

Understanding of new FIGO staging of endometrium

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Point 1

"Histopathological findings are central features of 2023 FIGO staging of endometrial carcinoma"

Risk stratification and management based on

- Routine H&E examination
- Molecular classification

Clinician's side

Pathologist's side





Compromised tissue preservation can affect pathological evaluation

Retraction artifact may affect LVSI evaluation and counting

If surgical specimen needs to be used for molecular classification (endometrial biopsy/curettage not available)

Antigen degradation may affect IHC result Example: p53 pattern: Wild-type VS Abnomal

DNA degradation may affect mutation/molecular studies

Clinicians' contribution to specimen handling is important for pathological evaluation



Point 2

The use of molecular classification

- Costs of testing may be a limitation for the use molecular classification in limited-resource settings
- Testing for POLE pathogenic mutation costs higher than IHC (MMR, p53)
- Testing for 5 hotspot mutations (2020 WHO) may detect over >90% of POLEmut endometrial carcinoma (PMID: 37229628)
 - Sanger sequencing for hotspots: much lower cost than NGS
- May this make molecular classification useful for stage I-II endometrial cancer in limited-resource settings?

Stage I-II 2021 ESGO/ESTRO/ESP guidelines



Point 3

Adnexal involvement in endometrial cancer

- Synchronous endometrial and ovarian low-grade endometrioid CAs with 'stage IA3' conditions have favorable prognosis (although clonally related or likely representing ovarian metastasis)
 - Limited ability for widespread metastasis
- Intraluminal tumor fragment is not considered for staging (not stage IIIA1)
- "Tubal intramucosal spread has controversial prognostic significance"
- For patients with low-grade endometrioid CA, myometrial invasion <inner half, without substantial LVSI
 - Tubal mucosal involvement, without muscular wall invasion
 - Some had previous tubal sterilization
- Is there any choice for conservative management?

