高雄榮民總醫院 甲狀腺癌臨床診療指引

2022 年第一版

癌症中心甲狀腺癌醫療團隊擬定

2019年第一版為參考美國癌症聯合委員會於2018年3月出版之甲狀腺癌臨床指引、2015年美國甲狀腺協會治療指引及其他參考文獻,於2022.02.10由甲狀腺癌團隊相關人員劉絮穎、梁宗榮、佘筠瑩、朱志勳、諶弘遠、邱宇莉、 李恒昇、許碩修、張國楨等討論後共同修訂。

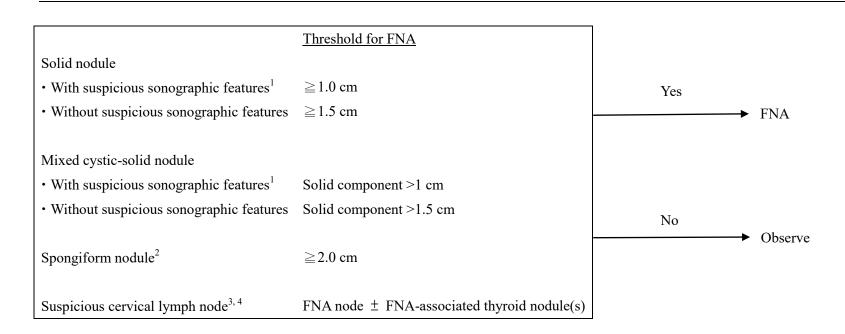
注意事項:這個診療準則主要作為醫師和其他保健專家診療癌症病人參考之用。

假如你是一個癌症病人,直接引用這個研究資訊及診療準則並不恰當。

只有你的醫師才能決定給你最恰當的治療。

上一版	新版
內容無放射碘 refractory 的治療方式	新增加放射碘 refractory 的治療方式

P. 03	甲狀腺結節評估
P. 04-05	甲狀腺細胞病理學報告
P. 06	乳突癌
P. 07	<u> 單葉切除術後發現之乳突癌</u>
P. 08	<u>濾泡癌</u>
P. 09	術後評估與治療
P. 10	美國甲狀腺協會危險分層系統
P. 11	促甲狀腺激素抑制原則
P. 12	術後放射線碘治療 (Initial Postoperative RAI therapy)
P. 13	放射線治療
P. 14	追蹤
P. 15	<u>復發</u>
P. 16	<u>髓質癌</u>
P. 17-18	分化不良癌
P. 19	化學治療 & 標靶治療處方
P. 20-21	参考 資料



¹Suspicious sonographic features: Hypoechoic, microcalcifications, infiltrative margins, and taller than wide in transverse plane. Sonographic features associated with a low risk of malignancy including spongiform nodules, isoechoic or hyperechoic solid nodules, and mixed solid-cystic nodules without any of the suspicious features listed above.

²Aggregation of multiple microcystic components in more than 50% of the volume of the nodule.

³Tg washout may be helpful in diagnosis of lymph node metastases.

⁴Suspicious lymph node features may include hypoechoic, rounded, absence of fatty hilum, cystic or partially cystic, and/or microcalcifications.

