

高雄榮民總醫院

鼻咽癌診療原則

2022年03月02日第一版

鼻咽癌醫療團隊擬訂

注意事項：這個診療原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，直接引用這個診療原則並不恰當，只有你的醫師才能決定給你最恰當的治療。

會議討論

上次會議：2021/04/28

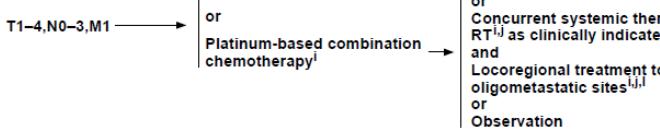
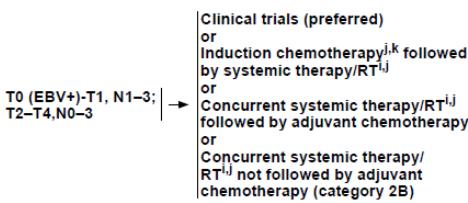
本共識與上一版的差異

上一版	新版
<ol style="list-style-type: none">Workup 中的 Multidisciplinary consultation 加上 會診項目 Fertility/reproductive, smoking cessation, ophthalmologic and endocrine evaluation if indicatedClinical staging 增加 T0 (EBV+)[Clinical T0 (EBV+)- T1, N1-3 or T2-4, any N, M0] 的部分，經過治療後如果 CR，刪除 adjuvant C/T[Clinical T0 (EBV+)- T1, N1-3 or T2-4, any N, M0] Primary treatment 的部分，將 induction CT + CCRT or RT 的優先順序移至 CCRT ± Adjuvant CT 之前。	<ol style="list-style-type: none">治療方面根據 EBV-associated NPC 討論，將 [Clinical T0 (EBV+)- T1, N1-3 or T2-4, any N, M0] 的部分調整成： (1) [Clinical T2, N0, M0] 做 Definitive RT ± concurrent systemic therapy if high risk features (2) [Clinical T1-2, N1, M0 or T3, N0] 做 concurrent systemic therapy/RT 優先，除非 high risk features 才考慮 induction or adjuvant chemotherapy。 (3) [Clinical T3-4, N1-3, M0 or any T, N2-3, M0] 做 induction chemotherapy followed by concurrent systemic therapy/RT 優先 (category 1)；然後是 concurrent systemic therapy/RT followed by adjuvant chemotherapy；純粹 CCRT 排最後 (category 3)。遠端轉移 M1 治療特別將 oligometastatic disease、widely metastasis with good PS (0-2)、widely metastasis with poor PS (3-4) 分開討論。Recurrent, Unresectable, Metastatic Disease 增加治療 Cisplatin/Gemcitabine + PD-1 inhibitor (eg, Pembrolizumab or Nivolumab)。

CLINICAL STAGING

TREATMENT OF PRIMARY AND NECK

FOLLOW-UP



NCCN Guidelines Version 1.2022
Cancer of the Nasopharynx

ⁱ See Principles of Radiation Therapy (NASO-A).
^j See Systemic Therapy for Nasopharyngeal Cancers (NASO-B).

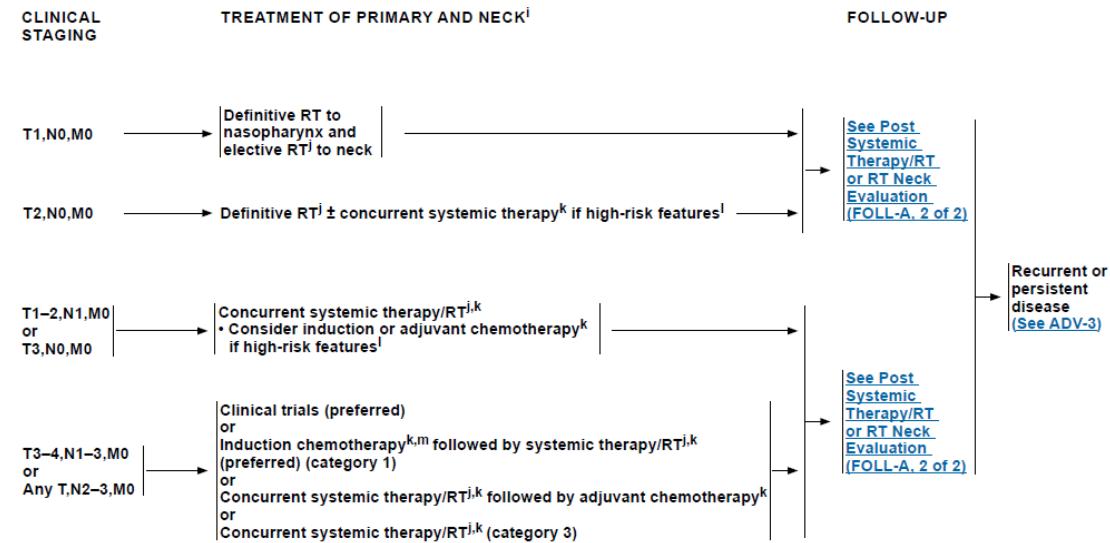
^k See Discussion on induction chemotherapy.
^l Can be used for select patients with distant metastases or for patients with symptoms in the primary or a

