Sex Hormones Production in Both Genders Skin Diseases

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Abstract
The skin is the largest organ in the body constituting 15% of the total body weight. It is therefore paramount to address skin diseases. Of major importance in the etiopathogenesis and pathophysiology of many of these diseases is the gender difference in the skin. Studies on endocrine effects on the skin have revealed that several important physiologic activities of the skin are either partly or wholly under the control of hormones secreted by different endocrine glands.

Keywords: gender; skin; dermatoses; sex hormones.

Introduction
The skin is an endocrine organ involved in the peripheral conversion of sex hormones to active forms (testosterone and DHT in males and estriadiol in females). Skin structures such as sebocytes, sweat glands and dermal papilla hair cells express enzymes that convert DHEA and androstenedione from adrenal cortex into testosterone and DHT. Aromatases convert testosterone into estradiol in the skin. The sebaceous glands, the outer and inner root sheath cells of anagen terminal hair follicles and dermal papilla cells express aromatases. Various structures in the skin show enzyme activity and express sex hormone receptors [1]. Epidermal keratinocytes contains androgen receptors (AR), estrogen receptor-beta (ERβ), and the enzymes 17β hydroxysteroid dehydrogenase (HSD) and 5α-reductase. Melanocytes express the AR, ERs and ERβ receptors. A disease is an aberration in the normal. This could be in the form of changes in structure or function. The functions of the skin have to be considered in relation to its structure i.e. the structure definitely determines its function. Structural and physiological changes in the skin can occur as a result of infection, genetic changes, physical factors (such as heat), nutritional diseases, metabolic derangements, cosmetic use, drug use, organ malfunctions, environmental factors (specific for the individual, for example, occupation, social lifestyle, geographical location, antibiotics use, the use of cosmetics and soaps), and endocrine changes.

Sex Hormones Production in Both Genders

The study of skin diseases is often related to age groups and environmental factors while their relationship to gender has not been fully elucidated. Differences in the structures and functions of the skin in both genders underline the gender differences observed in many skin diseases. Various structures in the skin are under the influence of endocrine activities in the skin or outside the skin. Sex hormones – androgens and estrogens – influence the skin structure and functions. In young men, testes produce testosterone and in young women, ovaries are the main source of estradiol (Figure 1) [1]. Sex steroids are produced by the hypothalamic-pituitary-gonadal axis, the adrenals (which produce dehydroepiandrosterone (DHEA) and androstenedione – precursors of androgens and estrogens in peripheral tissues), and by peripheral tissues (Figure 2) [1].

With aging, gender differences occur in serum concentrations of these hormones. In men, gonadal production of androgens slowly decreases and by 65 years, peripheral tissues such as skin produce 40% of the total androgens [2].

In women, the peripheral tissues produce almost all the sex steroids from DHEA after menopause [3]. In both genders, a gradual decline in DHEAS and DHEA occurs with age and this is thought to contribute to the degenerative changes seen with aging [4-8].

Effect of Sex Hormones on the Structure and Functions of the Skin in Both Gender

The effect of sex hormones differs in men and women [9]. Androgens affect several functions of the human skin, such as sebaceous gland growth and differentiation, hair growth, epidermal barrier homeostasis and wound healing [10]. Facial, axillary and pubic hairs respond to androgens while the eyebrows and eyelashes do not. Hence, masculinizing syndromes in women are usually accompanied by hirsutism [11]. Removal of testicular hormones in men results in reduction of beard growth, and re-growth of beard occurs when treated with androgens [12]. Excess testosterone in females results in androgenetic-dependent alopecia while excess testosterone (by increasing sebum production) in males predisposes to severe acne and removal of testosterone results in reversal of male-pattern hair loss [1,13,14].

Wound healing is slower in males. Trauma results in changes in sex steroid concentrations with higher estrogen concentrations in both sexes and decreased testosterone levels in males [15-18]. At physiological levels, 5α-dihydrotestosterone (5α-DHT) decrease wound immune function and impair wound healing after trauma and hemorrhage by increasing proinflammatory cytokines and decreased tumor growth factor-β at the wound site [19,20]. The male skin is thicker than female skin [21]. Females, however, have thicker subcutaneous tissues [22]. Female skin becomes thinner than male skin with aging due to reduction in estrogen [23,24]. Estrogen therapy increases skin thickness [25].

