Protocol for the Examination of Specimens From Patients With Sarcoma of the Uterus

Protocol applies to sarcomas of the uterus.

Based on AJCC/UICC TNM, 7th edition, and FIGO 2009 Annual Report
Protocol web posting date: January 2016

Procedures
- Hysterectomy
- Myomectomy

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For the Members of the Cancer Committee, College of American Pathologists
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CAP Uterine Sarcoma Protocol Revision History

Version Code
The definition of version control and an explanation of version codes can be found at www.cap.org (search: cancer protocol terms).

Version: UterineSarcoma 3.1.0.0

Summary of Changes
The following changes have been made since the December 2013 release.

The following data elements were modified:
- Histologic Type
- Peritoneal Ascitic Fluid
- Lymph Notes
- Distant Metastasis (changed to required only if confirmed pathologically)

The following data element was added:
- FIGO Stage (not required)
Surgical Pathology Cancer Case Summary

Protocol web posting date: January 2016

UTERUS: Hysterectomy and Myomectomy, With or Without Other Organs or Tissues

Select a single response unless otherwise indicated.

Specimen (select all that apply)
___ Uterine corpus
___ Cervix
___ Right ovary
___ Left ovary
___ Right fallopian tube
___ Left fallopian tube
___ Left parametrium
___ Right parametrium
___ Peritoneum
___ Vaginal cuff
___ Omentum
___ Other (specify): ___________________________
___ Not specified

Procedure (select all that apply)
___ Supracervical hysterectomy
___ Simple hysterectomy
___ Radical hysterectomy
___ Myomectomy
___ Right oophorectomy
___ Left oophorectomy
___ Right salpingectomy
___ Left salpingectomy
___ Right salpingo-oophorectomy
___ Left salpingo-oophorectomy
___ Omentectomy
___ Peritoneal biopsies
___ Other (specify): ____________________________
___ Not specified

Lymph Node Sampling (select all that apply)
___ Performed
   ___ Pelvic lymph nodes
   ___ Para-aortic lymph nodes
   ___ Other (specify): ____________________________
___ Not performed
___ Not known

Specimen Integrity
___ Hysterectomy specimen (intact)
___ Hysterectomy specimen without cervix
___ Morcellated hysterectomy specimen
___ Myomectomy (intact)
___ Morcellated myomectomy specimen
___ Other (specify): ____________________________

+ Data elements preceded by this symbol are not required. However, these elements may be clinically important but are not yet validated or regularly used in patient management.
Tumor Site
___ Fundus
___ Lower uterine segment/isthmus
___ Cervix
___ Other (specify): ______________________________

Tumor Size
Greatest dimension: ___ cm
+ Additional dimensions: ___ x ___ cm
___ Cannot be determined

Histologic Type (select all that apply) (Notes B, C, D)
___ Leiomyosarcoma
   + ___ Epithelioid type
   + ___ Myxoid type
   + ___ Other (specify): ____________________________________
___ Low-grade endometrial stromal sarcoma
___ Low-grade endometrial stromal sarcoma with:
   ___ Smooth muscle differentiation
   ___ Sex cord elements
   ___ Glandular elements
   ___ Other (specify): ____________________________________
___ High-grade endometrial stromal sarcoma
___ Undifferentiated uterine/endometrial sarcoma
___ Adenosarcoma
___ Adenosarcoma with:
   ___ Rhabdomyoblastic differentiation
   ___ Cartilagenous differentiation
   ___ Osseous differentiation
   ___ Other heterologous element (specify): ______________________________
___ Adenosarcoma with sarcomatous overgrowth
___ Other (specify): ____________________________________

# Low-grade endometrial stromal sarcoma is distinguished from benign endometrial stromal nodule by depth of myometrial invasion ≥3 mm, lymphovascular invasion, or ≥3 foci of myometrial invasion of any depth.

Minor marginal irregularity in the form of tongues ≤3 mm is allowable for an endometrial stromal nodule. This protocol does not apply to endometrial stromal nodule.

