Gazelle Eye like Facial Melanosis (Clinico-Histopathological Study)

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Abstract

**Background:** There are many causes of facial melanosis like melasma, frictional melanosis, acanthosis nigricanis, lichen planus actinicus, and others. But Gazelle eye like facial melanosis was not recognized as a special disease with characteristic features.

**Objective:** To report a new recognized facial pigmentation that is not uncommonly seen among adults especially females which is locally known as a Gazelle eye like facial melanosis.

**Patients and methods:** This case series descriptive study with histopathological examination was conducted in the Department of Dermatology, Baghdad Teaching Hospital, Baghdad, Iraq during the period from January 2009-December 2013. One hundred patients with gazelle eye like facial pigmentation were collected and evaluated by clinical and histopathological examinations. History was obtained from each patient regarding all socio-demographic aspects related to the disease. Also, clinical assessment was done including Wood’s light examination. Incisional or punch biopsy was performed from 10 patients for histopathological examination.

**Results:** One hundred patients were recorded and examined: 88(88%) females and 12(12%) males with a female to male ratio: 6.5:1. Their ages ranged from 16-48 (28.7 ± 5) years. The duration of the disease ranged from 5-15(6 ± 3.2) years. All patients had characteristic pigmentation that started around the eyes in a symmetrical fashion and descended down to the cheeks which ended with a tail like and there was a well demarcated white band across the cheek’s pigmentation. The pigmentation was dark brown in color that is not delineated from surrounding normal face skin with negative Wood’s light examination. Histopathological study revealed mainly dermal melanosis as many melanophages were seen in the superficial dermis with basal melanosis of the epidermis. On systemic review all patients were apparently healthy apart from facial melanosis as the main complaint.

**Conclusions:** We think this is a new entity that commonly seen in the clinical practice as a cause of facial melanosis but it’s not well recognized. It has a characteristic location, distribution and configuration that deserve the name “Gazelle eye like facial melanosis”.

**Keywords:** Gazelle eye; Facial melanosis; Pigmentation; Dermal melanosis

Introduction

Facial melanosis is a major cosmetic problem among people especially in people dark complexion [1,2], and from daily clinical practice we can categorize the commonest causes of facial melanosis as follow [1-3]:

1- **Melasma:** It is probably the commonest cause of facial pigmentation and it has a characteristic light to dark brown color with stuck up on appearance [1-8]. It could be defined as chronic disfiguring dermatological disease of females and males although much more common among females that appear during the active reproductive period with the presence of provoking factors like hormonal changes and sunlight that induce brown hyperpigmentation of the face. This may take many shapes but much more commonly a butterfly and a mask appearance [6-13].

2- **Frictional Melanosis:** As there is a common habit among people to do rubbing and friction of their face in order to induce whitening of the skin [14] but overtime there is a reactionary hypermelanosis of the epidermis and dermis that appeared clinically as a deep dark brown in color [14,15]. The pigmentation localized mainly over the bony areas of the face usually on the temples, forehead, and cheeks and might resemble melasma [16-18]. It affects young females and males and appears as a dark brown pigmentation with violaceous hue which is distributed on the butterfly area of the face: cheeks, nose, around the eye brows with or without scattered typical papules of lichen planus [1-3]. While the histopathological picture is typical for lichen planus like: basal liquefaction, band like lymphocytic infiltrations of the dermal layer with melanophages [1-3,16-18].

3- **Lichen Planus Actinicus (butterfly like pigmentation):** Is also one of the commonest causes of facial melanosis especially in chronic cases and might resemble melasma [16-18]. It affects young females and males and appears as a dark brown pigmentation with violaceous hue which is distributed on the butterfly area of the face: cheeks, nose, around the eye brows with or without scattered typical papules of lichen planus [1-3]. While the histopathological picture is typical for lichen planus like: basal liquefaction, band like lymphocytic infiltrations of the dermal layer with melanophages [1-3,16-18].

4- **Acanthosis Nigricans:** Acanthosis nigricans of the face which could be localized to the face only or part of the generalized ordinary acanthosis nigricans. The skin is darkly pigmented thickened
hyperkeratotic that usually affects the cheeks, around the eyes and forehead with or without skin tags. The histopathological features are that of acanthosis nigricans [19-24].

5-Postinflammatory Hyperpigmentation: Any inflammatory skin lesion that affects the face can be complicated by hypermelanosis as it resolves like: eczema, contact dermatitis, acne, lupus erythematosus etc. [1-3,13,25].

6-Miscellaneous: Phytodynamic dermatitis, Riehls melanosis, Naevus of Ota, drug induced hyperepigmentation, and others [1-3,25-29].

In addition there is not uncommon facial pigmentation that is locally known as Gazelle eye like facial melanosis that is commonly encountered in clinical practice but often misdiagnosed and commonly confused with melasma and does not respond to ordinary therapy options for melasma [1-3,6,8-14].

And after extensive searching in the medical literatures, we didn’t find a report about this type of facial pigmentation.

For this reason, the present work was arranged to evaluate this new unrecognized cause of facial pigmentation by doing clinical and histopathological studies.

Patients and methods

This is a case series descriptive and histopathological study that was conducted in the Department of Dermatology, Baghdad Teaching Hospital, Baghdad; Iraq during the period from January 2009-October 2013. One hundred patients with Gazelle eye like facial pigmentation were included in this work.

History was obtained from each patient regarding all socio-demographic aspects related to the disease including: age, age of onset, gender, duration of disease, previous medical history, drugs ingestion, hair epilation, history of friction, seasonal variation, menstrual cycle, marital status, pregnancy and hormonal intake, history of using sun block agents, marital status, occupation and family history of the same condition.

