

# 食道癌診療指引

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參考資料：NCCN Guidelines Version

Esophageal and Esophagogastric Junction Cancers V1.2015

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☆：選項

EUS：Endoscopic ultrasound

EMR：Endoscopic mucosal resection

BSC：Best supportive care

CCRT：Chemoradiation

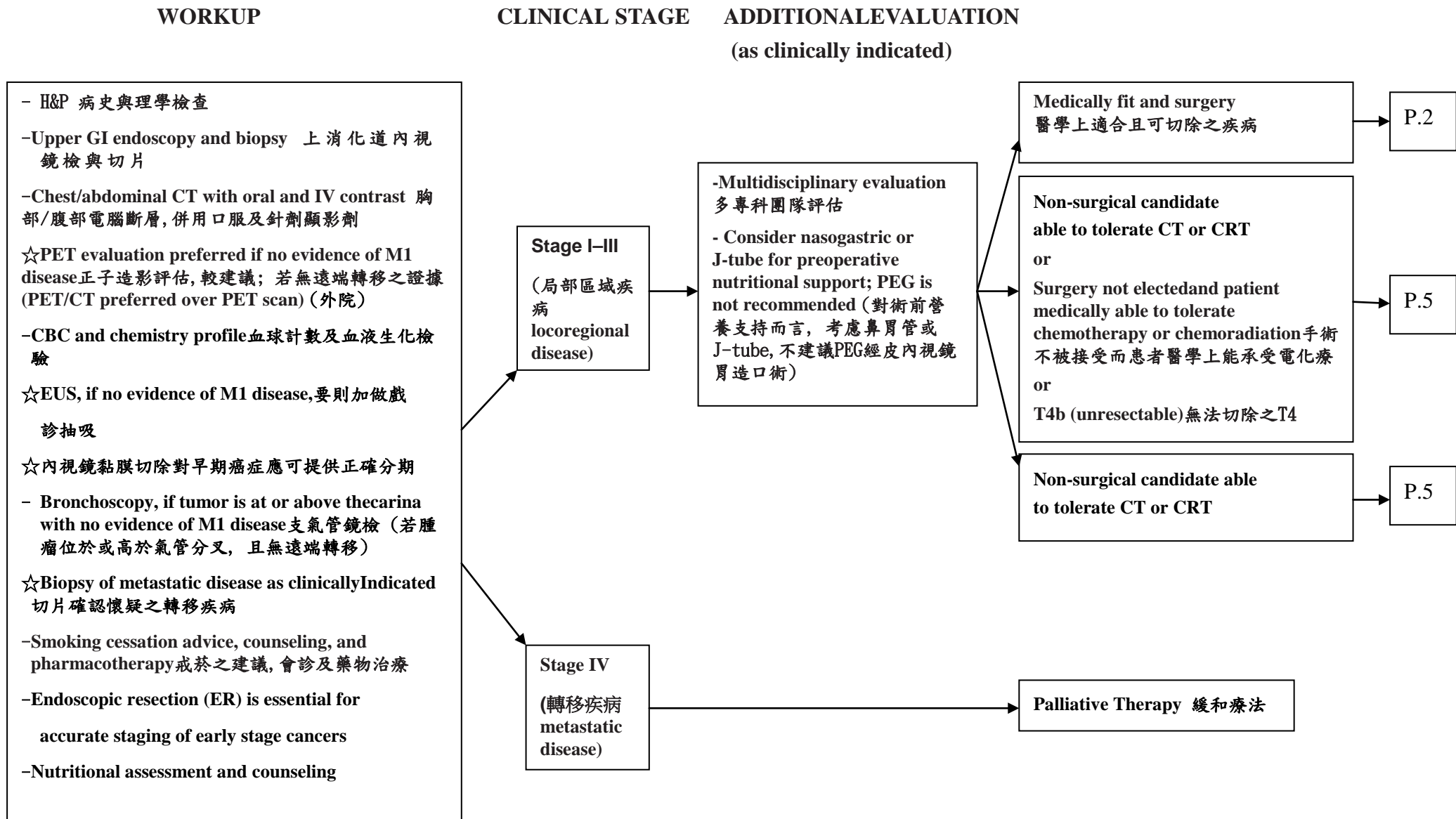
C/T：Chemotherapy

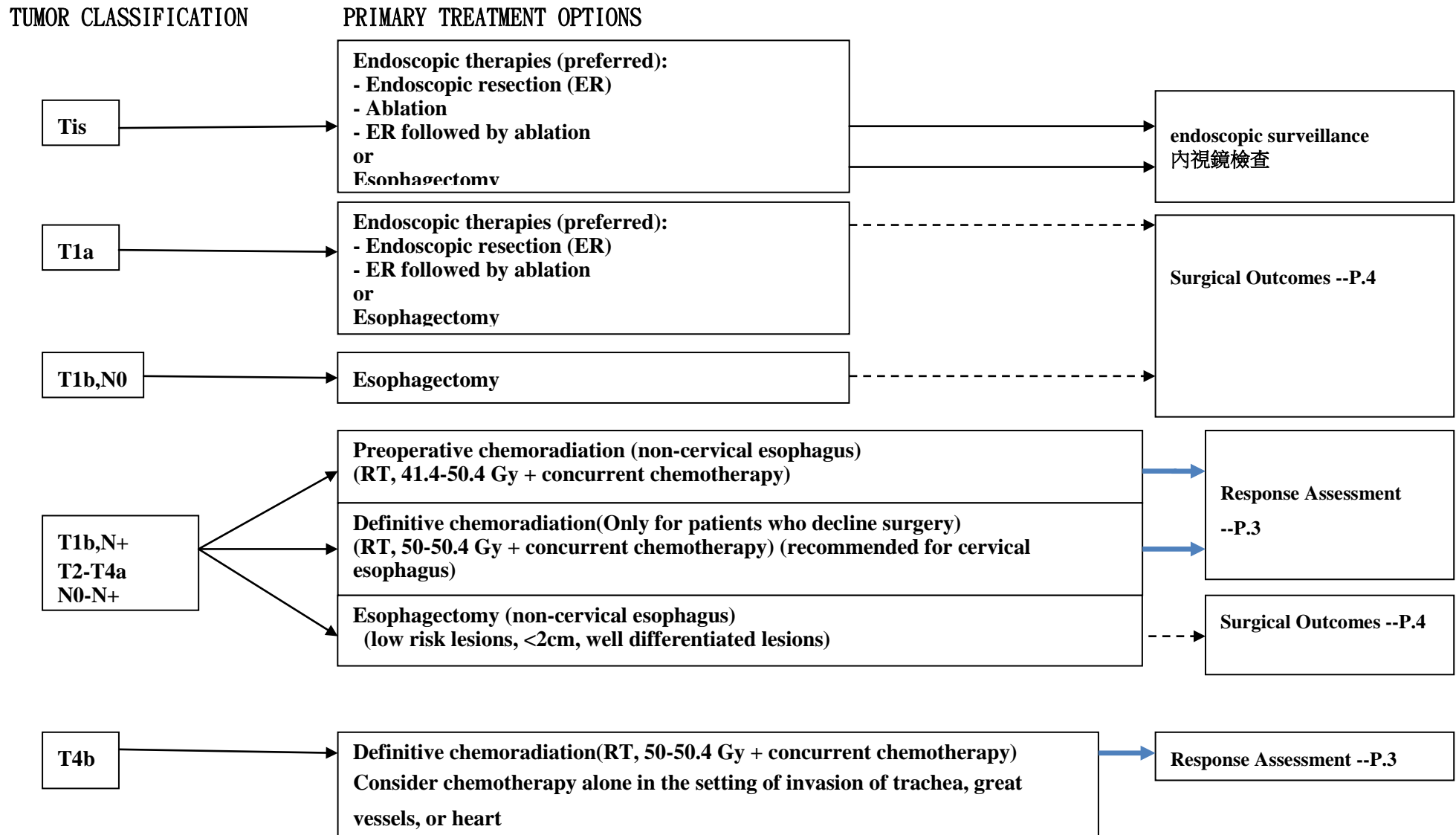