Histologic Grade (select grade as specifically applicable to histologic type)

Leiomyosarcoma (Note D)
___ Not applicable

Endometrial Stromal Sarcoma (Note C)
___ Low grade
___ High grade
___ Cannot be assessed

Adenosarcoma (select all that apply) (Note B)
___ Low grade
___ High grade
___ With sarcomatous overgrowth
___ Cannot be assessed

+ Data elements preceded by this symbol are not required. However, these elements may be clinically important but are not yet validated or regularly used in patient management.
Myometrial Invasion (only for adenosarcoma) (select all that apply) (Note D)
____ Not identified
____ Present
   Depth of invasion: ____ mm
   OR, if exact depth of invasion cannot be determined, state:
   ____ Extent of myometrial invasion cannot be determined (explain): _______________________
   ____ <50% myometrial invasion
   ____ ≥50% myometrial invasion
   Myometrial thickness: ____ mm
   ____ Myometrial thickness cannot be determined (explain): _______________________

Involvement of Cervix
____ Cannot be determined
____ Not involved
____ Tumor involves the glandular surface of the cervix only
____ Tumor invades the cervical stromal connective tissue

Extent of Involvement of Other Organs (select all that apply)
____ Not applicable
____ Right ovary
   ____ Involved
   ____ Not involved
____ Left ovary
   ____ Involved
   ____ Not involved
____ Right fallopian tube
   ____ Involved
   ____ Not involved
____ Left fallopian tube
   ____ Involved
   ____ Not involved
____ Vaginal cuff
   ____ Involved
   ____ Not involved
____ Right parametrium
   ____ Involved
   ____ Not involved
____ Left parametrium
   ____ Involved
   ____ Not involved
____ Omentum
   ____ Involved
   ____ Not involved
____ Other (specify): _______________________

Margins
____ Cannot be assessed
____ Uninvolved by sarcoma
   + Distance of sarcoma from closest margin: ____ mm
   + Specify margin: _______________________
____ Involved by sarcoma
   Specify margin(s): _______________________

Data elements preceded by this symbol are not required. However, these elements may be clinically important but are not yet validated or regularly used in patient management.
Lymph-Vascular Invasion
___ Not identified
___ Present
___ Indeterminate

+ Peritoneal Ascitic Fluid
+ ___ Not performed/unknown
+ ___ Negative for malignancy (normal/benign)
+ ___ Atypical and/or suspicious (explain): ______________________________
+ ___ Malignant (positive for malignancy)
+ ___ Unsatisfactory/nondiagnostic (explain): _____________________________

+ Peritoneal Washing
+ ___ Negative for malignancy
+ ___ Atypical and/or suspicious (explain): ______________________________
+ ___ Malignant (positive for malignancy)
+ ___ Unsatisfactory/nondiagnostic (explain): _____________________________

Pathologic Staging (pTNM)

TNM Descriptors (required only if applicable) (select all that apply)
___ m (multiple primary tumors)
___ r (recurrent)
___ y (posttreatment)

Leiomyosarcoma, Endometrial Stromal Sarcoma, and Undifferentiated Endometrial Sarcoma/Uterine Sarcoma

Primary Tumor (pT)
___ pTX: Primary tumor cannot be assessed
___ pT0: No evidence of primary tumor
pT1: Tumor is limited to the uterus
___ pT1a: Tumor is 5 cm or less (≤5 cm) in greatest dimension
___ pT1b: Tumor is greater than 5 cm (>5 cm) in greatest dimension
pT2: Tumor extends beyond the uterus, but is within the pelvis (tumor extends to extrauterine pelvic tissue)
___ pT2a: Tumor involves the adnexa
___ pT2b: Tumor involves other pelvic tissue
pT3: Tumor invades abdominal tissues (not just protruding into the abdomen)
___ pT3a: Tumor invades abdominal tissues at one site
___ pT3b: Tumor invades abdominal tissues at more than one site
___ pT4: Tumor invades bladder mucosa and/or rectum

