

Clinico-Epidemiology Study of Facial Hypermelanosis at Griya Satya Clinic in January 2017-December 2019

Satya Wydy Yenny¹, Yosep Prabowo²

^{1,2}Andalas University, Medical Faculty, Department of Dermatovenereology / Dr M. Djamil Hospital PerintisKemerdekaan, Padang, West Sumatra 25171, Indonesia

¹satyawidyayenny[at]med.unand.ac.id

²yosepina.dr2[at]gmail.com

Abstract: ***Background:** Hyperpigmentation is a condition characterized by the accumulation of pigment in the epidermis and dermis. Facial pigmentations are the most cosmetically important. They are common in middle-aged women, and are related to endogenous (hormones) and exogenous factors (such as use of cosmetics and perfumes, and exposure to sun radiation). Melasma is a hypermelanosis condition that often causes cosmetic disturbances in a person, so it is necessary to establish an appropriate diagnosis and management. **Objective:** The aim of this study was identify clinico-epidemiology of patient with facial hypermelanosis at Griya Satya Clinic in January 2017-December 2019. **Method:** The study was done with retrospective design by collecting data from medical records of patient with facial hypermelanosis at Griya Satya Clinic in January 2017-December 2019. All patients were diagnosed based on anamnesis and clinical presentation. **Result:** In this study, all patients with facial hypermelanosis were female. There were 133 patients (13.54%) of all female patients who visited during the study period. This type of facial hypermelanosis is the most common were melasma 87 patients (65.41%), post inflammatory hyperpigmentation (PIH) 38 patients (28.57%), lentigo solaris 5 patients (3.76%), and freckles 3 patients (2.26%). **Conclusion:** Facial hypermelanosis is a clinical feature of a diverse group of disorders, the most common of which is melasma. Enforcement diagnosis and appropriate treatment are needed in the management of melasma, as well as proper education to the patient because of this melasma is a chronic and difficult to treat disease.*

Keywords: Facial Hypermelanosis, Melasma, Postinflammatory Hyperpigmentation, Retrospective Study

1. Introduction

The colour of normal skin comes from a mixture of pigments, of which the predominant is melanin. Disorders of hyperpigmentation comprise a large group of skin conditions characterized by an increase in melanin production, increase in density of active melanocytes, abnormal melanin distribution and/or deposition of exogenous pigments.¹ Facial melanoses are commoner in Fitzpatrick skin types III and IV and form a major portion of patients visiting a dermatologist. Though the pathogenesis is not clearly understood in many cases, both light and photosensitizing chemicals seem to play an important role.²

Melasma is a disease hypermelanosis is most commonly encountered and usually found on the face most frequent sun exposure. Pathogenesis of this disease is not very clear, but the effects genetic and hormonal combined with UV radiation plays a very important role as originator. This disease is rarely reported on people who have not yet puberty and are seen more often in women especially in their reproductive age, but it can also be about teenagers, parents who are on medication, and sometimes on men who are usually idiopathic.³ This study was conducted to identify the clinic-epidemiology profile of patient with facial hypermelanosis at Griya Satya Clinic Padang period between January 2017 to December 2019.

2. Method

This retrospective study design was done by collecting data from the medical records from all patient diagnosed facial hypermelanosis at Griya Satya Clinic between January 2017 to December 2019. All the patients were diagnosed

depending on the anamnesis and the clinical finding. All the patient were classified by sex, age, and diagnosed.

3. Result

There were total of 1419 patients who visited the Griya Satya Clinic from January 2017-December 2019, 437 males (30,80%) and 982 females (69,20%). In this study, all patients with facial hypermelanosis were female. There were 133 patients (13.54%) of all female patients who visited during the study period. This type of facial hypermelanosis is the most common were melasma 87 patients (65.41%), postinflammatory hyperpigmentation (PIH) 38 patients (28.57%), lentigo solaris 5 patients (3.76%), and freckles 3 patients (2.26%).

Table 1: Prevalence of patients visiting Griya Satya Clinic between January 2017 and December 2019

Year	Male	Female
2017	156	381
2018	159	345
2019	122	256
Percentage (%)	30,80%	69,20%

Table 2: Clinico-epidemiology profile of facial hypermelanosis between January 2017 and December 2019

No	Clinical Profile	Patient (n)	Percentage (%)
1	Type of Facial Hypermelanosis		
	Melasma	87	65,4
	PIH	38	28,57
	Lentigo solaris	5	3,76
	Freckles	3	2,26
2	Age		
	Melasma		

Volume 9 Issue 12, December 2020

www.ijssr.net

Licensed Under Creative Commons Attribution CC BY

	Young adult	: 15-24	-	-
	Adult	: 25-64	86	98,85
	Elderly	: >65	1	1,15
	PIH			
	Young adult	: 15-24	26	68,42
	Adult	: 25-64	12	31,58
	Elderly	: >65	-	-
	Lentigo Solaris			
	Young adult	: 15-24	-	-
	Adult	: 25-64	5	100
	Elderly	: >65	-	-
	Freckles			
	Young adult	: 15-24	2	66,67
	Adult	: 25-64	1	33,33
	Elderly	: >65	-	-

4. Discussion

Facial hypermelanosis carries immense psychological impact owing to its evident cosmetic disfigurement and social stigmas attached to it. Moreover, people of Asian and African descent constitutively have a darker phenotype which is more vulnerable to pigmentation. Facial hypermelanosis encompasses a myriad of clinical entities which are commonly encountered: melasma, peri-orbital melanosis, postinflammatory hyperpigmentation (PIH), lichen planus pigmentosus, Riehl's melanosis, freckles and lentigenes, exogenous ochronosis, acanthosis nigricans, erythema dyschromicum perstans, and uncommonly poikiloderma of Civatte, erythromelanosis follicularis of face and neck, nevus of Ota.^{4,5}

Other conditions that may result in abnormal pigmentation of the facial region include Mongolian spots, late-stage failure of cardiopulmonary or renal system and drug/heavy metal induced pigmentation such as with iron, silver, gold, chloramphenicol, tetracycline, amiodarone, pirfenidone, antimalarials and antipsychotics.⁶ Systemic disorders including Addison disease, haemochromatosis and porphyria cutanea tarda can also lead to abnormal pigmentation of face.⁷ The diagnosis and differentiation of these conditions are based on history and clinical examination supplemented in some cases by Wood's lamp and histopathological evaluation.

