HISTOMORPHOLOGICAL SPECTRUM OF UTERINE LEIOMYOMA VARIANTS- ONE YEAR RETROSPECTIVE STUDY IN A TERTIARY CARE CENTRE.

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Abstract

Background: Benign leiomyomas were the most common pelvic tumor in women (1,2,3). Most myometrial tumors continue to be a major cause of morbidity and leading indication for hysterectomy in premenopausal women. And some of the leiomyoma variants, such as mitotically active leiomyoma, symplastic leiomyoma may simulate leiomyosarcoma or other malignant tumors as these tumors carry better prognosis and require detailed morphological study. In this study, we ascertained the frequency with which these variants were diagnosed and the difficulties that were encountered during the diagnosis.

Materials and methods: A retrospective analysis of hysterectomy and myomectomy specimen that were received in our department, Institute of Social Obstetrics and Kasturba Gandhi Hospital for women and children, from April 2016 to April 2017 was performed. The details regarding patient age, clinical presentation were collected from the medical records.

Results: During our study period, we received 1879 hysterectomy specimens and 664 specimens were found to have fibroid. We also received 29 myomectomy specimens and all these specimens were subjected for routine histopathological examination. Typical leiomyomas constituted to 88.6% and other variants like mitotically active leiomyoma, symplastic leiomyoma, cellular leiomyoma, neurilemmoma like variant, fatty degeneration, myxoid degeneration, hyaline degeneration, calcification and hydropic degeneration constituted 11.4%.

Conclusion: Variants of leiomyoma were relatively less, but it is important to differentiate them from malignant myometrial neoplasms, as they have good prognosis. Tumor size, cytological and nuclear atypia, mitosis and coagulative necrosis were the most important histological features for differentiation.
changes provide the basis for the classification of most benign smooth muscle tumors of uterus. However, diagnostic dilemmas were encountered with leiomyoma variants because of their wide spectrum of gross and microscopic appearances often causing concern for malignancy. Thus, a correct classification was important as currently prognosis of its malignant counterpart, leiomyosarcoma, is dismal even when confined to the uterus at time of diagnosis.

**Objective:**
The purpose of this study was to evaluate the histomorphological features of uterine leiomyoma variants from hysterectomy and myomectomy specimens.

**Materials and Methods:**
In this study, we performed retrospective analysis of hysterectomy and myomectomy specimen from April 2016 to April 2017 in Institute of Social Obstetrics and Kasturba Gandhi Hospital for women and children and details regarding patient age, clinical presentation were collected.

A detailed macroscopic examination, including size, appearance and external surface were noted. The specimens were allowed to fix in 10% neutral buffered formalin for 24 to 48 hours. Multiple parallel sections were made and their cut surface was examined. The tissue bits from representative areas were taken for histopathological examination. Multiple sections of five microns thickness were cut and routinely stained with hematoxylin and eosin.

**Results:**
We received 3319 specimens, among which 1879 were hysterectomy specimen done for varying indications and 29 were myomectomy specimen (including 4 cases of pregnancy associated fibroid).

Among 1879 hysterectomy specimen, 664 specimens were found to have fibroid and most of them were intramural in location and few located subserosally. Most fibroids were diagnosed clinically and some were incidental findings. Patients who underwent myomectomy (29 specimen) were in their reproductive age and found to have fundal fibroid, thus hysterectomy was deferred.

Most patients were in the premenopausal age group and had varied clinical presentation ranging from menorrhagia, mass abdomen, pain abdomen, and dysmenorrhea. Patients with fundal fibroid presented with primary infertility, recurrent abortions and 4 cases of fibroid complicating pregnancy were noted.

