Cystic fibrosis and estrogens: a perfect storm

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Irreversible destruction and widening of the airways due to acquired infections or genetic mutations as well as those of unknown cause are more severe in females. Differences between male and female anatomy, behavior, and hormonal state have been proposed to explain the increased incidence and severity in females with airway disease such as cystic fibrosis (CF); however, a mechanism to explain a sex-related difference has remained elusive. In this issue of the JCI, Coakley et al. report that elevations in the major estrogen hormone in humans — 17β-estradiol — reduce Ca2+-activated Cl− secretion by airway epithelial cells in culture, thereby disrupting ion and water balance (see the related article, doi:10.1172/JCI33893). They measure a similar diminution of nasal epithelial Ca2+-activated Cl− secretion in women with CF during the menstrual cycle phase at which 17β-estradiol level is at its highest. These data suggest that for about one week of a four-week menstrual cycle, women with CF will have a reduced ability to efficiently clear airway secretions, the buildup of which is a hallmark of CF. The authors suggest that these data warrant the testing of antiestrogen therapy in females with CF and propose an alternative avenue for CF therapeutic development.

Does female sex impose genetic, hormonal, and/or behavioral constraints on lung function, including a predilection to the airway destruction and widening known as bronchiectasis? In patients with cystic fibrosis (CF), an autosomal recessive inherited disorder resulting from mutations in the CFTR that very often is associated with bronchiectasis, CFTR genotype, acquisition of airway pathogens, and environmental factors conspire to adversely affect disease outcome and survival. However, women in whom no specific genetic disease has been identified have long been recognized as being more vulnerable than men to inflammatory disorders. For example, Mycobacterium avium intracellulare (MAI) pulmonary infection (1), asthma, inflammatory bowel disease, sarcoidosis, Sjögren syndrome, and rheumatoid arthritis are predominant in women (2). Idiopathic bronchiectasis behaves differently in women than it does in men with regard to locale, incidence,coinfecting pathogens (such as MAI), and etiology. Hormonal effects have long been suspected to contribute to periodic (cata menial; exacerbated at the time of menstruation) pneumothoraces in women (3). In an effort to explain these observations, estrogens have been intensely studied.

Bronchiectasis is defined as a progressive abnormality of conducting bronchial airways that results in dilatation, thinning of capillary walls, and impaired mucociliary clearance (clearance of mucus by airway epithelial cell cilia). Whether the inciting event is an infection that is poorly controlled, an impairment of the immune system, or a local anatomic variant leading to inadequate mucociliary clearance, it is often recognized long after it has occurred. Treatment of infection and augmentation of mucociliary clearance can stabilize disease, but these approaches are not always effective, and certain pathogens are notorious for being difficult to eradicate. Superimpose the above triggers on a person with CF, and you have a perfect storm.

**CF results from dysfunction in a cAMP-regulated Cl− channel, the CFTR**

CF is a systemic disorder that develops in the gastrointestinal tract prior to birth but does not manifest in the lungs until after birth. This delay in onset of airway obstruction opens a therapeutic window. CFTR is critically important to the airways and sinuses because it acts as a central regulator of periciliary ion and water content. Reducing or eliminating CFTR at the apical membrane of airway epithelial cells through genetic mutation, necrotizing infections, or experimentally by siRNA disrupts cAMP-mediated Cl− secretion and allows excessive epithelial Na+ channel–mediated (ENaC-mediated) Na+ reabsorption (Figure 1). The net result is depleted airway surface liquid depth, poor ciliary function, impaired mucociliary clearance, and increased bacterial infections. It has long been hypothesized that parallel, non-CFTR–mediated Cl− conductance pathways (Figure 1) might serve in a redundant capacity to carry Cl− and promote fluid secretion when CFTR is absent. A proposed member of an alternative pathway of Cl− conductance is the Ca2+-activated Cl− channel (CaCC) (4–6), which is the focus of the study reported by Coakley et al. (7) in this issue of the JCI. Unlike the outwardly rectifying Cl− channel (ORCC), which depends on CFTR to function (8), the CaCC is independent of CFTR and responsive to intracellular Ca2+ concentration. The hypothesis put forward by Coakley et al. is that the sensitivity of the CaCC is inversely proportional to the level of circulating 17β-estradiol, and as a result, higher 17β-estradiol levels adversely interfere with CaCC-mediated Cl− transport across the surface of airway epithelial cells (Figure 1).

The results of previously reported experiments performed at relatively high 17β-estradiol concentrations suggest that there is precedent for suggesting that estrogens and related molecules would have an impact on airway ion transport. The most common CF-associated mutation is the deletion of phenylalanine at residue 508 in CFTR (ΔF508 CFTR), and 17β-estradiol at nM concentrations has been shown to rescue ΔF508 CFTR from proteasomal degradation and increase CFTR channel activity (9). These authors identified the 17β-estradiol target as Na+/H+ exchanger–regulator factor 1 (NHERF1). Raising 17β-estradiol levels in the medium increased the levels of NHERF1, which facilitated the trafficking of mutant CFTR to the epithelial cell surface. Another group has studied CFTR expression in a rat model of human ovarian hyperstimulation syndrome (OHSS) (10),

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**Nonstandard abbreviations used:** CaCC, Ca2+-activated Cl− channel; CF, cystic fibrosis; ENaC, epithelial Na+ channel; FEV1, forced expiratory volume in 1 second; NPD, nasal potential difference; UTP, uridine triphosphate.