RT：Radiotherapy

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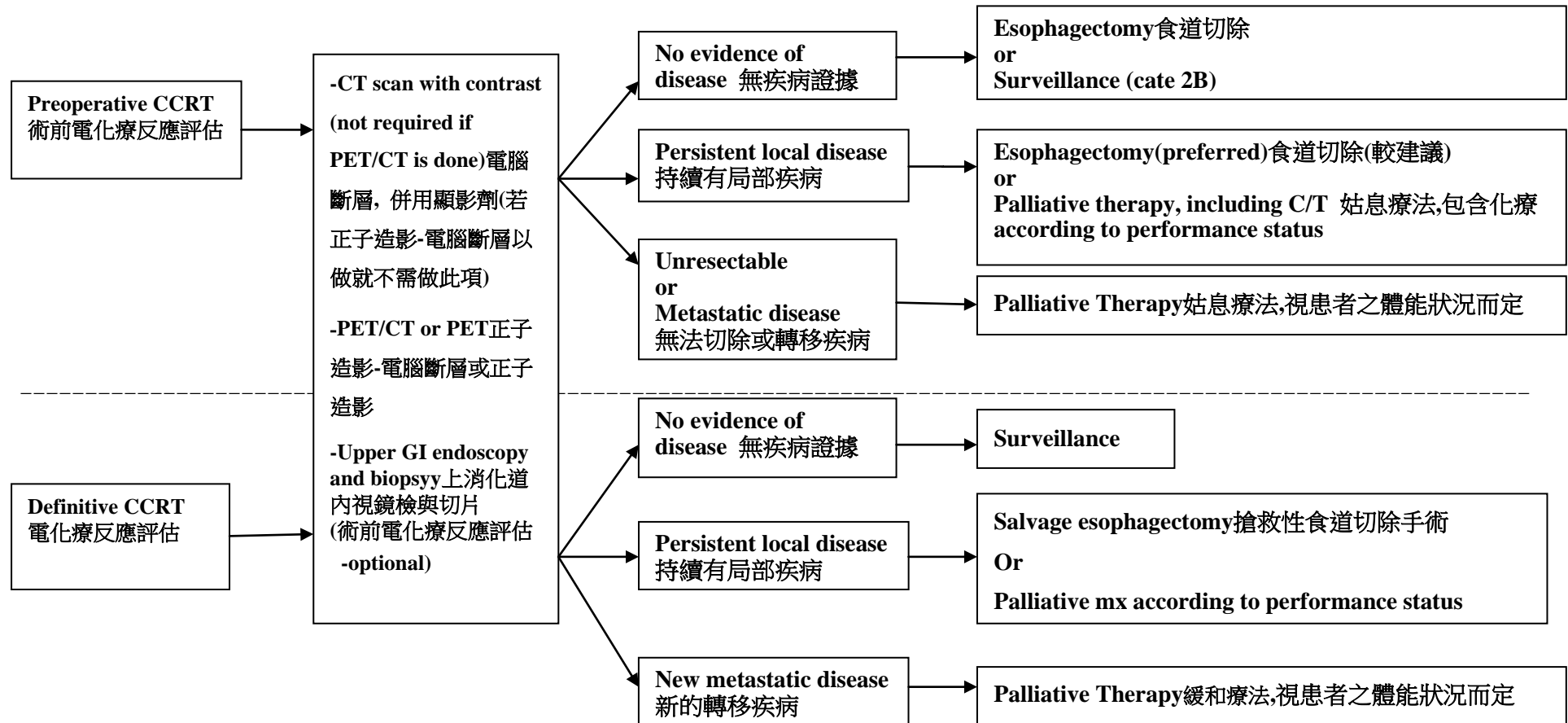
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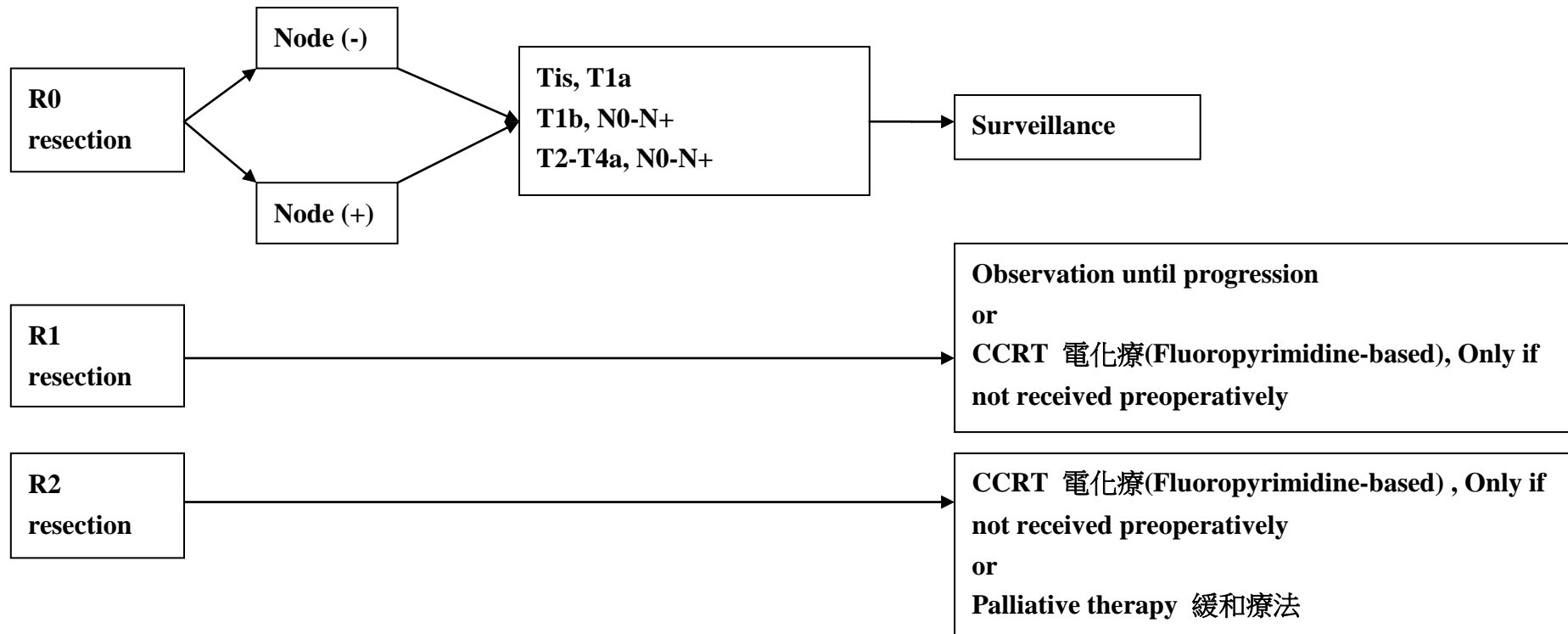




