Intravascular Papillary Endothelial Hyperplasia Mimicking Vaginal Malignant Tumor in a Post-Radiotherapy Patient: A Case Report and Literature Review

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Abstract: Intravascular papillary endothelial hyperplasia (IPEH) is a rare benign lesion that occurs in the soft tissue of any site, including the central nervous system, extremities, head, and female reproductive system. But no case occurred in vagina was reported previously. Here, we report a case of vaginal IPEH in a 54-year-old woman who has a history of radical hysterectomy for endometrial cancer with postoperative radiotherapy. She was positive for high-risk human papillomavirus (HR-HPV) with cytology result as low-grade squamous intraepithelial lesion (LSIL), and pathology from colposcopy-directed biopsy (CDB) presented suspicious high-grade squamous intraepithelial lesion (HSIL). The colposcopy showed an 8mm neoplasm at the right apical angle of vagina. The diagnosis of IPEH was confirmed by biopsy pathology and immunohistochemistry. As the lesion was benign and rather small, we discharged the patient with annual follow-up. The case illuminated that attention should be paid to “HSIL” patients after radiotherapy to exclude the incidence of benign IPEH.

Keywords: case report, intravascular papillary endothelial hyperplasia, radiotherapy, vagina, endometrial neoplasms

IPEH, also known as Masson’s tumor, is a rare hemangioma that occurred mainly in vessels, but it was also reported in head, extremities, breast, female vulva, etc.¹–⁴ It is considered as a benign tumor originated in vascular endothelium that presented excessive hyperplasia and papillary formation. Most cases occur without a definite etiology, and about 30% of cases are post-trauma and have a history of vascular diseases.⁵ The extracranial Masson’s tumor presents a slow-growing and asymptomatic characteristic with the clinical feature as a small, firm, non-pulsatile, slightly elevated mass with a red or blue coloration. Here, we report a case with colposcopic finding as suspicious HSIL but diagnosed as benign IPEH by pathology and immunohistochemistry.

Case Report

A 54-year-old woman with a history of endometrial carcinoma presented with suspicious HISL came to our hospital. She had hysterectomy and bilateral salpingo-ooophrectomy in 2016 with regular chemoradiotherapy (25 radiotherapies with 2 brachytherapy). Her postoperative follow-up showed HR-HPV positive, and the
ThinPrep cytologic test (TCT) result was LSIL. The pathology from CDB by another hospital implied suspicious HISL in April 2019. In July 2020, the colposcopy in our center detected a neoplasm at the right apical angle of vagina. The patient was then recommended for laser treatment in case of malignancy. During the operation, an 8-millimeter-diameter neoplasm at the right apical angle of vagina was seen (Figure 1). The subsequent pathologic result of CDB presented IPEH, and the immunohistochemical result was CD31(+), CD34(+), AE1/AE3(-), and Ki-67(+, 5%) (Figure 2). After reconfirmation with pathologists, we affirmed the neoplasm was a benign hemangioma, which was dispensable for extra therapy. The woman was then discharged with regular follow-up.

**Discussion**

IPEH was first reported in 1923 by Pierre Masson, the histologic hallmark of which was a constitutive thrombus within the vascular with endothelial cell proliferation.\(^4,6\) It can be divided into three types. The primary form occurred in dilated vessels, the secondary form presented in variceal vessels, hemangioma, inflammatory granuloma, vascular

![Figure 1](https://doi.org/10.2147/OTT.S317160)

**Figure 1** Colposcopes view of the lesion. (A) An 8-millimeter-diameter intravascular papillary endothelial hyperplasia in the right apical angle of the vagina with medium amount of blood. The lesion is red to purple coloration, and a medium amount of blood was seen. (B) Iodine staining of the lesion.

