing miR-9 from gaining access to that site [50]. ASO ANO1 has increased ANO1 expression, chloride activity, and mucus clearance in primary human CF cells and CF mice. Recent studies have

suggested that ANO1 plays an essential role in mucus production [51]. To date, we have not observed any significant increase in mucin 5AC (MUC5AC, the main component of respiratory mucus produced by goblet cells) or mucin 5B (MUC5B, gel-forming mucin that plays a key role in mucociliary clearance) expression in vitro.

Currently, ANO1 activation as a therapeutic target is subject to controversial opinions. Centeio et al. found an upregulation of ANO1 expression in submucosal glands, airway smooth muscles, and pulmonary blood vessels in CF and asthmatic inflamed lungs [52]. Moreover, activating ANO1 with Eact, which also activates other channels, induced mucus production in airway goblet cells and bronchoconstriction in ovalbumin-sensitized asthmatic mice [25], whereby activating ANO1 could worsen the pathology in inflammatory airway diseases. Instead, the authors demonstrated that ANO1 inhibition by niclosamide significantly reduced goblet cell metaplasia and mucus production in asthmatic mice, as well as inhibited MUC5AC expression in Calu-3 human submucosal cells, suggesting that ANO1 inhibition might be beneficial in inflammatory airway diseases [52]. It is also important to note that a transcriptome meta-analysis revealed that CF and asthma pathways are highly divergent [53]. Furthermore, the same group reported a defect of mucus secretion and accumulation in secretory cells in 2018 when ANO1 was knocked out in ciliated airway epithelial cells and intestinal goblet cells, highlighting the vital role of ANO1 in mucus secretion [51]. Another study showed that ANO1 inhibitors reduced both mucus secretion and airway hyperactivity [54].

On the other hand, using a human respiratory basal cell line (BCi-NS1.1), Simões et al. showed that MUC5AC production does not require ANO1, and their simultaneous upregulation is only circumstantial under cell proliferation [55]. The authors also showed a decrease in ASL when inhibiting ANO1. Furthermore, while replying to Olschewski et al.'s concerns on increasing ANO1 activity, Danahay et al. declared that positive modulation of ANO1 induces no bronchospasm in the conscious sheep model nor affects vascular smooth muscle contraction (unpublished observations) [56][57]. Overall, ANO1's possible role in mucus production remains obscure and evokes controversial opinions over the beneficial or deleterious results of stimulating the channel in CF.

References

1. Cutting, G.R. Cystic fibrosis genetics: From molecular understanding to clinical application. *Nat. Rev. Genet.* 2015, 16, 45–56.
2. Lopes-Pacheco, M. CFTR Modulators: The Changing Face of Cystic Fibrosis in the Era of Precision Medicine. *Front. Pharmacol.* 2019, 10, 1662.
3. Li, H.; Salomon, J.J.; Sheppard, D.N.; Mall, M.A.; Galietta, L.J. Bypassing CFTR dysfunction in cystic fibrosis with alternative pathways for anion transport. *Curr. Opin. Pharmacol.* 2017, 34, 91–97.
4. Mall, M.A.; Galietta, L.J. Targeting ion channels in cystic fibrosis. *J. Cyst. Fibros.* 2015, 1, 561–570.