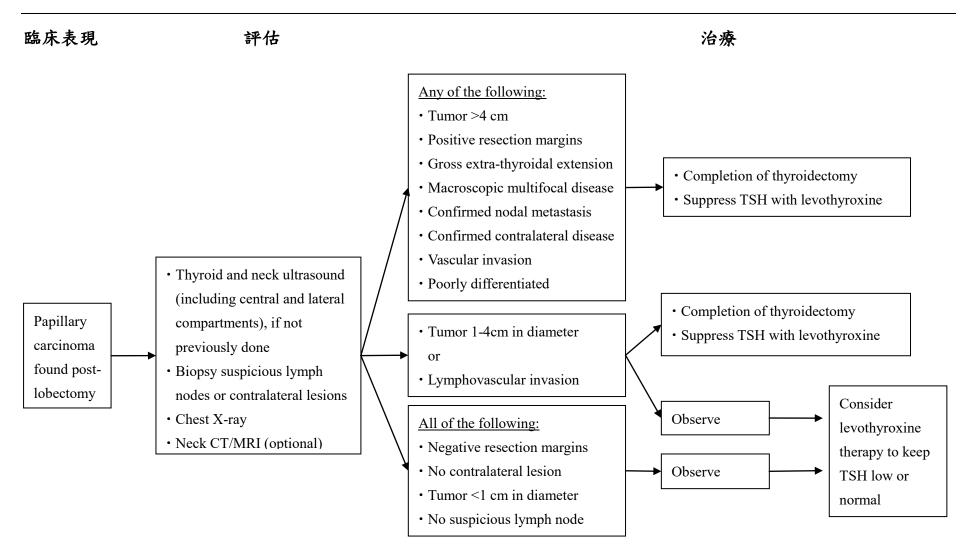
Be	nign form 良性的格式				
1.	Nondiagnostic				
	Specimen processed and examined, but nondiagnostic due to insufficient cellularity.				
2.	Benign				
	Presence of some benign-appearing follicular cells				
3.	BENIGN.				
	Consistent with lymphocytic (Hashimoto's) thyroiditis.				
	Numerous polymorphic lymphoid cells and scattered follicular cells. Lymphocytic (Hashimoto's) thyroiditis is considered. Suggest clinical				
	correlation.				
At	ypia 不確定的格式				
4.	Atypia of Undetermined Significance, favor reactive				
	Presence of groups of proliferative benign-looking follicular cells.				
	Suggest follow-up and repeat FNA cytology in the next visit.				
5. Atypia of Undetermined Significance, neoplasm cannot be excluded					
	Presence of groups/sheets of proliferating follicular cells with obvious nucleoli/ with focal papillary arrangement, the possibility of malignancy cannot				
	be excluded totally.				
	Suggest further evaluation and either repeat FNA cytology 3 months later or do biopsy if clinically indicated.				
Su	spicious for a Follicular Neoplasm. 針對濾泡型腫瘤				
6.	Suspicious for a Follicular Neoplasm.				
	Cellular aspirate of follicular cells with a predominantly microfollicular architecture, scattered isolated cells, and scant colloid.				

Sus	Suspicious for malignancy 疑似惡性			
7.	Suspicious for malignancy			
	Presence of some sheets of proliferating follicular cells, of which some having intranuclear pseudoinclusions. Suspicious for papillary carcinoma.			
	Presence of some atypical follicular cells with obvious nucleoli and a few pseudo-inclusion nuclei, of which the possibility of papillary carcinoma			
	should be considered. Suggest further evaluation and do biopsy.			
8.	Suspicious for malignancy, particularly medullary carcinoma cannot be excluded.			
	Presence of some small groups of small hyperchromatic cells, of which the possibility of malignancy, particularly medullary carcinoma cannot be			
	excluded.			
	Suggest do biopsy if clinically indicated.			
Ma	lignant 確定惡性			
9.	Positive for malignant cells.			
	The sections show groups of atypical follicular cells with obvious pseudo-inclusions, which is consistent with papillary carcinoma.			

細針抽吸結果 評估	手術方式考量	治療
Palliary Carciinoma or Atypia of undetermined significance (AUS) ¹ • Thyroid and neck ultrasound (including central and lateral compartments), if not previously done • CT/MRI for fixed, bulky, or substernal lesions • Chest X-ray • Consider evaluation of vocal cord mobility	Indications for total thyroidectomy (any present): • Known distant metastases • Bilateral nodularity • Extrathyroidal extension • Tumor >4 cm in diameter • Cervical lymph node metastases • Poorly differentiated • Consider for radiation history Indications for total thyroidectomy <u>or</u> lobectomy, If all present: • No prior radiation • No distant metastases • No cervical lymph node metastases • No extrathyroidal extension • Tumor <4 cm in diameter	 Total thyroidectomy Perform neck dissection of involved compartments for clinical apparent/biopsy-proven disease Suppress TSH with levothyroxine Total thyroidectomy + TSH suppression with levothyroxine or Lobectomy + isthmusectomy (consider TSH suppression)

