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ENDOMETRIAL STROMAL SARCOMA- A CASE REPORT

MRS.M.SARANYA, ASST PROF

SREE BALAJI COLLEGE OF NURSING,BHARATH INSTITUTE OF HIGHER EDUCATION AND RESEARCH,CHROMEPET, CHENNAI

ABSTRACT:

Endometrial stromal sarcomas are rare malignant tumours of the uterus. Most details about this tumour is obtained from previous case reports. In most of the cases, diagnosis is made only postoperatively. A 29 year old female, came with complaints of heavy menstrual bleeding, lower abdominal pain, mass in lower abdomen for 2 months. Clinical examination and investigations showed bulky irregularly enlarged uterus probably Fibroid Uterus. Patient was planned for Myomectomy. Intraoperatively, uterus was irregularly enlarged with cauliflower like growth lymphnode dissection with infracolic omentectomy was done. Histopatholgy examination revealed as Low Grade Endometrial Stromal of size 7 x 5 cm over fundus suggestive of Carcinoma. Total abdominal hysterectomy with bilateral salphingo oophorectomy with bilateral pelvic Sarcoma with Negative Lymph Node Spread. Postoperatively, Patient was evaluated for ER sensitivity and started on Tab. Anastrazole.

KEY WORDS:

Uterine sarcoma, Endometrial stromal sarcoma, Malignant uterine tumour

INTRODUCTION:

Endometrial stromal sarcomas constitute around 0.2% of all uterine malignancies. They constitute second most common mesenchymal uterine tumours. It usually involves age group of 40-58 years and most women are premenopausal. World health organisation classifies ESS into 1) Endometrial Stromal Nodule 2)Low Grade ESS 3) High Grade ESS 4)Undifferentiated ESS. It usually presents late with local metastasis. Distant

metastasis can occur even 20 years after initial diagnosis. Surgery and hormonal therapy are mainstay of treatment. Prognosis is good in case of early diagnosis.

CASE REPORT:

A 29 year old female was admitted in Department of Obstetrics and Gynaecology, Sree Balaji Medical College and Hospital, chromepet with complaints of heavy menstrual bleeding, lower abdominal pain, mass in lower abdomen for 2 months. Clinical examination revealed uterus enlarged upto 14-16 weeks size, firm, mobile from side to side, irregular margins with irregular surface. Ultrasonogram of abdomen showed uterus of size 10.3*8.0*7.8 cm, intramural fibroid of size 6.7*6 cm in anterior wall. Patient was planned for Myomectomy. Intraop findings-uterus irregularly enlarged, cauliflower like growth of size 7 x 5 cm seen over fundus, edematous right fallopian tube. Bilateral ovaries were cystic and unhealthy. On suspicion of malignancy, Total Abdominal Hysterectomy with Bilateral Salphingo Oophorectomy with Bilateral Pelvic Lymphnode Dissection with Infracolic Omentectomy was done. Histopatholgy examination report came as Low Grade Endometrial Stromal Sarcoma with Negative Tumour Deposit. Patient underwent Immuno histochemistry studies and found to be ER positive. Patient was started on Tab. Anastrazole 1 mg OD. Patient is in regular follow up.



DISCUSSION:

Endometrial stromal sarcoma (ESS)is second most common Malignant Pure Mesenchymal Uterine Tumor(MPMUT) constituting about 0.2% of all uterine tumours.1 It arises from submesothelial pleuripotent

mullerian cells.2 ESS is classified into i)Endometrial Stromal Nodule(ESN) ii)Low Grade ESS(LGESS) iii)High Grade ESS iv)Undifferentiated ess (UES).3 Cytogenetic Abnormality of ESS is recurrent translocation .4 Low grade tumours are indolent tumours with favorable prognosis LGESS occurs in age group of 28-67 years and presents as uterine enlargement, pelvic mass, abnormal uterine bleeding.6 High grade tumours have features intermediate between LGESS and UES.7 Microscopically, HGESS have round cells with spindle cell fibromyxoid component.6 Undifferentiated ESS usually present as postmenopausal bleeding.

