Review Article

Gender Differences, Aging and Hormonal Status in Mucosal Injury and Repair

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ABSTRACT: As the “baby boomers” age, the percentage of the population over sixty-five years of age is increasing rapidly. Chronic disease management is an important component in the care of the elderly. The effects of aging on different organ systems are also pertinent; such as the weakening homeostatic response to injury in the older individuals. Mucosal surfaces have the largest combined surface area in the body and are the site of important host-microbe interactions, especially in the gut which is prone to injury, both from local and systemic insult. This susceptibility has been known to increase with age. Therefore it is important to understand the interplay between aging, injury and recovery at the mucosal surface. Sex hormones play an important role in the maintenance of the mucosal barrier function as well as the mucosa associated immune function in both genders. Menopause in women is a defined time period in which major hormonal changes occur such as a decline in systemic estradiol levels. The differential levels of sex hormones contribute to the sexual dimorphism seen in response to injury at the mucosal surface, prior to and following menopause. Thus the effect of sex hormone and aging on mucosal mechanisms in response to injury is an important area of investigation.

Key words: aging, mucosal injury, gut mucosa, epithelial barrier function, microbial translocation, estradiol, hormones, menopause

By 2050, the total US population ages 65 and older is estimated to reach 89 million [1]. The prevalence of chronic diseases is very high in this age group, the likely consequence of which will include increased health care costs [2, 3]. Among the most prevalent chronic conditions are cardiovascular disorders, cancer, respiratory disease, arthritis and gastrointestinal disease [4-7]. All these diseases have an underlying immune dysregulation related etiology, which may cause persistent tissue damage. The increased propensity to tissue damage and reduced rate of tissue repair in the elderly contribute to disease progression [7, 8]. Thus, the study of injury and repair in the aging population is significant.

Mucosal surfaces, particularly those of the urogenital, respiratory and oro-gastro-intestinal (OGI) tracts, are often affected by chronic diseases and challenged by pathologies, due to factors such as infection and chemical insult. For example, sepsis associated with pneumonia and urinary tract infections (UTIs) is predominantly a disease of the aged, with increased incidence and mortality occurring in older individuals [9, 10]. Additionally, disorders like dysphagia and gastro-esophageal reflux disease, present special management challenges and often led to complications such as impairment of nutritional status and a reduction in the quality of life [11, 12]. There is substantial evidence that the aging mucosal surfaces suffer both structural and functional defense defects, thus changing the homeostatic baseline and contributing to compounded pathology when challenged with disease [13, 14]. To be able to facilitate, via pharmaceutical intervention, enhancement of the body’s mucosal compartments and their repair following...
injury in an aging population, a better understanding of age-related changes is required. This review intends to address the current knowledge of the influence of age and gender on mucosal injury and repair; and specifically the contribution of sex hormones to these processes.

Role of sex hormones in immune modulation

In the aging female population increased susceptibility to mucosal injury comes on the heels of menopause-induced hormonal alterations. During the reproductive years, females exhibit a more robust humoral and cellular immune response as compared to age matched males or post-menopausal counterparts [15]. Immunosenescence in the aging population is thought, in part, to be a result of altered hormonal status and decreased production of estrogen (17β-estradiol; E2) [16]. The major effects of E2 are mediated through two receptors, ERα and ERβ, both of which are expressed on a variety of cell types including, but not limited to, immune cells, epithelial cells and muscle cells [17-19]. Some data also suggest a spatial difference in immune-regulation by the two receptors with respect to tissue type [20]. Estrogen has been shown to regulate many facets of the immune response such as immune cell differentiation, cytokine production and regulation of Ca++ mobilization and release of inducible nitric oxide synthase within leukocytes [18, 21]. Thus, it is likely that decreased E2 production with aging substantially impacts mucosal health and recovery via loss of its immune-modulatory effects [22-24].

