

# 子宮內膜癌診療指引

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參考資料：

Uterine Neoplasms NCCN Guidelines V1. 2018

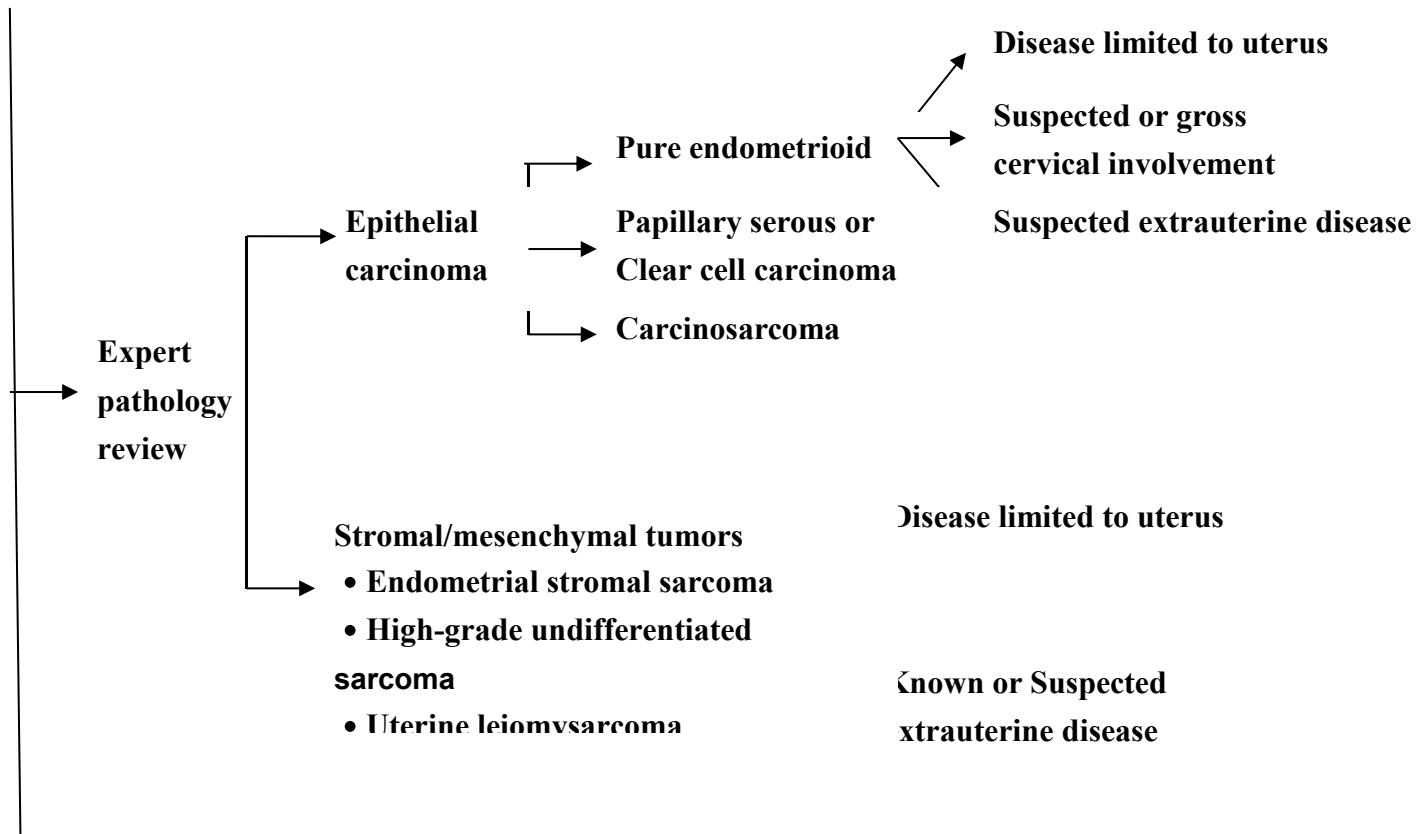
2011 年國家衛生研究院-婦癌臨床診療指引

全民健康保險藥品給付規定一百零五年版(30051\_2)