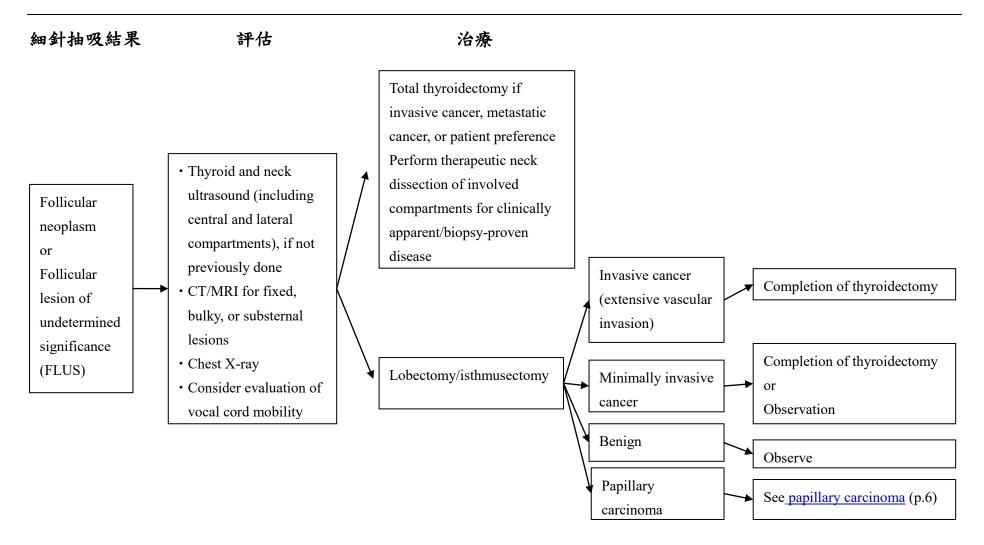
乳突癌 (Papillary Carcinoma)

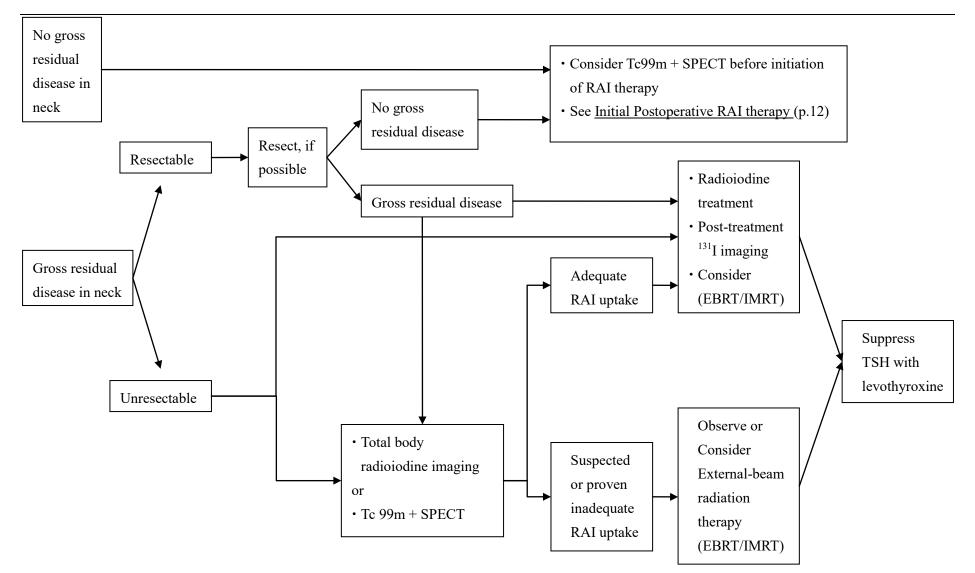
¹AUS with high clinical suspicion of malignancy may consider lobectomy or total thyroidectomy for definitive diagnosis/treatment.