RESPONSEASSESSMENTOUTCOMEADJUVANT TREATMENT

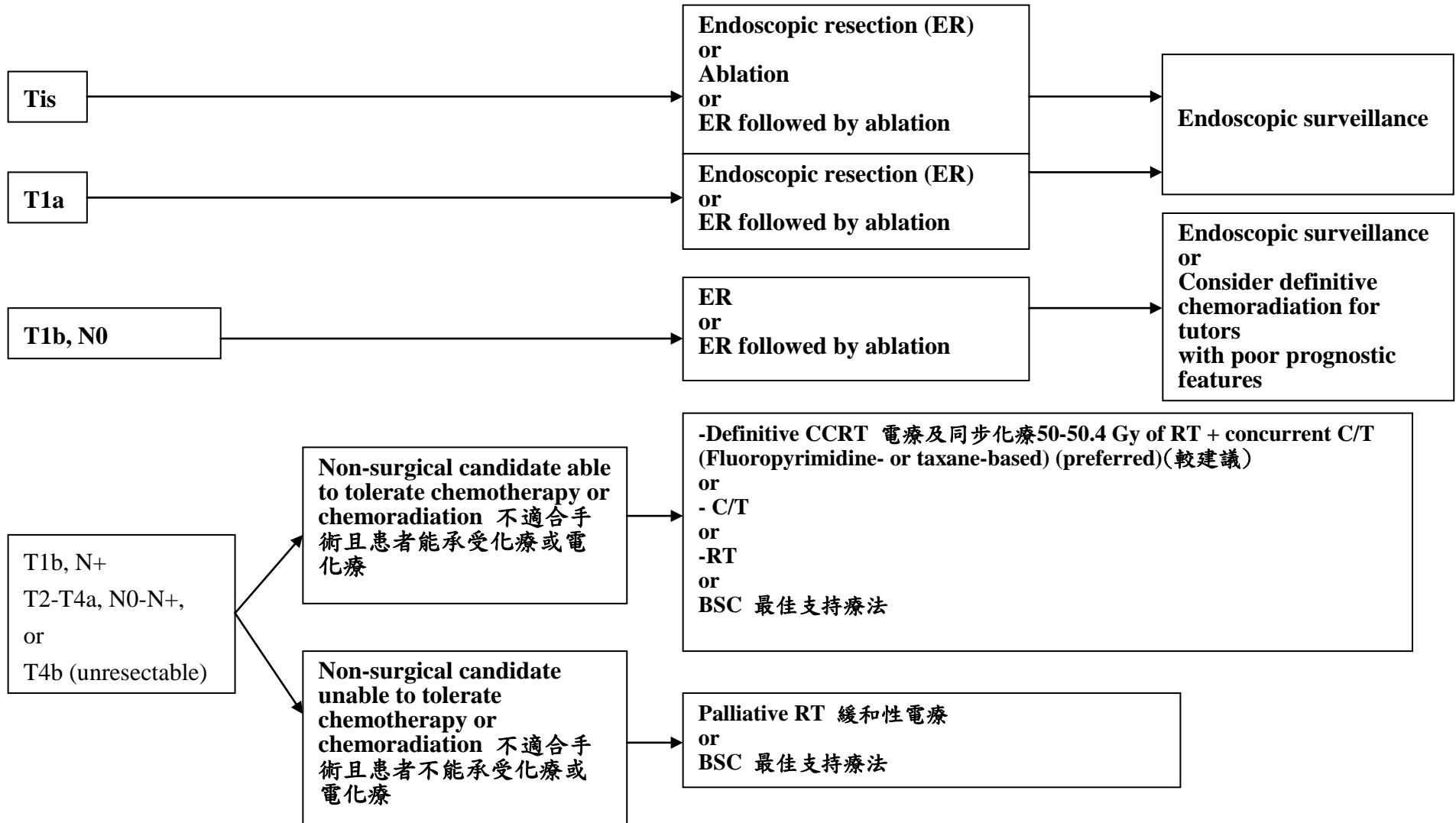


**SURGICAL OUTCOMES/CLINICAL      PATHOLOGIC FINDINGS      POSTOPERATIVE TREATMENT**  
(Patients Have Received Preoperative Chemoradiation or Chemotherapy)