Skin surface pH, and sweat rate are higher in males [9]. With the higher skin surface pH, the resultant effect is that microbial diversity is often lower in men as they are hindered by more acidic environments [26]. Trans-epidermal water loss (TEWL), erythema index and melanin index were found to be higher in men living in hot areas while women have higher skin hydration and elasticity [27]. Young men have higher SC hydration in comparison with women until the age of 40 when it progressively decreases while it remains stable or even increase in women with age [28].

Sex steroids have a differential effect on the immune system in men and women. Androgens are anti-inflammatory and depress both cellular and humoral immunity resulting in men having higher susceptibility to skin infections [29-31].
Men are more prone to bacterial and viral infections though they have less microbial diversity in their skin than women due to more acidic skin pH [9,26]. Estrogens stimulate the humoral immunity and cause development of auto reactive B cells, and inhibit apoptosis which results in survival of auto reactive T cells. Autoimmune diseases such as SLE and allergic contact dermatitis (related to occupation) occur more in women as a result of these effects [15].

Gender Related Dermatoses

Research studies involving skin physiology are very few in Nigeria. Not many studies have been performed to determine the relationship between gender and skin diseases. In a study by Oninla et al. [30] at Ille-Ife, gender predilection was studied among 1,013 patients presenting with 1,097 dermatoses [30]. A comparison was made with a study reporting the prevalence of skin diseases in both genders. Viral warts, folliculitis/carbunculosis, Hansen’s disease, elephantiasis were found to have a significant relationship with the male gender [31]. As mentioned earlier, bacterial and viral infections are more common in males. Infections seem more in females but not significant in them were can did an intertrigo, herpes zoster and onchodermatitis. Few studies have been done in Nigeria on the effect of gender on skin diseases. Studies of dermatological conditions in relation to measured hormone levels as well as skin physiological changes by hormones in both genders are needed.

