

## Case Report

## Uterine Primitive Neuroectodermal Tumor

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## Abstract

Primitive neuroectodermal tumors are fairly rare in uterus. A case of uterine body primitive neuroectodermal tumor in a 32-year-old Iranian woman is presented. The patient was admitted with abdominal pain and fever and underwent emergency exploratory surgery with total abdominal hysterectomy, bilateral salpingo-oophorectomy, and pelvic lymph node dissection. Posterior wall of the uterus was necrotic and ruptured and a huge tumor disrupted the uterine body. The tumor was strongly positive for CD99, NSE, and chromogranin; No reaction was seen for CD10, CD45 and myogenin. To the best of our knowledge, this is the first report of an uterine body primitive neuroectodermal tumor and the second report of uterine primitive neuroectodermal tumor from Iran.

**Keywords:** Primitive neuroectodermal tumor, uterus

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## Introduction

In 1973, Hart and Earl described a group of small round cell tumors with variable degrees of ependymal, glial, and neural differentiation which are considered to originate from fetal neuroectodermal cells.<sup>1</sup> Ewing's sarcoma and primitive neuroectodermal tumor (PNET) stand for a single group of bone and soft-tissue tumors in which PNET with evidence of neural differentiation lies at one end of the spectrum and undifferentiated Ewing's sarcoma lies at the other.<sup>2,3</sup> Both have similar phenotype and share an identical chromosomal translocation. PNET rarely involves the female genital tract<sup>4</sup>; however a few cases have been reported to involve the ovary, uterine corpus, uterine cervix, and vulva.<sup>3,5-7</sup> Here, we report a case of PNET in the uterine body.

## Case Report

In September 2011, a 32-year old Iranian woman, gravid 3, para 2, live 1, death 1, ectopic pregnancy 1 (G3P2L1D1EP1) presented with abdominal pain and fever since 2 days before admission to our emergency room. She suffered from abnormal vaginal bleeding since 4 years before admission; a fractional dilation and curettage (D and C) which was performed two weeks before admission was reported as PNET involving endometrium.

Family history was negative for malignancy; there was a history of cesarean section (C/S), laparoscopy (due to ectopic pregnancy) and D and C. Drug history was negative. The Papanicolaou smear

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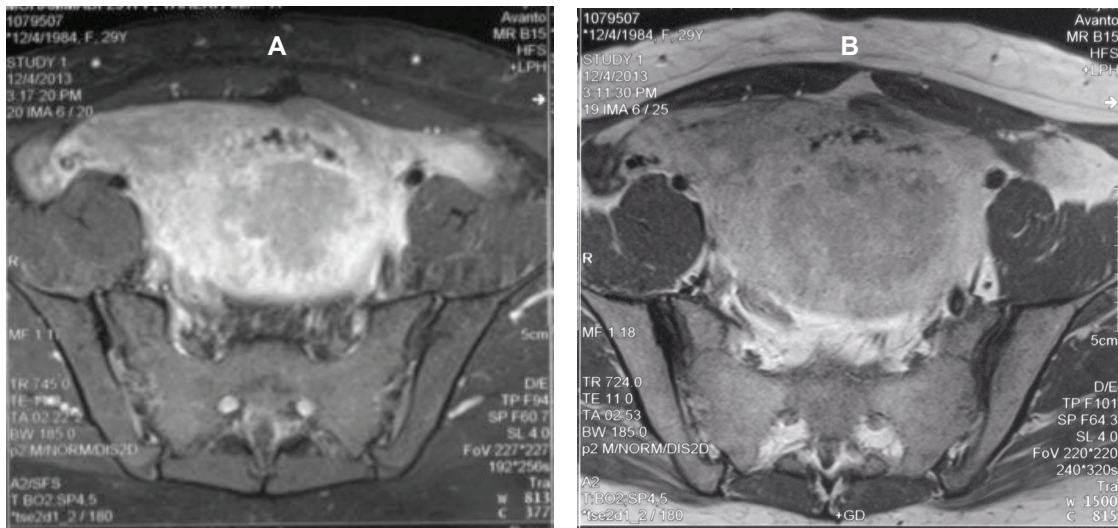
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was negative for malignancy. Ultrasound examination reported an enlarged uterus (12.5 × 11 × 8 cm) and solid mass 9 × 6.5 cm in the anterior and lateral walls of the uterus. Two 54 × 37 mm and 29 × 25 mm cysts with septa were reported in the left and right ovaries, respectively.

