A Clinicopathological Study of Spectrum of Pigmented Skin Lesions in Southern India: A Three Year Experience at a Tertiary Care Centre with Review of Literature

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Abstract

Pigmented skin lesions refer to lesions that are black, brown or blue in color. These lesions include both melanocytic and non-melanocytic lesions. A number of pigmented lesions are difficult to classify because of wide spectrum of histological appearances and raise the possibility of melanoma. With this study we intended to evaluate the spectrum of pigmented skin lesions and to correlate the clinical diagnosis with the histological diagnosis. In this retrospective study, 75 cases of pigmented skin lesions were reviewed on hematoxylin and eosin stained paraffin embedded tissue sections from June 2017- May 2020 in the Department of Pathology at a Tertiary Care Hospital and analyzed according to age, gender, site of occurrence and histological types. Out of the 75 cases evaluated there were 23 melanocytic lesions and 52 non-melanocytic lesions. Overall, benign melanocytic nevi (13 cases) were commonest lesions followed by seborrheic keratosis (11 cases). The lesions presented from 1st to 9th decade with slight female predominance. The most common site involved was head and neck. Clinicohistopathological correlation showed positive correlation in 55(73.3%) cases and negative correlation in 20 cases (26.6%). Pigmented skin lesions are common presenting problem, while majority are benign a small minority can be malignant. A histological interpretation by pathologist is essential to correctly diagnose these lesions in order not to miss a small percentage of malignant tumors and to differentiate melanocytic lesions from its nonmelanocytic mimickers.

Keywords: Pigmented skin lesions, Benign melanocytic nevi, Malignant melanoma, Seborrheic keratosis, Basal cell carcinoma, Dermatofibrosarcoma protuberans.

INTRODUCTION

Pigmented skin lesions (PSLs) refer to lesions that are black, brown or blue in color [1]. These lesions include both melanocytic and non-melanocytic lesions. A number of pigmented lesions are difficult to classify because of wide spectrum of histological appearances and raise the possibility of a melanoma [2]. Melanocytic lesions can develop from abnormal migration of melanocytes from neural crest to the skin during embryogenesis, impaired melanosome transfer to keratinocytes and alteration in melanin synthesis [3]. Melanocytic proliferations include the three related cell types: melanocytes, nevus cells or melanoma cells, each of which may be present in epidermis or dermis and infrequently in subcutis. Benign neoplasms of nevus cells are called melanocytic nevi, while malignant neoplasms are called malignant melanoma [4]. Non-melanocytic lesions can be keratinocytic, vascular, or reactive [1]. Some of these include seborrheic keratosis, basal cell carcinoma, actinic keratoses, dermatofibrosarcoma protuberans, lichen planus pigmentosus, and vasoformative lesions. A careful histopathological evaluation is essential in making the correct diagnosis of the PSL. The study was conducted to understand the spectrum of pigmented skin lesions with reference to age, sex, site and histopathological diagnosis in Southern India.

MATERIAL AND METHODS

This was a retrospective study of the histologically diagnosed pigmented skin lesions over a
span of three years from June 2017 to May 2020. Out of 1108 skin biopsies received over the three years, 75 clinically and histologically diagnosed PSLs were selected for the study. Data including demographic details, clinical diagnosis and clinical evidence of pigmentation were retrieved from histopathology requisition forms and patient case sheets. The hematoxylin & eosin stained slides of skin biopsies were reviewed and analyzed according to age, gender, site of occurrence and histological types.

