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HEMANGIOMA OF INFANCY A RARE CASE REPORT IN PAEDIARTIC PATIENT

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ABSTRACT

Article History: Received 12th March, 2018 Received in revised form 10th April, 2018 Accepted 7th May, 2018 Published online 28th June, 2018 Infantile hemangioma is a common vascular tumour that, unlike congenital hemangioma, arise after birth at first 8 weeks of life and involute gradually around the age of 5. It often described as an isolated lesion but it can be associated with numerous syndromes like PHACE(S), kassaberich merit syndromes. Hereby we report a case of superficial infantile capillary hemangioma.

Key words:

Hemangioma, Infant, Lesion

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INTRODUCTION

Infantile hemangioma considered to be a most common tumour of infancy occurring around $5-10 \%^{1}$. It is a beningn vascular neoplasm with endothelial cell proliferation. Infantile hemangioma clinically can be classified into superficial, deep and mixed variants. Other type of hemangiomas which are congenital in nature further classified into RICH (rapidly involuting congenital hemangioma) and NICH (non involuting congenital hemangioma). Both types of congenital hemangioma are fully developed at birth, unlike infantile hemangioma, and either quickly involute (RICH) or remainlargely unchanged with time (NICH). Although infantile hemangioma not present at birth it can have minor presentation at birth like a telangiectasia's macule and it follows a pattern of a rapid growth phase followed by a gradual involution phase.

Case presentation

A 4-month-old female infant with her parents presented at our institution with chief complaint of a reddish patch in the right side lower lip region since 1 month. The lesion started during the third month of life. Initially it was started as a smaller lesion and within a week rapidly increased to the present size, after that there was no tendency for progression in the size of the lesion. There was no history of any complication during mother pregnancy period. On extraoral examination a raised dark reddish plaque present at the right angle of mouth involving vermillion border of lower lip. Lesion extended into the oral cavity involving the right lateral labial mucosa of lower lip till the vestibular area. Intraorally lesion appeared erythematous with diffuse borders. On palpation the extraoral lesion was firm to soft in consistency and non-tender without any blanching and bleeding tendency. Intraoral lesion was non palpable and non-tender. Infant was referred to paediatric department to rule out any systemic conditions.

Based on the clinical features it was provisionally diagnosed as infantile hemangioma capillary variant presumably. The parents were counselled about the nature of the lesion and it had to be wait and watch for the regression of the lesion. No interventional treatment was required for this lesion unless a rapid proliferation begins.



Fig 1 Extraoral photograph depicting a raised reddish to purple plaque involving right angle of mouth and vermillion border of lower lip



Fig 2 Intraoral photograph depicting a reddish erythematous lesion involving right side labial mucosa of lower lip extending into the vestibular region.

DISCUSSION

Infantile hemangioma is a common subtype of hemangioma and it cannot be diagnosed prenattally because it deveopls after birth only. It typically begins to appear a few days to weeks after birth and follows nascent, proliferative, plateau, and involutional stages.² Congenital hemangiomas, unlike infantile hemangioma, are fully developed at birth andcan classified as RICH or NICH.³

Clinically infantile hemangiomas often present as a raised bright red plaques like our patient but it can have variable clinical behavior which are associated with syndromes like PHACE.⁴ Eventhough infantile and congenital variants are considered to be a separate lesions but some reports suggesting that both are on a same spectrum of vascular tumor.⁵

However ifantile hemngiomas standsout as a separate entity from congenital hemangiomas and other vascular malformation with strong expression of GLUT-1 immunohistochemical marker in their proliferating endothelial cells. GLUT-1(glucose transporter-1) expressed in placental trophoblast which led to the hypothesis of ifantile hemangiomas can arise from trophoblast.

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