

# 子宮內膜癌診療指引

2010 年 01 月制定 2011 年 12 月修訂

2012 年 09 月修訂 2013 年 01 月修訂

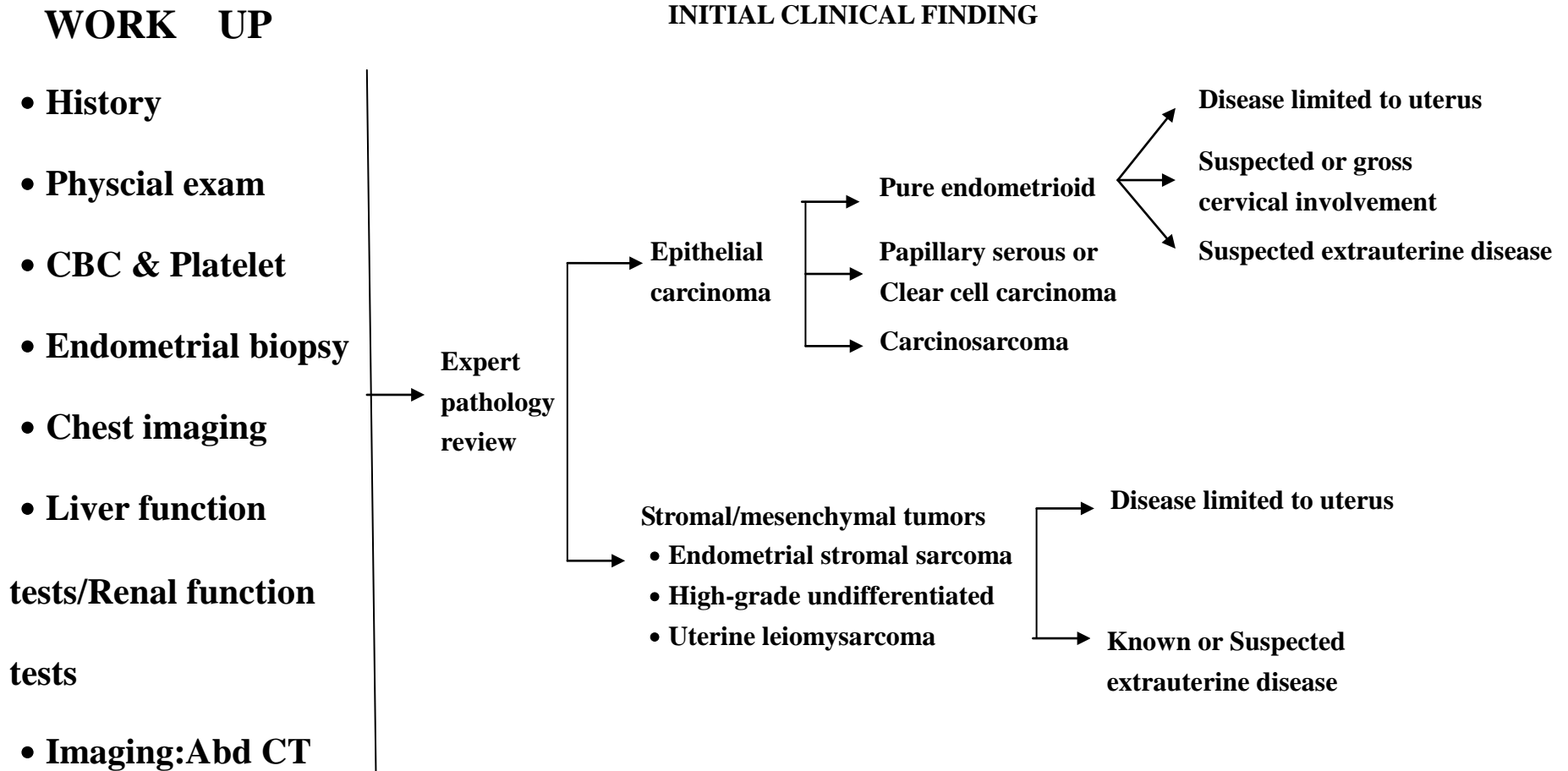