Note: All recommendations are category 2A unless otherwise indicated.
Clinical Trials: NCCN believes that the best management of any patient with cancer is in a clinical trial. Participation in clinical trials is especially encouraged.

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Compared 2022 to 2021

1. T2N0 降階為definitive RT or CCRT
2. T1-2, N1, M0 or T3, N0, M0 首選 CCRT
3. Advanced stage 建議IC+ CCRT (category 1); CCRT only (category 3)



ⁱ The recommendations are based on clinical trial data for those with EBV-associated nasopharynx cancer.

^j See Principles of Radiation Therapy (NASO-A).

^k See Systemic Therapy for Nasopharyngeal Cancers (NASO-B).

^l High risk features include bulky tumor volume, high serum EBV DNA copy number.

^m See Discussion on induction chemotherapy.

Note: All recommendations are category 2A unless otherwise indicated.
Clinical Trials: NCCN believes that the best management of any patient with cancer is in a clinical trial. Participation in clinical trials is especially encouraged.

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Nasopharyngeal cancer

Clinical staging AJCC 8th

Nasopharyngeal cancer TNM staging AJCC UICC 8th edition

Primary tumor (T)		Prognostic stage groups			
T category	T criteria	Then T is...	And N is...	And M is...	Then the stage group is...
TX	Primary tumor cannot be assessed	Tis	N0	M0	0
T0	No tumor identified, but EBV-positive cervical node(s) involvement	T1	N0	M0	I
Tis	Tumor <i>in situ</i>	T1, T0	N1	M0	II
T1	Tumor confined to nasopharynx, or extension to oropharynx and/or nasal cavity without parapharyngeal involvement	T2	N0	M0	II
T2	Tumor with extension to parapharyngeal space, and/or adjacent soft tissue involvement (medial pterygoid, lateral pterygoid, prevertebral muscles)	T2	N1	M0	III
T3	Tumor with infiltration of bony structures at skull base, cervical vertebra, pterygoid structures, and/or paranasal sinuses	T1, T0	N2	M0	III
T4	Tumor with intracranial extension, involvement of cranial nerves, hypopharynx, orbit, parotid gland, and/or extensive soft tissue infiltration beyond the lateral surface of the lateral pterygoid muscle	T2	N2	M0	III
		T3	N0	M0	IVA
		T3	N1	M0	IVB
		T4	N0	M0	IVB
		T4	N1	M0	IVB
		T4	N2	M0	IVB
		Any T	N3	M0	IVB
		Any T	Any N	M1	IVB

tumor, node, metastasis; AJCC: American Joint Committee on Cancer; UICC: Union for International Cancer Control; EBV: Epstein-Barr virus.

with permission of the American College of Surgeons, Chicago, Illinois. The original source for this information is the AJCC Cancer Staging Manual, Eighth Edition (2017) published by Springer International Publishing. Corrected at 4th printing, 2018.

UpToDate®

Carcinoma of Nasopharynx

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WORK-UP

- History & PE & NP scopy
 - NP biopsy ± Neck FNA
 - Image
 - MRI* or CT* of H&N or PET/CT
 - Chest X-ray * (if PET/CT not done)
 - Bone scan * (if PET/CT not done)
 - Abd. Sono *
 - ± PET scan ± Chest CT
 - EBV status: viral load, ± EB-EA/NA, ± EB-VCA IgG/IgA
 - Dental evaluation*
 - Panorex ± teeth extraction
 - Hearing evaluation
 - PTA, tympanogram
 - Multidisciplinary consultation
 - (± Fertility/reproductive, smoking cessation, ophthalmologic and endocrine evaluation if indicated)
- (* 期別之相關之主要檢查)

STAGING & TREATMENT

- [cT1,N0,M0]
詳見 Page 2
- [cT2,N0,M0]
詳見 Page 2
- [cT1-2, N1, M0 or cT3, N0, M0]
詳見 Page 3
- [T3, N1, M0 or T4, N0-1, M0, Any T, N2-3, M0]
詳見 Page 4
- [M1]
詳見 Page 5