單葉切除術後發現之乳突癌(Papillary carcinoma found post-lobectomy)

濾泡癌(Follicular Carcinoma)





術後評估與治療(Postsurgical Evaluation & Treatment)

美國甲狀腺協會危險分層系統(ATA risk stratification system with Proposed Modifications)

1				
	Papillary thyroid cancer (with all of the following):			
	• No local or distant metastases;			
	All macroscopic tumor has been resected			
	No tumor invasion of loco-regional tissues or structures			
任日吟	• The tumor does not have aggressive histology (e.g., tall cell, hobnail variant, columnar cell carcinoma)			
低風險	• If 131-I is given, there are no RAI-avid metastatic foci outside the thyroid bed on the first posttreatment whole-body RAI scan			
Low risk	No vascular invasion			
	• Clinical N0 or ≤ 5 pathologic N1 micrometastases (<0.2 cm in largest dimension) ^a			
	Intrathyroidal, encapsulated follicular variant of papillary thyroid cancer ^a			
	Intrathyroidal, well differentiated follicular thyroid cancer with capsular invasion and no or minimal (<4 foci) vascular invasion ^a			
	Intrathyroidal, papillary microcarcinoma, unifocal or multifocal, including BRAF ^{V600E} mutated (if known) ^a			
	Microscopic invasion of tumor into the perithyroidal soft tissues			
	RAI-avid metastatic foci in the neck on the first posttreatment whole-body RAI scan			
中度風險	Aggressive histology (e.g., tall cell, hobnail variant, columnar cell carcinoma)			
Intermediate risk	Papillary thyroid cancer with vascular invasion			
	Clinical N1 or >5 pathologic N1 with all involved lymph nodes <3 cm in largest dimension ^a			
	Multifocal papillary microcarcinoma with ETE and BRAF ^{V600E} mutated (if known) ^a			
	Macroscopic invasion of tumor into the perithyroidal soft tissues (gross ETE) 腫瘤沒有完整切除			
	Incomplete tumor resection			
高風險	Distant metastases ^a Proposed modifications, not present in the			
High risk	Postoperative serum thyroglobulin suggestive of distant metastases original 2009 initial risk stratification system			
	Pathologic N1 with any metastatic lymph node \$3 cm in largest dimension ^a			
	Follicular thyroid cancer with extensive vascular invasion (> 4 foci of vascular invasion) ^a			

促甲狀腺激素抑制原則[Principles of Thyroid-Stimulating Hormone (TSH) Suppression]

- Because TSH is a trophic hormone that can stimulate the growth of cells derived from thyroid follicular epithelium, the use of levothyroxine to maintain low TSH levels is considered optimal in treatment of patients with papillary, follicular, or Hurthle cell arcinoma. However, data are lacking to permit precise specification of the appropriate serum levels of TSH.
- Given the potential toxicities associated with TSH-suppressive doses of levothyroxine—including cardiac tachyarrhythmias (especially in the elderly) and bone demineralization (particularly in post-menopausal women) as well as frank symptoms fo thyrotoxicosis—the risk and benefit of TSH-suppressive therapy must be balanced for each individual patients.
- Patients whose TSH levels are chronically suppressed should be counseled to ensure adequate daily intake of calcium and vitamin D.

術後放射線碘治療 (Initial Postoperative RAI therapy)

Papillary Carcinoma	Follicular carcinoma
RAI not tpyically recommended (if all present) :	RAI not tpyically recommended (if all present) :
Classic papillary thyroid carcinoma (PTC)	• Primary tumor <2 cm
• Primary tumor < 2 cm	• Intrathyroidal
• Intrathyroidal	No vascular invasion
• Unifocal or multifocal	• Clinical N0
RAI selectively recommended (if any present):	RAI selectively recommended (if any present):
Primary tumor 2-4cm	Primary tumor 2-4cm
High-risk histology	Minor vascular invasion
Lymphovascular invasion	Cervical lymph node metastases
Cervical lymph node metastases	RAI typically recommended (if any present):
• Macroscopic multifocality (one focus >1 cm)	Gross extrathyroidal extension
RAI typically recommended (if any present):	• Primary tumor >4 cm
Gross extrathyroidal extension	Extensive vascular invasion
• Primary tumor >4 cm	Known or suspected distant metastases at presentation
Known or suspected distant metastases at presentation	(if amenable to RAI)
(if amenable to RAI)	

