



Protocol for the Examination of Specimens From Patients With Carcinoma and Carcinosarcoma of the Endometrium

Version: 5.0.0.0

Protocol Posting Date: December 2023

CAP Laboratory Accreditation Program Protocol Required Use Date: September 2024

The changes included in this current protocol version affect accreditation requirements. The new deadline for implementing this protocol version is reflected in the above accreditation date.

For accreditation purposes, this protocol should be used for the following procedures AND tumor types:

Procedure	Description
Hysterectomy	
Tumor Type	Description
Carcinoma	Includes carcinomas, carcinosarcomas (malignant mixed Müllerian tumor), and neuroendocrine carcinomas arising in the endometrium

This protocol is NOT required for accreditation purposes for the following:

Procedure
Endometrial biopsy
Endometrial curettage
Primary resection specimen with no residual cancer (e.g., following previous biopsy or curettage)
Cytologic specimens

The following tumor types should NOT be reported using this protocol

Tumor Type
Carcinomas arising in the uterine cervix (consider the Uterine Cervix protocol)
Uterine sarcoma, including adenosarcoma (consider the Uterine Sarcoma protocol)
Lymphoma (consider the Hodgkin or non-Hodgkin Lymphoma protocols)

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With guidance from the CAP Cancer and CAP Pathology Electronic Reporting Committees.

* Denotes primary author.

Accreditation Requirements

This protocol can be utilized for a variety of procedures and tumor types for clinical care purposes. For accreditation purposes, only the definitive primary cancer resection specimen is required to have the core and conditional data elements reported in a synoptic format.

- Core data elements are required in reports to adequately describe appropriate malignancies. For accreditation purposes, essential data elements must be reported in all instances, even if the response is “not applicable” or “cannot be determined.”
- Conditional data elements are only required to be reported if applicable as delineated in the protocol. For instance, the total number of lymph nodes examined must be reported, but only if nodes are present in the specimen.
- Optional data elements are identified with “+” and although not required for CAP accreditation purposes, may be considered for reporting as determined by local practice standards.

The use of this protocol is not required for recurrent tumors or for metastatic tumors that are resected at a different time than the primary tumor. Use of this protocol is also not required for pathology reviews performed at a second institution (i.e., secondary consultation, second opinion, or review of outside case at second institution).

Synoptic Reporting

All core and conditionally required data elements outlined on the surgical case summary from this cancer protocol must be displayed in synoptic report format. Synoptic format is defined as:

- Data element: followed by its answer (response), outline format without the paired Data element: Response format is NOT considered synoptic.
- The data element should be represented in the report as it is listed in the case summary. The response for any data element may be modified from those listed in the case summary, including “Cannot be determined” if appropriate.
- Each diagnostic parameter pair (Data element: Response) is listed on a separate line or in a tabular format to achieve visual separation. The following exceptions are allowed to be listed on one line:
 - Anatomic site or specimen, laterality, and procedure
 - Pathologic Stage Classification (pTNM) elements
 - Negative margins, as long as all negative margins are specifically enumerated where applicable
- The synoptic portion of the report can appear in the diagnosis section of the pathology report, at the end of the report or in a separate section, but all Data element: Responses must be listed together in one location

Organizations and pathologists may choose to list the required elements in any order, use additional methods in order to enhance or achieve visual separation, or add optional items within the synoptic report. The report may have required elements in a summary format elsewhere in the report IN ADDITION TO but not as replacement for the synoptic report ie, all required elements must be in the synoptic portion of the report in the format defined above.

Summary of Changes

v 5.0.0.0

- FIGO staging 2023 update
- Peritoneal/Ascitic Fluid and The International System for Reporting Serous Fluid Cytology updates
- Lymphatic and/or Vascular Invasion update to include LVSI FIGO terminology
- Explanatory note updates to D, G, H, I, K, L, M, and N

RETIRED

Reporting Template**Protocol Posting Date: December 2023****Select a single response unless otherwise indicated.****CASE SUMMARY: (ENDOMETRIUM)****Standard(s):** AJCC-UICC 8, FIGO Cancer Report 2023**CLINICAL****+Clinical History (Note [A](#)) (select all that apply)**

- Lynch syndrome
 Other (specify): _____

SPECIMEN (Note [B](#))**Procedure (select all that apply)***For information about lymph node sampling, please refer to the Regional Lymph Node section.*

- Total hysterectomy and bilateral salpingo-oophorectomy
 Radical hysterectomy
 Simple hysterectomy
 Supracervical hysterectomy
 Bilateral salpingo-oophorectomy
 Right salpingo-oophorectomy
 Left salpingo-oophorectomy
 Salpingo-oophorectomy, side not specified
 Right oophorectomy
 Left oophorectomy
 Oophorectomy, side not specified
 Bilateral salpingectomy
 Right salpingectomy
 Left salpingectomy
 Salpingectomy, side not specified
 Vaginal cuff resection
 Omentectomy
 Peritoneal biopsy(ies)
 Peritoneal washing
 Other (specify): _____

+Hysterectomy Type

- Abdominal
 Vaginal
 Vaginal, laparoscopic-assisted
 Laparoscopic
 Laparoscopic, robotic-assisted
 Other (specify): _____
 Not specified

+Specimen Integrity

- Intact
 Opened
 Morcellated
 Other (specify): _____

TUMOR**+Tumor Site (select all that apply)**

- Endometrium: _____
 Lower uterine segment: _____
 Endometrial polyp: _____
 Other (specify): _____
 Cannot be determined: _____

+Tumor Size

- Greatest dimension in Centimeters (cm): _____ cm
+Additional Dimension in Centimeters (cm): ____ x ____ cm
 Cannot be determined (explain): _____

Histologic Type (Note C)

- Endometrioid carcinoma, NOS
 POLE-ultramutated endometrioid carcinoma
 Mismatch repair-deficient endometrioid carcinoma
 p53-mutant endometrioid carcinoma
 No specific molecular profile (NSMP) endometrioid carcinoma
 Serous carcinoma
 Carcinosarcoma
 Mucinous carcinoma, intestinal type
 Clear cell adenocarcinoma, NOS
 Small cell neuroendocrine carcinoma
 Large cell neuroendocrine carcinoma
 Mixed neuroendocrine non-neuroendocrine carcinoma
 Mixed cell adenocarcinoma (specify types and percentages): _____
 Dedifferentiated carcinoma
 Undifferentiated carcinoma, NOS
 Mesonephric adenocarcinoma
 Squamous cell carcinoma, NOS
 Mesonephric-like adenocarcinoma
 Other histologic type not listed (specify): _____

+Histologic Type Comment: _____

Histologic Grade# (Note D)

International Federation of Gynecology and Obstetrics (FIGO) Grading System applies to endometrioid and mucinous carcinomas only. Serous, clear cell, transitional, small cell and large cell neuroendocrine carcinomas, undifferentiated / dedifferentiated carcinomas, and carcinosarcomas are generally considered to be high grade and it is not recommended to assign a histologic grade to these tumor types

- Not applicable
- FIGO grade 1
- FIGO grade 2
- FIGO grade 3
- Other (specify): _____
- Cannot be assessed (explain): _____

+Two-Tier Grading System (for endometrioid and mucinous carcinomas only) (Note [D](#))