5. Moore, P.J.; Tarhan, R. The epithelial sodium channel (ENaC) as a therapeutic target for cystic fibrosis lung disease. *Expert Opin. Ther. Targets* 2018, 22, 687–701.
6. Balazs, A.; Mall, M.A. Mucus obstruction and inflammation in early cystic fibrosis lung disease: Emerging role of the IL-1 signaling pathway. *Pediatr. Pulmonol.* 2019, 54 (Suppl. 3), S5–S12.
7. Cross, N.L. Initiation of the activation potential by an increase in intracellular calcium in eggs of the frog, *Rana pipiens*. *Dev. Biol.* 1981, 85, 380–384.
8. Miledi, R. A calcium-dependent transient outward current in *Xenopus laevis* oocytes. *Proc. R. Soc. Lond. B Biol. Sci.* 1982, 215, 491–497.
9. Frings, S.D.; Reuter, B.; Kleene, S.J. Neuronal Ca²⁺-activated Cl-channels—Homing in on an elusive channel species. *Prog. Neurobiol.* 2000, 60, 247–289.
10. Leblanc, N.; Ledoux, J.; Saleh, S.; Sanguinetti, A.; Angermann, J.; O'Driscoll, K.; Britton, F.; Perrino, B.A.; Greenwood, I.A. Regulation of calcium-activated chloride channels in smooth muscle cells: A complex picture is emerging. *Can. J. Physiol. Pharmacol.* 2005, 83, 541–556.
11. Kidd, J.F.; Thorn, P. Intracellular Ca²⁺ and Cl-channel activation in secretory cells. *Annu. Rev. Physiol.* 2000, 62, 493–513.
12. Kunzelmann, K.; Tian, Y.; Martins, J.R.; Faria, D.; Kongsuphol, P.; Ousingsawat, J.; Thevenod, F.; Roussa, E.; Rock, J.; Schreiber, R. Anoctamins. *Pflug. Arch.* 2011, 462, 195–208.
13. Dibattista, M.; Amjad, A.; Maurya, D.K.; Sagheddu, C.; Montani, G.; Tirindelli, R.; Menini, A. Calcium-activated chloride channels in the apical region of mouse vomeronasal sensory neurons. *J. Gen. Physiol.* 2012, 140, 3–15.
14. Caputo, A.; Caci, E.; Ferrera, L.; Pedemonte, N.; Barsanti, C.; Sondo, E.; Pfeffer, U.; Ravazzolo, R.; Zegarra-Moran, O.; Galietta, L.J. TMEM16A, a membrane protein associated with calcium-dependent chloride channel activity. *Science* 2008, 322, 590–594.
15. Schroeder, B.C.; Cheng, T.; Jan, Y.N.; Jan, L.Y. Expression cloning of TMEM16A as a calcium-activated chloride channel subunit. *Cell* 2008, 134, 1019–1029.
16. Yang, Y.D.; Cho, H.; Koo, J.Y.; Tak, M.H.; Cho, Y.; Shim, W.S.; Park, S.P.; Lee, J.; Lee, B.; Kim, B.M.; et al. TMEM16A confers receptor-activated calcium-dependent chloride conductance. *Nature* 2008, 455, 1210–1215.
17. Knowles, M.R.; Clarke, L.L.; Boucher, R.C. Activation by extracellular nucleotides of chloride secretion in the airway epithelia of patients with cystic fibrosis. *N. Engl. J. Med.* 1991, 325, 533–538.
18. Frizzell, R.A.; Rechkemmer, G.; Shoemaker, R.L. Altered regulation of airway epithelial cell chloride channels in cystic fibrosis. *Science* 1986, 233, 558–560.

19. Blouquit, S.; Regnier, A.; Dannhoffer, L.; Fermanian, C.; Naline, E.; Boucher, R.; Chinet, T. Ion and fluid transport properties of small airways in cystic fibrosis. *Am. J. Respir. Crit. Care Med.* 2006, 174, 299–305.

20. Benedetto, R.; Ousingsawat, J.; Wanitchakool, P.; Zhang, Y.; Holtzman, M.J.; Amaral, M.; Rock, J.R.; Schreiber, R.; Kunzelmann, K. Epithelial Chloride Transport by CFTR Requires TMEM16A. *Sci. Rep.* 2017, 7, 12397.

21. Benedetto, R.; Centeio, R.; Ousingsawat, J.; Schreiber, R.; Janda, M.; Kunzelmann, K. Transport properties in CFTR-/- knockout piglets suggest normal airway surface liquid pH and enhanced amiloride-sensitive Na(+) absorption. *Pflugers Arch.* 2020, 472, 1507–1519.

22. Ruffin, M.; Voland, M.; Marie, S.; Bonora, M.; Blanchard, E.; Blouquit-Laye, S.; Naline, E.; Puyo, P.; Le Rouzic, P.; Guillot, L.; et al. Anoctamin 1 dysregulation alters bronchial epithelial repair in cystic fibrosis. *Biochim. Biophys. Acta* 2013, 1832, 2340–2351.

23. Quinton, P.M. Role of epithelial HCO3(–) transport in mucin secretion: Lessons from cystic fibrosis. *Am. J. Physiol. Cell Physiol.* 2010, 299, C1222–C1233.

24. Danahay, H.; Gosling, M. TMEM16A: An Alternative Approach to Restoring Airway Anion Secretion in Cystic Fibrosis? *Int. J. Mol. Sci.* 2020, 21, 2386.

25. Ramu, Y.; Xu, Y.; Shin, H.G.; Lu, Z. Counteracting suppression of CFTR and voltage-gated K⁺ channels by a bacterial pathogenic factor with the natural product tannic acid. *eLife* 2014, 3, e03683.

26. Genovese, M.; Borrelli, A.; Venturini, A.; Guidone, D.; Caci, E.; Viscido, G.; Gambardella, G.; di Bernardo, D.; Scudieri, P.; Galletta, L.J.V. TRPV4 and purinergic receptor signalling pathways are separately linked in airway epithelia to CFTR and TMEM16A chloride channels. *J. Physiol.* 2019, 597, 5859–5878.