Like females, aging males also experience a profound reduction in the levels of sex hormones, specifically androgens. Though androgen decline in males is associated with similar defects in mucosal healing and repair, it should be stressed that “male menopause” follows a very different progression pattern as compared to female menopause. It has been documented that from the ages of 25-75 years a healthy male experiences approximately 30% loss of circulating testosterone, with over 50% of males over the age of 65 years meeting endocrine criteria for hypogonadism [25, 26]. In addition to testosterone’s direct immune-modulatory function, it also acts as a pro-hormone, converted to both 5α-hydrotestosterone (DHT) and E2 [27]. Because E2 also contributes greatly to sex hormone regulation of immune response in males, the effects of androgen loss with age is compounded. Independently, androgens contribute to both pro- and anti-inflammatory states, modulating cytokines such as IL-1, IL-2, IL-6 and TNFα in a variety of cell types including macrophages, Kupffer cells, fibroblasts and splenocytes [28-30]. Thus, in males, it is proposed that the major actions of testosterone are mediated though both aromatization to E2 and E2-independent mechanisms; however, it is also the balance between testosterone and E2 which may be responsible for the immune regulation in mucosal healing following injury.

Gut mucosal function and aging

The oro-gastro-intestinal (OGI) tract carries out the functions of food processing and digestion, nutrient absorption, and expulsion of waste. Along with its role in digestion, the gut also harbors about 90% of the body’s lymphocytes within the gut associated lymphoid tissue (GALT) [31]. The intestinal structure is comprised of simple columnar epithelium, mucosa, submucosa, smooth muscle and serosa. The absorptive surface area of the intestine is increased by plicae circulares, villi, and microvilli. Glandular epithelium is present along the whole length of the gut in the form of goblet cells, which secrete mucous that lubricates the passage of food and protects the tissue from digestive enzymes (Figure 1). Changes in the microenvironment of the small intestine are associated with alterations in the composition, pH, and thickness of the mucous layer [32, 33]. Villi are in vagnations of the mucosa and increase the overall surface area of the intestine. The next layer is the muscularis mucosa, a layer of smooth muscle that aids in the action of continued peristalsis along the gut. The submucosa contains nerves, blood vessels, and elastic fiber with collagen that stretches with increased capacity but maintains the shape of the intestine. Surrounding this is the muscularis externa comprised of longitudinal and smooth muscle that helps with continued peristalsis and...
the movement of digested material out of and along the gut. Lastly there is the serosa, which is made up of loose connective tissue and is coated in mucus so as to prevent friction damage from the intestine rubbing against other tissue.

Figure 2. Fluorescent immunohistochemical staining of jejunum tight junction proteins. (A). Goblet cell diameter is marked in yellow. Nucleus: Blue Dapi, Cell Membrane: Orange mask. Image was obtained at 100X oil immersion confocal microscope. (B and C) IHC demonstrating the expression of Tight junction proteins, occluding (red) and ZO1 (Green) in small intestinal tissue (60X).

One of the key characteristics of the GI tract epithelium is the rapid proliferation of cells that differentiate from immature stem cells, within the crypt, to terminally maturated cells, which move up the villus. In the murine model it has been observed that a state of hyperproliferation, not hypoproliferation, occurs in the gut mucosal epithelial cells of older (aged) rats compared to younger rats fed the same diet and with the same living conditions [34]. This increased turnover of epithelial cells was hypothesized to be due to increased loss of epithelial cells at the tips of the intestinal villi. Furthermore, abnormalities of the proliferative and differentiation responses became more evident when gastrointestinal tissues were stimulated by systemic injury.

Mucosal barrier function is essential to prevent potentially harmful pathogens within the gastrointestinal lumen, respiratory tract, and urogenital tract from gaining access to the body [35]. Increased microbial translocation into mucosal tissue and the blood stream results in increased systemic inflammatory cytokine production [36-39]. Cellular tight junctions (TJs) are dynamic structures located in the most apical region of cell-cell contact points and play a critical role in maintenance of epithelial barrier function, cell polarity, and intercellular adhesion [40, 41]. Tight junction proteins include zona occludens, occludin (OCLN), claudins (CLDNs) and others (Figure 2).