- Low grade (encompassing FIGO 1 and 2)
- High grade (FIGO 3)
- Other (specify): _____
- Cannot be assessed: _____
- Not applicable

Myometrial Invasion (required only if applicable) (Note [E](#))

- Not applicable
- Not identified
- Present

Depth of Myometrial Invasion

- Specify in Millimeters (mm): _____ mm
- Other (specify): _____
- Cannot be determined (explain): _____

Myometrial Thickness

- Specify in Millimeters (mm): _____ mm
- Other (specify): _____
- Cannot be determined (explain): _____

Percentage of Myometrial Invasion

- Specify Percentage: _____ %
- Estimated to be less than 50%
- Estimated to be 50% or greater
- Cannot be determined (explain): _____
- Cannot be determined (explain): _____

+Adenomyosis

- Not identified
- Present, uninvolved by carcinoma
- Present, involved by carcinoma
- Cannot be determined: _____

Uterine Serosa Involvement

- Not identified
- Present
- Cannot be determined (explain): _____

Lower Uterine Segment Involvement (Note [F](#))

- Not identified
- Present, superficial (non-myoinvasive)

- Present, myoinvasive
- Present: _____
- Cannot be determined (explain): _____

Cervical Stroma Involvement (required only if applicable) (Note G)

- Not applicable
- Not identified
- Present

+Depth of Cervical Stroma Invasion

- Specify in Millimeters (mm): _____ mm
- Other (specify): _____
- Cannot be determined: _____

+Cervical Stroma Thickness

- Specify in Millimeters (mm): _____ mm
- Other (specify): _____
- Cannot be determined: _____
- Cannot be determined (explain): _____

Other Tissue / Organ Involvement# (Note G) (select all that apply)

Any organ not selected is either not involved or was not submitted.

- Not applicable
- Not identified
- Right ovary
- Left ovary
- Ovary (side not specified)
- Right fallopian tube
- Left fallopian tube
- Fallopian tube (side not specified)
- Vagina
- Right parametrium
- Left parametrium
- Parametrium (side not specified)
- Pelvic wall
- Bladder wall

Tumor must involve the mucosal surface of the bladder or intestine(s)

- Bladder mucosa##
- Rectal wall
- Bowel mucosa##
- Other organs / tissue (specify): _____
- Cannot be determined (explain): _____

+Peritoneal / Ascitic Fluid (Note H)

- Not submitted / unknown
- Negative for malignant cells
- Malignant cells present
- Results pending

+The International System for Reporting Serous Fluid Cytopathology Negative for malignancy (NFM)*# If malignancy cannot be excluded, cells are classified as "suspicious for malignancy".* Atypia of undetermined significance (AUS) (explain)#: _____ Suspicious for malignancy (SFM) (explain)#: _____ Malignant (MAL) Cannot be determined: _____ Results pending**Lymphatic and / or Vascular Invasion# (Note [I](#))***# Lymphatic and / or Vascular Invasion (LVI) is equivalent to the FIGO term Lymphatic and / or Vascular Space Invasion (LVSI).* Not identified Present+ Focal (less than 5 vessel involvement) (specify location, if possible): _____+ Extensive / substantial (greater than or equal to 5 vessel involvement) (specify location, if possible): _____ Equivocal (explain): _____ Cannot be determined: _____**+Tumor Comment:** _____**MARGINS (Note [J](#))****Margin Status***Margin section is required only if cervix and / or parametrium / paracervix is involved by carcinoma.* Not applicable All margins negative for invasive carcinoma**+Closest Margin(s) to Invasive Carcinoma (select all that apply)** Ectocervical / vaginal cuff (specify location, if possible): _____ Parametrial / paracervical (specify location, if possible): _____ Other (specify): _____ Cannot be determined: _____**+Distance from Invasive Carcinoma to Closest Margin***Specify in Millimeters (mm)* Exact distance: _____ mm Greater than: _____ mm At least: _____ mm Less than: _____ mm Less than 1 mm Other (specify): _____ Cannot be determined: _____ Invasive carcinoma present at margin**Margin(s) Involved by Invasive Carcinoma (select all that apply)** Ectocervical / vaginal cuff (specify location, if possible): _____ Parametrial / paracervical (specify location, if possible): _____ Other (specify): _____ Cannot be determined: _____ Other (specify): _____

___ Cannot be determined (explain): _____

+Margin Comment: _____

REGIONAL LYMPH NODES (Note [K](#))

Regional Lymph Node Status#

Lymph nodes designated as pelvic (parametrial, obturator, internal iliac (hypogastric), external iliac, common iliac, sacral, presacral) and para-aortic are considered regional lymph nodes. Any other involved nodes should be categorized as metastases (pM1) and reported in the distant metastasis section. Presence of isolated tumor cells no greater than 0.2 mm in regional lymph node(s) is considered N0 (i+).

___ Not applicable (no regional lymph nodes submitted or found)

___ Regional lymph nodes present

___ All regional lymph nodes negative for tumor cells

___ Tumor present in pelvic lymph node(s)

Macrometastases (greater than 2 mm), Micrometastases (greater than 0.2 mm to 2 mm), Isolated Tumor Cells (ITC: less than or equal to 0.2 mm or single cells or clusters of cells less than or equal to 200 cells in a single lymph node cross section). If pelvic and / or para-aortic lymph nodes are submitted and positive for tumor cells, reporting the number of nodes with or without macrometastases and micrometastases is required. Reporting isolated tumor cells is required only in the absence of macrometastasis or micrometastasis.

Pelvic Lymph Nodes (required only if present)

Total Number of Pelvic Nodes with Macrometastasis (greater than 2 mm) (sentinel and non-sentinel)

___ Exact number: _____

___ At least: _____

___ Other (specify): _____

___ Cannot be determined (explain): _____

+Number of Pelvic Sentinel Nodes with Macrometastasis

___ Exact number: _____

___ At least: _____

___ Other (specify): _____

___ Cannot be determined (explain): _____

Total Number of Pelvic Nodes with Micrometastasis (greater than 0.2 mm up to 2 mm and / or greater than 200 cells) (sentinel and non-sentinel)

___ Exact number: _____

___ At least: _____

___ Other (specify): _____

___ Cannot be determined (explain): _____

+Number of Pelvic Sentinel Nodes with Micrometastasis

___ Exact number: _____

___ At least: _____

___ Other (specify): _____

___ Cannot be determined (explain): _____

Total Number of Pelvic Nodes with Isolated Tumor Cells (0.2 mm or less and not more than 200 cells) (reported only if applicable)#

Reporting the number of lymph nodes with isolated tumor cells is required only in the absence of macrometastasis or micrometastasis in other lymph nodes.