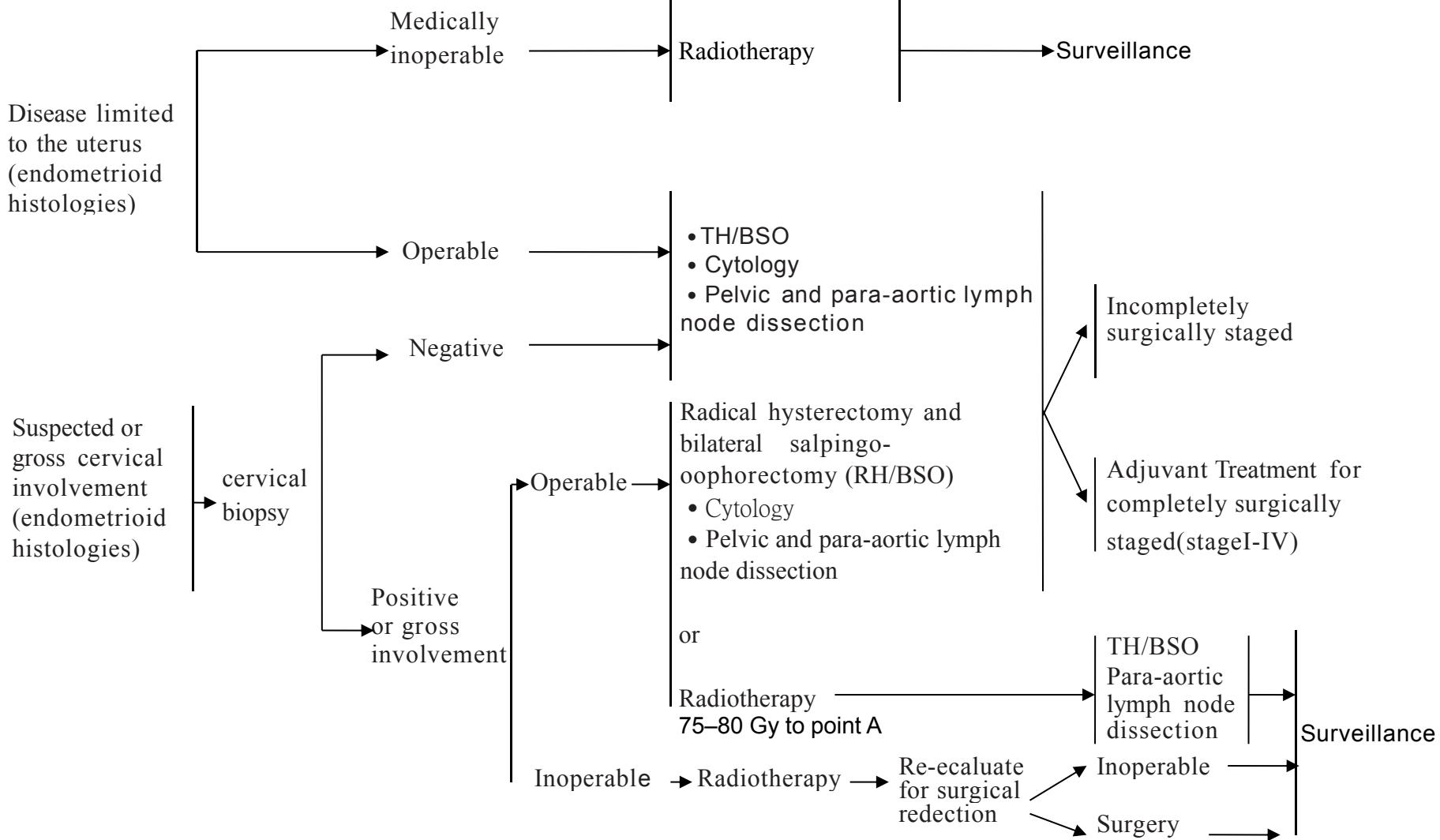
## WORK UP

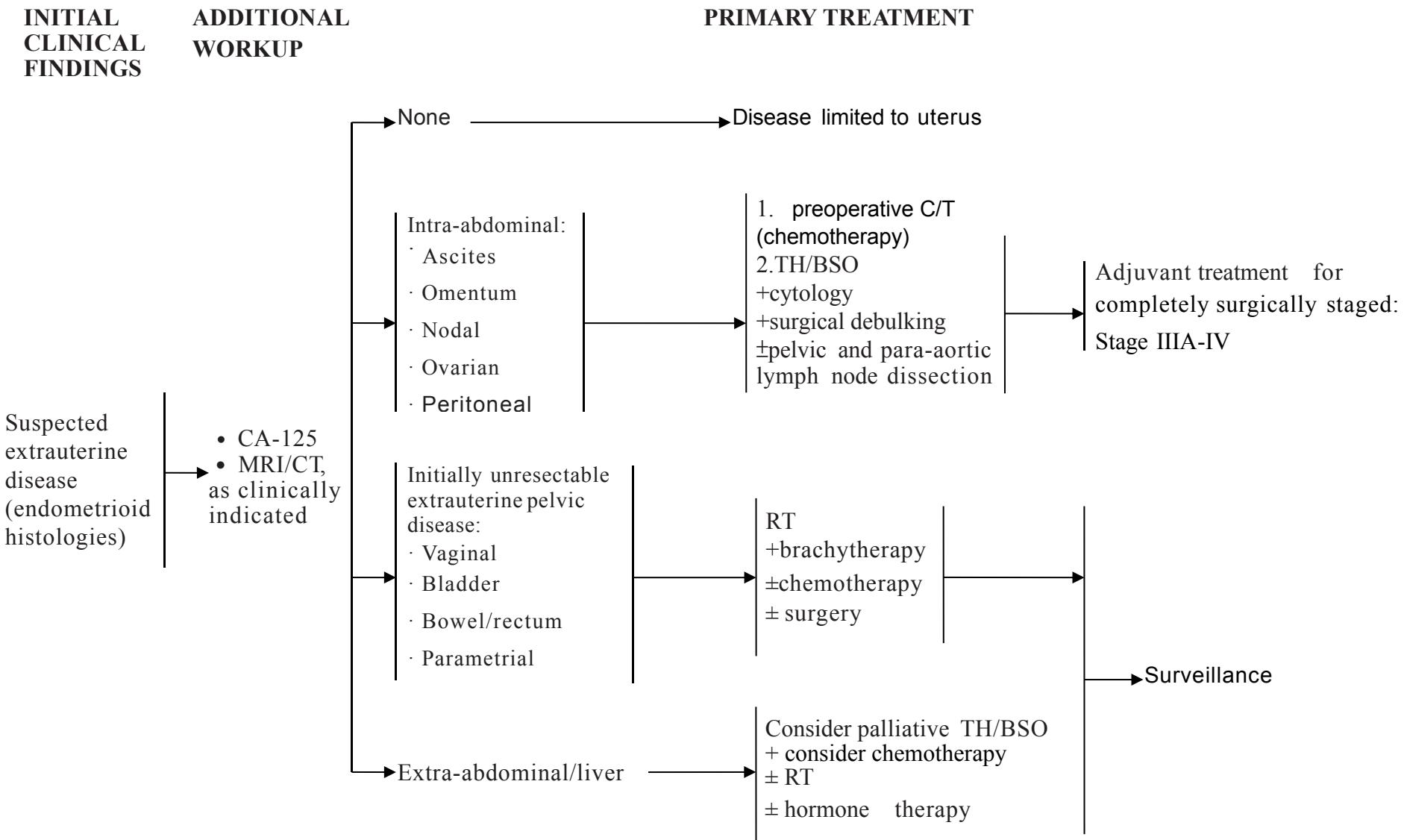
- History
- Physical exam
- CBC & Platelet
- Endometrial biopsy
- Chest imaging
- Liver function
- tests/Renal function tests
- Imaging :
   