RESULTS
In the present study, out of 1108 skin biopsies received, 6% (75) were diagnosed as PSLs. These 75 PSLs were further categorized, into melanocytic 23 (30.6%) and non-melanocytic 52 (69.4%) pigmented lesions (Table-1).

| Lesions                        | No of cases (n=75) (%) |
|--------------------------------|-----------------------|
| **Melanocytic Lesions**        |                       |
| Benign melanocytic nevi        | 13 (17.3)             |
| Malignant Melanoma             | 10 (13.3)             |
| **Non Melanocytic Lesions**    |                       |
| Dermatofibrosarcoma protuberans| 2 (2.7)               |
| Keratoacanthoma                | 1 (1.3)               |
| Post Inflammatory hyperpigmentation | 5 (6.7)              |
| Dermatofibroma                 | 10 (13.3)             |
| Seborrheic keratosis           | 11 (14.7)             |
| Basal cell carcinoma           | 7 (9.4)               |
| Lichen planus pigmentosus      | 8 (10.7)              |
| Lobular Capillary Hemangioma   | 4 (5.3)               |
| Cavernous Hemangioma           | 3 (4)                 |
| Angiokeratoma                  | 1 (1.3)               |
| **Total**                      | 75 (100)              |

Out of these, 31 (41.3%) cases were males, 44 cases (58.7%) were females, with male: female ratio of 1:1.4 (Table-2).

| Lesions                        | Males (%) | Females (%) |
|--------------------------------|-----------|-------------|
| **Melanocytic Lesions**        |           |             |
| Benign Melanocytic nevi        | 6 (8)     | 7 (9.3)     |
| Malignant Melanoma             | 4 (5.3)   | 6 (8)       |
| **Non Melanocytic Lesions**    |           |             |
| Dermatofibrosarcoma protuberans| -         | 2 (2.7)     |
| Keratoacanthoma                | 1 (1.3)   | -           |
| Post- Inflammatory hyperpigmentation | 4 (5.3)   | 1 (1.3)     |
| Dermatofibroma                 | 4 (5.3)   | 6 (8)       |
| Seborrheic keratosis           | 3 (4)     | 8 (10.7)    |
| Basal cell carcinoma           | 3 (4)     | 4 (5.3)     |
| Lichen planus pigmentosus      | 4 (5.3)   | 4 (5.3)     |
| Lobular capillary hemangioma   | 2 (2.7)   | 2 (2.7)     |
| Cavernous hemangioma           | -         | 3 (4)       |
| Angiokeratoma                  | -         | 1 (1.3)     |
| **Total**                      | 31 (41.3) | 44 (58.7)   |

The age of the patients ranged from 4 months old – 84 years Maximum cases presented in 3rd decade of life (21.3%). The commonest lesions in the third decade being, benign melanocytic nevi followed by lichen planus pigmentosus (Table-3).

Also, benign melanocytic nevi were the most common among both sexes (17.3%) followed by seborrheic keratosis (14.7%) (Table-2). Of the 13 benign nevi, there were 9 (69.2%) cases of intradermal nevi, 3 (23.1%) cases of compound nevi and 1 (7.7%) case of congenital melanocytic nevus. There were 10 (13.3%) cases of malignant melanoma with slight female predilections (Table-2) with increased occurrence in 5th, 6th and 8th decades (Table-3). Among the 53 non melanocytic pigmented lesions, seborrheic keratosis was most common 11 (20.7%) followed by dermatofibroma 10 (18.8%). Both showed increased incidence in females (Table-2).
Most commonly involved site was the head and neck region including the face, 29(38.6\%) (Table 4a & 4b). While the benign melanocytic lesions were distributed predominantly over head, face, abdomen and back, lesions of malignant melanoma showed variable distribution (Table-4a).