___ Not applicable

___ Exact number: _____

At least: _____
 Other (specify): _____
 Cannot be determined (explain): _____

+Number of Pelvic Sentinel Nodes with ITCs

Exact number: _____
 At least: _____
 Other (specify): _____
 Cannot be determined (explain): _____

Laterality of Pelvic Node(s) with Tumor (reported only if applicable) (select all that apply)

Not applicable
 Right sentinel: _____
 Right non-sentinel: _____
 Left sentinel: _____
 Left non-sentinel: _____
 Cannot be determined: _____

+Size of Largest Pelvic Nodal Metastatic Deposit

Specify in Millimeters (mm)

Specify exact size: _____ mm
 Less than: _____ mm
 Greater than: _____ mm
 Other (specify): _____
 Cannot be determined (explain): _____

Tumor present in para-aortic lymph node(s)

Para-aortic Nodes (required only if present)

Total Number of Para-aortic Nodes with Macrometastasis (greater than 2 mm) (sentinel and non-sentinel) (reported only if applicable)

Not applicable
 Exact number: _____
 At least: _____
 Other (specify): _____
 Cannot be determined (explain): _____

+Number of Para-aortic Sentinel Nodes with Macrometastasis

Exact number: _____
 At least: _____
 Other (specify): _____
 Cannot be determined (explain): _____

Total Number of Para-aortic Nodes with Micrometastasis (greater than 0.2 mm up to 2 mm and / or greater than 200 cells) (sentinel and non-sentinel) (reported only if applicable)

Not applicable
 Exact number: _____
 At least: _____
 Cannot be determined (explain): _____

+Number of Para-aortic Sentinel Nodes with Micrometastasis

Exact number: _____
 At least: _____
 Other (specify): _____
 Cannot be determined (explain): _____

Total Number of Para-aortic Nodes with Isolated Tumor Cells# (0.2 mm or less and not more than 200 cells) (required only if applicable)

Reporting the number of lymph nodes with isolated tumor cells is required only in the absence of macrometastasis or micrometastasis in other lymph nodes.

- Not applicable
- Exact number: _____
- At least: _____
- Cannot be determined (explain): _____

+Number of Para-aortic Sentinel Nodes with ITCs

- Exact number: _____
- At least: _____
- Other (specify): _____
- Cannot be determined (explain): _____

Laterality of Para-aortic Node(s) with Tumor (required only if applicable) (select all that apply)

- Not applicable
- Right sentinel: _____
- Right non-sentinel: _____
- Left sentinel: _____
- Left non-sentinel: _____
- Cannot be determined: _____

+Size of Largest Para-aortic Nodal Metastatic Deposit

Specify in Millimeters (mm)

- Specify exact size: _____ mm
- Less than: _____ mm
- Greater than: _____ mm
- Other (specify): _____
- Cannot be determined (explain): _____
- Other (specify): _____
- Cannot be determined (explain): _____

Lymph Nodes Examined

Total Number of Pelvic Nodes Examined (sentinel and non-sentinel)

- Exact number: _____
- At least: _____
- Other (specify): _____
- Cannot be determined (explain): _____

Number of Pelvic Sentinel Nodes Examined (required only if applicable)

- Not applicable
- Exact number: _____
- At least: _____
- Other (specify): _____
- Cannot be determined (explain): _____

Total Number of Para-aortic Nodes Examined (sentinel and non-sentinel)

- Exact number: _____
- At least: _____
- Other (specify): _____
- Cannot be determined (explain): _____

Number of Para-aortic Sentinel Nodes Examined (required only if applicable)

- Not applicable
 Exact number: _____
 At least: _____
 Other (specify): _____
 Cannot be determined (explain): _____

+Regional Lymph Node Comment: _____

DISTANT METASTASIS**Distant Site(s) Involved, if applicable# (Note [L](#)) (select all that apply)**

This excludes metastasis to pelvic or para-aortic lymph nodes, vagina, uterine serosa, or adnexa

- Not applicable
 Inguinal lymph node(s): _____
 Omentum: _____
 Extrapelvic peritoneum: _____
 Lung: _____
 Liver: _____
 Bone: _____
 Other (specify): _____
 Cannot be determined: _____

pTNM CLASSIFICATION (AJCC 8th Edition) (Note [L](#))

Reporting of pT, pN, and (when applicable) pM categories is based on information available to the pathologist at the time the report is issued. As per the AJCC (Chapter 1, 8th Ed.), it is the managing physician's responsibility to establish the final pathologic stage based upon all pertinent information, including but potentially not limited to this pathology report.

Modified Classification (required only if applicable) (select all that apply)

- Not applicable
 y (post-neoadjuvant therapy)
 r (recurrence)

pT Category

- pT not assigned (cannot be determined based on available pathological information)
 pT0: No evidence of primary tumor
pT1: Tumor confined to the corpus uteri, including endocervical glandular involvement
 pT1a: Tumor limited to endometrium or invading less than half the myometrium
 pT1b: Tumor invading one half or more of the myometrium
 pT1 (subcategory cannot be determined)
 pT2: Tumor invading the stromal connective tissue of the cervix but not extending beyond the uterus.
 Does NOT include endocervical glandular involvement.
pT3: Tumor involving serosa, adnexa, vagina, or parametrium
 pT3a: Tumor involving serosa and / or adnexa (direct extension or metastasis)
 pT3b: Vaginal involvement (direct extension or metastasis) or parametrial involvement
 pT3 (subcategory cannot be determined)
Tumor must involve the mucosal surface
 pT4: Tumor invading bladder mucosa and / or bowel mucosa (bullous edema is not sufficient to

classify a tumor as T4)#

T Suffix (required only if applicable)

- Not applicable
 (m) multiple primary synchronous tumors in a single organ

pN Category

- pN not assigned (no nodes submitted or found)
 pN not assigned (cannot be determined based on available pathological information)
 pN0: No regional lymph node metastasis
 pN0(i+): Isolated tumor cells in regional lymph node(s) no greater than 0.2 mm
pN1: Regional lymph node metastasis to pelvic lymph nodes
Even one metastasis greater than 2.0 mm would qualify the classification as pN1a and pN2a.
 pN1mi: Regional lymph node metastasis (greater than 0.2 mm but not greater than 2.0 mm in diameter) to pelvic lymph nodes#
 pN1a: Regional lymph node metastasis (greater than 2.0 mm in diameter) to pelvic lymph nodes
 pN1 (subcategory cannot be determined)
pN2: Regional lymph node metastasis to para-aortic lymph nodes, with or without positive pelvic lymph nodes
 pN2mi: Regional lymph node metastasis (greater than 0.2 mm but not greater than 2.0 mm in diameter) to para-aortic lymph nodes, with or without positive pelvic lymph nodes#
 pN2a: Regional lymph node metastasis (greater than 2.0 mm in diameter) to para-aortic lymph nodes, with or without positive pelvic lymph nodes
 pN2 (subcategory cannot be determined)

N Suffix (required only if applicable)

Suffix (sn) is added to the N category when metastasis is identified only by sentinel lymph node biopsy. If after a sentinel node biopsy, the patient then undergoes a complete lymph node dissection, the (sn) suffix is not used.

