ORIGINAL RESEARCH

A clinical and epidemiological investigation of a facial hyperpigmentation disease

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ABSTRACT

Aim: A clinical and epidemiological investigation of a facial hyperpigmentation disease.

Material and methods: A total of 100 patients diagnosed with facehyperpigmentationwereincluded in this research. Upon receiving ethical permission from the institutional ethics committee and gaining agreement from the patient, all patients seeking treatment for face pigmentation problems in the out-patient department were included in the research. Following the collection of demographic data, a comprehensive clinical history was recorded, including information on the age at which the condition was first seen, the age at which symptoms began, the length of time the illness has been present, and any relevant family medical history.

Results: The average age of the participants in the research was 29.01 ± 3.25 years. Among the 100 patients, the majority, namely 75%, were females, while just 25% were men. A higher proportion of females was observed, with a female to male ratio of 3:1. The predominant form of face hyperpigmentation seen in our research was melasma, accounting for 52% of cases. Post inflammatory hyperpigmentation (PIH) occurs in 15% of cases, whereas ephilides occur in 7%. Rehl's melanosis and drug-induced melanosis were seen in 6% of the patients each. Thirteen percent of patients had thyroid problems. In our analysis, post-inflammatory hyperpigmentation accounted for 13% of cases with changed face pigmentation, making it the second most prevalent cause. There was a small majority of females. The primary cause in the majority of cases was acne vulgaris. In our research, 14% of patients had a family history of PIH.

Conclusion: The most often reported age group was between 20 to 40 years old, with a higher proportion of females. The most frequently detected types of face hyperpigmentation were melasma (52%), post-inflammatory hyperpigmentation (15%), and ephilides (7%).

Keywords: PIH, Face hyperpigmentation, Melasma, Ephilides

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INTRODUCTION

Facial pigmentary diseases refer to a diverse collection of conditions characterized by changes in the pigmentation of the face, which are immediately noticeable and may cause esthetic deformity. The skin may have pigmentation abnormalities that are either characterized by excessive melanin production (hyper-melanotic) or insufficient melanin production (hypo-melanotic). Hyper-melanotic illnesses include a wide range of conditions, such as melasma, lichen planus pigmentosus (LPP), Riehl' s melanosis, and periorbital hyperpigmentation (POH).¹Dark skin phenotypes have a greater concentration of melanin, a higher ratio of eumelanin to pheomelanin, and a more efficient distribution of melanin for shielding against ultraviolet (UV) radiation. The quantity and

distribution of melanin in individuals with darker skin tones are significant biological characteristics. Melanin is not a single substance, but rather a combination of biopolymers that are produced by melanocytes found in the basal layer of the epidermis. Melanins are categorized into two categories, eumelanin and pheomelanin, based on their chemical makeup. Various studies have shown that persons with darker skin possess a greater quantity of total melanin and a higher concentration of eumelanin compared those skin.2to with lighter ⁵Melaninisthekeyfactorofcolourintheskin. The concentration of epidermal melanin in melanosomes doubleindarkerskin types is compared with lightlypigmentedskintypes.5 In addition. melanosomedegradation with in the keratinocyte is

slower in darkly pigmented skin when compared with lighter skin types.6The melanin content and melanosomal dispersion patternis thought to confer from damage protection induced byUVradiation.^{1,3}While the heightened presence of melanin offers defense against the detrimental impacts of UV radiation, such as photodamage and skin malignancies, it also renders darkly pigmented skin more post-inflammatory susceptible to dyspigmentation. The current research aimed to assess the clinical characteristics of individuals with face hyperpigmentation.

MATERIAL AND METHODS

This research was done at the dermatology department in partnership with the community medicine department. It was an observational study conducted in a hospital setting. A total of 100 patients diagnosed with face hyperpigmentation were include in this research. Upon receiving ethical permission from the institutional ethics committee and gaining agreement from the patient, all patients seeking treatment for face pigmentation problems in the out-patient department were included in the research. Following the collection of demographic data, a comprehensive clinical history was recorded, including information on the age at which the condition was first seen, the age at which symptoms began, the length of time the illness has been present, and any relevant family medical history. The data on several predisposing variables, including sun exposure, pregnancy, cosmetic usage, ovarian tumor, atopy, iron deficiency, and other endocrine illnesses, were documented. Relevant tests were conducted as necessary to exclude the possibility of these causes.

STATISTICAL ANALYSIS

The data thus collected was entered in M S excel sheet and analysed by using SPSS 25.0 version. The

qualitativedata was presented as percentages and quantitative datawaspresented as mean and standarddeviation.

RESULTS

We included a total of 100 individuals diagnosed with face hyperpigmentation for our research. Among the 100 patients, the largest proportion, namely 32%, belonged to the age group of 20 to 30 years, while the next largest proportion, 24%, belonged to the age group of 30 to 40 years. 17% of the individuals belonged to the age range of 40 to 50 years. The voungest individual belonged to the age group of less than 10 years. The average age of the participants in the research was 29.01±3.25 years. Among the 100 patients, the majority, namely 75%, were females, while just 25% were men. A higher proportion of females was observed, with a female to male ratio of 3:1. The predominant form of face hyperpigmentation seen in our research was melasma, accounting for 52% of cases. Post inflammatory hyperpigmentation (PIH) occurs in 15% of cases, whereas ephilides occur in 7%. Rehl's melanosis and drug-induced melanosis were seen in 6% of the patients each. Exposure to sunlight and the use of cosmetics were often described as factors that trigger melasma, post-inflammatory hyperpigmentation (PIH), ephilides, lichen planus pigmentosus (LPP), and Rehl's melanosis. In our investigation, we discovered a familial history of pigmentary disorders, including melasma, postinflammatory hyperpigmentation (PIH), and ephilides. Thirteen percent of patients had thyroid problems. In our analysis, post-inflammatory hyperpigmentation accounted for 13% of cases with changed face pigmentation, making it the second most prevalent cause. There was a small majority of females. The primary cause in the majority of cases was acne vulgaris. In our research, 14% of patients had a family history of PIH.