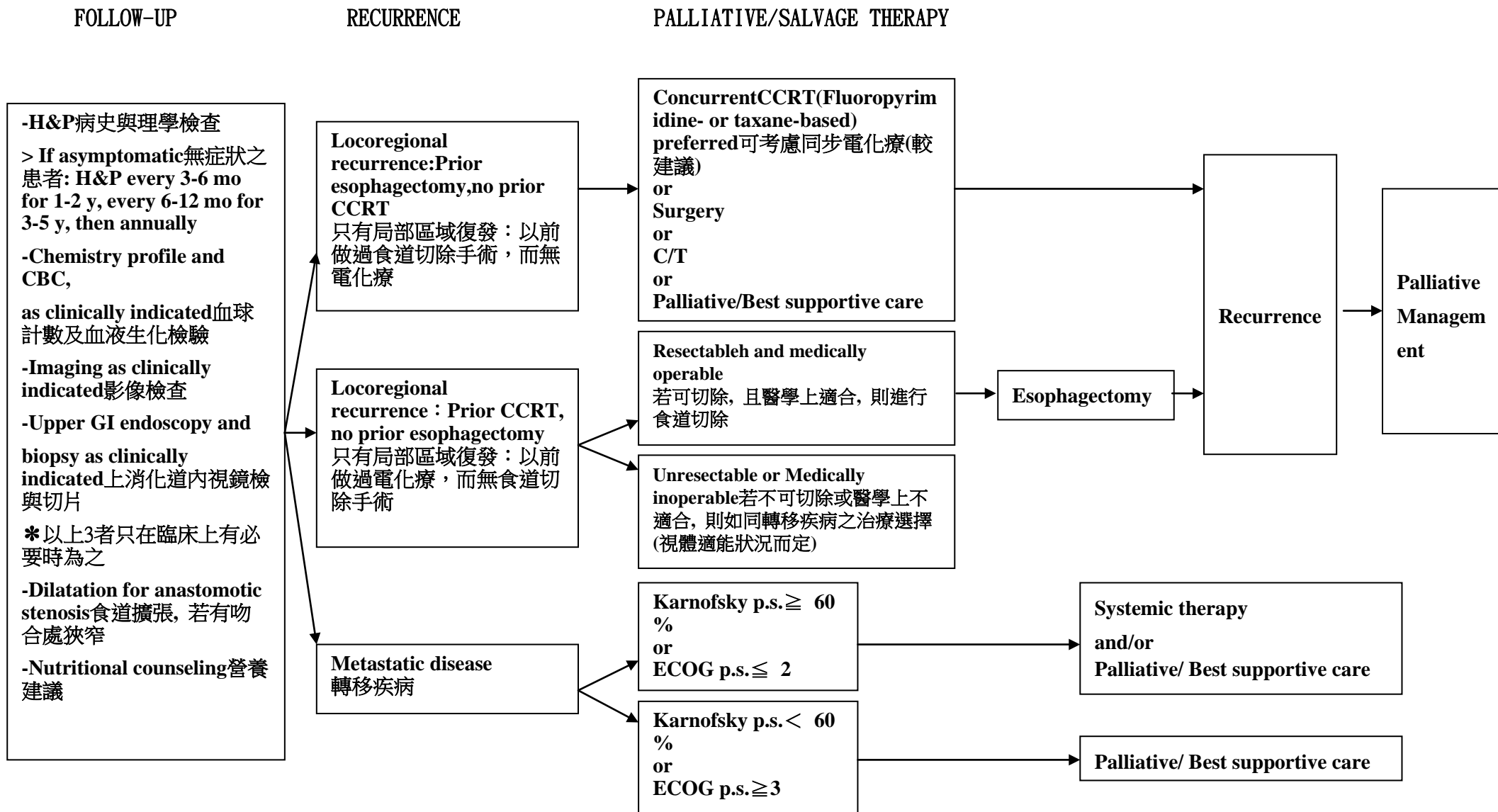


**R0 : No cancer at resection margins    R1 : Microscopic residual cancer    R2 : Macroscopic residual cancer or M1b**

PRIMARY TREATMENT FOR NON-SURGICAL CANDIDATE







## Chemotherapy Regimens

### Preoperative Chemoradiation

- **Paclitaxel and carboplatin (optinal)**  
Paclitaxel 50 mg/m<sup>2</sup>IV on Day 1  
CarboplatinAUC 2 IV on Day 1  
Weekly for 5 weeks
- **Cisplatin and fluoropyrimidine (5-FU or capecitabine)**  
Cisplatin 75-100 mg/m<sup>2</sup>IV on Days 1 and 29  
5-FU 750-1000 mg/m<sup>2</sup>IV continuous infusion  
over 24 hours daily on Days 1-4 and 29-32 35-Day cycle  
  