References

1. Farage MA, Miller KW, Zouboulis CC, Piérard GE, Maibach HI. Gender Differences in Skin Aging and the Changing Profile of the Sex Hormones with Age. J Steroids Horm Sci. 2012;3(2):1-16.
2. Labrie F. DHEA, important source of sex steroids in men and even more in women. Prog Brain Res. 2010;182:97-148. doi.org/10.1016/S0079-6123(10)82004-7
3. Labrie F, Bélanger A, Luu-The V, Labrie C, Simard J, Cusan L, et al. DHEA and the intracrine formation of androgens and estrogens in peripheral targettissues: its role during aging. Steroids. 1998;63(5):322-8. doi.org/10.1016/S0039-128X(98)00007-5.
4. Bélanger A, Candas B, Dupont A, Cusan L, Diamond P, Gomez JL, et al. Changes in serum concentrations of conjugated and unconjugated steroids in 40- to 80-year-old men. J Clin Endocrinol Metab. 1984;79(4):1086-90. doi.org/10.1210/jcem.79.4.7962278
5. Orentreich N, Brind JL, Rizer RL, Vogelman JH. Age changes and sex differences in serum dehydroepiandrosterone sulfate concentrations throughout adulthood. J Clin Endocrinol Metab. 1984;59(3):551-5. doi.org/10.1210/jcem.59-3-551
6. Feldman HA, Longcope C, Derby CA, Johannes CB, Araujo AB, Coviello AD, et al. Age trends in the level of serum testosterone and other hormones in middle-aged men: longitudinal results from the Massachusetts male aging study. J Clin Endocrinol Metab. 2002;87(2):589-98. doi.org/10.1210/jcem.87.2.8201
7. Burger HG, Dudley EC, Cui J, Dennerstein L, Hopper JL. A prospective longitudinal study of serum testosterone, dehydroepiandrosterone sulfate, and sex hormone-binding globulin levels through the menopause transition. J Clin Endocrinol Metab. 2000;85(8):2832-8. doi.org/10.1210/jcem.85.8.6740
8. Sulcová J, Hill M, Hampl R, Stárka L. Age and sex related differences in serum levels of unconjugated dehydroepiandrosterone and its sulphate in normal subjects. J Endocrinol. 1997;154(1):57-62. doi.org/10.1677/joe.0.1540057
9. Giacomoni PU, Mammone T, Teri M. Gender-linked differences in human skin. J Dermatol Sci. 2009;55(3):144-9. doi.org/10.1016/j.jdermsci.2009.06.001
10. Zouboulis CC, Degitz K. Androgen action on human skin – from basic research to clinical significance. Exp Dermatol. 2004;13:5-10.
11. Moschella SL, Hurley HG. Dermatology. Philadelphia: WB Saunders Company, USA; 1985.
12. Fitzpatrick TB, Zur Eisern A, Wolff K, Freedberg IM, Austen KF. Dermatology in general medicine. New York: McGraw-Hill, USA; 1987.
13. Boisselle A, Tremblay RR. New therapeutic approach to the hirsute patient. Fertil Steril. 1979;32(3):276-9.
14. Chen W, Thiboutot D, Zouboulis CC. Cutaneous androgen metabolism: basic re. doi.org/10.1046/j.1523-1747.2002.00613.x
15. Fimmel S, Zouboulis CC. Influence of physiological androgen levels on wound healing and immune status in men. Aging Male. 2005;8(3):166-74
16. Christoff N, Carli A, Benassayag C, Bleichner G, Vaxelaire JF, Nunez EA. Relationship between changes in serum estrone levels and outcome in human males with septic shock. Circ Shock. 1992;36(4):249-55
17. Fourrier F, Jallot A, Leclerc L, Jourdain M, Racadot A, Chagnon JL, et al. Sex steroid hormones in circulatory shock, sepsis syndrome, and septic shock. Circ Shock. 1994;43(4):171-8.
18. Lephart ED, Baxter CR, Parker CR Jr. Effect of burn trauma on adrenal and testicular steroid hormone production. J Clin Endocrinol Metab. 1987;64(4):842-8.
19. Dao K, Kazin RA. Gender differences in skin: a review of the literature. Gend Med. 2007;4(4):308-28. doi.org/10.1016/S1550-53.21500.2007.00613.x
20. Nitsch SM, Wittmann F, Angele P, Wichmann MW, Hatz R, Hernandez/Richter T, et al. Physiological levels of 5 alpha-dihydrotestosterone depress wound immune function and impair wound healing following traumahemorrhage. Arch Surg. 2004;139(2):157-63. 10.1001/archsurg.139.2.157
21. Seidenari S, Pagnoni A, Di Nardo A, Giannetti A. Echographic evaluation with image analysis of normal skin: Variations according to age and sex. Skin Pharmacol. 1994;7(4):201-9. doi.org/10.1159/000211295
22. Sjöström L, Smith U, Krotkiewski M, Björntorp P. Cellularity in different regions of adipose tissue in young men and women. Metabolism. 1972;21(12):1143-53. doi.org/10.1006/jmsb.1987.00957(97)090109-6
23. Leveque JL, Corcuff P, de Rijal J, Agache P. In vivo studies of the evolution of physical properties of the human skin with age. Int J Dermatol. 1984;23(5):322-9.
24. Punnonen R. Effect of castration and peroral estrogen therapy on the skin. Acta Obstet Gynecol Scand Suppl. 1972;21:3-44.

25. Zeeuwen PL, Boekhorst J, van den Bogaard EH, de Koning HD, van de Kerkhof PM, Saulnier DM, et al. Microbiome dynamics of human epidermis following skin barrier disruption. Genome Biol. 2012;13(11):R101.

26. Firooz A, Sadr B, Babakoohi S, Sarraf-Yazdy M, Fanian F, KazerouniTimsar A, et al. Variation of biophysical parameters of the skin with age, gender, and body region. ScientificWorldJournal.2012;2012:386936.dx.doi.org/10.1100/2012/386936

27. Luebberding S, Krueger N, Kerscher M. Skin physiology in men and women: in vivo evaluation of 300 people including TEWL, SC hydration, sebum content and skin surface pH. Int J Cosmet Sci. 2013;35(5):477-83.

28. Kayhan M, Unluoglu I, Kayhan S, Kayhan U. Assessment of clinical diagnosis, age and gender differences of elderly patients applying to dermatology clinic of a secondary health institute in family medicine aspect. Biomedical Research. 2016.

29. Oninla OA, Oninla SO, Oke OO, Oripelaye MM, Olarewaju FO, Fabusuyi OT. Gender Differences in Dermatoses at Obafemi Awolowo University Teaching Hospitals’ Complex, Ile-Ife. BJMMR.2016;17(4):1-12.

30. Kanerva L, Jolanki R, Toikkanen. Frequencies of occupational allergic diseases and gender differences in Finland. J Int Arch Occup Environ Health. 1994;66(2):111.

31. Atraide DD, Akpa MR, George IO. The pattern of skin disorders in a Nigerian tertiary hospital. J Public Health Epidemiol. 2011;3(4):177-81.