2013 年 08 月修訂 2014 年 12 月修訂

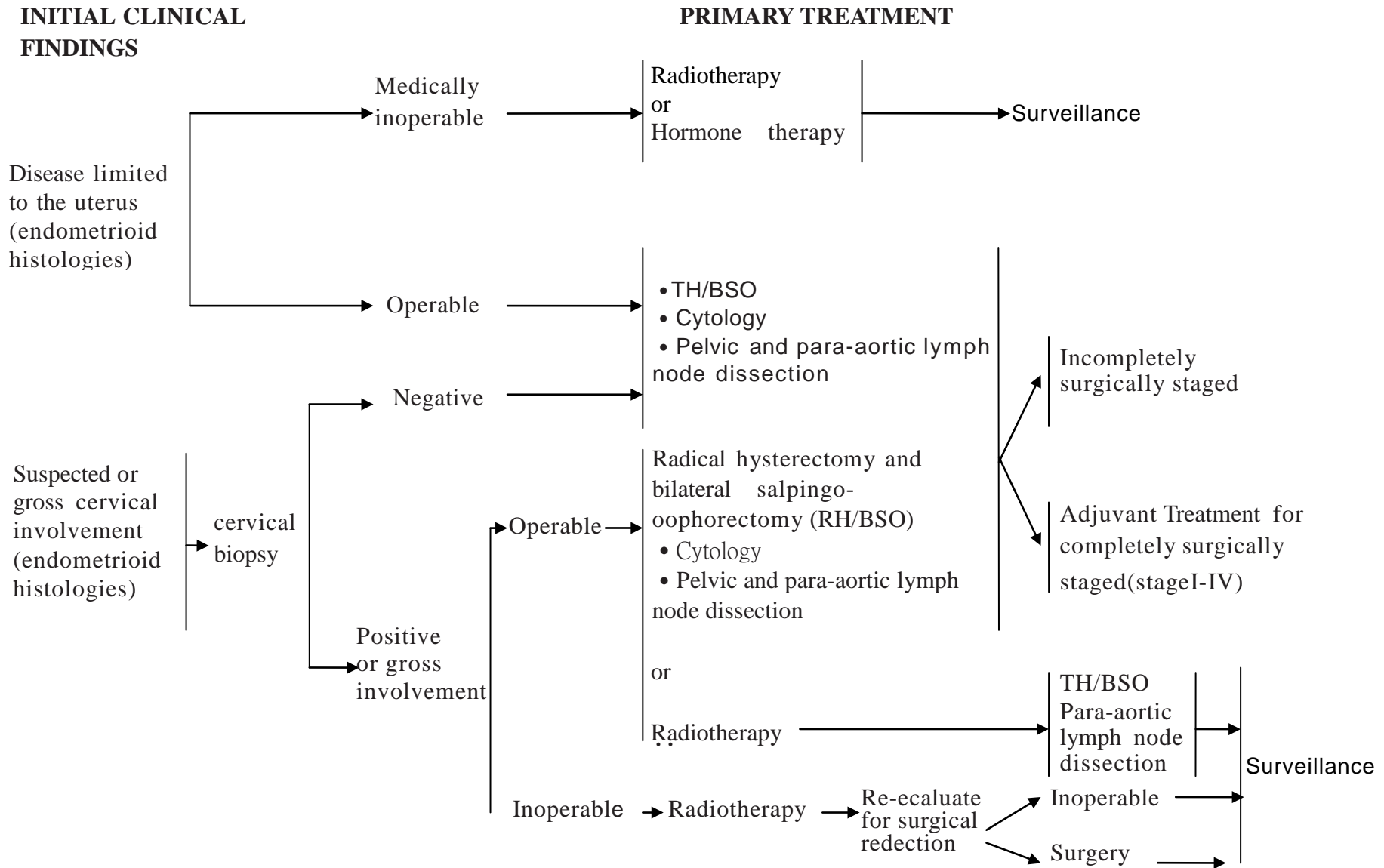
參考資料：

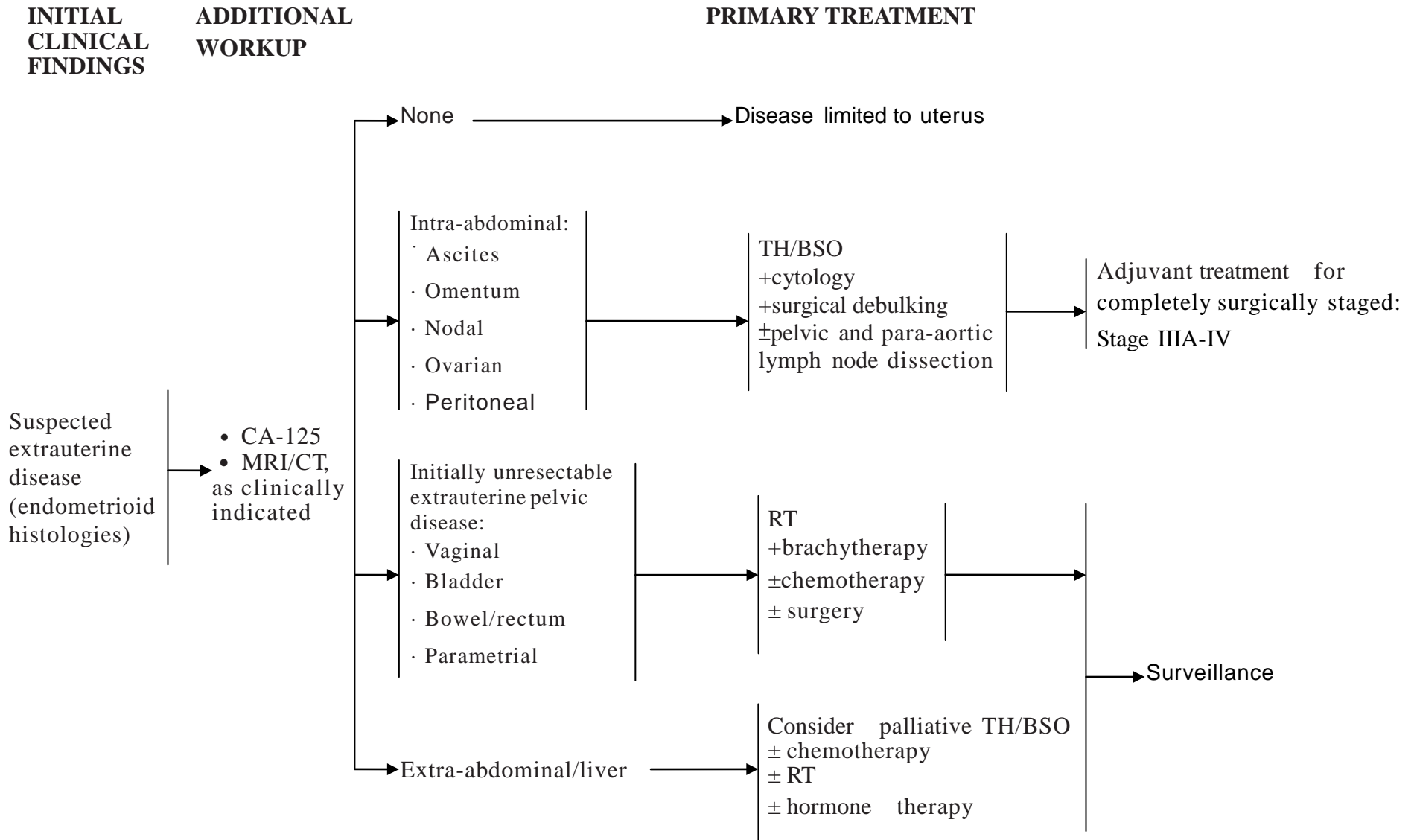
Uterine Neoplasms NCCN Guidelines V1.2015

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一百零三年版(22868\_1)