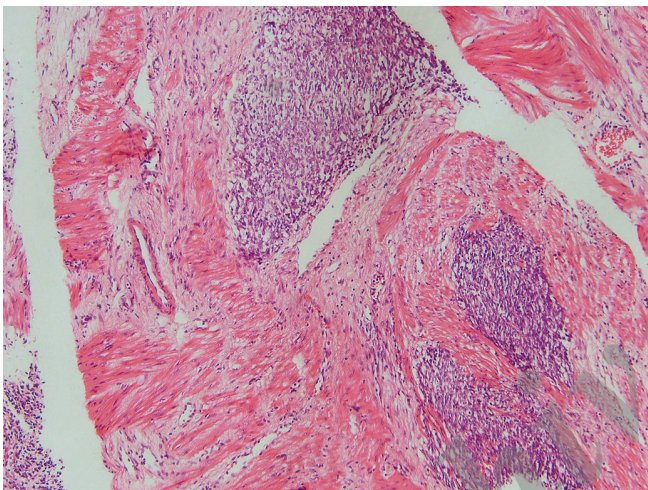
Magnetic resonance imaging (MRI) showed a bulky intramural, partially enhancing mass on the right side of fundus of uterus. (Figure 1). An enhancing bony lesion in the right aspect of sacrum was also reported which suggested probable bony metastasis; however, bone scan reported no evidence of bony metastasis and exploratory laparotomy did not reveal any evidence for such a diagnosis. Tumor markers including CA 125, AFP, and CEA were in the normal range. The patient subsequently underwent emergency exploratory surgery with total abdominal hysterectomy, bilateral salpingo-oophorectomy, and pelvic lymph node dissection. The posterior wall of the uterus was necrotic and ruptured and a huge tumor disrupted the uterine body.

Microscopically, there was a small round cell tumor growing as solid nests throughout the myometrium and replacing the endometrium with large areas of necrosis. There was no cellular pleomorphism or evidence of rosette formation. Immunohistochemically, the tumor cells were positive for CD99, NSE and chromogranin; no reaction was seen for CD10, CD45, and myogenin. Thus, small round cell tumors like endometrial stromal sarcoma and rhabdomyosarcoma were ruled out<sup>8,9</sup> (Figures 2–4).

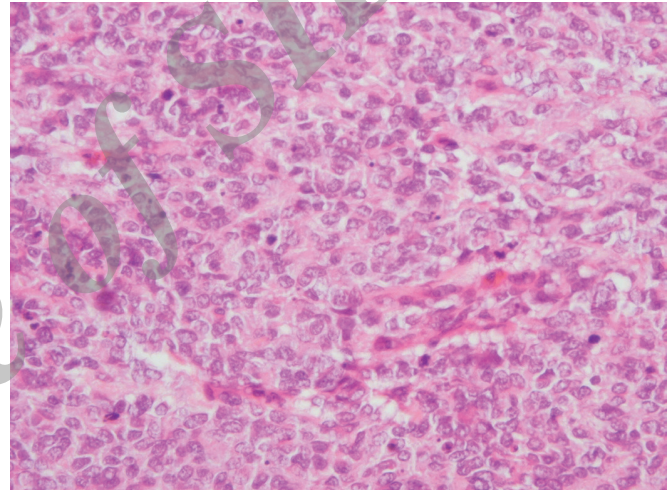
The patient underwent chemotherapy with Holoxan (1.5 gr/m<sup>2</sup>) and Mesna (400 mg TDS) Cisplatin (80 mg/m<sup>2</sup>) for 4 days followed by radiotherapy. Two other chemotherapy cycles with the same diet were performed after radiotherapy. In February 2013, the patient was re-admitted due to ascites and tumor recurrence was confirmed by laparoscopy. The patient was treated with Paclitaxol (180 mg/m<sup>2</sup>) and Carboplatin AUC7 (600 mg) for 6 cycles. In November 2013, she came back with ascites and tumor recurrence and peritoneal seeding, diagnosed by ascites tap; because of poor prognosis, the patient was given palliative treatment. She is alive now, but her prognosis is quite poor.