27. Gianotti, A.; Ferrera, L.; Philp, A.R.; Caci, E.; Zegarra-Moran, O.; Galletta, L.J.; Flores, C.A. Pharmacological analysis of epithelial chloride secretion mechanisms in adult murine airways. *Eur. J. Pharmacol.* 2016, 781, 100–108.

28. Danko, T.; Hargitai, D.; Pataki, A.; Hakim, H.; Molnar, M.; Zsembery, A. Extracellular alkalinization stimulates calcium-activated chloride conductance in cystic fibrosis human airway epithelial cells. *Cell Physiol. Biochem.* 2011, 27, 401–410.

29. McCarty, N.A.; McDonough, S.; Cohen, B.N.; Riordan, J.R.; Davidson, N.; Lester, H.A. Voltage-dependent block of the cystic fibrosis transmembrane conductance regulator Cl⁻ channel by two closely related arylaminobenzoates. *J. Gen. Physiol.* 1993, 102, 1–23.

30. Reddy, M.M.; Quinton, P.M. Effect of anion transport blockers on CFTR in the human sweat duct. *J. Membr. Biol.* 2002, 189, 15–25.

31. Fischer, H.; Illek, B.; Sachs, L.; Finkbeiner, W.E.; Widdicombe, J.H. CFTR and calcium-activated chloride channels in primary cultures of human airway gland cells of serous or mucous phenotype. *Am. J. Physiol. Lung Cell Mol. Physiol.* 2010, 299, L585–L594.

32. Cabrita, I.; Benedetto, R.; Schreiber, R.; Kunzelmann, K. Niclosamide repurposed for the treatment of inflammatory airway disease. *JCI Insight.* 2019, 4, e128414.

33. Scott-Ward, T.S.; Li, H.; Schmidt, A.; Cai, Z.; Sheppard, D.N. Direct block of the cystic fibrosis transmembrane conductance regulator Cl(-) channel by niflumic acid. *Mol. Membr. Biol.* 2004, 21, 27–38.

34. Yu, B.; Zhu, X.; Yang, X.; Jin, L.; Xu, J.; Ma, T.; Yang, H. Plumbagin Prevents Secretory Diarrhea by Inhibiting CaCC and CFTR Channel Activities. *Front. Pharmacol.* 2019, 10, 1181.

35. Pyle, L.C.; Fulton, J.C.; Sloane, P.A.; Backer, K.; Mazur, M.; Prasain, J.; Barnes, S.; Clancy, J.P.; Rowe, S.M. Activation of the Cystic Fibrosis Transmembrane Conductance Regulator by the Flavonoid Quercetin. *Am. J. Respir Cell Mol. Biol.* 2010, 43, 607–616.

36. Zhang, S.; Smith, N.; Schuster, D.; Azbell, C.; Sorscher, E.J.; Rowe, S.M.; Woodworth, B.A. Quercetin increases cystic fibrosis transmembrane conductance regulator-mediated chloride transport and ciliary beat frequency: Therapeutic implications for chronic rhinosinusitis. *Am. J. Rhinol. Allergy* 2011, 25, 307–312.

37. Nasal Potential Studies Utilizing Cystic Fibrosis Transmembrane Regulator (CFTR) Modulators. Available online: <https://ClinicalTrials.gov/show/NCT01348204> (accessed on 1 October 2021).

38. Deterding, R.; Retsch-Bogart, G.; Milgram, L.; Gibson, R.; Daines, C.; Zeitlin, P.L.; Milla, C.; Marshall, B.; Lavange, L.; Engels, J.; et al. Safety and tolerability of denufosal tetrasodium inhalation solution, a novel P2Y2 receptor agonist: Results of a phase 1/phase 2 multicenter study in mild to moderate cystic fibrosis. *Pediatr. Pulmonol.* 2005, 39, 339–348.

39. Deterding, R.R.; Lavange, L.M.; Engels, J.M.; Mathews, D.W.; Coquillette, S.J.; Brody, A.S.; Millard, S.P.; Ramsey, B.W. Phase 2 randomized safety and efficacy trial of nebulized denufosal tetrasodium in cystic fibrosis. *Am. J. Respir. Crit. Care Med.* 2007, 176, 362–369.

40. Goss, C.H.; McKone, E.F.; Mathews, D.; Kerr, D.; Wanger, J.S.; Millard, S.P. Experience using centralized spirometry in the phase 2 randomized, placebo-controlled, double-blind trial of denufosal in patients with mild to moderate cystic fibrosis. *J. Cyst. Fibros.* 2008, 7, 147–153.

41. Kellerman, D.; Rossi Mospan, A.; Engels, J.; Schaberg, A.; Gorden, J.; Smiley, L. Denufosal: A review of studies with inhaled P2Y(2) agonists that led to Phase 3. *Pulm. Pharmacol. Ther.* 2008, 21, 600–607.