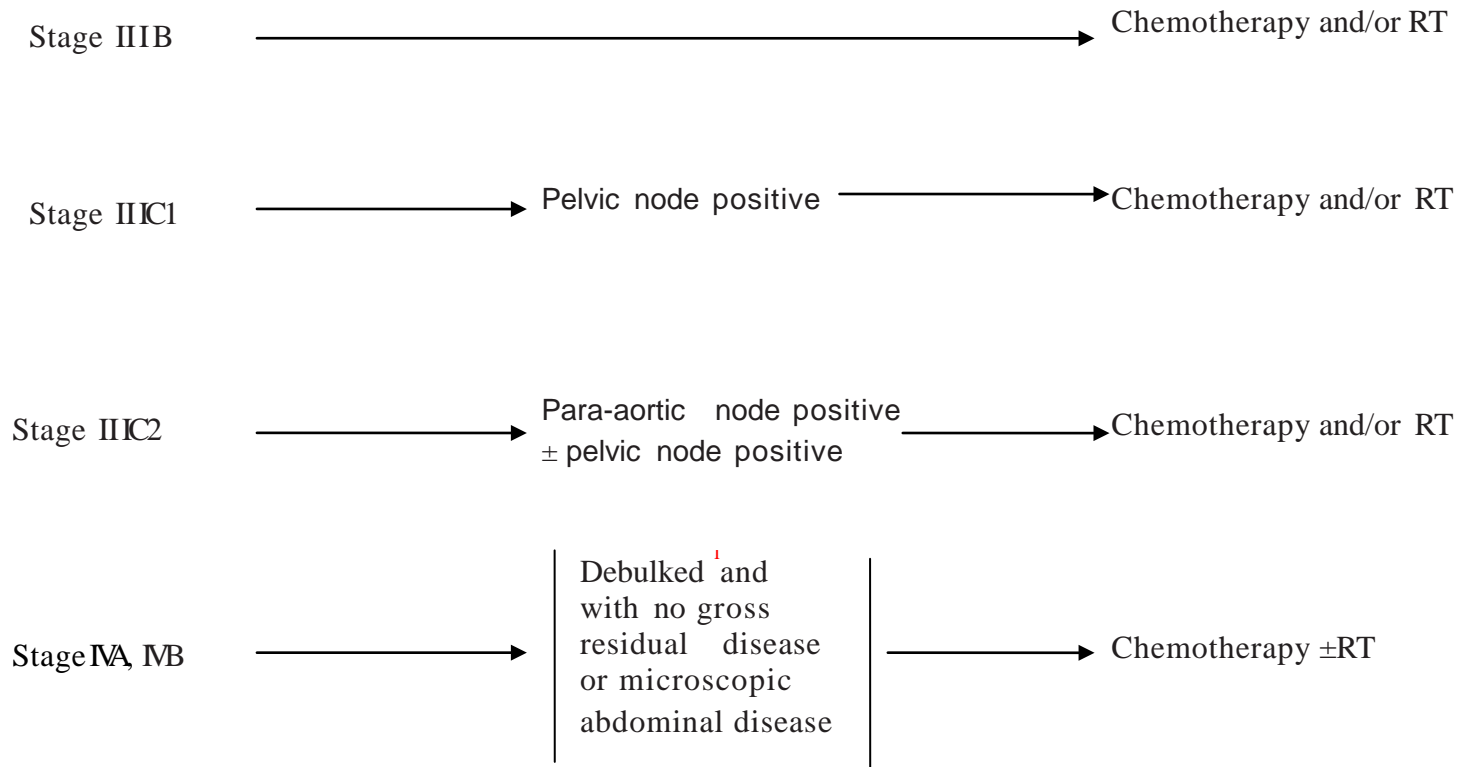


CLINICAL FINDINGS (completely surgically staged)	ADVERSE RISK FACTORS	HISTOLOGIC GRADE/ADJUVANT TREATMENT		
		G1	G2	G3
Stage IA ( $< 50\%$ ) myometrial invasion	Adverse risk factors not present	Observe	Observe or Vaginal brachytherapy	Observe or Vaginal brachytherapy
	Adverse risk factors present	Observe or Vaginal brachytherapy	Observe or Vaginal brachytherapy and/or pelvic RT	Observe or Vaginal brachytherapy and/or pelvic RT
Stage IB ( $\geq 50\%$ ) myometrial invasion	Adverse risk factors not present	Vaginal brachytherapy	Vaginal brachytherapy and/or pelvic RT	pelvic RT and/or Vaginal brachytherapy
	Adverse risk factors present	Vaginal brachytherapy and/or pelvic RT	Vaginal brachytherapy and/or pelvic RT	Pelvic RT and/or Vaginal brachytherapy $\pm$ chemotherapy
Stage II		pelvic RT and Vaginal brachytherapy	pelvic RT +Vaginal brachytherapy	pelvic RT +Vaginal brachytherapy $\pm$ chemotherapy
Stage IIIA		chemotherapy $\pm$ RT or RT $\pm$ chemotherapy or pelvic RT $\pm$ Vaginal brachytherapy	chemotherapy $\pm$ RT or RT $\pm$ chemotherapy or pelvic RT $\pm$ Vaginal brachytherapy	chemotherapy $\pm$ RT or RT $\pm$ chemotherapy or pelvic RT $\pm$ Vaginal brachytherapy