Cisplatin 30 mg/m<sup>2</sup>IV on Day 1 (optional)  
Capecitabine 800 mg/m<sup>2</sup>PO BID on Days 1-5  
Weekly for 5 weeks

Definitive Chemoradiation

- Cisplatin and fluoropyrimidine (5-FU or capecitabine)

Cisplatin 75-100 mg/m<sup>2</sup>IV on Day 1

5-FU 750-1000 mg/m<sup>2</sup>IV continuous infusion over 24 hours daily on Days 1-4

Cycled every 28 days for 2-4 cycles for 2 cycles with radiation followed by 2 cycles without radiation

Cisplatin 30 mg/m<sup>2</sup>IV on Day 1 (optional)

Capecitabine 800 mg/m<sup>2</sup>PO BID on Days 1-5

Weekly for 5 weeks

- Paclitaxel or docetaxel and cisplatin (optional)

Paclitaxel 60 mg/m<sup>2</sup>IV on Days 1, 8, 15, and 22

Cisplatin 75 mg/m<sup>2</sup>IV on Day

Given for 1 cycle

Docetaxel 60 mg/m<sup>2</sup>IV on Days 1 and 22

Cisplatin 60-80 mg/m<sup>2</sup>IV on Days 1 and 22

Given for 1 cycle

Docetaxel 20-30 mg/m<sup>2</sup> IV on Day 1

Cisplatin 20-30 mg/m<sup>2</sup>IV on Day 1

Weekly for 5 weeks

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## References

1. vanMeertenE, MullerK, TilanusHW, et al. Neoadjuvant concurrent chemoradiation with weekly paclitaxel and carboplatin for patients with esophageal cancer: a phase II study. *Br J Cancer* 2006;94:1389-1394.
2. GaastAV, vanHagen P, HulshofM, et al. Effect of preoperative concurrent chemoradiotherapy on survival of patients with resectable esophageal or esophagogastric junction cancer: Results from a multicenter randomized phase III study. *J Clin Oncol (Meeting Abstracts)* 2010;28:4004-.
3. TepperJ, KrasnaMJ, NiedzwieckiD, et al. Phase III trial of trimodality therapy with cisplatin, fluorouracil, radiotherapy, and surgery compared with surgery alone for esophageal cancer: CALGB9781. *J Clin Oncol* 2008;26:1086-1092.
4. LeeSS, KimSB, ParkSI, et al. Capecitabine and cisplatin chemotherapy (XP) alone or sequentially combined chemoradiotherapy containing XP regimen in patients with three different settings of stage IV esophageal cancer. *Jpn J Clin Oncol* 2007;37:829-835.
5. BedenneL, MichelP, BoucheO, et al. Chemoradiation followed by surgery compared with chemoradiation alone in squamous cancer of the esophagus: FFCD9102. *J Clin Oncol* 2007;25:1160-1168.
6. MinskyBD, PajakTF, Ginsberg RJ, et al. INT 0123 (Radiation Therapy Oncology Group 94-05) phase III trial of combined-modality therapy for esophageal cancer: high-dose versus standard-dose radiation therapy. *J Clin Oncol* 2002;20:1167-1174
7. UrbaSG, OrringerMB, IannettoniM, et al. Concurrent cisplatin, paclitaxel, and radiotherapy as preoperative treatment for patients with locoregional esophageal carcinoma. *Cancer* 2003;98:2177-2183.
8. LiQQ, LiuMZ, HuYH, et al. Definitive concomitant chemoradiotherapy with docetaxel and cisplatin in squamous esophageal carcinoma. *Dis Esophagus* 2010;23:253-259.
9. DayFL, LeongT, NganS, et al. Phase I trial of docetaxel, cisplatin and concurrent radical radiotherapy in locally advanced esophageal cancer. *Br J Cancer* 2011;104:265-271.