42. Moss, R.B. Pitfalls of drug development: Lessons learned from trials of denufosal in cystic fibrosis. *J. Pediatr.* 2013, 162, 676–680.

43. Accurso, F.J.; Moss, R.B.; Wilmott, R.W.; Anbar, R.D.; Schaberg, A.E.; Durham, T.A.; Ramsey, B.W.; T.-I.S. Group. Denufosal tetrasodium in patients with cystic fibrosis and normal to mildly impaired lung function. *Am. J. Respir. Crit. Care Med.* 2011, 183, 627–634.

44. Ratjen, F.; Durham, T.; Navratil, T.; Schaberg, A.; Accurso, F.J.; Wainwright, C.; Barnes, M.; Moss, R.B.; T.-S.I. Group. Long term effects of denufosal tetrasodium in patients with cystic fibrosis. *J. Cyst. Fibros.* 2012, 11, 539–549.

45. Lancovutide (Moli1901) Inhalation Solution Study in Adolescents and Adults with Cystic Fibrosis. Available online: <https://ClinicalTrials.gov/show/NCT00671736> (accessed on 1 October 2021).

46. Danahay, H.L.; Lilley, S.; Fox, R.; Charlton, H.; Sabater, J.; Button, B.; McCarthy, C.; Collingwood, S.P.; Gosling, M. TMEM16A Potentiation: A Novel Therapeutic Approach for the Treatment of Cystic Fibrosis. *Am. J. Respir. Crit. Care Med.* 2020, 201, 946–954.

47. Lu, B.D.; Corey, A.; Kelley, T.J. Resveratrol restores intracellular transport in cystic fibrosis epithelial cells. *Cells* 2020, 318, L1145–L1157.

48. Non-pulmonary Contributors of Exercise Intolerance in Patients with Cystic Fibrosis. Available online: <https://ClinicalTrials.gov/show/NCT04166396> (accessed on 1 October 2021).

49. Mechanisms for Vascular Dysfunction and Exercise Tolerance in CF. Available online: <https://ClinicalTrials.gov/show/NCT02690064> (accessed on 1 October 2021).

50. Sonneville, F.; Ruffin, M.; Coraux, C.; Rousselet, N.; Le Rouzic, P.; Blouquit-Laye, S.; Corvol, H.; Tabary, O. MicroRNA-9 downregulates the ANO1 chloride channel and contributes to cystic fibrosis lung pathology. *Nat. Commun.* 2017, 8, 710.

51. Benedetto, R.; Cabrita, I.; Schreiber, R.; Kunzelmann, K. TMEM16A is indispensable for basal mucus secretion in airways and intestine. *FASEB J.* 2019, 33, 4502–4512.

52. Centeio, R.; Ousingsawat, J.; Cabrita, I.; Schreiber, R.; Talbi, K.; Benedetto, R.; Dousova, T.; Verbeken, E.K.; De Boeck, K.; Cohen, I.; et al. Mucus Release and Airway Constriction by TMEM16A May Worsen Pathology in Inflammatory Lung Disease. *Int. J. Mol. Sci.* 2021, 22, 2888.

53. Clarke, L.A.; Botelho, H.M.; Sousa, L.; Falcao, A.O.; Amaral, M.D. Transcriptome meta-analysis reveals common differential and global gene expression profiles in cystic fibrosis and other respiratory disorders and identifies CFTR regulators. *Genomics* 2015, 106, 268–277.

54. Huang, F.; Zhang, H.; Wu, M.; Yang, H.; Kudo, M.; Peters, C.J.; Woodruff, P.G.; Solberg, O.D.; Donne, M.L.; Huang, X.; et al. Calcium-activated chloride channel TMEM16A modulates mucin secretion and airway smooth muscle contraction. *Proc. Natl. Acad. Sci. USA* 2012, 109, 16354–16359.

55. Simoes, F.B.; Quaresma, M.C.; Clarke, L.A.; Silva, I.A.; Pankonien, I.; Railean, V.; Kmit, A.; Amaral, M.D. TMEM16A chloride channel does not drive mucus production. *Life Sci. Alliance*

2019, 2, e201900462.

56. Danahay, H.L.; Morris, D.G.; Gosling, M. Reply to Olschewski et al.: TMEM16A Potentiation: Possible Drawbacks. *Am. J. Respir Crit. Care Med.* 2020, 202, 905–906.
57. Olschewski, A.; Nagaraj, C.; Zabini, D.; Nagy, B.M.; Kwapiszewska, G.; Olschewski, H. TMEM16A Potentiation: Possible Drawbacks. *Am. J. Respir Crit. Care Med.* 2020, 202, 904–905.

Retrieved from <https://encyclopedia.pub/entry/history/show/36372>