Squamous cell carcinoma in situ of the cervix and placental site nodule: Case report

María Constanza Gómez¹, María Claudia Abaúnza², Elga Johanna Vargas² and Inés Acosta¹ ¹NOVOPAT Laboratorio de Patología y Citología, Bogotá, Cundinamarca, Colombia; ²Surgical Pathology Research Group, Faculty of Medicine, Universidad de La Sabana, Chía, Cundinamarca, Colombia

Abstract

An asymptomatic 24-year-old woman underwent a colposcopy, cervical biopsy, and subsequently, a conization for a cervical squamous cell carcinoma in situ with glandular extension. Simultaneously, an endometrial biopsy was carried out in which, incidentally, a placental site nodule was diagnosed, a rare non-neoplastic lesion originating in the intermediate trophoblast. Given the coexistence of these two entities, it was necessary to make a differential diagnosis between them and also with other pathologies of the trophoblast such as an exaggerated placental site, placental site trophoblastic tumor, and epithelioid trophoblastic tumor.

KEY WORDS: Placental disease. Trophoblast. Trophoblast tumor. Placental site nodule. Intermediate trophoblast. Extravillous trophoblast.

Introduction

Placental site nodule (PSN) is a non-neoplastic infrequent lesion originating from the intermediate trophoblast, which theoretically represents a non-involuted portion of placental tissue. This lesion consists of a circumscribed nodule or plate, with abundant hyalinized stroma and trophoblastic cells of the intermediate trophoblast type^{1,2}. We present the case of a patient with coexistence of squamous cell carcinoma in situ with glandular extension of the cervix and PSN.

Case report

A 24-year-old female resident of a rural area of Colombia attended the local hospital in October 2013 owing to the finding of a high-grade squamous intraepithelial lesion (HSIL) on her Pap smear. She had no relevant medical history. Pregnancies 1, deliveries 1, live births 1. Date of last delivery: September 2010. She was having birth control with subdermal implant since one year prior. Physical examination was normal.

She underwent colposcopy and cervical biopsy, with a squamous cell carcinoma in situ being found, which led to the performance of a conization that confirmed the HSIL diagnosis: severe dysplasia and squamous cell carcinoma in situ with glandular extension on six out of 16 sections, with involvement of the endocervical resection margin (Fig. 1 A and B). Simultaneously, she underwent endometrial biopsy, with proliferative endometrium fragments and a PSN being observed.

By late 2014, the subdermal implant was removed and the control Pap smear reported a low grade squamous intraepithelial lesion, with cytopathic changes consistent with human papillomavirus infection, which was classified as genotype 16. Two months later, a second colposcopy and cervical biopsy were performed, with the latter being negative for squamous

Correspondence:

María Claudia Abaúnza Autopista Norte de Bogotá, km 7 Chía, Cundinamarca, Colombia E-mail: maria.abaunza@unisabana.edu.co

Date of modified version reception: 25-02-2016 Date of acceptance: 07-04-2016



M.C. Gómez, et al.: Cervical carcinoma and placental nodule

Figure 1. A: squamous cel carcinoma in situ of the cervix (hematoxylin & eosin, x10). **B:** glandular extension (hematoxylin & eosin, x20).

intraepithelial lesion. The patient is currently breastfeeding her second child.

Discussion

PSN occurs in childbearing age women^{2,3} and, usually, it is an incidental finding of endocervical curettages, cervical and endometrial biopsy and hysterectomy^{1,4,5}. It is mainly located at the endometrium or the cervix, and rarely at the uterine tube. It is usually detected several months or years after a pregnancy, with an average of 3 years⁶⁻⁸.