Recommended Dosage :

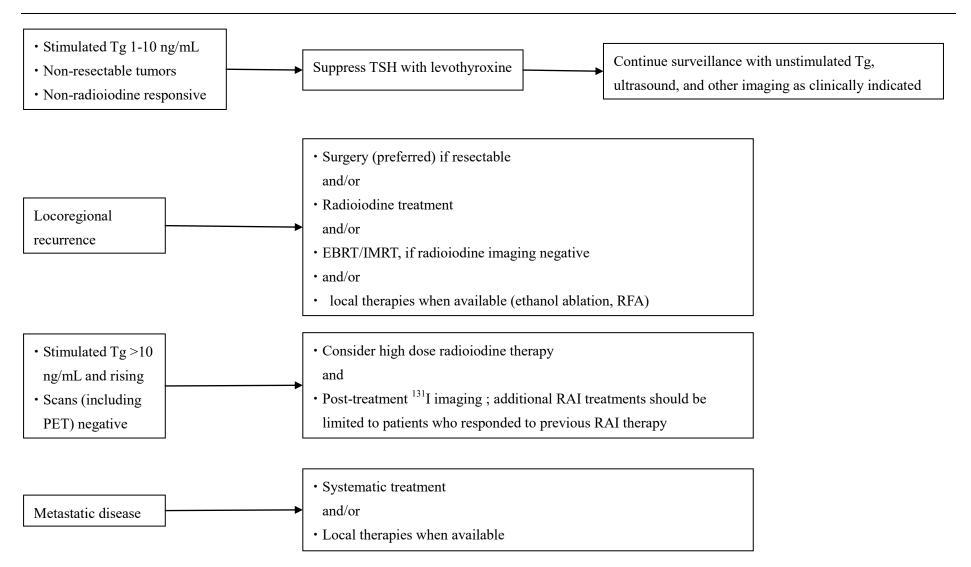
- RAI not typically recommended: 0-30 mCi
- RAI selectively recommended: 30-150 mCi
- RAI typically recommended: 100-200 mCi

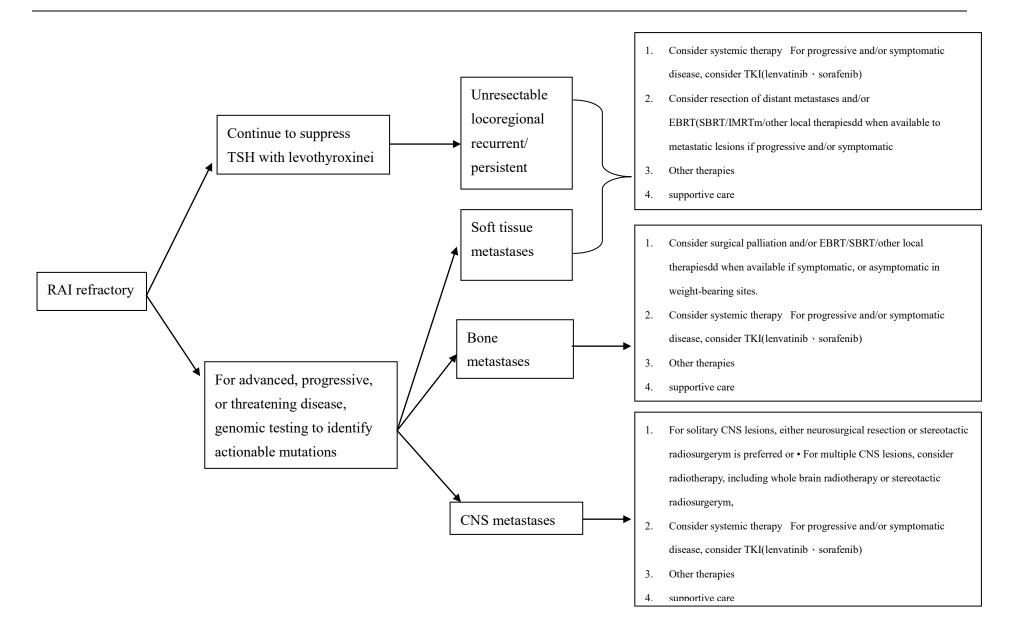
放射線治療 (External Beam Radiation Therapy/Intensity Modulated Radiation Therapy)