- Not applicable
 (sn)

pM Category (required only if confirmed pathologically)

- Not applicable - pM cannot be determined from the submitted specimen(s)
 pM1: Distant metastasis (includes metastasis to inguinal lymph nodes, intraperitoneal disease, lung, liver, or bone). (It excludes metastasis to pelvic or para-aortic lymph nodes, vagina, uterine serosa, or adnexa)

FIGO STAGE

+FIGO Stage (2023 staging for cancer of the endometrium) (Note [M](#))

- I: Confined to the uterine corpus and ovary
 IA: Disease limited to the endometrium OR non-aggressive histological type, i.e., low-grade endometrioid, with invasion of less than half of the myometrium with no or focal lymphovascular space involvement (LVSI) OR good prognosis disease
 IA1: Non-aggressive histological type limited to an endometrial polyp OR confined to the endometrium
 IA2: Non-aggressive histological types involving less than half of the myometrium with no or focal LVSI
 IA3: Low-grade endometrioid carcinomas limited to the uterus and ovary

- + IAm (POLEmut): POLE mutated endometrial carcinoma, confined to the uterine corpus or with cervical extension, regardless of the degree of LVSI or histological type
- IB: Non-aggressive histological types with invasion of half or more of the myometrium, and with no or focal LVSI
- IC: Aggressive histological types limited to a polyp or confined to the endometrium
- II: Invasion of cervical stroma without extrauterine extension OR with substantial LVSI OR aggressive histological types with myometrial invasion
- IIA: Invasion of the cervical stroma of non-aggressive histological types
- IIB: Substantial LVSI of non-aggressive histological types
- IIC: Aggressive histological types with any myometrial involvement
- + IICm (p53abn): p53 abnormal endometrial carcinoma confined to the uterine corpus with any myometrial invasion, with or without cervical invasion, and regardless of the degree of LVSI or histological type
- III: Local and / or regional spread of the tumor of any histological subtype
- IIIA: Invasion of uterine serosa, adnexa, or both by direct extension or metastasis
- IIIA1: Spread to ovary or fallopian tube (except when meeting stage IA3 criteria)
- IIIA2: Involvement of uterine subserosa or spread through the uterine serosa
- IIIB: Metastasis or direct spread to the vagina and / or to the parametria or pelvic peritoneum
- IIIB1: Metastasis or direct spread to the vagina and / or the parametria
- IIIB2: Metastasis to the pelvic peritoneum
- IIIC: Metastasis to pelvic or para-aortic lymph nodes or both
- IIIC1: Metastasis to the pelvic lymph nodes
- IIIC1i: Micrometastasis (to pelvic nodes)
- IIIC1ii: Macrometastasis (to pelvic nodes)
- IIIC2: Metastasis to para-aortic lymph nodes up to the renal vessels, with or without metastasis to the pelvic lymph nodes
- IIIC2i: Micrometastasis (to para-aortic lymph nodes up to the renal vessels, with or without metastasis to the pelvic nodes)
- IIIC2ii: Macrometastasis (to para-aortic lymph nodes up to the renal vessels, with or without metastasis to the pelvic nodes)
- IV: Spread to the bladder mucosa and / or intestinal mucosa and / or distant metastasis
- IVA: Invasion of the bladder mucosa and / or intestine / bowel mucosa
- IVB: Abdominal peritoneal metastasis beyond the pelvis
- IVC: Distant metastasis, including metastasis to any extra- or intra-abdominal lymph nodes above the renal vessels, lungs, liver, brain or bone

+FIGO Modified Classification

- Not performed
- mPOLEmut (POLE mutation)
- mMMRd (mismatch repair deficiency)
- mNSMP (no specific molecular profile)
- mp53abn (p53 abnormal)

ADDITIONAL FINDINGS (Note [N](#))

+Additional Findings (select all that apply)

- None identified

___ Atypical hyperplasia / endometrial intraepithelial neoplasia (EIN)

___ Other (specify): _____

SPECIAL STUDIES

For reporting molecular testing, immunohistochemistry, and other cancer biomarker testing results, the CAP gynecologic origin biomarker template should be used. Pending biomarker studies should be listed in the Comments section of this report.

COMMENTS

Comment(s): _____

RETIRED

Explanatory Notes

A. Clinical History and Biomarker Testing

Colon carcinoma is the most common malignancy in hereditary nonpolyposis colon cancer [HNPCC; Lynch syndrome (LS)], which is caused by germline mutations in DNA mismatch repair genes (*MLH1*, *MLH2*, *MLH6*, and *PMS2*). However, endometrial carcinoma develops before colon carcinoma in >50% of women with HNPCC.^{1,2,3,4} 3% to 5% of endometrial carcinomas can be attributed to Lynch syndrome (LS). Patients with LS have a 40-60% lifetime risk for endometrial and colon cancer.⁵ Histopathologic features suggestive of HNPCC/LS-related carcinoma are well characterized in the colon, but not as well in the uterus. While lower uterine segment tumors and high-grade tumors in the endometrium seem to have a higher rate of being LS-associated tumors, tumor morphology and anatomic location of tumor cannot be used to select patients for screening for LS. Many LS-associated endometrial carcinomas are seen in probands that do not meet Bethesda or Amsterdam personal/family history criteria for Lynch Syndrome. However, when examining an endometrial carcinoma in a patient under 50 years of age or with a personal or family history of colon carcinoma, it is important to consider the possibility of an HNPCC/LS-related endometrial carcinoma.

According to the NCCN guidelines, there should be universal testing of endometrial carcinomas for mismatch repair (MMR) proteins/microsatellite instability (MSI). This can be tested on the hysterectomy specimen or the pre-surgical biopsy. Testing for defective DNA mismatch repair proteins by immunohistochemistry is the most cost-effective method (*MLH1*, *MSH2*, *MSH6*, and *PMS2* antibodies are commercially available).⁶ Loss of *MSH2* or *MSH6* expression essentially always indicates Lynch syndrome. HNPCC/LS-related endometrial carcinoma is predominantly associated with *MSH2* mutations and *MSH6* mutations.^{1,2,3,4} *PMS2* loss is often associated with loss of *MLH1* and is only independently meaningful if *MLH1* is intact. *MLH1* hypermethylation analysis should be completed on tumors that show loss of *MLH1* on IHC to help triage appropriate cases for germline testing. There should be genetic counseling and testing for all other MMR abnormalities. PCR assays can be used to detect high levels of microsatellite alterations (MSI), a condition that is definitional for defective DNA mismatch repair. This testing is performed on paraffin-embedded tissue and compares the results of tumor DNA to those of non-neoplastic tissues from the same patient.

In addition, Estrogen Receptor (ER) testing is recommended for stage III, IV, and recurrent disease and may be requested by the treating clinician in order to predict response to endocrine therapy. HER2 immunohistochemistry (with reflex test to HER2 FISH for equivocal IHC) should be considered for serous endometrial cancer. Please refer to the CAP endometrial cancer biomarker reporting template on www.cap.org/cancerprotocols for further details.

References

1. Aarnio M, Sankila R, Pukkala E, Salovaara R, Aaltonen LA, de la Chapelle A, Peltomäki P, Mecklin JP, Järvinen HJ. Cancer risk in mutation carriers of DNA-mismatch-repair genes. *Int J Cancer*. 1999;81:214-218.
2. Watson P, Vasen HF, Mecklin JP, Järvinen H, Lynch HT. The risk of endometrial cancer in hereditary nonpolyposis colorectal cancer. *Am J Med*. 1994;96:516-520.
3. Wijnen J, de Leeuw W, Vasen H, et al. Familial endometrial cancer in female carriers of *MSH6* germline mutations. *Nat Genet*. 1999;23:142-144.
4. Charames GS, Millar AL, Pal T, Narod S, Bapat B. Do *MSH6* mutations contribute to double primary cancers of the colorectum and endometrium? *Hum Genet*. 2000;107:623-629.