Regional Lymph Nodes (pN) (select all that apply)
___ pNX: Cannot be assessed
___ pN0: No regional lymph node metastasis
___ pN1: Regional lymph node metastasis to pelvic lymph nodes
___ No nodes submitted or found

Pelvic lymph nodes:
___ No pelvic nodes submitted or found

Number of Pelvic Lymph Nodes Examined
Specify: ______
___ Number cannot be determined (explain): _____________________________

+ Data elements preceded by this symbol are not required. However, these elements may be clinically important but are not yet validated or regularly used in patient management.
Number of Pelvic Lymph Nodes Involved
Specify: ____
___ Number cannot be determined (explain): ________________________________

Para-aortic lymph nodes:
___ No para-aortic nodes submitted or found

Number of Para-aortic Lymph Nodes Examined
Specify: ____
___ Number cannot be determined (explain): ________________________________

Number of Para-aortic Lymph Nodes Involved
Specify: ____
___ Number cannot be determined (explain): ________________________________

Other lymph nodes:
Specify site: ________________________________

Number of Other Lymph Nodes Examined
Specify: ____
___ Number cannot be determined (explain): ________________________________

Number of Other Lymph Nodes Involved
Specify: ____
___ Number cannot be determined (explain): ________________________________

Distant Metastasis (pM) (required only if confirmed pathologically in this case)
___ pM1: Distant metastasis (excluding adnexa, pelvic and abdominal tissues)
   Specify site(s), if known: ________________________________

+ FIGO Stage
  + I: Tumor limited to uterus
    + ___ IA: Less than or equal to 5 cm
    + ___ IB: More than 5 cm
  + II: Tumor extends beyond the uterus, within the pelvis
    + ___ IIA: Adnexal involvement
    + ___ IIB: Involvement of other pelvic tissues
  + III: Tumor invades abdominal tissues (not just protruding into the abdomen)
    + ___ IIIA: 1 site
    + ___ IIIB: More than 1 site
    + ___ IIIC: Metastasis to pelvic and/or para-aortic lymph nodes
  + IV: Tumor invades bladder and/or rectum and/or distant metastasis
    + ___ IVA: Tumor invades bladder and/or rectum
    + ___ IVB: Distant metastasis

+ Data elements preceded by this symbol are not required. However, these elements may be clinically important but are not yet validated or regularly used in patient management.
Adenosarcoma

Primary Tumor (pT)
___ pTX: Primary tumor cannot be assessed
___ pT0: No evidence of primary tumor
pT1: Tumor is limited to the uterus
___ pT1a: Tumor is limited to the endometrium/endocervix without myometrial invasion
___ pT1b: Tumor invades less than or equal to 50% (≤50%) total myometrial thickness
___ pT1c: Tumor invades greater than 50% (>50%) total myometrial thickness
pT2: Tumor extends beyond the uterus, but is within the pelvis (tumor extends to extrauterine pelvic tissue)
___ pT2a: Tumor involves the adnexa
___ pT2b: Tumor involves other pelvic tissue
pT3: Tumor invades abdominal tissues (not just protruding into the abdomen)
___ pT3a: Tumor invades abdominal tissues at one site
___ pT3b: Tumor invades abdominal tissues at more than one site
___ pT4: Tumor invades bladder mucosa and/or rectum

Regional Lymph Nodes (pN)
___ pNX: Cannot be assessed
___ pN0: No regional lymph node metastasis
___ pN1: Regional lymph node metastasis to pelvic lymph nodes
___ No nodes submitted or found

Pelvic lymph nodes:
___ No pelvic nodes submitted or found

Number of Pelvic Lymph Nodes Examined
Specify: ____
___ Number cannot be determined (explain): ______________________

Number of Pelvic Lymph Nodes Involved
Specify: ____
___ Number cannot be determined (explain): ______________________

Para-aortic lymph nodes:
___ No para-aortic nodes submitted or found

Number of Para-aortic Lymph Nodes Examined
Specify: ____
___ Number cannot be determined (explain): ______________________