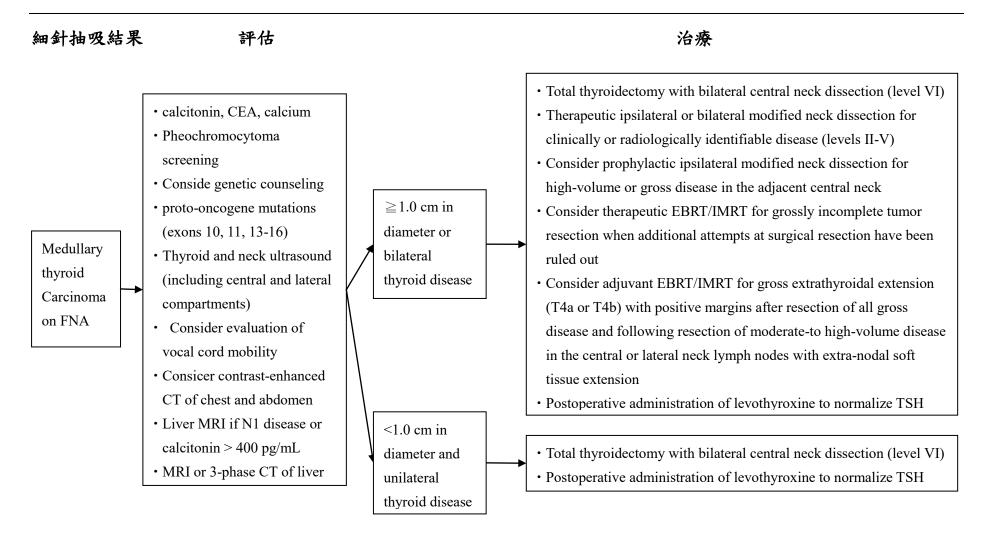
- There is no major prospective, randomized trials in external beam radiotherapy but some retrospective studies showed beneficial in local control or cancer-specific survival.
- Indication of considering R/T
 - gross residual tumor/unresectable tumor, esp in no/poor RAI uptake tumor
 - extrathyroidal extension or extranodal extension.
- Dosage:
 - low-risk : 54 Gy,
 - microscopic disease: 59.4–63 Gy,
 - positive margins: 63–66 Gy,
 - macroscopic disease: 66–70 Gy.

- Physical examination, TSH and Tg measurement + antithyroglobulin antibodies
- Neck ultrasound every 6-12 months
- Chest X-ray every 6-12 months
- For patients underwent total thyroidectomy and postoperative RAI therapy, consider radioiodine imaging in the first year postoperatively
- Consider TSH-stimulated radioiodine imaging in high-risk patients, patients with previous RAI-avid metastases, or patients with abnormal Tg levels (either TSH-suppressed or TSH-stimulated), rising antithyroglobulin antibodies, or abnormal ultrasound during surveillance.
- Consider FDG-PET/CT for patients with abnormal Tg level but negative ¹³¹I imaging.

