

Pregnancy Tumor: A Case Report

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Received Date: February 11, 2020; Published Date: February 17, 2020

Abstract

Skin tumors can be developed during pregnancy. Pyogenic granuloma is part of it; achromic melanoma is the main differential diagnosis. We report a case.

Keywords: Hemangiomas; Dermoscopy; Pyogenic

Introduction

During pregnancy, some skin tumors can occur accidentally due to hormonal changes. Pyogenic granuloma is part of it by acting on the vascular system and deserves special attention in treatment. The diagnosis is based on the clinic, dermoscopy and histology. We report a case.

Case Report

28-year-old woman, with no pathological history, pregnant

with 34 weeks amenorrhea. She presented during the third trimester of pregnancy a budding lesion on the little finger of the right hand painful gradually increasing size and becoming bleeding on contact. Note that the patient does not report the notion of trauma. The clinical examination revealed a well-defined budding tumor of regular contours surrounded by an epidermal collar about 1.5cm in diameter, pediculate-based, purplish red, at the level of the palmar extremity of the little finger of the right hand (Figure 1).

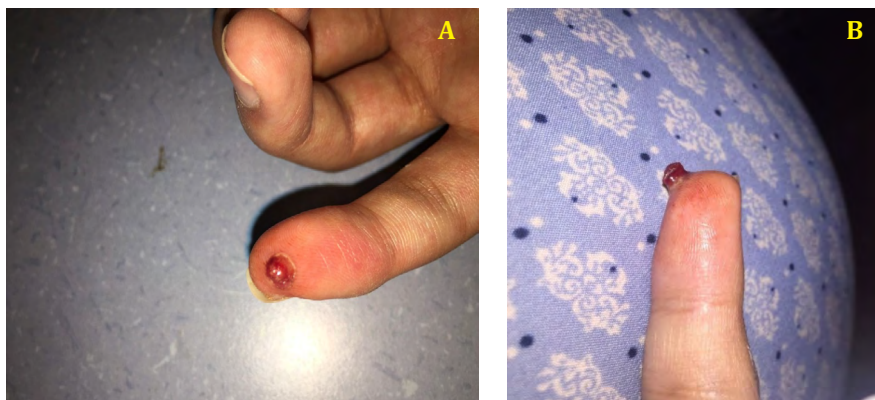


Figure 1: A budding tumor of regular contours surrounded by an epidermal collar (A) about 1,5cm in diameter, pediculate-based (B), purplish red, at the level of the little finger of the right hand.

The dermoscopy was not very conclusive, revealing vascular lagoons associated with a whitish collar on the periphery (Figure 2).



Figure 2: Dermoscopic image revealing vascular lagoons associated with a whitish collar on the periphery.

The somatic exam was normal. A pyogenic granuloma was strongly suspected, but a malignant tumor such as an

achromic melanoma could not be ruled out. An exeresis was made (Figure 3) and the histological result was compatible with a botryomycoma. No occurrence was noted after two years (Figure 4).

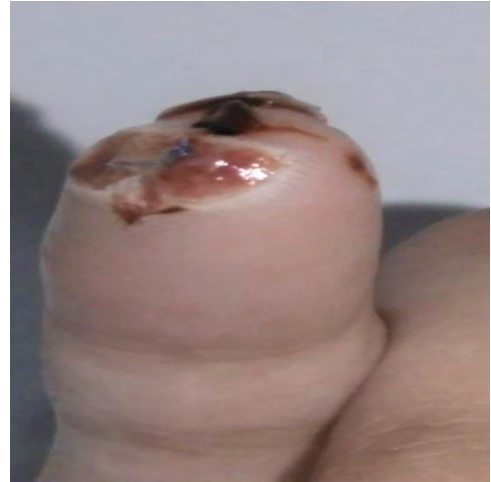
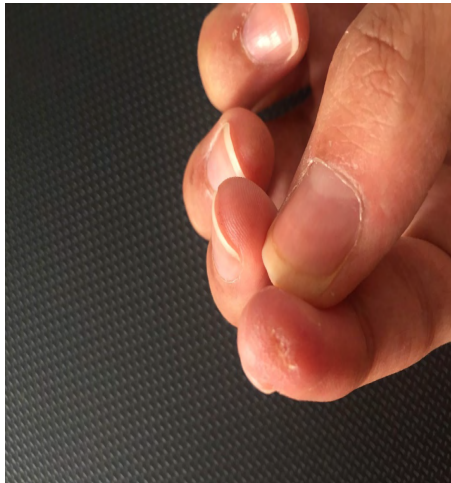


Figure 3: Post-operative image.