Abd CT or MRI
- genetic counseling

## INITIAL CLINICAL FINDING



**INITIAL CLINICAL FINDINGS**





CLINICAL FINDINGS (completely surgically staged)	ADVERSE RISK FACTORS	HISTOLOGIC GRADE/ADJUVANT TREATMENT		
		G1	G2	G3
Stage IA (< 50%) myometrial invasion	Adverse risk factors not present	Observe	Observe or Vaginal brachytherapy	Observe or Vaginal brachytherapy
	Adverse risk factors present	Observe or Vaginal brachytherapy	Observe or Vaginal brachytherapy and/or pelvic RT(category 2B)	Observe or Vaginal brachytherapy and/or pelvic RT
Stage IB (≥ 50%) myometrial invasion	Adverse risk factors not present	Observe or Vaginal brachytherapy	Observe or Vaginal brachytherapy	pelvic RT and/or Vaginal brachytherapy or Observe
	Adverse risk factors present	Observe or Vaginal brachytherapy and/or Pelvic RT	Observe or Vaginal brachytherapy and/or Pelvic RT	Pelvic RT and/or Vaginal brachytherapy±chemotherapy (category 2B for chemotherapy)
Stage II		pelvic RT and Vaginal brachytherapy	pelvic RT +Vaginal brachytherapy	pelvic RT +Vaginal brachytherapy ±chemotherapy(category 2B)
Stage IIIA		RT or CT pelvic RT ±Vaginal brachytherapy	chemotherapy±RT or RT ±chemotherapy or pelvic RT ±Vaginal brachytherapy	chemotherapy±RT or RT ±chemotherapy or pelvic RT ±Vaginal brachytherapy
Adverse Risk Factors: Age, Lymphovascular invasion, tumor size				

**CLINICAL FINDINGS**  
(completely surgically staged)

**ADJUVANT TREATMENT**

Stage IIIB → Chemotherapy or RT or CCRT

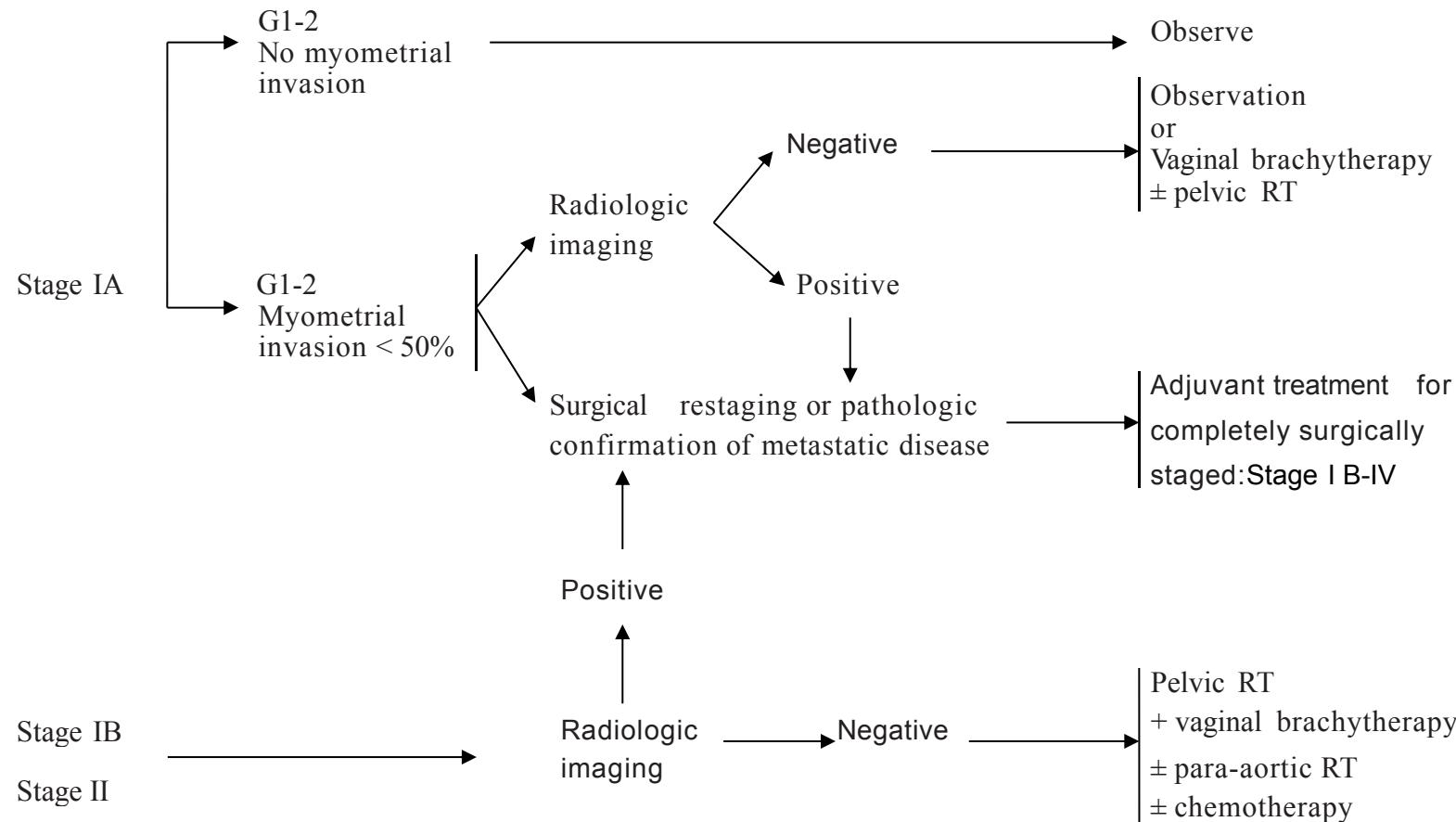
Stage IIC1 → Pelvic node positive → Chemotherapy and/or RT