**Table 1:** Clinical presentation of the patients

<table>
<thead>
<tr>
<th>Clinical presentation</th>
<th>Number of women</th>
<th>Percentage of women</th>
</tr>
</thead>
<tbody>
<tr>
<td>Menorrhagia</td>
<td>293</td>
<td>42.27%</td>
</tr>
<tr>
<td>Mass abdomen</td>
<td>89</td>
<td>12.84%</td>
</tr>
<tr>
<td>Pain abdomen</td>
<td>52</td>
<td>7.50%</td>
</tr>
<tr>
<td>Dysmenorrhea</td>
<td>41</td>
<td>5.91%</td>
</tr>
<tr>
<td>Primary infertility</td>
<td>2</td>
<td>0.33%</td>
</tr>
<tr>
<td>Recurrent abortion</td>
<td>3</td>
<td>0.43%</td>
</tr>
<tr>
<td>Pregnancy associated</td>
<td>4</td>
<td>0.57%</td>
</tr>
<tr>
<td>Asymptomatic/incidental</td>
<td>209</td>
<td>30.15%</td>
</tr>
</tbody>
</table>

**Table 2:** Histomorphological variants of leiomyoma.

<table>
<thead>
<tr>
<th>Variant of leiomyoma</th>
<th>Number of cases</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Typical leiomyoma</td>
<td>614</td>
<td>88.6%</td>
</tr>
<tr>
<td>Mitotically active</td>
<td>3</td>
<td>0.43%</td>
</tr>
<tr>
<td>Hydropic change</td>
<td>1</td>
<td>0.14%</td>
</tr>
<tr>
<td>Fatty degeneration</td>
<td>9</td>
<td>1.29%</td>
</tr>
<tr>
<td>Hyaline degeneration</td>
<td>44</td>
<td>6.34%</td>
</tr>
<tr>
<td>Calcific degeneration</td>
<td>12</td>
<td>1.73%</td>
</tr>
<tr>
<td>Symplastic leiomyoma</td>
<td>2</td>
<td>0.28%</td>
</tr>
<tr>
<td>Cellular leiomyoma</td>
<td>3</td>
<td>0.43%</td>
</tr>
<tr>
<td>Neurilemmoma like</td>
<td>1</td>
<td>0.14%</td>
</tr>
</tbody>
</table>
Myxoid degeneration | 4 | 0.62%

Fig 1: Gross image of symplastic leiomyoma showing well circumscribed mass which was grey white, firm without areas of necrosis and hemorrhage.

Fig 2: Gross image of leiomyoma with myxoid areas.
Fig 3: Symplastic leiomyoma-Scanner view (40X) showing well circumscribed neoplasm arranged in fascicles & bundles. Inset showing pleomorphic bizarre hyperchromatic nuclei.
Fig 4:- Cellular leiomyoma- Low power view (100 X) showing cellular neoplasm composed of spindle cells.

Fig 5:- High power view (400 X) showing calcific degeneration of leiomyoma.
Fig6: High power view (400 X) showing myxoid degeneration of leiomyoma.

Fig7: Low power view (100 X) showing hydropic degeneration of leiomyoma.
Fig8: High power view (400 X) showing fatty degeneration of leiomyoma.

Fig9: Low power view (100 X) showing neurilemmoma like variant of leiomyoma.
**Discussion:**
Leiomyoma was the most common uterine neoplasms.\(^{(6,7)}\) Based on location, they have been classified as submucosal, intramural, subserosal leiomyoma. They were noted clinically in 20-30% of women over 30 years of age.\(^{(8,9)}\) Most leiomyomas were detected in middle aged women.
In this study, the mean age of diagnosis was around 23-45 years. Most of the patients were asymptomatic and some had clinical presentation ranging from menorrhagia, pain abdomen, mass abdomen, dysmenorrheal and few presented with primary infertility, recurrent abortions. Four cases were found to be associated with pregnancy and these patients underwent myomectomy during their second trimester. Premenopausal patients with fibroid underwent hysterectomy.