FOLLOW-UP

- [Post-Tx within 6 months]
 - Post-Tx baseline MRI and/or CT, EBV viral load,
 - Every 2-3 months: PE, NP scopy± Neck Sono
 - [0.5-3 years]
 - Every 3-4 months: PE, NP scopy+/- EBV viral load,
 - Every 1 yr: ± EB-EA/NA ± EB-VCA IgG/IgA, MRI and/or CT, CxR, WBBS & Abd. Sono as indicated, ±TSH, free T4*
 - [3-5 years] → Every 4-6 months: PE, NP scopy
 - [5 years later]
 - Every 6-12 months: PE, NP scopy
- (*if RT, 6-12 months)

Carcinoma of Nasopharynx

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Clinical T1,N0,M0

Primary treatment

Definitive RT to nasopharynx
and elective RT to neck

Follow-up

Clinical T2,N0,M0

Primary treatment

Definitive RT ± concurrent
systemic therapy if high-risk
features @

Follow-up

@Bulky tumor volume, high serum EBV DNA copy number

Carcinoma of Nasopharynx

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Clinical T1-2, N1, M0 or T3, N0, M0

Primary treatment

CCRT ± induction or adjuvant CT 註

1-3 if high risk features @

~~2-3 cycles for locoregional advanced disease~~ 若只打1 cycle 且與後續CCRT間隔小於 2 weeks，視為CCRT only。

Response and salvage treatment

Complete clinical response

Follow-up

Residual disease
or clinically
suspicious residue

Surgery if
operable* #

Adjuvant CT
註3

@Bulky tumor volume, high serum EBV DNA copy number

Salvage neck dissection is indicated if residual neck disease.

* Salvage nasopharyngectomy is indicated for operable residual primary tumor

Carcinoma of Nasopharynx

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Clinical T3, N1, M0 or T4, N0-1, M0, Any T, N2-3, M0

Primary treatment

Clinical trials

Induction CT + CCRT 註1-3

~~2-3 cycles for locoregional advanced disease (cT4 or cN3 or 視病情需求, 如: EBV titer > 1000, bulky T3, advanced N2\$)~~; 若只打1 cycle 且與後續CCRT間隔小於 2 weeks, 視為CCRT only。

CCRT ± Adjuvant CT 註1-3

High risk for distant failure (ex. cT4 or cN3 or 視病情需求)
建議加打 2-3 cycles of adjuvant CT。

Definitive RT 註1

Poor medical condition or patient's preference。

Response and salvage treatment

Complete clinical response

Follow-up

Residual disease or clinically suspicious residue

Surgery if operable*

Adjuvant CT 註3

Salvage neck dissection is indicated if residual neck disease.

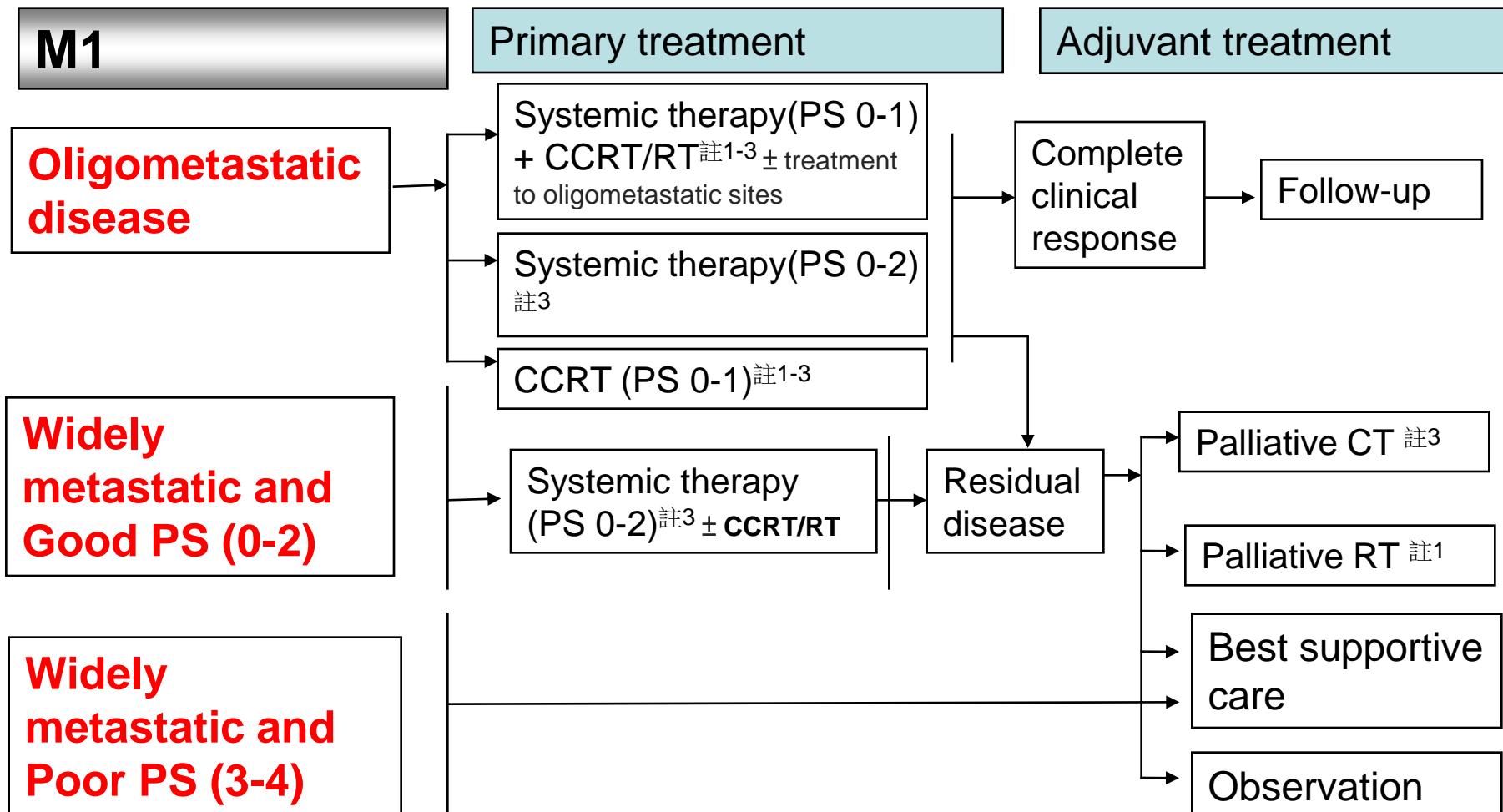
* Salvage nasopharyngectomy is indicated for operable residual primary tumor.