Stage IIC2 → Para-aortic node positive  
± pelvic node positive → Chemotherapy and/or RT

Stage IV<sub>A</sub>, IVB → Debulked <sup>1</sup> and  
with no gross  
residual disease  
or microscopic  
abdominal disease → Chemotherapy ± RT

**CLINICAL FINDINGS**  
(Incompletely surgically staged)

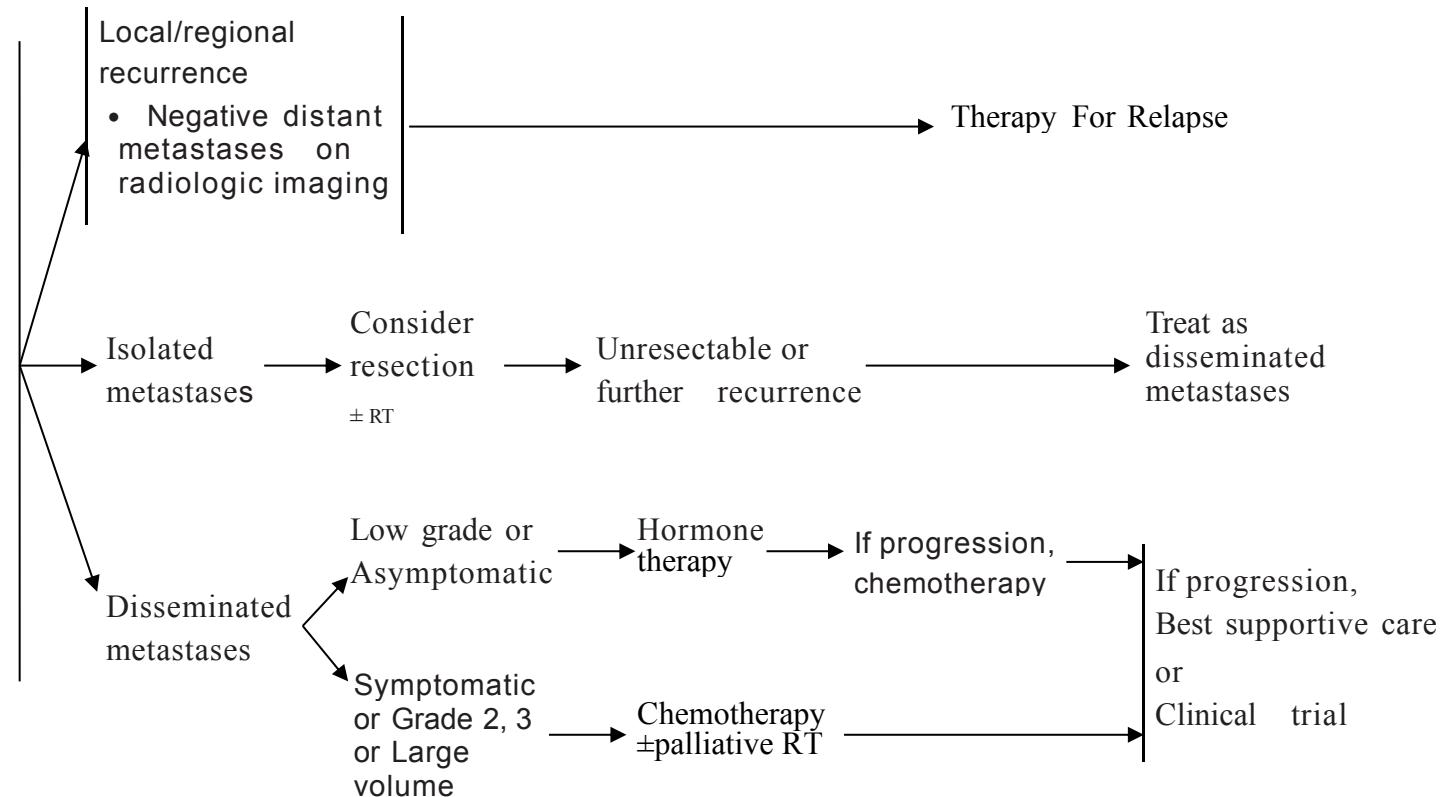
**ADJUVANT TREATMENT**



## SURVEILLANCE

- Physical exam every 3-6 mo for 2 y, then 6 mo or annually
- Vaginal cytology
- Patient education regarding symptoms, **lifestyle**, obesity
- CA-125 (optional)
- Chest x-ray annually
- CT/MRI as clinically indicated
- Consider genetic counseling/testing for young patients (< 55y) with a significant family history and/or selected pathologic risk features

## CLINICAL PRESENTATION



## CLINICAL PRESENTATION

Local/regional recurrence  
 • Negative distant metastases on radiologic imaging

No prior RT to site of recurrence

Prior RT to site of recurrence

Previous brachytherapy only

Previous external-beam RT

## THERAPY FOR RELAPSE

RT  
+brachytherapy

And/or  
Surgical exploration of pelvis + resection (category 3)

Surgical exploration of pelvis + resection ± IORT or  
Hormone therapy or  
Chemotherapy