Imaging studies of ESS are usually nonspecific. Positive labelling for CD, PR, ER with negativity for CD1,CD34 is confirmative diagnosis. Mainstay of treatment is total abdominal hysterectomy with bilateral salphingo hormonal therapy is helpful. It includes medroxyprogesterone acetate, Gnrh agonists, aromatase inhibitors. In earlier studies, oral medroxyprogesterone acetate was used. In recent times, combination containing Doxorubicin is used. Ifosfamide is helpful in chemoresistant sarcomas. Tamoxifen, estrogens are contraindicated as they cause stimulative effect on disseminated endometrial stromal cells. Dydrogestrone is hormonally active, non androgenic, synthetic steroid. It lacks estrogenic, androgenic, glucocorticoid, mineralocorticoid properties. Dydrogestrone is preferred to other progestins due to lack of side effects. Long term use of Gnrh agonists can cause osteoporosis. Anastrazole is the preferred oophorectomy. In case of pregnancy associated with sarcomas, laproscopy can be done safely. Abdominal surgery can also be done safely in case of pregnancy associated ESS. In case of inoperable, recurrent tumours, aromatase inhibitor. They cause muscle pain, nausea, vomiting.

CONCLUSION:

There is noncompliance of Letrozole while usage in breast cancer patients. 5 year survival rate of patients with stage I and II is 90%, stage III and IV is 50%. Follow up is done once in 3 months for first year, half yearly for next 4 years, thereafter yearly follow up.

REFERENCES:

- 1. D" Angeloe, Prat J. Uterine Sarcomas: A Review. Gynaecological Oncol 2010;116;131-9.
- 2. Baiocchi G, Kavanagh JJ, Wharton JT. Endometrial Stromal sarcomas arising from ovarian and extraovarian endometriosis:report of two cases and review of literature. Gynaecological Oncol 1990;36:147-51.
- 3. Oliva E. Cellular mesenchymal tumours of uterus: A review emphasizing recent Observatios. Int J Gynecol Pathol 2014;33:374-384.
- 4. Nucci Mr, Harburger D, Koontz J, Molecular analysis of jazf1-jjaz1 gene fusion by rt pcr and fish in endometrial stromal neoplasms. Am J Surg Pathol 2007;31:65-70. 5.

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- 5.Chang Kl, Crabtree Gs, Lim Tan Sk. Primary uterine endometrial stromal neoplasms. A clinicopatholgic study of 117 cases. Am J Surg Pathol 1990;14:415-38.
- 6. Lee Ch, Marino-Enriquez A, Ou W. The clinicopathologic features of ywhae-fam22 endometrial stromal sarcomas: A histologically high-grade and clinically aggressive tumor. Am J surgpathol 2012;36:641–53
- 7. Kurihara S, Oda Y, Ohishi Y. Endometrial stromal sarcomas and related high-grade Sarcomas: Immunohistochemical and molecular genetic study of 31 cases. Am J Surg Pathol 2008;32:1228–38
- 8. Tanner EJ, Garg K, Leitao MM Jr. High grade undifferentiated uterine sarcoma: Surgery, treatment, and survival outcomes. Gynecol Oncol 2012;127:27–31.
- 9. Mourra N, Tiret E, Parc Y, de Saint-Maur P, Parc R, Flejou JF et al. Endometrial stromal sarcoma of the rectosigmoid colon arising inextragonadal endometriosis and revealed by portal vein thrombosis. Arch Pathol Lab Med 2001;125:1088-90.
- 10.Koyama T, Togashi K, Konishi I, Kobayashi H, Ueda H, Kataoka ML et al. MR imaging of endometrial stromal sarcoma: correlation with pathologic findings. AJR Am J Roentgenol 1999;173:767-72. 12