Number of Para-aortic Lymph Nodes Involved
Specify: ____
___ Number cannot be determined (explain): ______________________

Other lymph nodes (specify site): ______________________:

Number of Lymph Nodes Examined
Specify: ____
___ Number cannot be determined (explain): ______________________

Number of Lymph Nodes Involved
Specify: ____

+ Data elements preceded by this symbol are not required. However, these elements may be clinically important but are not yet validated or regularly used in patient management.
Number cannot be determined (explain): ____________________________

Distant Metastasis (pM) (required only if confirmed pathologically in this case)

___ pM1: Distant metastasis (excluding adnexa, pelvic and abdominal tissues)
  Specify site(s), if known: ______________________________

+ FIGO Stage
+ I: Tumor limited to uterus
  + ___ IA: Tumor limited to endometrium/endocervix with no myometrial invasion
  + ___ IB: Less than or equal to half myometrial invasion
  + ___ IC: More than half myometrial invasion
+ II: Tumor extends beyond the uterus, within the pelvis
  + ___ IIA: Adnexal involvement
  + ___ IIB: Tumor extends to extraperitoneal pelvic tissue
+ III: Tumor invades abdominal tissues (not just protruding into the abdomen).
  + ___ IIIA: 1 site
  + ___ IIIB: More than 1 site
  + ___ IIIC: Metastasis to pelvic and/or para-aortic lymph nodes
+ IV: Tumor invades bladder and/or rectum and/or distant metastasis
  + ___ IVA: Tumor invades bladder and/or rectum
  + ___ IVB: Distant metastasis

+ Ancillary Studies (Note B)
+ Specify: _____________________________________________
  + _____ Not performed

+ Comment(s)
Explanatory Notes

A. Carcinosarcoma
Carcinosarcoma (malignant mixed mullerian tumor) is excluded from the uterine sarcoma diagnostic category as it is considered in tumors of the endometrial epithelium.

B. Adenosarcoma
According to World Health Organization (WHO) criteria, mitotic activity in the mesenchymal component in excess of 2 or more per 10 high-power fields (HPFs) is required for a diagnosis of adenosarcoma, but others use a cutoff of 4 per 10 HPFs. However, given the multiple and well-known problems associated with counting mitotic figures and the fact that the number of mitoses may be variable from area to area, in practice, if the characteristic leaf-like architecture of adenosarcoma is present with periglandular cuffing resulting in a cambium layer, a diagnosis of adenosarcoma should be strongly considered with mitotic counts <2 per 10 HPFs or even in the absence of mitotic figures. In adenosarcomas without sarcomatous overgrowth, it is recommended to record on the pathology report whether the stromal component is morphologically “low grade” or “high grade.” Even though there are no studies showing that this is of prognostic significance, anecdotal evidence suggests that even a small focus of “high-grade” sarcoma may result in an adverse behavior. It is suggested that the parameter of nuclear atypia be used to distinguish between low grade and high grade. In low-grade neoplasms, the atypia should be akin to that seen in low-grade endometrial stromal sarcoma. Sarcomatous overgrowth in adenosarcoma is defined as the presence of pure sarcoma, usually high grade and without an epithelial component, occupying at least 25% of the tumor.

Adenosarcomas rarely exhibit lymphovascular invasion unless associated with deep myometrial invasion or sarcomatous overgrowth.

The depth of myometrial invasion is important in the substaging of stage I adenosarcomas (tumor confined to the uterus). Stage IA tumors are limited to the endometrium or endocervix with no myometrial involvement, stage IB equates to less than or half of myometrial invasion, and stage IC equates to more than one-half myometrial invasion. This staging system is similar to the 1988 FIGO staging system for carcinomas of the uterine corpus. Since low-grade endometrial stromal sarcoma and leiomyosarcoma are predominantly myometrial-based lesions, myometrial invasion per se is not used in the staging of these neoplasms.