## 食道癌放射治療線治療

放射治療的適應症：

- (一) 第 II-III 期並可進行根治手術者：手術前進行新輔助合併化學放射治療 (neoadjuvant CCRT)，或在手術後進行輔助性合併化學放射治療 (adjuvant CCRT)。
- (二) 第 I-III 期無法進行根治手術者：合併化學放射治療(CCRT)或單純治癒性放射治療(definitive curative radiotherapy alone)(拒絕或不適化療者)
- (三) 第 IV 期：針對轉移部位(如骨骼、腦等部位)或原發部位施行緩解性放射治療。

二、放射治療執行情序(procedures)：

(一) 電腦斷層模擬攝影(CT-based simulation)

1. 仰臥、雙手置於頭頂，並以真空氣墊(vaccum pillow) or alpha cradle 固定姿勢
2. 以雷射光於病人腹部、身體兩側劃上等中心(isocenter)記號
3. 透過靜脈注射顯影劑，可加強判讀腫瘤侵犯之範圍，但如果病患腎功能差(creatinine>2.0 mg/dl)或其他禁忌症為例外。
4. 每 3-5 毫米擷取一張電腦斷層影像
5. 將影像傳送至電腦治療計劃系統(radiation treatment plan, RTP system)

(二) 描繪標靶體積 (contouring target volume)

1. 標靶體積(Gross Target Volume, GTV):應包含由 CT 影像及可判讀之主要腫瘤(primary tumor)及臨床上呈陽性之淋巴腺。
2. 臨床標靶體積(Clinical Target Volume, CTV): 為包含 GTV 可能侵犯之範圍，由 GTV 頭腳方向(cranio-caudal directions) 加上 3~5 公分、左右前後側加上 0.5~1.0 公分之三度空間之範圍。
3. 計畫標靶體(Planning Target Volume, PTV):  $CTV + 0.5-1\text{ cm}$

(三) Organs At Risk (OAR):

1 脊索(Spinal cord)

(1) 圈選範圍：所有含 PTV 的橫切面(axial planes)影像圈選畫脊索，外加頭腳方向

要再加圈選至少 1 公分之脊索。另再組成範圍 5 毫米的 PRV。

(2) 劑量限制：最高劑量 < 50 Gy。

2 肺(Lung)

(1) 圈選範圍：整個肺部(不包含 CTV)，分別圈選左、右及整個肺部。

(2) 劑量限制：整個肺部之平均劑量建議 < 20 Gy、整個肺部之  $V_{20Gy} < 35\%$  (20 Gy 劑量之體積，應低於肺總體積之 35%)，如肺功能較差之病患，整個肺部之  $V_{20Gy} < 30\%$ 。

### 3 肝臟(Liver)

- (1) 圈選範圍：胸腔內之食道(不含 PTV)。
- (2) 劑量限制：1/2 之體積應低於 35 Gy、2/2 之體積應低於 30 Gy。

### 4 腎臟(Kidney)

- (1) 圈選範圍：分別圈選左、右可看到之腎臟。
- (2) 劑量限制：1/3 之體積應低於 50 Gy、2/3 之體積應低於 32 Gy、所有體積應低於 23Gy。

### 5 心臟(Heart)

- (1) 圈選範圍：影像中可圈選之心臟組織。
- (2) 劑量限制：1/3 之體積應低於 60 Gy、2/3 之體積應低於 45 Gy、所有體積應低於 40 Gy。

(四) 劑量處方 (dose prescription) :

1. 手術前進行新輔助合併化學放射治療 (neoadjuvant CCRT) : 每分次 1.8-2.0 格雷, 每週五~六分次, 五週總劑量為 45-50.4 格雷(1.8-2.0 fraction dose, 5-6 fractions per week, total dose 45-50.4Gy over 5 weeks)
2. 手術後進行輔助性合併化學放射治療 (adjuvant CCRT): 每分次 1.8-2.0 格雷, 每週五~六分次, 五週總劑量為 45-50.4 格雷(1.8-2.0 fraction dose, 5-6 fractions per week, total dose 45-50.4Gy over 5 weeks), 根治性病患可縮小範圍(GTV)續照至 50-66Gy
3. 合併化學放射治療(CCRT)或單純治癒性放射治療(definitive curative radiotherapy alone): 每分次 1.8-2.0 格雷, 每週五~六分次, 五週總劑量為 45-50.4 格雷(1.8-2.0 fraction dose, 5 fractions per week, total dose 45-50.4Gy over 5 weeks), 根治性病患可縮小範圍(GTV)續照至 50-66Gy

(五) 體外放射治療技術 (external radiotherapy technique) :

1. 三維順形放射治療(3-Dimension Conformal Radiation Therapy, 3D-CRT)
2. 強度調控放射治療(Intensity Modulation Radiation Therapy, IMRT)
3. 影像導引放射治療(Image-guided Radiation Therapy, IGRT)

(六) 治療驗證(Treatment Verification)

1. 三度空間放射治療或強度調控放射治療: 治療前及每周應由放射師拍攝正交之驗證片(orthogonal verification films)來驗證照野之中心點。
2. 影像導引放射治療(IGRT): 如放射治療設備備有影像導引功能, 治療前及每周應由放射師拍攝電腦斷層影像或正交之驗證片確認治療範圍。



參考資料:

1. NCCN clinical practice guideline in Esophageal cancer (V.2.2011)
2. Principles and practice of radiation oncology, 5<sup>th</sup> ed.
3. International Commission on Radiation Units and Measurements. ICRU Report No 50:Prescribing, Recording and Reporting Photon Beam Therapy. Bethesda, MD: ICRUPublications 1993.
4. International Commission on Radiation Units and Measurements. ICRU Report No 62:Prescribing, Recording and Reporting Photon Beam Therapy (Supplement to ICRU Report 50).Bethesda, MD: ICRU Publications 1999.