Disorders of the gastrointestinal tract, including increased incidence of diarrhea and constipation, are common in elderly people; however, the molecular mechanisms of aging that contribute to the vulnerability of the gastrointestinal tract have not been fully elucidated. Studies suggest that patients with gastrointestinal disorders have increased intestinal permeability. A study utilizing the baboon model has shown that gastrointestinal permeability was higher in colonic biopsies in aging monkeys [42]. Tight junction protein expression was decreased including zona occluden 1 (ZO-1), OCLN, and junctional adhesion molecule-A (JAM-A). Claudin 2 (CLDN2), a pore forming tight junction protein, expression was increased. Inflammatory cytokines interferon gamma (IFN-γ), interleukin 6 (IL-6), and interleukin 1 beta (IL-1β) were also found to be increased in colonic biopsies from old baboons compared to young baboons, and have previously been shown to directly hinder tight junction complex formation [14, 42, 43]. IL-1β also contributes to a disruption of tight junction integrity [44]. IL-1β levels are consistently higher in older individuals as well as during the course of systemic inflammation and injuries like severe burns. IL-1 receptor (IL1R) also plays an important role in the maintenance of epithelial integrity in aging [43]. These studies indicate that increased colonic permeability via age-associated remodeling of intestinal epithelial tight junction proteins may be an important component of gastrointestinal dysfunction.
Collagen provides the connective tissue backbone to the OGI mucosa (Figure 3). The effect of aging on connective tissue structure, as it pertains to mucosal tissue injury and repair, has been extensively investigated in the rodent model. Connective tissue and collagen deposition in the rat gastric mucosa increases with aging [46]. Studies have shown that age-induced changes in gastric connective tissue structure lead to a decreased capacity for tissue repair in response to gastric acid [45]. Furthermore, accumulation of oxidative products, observed in the stomach of older rats, is hypothesized to contribute to thickening of connective tissue deposition and replacement of mucosal tissue in the lower part of the gastric mucosa [46]. It is known that the intestinal milieu of inflammatory cytokines and systemic circulation of sex hormones play a significant role in both fibroblast and keratinocyte migration and proliferation. These cell types are critical in the deposition and restructuring of collagen throughout the body. Changes in collagen deposition, composition, and restructuring in aging may impact not only function of the OGI mucosa but also response to injury.

Sex hormones and mucosal barrier function

Mucosal surfaces are unique anatomical niches, as they are an interface between a sterile internal environment and a contaminated external environment [47, 48]. Mucosal sites require contact with the external environment to perform nutrient absorption in the small intestine, gas exchange in the lungs, water reabsorption in the colon etc. For this reason, one of the key functions of the epithelial cells which coat mucosal surfaces is to maintain barrier functions from the antigens of the external environment. The epithelial layer possesses polarity and requires close contact between the cells to function as an anatomical barrier. Maintenance of the barrier is critical for mucosal and systemic health, and is first to be damaged during external injury. Rapid healing of the epithelium and revival of barrier function is important in preventing ongoing immune activation and further infection at mucosal sites [49, 50]. With age, this process is slowed and results in some of the co-morbidities such as endometriosis, increased incidence of UTIs, diarrhea, as well as increased prevalence of pulmonary hypertension and protracted recovery from lung disease, associated with mucosal injuries in the aging population.

Disruption of TJ complexes is associated with a variety of human diseases including genital and gastric cancers, inflammatory bowel disease, and HIV infection [51-53]. Interestingly, sex hormones such as E2 play an important role in the maintenance of tight junctions. E2 levels decrease dramatically during the course of the menopausal transition [54-57]. E2 affects TJ formation, which can alter the level of bacterial translocation from any mucosal surface [58, 59]. A recent study showed that increased E2 in rats induced the expression of TJ protein OCLN via binding of ERβ [59]. This resulted in decreased intestinal epithelial permeability [59] and microbial translocation. Conversely, reduced levels of E2, as occurs with aging, particularly during female menopause, can potentially increase epithelial permeability and microbial translocation (Figure 4). No significant disparities are observed between the genders with regards to homeostatic epithelial barrier permeability [27]. Therefore, an E2-dependent mechanism for maintenance of the epithelial barrier may contribute equally in both males and females to mucosal integrity.
Gender differences in OGI mucosal injury

Recently, several large-scale, cross-sectional epidemiological studies have been performed to determine whether there is gender-based skewing in populations of individuals with gastro-esophageal reflux diseases (GERD) [60-62]. The three conditions that comprise the majority of this spectrum include Barrett's esophagus (BE), erosive reflux disease (ERD), and nonerosive reflux disease (non-ERD). Both Barrett’s esophagus and ERD are associated with erosive esophagitis and mucosal injury [61, 63]. Furthermore, it is generally accepted that there is a sequential progression from reflux erosive esophagitis, to Barrett's esophagus, and finally to esophageal adenocarcinoma. Reflux erosive esophagitis results from exposure of the esophageal epithelium to the refluxed gastroduodenal contents. A male-predominant gender bias exists across the spectrum, although the ratios become higher with the progression towards esophageal adenocarcinoma. Meanwhile, non-erosive reflux diseases generally affect women more than men [60, 62]. These data allude to the gender differences in the vulnerability or resistance of the esophageal epithelium to caustic compounds of the gastroduodenal contents in males and females.