In most adenosarcomas with a low-grade stromal component without sarcomatous overgrowth, the stromal element expresses estrogen receptor (ER), progesterone receptor (PgR), CD10, and WT1; is negative (“wild-type”) with p53; and exhibits a low MIB1 proliferation index. Thus, the immunophenotype resembles that of low-grade endometrial stromal sarcoma. Smooth muscle actin and desmin may also be positive. In areas of high-grade sarcoma and of sarcomatous overgrowth, the mesenchymal component exhibits a higher MIB1 proliferation index and may be p53 positive/aberrant. There is usually loss of expression of the cell differentiation markers ER, PgR, and CD10, the immunophenotype being similar to that of an undifferentiated sarcoma. Rhabdomyosarcomatous elements in adenosarcomas express desmin and sometimes the skeletal muscle markers myogenin and myoD1. Sex cord-like elements may express inhibin and calretinin.

C. Endometrial Stromal Sarcoma
Low-grade endometrial stromal sarcoma, in contrast to endometrial stromal nodule, demonstrates myometrial invasion from the nodule or tumor mass of ≥3 mm, lymphovascular invasion, or ≥3 foci of myometrial invasion of any depth.

Even though in the past endometrial stromal sarcomas (ESS) were classified as low grade (LG) and high grade (HG) based on mitotic activity, the largest and most comprehensive review of these tumors by Chang and colleagues in 1990 showed that mitotic activity was not predictive of outcome in stage I tumors. Thus, the diagnosis of HG-ESS was discouraged in those tumors that resemble proliferative-phase endometrial stroma but in which the mitotic index exceeded 10 per 10 HPF. Currently many expert gynecologic pathologists, without any proven basis outside of personal experience, make the diagnosis of HG-ESS when there is a transition from high-grade undifferentiated sarcoma to areas that can be recognized as conventional LG-ESS. However, recently, a subset of cases previously diagnosed as HG-ESSs has been histologically and genetically defined by Lee, Nucci.
and colleagues.\textsuperscript{10,11} In these tumors, the high-grade areas are characterized by cells with a round cell-epithelioid appearance and high-grade cytologic features which often are associated with areas that have the appearance of the fibroblastic variant of low-grade conventional ESS.\textsuperscript{10} These tumors have been shown to have a novel genetic fusion between \textit{YWHAE} and \textit{FAM22A/B} and harbor t(10;17)(q22;p13). The high-grade areas of the tumor express cyclin D1 but lose CD10, ER, and PgR expression (in contrast to the conventional low-grade areas) consistent with a high-grade sarcoma.\textsuperscript{10} It is important to recognize these tumors as they have an intermediate prognosis between LG-ESS and undifferentiated uterine sarcoma (UUS) and appear not to respond to the usual treatment for low-grade ESS.

Low-grade ESS, high-grade ESS, and UUS all exist and should be separately diagnosed, although UUS should be a diagnosis of exclusion (leiomyosarcomas and other high-grade sarcomas, for example rhabdomyosarcoma, should be excluded). Molecular testing is diagnostically unnecessary in conventional ESS and in USS, but is useful in confirming the diagnosis of HG-ESS in tumors with a round cell-epithelioid appearance that can be associated with areas that have the appearance of the fibroblastic variant of conventional LG-ESS.

D. Leiomyosarcoma

By definition, uterine leiomyosarcoma (LMS) is a highly malignant neoplasm with survival rates depending upon the extent of spread. For tumors confined to the uterine corpus, size plays a significant role in prognosis. Despite differences in survival rates, it is clear that stage is a significant factor related to outcome. Histologic grade, however, has not been consistently identified as a significant prognostic parameter.\textsuperscript{12} The utility of grading uterine LMS is controversial, and no universally accepted grading system exists.\textsuperscript{5} In 2011, Veras et al\textsuperscript{13} tried to characterize "low-grade uterine leiomyosarcomas" as a clinicopathological entity but came to the conclusion that this can be diagnosed only retrospectively at present.\textsuperscript{13} Furthermore, when the Stanford criteria are strictly applied, all tumors classified as leiomyosarcomas should be regarded intrinsically as high grade.\textsuperscript{13,14}