## ADDITIONAL THERAPY

Disease confined to vagina

Pelvic lymph node  
Extra-vaginal disease

Para-aortic or common iliac lymph node

Upper abdominal/peritoneal  
Microscopic residual

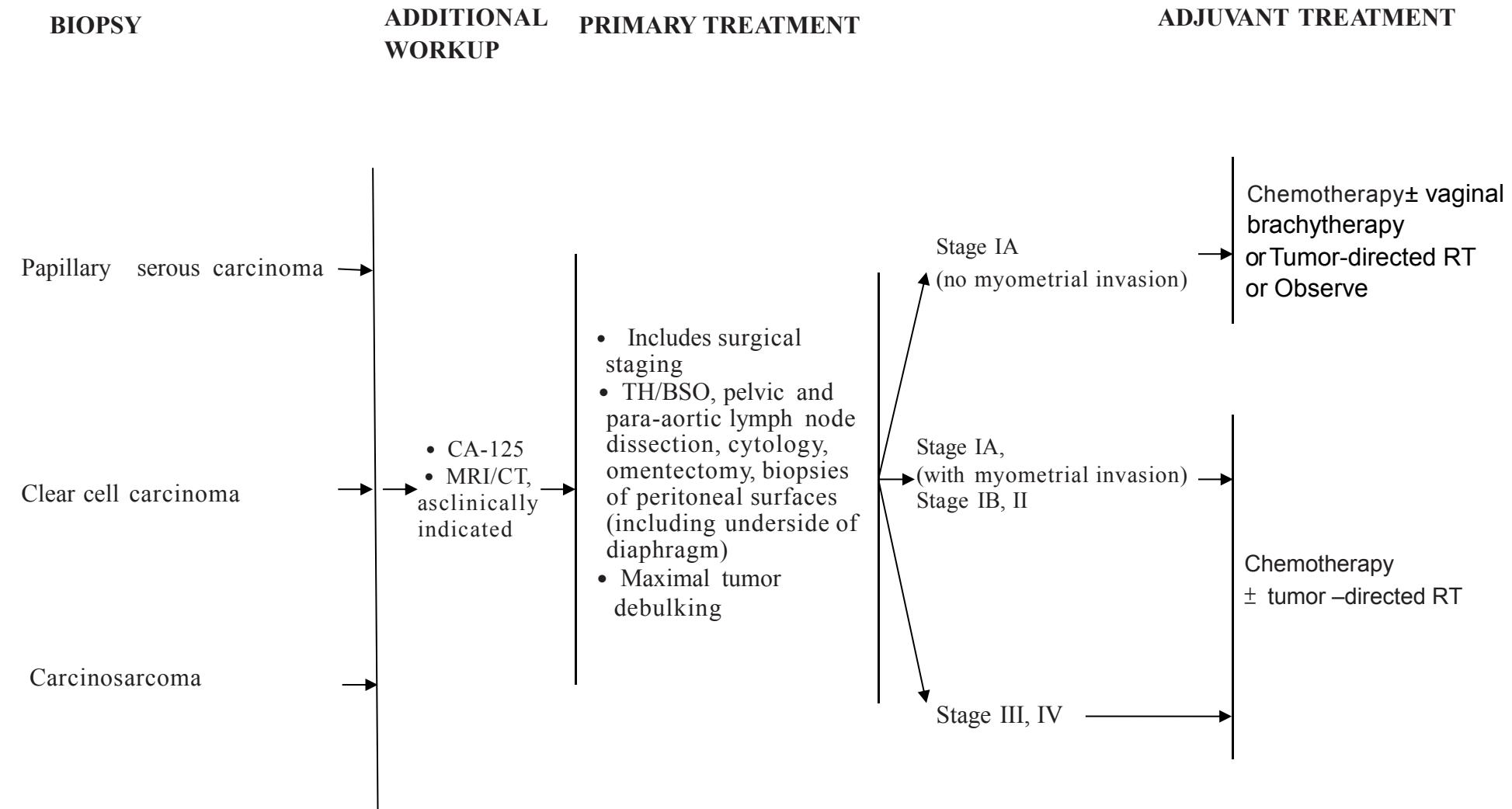
Gross upper abdominal residual disease

Tumor-directed RT  
±brachytherapy  
±chemotherapy

Tumor-directed RT  
±brachytherapy  
±chemotherapy<sup>p</sup>

Chemotherapy<sup>p</sup>  
±tumor-directed RT

Therapy For Relapse



## High Risk Patients

1. Papillary serous carcinoma
2. Clear cell carcinoma(Papillary carcinoma)
3.  $\geq$  Stage IIIA

## Risk Factor

1. Deep myometrial invasion
2. Grade III disease Stage II A&IIB
3. Lympho vascular space invasion(LVSI)
4. Age  $> 70$  age

## HORMONE THERAPY

不適用於：1.serous adenocarcinoma, clear cell adenocarcinoma, or carcinosarcoma

適用於：1.Progesterone receptor(+) 2.Well differentiation 3.Low grade

1.Prgestational agents

2.Tamoxifen 20mg/day

3.Aromatase inhibitors (自費)

4.Megestrol/tamoxifen (alternating using)

5. Medroxyprogesterone 80mg BID for 3wks 改用Tamoxifen 20mg BID

6.Megestrol 60 mg/day

## ADJUVANT CHEMOTHERAPY REGIMENS

1.Cisplatin/ Doxorubicin (每三週一次)

Cisplatin (Abiplatin) inj.(50 mg/m<sup>2</sup>) 稀釋於 N/S 500 ml IVD for 2 hours.