5. Cho KR, Cooper K, Croce S: International Society of Gynecological Pathologists (ISGyP): Guidelines from the special techniques and ancillary studies group. *Int J Gynecol Pathol*. 2019; 38(suppl 1):S114-122.
6. Mills AM, Liou S, Ford JM, Berek JS, Pai RK, Longacre TA. Lynch syndrome screening should be considered for all patients with newly diagnosed endometrial cancer. *Am J Surg Pathol*. 2014;38:1501-1509/

B. Specimen Type

In rare occasions when an endometrial carcinoma is not suspected, the pathologist may receive a supracervical hysterectomy specimen removed by laparoscopy. It has been reported that hysterectomies performed using certain laparoscopic techniques result in the finding of venous tumor emboli that are likely to be iatrogenic.¹ The FDA discourages morcellation for removal of uterus in women with suspected or known uterine cancer because there is risk of spreading tumor cells to the pelvis and peritoneal cavity. Therefore, if applicable, reporting of such a procedure is recommended (and listed under Specimen Integrity in the case summary).

References

1. Logani S, Herdman AV, Little JV, Moller KA. Vascular 'pseudo invasion' in laparoscopic hysterectomy specimens: a diagnostic pitfall. *Am J Surg Pathol*. 2008;32:560-565.

C. Histologic Type

Endometrial endometrioid carcinoma (EEC) displays varying proportions of glandular, papillary, and solid architecture, with the malignant cells showing endometrioid differentiation.¹ There are four molecular subtypes: POLE-ultramutated endometrioid carcinoma, mismatch repair-deficient endometrioid carcinoma, p53-mutant endometrioid carcinoma, and no specific molecular profile (NSMP) endometrioid carcinoma.² For a diagnosis of endometrioid carcinoma it is essential to have invasive endometrial carcinoma with endometrioid differentiation and desirable to have some degree of squamous, secretory, or mucinous differentiation. In high-grade tumors, squamous differentiation strongly favors endometrioid carcinoma over other histological types. Loss of immunoreactivity for ARID1A, PTEN, or one of the mismatch repair proteins favors high-grade EEC. Abnormal p53 expression is reported in 2–5% of low-grade and 20% of high-grade EECs.¹

For a diagnosis of serous carcinoma, it is essential to have a cytologic high-grade endometrial carcinoma with complex papillary and/or glandular architecture and desirable to have abnormal p53 and diffuse p16 immunohistochemistry. The vast majority of serous carcinoma tumors demonstrate *TP53* mutations.³ *ERBB2* (*HER2*) amplification is present in 30% of cases, frequently distributed heterogeneously.⁴ In the total cancer genome atlas (TCGA) cohort, all serous carcinomas were within the copy-number-high subgroup.²

To distinguish clear cell carcinoma from histological mimics it is important to adhere to architectural and cytological criteria. An admixture of tubulocystic, papillary, and/or solid patterns with clear to eosinophilic cuboidal, polygonal, hobnail, or flat cells is required. Confirmation by immunoreactivity, usually in the majority of cells, for stains such as HNF1 β , napsin A, and AMACR (P504S) is desirable.

Undifferentiated carcinoma of the endometrium is an epithelial malignancy with no overt cell lineage differentiation. Dedifferentiated carcinoma is composed of an undifferentiated carcinoma and a differentiated component. Almost 40% of monomorphic undifferentiated carcinomas contain a second

component of differentiated carcinoma, which is most frequently a FIGO grade 1 or 2 endometrioid carcinoma. In rare situations, undifferentiated carcinoma may be associated with a high-grade carcinoma (e.g., FIGO grade 3 endometrioid carcinoma and serous carcinoma).^{5,6} A discohesive cell morphology, lack, or focal PAX8 positivity by IHC, typically very focal staining for EMA and keratin (particularly CK8/18), and <10 % reactivity for neuroendocrine markers are features that support a diagnosis of undifferentiated carcinoma. Diffuse strong staining with pan-cytokeratin should not be present. Tumor cells express vimentin but not ER, PR, or E-cadherin. Half to two-thirds of dedifferentiated and half of undifferentiated carcinomas are mismatch repair–deficient/microsatellite unstable. About one-third of endometrial undifferentiated carcinomas show loss of SMARCA4 (BRG1) expression.

Carcinosarcoma is a biphasic tumor composed of high-grade carcinomatous and sarcomatous components. The carcinomatous component most often shows endometrioid or serous differentiation, but clear cell and undifferentiated carcinoma may be encountered. The mesenchymal component most commonly consists of high-grade sarcoma NOS, but heterologous elements (including rhabdomyosarcoma, chondrosarcoma, and rarely osteosarcoma) may be seen.¹

Mixed cell adenocarcinomas are endometrial carcinoma with two distinct histological types, in which at least one component is either serous or clear cell. Any amount of serous or clear cell carcinoma that can be confidently recognized on routine H&E sections in an endometrioid carcinoma qualifies for a mixed carcinoma. These are graded as high-grade carcinoma irrespective of the relative percentages of serous or clear cell carcinoma present. Dedifferentiated carcinoma and carcinosarcoma are not mixed carcinomas. Immunohistochemical demonstration of the two distinct carcinoma types is desirable.¹

Other types: Mesonephric adenocarcinoma is an adenocarcinoma originating from mesonephric remnants. Mesonephric-like adenocarcinoma, a newly described entity, is an adenocarcinoma resembling mesonephric differentiation and limited data suggest an aggressive behavior.^{7,8} Primary squamous carcinoma is a carcinoma with exclusive squamous differentiation. Primary gastric (gastrointestinal)–type mucinous carcinoma is a carcinoma with mucinous gastric/gastrointestinal features. The diagnosis of these rare carcinomas is based on morphology and it is important to exclude an endometrioid component, a cervical origin, (and/or metastasis from the gastrointestinal tract in cases of mucinous carcinoma), before rendering these as the histologic type. Small cell and large cell neuroendocrine carcinomas (NEC) account for less than 1% of all gynecological malignancies. Endometrial NEC are typically seen in postmenopausal women and the etiology is unknown. Abnormal mismatch repair protein expression has been described in endometrial SCNECs. In mixed NEC and non-NEC the percentages of individual tumor types should be given.

Stromal invasion, which is defined by loss of intervening stroma (a confluent glandular, cribriform, or labyrinthine pattern), altered fibroblastic stroma (desmoplastic stromal reaction), a complex (mostly villoglandular) or a non-squamous solid architecture distinguishes well differentiated EEC from endometrial atypical hyperplasia / endometrioid intraepithelial neoplasia. It may be difficult to distinguish EEC with mucinous differentiation from atypical mucinous glandular proliferations; cribriform or confluent architecture and cytological atypia are distinguishing features.⁹

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D. Histologic Grading

The International Federation of Gynecology and Obstetrics (FIGO) grading system for carcinomas of the uterine corpus is only officially designated for endometrioid and non-intestinal type mucinous carcinomas and is based on architectural features as follows:^{1,2}

Grade 1	5% or less non-squamous solid growth pattern
Grade 2	6% to 50% non-squamous solid growth pattern
Grade 3	>50% non-squamous solid growth pattern

Severe cytologic atypia in the majority of cells (>50%), which exceeds that which is routinely expected for the architectural grade, increases the tumor grade by 1.^{2,3} Generally, most tumors can be graded on architecture alone; cytologic atypia should be pronounced from a low power. In addition, the following guidelines should be used in grading:

1. The squamous component of endometrioid adenocarcinoma should not be graded because the degree of differentiation typically parallels that of the glandular component.²
2. Because non-intestinal type mucinous carcinomas are closely related to endometrioid carcinomas, they can be graded by the same criteria. However, FIGO grading should NOT be used when endometrioid or mucinous differentiation is in doubt or cannot be established.³
3. Serous, clear cell, transitional, small cell, and large cell neuroendocrine carcinomas, undifferentiated/dedifferentiated carcinomas, and carcinosarcomas are considered to be high grade and a FIGO grade should not be assigned to these tumor types.^{2,3} When the case summary is being completed, these should be designated as “not applicable” for histologic grade.