Clinical assessment was done regarding the following points: site, symmetry, border, color, size of lesions and morphological picture. Systemic evaluation for all patients was performed to exclude any underlying internal causes. All patients were examined with Wood’s light to differentiate whether it is dermal or epidermal or mixed melanosis. All patients had Fitzpatrick’s skin type III and IV. Incisional or punch biopsy was performed from 10 patients for histopathological evaluation after processing with Hematoxyllin and Eosin and Fontana masson stains.

Formal consent was taken from each patient after full explanation about the goal and nature of the present study, the nature of the disease, course, and the need for photographs. Also, ethical approval was performed by the Scientific Council of Dermatology and Venereology-Iraqi Board for Medical Specializations.

All patients were photographed by Sony-Cyber Digital, high sensitivity, 16.1 megapixels, 5 x optical zoom camera, in the same place with fixed illumination and distance.

Results

A total of 100 patients with periorbital hypermelanosis with cheek extension (Gazelle eye like facial pigmentation) were seen. There were 88(88%) females and 12(12%) males with a female to male ratio: 6.5:1. Their ages ranged from 16-48 years with a mean ± SD of 21.2 ± 4.4 years. The duration of the disease ranged from 5-15 years with a mean ± SD of 8 ± 3.2 years. While the age of onset was ranged from 13-40 years with a mean ± SD of 21.2 ± 4 years.

Regarding the age of onset of the disease were as follow: in 40(40%) patients between 10-19 years, 43(43%) patients between 20-29 years while in 17(17%) patients developed between 30-39 years. While the age distribution of patients was as follow as follow: 20(20%) patients had the disease between 10-19 years, 43(43%) between 20-29 years, 20(20%) patients between 30-39 years while in 17(17%) had between 40-50 years.

Family history of the same condition was positive in 37(36%) patients (4 males and 33 females). Negative history of the following factors: friction, drugs or pregnancy, oral contraceptive pills, seasonal variation. No effect of the hair epilation and using sun block agents in remission or exacerbation of the condition. No difference regarding the aggravating factors between married versus unmarried females. Also On systemic review all patients were apparently healthy apart from facial pigmentation of face as the main complaint of these patients.

Many of assumed provoking factors such as: fatigue was seen in 86(86%) patients, sunlight in 70(70%) patients.

All patients mentioned that the disease was recalcitrant to all standard optional therapies for melanosis like bleaching agents.

All patients had characteristic pigmentation that started around the eyes periorbital hyperpigmentation in very close symmetrical fashion and descended down to the cheeks which ended with a pointed tail like configuration. Also, there was a well demarcated linear white band across the middle cheek’s pigmentation that divided the pigmentation in two parts (Figures 1-4).

The pigmentation was uniform even and not well demarcated from normal surrounding face skin and it was dark brown in color with negative Wood’s light examination in all cases.

The histopathological study showed basal melanosis in the epidermis and dermal melanosis in a form of numerous melanophages seen in the superficial dermis in the H/E stain (Figure 5A). While the Fontana masson stain section showed also basal melanosis of the epidermis and dermal melanosis (Figure 5B).

Figure 1: Thirty five years old female with Gazelle eye like facial melanosis with characteristic periorbital pigmentation and cheeks extension.
The present study had shown that this entity has many characteristic features that could be differentiated from other causes of facial melanosis:

1. It has characteristic distinguishing location mainly the periorbital region and cheeks in a symmetrical fashion.

2. In contrast with what has been mentioned about periobital pigmentation only, gazelle eye like mostly occurs below ten years and increases in puberty and with age [1-3,13].

3. It has distinctive morphological features as it has a well demarcated linear band of hypopigmentation in the middle of the cheeks pigmentation and it extends downward as a pointed tail like configuration.

Figure 2: Twenty five years old male with Gazelle eye like facial melanosis.

Figure 3: Thirty eight years old female with Gazelle eye like facial melanosis.

Figure 4: Forty years old female with Gazelle eye like facial melanosis.

Figure 5A: Biopsy from patient with Gazelle eye like facial melanosis stained by H&E stain, showing basal hypermelanosis with numerous dermal melanophages (original magnification × 40).

Figure 5B: Fontana masson stain, showing basal and superficial dermal melanosis (original magnification × 40).

Discussion

Gazelle eye like facial melanosis although is not un common pigmentory problems among Iraqi population but unfortunately often confused and mixed up either with melasma especially dermal type [1-13] or with frictional melanosis [13-15].
4. The pigmentation was always dark-brown with no distinct demarcation from the surrounding normal face. Wood's light examination showed no contrast seen in all patients which does indicate dermal melanosis.

5. The histopathological pictures are mostly dermal melanosis.

6. There are no hormonal effects by menstrual period, pregnancy and contraceptive pills.

7. It is often not affected by sunlight exposure like melasma and lichen planus actinicus and doesn’t respond to routinely topical bleaching therapies [16-18,29].

The aetiopathogenesis of this pigmentation could not be well elaborated but as this disease appears in adult life, in addition to positive family history among other family member, these might suggest that this disease could have genetic and racial elements involved in its pathogenesis.

The management of this type of facial pigmentation is often very difficult as all bleaching agents might give only temporary relieving effects [1-3,6-14,16].

Accordingly, we suggest doing deep dermabrasion as an effective mode of therapy for this condition similar to the dermabrasion that could be carried out for dermal melasma which is resistant to the standard treatments [29].

Also, one of the targets of this work is to stimulate other dermatologists in other countries to observe this type of pigmentation and to be reported.

In conclusion, Gazelle eye like facial melanosis is a distinctive clinical and pathological entity that has many characteristic features.

References