Doxorubicin (Adriblastina) inj. (60 mg/m<sup>2</sup>) 稀釋於 N/S 5250ml IVD for 1.5 hours.

2.Carboplatin+Paclitaxel (每三週一次)用於Endometrium Ca(自費)

Carboplatin AUC of 5-7,IV on day 1

Paclitaxel 175 mg/m<sup>2</sup> IV over 3 hours on day 1

3.Carboplatin Auc 6 IV+docetaxel 75mg/m<sup>2</sup> for IV infusion 1 hours/every 3 weeks

4.Ifosfamide/paclitaxel ( for carcinosarcoma)

a.Ifosamide 1.2gm/m<sup>2</sup>/paclitaxel 135gm/m<sup>2</sup> IV for 3 days duration every 3 weeks

b.Ifosamide 1gm/m<sup>2</sup>/day/paclitaxel 135mg/m<sup>2</sup>/day in 5% G/NS 500ml IV over 4hours for 6 cycles

5.Temsirolimus (自費)

6.Bevacizumab(自費)

## ADJUVANT RADIOTHERAPY REGIMENS

### Pelvic RT

The pelvis is treated with external beam radiation therapy to 45-50Gy, in 25-28 daily fractions using 6-10 MV photon beams. IMRT techniques are recommended to better spare normal tissues.

Vaginal Brachytherapy: 用在 High intermediates side effect 少

HDR brachytherapy, when used as a boost to EBRT: 4-6Gy in 2-3 fractions prescribed to the vaginal surface.. When used alone: 6Gy x5 prescribed to the vaginal surface.

## REFERENCE

Decision Making in Radiation Oncology, Jiade J. Lu et al, 2011

<b>FIGO</b>		<b>PRIMARY TUMOR (T)</b>
TX		Primary tumor cannot be assessed
T0		No evidence of primary tumor
Tis	*	Carcinoma in situ (preinvasive carcinoma)
T1	I	Tumor confined to corpus uteri
T1a	IA	Tumor limited to endometrium or invades less than one-half of the myometrium
T1b	IB	Tumor invades one-half or more of the myometrium
T2	II	Tumor invades stromal connective tissue of the cervix but does not extend beyond uterus**
T3a	IIIA	Tumor involves serosa and/or adnexa (direct extension or metastasis)
T3b	IIIB	Vaginal involvement (direct extension or metastasis) or parametrial involvement
T4	IVA	Tumor invades bladder mucosa and/or bowel mucosa (bulous edema is not sufficient to classify a tumor as T4)
* FIGO staging no longer includes Stage 0 (Tis)		
** Endocervical glandular involvement only should be considered as stage I and not Stage II.		
<b>REGIONAL LYMPH NODES (N)</b>		
NX		Regional lymph nodes cannot be assessed
N0		No regional lymph node metastasis
N1	IIIC1	Regional lymph node metastasis to pelvic lymph nodes
N2	IIIC2	Regional lymph node metastasis to para-aortic lymph nodes, with or without positive pelvic lymph nodes
<b>DISTANT METASTASIS (M)</b>		
M0		No distant metastasis(no pathologic M0; use clinical M to complete stage group)
M1	IVB	Distant metastasis (includes metastasis to inguinal lymph nodes intraperitoneal disease, or lung, liver, or bone. It excludes metastasis to para-aortic lymph nodes, vagina, pelvic serosa, or adnexa)

STAGE			
GROUP	T	N	M
0*	Tis	N0	M0
I	T1	N0	M0
I	T1a	N0	M0
IB	T1b	N0	M0
II	T2	N0	M0
III	T3	N0	M0
IIIA	T3a	N0	M0
IIIB	T3b	N0	M0
IIIC1	T1 - T3	N1	M0
IIIC2	T1 - T3	N2	M0
IVA	T4	Any N	M0
IVB	Any T	Any N	M1
*FIGO no longer includes Stage 0 (Tis)			
Carcinosarcomas should be staged as carcinoma.			
Stage unknown			

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