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OHSS occurs as a complication of assisted reproduction treatments that stimulate the ovaries. Using RT-PCR, Western blotting, and electrophysiologic techniques, this group demonstrated that CFTR expression is upregulated in this syndrome. Further-more, estrogen but not progesterone stimulated cAMP-mediated Cl⁻ secretion. In both CF and non-CF airway epithelial cells, P2Y₁ receptors coexist in the apical membrane and are stimulated by ATP or other purinergic agonists to initiate increased intracellular Ca²⁺. Ca²⁺ signaling stimulates Cl⁻ secretion through the CaCC pathway. In this issue of the JCI, Coakley et al. (7) show that 17β-estradiol intersects with this pathway. Higher 17β-estradiol levels are sensed through estrogen receptors, and the authors show that Ca²⁺-activated Cl⁻ secretion is decreased in women with CF at times when 17β-estradiol levels are high. The net result in CF is a worsening degree of airway surface dehydration and decreased mucociliary clearance.
A surrogate marker of airway epithelial ion transport

The study reported by Coakley et al. (7) in this issue of the JCI employed the in vivo nasal potential difference (NPD) assay to measure the activity of Cl− channel pathways. NPD is a surrogate biomarker of Na+ and Cl− transport in the lower airways (14, 15). NPD measurements of Na+ reabsorption and the activity of various agonist-activated Cl− secretory pathways have informed therapeutic development of gene therapy vectors (15), Na+ channel inhibitors (16), alternative Cl− channel activators (17), and CFTR repair molecules (18). The test is performed in the clinic and requires the subject to be free of recent nasal steroids, upper respiratory infection, or topical adrenergic agents. Using this assay, it is possible to quantify the activity of the ENaC and several different Cl− conductance pathways, including CFTR.

Coakley et al. (7) recorded the NPD in non-CF and CF women during different phases of the menstrual cycle, during which estrogen levels naturally rise and fall. They modified the standardized NPD protocol, which is designed to measure CFTR through isoproterenol-induced increases in cAMP, by adding a perfusion with uridine triphosphate (UTP), an agonist of the G protein–coupled receptor P2Y2. UTP leads to increased intracellular Ca2+ concentration and subse- quent Ca2+-activated Cl− secretion. The authors detected as much as a 50% reduction in the UTP-stimulated NPD during the phase of the menstrual cycle when 17β-estradiol levels were at their highest in CF and non-CF women. But they did not observe an increase in the isoproterenol-mediated Cl− transport portion of the NPD in non-CF or CF female subjects during the periods of the menstrual cycle in which 17β-estradiol levels were high. The most likely explanation for this is that endogenous 17β-estradiol levels are not high enough to stimulate CFTR expression and function. Another possibility is that the variability of the NPD limits the ability to detect modest increases in nasal epithelial CFTR expression. Yet the NPD has been sensitive enough to show significant differences in the amiloride-insensitive Na+ potential (19) and now the UTP-activated Cl− secretory pathway. To further understand the effects of 17β-estradiol on ion conductance pathways, Coakley et al. (7) studied the effects of 17β-estradiol in human CF and non-CF airway epithelial cell cultures in vitro. They did not find a 17β-estradiol–mediated change in estrogen receptor activity. In addition, instead of a direct effect of 17β-estradiol on Ca2+-activated Cl− secretion, the study implicated upstream targets such as Ca2+ signaling rather than the CaCC itself. Estrogens have been studied in the context of ENaC activity, and Sweezy et al. (19) have observed stimulation of amiloride-insensitive Na+ potential difference, which, if experienced during the Coakley et al. study, would have been expected to lead to further polarization of the baseline NPD. In an earlier study, serum levels of progesterone and estrogen were measured during the menstrual cycles of 7 women with CF, and NPD values were simultaneously recorded. In this small study, estrogen varied between approximately 100 pM and 300 pM (well below the concentrations used in vitro in most studies) and progesterone varied from approximately 1–40 mM (19). Most studies use exogenous estrogens at very high levels, and the results under these artificial conditions may not extrapolate well to native conditions.