Adverse Risk Factors: Age, Lymphovascular invasion, tumor size

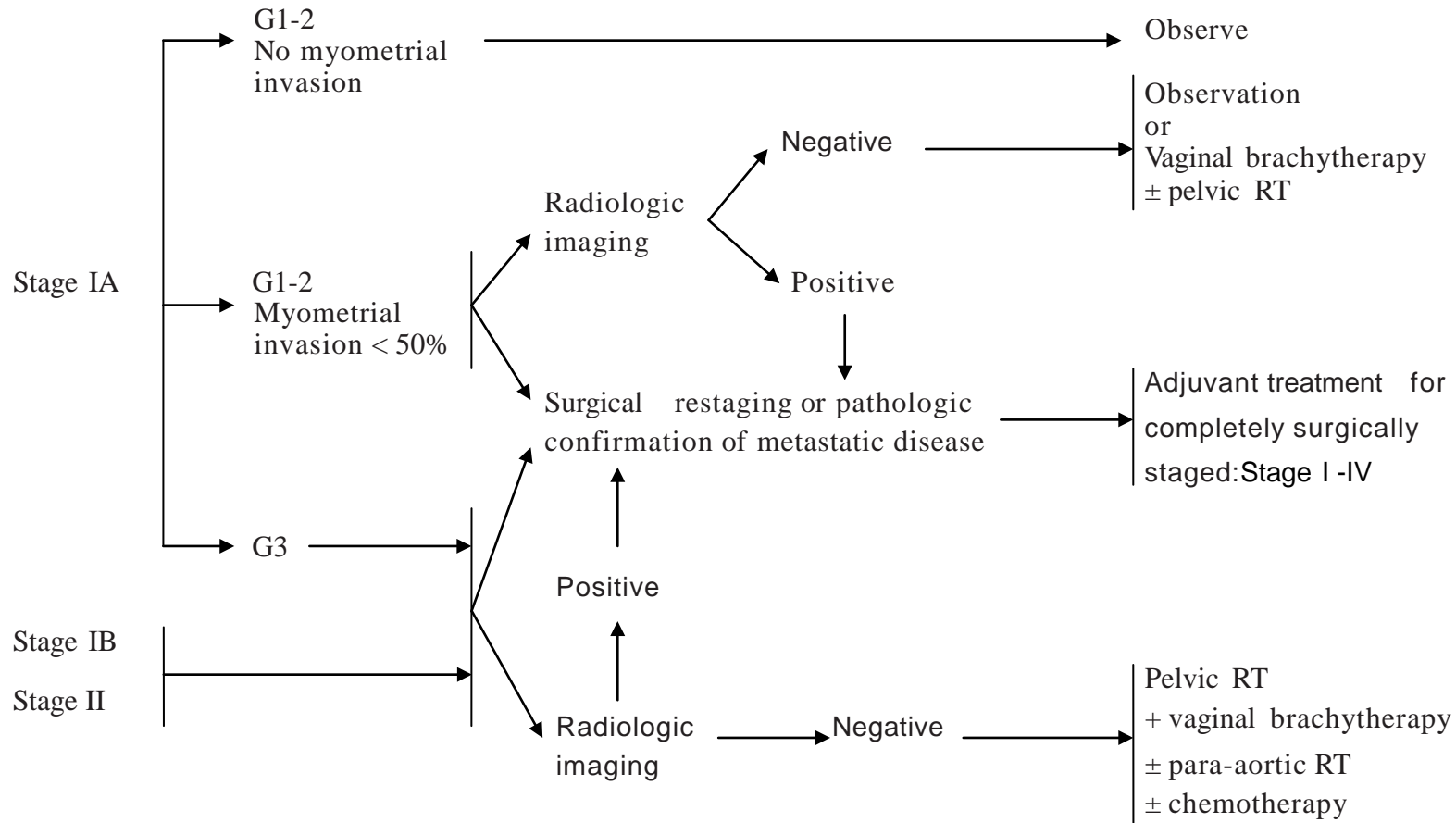
**CLINICAL FINDINGS**  
(completely surgically staged)