Table1: Distribution according to age

| Age group(in years) | Frequency | % |
|---------------------|-----------|----|
| <10 | 4 | 4 |
| 10to 20 | 11 | 11 |
| 20to 30 | 32 | 32 |
| 30to 40 | 24 | 24 |
| 40to 50 | 17 | 17 |
| >50 | 12 | 12 |

Table2: Distribution according to gender

| Gender | Frequency | % |
|--------|-----------|----|
| Male | 25 | 25 |
| Female | 75 | 75 |

Table 3: Distribution according to types of facial melanosis.

| Types official melanosis | Frequency | % |
|--------------------------|-----------|----|
| Melasma | 52 | 52 |
| PIH | 15 | 15 |
| Ephilides | 7 | 7 |

| LPP | 5 | 5 |
|---------------------|---|---|
| Rehl'smelanosis | 6 | 6 |
| Drug induced | 6 | 6 |
| Naevus | 3 | 3 |
| Contactdermatitis | 4 | 4 |
| Acanthosisnigricans | 1 | 1 |
| Others | 1 | 1 |

PIH: post inflammatory hyperpigmentation

Table4: Clinical characteristics of facial hyperpigmentation.

| Types of facial hyperpigmentation | Predisposing factors | Associated co morbid condition | Family history(%) |
|-----------------------------------|--|-----------------------------------|-------------------|
| Melasma | Sunlight, cosmetics, pregnancy | Hypothyroidism | 20 |
| PIH | Sunlight, dermatitis, pyoderma, trauma | Anemia | 14 |
| Ephilides | Sunlight | | 35 |
| LPP | Cosmetics | - | - |
| Rehl's melanosis | Sunlightand cosmetics | - | - |
| Druginduced | ATT | - | - |
| Naevus | - | - | - |
| Contact dermatitis | - | - | - |
| Acanthosisnigricans | - | Diabetes | - |
| Others | - | - | - |

DISCUSSION

We included a total of 100 participants diagnosed with face hyperpigmentation in our research. We included a total of 100 participants diagnosed with face hyperpigmentation in our research. Among the 100 patients, the largest proportion, namely 32%, belonged to the age group of 20 to 30 years, while the next largest proportion, 24%, belonged to the age group of 30 to 40 years. 17% of the individuals belonged to the age range of 40 to 50 years. The youngest individual belonged to the age group of under 10 years. The average age of the participants in the research was 29.01±3.25 years. Among the 100 patients, the majority, namely 75%, were females, while just 25% were men. Hassan et al comprised of patients of altered facial pigmentation. 208 ⁷Theyoungestpatient was a4-year-oldmale, and the oldest was 58-year-old female, with a meanage of 27.40 years. The maximum number of patients thatis, 118(56.73%) belonged to 21 to 40 years age group, followed by54 (25.96%)to <20yearsand 36(17.30%) to>40 years of age group. There were 71 males and 137females, with a female to male ratio of 1.92:1.Commonly observed facial hyperpigmentation type wasmelasma in our study i.e. 52%. It is followed hyperpigmentation in flammatory by post (PIH)in15% and ephilidesin7%. Rehl'smelanos is and druginduced mela no si s was seen in 6% each of thepatients. The average age of melasma patients was 31.78 years inour study, which was similar to 33.45 years in a study byAchar et al.⁸ It is against 42.3 years reported in a study from Singapore.9Wefound about 25% involvementofmen. It is comparable to 19.87% and 10% in differentstudies.8,10 About 70% of our patients with melasma described sunexposurea sex aggerating factor, similartopreviousstudies.¹¹ Thyroid

function seenin13% of patients, dys was hypothyroidism being commonest which was comparabletoprevious studies.8In 69% of patients, there was a history of association with the application no fvariouscosmeticproducts and to picalsteroids, available as over the counterfairness creams, leading to typical steroid facies. This associationof melasma with these cosmetic products has also beenreportedby etal and Grimes.^{8,12}Post-in flammatory Achar hyperpigmentation was the second most common cause of altered facial pigmentation in our study i.e. 13%. It showed a slight female predominance. Most common etiology was secondary to acne vulgaris. This finding was similar to a study by Taylor et al who evaluated acne in skin of colour and found that 65.3% f A frican -American, 52.7% of Hispanicand 47.4% of A sian patients developed acneinduced PIH.¹³PIH family history was seen in 14% of patients in ourstudy. Ranu et al and Sheth et al reported 42.2% and 63% patients with appositive family history of POH respectively.14,15PIH was the second most common cause of altered facialpigmentation in our study i.e. 15%. Acanthosis nigricansis characterized by dark, coarse, thickened skin with avelvetytexture.

CONCLUSION

The most often reported age group was between 20 to 40 years old, with a higher proportion of females. The most frequently detected types of face hyperpigmentation were melasma (52%), post-inflammatory hyperpigmentation (15%), and ephilides (7%).

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