Importance of the balance between Na+ and Cl− transport in regulation of airway surface liquid homeostasis

Which kind of ion channel is most important for periciliary ion and water content—CFTR, CaCC, or ENaC? This debate continues to rage. CFTR-null mice do not develop lung disease without a superimposing infection, yet mice overexpressing ENaC immediately suffer from airway inflammation (20). Unfortunately, the CaCC has not been convincingly cloned, although a candidate subunit has been identified (4–6), and knockout of the CaCC cannot be studied. Clearly, autosomal recessive classic CF is most serious for humans, followed by ENaC gain-of-function disease (ENaC is a product of the non–voltage-gated 1γ [SNNC1γ]) Can a channel or channels regulated by nucleotides and inhibited by high 17β-estradiol levels during specific phases of the menstrual cycle have an impact on the incidence and severity of airway disease in women? Coakley et al. (7) speculate that antiestrogen therapy (such as tamoxifen) should be tested in women with CF to promote maximal non–CFTR-mediated Cl− regulation. However, we must consider whether the known side effects, such as endometrial cancer, pulmonary embolism, deep-vein thrombosis, stroke, uterine abnormalities, and cataracts, outweigh the potential benefits of this proposed therapy. The authors predict that for one week out of every four of a woman’s menstrual cycle, airway mucociliary clearance will be compromised by a decrease in CaCC function. Is this cyclical compromise in mucociliary clearance responsible for diminished female survival in CF? A small study of 12 women with CF compared forced expiratory volume in 1 second (FEV1) at time of ovulation (high 17β-estradiol and low progesterone levels) to that during the luteal phase (high 17β-estradiol and high progesterone levels) and during the menstrual phase (low 17β-estradiol and low progesterone levels) (21). FEV1 was significantly higher during the luteal phase as compared with FEV1 during ovulation and menstruation. These authors interpreted their findings to suggest that lung function changes were related to progesterone levels, based on their earlier studies of delayed onset of puberty in CF girls (22).

Sex-based differences have been described for a number of aspects of CF. Some of these differences may be the result of hormonal exposures or genetic inheritance. Others may be sociological or behavior based. Over the past few decades, there has been an increased mortality reported for women with CF (23); however, this sex-related difference may be dissipating as more attention is given to aggressive therapy, as discussed in Coakley et al. (7). Of the causes of mortality, lung infections are described more often for women with CF (23), and death rates tend to increase for women with CF (23); however, this sex-related difference may be dissipating as more attention is given to aggressive therapy, as discussed in Coakley et al. (7). Of the causes of mortality, lung infections are described more often for women with CF (23), and death rates tend to increase for women with CF (23); however, this sex-related difference may be dissipating as more attention is given to aggressive therapy, as discussed in Coakley et al. (7). Of the causes of mortality, lung infections are described more often for women with CF (23), and death rates tend to increase for women with CF (23); however, this sex-related difference may be dissipating as more attention is given to aggressive therapy, as discussed in Coakley et al. (7). Of the causes of mortality, lung infections are described more often for women with CF (23), and death rates tend to increase for women with CF (23). In data collected from 1995 to 2005 in Ireland (24), both male and female adults with CF were more likely to die if they had worse lung function and were infected with either Pseudomonas aeruginosa or Burkholderia cepacia complex. FEV1 and infection with P. aeruginosa or B. cepacia are the most significant predictors of survival (24). Looking at the bulk of studies over the past two decades, women with CF have a worse prognosis overall; they participate less in aerobic exercise, ingest fewer kcals, perform less physical therapy, exhibit an accelerated decline in FEV1 with acquisition of P. aeruginosa, and show increased asthma reactivity. Mucociliary clearance in particular is an important defense mechanism in CF lung disease. Female anatomy may impose increased vulnerabilities toward impaired mucociliary clearance. For example, women have smaller lungs than men relative to their height, and comparison of females and males with similar lung volumes shows that female airways are smaller. The smaller
Therapeutic implications

Coakley et al. (7) demonstrate that 17β-estradiol affects Ca++ signaling, not CaCC conductance. The P2Y2 receptor agonist denulosof (25) activates CaCC conductance through the same pathway as UTP. These data point to an 17β-estradiol-mediated blockade of Ca++ signaling distal to the purinergic receptor, raising the concern that the investigatory product denulosof may have less efficacy for women during the high–17β-estradiol periovulatory period of their menstrual cycles. Antiestrogen therapy might restore the ability to respond to P2Y2 agonists but at the cost of potential side effects. One alternative to antiestrogen therapy might be alternative downstream releasers of Ca++ such as Moli1901 (lancuvotide) (17). Another might be activation of a third parallel alternative Cl− channel such as the pH- and voltage-activated Cl− channel 2 (CLC2) (Figure 1), which can be stimulated by a prostone agonist such as lubiprostone (26) or the related investigational drug cobi-prostone (www.sucampo.com/inthepipeline.html). Many of these agents are not yet approved for human use and are still in clinical development. Rather than attempting to compensate for loss of the CaCC, it might be possible to utilize ENaC antagonists and reduce the driving force for periciliary fluid reabsorption (27). Investigational ENaC antagonists in development for CF lung disease include long-acting amiloride analogs, prostanin inhibitor QAU145 (28), and INO-4995 (27).

In summary, female sex–based vulnerabilities in CF, such as pregnancy-related declines in lung function and compromised nutrition and accelerated declines in lung function beginning at puberty, have long been discussed. More recent studies suggest that the gap in lung function and prognosis between women and men is narrowing. If sex hormone cycling is leading to a significant reduction in airway mucociliary clearance, perhaps low-dose oral or patch contraceptives could be modified to reduce the disadvantage. The results of the current study by Coakley et al. (7) reinforce that there is clearly a pressing need to raise awareness of sex-related differences in lung disease.

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