4. In mixed carcinomas, the highest grade should be assigned.

The FIGO 2023 revisions adopted the WHO binary system (low versus high grade) for grading endometrioid and mucinous (non-intestinal type) carcinomas in recognition that FIGO grade 1 and 2 tumors behave similarly.⁴ Moreover, histological grading is of no consequence for women with MMR-deficient carcinoma. However, women who desire fertility-conserving therapy may benefit from the use of the FIGO 3-tiered system, which is also endorsed by FIGO. Either or both systems may be employed. Molecular profiling with subclassification is highly encouraged for high-grade endometrioid carcinomas to establish risk for recurrence and prognosis.⁴

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E. Myometrial Invasion

Assessing myometrial invasion may be difficult. Depth of invasion should be measured from the endomyometrial junction to the deepest point of invasion, which may not be easy because the endomyometrial junction in normal conditions is often irregular. In these cases, it is always helpful to look for compressed, non-neoplastic endometrial glands at the nearby endomyometrial junction or even at the base of the tumor. Carcinoma involving adenomyosis foci should not be interpreted as invasive carcinoma. However, the distinction between invasive carcinoma and carcinoma involving adenomyosis may be difficult, because in some cases invasive carcinoma may not elicit stromal response. In the absence of adenomyosis uninvolved by tumor in other sections of the specimen, a diagnosis of adenomyosis involved by adenocarcinoma should be made with caution. CD10 staining is not helpful in this differential diagnosis because stromal cells surrounding foci of invasive carcinoma are also frequently CD10 positive. There are no rules for determining how to measure the depth of invasion in the rare cases where myoinvasive carcinoma is only encountered in foci of adenomyosis involved by carcinoma. In such cases, it is advised that the distance from the adenomyotic focus to the deepest area of invasion be measured (Figure 1).¹ Therefore, if there is a tumor with a 2-mm focus of myoinvasion from a focus of adenomyosis in the deep myometrium, it is still considered as having <50% myometrial invasion (FIGO stage IA). In EEC with a MELF (microcystic, elongated and fragmented) pattern of invasion, desmoplasia alone should not be a criteria to measure the depth of invasion. Depth of invasion should be measured as the deepest extent with malignant cells present. LVI should not be used in measuring depth of myometrial invasion; only carcinoma infiltrating the myometrium is to be measured.²

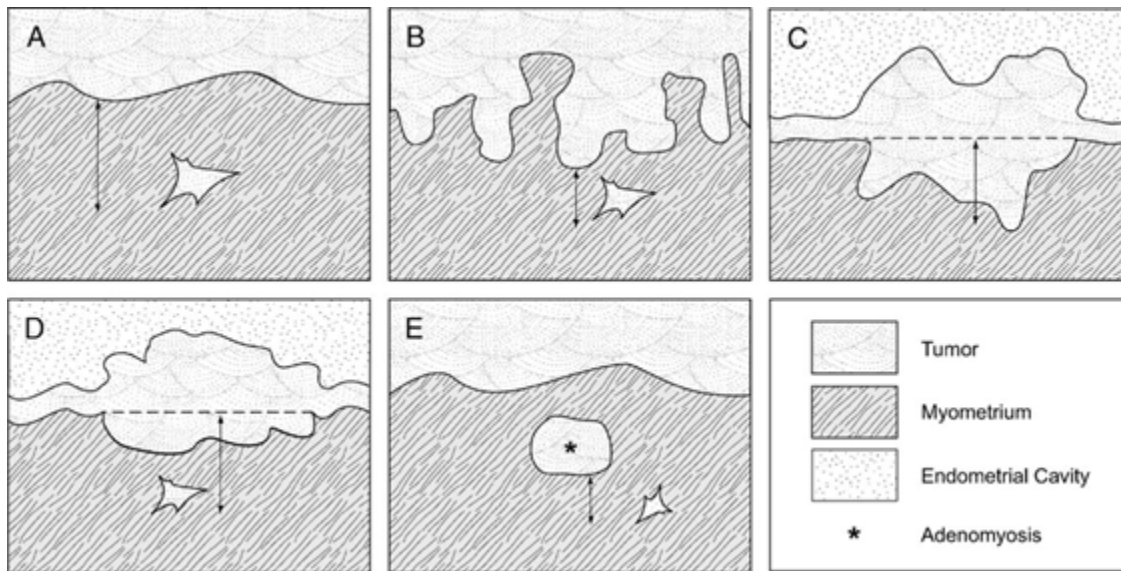


Figure 1. Schematic of measurement of depth of invasion in (A) tumor with a regular interface; (B) tumor with an irregular endomyometrial interface; (C) and (D) tumor with an exophytic growth; (E) tumor arising from adenomyosis. From Ali A, Black D, Soslow RA. Difficulties in assessing the depth of myometrial invasion in endometrial carcinoma. *Int J Gynecol Pathol.* 2007;26:115-123. Copyright © 2007, Wolters Kluwer Health. Reproduced with permission.

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F. Lower Uterine Segment Involvement

The prevalence of Lynch syndrome in patients with LUS endometrial carcinoma (29%) has been reported to be much greater than that of the general endometrial cancer patient population (1.8%) or in endometrial cancer patients younger than age 50 years (8% to 9%).¹

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G. Cervical, Adnexal, and Other Organ Involvement

The American Joint Committee on Cancer (AJCC)/FIGO staging system considers stage II disease only when there is invasion of the cervical stroma involvement is seen. Invasion can be identified by the appearance of tumor within stroma containing benign endocervical glands, or in regions deep to endocervical crypts. Involvement of the surface endocervical epithelium and/or endocervical glands (by either direct extension or drop metastases) does not have any prognostic significance and is not T2/Stage II.

There remain controversies regarding primary endometrial and ovarian tumors versus synchronous primary tumors, but clonal studies have uncovered that for the majority of low-grade endometrioid

carcinomas, the tumor arises in the endometrium and spreads to the ovaries.^{1,2} Most high-grade endometrial tumors are metastatic to ovaries rather than synchronous. However, due to favorable outcomes for these low-grade tumors, the World Health Organization (WHO)³, European Society of Gynecologic Oncology (ESGO), European Society for Therapeutic Radiology and Oncology (ESTRO), and European Society of Pathology (ESP)⁴ recommend treatment of concomitant endometrial and ovarian low-grade endometrioid carcinoma as if they were synchronous primary tumors. The FIGO category of stage IA3 was designed for this contingency, provided these cases do not show $\geq 50\%$ myometrial invasion, substantial LVSI, or additional metastases, and the tumor involves only one ovary without ovarian capsular invasion or extension. If the case does not fulfill these criteria, the ovarian tumor should be reported as extension of tumor to the ovary (AJCC pT3a or FIGO Stage IIIA1).