There have been reports that females are less affected than males by gastric or intestinal inflammation in response to chemical insult or bacterial infection [64-66]. In these studies E2 was demonstrated to have anti-inflammatory activity and thus contributes to tissue resistance in females. A recent study by Masaka et al. explored the potential role of E2 in controlling esophageal tissue damage [63]. Employing a chronic rat reflux esophagitis model, a significant male-predominant, gender-related difference in esophageal tissue damage in the presence of exogenous nitric oxide (NO) as an exacerbating factor, was found [67]. While in the baseline model of reflux esophagitis macroscopic esophageal ulcers and microscopic inflammatory cell infiltrates were only mildly observed in both genders, in males, exogenous NO exacerbation induced deep esophageal ulcers and intense inflammation with polymophonuclear cell and lymphocyte infiltrates. In contrast, in female reflux esophagitis models, treatment with NO rarely exacerbated the mild tissue damage observed at baseline. Further, exogenous 17β-estradiol binding and signaling through E2 receptors attenuated esophageal tissue damage in males and ovariectomised rats via a reduction of mast cell-mediated cytotoxicity and cytokine, specifically tumor necrosis factor alpha (TNFa), driven inflammation [63, 68]. Treatment with 17α-estradiol, which binds but does not induce downstream signaling, had no effect on tissue damage. This was the first study showing the prominent gender difference in the severity of esophageal tissue damage in a GERD-related animal model. Additional understanding of the causative luminal or genetic factors
in yielding the gender-related difference would be clinically relevant to predict the etiological factors involved in the pathogenesis of reflux esophagitis in humans [69, 70].

Sex hormones in tissue repair

Gender variation in wound healing has been observed at various mucosal sites throughout the body. Specifically, variations in healing rates have been attributed to levels of circulating sex hormones and their effect on modulation of inflammation as well as fibroblast and keratinocytes cellular proliferation, differentiation and growth [71-73]. It has been established that both dermal and mucosal wound healing is significantly altered by the effects of E2 signaling [74-76]. Kumral et al. recently showed that gastric and colonic tissue damage is alleviated by E2, via both ERα and ERβ mediation as well as direct antioxidant effects [76]. In addition to its role in inflammatory regulation and re-epithelialization, an epidemiological study by Tuo et al. linked higher circulating E2 levels in women to the elevated production of duodenal mucosa bicarbonate secretion (DBS), a compound central in duodenal mucosal protection against acid-induced injury [77]. Duodenal ulcers are less prevalent in pre-menopausal women, compared to age matched men or post-menopausal women [78, 79]. In their findings, ex vivo stimulation of duodenal tissues with 17β-estradiol did not result in a difference in the levels of DBS secretion. The authors hypothesized that this stemmed from the result that males and females express similar levels of ERα and ERβ on duodenal epithelial cell surface [77]. This highlights that the observed sex differences of DBS were likely due to the gender differences in circulating E2 levels rather than a dimorphism in expression levels of E2 receptors between different sexes, and can possibly be extrapolated to the other effects of E2.

Likewise, metabolites of testosterone, as well as other androgens [5α-dihydrotestosterone (DHT) and Dehydroepiandrosterone (DHEA)], have been shown to affect dermal wound closure by impairing re-epithelialization and inducing immunosuppressive effects [80]. As in dermal wounds, a study by Engeland et al. reported that in oral mucosal damage testosterone levels were inversely correlated with wound healing rates in pre-menopausal women and age matched males. Conversely, in post-menopausal women a positive correlation of testosterone levels and wound healing rates was observed [81]. It was hypothesized that the immunomodulatory role of testosterone in reducing IL-6, which is mitogenic to keratinocytes, contributes to the effect observed in pre-menopausal women and men. The effect observed in post-menopausal women was not linked to age specifically, but hormonal status. Authors put forth the idea that with the increased immune activation observed in post-menopausal women at baseline, higher levels of anti-inflammatory testosterone decreased this activation thus being beneficial to tissue healing [82, 83].