Exposure to ultraviolet rays have been found to be crucial in various studies as it leads to increased levels of various hormones: α -melanocyte-stimulating hormone, corticotrophin and also interleukin (IL)-1 that, in turn, results in increased melanin production. In PIH, skin inflammation results in the production of cytokines, prostaglandins, leukotrienes which further stimulate melanin synthesis. The most common cause attributed was acne. Multiple other studies have also reported similar findings.⁸

Most of the studies worldwide on facial hypermelanosis have reported a female predominance. Number of females with facial hyper melanosis is seeing an upwards trend among the out patients presenting to dermatologists and various contributing factors are: increased awareness, society and marriage pressures to look more beautiful, use of drugs and cosmetics, increased sun exposure and rising obesity and other hormonal abnormalities due to changing life styles.^{4,5} This is consistent with this study, that the facial

hypermelanosis most often found is melasma. In this study it was also found that of all patients who came for treatment with facial hypermelanosis at the Griya Satya Clinic were female and most were in the age range 25-64 years. This may have something to do with the influence of hormones and exposure to ultraviolet rays during his life. In this study, PIH was the second most common type of facial hypermelanosis found. The cause of PIH in this study was due to previous acne and the age range of 15-24 years was mostly. This is probably due to the increased activity of the sebum glands in that age range.

Although hyperpigmentation is typically not harmful, it can cause deleterious emotional and psychological impact on the health-related quality of life of affected individuals. Special considerations when evaluating individuals with skin of color with facial hyperpigmentation can improve both cutaneous disease and quality of life.⁹

Treatment protocols of different hyperpigmentary disorders depend on the cause and site of pigment present. Epidermal hyperpigmentation responds to bleaching creams, peeling, fractional ablative and non-ablative lasers, or intense pulsed light. Sun avoidance, sun protection (caps, umbrella), and sunscreens against UVA and UVB are essential in all cases. Treating the underlying condition helps to stop the progress of the disease and may lead to decreased pigmentation. When using topical and oral therapies, the duration of treatment usually takes 12 weeks to show results. With energy-based devices (lasers and IPL), it requires multiple sessions spaced 2, 4, or 8 weeks apart, according to different regimens.¹⁰

5. Conclusion

Facial hypermelanosis is a condition that more commonly affects individuals with skin of color. Hyperpigmentation, specifically occurring on the facial areas, can have deleterious effects on quality of life. Unfortunately, current treatment protocols are typically not curative and have limited efficacy. Sun-protective measures are the mainstay of both prevention and treatment, whereas a multitude of other modalities, including both non-invasive and invasive techniques, can be used depending on the etiology of dyspigmentation. The incidence of facial hypermelanosis in Griya Satya Clinic was 133 patients from January 2017-December 2019 with melasma was the most common type. More studies should be done to know the incidence of facial hypermelanosis in other hospital in Indonesia.

References

- [1] Bastonini E, Kovacs D, Picardo M. Skin pigmentation and pigmentary disorders: focus on epidermal/dermal cross-talk. *Annals of dermatology*. 2016;1;28(3):279-89.
- [2] Hassan I, Aleem S, Bhat YJ, Anwar P. A clinicoepidemiological study of facial melanosis. *Pigment Int*. 2015;2:34-40.
- [3] Rodrigues M, Pandya AG. Hypermelanoses. *Fitzpatrick's Dermatology*. 9th edition. United States of America: The McGraw-Hill Companies. 2019; 77:1351-89.
- [4] Bhatia V, Bubna AK, Subramanyam S, Veeraraghavan M, Rangarajan S, Sankarasubramanian A. Stubborn facial

- hypermelanosis in females: A clinicopathologic evaluation. *Pigment Int* [serial online]. 2016;3:83-9.
- [5] Cheng J, Vashi NA. Treatment strategies for hyperpigmentation. *Dermatoanthropol Ethnic Skin Hair*. 2017;4:417-36.
- [6] Abdel-Naser MB. The color of skin: gray diseases of the skin, nails, and mucosa. *Clin Dermatol*. 2019;37:507–15.
- [7] Bagherani N, Gianfaldoni S, Smoller BR. An overview on melasma. *J Pigment Disord*. 2015;2:218.
- [8] Nouveau S, Divya A, Kohli M, Bernerd F, Misra N, Nayak CS. Skin hyperpigmentation in Indian population: Insights and best practice. *Indian J Dermatol*. 2016;61:487-95.
- [9] Ikino JK, Nunes DH, Silva VP, Frode TS, Sens MM. Melasma and assessment of the quality of life in Brazilian women. *An Bras Dermatol*. 2015;90(2):196–200.
- [10] Chang MW. Disorders of hyperpigmentation. Chapter 67. In: Bolognia JL, Jorizzo JL, Schaffer JV, editors. *Dermatology*. 3rd ed. Amsterdam: Elsevier; 2012. p. 1049–74.

Author Profile



Satya WydyaYenny received the dermatologist degrees in Medical Faculty, Indonesia University. She now as the head of Dermatology Residency Program in Andalas University

Yosep Prabowo a resident of dermatology and venereology, Medical Faculty, Andalas University