When macroscopically visible, a yellow or hemorrhagic nodule is appreciated, with a diameter of 1 to 14 mm, although it rarely exceeds 4 mm; it can occur as multiple nodules^{2,4-6}. Microscopically, a circumscribed plaque or nodule composed of hyalinized stroma with intermediate trophoblast-type cells distributed in groups or cords, or individually displayed with

Figure 2. A: *PSN composed of intermediate type trophoblastic cells with hyperchromatic nuclei and vacuolated cytoplasms. Hyaline stroma adjacent areas (hematoxylin & eosin, x40).* **B:** *trophoblastic cells nuclei spread on large areas of hyaline stroma. Lymphocytes are appreciated at the lesion periphery (hematoxylin & eosin, x40).*

absence or sparse atypical mitoses. The cells vary in size; the small ones are mono- or binucleated, with a clear, glycogen-rich cytoplasm, and the large ones have broad acidophilic or amphiphilic cytoplasms, with irregular and hyperchromatic nuclei (Fig. 2 A). Multinucleated trophoblastic cells can occasionally be observed. Towards the periphery, there is a circumferential inflammatory infiltrate composed of lymphocytes and plasmocytes^{2,3,5-8} (Fig. 2 B).

PSNs are positive for PLAP, p63, inhibin α and cytokeratin 18, with focal or negative expression for hPL and CD146 (Mel-CAM). They are usually negative for β -hCG and their Ki67 proliferation index ranges between 1 and 5% (Fig. 3 A and E). Given their morphology and immunohistochemical (IHC) characteristics, PSNs originate from the intermediate trophoblast of the chorionic type, and it is suggested that they are the benign counterpart of the epithelioid traphoblastic tumor (ETT)^{1,2,8,9}.

Most important differential diagnoses include exaggerated placental site (EPS), placental site trophobalstic tumor (PSTT), ETT and squamous cell carcinoma of the cervix^{5,8-10} (Table 1). EPS is diagnosed after a normal pregnancy, an ectopic pregnancy or a



Figure 3. PSN trophoblastic cells are positive for P63 (IHC, x20) (**A**), PLAP (IHC, x40) (**B**), inhibin (IHC, x40) (**C**) and keratin (IHC, x40) (**D**). **E:** Ki67proliferation index is lower than 1% (IHC, x40).

	EPS	PSN	PSTT	ETT	SCC
Histogenesis	Implantation site intermediate trophoblast	Chorionic type intermediate trophoblast	Implantation site intermediate trophoblast	Chorionic type intermediate trophoblast	Epithelial lesion. Non-trophoblastic
Morphological findings					
Growth pattern	Not arranged in nodules	Circumscribed nodule(s)	Poorly-defined and infiltrating mass	Falsely circumscribed mass with focal infiltration	Infiltrating mass
Cellularity	High	Low	High	High	High
Cytological atypia	Generalized	Focal/sparse	Generalized	Generalized	Generalized
Mitotic figures	Absent	Absent to rare	Common	Common	Common
Stroma	Fibrin deposit	Abundant hyalinization	Deposit of fibroid material	Deposit of hyaline material	Sparse
Necrosis	Absent	Occasional, central	Common, coagulative	Geographic and extensive	Geographic
Chorionic villi	Present	Absent	Absent	Absent	Absent
Immunohistochemistry					
CK 18	Positive. Diffuse	Positive. Diffuse	Positive. Diffuse	Positive. Diffuse	Negative
CD146 (Mel-CAM)	Positive. Diffuse	Negative/positive focal	Positive. Diffuse	Negative/positive focal	Negative

Table 1. PSN differential diagnoses

	EPS	PSN	PSTT	ETT	SCC
PLAP	Negative	Positive. Diffuse	Negative	Positive. Diffuse	Negative
Inhibin alpha	Positive	Positive	Positive	Positive	Negative
hPL	Positive. Diffuse	Negative/positive focal	Positive. Diffuse	Negative/positive focal	Negative
P63	Negative	Positive. Diffuse	Negative	Positive. Diffuse	Positive. Diffuse
P16 (nuclear)	Negative	Negative	Negative	Negative	Positive
Ki-67	< 1%	1-5%	> 10%	10-25%	10-50%

Table 1. PSN diff	erential diagnoses	(Continued)
-------------------	--------------------	-------------

EPS: exaggerated placental site; PSN: placental site nodule; PSTT: placental site trophoblastic tumor; ETT: epithelioid trophoblastic tumor; SCC: squamous cell carcinoma.

molar pregnancy; it doesn't arrange in nodules, it is usually associated with chorionic villi and its Ki-67 proliferation index is lower than 1%^{8,10}. PSTT is macroscopically visible, and its histology is characterized by myometral infiltration, areas of coagulation necrosis, tumor cells with atypical nuclei, frequent mitosis and Ki-67 proliferation index higher than 10%^{5,7,8,10}. Both EPS and PSTT are negative for p63 and diffusely positive for hPL¹¹. ETT shows IHC findings similar to those of PSN; however, they are neoplasms of larger size, infiltrating growth with high cellularity, pleomorphism, atypical mitoses, geographic necrosis and high cell proliferation index (10-25%)^{1,5,8,10.11}.