| Lesions                                | <21yr | 21-30yr | 31-40yr | 41-50yr | 51-60yr | 61-70yr | >70yr |
|----------------------------------------|-------|---------|---------|---------|---------|---------|-------|
| Benign melanocytic nevi (n=13) (%)     | 2(15.3)| 7(53.8)| 1(7.7)  | 1(7.7)  | 1(7.7)  | 1(7.7)  | -     |
| Malignant melanoma (n=10) (%)          | -     | -       | -       | 1(10)   | 5(50)   | -       | 4(40) |
| Dermatofibrosarcoma protubersans (n=2) | 1(50) | -       | 1(50)   | -       | -       | -       | -     |
| Keratoacanthoma (n=1) (%)              | -     | -       | -       | -       | 1(100)  | -       | -     |
| Post inflammatory hyperpigmentation (n=6) (%) | 1(20) | 1(20)   | 1(20)   | 1(20)   | 1(20)   | 1(20)   |
| Dermatofibroma (n=10) (%)              | 1(10) | 2(20)   | 4(40)   | 2(20)   | --       | 1(10)   |
| Seborrheic keratosis (n=11) (%)        | -     | 1(9.1)  | -       | 3(27.3) | 4(36.4) | 2(18.2) | 1(9.1) |
| Basal cell carcinoma (n=7) (%)         | -     | -       | -       | 1(14.3) | 1(14.3) | -       | 5(71.4)|
| Lichen planus pigmentosus (n=8) (%)   | 1(12.5)| 3(37.5)| 2(25)  | 2(25)   | -       | -       | -     |
| Lobular capillary hemangioma (n=4) (%) | -     | 2(50)   | 1(25)   | -       | -       | 1(25)   | -     |
| Cavernous hemangioma (n=3) (%)         | -     | 1(33.33)| 1(33.33)| 1(33.33)| -       | -       | -     |
| Angiokeratoma(n=1) (%)                 | -     | -       | -       | -       | 1(100)  | -       | -     |

Table-4a: Distribution of melanocytic lesions on various sites

| Site          | Benign Melanocytic nevi (n=13) (%) | Malignant Melanoma (n=10) (%) |
|---------------|------------------------------------|-------------------------------|
| Eyebrow       | 1(7.7)                            | -                             |
| Eyelid        | 2(15.3)                           | -                             |
| Forehead      | 1(7.7)                            | -                             |
| Ala of nose   | 2(15.3)                           | -                             |
| Cheek         | 1(7.7)                            | -                             |
| Chin          | 1(7.7)                            | -                             |
| Temporal area | 1(7.7)                            | -                             |
| Scalp         | 1(7.7)                            | -                             |
| Neck          | -                                 | 1(10)                         |
| Back          | 1(7.7)                            | 1(10)                         |
| Abdomen       | 1(7.7)                            | 2(20)                         |
| Shoulder      | 1(7.7)                            | 3(30)                         |
| Labia         | -                                 | 1(10)                         |
| Sole          | -                                 | 2(20)                         |
| Total         | 13                                | 10                            |

Table-4b: Distribution of non melanocytic lesions on various sites

| Site          | DFSP (n=2) (%) | KA (n=1) (%) | PIH (n=5) (%) | DF (n=10) (%) | SK (n=11) (%) | BCC (n=7) (%) | LPP (n=8) (%) | LCH (n=4) (%) | CH (n=3) (%) | AK (n=1) (%) |
|---------------|---------------|-------------|--------------|--------------|--------------|--------------|--------------|--------------|-------------|--------------|
| Forehead      | -             | -           | -            | -            | 1(14.3)      | -            | -            | -            | -           | -            |
| Ala of nose   | -             | -           | -            | -            | 1(9.1)       | 1(14.3)      | -            | -            | -           | -            |
| Cheek         | -             | -           | -            | -            | 1(9.1)       | 1(14.3)      | -            | -            | -           | -            |
| Temporal area | -             | -           | 1(10)        | 1(9.1)       | 3(42.8)      | -            | -            | -            | -           | -            |
| Scalp         | 1(50)         | -           | -            | 2(18.2)      | -            | -            | 1(25)        | -            | -           | -            |
| Ear           | -             | -           | -            | 2(18.2)      | -            | -            | -            | -            | -           | -            |
| Neck          | 1(50)         | 2(40)       | -            | -            | -            | -            | -            | -            | -           | -            |
| Chest         | -             | -           | -            | -            | -            | -            | 1(33.3)      | -            | -           | -            |
| Back          | -             | -           | -            | 1(9.1)       | 2(25)        | -            | -            | -            | -           | -            |
| Abdomen       | -             | 1(20)       | 2(18.1)      | -            | -            | -            | -            | -            | -           | -            |
| Forearm       | -             | -           | 5(50)        | -            | -            | 1(12.5)      | -            | -            | -           | -            |
| Hand          | -             | -           | -            | -            | -            | -            | 2(25)        | 1(25)        | -            | -            |
| Shoulder      | -             | -           | -            | -            | -            | 1(12.5)      | -            | -            | -           | -            |
| Penis         | -             | -           | -            | -            | 1(12.5)      | -            | -            | -            | -           | -            |
| Thigh         | -             | 1(20)       | 1(10)        | 1(9.1)       | 2(28.6)      | -            | -            | -            | -           | -            |
| Knee          | -             | -           | -            | 1(9.1)       | -            | -            | -            | -            | -           | -            |
| Gluteal       | -             | 1(20)       | 1(10)        | -            | -            | -            | -            | -            | -           | -            |
| Leg           | -             | -           | 1(10)        | -            | 1(12.5)      | 2(66.7)      | -            | -            | -           | -            |
| Scrotum       | -             | 1(100)      | -            | -            | -            | -            | 1(100)       | -            | -           | -            |
| Foot          | -             | 1(10)       | -            | -            | 2(50)        | -            | -            | -            | -           | -            |
| Total         | 2             | 1           | 5            | 10           | 11           | 7            | 8            | 4            | 3           | 1            |