Conventional uterine LMS is a cellular tumor composed of fascicles of spindle-shaped cells exhibiting smooth muscle differentiation with moderate to severe pleomorphism. Usually coagulative tumor cell necrosis (CTCN) is present and mitoses exceed 10 to 15 per 10 HPF.\textsuperscript{14} Two LMS subtypes included in the WHO classification deserve special attention as their pathologic features differ from those of ordinary spindle cell LMS. Epithelioid leiomyosarcoma (E-LMS) is composed predominantly of round or polygonal cells with eosinophilic to clear cytoplasm exhibiting nested, plexiform or cored growth patterns. Nuclear atypia may be only mild and necrosis may be absent. Mitotic rate is generally \( \leq 3 \) per 10 HPF, and most tumors infiltrate adjacent myometrium. Myxoid leiomyosarcoma (M-LMS) may be grossly gelatinous, microscopically hypocellular with a predominant myxoid stroma, and often has a low mitotic rate. In the absence of severe cytologic atypia and high mitotic activity, both epithelioid and myxoid LMS are diagnosed as sarcomas based on their infiltrative borders.\textsuperscript{12}

Ancillary Studies in the Differential Diagnosis

Immunoreactivity for smooth muscle actin, muscle specific actin, calponin, desmin, h-caldesmon, and heavy chain smooth muscle myosin are commonly seen in uterine LMS. Desmin expression may be focal.\textsuperscript{15-17} Similarly, E-LMS and M-LMS may demonstrate lesser degrees of immunoreactivity for these markers. Cell cycle related markers Ki-67, p53, and p16 are usually overexpressed in LMS compared to leiomyoma.\textsuperscript{18} Cytokeratins and EMA may be focally positive in LMS, especially in the epithelioid variant.

E. Undifferentiated Uterine/Endometrial Sarcoma

Undifferentiated uterine/endometrial sarcoma (UUS) is a high-grade sarcoma that lacks specific differentiation. Histopathologically these tumors show marked cellular pleomorphism and abundant mitotic activity with atypical forms. They lack the typical growth pattern and vascularity of low-grade ESS and displace the myometrium in contrast to the infiltrative pattern of low-grade ESS. They often resemble the sarcomatous component of a carcinosarcoma. These sarcomas are most often aneuploid with an S-phase fraction greater than 10%, and are negative for ER and PgR.\textsuperscript{10} Nucci et al proposed that high-grade ESS with the novel fusion gene \textit{YWHAE-FAM22} should be distinguished from undifferentiated uterine/endometrial sarcoma.\textsuperscript{16}

F. Other

Other differential diagnostic considerations included in spindle/sarcomatous lesions primary to the uterus include perivascular epithelioid cell tumor (PEComa). PEComa belongs to a group of tumors characterized by both
melanocytic and smooth muscle differentiation, and should be recognized separately from smooth muscle
tumors.\textsuperscript{19,21}

The TNM staging system for uterine sarcoma endorsed by the American Joint Committee on Cancer (AJCC) and
the parallel system formulated by the International Federation of Gynecology and Obstetrics (FIGO) are
recommended, as shown below.

According to AJCC/International Union Against Cancer (UICC) convention, the designation “T” refers to a primary
tumor that has not been previously treated. The symbol “p” refers to the pathologic classification of the TNM, as
opposed to the clinical classification, and is based on gross and microscopic examination. pT entails a resection
of the primary tumor or biopsy adequate to evaluate the highest pT category, pN entails removal of nodes
adequate to validate lymph node metastasis, and pM implies microscopic examination of distant lesions. Clinical
classification (cTNM) is usually carried out by the referring physician before treatment during initial evaluation of
the patient or when pathologic classification is not possible.

