



ADULT ACNE: ITS CLINICOEPIDEMIOLOGICAL FEATURES

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ABSTRACT

Although acne is principally a disorder of adolescence, the prevalence of adult acne is increasing. Adult acne has been defined as the presence of acne beyond the age of 25 years. Acne in adult women may have different clinical features i.e., more involvement of lower face, association with hair loss, premenstrual flare, signs and symptoms of insulin resistance, dyslipidaemias. Various studies have shown that a significant percentage of adult women with acne failed to respond to treatment with systemic antibiotics and isotretinoin which indicates a need for treatment alternatives with improved effectiveness and acceptable side effects for resistant acne. This study aims to study the different clinicoepidemiological features of adult acne.

KEYWORDS : acne, adult, hirsutism**INTRODUCTION**

Acne vulgaris is the most common skin disease of adolescents and young adults with reported prevalence of nearly 80% [1].

Adult acne has been defined as the presence of acne beyond the age of 25 years. [2] Acne persisting beyond the age of 25 years is called persistent adult acne and acne developing for the first time after the age of 25 years is called late-onset adult acne. Both the types are more common in women. Late onset acne can be further subdivided into chin acne, which occurs around the chin and perioral area, is inflammatory and flares premenstrually and sporadic acne which occurs suddenly in adult life with no distinguishing features. It has been reported that patients with adult acne have a 40% prevalence of psychiatric comorbidity. [3, 4, 5]

Aims and objectives

To study the different clinicoepidemiological features of adult acne

Study design

The study was performed in accordance with Good Clinical Practices and after clearance from the ethical committee. This observational study was conducted in the outpatient department of a tertiary care hospital. Patients were recruited for the study from July 2018 to July 2020. Seventy two adult female patients presenting with acne fulfilling the inclusion and exclusion criteria were enrolled for the study. Epidemiological profile, historical details including gynaecological history, associated illnesses, clinical grading and pictographic record, complete physical examination, laboratory findings and ultrasonography (USG) pelvis were recorded.

Study period: July 2018 to July 2020

Inclusion criteria:

1. Patients above the age of 25yrs both males and females.
2. Patients experiencing onset of lesions first after the age of 25yrs
3. Patients of adolescent acne persisting into adulthood i.e. persisting beyond 25yrs of age.

Exclusion Criteria:

1. Patients receiving any systemic steroid, ATT, any other drugs known to cause acne.
2. Patients with occupational acne, chloracne.
3. Patients receiving any form of treatment for acne in the past two months.

A total of 72 patients of adult acne were included in the study after taking written informed consent. The sample size of 72 was calculated using OpenEpi software for the prevalence of 25 % (Prevalence of acne among adults more than 25 year old) at 95% confidence level,

80% power of study and 10% of absolute precision. 05 male patients presented with adult acne however were not included in the study as they did not consent to be a part of the study. A detailed history and examination was carried out for each patient, including a medical and family history. Information about the extent and site of involvement, aggravating factors including drug intake, sun exposure, application of cosmetics, stress, premenstrual flare, hairloss, associated comorbidity, signs of hyperandrogenism, menstrual irregularities. Gynaecological history of patients was noted. At the beginning of study all patients were assessed for presence of stress based on a semi structured perceived stress questionnaire (PSS). The PSS measures the degree to which situations in one's life are appraised as stressful. [6] Data was analysed using EPI-info software. Hairloss was graded on basis of Sinclair pictographic 5-point scale.[7]

For evaluation of hirsutism a modified Ferriman Gallwey scale was used which quantified the presence of terminal hair over nine body parts: the upper lip, chin, chest, upper and lower abdomen, thighs, upper and lower back, and upper arms. They were divided into five classes and grades zero to four (0-4) assigned to them as follows: (0) in case of no hair, (1) only vellus hair, (2) terminal hair lower than 10, (3) more than 10 terminal hairs but restricted, and (4) high terminal hairs and countless. The subjects with a score of 8 or higher were defined as hirsute. [8,9,10]

Clinical assessment of each patient included type of acne lesions, distribution, severity and grading of acne was carried out. For the purpose of grading, acne Score (Hayashi et al 2008) was calculated based on the lesion count i.e. inflammatory eruptions on half of the face. [11]

Rosacea was excluded on the basis of clinical examination. Associated findings such as obesity, hirsutism and alopecia indicating hormonal imbalance were also noted.

All the patients were screened on the 2nd day of the menstrual cycle for hormonal assay and transabdominal ultrasound sonography (USG) pelvis. Fasting blood sample was collected. Following set of investigations were done for all patients.

- Baseline blood pressure recording
- Complete blood count
- Liver and kidney function tests
- Lipid profile
- Blood sugar (fasting and postprandial)
- Thyroid profile
- serum Leutinizing Hormone, Follicle Stimulating Hormone, Testosterone (total), prolactin levels

Other features observed in decreasing order were hairfall (80.6%), premenstrual flare (70.8%), dandruff (52.8%), acanthosis nigricans (22.2%), menstrual irregularities (15.3%), hirsutism (15.28%). In 2001, Stoll et al studied the effect of the menstrual cycle on acne, 44 percent of respondents experienced perimenstrual acne flare. [16] Niti Khunger et al. reported clinical features suggestive of hyperandrogenism such as premenstrual flare (11.7%), hirsutism (5.7%) and alopecia (1.8%). [12]

In the present study, FPHL was observed in 69% of patients. Approx. 60% patients had grade 1 FPHL as per Sinclair grading. Acne scoring was done as per Hayashi scoring. 93.1 % of patients did belong to the moderate category as per Hayashi scoring. No significant correlation was observed amongst Hayashi acne scoring and Sinclair grading of FPHL.