DFSP-Dermatofibrosarcoma protubersans, KA – Keratoacanthoma, PIH – Post inflammatory hyperpigmentation, DF –Dermatofibroma, SK – Seborrheic keratosis, BCC- Basal cell carcinoma, LPP- Lichen planus pigmentosus, LCH- Lobular capillary hemangioma, CAVH- Cavernous hemangioma, AK - Angiokeratoma

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Out of 75 cases, 55 (73.3%) cases were consistent with both clinical and histopathological diagnosis, while 20 (26.6%) cases were inconsistent (Table-5).

### Table-5: Clinical and histopathological correlation of pigmented skin lesions

| Lesions                                      | Consistent with clinical diagnosis | Inconsistent with clinical diagnosis |
|----------------------------------------------|------------------------------------|-------------------------------------|
| Benign melanocytic nevi (n=13) (%)           | 11 (84.6%)                         | 2 (15.4%) Seborrhic keratosis        |
| Malignant melanoma (n=10) (%)                | 7 (70%)                            | 1 (10%) Basal cell carcinoma         |
| Dermatofibrosarcoma protrubers (n=2) (%)     | -                                  | 1 (50%) Neurofibroma                |
| Keratoacanthoma (n=1) (%)                    | -                                  | 1 (100%) Squamous cell carcinoma    |
| Post inflammatory hyperpigmentation (n=5) (%) | 5 (100%)                           | -                                   |
| Dermatofibroma (n=10) (%)                    | 5 (50%)                            | 2 (20%) Dermatofibrosarcoma protrubers, 3 (30%) Nevi |
| Seborrheic keratosis (n=11) (%)              | 9 (81.8%)                          | 2 (18.2) Nevi                       |
| Basal cell carcinoma (n=7) (%)               | 5 (71.4%)                          | 2 (28.6%) Nevi                      |
| Lichen planus pigmentosus (n=8) (%)         | 6 (75%)                            | 1 (12.5) Pityriasis rosea, 1 (12.5) Ashy dermatosis |
| Lobular capillary hemangioma (n=4) (%)      | 4 (100%)                           | -                                   |
| Capillary hemangioma (n=3) (%)              | 3 (100%)                           | -                                   |
| Angiokeratoma(n=1) (%)                       | -                                  | 1 (100%) Hemangioma                 |