Pathologic staging is usually performed after surgical resection of the primary tumor. Pathologic staging depends
on pathologic documentation of the anatomic extent of disease, whether or not the primary tumor has been
completely removed. If a biopsied tumor is not resected for any reason (eg, when technically infeasible) and if the
highest T and N categories or the M1 category of the tumor can be confirmed microscopically, the criteria for
pathologic classification and staging have been satisfied without total removal of the primary cancer.

It is important to note that in uterine sarcoma, as in cancer of other organs, the validity of T stage depends upon
the adequacy and completeness of the surgical staging.

| TNM Classification and FIGO Staging System for Leiomyosarcoma, Endometrial Stromal Sarcoma and
Undifferentiated Uterine Sarcoma |
<table>
<thead>
<tr>
<th></th>
<th></th>
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</thead>
<tbody>
<tr>
<td>Primary Tumor</td>
<td>FIGO</td>
</tr>
<tr>
<td>Category</td>
<td>Stage</td>
</tr>
<tr>
<td>---------</td>
<td>-------</td>
</tr>
<tr>
<td>pTX</td>
<td>[-]</td>
</tr>
<tr>
<td>pT0</td>
<td>[-]</td>
</tr>
<tr>
<td>pT1</td>
<td>[I]</td>
</tr>
<tr>
<td>pT1a</td>
<td>[IA]</td>
</tr>
<tr>
<td>pT1b</td>
<td>[IB]</td>
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<td>[II]</td>
</tr>
<tr>
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<td>[IIIA]</td>
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<tr>
<td>pT2b</td>
<td>[IIIB]</td>
</tr>
<tr>
<td>pT3</td>
<td>[III]</td>
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<tr>
<td>pT3a</td>
<td>[IIIA]</td>
</tr>
<tr>
<td>pT3b</td>
<td>[IIIB]</td>
</tr>
<tr>
<td>pT4</td>
<td>[IVA]</td>
</tr>
</tbody>
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Regional Lymph Nodes (pN)*

<table>
<thead>
<tr>
<th>Category</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>pNX</td>
<td>Cannot be assessed</td>
</tr>
<tr>
<td>pN0</td>
<td>No regional lymph node metastasis</td>
</tr>
<tr>
<td>pN1</td>
<td>Regional lymph node metastasis to pelvic lymph nodes</td>
</tr>
</tbody>
</table>

* Regional lymph nodes include the pelvic, obturator, internal iliac (hypogastric), external iliac, common iliac, para-
aortic, presacral, and parametrial lymph nodes.

Distant Metastasis (pM)

<table>
<thead>
<tr>
<th>Category</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>pM0</td>
<td>No distant metastasis (no pathologic M0; use clinical M to complete stage group)</td>
</tr>
<tr>
<td>pM1</td>
<td>[IVB] Distant metastasis (excluding adnexa, pelvic and abdominal tissues)</td>
</tr>
</tbody>
</table>
### Adenosarcoma

<table>
<thead>
<tr>
<th>TNM Category</th>
<th>FIGO Stage</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>pTX</td>
<td>[--]:</td>
<td>Primary tumor cannot be assessed</td>
</tr>
<tr>
<td>pT0</td>
<td>[--]:</td>
<td>No evidence of primary tumor</td>
</tr>
<tr>
<td>pT1</td>
<td>[I]:</td>
<td>Tumor is limited to the uterus</td>
</tr>
<tr>
<td>pT1a</td>
<td>[IA]:</td>
<td>Tumor is limited to the endometrium/endocervix without myometrial invasion</td>
</tr>
<tr>
<td>pT1b</td>
<td>[IB]:</td>
<td>Tumor invades less than or equal to 50% (≤50%) total myometrial thickness</td>
</tr>
<tr>
<td>pT1c</td>
<td>[IC]:</td>
<td>Tumor invades greater than 50% (&gt;50%) total myometrial thickness</td>
</tr>
<tr>
<td>pT2</td>
<td>[II]:</td>
<td>Tumor extends beyond the uterus, but is within the pelvis (tumor extends to extrauterine pelvic tissue)</td>
</tr>
<tr>
<td>pT2a</td>
<td>[IIA]:</td>
<td>Tumor involves the adnexa</td>
</tr>
<tr>
<td>pT2b</td>
<td>[IIB]:</td>
<td>Tumor involves other pelvic tissue</td>
</tr>
<tr>
<td>pT3</td>
<td>[III]:</td>
<td>Tumor invades abdominal tissues (not just protruding into the abdomen)</td>
</tr>
<tr>
<td>pT3a</td>
<td>[IIIA]:</td>
<td>Tumor invades abdominal tissues at one site</td>
</tr>
<tr>
<td>pT3b</td>
<td>[IIIB]:</td>
<td>Tumor invades abdominal tissues at more than one site</td>
</tr>
<tr>
<td>pT4</td>
<td>[IVA]:</td>
<td>Tumor invades bladder mucosa and/or rectum</td>
</tr>
</tbody>
</table>