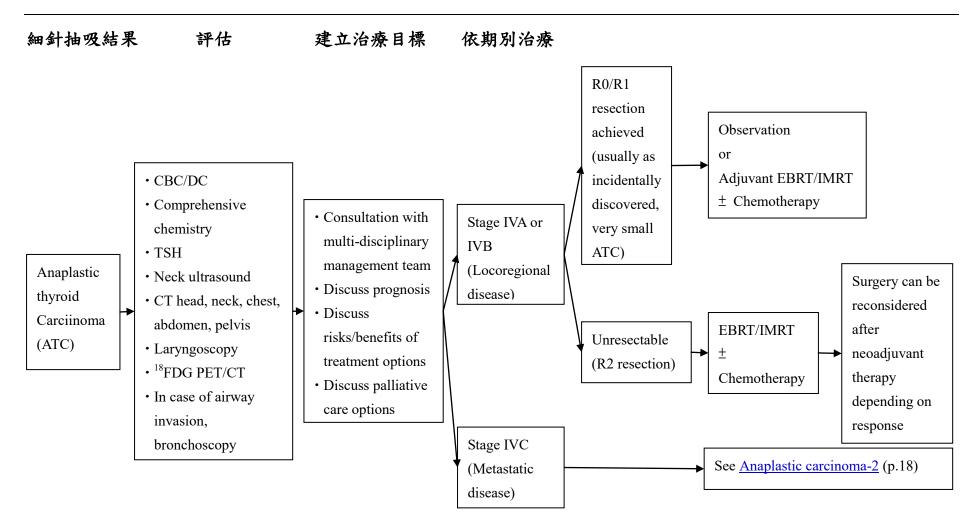
復發 (Recurrent Disease)

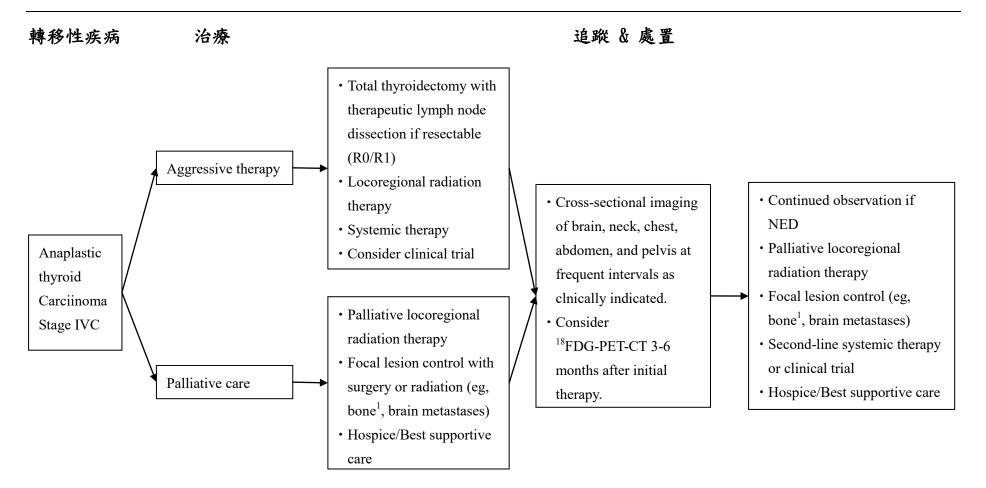






分化不良癌 (Anaplastic Carcinoma) -1





¹Consider use of intravenous bisphosphonates or denosumab. Denosumab and intravenous bisphosphonates can be associated with severe hypocalcemia; patients with hypoparathyroidism and vitamin D deficiency are at increase risk.

Metastasis 第一線治療

1. RAI therapy

2. 標靶治療處方 (Kinase Inhibitor Therapy or metastasis)

Regimen	Agents/Dosages	Frequency
Sorafenib (Nexavar)	400mg Oral	Twice daily (BID)
Lenvatinib(Lenvima)	24mg Oral	Once daily (QD)

3. 化學治療處方 (Chemotherapy Regimen or metastasis)

Regimen	Agents/Dosages	Frequency
Paclitaxel (weekly)	60-90 mg/m ² IV	Weekly
Doxorubicin (weekly)	20 mg/ m ² IV	Weekly
Docetaxel (3-week)	60 mg/ m ² IV	<u>(3-week)</u>
Docetaxel + Carboplatin	75 mg/ m ² IV+Auc x 5 mg/CCR	<u>(3-week)</u>
<u>(CCr < 60)</u>		
Docetaxel + Cisplatin	$75 \text{ mg/m}^2 \text{ IV} + 60 \text{ mg/m}^2 \text{ IV}$	<u>(3-week)</u>
(3-week cycle)		

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