### DISCUSSION

In our study, the pigmented skin lesions constituted 6% of all skin biopsies received over three years. Majority of them were non-melanocytic pigmented lesions 52 (69.4%) as compared to melanocytic lesions which were 23 (30.6%) (Table-1). Similar findings were reported in study by Prasad et al., [6]. However, in studies by Leishram et al., [5], Bohra et al., [7] and Parvathi et al [8] the incidence of PSLs of melanocytic origin was more. In our study, there was an increased presentation of PMLs in females 44 cases (58.7%) as compared to males 31 (41.3%) with male to female ratio of 1:1.4 (Table-2). Overall, benign melanocytic nevi were the most common lesion 13 (17.3%) which showed increased preponderance among females 7 (9.3%), similar to observations of Leishram et al., [5], Bohra et al., [7], Parvathi et al., [8] and Schafer et al., [9]. Most PSLs presented below 50 years of age. The findings were similar to Leishram et al., [5], Parvathi et al., [8], Mackie et al., [10] and dissimilar with Youl et al., [11]. Most commonly involved site was the head and neck region including the face 29 (38.6%) (Table 4a & 4b) which was in concordance with Leishram et al., [5], Bohra et al., [7] and Parvathi et al., [8]. Malignant lesions constituted 17 (22.6%) of the study which included 10 (13.3%) cases of malignant melanoma and 7 (9.3%) cases of basal cell carcinoma. Clinical and histological concordance was achieved in 55 (73.3%) cases which was comparable with Parvathi et al., [8].

Melanocytic nevi are benign neoplastic proliferation or hamartomas of melanocytes [12]. Melanocytic nevi are not clinically apparent at birth with most nevi appearing in adolescence and early adulthood. Melanocytic nevi are characterized by the presence of nevus cells, which, though being melanocytes, differ from them by at least being partially arranged in clusters or nests, with the propensity to be round rather than having dendritic cell shape and a tendency to retain pigment in their cytoplasm. They follow a rather predictable evolution which is believed to begin as junctional nevi, become compound nevi and then intradermal nevi [13]. Rarely there can be a dramatic event such as spontaneous regression, activation or malignant transformation [14]. They may be papillomatous, pedunculated or flat and are more common in the skin of head, neck and trunk [15]. Histologically they are characterized by small nests or bundles of melanocytes in upper dermis with tendency to cluster around pilosebaceous units [16]. The degree of pigmentation and cellularity may vary considerably with lower half of the lesion being less cellular and less pigmented and more spindled with fibrillar cytoplasm [17] with ‘Wagner -Meissner’ type structures representing expression of neural component of nevus.

In the present study among the melanocytic pigmented lesions, we noted maximum number of intradermal naevi followed by compound nevi and congenital melanocytic nevus that was similar to study by Prasad et al., [6] and Shoko et al., [13]. All intradermal nevi were seen in the 2nd to 5th decade of life. These were distributed over facial region including cheeks, chin, ala of nose, forehead and eyelid varying in size from 0.4 – 1.5cm with flat to papillary lesions presenting with tan to light brown pigmentation. Microscopically small nests of nevus cells were present in upper dermis surrounding pilosebaceous units with focal pigmentation with lower half of the lesion being less cellular and less pigmented and more spindled (Fig-1).

Compound melanocytic nevi show high incidence in 2nd decade [18], are 3-6mm papules, slightly elevated with homogenous light to dark brown pigment and features of both junctional and intradermal nevi. This lesion was diagnosed in three cases, all in 3rd decade of life with distribution over cheek, back and abdomen. The lesions were slightly elevated.
papulonodular, light brown in color varying in size from 1.5 – 3.5 cm. Histologically there were epidermal and dermal component of nevus cells showing maturation towards depth of the lesion (Fig-2). Two of these lesions were clinically diagnosed as seborrheic keratosis because of epidermal hyperplasia over the lesion.

Congenital melanocytic nevi are reported in about 1% of live births [19]. It is hypothesized that morphological error occurs in the neuroectoderm during embryogenesis, leading to unregulated growth of melanoblasts, which are the precursor cells of melanocytes [20]. A single case of congenital melanocytic nevus was reported in our study in a 65 years old male as a dark brown irregular macule over left side of forehead and eyebrow measuring approximately 5 cm x 6 cm in size. It was present since birth. Microscopically, it was characterized by melanocytes clustered around follicles, adnexa, nerves and between collagen fibers at base of lesion.