#### Regional Lymph Nodes (pN) #

| pNX          | Cannot be assessed |
| pN0          | No regional lymph node metastasis |
| pN1          | Regional lymph node metastasis to pelvic lymph nodes |

# Regional lymph nodes include the pelvic, obturator, internal iliac (hypogastric), external iliac, common iliac, para-aortic, presacral, and parametrial lymph nodes.

#### Distant Metastasis (pM)

| pM0          | No distant metastasis (no pathologic M0; use clinical M to complete stage group) |
| pM1          | Distant metastasis (excluding adnexa, pelvic and abdominal tissues) |

### TNM Stage Groupings

- **Stage 0**: Tis N0 M0
- **Stage IA #**: T1a N0 M0
- **Stage IB #**: T1b N0 M0
- **Stage IC # #**: T1c N0 M0
- **Stage II**: T2 N0 M0
- **Stage IIIA**: T3a N0 M0
- **Stage IIIB**: T3b N0 M0
- **Stage IIIC**: T1-T3 N1 M0
- **Stage IVA**: T4 Any N M0
- **Stage IVB**: Any T Any N M1

# Stage IA and IB for adenosarcoma differ from those applied to leiomyosarcoma and endometrial stromal sarcoma.

# # Stage IC does not apply for leiomyosarcoma and endometrial stromal sarcoma.

### TNM Descriptors

For identification of special cases of TNM or pTNM classifications, the "m" suffix and "y," "r," and "a" prefixes are used. Although they do not affect the stage grouping, they indicate cases needing separate analysis.
The "m" suffix indicates the presence of multiple primary tumors in a single site and is recorded in parentheses: pT(m)NM.

The "y" prefix indicates those cases in which classification is performed during or after initial multimodality therapy (ie, neoadjuvant chemotherapy, radiation therapy, or both chemotherapy and radiation therapy). The cTNM or pTNM category is identified by a "y" prefix. The ycTNM or ypTNM categorizes the extent of tumor actually present at the time of that examination. The "y" categorization is not an estimate of tumor before multimodality therapy (ie, before initiation of neoadjuvant therapy).

The "r" prefix indicates a recurrent tumor when staged after a documented disease-free interval and is identified by the "r" prefix: rTNM.

The "a" prefix designates the stage determined at autopsy: aTNM.

Additional Descriptors

Residual Tumor (R)
Tumor remaining in a patient after therapy with curative intent (eg, surgical resection for cure) is categorized by a system known as R classification, shown below.

RX Presence of residual tumor cannot be assessed
R0 No residual tumor
R1 Microscopic residual tumor
R2 Macroscopic residual tumor

For the surgeon, the R classification may be useful to indicate the known or assumed status of the completeness of a surgical excision. For the pathologist, the R classification is relevant to the status of the margins of a surgical resection specimen. That is, tumor involving the resection margin on pathologic examination may be assumed to correspond to residual tumor in the patient and may be classified as macroscopic or microscopic according to the findings at the specimen margin(s).

References


Bibliography