The mean size of leiomyoma was between 1-15cm. Most specimens revealed characteristic macroscopic appearance. Typical leiomyomas constituted about 88.4% and variants constituted to about 11.6% which includes mitotically active leiomyoma, cellular leiomyoma, atypical/symplastic leiomyoma, leiomyoma with fatty degeneration, calcific degeneration, hydropic degeneration, hyaline degeneration, neurilemmoma like leiomyoma.

Most of these variants carry no clinical significance, yet their correct diagnosis and differentiation from malignant myometrial neoplasms were important because of their good prognosis.

In this study, variants of leiomyoma constituted about 11.6%. Hyaline degeneration was the common finding and it was associated with large tumor size.

Other degenerative changes include red degeneration, calcific degeneration, fatty degeneration and hydropic degeneration.

Mitotically active leiomyoma:-
We reported three cases of mitotically active leiomyoma. All these cases were reported in premenopausal women and they found to have 6-9 mitosis/10 high power field. Thus, mitotically active leiomyomas usually have more than or equal to 5 mitotic figures per 10 high power field.\(^{(10,11,12,13)}\) It is imperative that this diagnosis not be used for neoplasms that exhibit moderate to severe nuclear atypia, for those that contain abnormal mitotic figure or for those that demonstrate zones of tumor cell necrosis.

Cellular leiomyoma:-
A cellular leiomyoma is one in which the cellularity is significantly greater than the surrounding myometrium. We reported three cases of cellular leiomyoma.

The isolated finding of hypercellularity may suggest a diagnosis of leiomyosarcoma, but cellular leiomyoma lacks tumor cell necrosis, has few mitotic figures and lacks the moderate to severe cytological atypia seen in leiomyosarcoma. A cellular leiomyoma composed of small cells with scant cytoplasm can be confused with an endometrial stromal tumor.

Features that help distinguish a cellular leiomyoma from a stromal tumor are the spindled shape of the cell, the fusiform shape of the nuclei, the reticulin pattern and absence of a plexiform vasculature.

Reticulin fibres tend to parallel the fascicles of cells in a leiomyoma, but reticulin network surrounds individual tumor cells in an endometrial stromal nodule.

Large thick walled muscular vessels serve to distinguish a cellular leiomyoma from an endometrial stromal tumor.\(^{(14)}\)

Symplastic / atypical leiomyoma:-
As an isolated finding, cytological atypia, even when severe, is an unreliable criterion for the diagnosis of clinically malignant uterine smooth muscle tumors because it can be seen in clinically benign otherwise banal smooth muscle neoplasm.\(^{(15)}\)

We reported two cases of symplastic leiomyoma. Both the patients were in the premenopausal age group. Their clinical presentation was dysmenorrhea and menorrhagia. Gross examination of both the specimens revealed a well circumscribed grey white, firm, homogenous mass. There were no areas of necrosis, hemorrhage. Microscopic examination revealed a well circumscribed neoplasm arranged in fascicles and bundles composed of spindle shaped cells with moderate eosinophilic cytoplasm with pleomorphic hyperchromatic nuclei. Extensive sampling was done.
and it revealed mitosis less than five mitotic figures per 10 high power field. There were no areas of necrosis, hemorrhage or zone of tumor cell necrosis seen.

Neurilemmoma like leiomyoma:-
A case of neurilemmoma like leiomyoma was diagnosed in 37 year old female, who presented with excessive bleeding. The tumor was intramural in location, measured 4.5 cm in diameter. Histological examination showed fascicular pattern of leiomyoma with areas showing nuclear palisading mimicking a schwannoma (16).

Outcome:-
Most of the patients were followed up and their follow up period was uneventful.

Conclusion:-
Most subtypes of leiomyoma were chiefly of clinical interest in that they mimic malignancy in one or more respects. Though the frequency with which they occur remains less, their correct diagnosis was essential and they need to be differentiated from malignant myometrial neoplasms as most of these variants carry good prognosis and of less clinical significance.

References:-