In our study, we reported ten cases of malignant melanoma. All cases presented in late adulthood from 4th – 9th decade of life and were distributed in both sun exposed and non-sun exposed areas including shoulder, neck, back, abdomen, labia and sole of feet, the latter location particularly being more common in women [21]. Also, there was an overall female predilection similar to observations by Parvathi et al., [8] and Leishram et al., [5]. All lesions varied in size from 2– 3.5 cm with brown black pigmentation and irregular notched borders, unlike smooth round and uniform borders of benign melanocytic nevi (Fig-3). Melanoma is notorious for its great microscopic variability [22]. Histologically all lesions exhibited junctional activity, prominent melanin pigment, marked cytological atypia, large eosinophilic nucleoli, numerous mitotic figures and invasion of surrounding tissue (Fig-3).

Seborrheic keratosis is a benign keratinocytic pigmented skin lesion. It develops from the proliferation of keratinocytes of the epidermis [6]. Its common locations are the face, scalp, trunk, abdomen, back, and upper extremities. Unusual sites are conjunctiva, nipple, and areola [9]. In our study, among the non-melanocytic pigmented lesions, seborrheic keratosis was the commonest with 11 reported cases. Females were most commonly affected. Similar observations were reported by Leishram et al., [5] and Prasad et al., [6]. The most common age group affected was between the 3rd-8th decade. The lesions were brown black in color measuring 0.5–2.6 cm in size and distributed over scalp, face, shoulder, abdomen and lower limb. Microscopically, the lesions exhibited hyperplastic stratified squamous epithelium with multiple horn cysts and melanin pigment (Fig-4). Of the 11 cases, 02 cases were clinically not consistent as they mimicked nevi due to presenting as brownish black lesion occurring on sun exposed parts. In our study, the most common age group reported was in the 6th decade, similar to Parvati et al., [8].

Fig-1: Intradermal nevus showing upper dermis containing nests and cords of nevus cells with variable pigmentation (H&E)

Fig-2: Compound nevus with junctional activity and nests and cords of nevus cells with variable pigmentation (H&E)

Fig-3: Malignant melanoma with junctional activity, melanin pigment and dermal involvement by atypical melanocytes without maturation (H&E)
Basal cell carcinoma is most frequent form of skin cancer primarily occurring on sun exposed skin due to UV exposure. It has been observed that they may also occur uncommonly on sun protected areas like legs in association with venous stasis, trauma, immunosuppression, arteriovenous malformations and X-ray exposure [15]. In our study we reported 7 cases of basal cell carcinoma out of which 5 were distributed over face and 2 cases presented over thigh. All cases, in our study affected individuals between 6th-9th decade with female preponderance. The lesions presented as brown black nodules or papules measuring 1–2.5cm in size. Two cases out of five were clinically misdiagnosed as nevi due to more regular uniform nodular pigmented lesion. Microscopically, proliferating nests of basaloid cells arising from epidermis and extending into superficial dermis were noted with cells showing prominent palisading surrounded by lose myxoid stroma. Focal cleft like spaces around nests of cells were also observed. There were scattered dermal melanophages with intracytoplasmic melanin between tumor nests (Fig-5). Basal cell carcinoma were also reported in studies by Leishram et al., [5], Prasad et al., [6] and Parvathi et al., [8].

We reported 02 cases of dermatofibrosarcoma protuberans (DFSP). Though trunk and extremities are the most common locations [23], the lesions in our study presented on the neck and scalp. Keeping in consistency that they occur commonly in early to mid-adulthood [23], the tumors in our study presented in 3rd and 4th decade as purple red raised lesions measuring 1.4 and 3.5cm each. Microscopically, proliferation of spindle-shaped cells were seen in the deep dermis with focal infiltration in the subcutaneous fat with sparing of the epidermis. The spindled tumor cells were uniform in appearance with elongated nuclei and little or no pleomorphism and were arranged in a storiform manner. Scattered melanophages were noted (Fig-6). Cases of DFSP were also reported by Cresta et al., [24].