**ADJUVANT TREATMENT**



CLINICAL FINDINGS  
(Incompletely surgically staged)

ADJUVANT TREATMENT



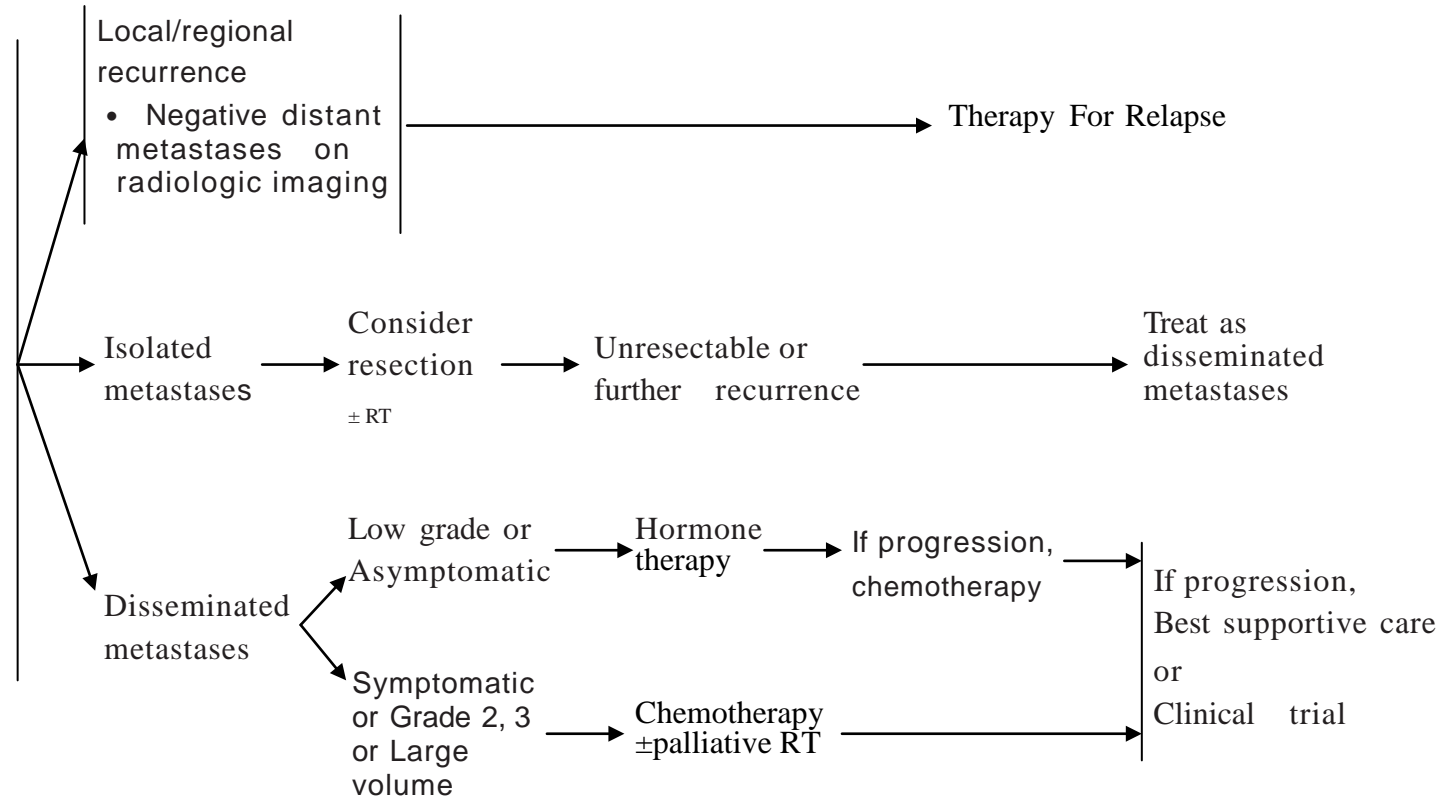


SURVEILLANCE

CLINICAL PRESENTATION

THERAPY FOR RELAPSE

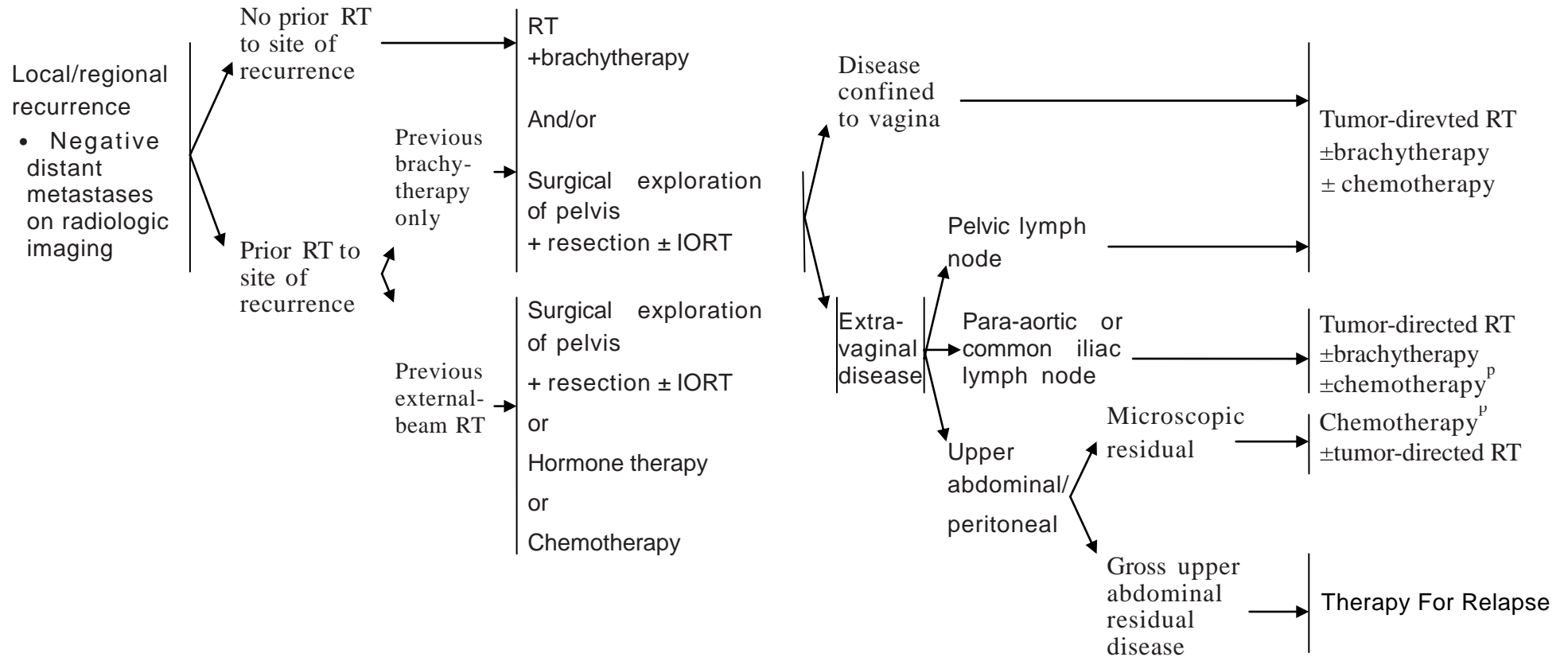
- Physical exam every 3-6 mo for 2 y, then 6 mo or annually
- Vaginal cytology
- Patient education regarding symptoms
- CA-125 (optional)
- Chest x-ray annually
- CT/MRI as clinically indicated
- Consider genetic counseling/testing for young patients (< 55y) with a significant family history and/or selected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