A



B

Figure 4: Control after 3 months (A) and after 2 years (B) without recurrence.

Discussion

Pyogenic granuloma, also called botryomycoma or lobular capillary hemangioma, is a benign tumor of vascular origin of the superficial dermis, but hypodermic forms have also been reported. It can be found all over the skin in tegument and mucous membrane [1]. It occurs as a result of chronic irritation, trauma or hormonal change during pregnancy [2]. The usual clinical presentation is that of a single budding tumor of small size, brittle, which can bleed on contact, of

preferential localization on the upper part of the body or the extremities presenting a great risk of recurrence. In dermoscopy, it typically has dots or globules of pink or light red color separated from each other by thick white lines (collagen septum). A white collar is often present at the periphery [3]. The pathology can be found at all ages but is more common in children. Its prevalence varies from 0.2% to 9.6% in pregnant women and frequently develops between the beginning and mid-gestation, between months 2 and 5 of pregnancy related to hormonal changes [4,5].

Localization in the gingiva is the most frequently described during pregnancy. Our patient's tumor appeared during her third trimester at his finger. The high level of sex hormones (estrogen and progesterone) seems to play an important role in the pathogenesis of botriomycoma after the first trimester of pregnancy.

Indeed, these hormones stimulate the expression of angiogenic factors in tissues which have an important role in vascular morphogenesis and are found in large quantities in pyogenic granulomas during pregnancy and in small quantities after childbirth [6-9]. In our case, the hormonal imbalance due to pregnancy was considered the etiological factor due to the absence of others. Histology confirms the diagnosis. The treatment is based on surgical excision, which illustrates the case of our patient. Other invasive options such as curettage, electrocoagulation or the application of silver nitrate are possible. The pulsed dye laser and the long pulse Yag laser can be effective on small lesions, with good esthetic results, but require suitable equipment and training [10,7]. The topical treatments used so far are Imiquimod, but it is often poorly tolerated and its efficacy is inconsistent, as is Timolol which is a non-cardioselective beta-blocker already recognized for its effectiveness in the treatment of hemangiomas of the child.

Conclusion

Several types of skin tumors develop or change during pregnancy, but fortunately the majority are benign and of no consequence to fetal and maternal health. In our case, achromic melanoma is the main differential diagnosis, if in doubt; surgical excision with histological study is required.

References

1. Fortna RR, Junkins-Hopkins JM (2007) A case of lobular capillary hemangioma (pyogenic granuloma), localized to the subcutaneous tissue, and review of the literature. *Am J Dermatopathol* 29(4): 408-411.
2. Akyol MU, Yalçiner EG, Dogan AI (2011) Pyogenic granuloma (lobular capillary hemangioma) of the tongue. *Int J Pediatr Otorhinolaryngol* 58(3): 239-241.
3. Zaballos P, Carulla M, Ozdemir F, Zalaudek I, Banus J, et al. (2010) Dermoscopy of pyogenic granuloma: a morphological study. *Br J Dermatol* 163(6): 1229-1237.
4. Ramos-E-Silva M, Martins NR, Kroumpouzou G (2016) Oral and vulvovaginal changes in pregnancy. *Clin Dermatol* 34(3): 353-358.
5. Fernandez A, Hamilton J, Nach R (2014) Two cases of pyogenic granuloma in pregnancy. *Ear, Nose Throat J* 93(8): 302-304.
6. Jafarzadeh H, Sanatkhan M, Mohtasham N (2006) Oral pyogenic granuloma: a review. *J Oral Sci* 48(4): 167-165.
7. Patil K, Mahima VG, Lahari K (2006) Extralingival pyogenic granuloma. *Indian J Dent Res* 17(4): 199-202.
8. Silverstein LH, Burton CH, Singh BB (1995) Oral pyogenic granuloma in pregnancy. *Int J Gyn Obst* 49(3): 331-332.
9. Choube S, Joshi P, Rawlani SM, Chawla R, Rathi V, et al. (2018) Pyogenic Granuloma -A Case Report. *Scientif J* 2: 1-4.
10. Gordón-Núñez MA, de Vasconcelos Carvalho M, Benevenuto TG, Lopes MF, Silva LM, et al. (2010) Oral pyogenic granuloma: a retrospective analysis of 293 cases in a Brazilian population. *J Oral Maxillofac Surg* 68(9): 2185-2188.