<b>PRIMARY TUMOR (T)</b>	
<b>TX</b>	<b>Primary tumor cannot be assessed</b>
<b>T0</b>	<b>No evidence of primary tumor</b>
<b>Tis</b>	<b>High-grade dysplasia *</b>
<b>T1</b>	<b>Tumor invades lamina propria, muscularis mucosae, or submucosa</b>
<b>T1a</b>	<b>Tumor invades lamina propria or muscularis mucosae</b>
<b>T1b</b>	<b>Tumor invades submucosa</b>
<b>T2</b>	<b>Tumor invades muscularispropria</b>
<b>T3</b>	<b>Tumor invades adventitia</b>
<b>T4</b>	<b>Tumor invades adjacent structures</b>
<b>T4a</b>	<b>Resectable tumor invading pleura, pericardium, or diaphragm</b>
<b>T4b</b>	<b>Unresectable tumor invading other adjacent structures, such as aorta, vertebral body, trachea, etc.</b>
<p><b>*High-grade dysplasia includes all non-invasive neoplastic epithelium that was formerly called carcinoma in situ, a diagnosis that is no longer used for columnar mucosae anywhere in the gastrointestinal tract.</b></p>	

<b>REGIONAL LYMPH NODES (N)</b>	
<b>NX</b>	<b>Regional lymph nodes cannot be assessed</b>
<b>N0</b>	<b>No regional lymph node metastasis</b>
<b>N1</b>	<b>Regional lymph node metastases involving 1 to 2 nodes</b>
<b>N2</b>	<b>Regional lymph node metastases involving 3 to 6 nodes</b>
<b>N3</b>	<b>Regional lymph node metastases involving 7 or more nodes</b>

<b>DISTANT METASTASIS (M)</b>	
<b>M0</b>	<b>No distant metastasis (no pathologic M0; use clinical M to complete stage group)</b>
<b>M1</b>	<b>Distant metastasis</b>

<b>ANATOMIC STAGE • PROGNOSTIC GROUPS</b>					
<b>Squamous Cell Carcinoma</b>					
<b>GROUP</b>	<b>T</b>	<b>N</b>	<b>M</b>	<b>Grade</b>	<b>Tumor Location**</b>
<b>0</b>	<b>Tis(HGD)</b>	<b>N0</b>	<b>M0</b>	<b>1</b>	<b>Any</b>
<b>IA</b>	<b>T1</b>	<b>N0</b>	<b>M0</b>	<b>1,X</b>	<b>Any</b>
<b>IB</b>	<b>T1</b>	<b>N0</b>	<b>M0</b>	<b>2-3</b>	<b>Any</b>
	<b>T2-3</b>	<b>N0</b>	<b>M0</b>	<b>1,X</b>	<b>Lower,X</b>
<b>IIA</b>	<b>T2-3</b>	<b>N0</b>	<b>M0</b>	<b>1,X</b>	<b>Upper,middle</b>
	<b>T2-3</b>	<b>N0</b>	<b>M0</b>	<b>2-3</b>	<b>Lower,X</b>
<b>IIB</b>	<b>T2-3</b>	<b>N0</b>	<b>M0</b>	<b>2-3</b>	<b>Upper,middle</b>
	<b>T1-2</b>	<b>N1</b>	<b>M0</b>	<b>Any</b>	<b>Any</b>
<b>IIIA</b>	<b>T1-2</b>	<b>N2</b>	<b>M0</b>	<b>Any</b>	<b>Any</b>
	<b>T3</b>	<b>N1</b>	<b>M0</b>	<b>Any</b>	<b>Any</b>
	<b>T4a</b>	<b>N0</b>	<b>M0</b>	<b>Any</b>	<b>Any</b>
<b>IIIB</b>	<b>T3</b>	<b>N2</b>	<b>M0</b>	<b>Any</b>	<b>Any</b>
<b>IIIC</b>	<b>T4a</b>	<b>N1-2</b>	<b>M0</b>	<b>Any</b>	<b>Any</b>
	<b>T4b</b>	<b>Any</b>	<b>M0</b>	<b>Any</b>	<b>Any</b>
	<b>Any</b>	<b>N3</b>	<b>M0</b>	<b>Any</b>	<b>Any</b>
<b>IV</b>	<b>Any</b>	<b>Any</b>	<b>M1</b>	<b>Any</b>	<b>Any</b>