Tumor invading the fallopian tube also constitutes pT3a or FIGO Stage IIIA1, but the presence of only intraluminal, unattached tumor should not be considered tubal involvement. The presence of intramucosal carcinoma must be distinguished from serous tubal intraepithelial carcinoma (STIC) through immunohistochemical markers or molecular studies whenever possible. The staging and significance of intramucosal involvement by non-STIC tumor alone remains controversial.⁵

Uterine serosa is involved when the tumor reaches submesothelial connective tissue or the mesothelial layer, whether or not tumor cells appear on the serosal surface.⁵ To qualify for Stage IV disease, involvement of the bladder or bowel must show invasion of the mucosal layer. Peritoneal involvement beyond the pelvic brim also constitutes Stage IV disease (FIGO IVB).

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H. Peritoneal Washings or Ascites Fluid

The prognostic significance of presence of tumor cells in peritoneal washings or ascites fluid in uterine cancer is controversial. There are studies that indicate either a worse prognosis or no alteration of prognosis on the basis of positive cytology. Consequently, staging systems no longer utilize positive cytology to alter stage. Use of the terminology of the International System (TIS)¹ for reporting serous cytopathology is optional, but is supported by the American Society of Cytopathology and the International Academy of Cytology for consistency with cytology specimens. When the cytology report of collected fluid is available for correlation with the specimen, reporting the presence or absence of tumor cells in peritoneal fluid is strongly encouraged.

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I. Lymphatic and/or Vascular Invasion

The presence or absence of LVI/LVSI (lymphatic/vascular space invasion) should be recorded in the pathology report. Before diagnosing LVSI, mimics should be excluded, such as retraction, MELF pattern of invasion, and artifactual displacement of tumor cells. MELF (microcystic, elongated, fragmented) is a pattern of myometrial infiltration by low-grade endometrial carcinoma whereby the infiltrating cells are attenuated, elongated, dilated, and/or fragmented and may mimic LVSI.¹ Immunohistochemistry is of limited use in the identification of LVSI. The presence of actual tumor emboli within the vessels is required for a diagnosis of LVSI. Studies have shown extensive LVI to be a strong independent prognostic factor for pelvic regional recurrence, distant recurrence, and overall survival. When present, the extent of LVSI may be semi-quantified as low focal (less than 5-vessel involvement) or as extensive (greater than or equal to 5-vessel involvement).^{1,2} Practice guidelines differ regarding the definition of significant lymphatic and/or vascular invasion. In 2021, the WHO defined extensive/substantial LVSI as 5 or more involved vessels.² The same number for “substantial” involvement is adopted by FIGO and the 2021 ESGO/ESTRO/ESP guidelines.^{3,4} For purposes of clarity, this protocol uses the WHO and FIGO terminology of “extensive/substantial” for 5 or more vessel involvement. Previous versions defined 3 or more vessel involvement as “extensive”. The vessel count is based on the single slide with the largest number of involved vessels and not the sum of vessel involvement on all slides. The location of LVSI (e.g., deep myometrial, cervical, adnexal, parametrial, etc.) may allow future studies to assess their significance.⁴ Studies have shown extensive LVSI to be a strong independent prognostic factor for pelvic regional recurrence, distant recurrence, and overall survival.^{5,6,7,8,9,10}

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J. Margins

The parametrial/paracervical soft tissue and the vaginal cuff are the only true margins in total hysterectomy specimens. These margins should be reported if the cervix and/or parametrium/paracervix is involved by carcinoma. If not, reporting the status of the vaginal and parametrial margins in a hysterectomy specimen is optional.

K. Lymph Node Status

Lymph nodes that are considered “regional” are the **pelvic** (parametrial, obturator, internal iliac/hypogastric, external iliac, common iliac, sacral, presacral) and **para-aortic** nodes. Any other involved nodes should be categorized as metastases (pM1) and reported in the distant metastasis section.

FIGO staging also incorporates the concept of micrometastasis for lymph nodes, distinguishing micrometastasis (i) from macrometastasis (ii) in Stage IIIC disease. The definition of micro- and macrometastasis is the same for AJCC and FIGO. The presence of isolated tumor cells (ITC) no greater than 0.2 mm in regional lymph node(s) is considered N0 (i+). Isolated tumor cells do not upstage a patient and need only be reported in the absence of micro- or macrometastases. Micrometastases (pN1(mi)) are deposits greater than 0.2 mm but not greater than 2 mm, and macrometastases are greater than 2 mm. Those patients at intermediate or high risk for recurrence benefit from lymph node staging. Sentinel lymph node sampling is widely used for staging low or intermediate-risk patients, but is also an alternative to systematic lymphadenectomy in presumed early-stage cancers for higher-risk patients.¹ Sentinel nodes should be appropriately processed for maximal tumor detection, to include gross sectioning of the node in 2 mm segments, embedding the entire node, and performing serial sections (3 sections at 200-250 µm) to detect micrometastases.²

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L. pTNM Classification

The TNM staging system for endometrial cancer endorsed by the AJCC and the UICC¹, and the parallel system formulated by FIGO² are recommended.

According to AJCC/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.

The referring physician usually carries out clinical classification (cTNM) 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 (e.g., 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.

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 “y” prefix indicates those cases in which classification is performed during or after initial multimodality therapy (i.e., 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 (i.e., 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.

T Category Considerations

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

N Category Considerations

Isolated tumor cells (ITCs) are single cells or small clusters of cells not more than 0.2 mm in greatest dimension. Lymph nodes or distant sites with ITCs found by either histologic examination (e.g., immunohistochemical evaluation for cytokeratin) or non-morphological techniques (e.g., flow cytometry, DNA analysis, polymerase chain reaction [PCR] amplification of a specific tumor marker) should be so identified. There is currently no guidance in the literature as to how these patients should be coded; until more data are available, they should be coded as “N0(i+)” with a comment noting how the cells were identified.

Sentinel nodes should be sliced at 2.0 mm intervals. The sentinel nodes should undergo ultrastaging; currently, there is no universal ultrastaging protocol. However, all institutions undertaking sentinel lymph node examination should have a standard procedure in place for sentinel lymph nodes. Protocols used at the 2 largest cancer centers in the United States are as follows:

- 1) Memorial Sloan Kettering Cancer Center Protocol³: If the initial H&E-stained slide is negative for carcinoma and the endometrial cancer is myo-invasive or associated with vascular/lymphatic

invasion, 2 additional levels at 50 µm apart are examined, at each level 2 slides are obtained, one for H&E and the second for keratin cocktail IHC if the H&E-stained slide is negative.

- 2) The University of Texas M.D. Anderson Cancer Center Protocol⁴: If the H&E-stained slide is negative for tumor, 3 consecutive sections at 250 µm into the paraffin block are obtained (one for H&E and one of the remaining 2 is to be used for keratin cocktail IHC if the additional H&E-stained slide is negative).

There is little data to assign risk for nonsentinel lymph node metastasis based on the size of the metastasis in the sentinel lymph node. However, the size criteria for micrometastasis and macrometastasis is adopted from the experience in breast carcinoma. Micrometastasis is defined as a metastasis measuring greater than 0.2 mm but less than 2 mm.