Dermatofibromas account for almost 3% of dermatopathology and have a distinct predilection for the extremities, particularly the lower limbs, of young adults with females affected more than the males [25]. In our study, we reported 10 cases of dermatofibroma with females being affected more than males and majority presenting in the 4th decade. The upper limbs were involved commonly followed by lower limbs and unusual presentation of one case on the upper lip. The lesions were polyoidal to flat plaques measuring 0.5cm-1.5cm with a dusky brown color. Microscopically, these were poorly circumscribed tumors mainly located in the dermis with extension into the superficial subcutaneous tissue and composed of a variable admixture of fibroblast-like cells, histiocytes and blood vessels (Fig-7).

We reported 8 cases of lichen planus pigmentosus (LPP). Contrary to the literature that LPP involve sun exposed areas of skin and occur in children and young adults [26], in our study the lesions presented between 4th – 5th decade with no gender predilection and involved back, upper limbs, shoulder and penis. They presented as annular dark brown to violaceous plaques with slightly raised hypopigmented
borders. Histologically, there was hyperkeratosis, atrophic epidermis with vacuolar alteration of the basal layer and scarce lymphohistiocytic to lichenoid infiltrates in the dermis with pigment incontinence (Fig-8). Cases of LPP were also reported by Leishram et al., [5], Parvathi et al., [8] and Mruthyunjayappa et al., [27].

We reported a single case of keratoacanthoma in a 57year old male presenting as a flesh colored raised papule over scrotal skin. Originating in the pilosebaceous unit, keratoacanthomas are derived from an abnormality leading to hyperkeratosis of the infundibulum. They are associated with hair-bearing areas [29]. Considered by some to be a highly differentiated form of squamous cell carcinoma, histological examination revealed a circumscribed proliferation of well-differentiated keratinocytes with the epidermis extending over the tumor, with a central horn plug of keratin and lip-like, peripheral borders of the epidermis. Intraepidermal neutrophilic abscesses were visualized in addition to horn pearls.

Vascular lesions can also present as pigmented lesions due to presence of underlying blood. They may appear black with the naked eye, but under dermatoscopy appear red, purple or blue [1]. In our study, 10.6% cases were of vascular etiology. All lesions were present in females with variable age distribution. There was 01 case of angiokeratoma which presented as a 1–6 mm red–blue, hyperkeratotic papule characterized by hyperplasia and hyperkeratosis of the epidermis along with multiple dilated vessels in the superficial dermis. There were 04 cases of lobular capillary hemangioma which showed distribution over scalp, hand and foot as lobular proliferation of small blood vessels surrounded by a mixed cell population of fibroblasts, mast cells, lymphocytes and plasma cells. Feeding blood vessel was identified at the base of lesions. The 03 cases of cavernous hemangioma were distributed over chest and leg microscopically characterised as cavernous hemangioma were also described by Bohra et al., [7] in their study.

Another inflammatory pigmentary lesion reported was post inflammatory hyperpigmentation (PIH). It is an acquired hypermelanosis that occurs after cutaneous inflammatory conditions that more frequently affects skin-of-color patients [28]. We had 05 cases of PIH, presenting between 2nd – 7th decade and distributed over neck, thigh, gluteal region and abdomen as tan brown macules and papules with a male preponderance. Histologically, melanophages were present in the superficial dermis, along with a variably dense infiltrate of lymphohistiocytes around superficial blood vessels and in dermal papillae (Fig-8).

Fig-7: Dermatofibroma with proliferating fibroblasts, histiocytes with melanophages and blood vessels (H&E)

Fig-8: Post inflammatory hyperpigmentation and perivascular lymphohistiocytic cells in superficial dermis (H&E)

Fig-9: Cavernous hemangioma with dilated, thin walled capillaries and simple endothelial lining (H&E)
CONCLUSION

Both melanocytic and non-melanocytic lesions can present as PSLs among any age group. Histopathological confirmation is imperative whenever a pigmented lesion clinically mimicking melanoma is encountered as many of these may turn out to be non-melanocytic and other benign melanocytic lesions on biopsy giving reassurance to the patient. A complete clinical diagnosis assists the histopathologist to give an accurate report.

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