Estrogen and Urinary Tract Infections

While UTIs are most prevalent in females aged 18-24, a significant number of women over 50 still contract UTIs [84]. Recurrent infections in healthy, aging women ages 50-70 have been linked to decreased levels of estradiol [85]. After menopause, decreased levels of E2 cause vulvovaginal atrophy in 25-50% of women [13]. Symptoms such as vaginal dryness, itching, increase in vaginal pH, urinary frequency and incontinence, contribute to the impairment of defenses against incoming pathogens at the urogenital mucosa [86]. Thus estradiol supplementation has been considered as a way to decrease the risk of recurrent infections in the postmenopausal population and has demonstrated moderate success [87]. One of the mechanisms by which estradiol therapy in post-menopausal women has proven successful may be related to tight junction formation enhancement by E2. Numerous studies have shown that estrogen treatment, in vitro, increases tight junction protein expression including ZO1 and CLDN in the vaginal epithelium [88]. In both, a urothelial cell line and exfoliated bladder cells from postmenopausal women, estradiol treatment increased transcripts of ZO1 and OCLN as well as e-cadherin protein [86]. This demonstrates that estrogen’s beneficial effects on tight junction proteins may occur on mucosal surfaces outside of the vagina [86]. Estrogen-mediated restoration of a diminished antimicrobial response in post-menopausal women could also contribute to the decreases seen in UTIs following treatment. When post-menopausal women were given estradiol supplementation for two weeks, 75% showed increases in at least three antimicrobial peptides in urinary tract cells. The most highly increased peptides were beta-defensin 3 (hBD3), beta-defensin 1 (hBD1), and RNase 7 [86].

An alternate mechanism by which estradiol supplementation can contribute to urogenital mucosal health in post-menopausal women is via its role in vaginal pH regulation. A number of theories have been proposed for the contribution of E2 toward vaginal pH control, including direct effects on the epithelium and altering vaginal microflora populations [89, 90]. In healthy, young, menstruating women vaginal microflora is dominated by Lactobacilli which produce lactic acid, hydrogen peroxide, and various bacterial proteins which together inhibit overgrowth of any pathogenic bacterial species [91]. Thus, the vaginal microflora confers
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**Sex Hormones in Lung Disease**

While the prevalence of chronic obstructive pulmonary disease (COPD) is greater in men, women appear to be more sensitive to the effects of cigarette smoke in developing COPD, often developing COPD after smoking less than men [92]. This may be related to an E2-mediated difference in metabolism of cigarette smoke toxins at the lung mucosa. Estradiol increases the activity of cytochrome P450 (CPY) enzymes, which break down toxins but create harmful metabolites which can be more toxic than their precursors [93]. Because complete metabolism of these toxins requires a rate limiting process which is not increased by E2, harmful intermediate metabolites may remain in the lungs of women smokers longer than their male counterparts, increasing lung damage and resulting in poorer clinical outcomes.

A gender discrepancy in outcomes of cystic fibrosis (CF) patients has been seen clinically, with females having much higher early mortality rates than males [94]. This may be due to an E2-mediated increase in mucus production, a principal determinant of mortality in CF patients. Estradiol has been shown to increase mucin gene expression in lung epithelium, which may result in increased mucus overall [95]. Additionally, female CF patients show lowest lung function just before ovulation, suggesting that this surge of E2 may be enhancing mucus production and thereby decreasing lung function [96].

Taken together, these studies demonstrate the wide variety of ways E2 modulates lung function and affects disease morbidity. In some instances, modulation of sex hormones may be a viable clinical intervention worthy of further investigation.

**Summary**

Aging and sex differences play an important role in the development of mucosal injury as well as its repair. The role of sex hormones is controversial at best. The part that is clear is that aging mucosal surfaces are extremely susceptible to injury. New studies indicate that the microbiome may also play an important role in mucosal health. This too is a rapidly developing area of research that is poorly understood and likely plays a role in healthy aging. The molecular mechanisms that regulate aging are poorly understood. Taken together comprehensive studies using relevant animal models are needed to better understand the interplay between mucosal injury, gender, sex hormones and aging.

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