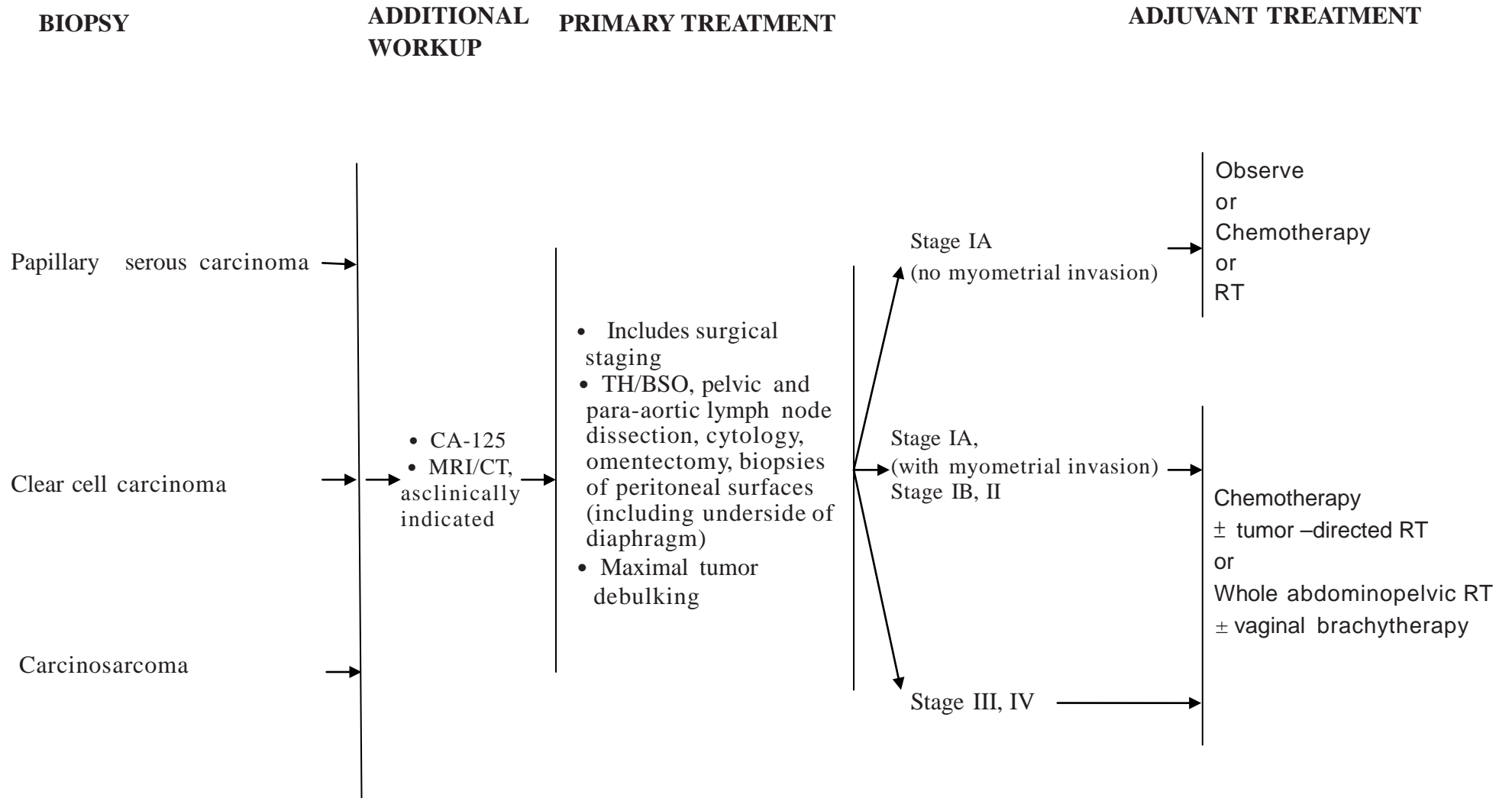


CLINICAL PRESENTATION

THERAPY FOR RELAPSE

ADDITIONAL THERAPY





**High Risk Patients**

1. Serous carcinoma
2. Clear cell carcinoma(Papillary carcinoma)
3. Stage III

**Risk Factor**

1. Deep myometrial invasion
2. Grade III disease
3. Lympho vascular space invasion(LVSI)
4. Age > 70 age

## HORMONE THERAPY

- Progestational agents
- Tamoxifen
- Aromatase inhibitors

## ADJUVANT CHEMOTHERAPY REGIMENS

- Cisplatin/ Doxorubicin

Cisplatin (Abliplatin) 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.

- Plan B: Carboplatin+Paclitaxel

Carboplatin      AUC of 5-7,IV on day 1

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

- R/T concomitant Cisplatin **50 mg/m<sup>2</sup>**

Follow by

Carboplatin      AUC of 5-7,IV on day 1

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

## **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

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 (bullous 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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