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DERMOSCOPY AND SKIN IMAGING

DERMOSCOPIC-HISTOLOGICAL CORRELATION IN LICHEN PLANUS PIGMENTOSUS: UTILITY OF DERMOSCOPY IN ASSESSING DISEASE SEVERITY

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Background: Lichen planus pigmentosus (LPP) is characterized by macular pigmentation of varying shades involving the exposed areas, especially the face and neck. Dermoscopy, a non-invasive technique aids in diagnosis and prognostication of various dermatoses. In this prospective study, we present a dermoscopic-histological correlation in cases of inactive LPP.

Methodology: Out of 169 treatment-naïve Indian patients with Aquired dermal pigmentation, 67 patients were included with the closest diagnosis of inactive LPP, based on involvement, absence of preceding/concurrent erythema/edema & minimal-to-absent infiltrate on histology. Dermoscopic images were evaluated for - alteration of pseudoreticular network, presence of dermoscopic pigmented structures (DPS) – dots, globules and clods, their color and peri-adnexal arrangement. Histology was graded for dermal density and depth of macrophages. Spearman's coefficient (P<0.05) was employed for statistical correlation.

Results: Mean age & disease duration were 40.2±11.5 years and 4.15± 1.82 years respectively (51 females, 16 males). Dermoscopic findings - exaggeration of pseudoreticular network (41.9%); pigmented dots (100%), globules (84%), clods (53.2%), periadnexal involvement (30.6%). Color of DPS - brownish-black (88.7%), slate grey-bluish (38.7%). Histological findings – dermal macrophage density - severe in 64.5%, and dermal depth showed: <2mm-19.3%; 2-3 mm-37.1%; >3 mm-43.5%. Significant correlation found between size of DPS and dermal melanophages density & color of DPS and dermal depth of pigment incontinence, with deeper involvement (>3mm) correlating with slate grey-to-bluish hue.

Conclusion: The density of melanophages in the dermis & depth of dermal involvement with pigment incontinence are the two primary histological determinants of severity of LPP & response to therapy can be detected dermoscopically based on the size and color of DPS. A validated clinico-dermoscopic-histologic score needs to be evolved for LPP severity for therapeutic trials.