About 22% patients had family history of DM, 4.2% had Positive family history of PCOS and hirsutism each, only 2.8% had family history of acne. Studies have shown that increased serum levels of IGF-1 have been observed in adult women and men with acne, giving rise to the possibility of hyperinsulinemia in acne. As insulin resistance is usually familial, serum insulin levels should be measured in patients of treatment resistant acne especially in patients with positive family history of DM. [17]

In the present study, semistructured questionnaire stress scale was devised to know the level of stress in acne patients. 11.1 % had positive history of stress due to acne vulgaris. According to Goulden et al., who reported that in 71% of their patients acne flared with stress. [13] Kligman AM proposed that perhaps the major etiological factor in adult acne was increased levels of stress, leading to increase in adrenal androgens. [18]

In the present study, 12.5% had a positive history of use of cosmetics. While in a study by Niti Khunger et al. 40 patients (22%) out of 176 using some form of cosmetics reported aggravation due to cosmetic use. [12]

The present study reported that 37.5% of patients had dyslipidemia. Deranged thyroid function was observed in 4.16% of patients. 8.3% had abnormal USG out of which three patients 4.17% fulfilled the criteria for PCOD. Legro RS et al. and Nina Madnani et al had described that PCOS has been associated with metabolic complications including obesity, dyslipidemia, insulin-resistance (IR) and a risk of developing Type 2 diabetes mellitus (T2DM). [19, 20]

CONCLUSION

The present study was conducted with the objective of assessing the clinical spectrum of adult acne.

Acne in adult women may have different clinical features i.e. more involvement of lower face, association with hairloss, premenstrual flare, signs and symptoms of insulin resistance, dyslipidemias.

Dermatologists play a critical role in the diagnosis of PCOS as acne is a common cutaneous and presenting sign. It is important to look for potential symptoms and signs of hyperandrogenism and to exclude an underlying hormonal disorder by complete history, physical examination and relevant investigations.

All patients of adult acne may not have an underlying hormonal imbalance or hyperandrogenism.

Further studies with longer follow up are recommended to assess the treatment response in the management of adult acne patients.

REFERENCES:

1. Cunliffe WJ, Gould DJ. Prevalence of facial acne vulgaris in late adolescence and in adults. *Br Med J* 1979; 1:1109-10
2. Williams C, Layton AM. Persistent acne in women. Implications for the patient and for therapy. *Am J Clin Dermatol* 2006; 7: 281-90.
3. Henkel V, Moehrenschrager M, Hegerl U, Moeller HJ, Ring J, Worret WI. Screening for depression in adult acne vulgaris patients: Tools for the dermatologist. *J Cosmet Dermatol* 2002;1: 202-7
4. Bhamri S, Del Rosso JQ, Bhamri A. Pathogenesis of acne vulgaris: recent advances. *J Drugs Dermatol* 2009; 8:615-8.
5. Gupta MA, Gupta AK. The psychological comorbidity in acne. *Clin Dermatol* 2001; 19: 360-363. (10)
6. Gade S, Chari S, Gupta M. Perceived stress among medical students: To identify its sources and coping strategies. *Arch Med Health Sci.* 2014; 2:80-6.
7. Sinclair R, Jolley D, Mallari R, Magee J. The reliability of horizontally sectioned scalp biopsies in the diagnosis of chronic diffuse telogen hair loss in women. *J Am Acad Dermatol.* 2004; 51:189-99.

8. Hatch R, Rosenfield RL, Kim MH, Tredway D. Hirsutism: Implications, etiology, and management. *Am J Obstet Gynecol.* 1981; 140: 815-30.
9. Ferriman D, Purdie AW. Association of oligomenorrhea, hirsuties, and infertility. *Br Med J.* 1965;2: 69-72.
10. Ferriman D, Purdie AW. The aetiology of oligomenorrhea and/or hirsuties: A study of 467 patients. *Postgrad Med J.* 1983; 59:17-20
11. Hayashi N, Akamatsu H, Kawashima M. Acne Study Group. Establishment of grading criteria for acne severity. *J Dermatol* 2008; 35:255-60.
12. Khunger N, Kumar C. A clinico-epidemiological study of adult acne: Is it different from adolescent acne? *Indian J Dermatol Venereol Leprol* 2012; 78: 335-41
13. Goulden V, Clark SM, Cunliffe WJ. Post-adolescent acne: a review of clinical features. *Br J Dermatol* 1997; 136: 66-70. (9)
14. Poli F, Dreno B, Verschoore M. An epidemiological study of acne in female adults: Results of a survey conducted in France. *J Eur Acad Dermatol Venereol* 2001; 15:541-5
15. Capitano B, Sinagra JL, Bordignon V, Fei PC, Picardo M, Zouboulis C. Underestimated clinical features of postadolescent acne. *J Am Acad Dermatol* 2010; 63:782-8.
16. Stoll S, Shalita AR, Webster GF, et al. The effect of the menstrual cycle on acne. *J Am Acad Dermatol.* 2001; 45(6):957-960
17. Zoulden V, McGeown CH, Cunliffe WJ. The familial risk of adult acne: a comparison between first-degree relatives of affected and unaffected individuals. *Br J Dermatol* 1999; 141: 297-300. (8)
18. Kligman AM. Post-adolescent acne in women. *Cutis* 1991; 48: 75-7.
19. Legro RS, Kunselman AR, Dodson WC, Dunaif A. Prevalence and predictors of risk for type 2 diabetes mellitus and impaired glucose tolerance in polycystic ovary syndrome: A prospective, controlled study in 254 affected women. *J Clin Endocrinol Metab* 1999; 84: 165-9.
20. Madnani, Nina, et al. "Polycystic ovarian syndrome." *Indian Journal of Dermatology, Venereology, and Leprology* 79.3 (2013): 310.