![Figure 2](https://doi.org/10.2147/OTT.S317160)

**Figure 2** Pathology and immunohistochemistry result (200×). (A) Microscopic finding presents proliferation of erythrocytes within a dilated vascular structure. (B) Ki-67 labeling index was about 5%. Immunohistochemistry shows the tissue was positive for CD31 (C) and CD34 (D).
tumor, and the extravascular Masson’s tumor is extremely rare, which can occur in multiple sites throughout the body. The rare extravascular form has been reported to happen in periocular region which causes reversible vision loss and proptosis, mandible, and oral region. In the female reproductive system, three vulval cases, two ovarian cases and one cervical case of IPEH occurred have been reported, the common therapy was complete excision of the lesion (Table 1). In ovarian IPEH, the lesion was prominently larger and in one case it was mistaken with a malignant tendency. And the other case reported a mixed neoplasm of malignant mucinous borderline ovarian tumor and IPEH, which hinted the possibility of simultaneous incidence of malignant tumor and benign Masson’s tumor. In IPEH reported in the cervix, the patient was in gestation and the tumor enlarged fast, growing from 3cm to 7cm in 8 weeks under the environment of high levels of estrogen and progesterone. The lesion was positive for CD31, CD34, and factor VIII, negative for epithelial membrane antigen, and Ki-67 labeling index was 1%, which was consistent with our case. The pathological result was of the best diagnostic value to identify IPEH, and CD31 and CD34 in immunohistochemistry were used to remark the vascular endothelium. Considering the prominent symptoms as vaginal bleeding and pain, the excision of the lesion followed by suture of the anterior cervical lip was implemented during pregnancy, ensuing favorable maternal and fetal prognosis. The patient in our case was asymptomatic and the lesion was small, hence we did not perform excision.

Irradiation has proved to be the risk factor of its induction and IPEH in the central nervous system, bladder after radiotherapy has also been reported. IPEH sometimes can be misdiagnosed as radiotherapy-associated malignant tumor such as angiosarcoma. Two cases reported post-radiotherapy breast cancer patients with IPEH, in which the author differentiated IPEH and angiosarcomas by its clear border and non-invasive feature and macroscopic absence of atypia, necrosis, pleomorphism. However, adjuvant radiotherapy may be required in intracranial Masson’s tumor to prevent recurrence or to treat patients whose lesion cannot achieve radical excision. The patient in our case has undergone 25 courses of radiotherapy including 2 brachytherapy, which might be the predisposing cause of IPEH.

**Conclusion**

Radiotherapy can induce the occurrence of IPEH, which can be misdiagnosed as malignant tumor which results in inappropriate treatment. To differentiate it from other malignant tumors, biopsy pathology and immunohistochemistry need to be applied. The treatment of IPEH is mainly complete local excision, but asymptomatic small lesions can adapt conserved therapy and continuously following up.

| Reference         | Age (Years) | Region     | Symptom                                 | Size            | Treatment                                                                 |
|-------------------|-------------|------------|-----------------------------------------|-----------------|---------------------------------------------------------------------------|
| Kim et al (2014)  | 37          | Vulva      | A palpable mass with intermittent vulvar pain | 30×20mm         | Excision                                                                 |
| Beutler et al (2016) | 32         | Vulva      | Painless, black papule                   | 3×2mm           | Excision biopsy                                                           |
| Nwokolo et al (2020) | 35         | Vulva      | Tender mass with worsening pain          | 20×30mm         | Excision                                                                 |
| Sherring et al (2020) | 44        | Left ovary | Intermittent pelvic discomfort           | 200×180×95mm    | Hysterectomy, bilateral salpingooophorectomy, and omentectomy            |
| Sherring et al (2020) | 57        | Left ovary | Increasing abdominal girth               | 260×225×165mm   | Hysterectomy, bilateral salpingooophorectomy, and omentectomy            |
| Susini et al (2010) | 40         | Uterine cervix | Severe pain and vaginal bleeding         | 70mm            | Excision                                                                 |

**Abbreviations:** IPEH, intravascular papillary endothelial hyperplasia; HPV, human papillomavirus; NS, not state; HHV, human herpesvirus.
Ethical Approval and Consent to Participate

Institutional review board/ethics committee approval was obtained from Fudan University Obstetrics and Gynecology Hospital for this study (reference number: Kyy2018-81) and written informed consent was obtained from the patient for publication of this report and accompanying images.

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Disclosure

The authors declare that they have no conflicts of interest and nothing to disclose.

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