M Category Considerations

Metastases to the intrapelvic peritoneum, that do not extend beyond the pelvic brim, are considered T3 and not M1 disease. Metastases confined to the pelvis are excluded from M1 categorization. Distance metastases are required to be beyond the pelvic brim. In complex cases, it may be necessary to confer with the surgeon to determine the appropriate stage.

Primary Tumor (T)

FIGO

T Category	Stage	Definition
T1	I	Tumor confined to corpus uteri
T1a	IA	Non-aggressive tumor type limited# to a polyp, endometrium or invades less than one-half of the myometrium, with no or focal LVSI
T1b	IB	Non-aggressive tumor type invades one-half or more of the myometrium, with no or focal LVSI
	IC	Aggressive tumor types limited to a polyp or the endometrium
T2	II	Tumor invades stromal connective tissue of the cervix but does not extend beyond the uterus, or with extensive/substantial LVSI or aggressive histologic types with myometrial invasion
T3	III	Tumor of any histologic type involving serosa, adnexa, vagina, or parametrium, i.e., local and/or regional spread as specified in T3a and T3b, and in FIGO IIIA, IIIB and IIIC
T3a	IIIA	Tumor involving the uterine serosa and/or adnexa (direct extension or metastasis)
T3b	IIIB	Vaginal involvement (direct extension or metastasis), parametrial or pelvic peritoneal involvement
	IIIC	Metastasis to the pelvic or para-aortic lymph nodes or both
T4##	IV	Tumor invading bladder mucosa### and/or bowel mucosa### and/or distant metastasis
	IVA	Invasion of bladder mucosa and/or bowel mucosa
	IVB	Abdominal peritoneal metastasis beyond the pelvis
	IVC	Distant metastasis, including metastasis to any extra- or intra-abdominal lymph nodes above the renal vessels, lungs, liver, brain, or bone

Low-grade endometrioid carcinoma may involve the ovary (FIGO IA3)

Tumor must involve the mucosal surface; presence of bullous edema is not sufficient evidence to classify a tumor as T4.

Regional Lymph Nodes (N):# TNM Staging System

FIGO

N Category	Stage	Definition
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NX		Regional lymph nodes# cannot be assessed
NO		No regional lymph node metastasis
N0(i+)		Isolated tumor cells in regional lymph node(s) no greater than 0.2 mm
N1	IIIC1	Regional lymph node# metastasis to pelvic lymph nodes
N1mi	IIIC1i	Regional lymph node metastasis (greater than 0.2 mm but not greater than 2 mm in diameter) to pelvic lymph nodes
N1a	IIIC1ii	Regional lymph node metastasis (greater than 2 mm in diameter) to pelvic lymph nodes
N2	IIIC2	Regional lymph node# metastasis to para-aortic lymph nodes with or without positive pelvic lymph nodes
N2mi	IIIC2i	Regional lymph node metastasis (greater than 0.2 mm but not greater than 2 mm in diameter) to para-aortic lymph nodes, with or without positive pelvic lymph nodes
N2a	IIIC2ii	Regional lymph node metastasis (greater than 2 mm in diameter) to para-aortic lymph nodes, with or without positive pelvic lymph nodes

Regional lymph nodes include the pelvic, obturator, internal iliac (hypogastric), external iliac, common iliac, para-aortic, presacral, and parametrial lymph nodes. Even one metastasis >2.0 mm would qualify as pN1a or pN2a.

Distant Metastasis (M): TNM Staging System

FIGO

M Category	Stage	Definition
M0	IVA	No distant metastasis
M1	IVB	Abdominal peritoneal metastasis beyond the pelvis
	IVC	Distant metastasis (includes metastasis to abdominal lymph nodes [other IVC than para-aortic], and/or inguinal lymph nodes, intraperitoneal disease, lung, liver, or bone; excludes metastasis to vagina, pelvic serosa, or adnexa)

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M. FIGO Staging

FIGO staging currently emphasizes the recognition of non-aggressive vs aggressive histological tumor types and molecular subclassification for endometrial tumors. Non-aggressive tumor types include FIGO grades 1 and 2 endometrioid carcinoma (EEC) and mucinous carcinoma (other than gastro-intestinal type). These low-grade tumors have a good prognosis and generally do not require neoadjuvant therapy. All other histologic types are aggressive: grade 3 endometrioid carcinoma, serous carcinoma, clear cell carcinoma, mesonephric-like carcinoma, gastro-intestinal-type mucinous carcinoma, mixed carcinoma, carcinosarcoma, and undifferentiated carcinoma.

The addition of molecular classification to all FIGO stages is optional but encouraged, especially for high-grade endometrioid carcinoma. This predictive information is added as a subscript to the “m” (molecular) modifier of the FIGO stage. For example, Stage IAmpOLEmut is a POLE-mutated carcinoma confined to the uterus or with cervical extension, regardless of the degree of LVSI or histologic type. A metastatic serous carcinoma would be Stage IVmp53mut. POLE-mutated and p53-abnormal findings alter the FIGO stage of early disease; the others do not. Complete molecular classification of all endometrial tumors is encouraged and can be performed on biopsy specimens, but not all laboratories have access to all tests. A simple approach is to use 3 immunohistochemical (IHC) markers (p53, MSH6, PMS2) and one molecular marker (POLE mutational analysis) to group tumors into 4 categories: POLEmut, MMRd, NSMP, and p53abn.¹ A small number of cases may show results in more than one category (“multiple classifiers”). Those with POLEmut or MMRd along with p53abn should not be classified as p53abn. Investigations are still in progress but most of these cases retain the better prognosis and should be assigned as such. If the tumor is both POLE-mutated and MMR-deficient, consider genetic testing for Lynch syndrome.¹

Molecular Classification	Prognostic Category	Preferred Testing
mPOLE-mut	Favorable	POLE mutational analysis
mMMRd	Intermediate	Mismatch repair (MMR) protein analysis by IHC
mNSMP	Intermediate	All tests negative
mp53abn	Poor	p53 by IHC

Previously, there has been confusion regarding the definition of distal metastasis. FIGO clarifies the anatomic limit of the pelvis as the line between the anterior superior iliac spines.¹ In some cases, the pathologist may need to confer with the surgeon to determine the exact site of a peritoneal biopsy when considering metastatic disease.

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N. Additional Findings

Atypical Hyperplasia/Endometrioid Intraepithelial Neoplasia^{1,2,3}

It is essential to see a crowded architecture of cytologically altered glands that are distinct from both the background architecture and cytology of adjacent or entrapped normal glands from low power. The volume of crowded glands exceeds that of the stroma. In addition, there is nuclear atypia in the form of nuclear enlargement, pleomorphism, rounding, loss of polarity, and nucleoli.¹ A size of at least 1.0 mm is recommended. Loss of immunoreactivity for PTEN, PAX2, or mismatch repair proteins may be a helpful diagnostic tool.³ Common mimics such as metaplasia, basaloid, polyp, or dys-synchronous-phase endometrium must be excluded.

Proposed criteria distinguishing Well-Differentiated Endometrioid Adenocarcinoma from EIN/Atypical Endometrial Hyperplasia

- (1) Irregular infiltration of myometrium associated with an altered fibroblastic stroma (desmoplastic response), **or**
- (2) Confluent glandular pattern (cribriform growth, or complex folded mazelike epithelium), **or**

(3) Solid non-squamous epithelial growth

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