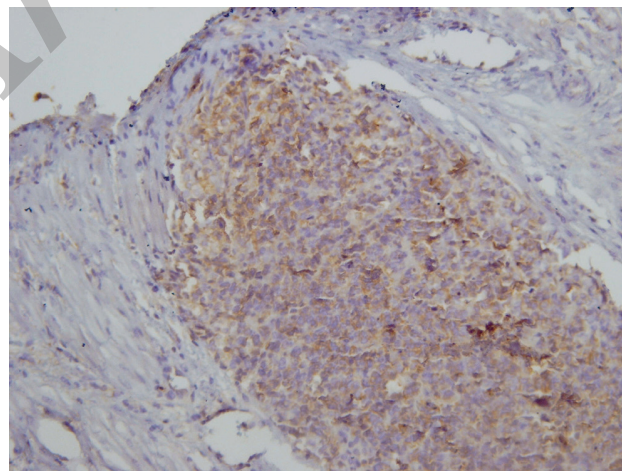
**Figure 1. A)** Magnetic resonance imaging (MRI) showed a bulky intramural, **B)** partially enhancing mass on the right side of fundus of uterus.



**Figure 2.** Several nests of tumor invading myometrium (Hematoxylin and Eosin,  $\times 100$ )



**Figure 3.** Tumoral cells are small to medium size, with plump nuclei, fine chromatin and small amounts of cytoplasm. Note frequent mitotic figures (Hematoxylin and Eosin,  $\times 400$ )



**Figure 4.** Immunohistochemical (IHC) staining for CD99. Tumor cells are diffusely positive ( $\times 200$ )



## Discussion

In 1918, South described this tumor but the term PNET was coined by Hart and Earle in 1973 to show a number of small round cell tumors which may be derived from neuroectodermal cell showing different signs of neural, glial and ependymal differentiation.<sup>1</sup> PNET of uterus has been rarely reported. Hendrickson and Scheithauer first reported this tumor in 1986.<sup>10</sup> It has been shown that PNET has high malignant feature which progresses rapidly and metastasizes widely.<sup>1,3,11</sup> The differential diagnosis of PNET in the uterus usually includes other uterine tumors including small cell carcinoma, sarcoma, and lymphoma.<sup>4,11</sup> Immunohistochemical staining and histological findings might help to distinguish this rare tumor from other more common malignancies.<sup>3,8</sup> PNET has a mesodermal origin and is associated with tumors of known Mullerian origin.<sup>12</sup>

Uterine PNET has been typically reported in 2 groups: adolescent girls and postmenopausal women; however our patient was a 32-year-old woman. In line with our case presentation, the most common reported findings in PNET are abnormal vaginal bleeding and uterine enlargement.<sup>4,11</sup>

PNET diagnosis is difficult and several differential diagnoses should be considered. Uterine PNET mostly shows various types of neuroectodermal elements like fibrillary background, ganglion cells astrocyte like cells, rosette like cells, rosetts, ependymal and medulloepithelial differentiation.<sup>13</sup> Unfortunately, many PNET cases have been described at advanced stages.<sup>6,10,11,13</sup>

Immunohistochemical staining can help with the diagnosis of PNET and determining its prognosis. CD99 is the most practical immunohistochemical marker for diagnosis of PNET.<sup>14</sup> CD99 is a monoclonal antibody of the surface protein MIC2 whose gene is located on pseudoautosomal region of X and Y chromosomes.<sup>8,9</sup> CD99 prognostic properties vary in different tumors. For example, CD99 presentation predicts good prognosis in diffuse large B-cell lymphoma (DLBCL) with the germinal center B-cell subtype and non-small cell lung cancer (NSCLC), while in DLBCL with non-GCB, CD99 is a marker of poor prognosis.<sup>15,16</sup>

Unfortunately, to our knowledge no study has evaluated CD99 in uterine PNET; however, it has been shown that atypical Ewing Sarcoma (including CD99, FLI1, HNK1, and CAV1 negative tumors) is associated with less favorable clinical outcome.<sup>17</sup> Our patient's tumor was strongly positive for CD99 (Figure 2). As a limitation of our case presentation, t(11/22) translocation as a precise marker for PNET was not evaluated in our case.

Without sufficient treatment, more than 90% of patients with Ewing's sarcoma die due to secondary hematogeneous metastases. However, with satisfactory cytotoxic treatment, the five-year survival rate can increase up to 60% in localized disease.<sup>18</sup> Rather than cytotoxic treatment, in the case of PNET, surgery should be considered as the first step. Total hysterectomy with bilateral salpingo-oophorectomy with or without chemotherapy and/or radiotherapy is the usual course of treatment provided.<sup>19</sup> To the best of our knowledge, this is the first report of uterine body PNET from Iran. Only one case of cervical PNET from our country has been reported before.<sup>14</sup>

## Consent

Written informed consent was obtained from the patient.

## Conflict of interests

We declare that we have no conflict of interest.

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