In our case, the most important differential diagnosis is cervical squamous cell carcinoma. Some patients assessed for cervical dysplasia or carcinoma in situ with a concomitant PSN can be wrongly diagnosed with infiltrating squamous cell carcinoma. In this case, IHC is useful because antibodies against HLA-G and CK18 are diffusely positive in trophoblastic lesions and negative in squamous cell carcinoma. Inhibin alpha is positive in PSN and negative in squamous cell carcinoma⁶.

PSNs are lesions that do not require treatment in addition to initial surgical resection¹⁰.

Conclusion

PSNs are sometimes diagnosed in women with Pap smear abnormal findings¹. In a study by Shih et al.⁶, this condition was observed in 29% of patients. Although

PSN is infrequent, pathologists should recognize its morphological features and obtain the necessary IHC markers to differentiate it in patients with concomitant HSIL, in order to avoid the infiltrating squamous cell carcinoma of the cervix misdiagnosis.

References

- Hui P, Baergen R, Cheung ANY, et al. Non neoplastic lesions. En: Kurman RJ, Carcangiu ML, Herrington CS, Young RH, editores. WHO Classification of tumours of female reproductive organs. Lyon, Francia: IARC; 2014. p. 162.
- Shih IM, Mazur MT, Kurman RJ. Gestational trophoblastic tumors and related tumors and related tumor-like lesions. En: Shih IM, Mazur MT, Kurman RJ, editores. Blaustein's Pathology of the female genital tract. 6th ed. New York, EE.UU.: Springer; 2011. p. 1108-21.
- Huettner PC, Gersell DJ. Placental site nodule: a clinicopathologic study of 38 cases. Int J Gynecol Pathol. 1994;13:191-8.
- Shih IM, Mazur MT, Kurman RJ. Gestational trophoblastic disease. En: Mills SE, Greenson JK, Hornick JL, Longacre TA, Reuter VE, editores. Sternberg's Diagnostic surgical pathology. 6th ed. Philadelphia, EE.UU.: Wolters Kluwer; 2015. p. 2297-319.
- Mali K, Fernandes G, More V, Satia MN. Placental site nodule. JPGO. 2014;1(10). Disponible en: http://www.jpgo.org/2014/10/placental-site-nodule.html
- Shih IM, Seidman JD, Kurman RJ. Placental site nodule and characterization of distinctive types of intermediate trophoblast. Human Pathol.1999:30:687-94.
- Young RH, Kurman RJ, Scully RE. Placental site nodules and plaques. A clinicopathologic analysis of 20 cases. Am J Surg Pathol. 1990:14: 1001-9.
- Kim KR. Gestational trophoblastic disease. En: Mutter GL, Prat J, editores. Pathology of the female reproductive tract. 3rd ed. Philadelphia, EE.UU.: Churchill Livingstone Elsevier; 2014. p. 803-9.
- Valverde D, Dulcey I, Nogales FF. Coexistence of placental site nodule and cervical squamous carcinoma in a 72-year-old woman. Int J Gynecol Pathol. 2013;32:335-7.
- Crum CP, Hirsch MS, Peters III WA, Quick CM, Laury AR. Gynecologic and Obstetric Pathology. A volume in the High Yield Pathology Series. Philadelphia, EE.UU.: Saunders Elsevier; 2016. p. 735-57.
- Shih IM, Kurman RJ. P63 expression is useful in the distinction of epithelioid trophoblastic and placental site trophoblastic tumors by profiling trophoblastic subpoblations. Am J Surg Pathol. 2004;28:1117-83.