\$ Diffuse LAP near the cricoid cartilage, big LAP ≥ 5 cm (2020/07/08 團隊會議增訂)

@ N3, T3-4N1-2, or stage IV (2020/07/08 團隊會議增訂)

Carcinoma of Nasopharynx

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Carcinoma of Nasopharynx

註1 高雄榮民總醫院 臨床診療指引 | Ver.1 修訂於2022.03.02 Page 6 (Ref. 1,5,6)

Principles of Radiotherapy

Definitive Radiotherapy

- Primary and gross adenopathy : 66 - 74 Gy (2.0-2.2 Gy/fraction)
- Neck uninvolved nodal stations : 44 - 58 Gy (1.6-2.0 Gy/fractions)
- Suspicious Neck lymph nodes : 59.4 Gy (2.2 Gy/fractions) (optional)
- Adaptive radiotherapy : direct CCRT, BW change more than 3-5 kg, high initial stage etc.(optional)

CCRT or RT

- RT alone if : Old age, Impaired renal function, Poor condition

Palliative RT

- Indicated in : Relieve local symptoms, Prevent debilitation such as spinal cord compression and pathological fracture, Achieve durable loco-regional control.

Carcinoma of Nasopharynx

註2 高雄榮民總醫院 臨床診療指引 | Ver.1 修訂於2022.03.02 Page 7 (Ref. 1,5-9)

Principles of Chemotherapy

Concurrent with RT

Regimen 1: q3w CDDP ± Cetuximab + RT 註5

- Cisplatin (80-100mg/ m²) q3w during R/T
- Cetuximab(400mg/ m²) loading dose first week, then Cetuximab(250mg/ m²) maintain dose D1 + Cisplatin (80-100mg/ m²) q3w D2 during R/T

Regimen 2: Weekly CDDP ± Cetuximab + RT 註5

- Cisplatin (30-40mg/ m²) weekly during R/T
- Cetuximab(400mg/ m²) loading dose first week, and then Cisplatin (30-40mg/ m²) weekly D1 + Cetuximab(250mg/ m²) maintain dose D2 during R/T

Regimen 3: q3w Carboplatin ± Cetuximab + RT 註5

- Carboplatin (AUC x 5mg) q3w during R/T
- Cetuximab(400mg/ m²) loading dose first week, then Cetuximab(250mg/ m²) maintain dose D1 + Carboplatin (AUC x 5mg) q3w D2 during R/T

Regimen 4: Weekly Cetuximab + RT 註5

- Cetuximab(400mg/ m²) loading dose first week, then Cetuximab(250mg/ m²) maintain dose during RT

Carcinoma of Nasopharynx

註3 高雄榮民總醫院 臨床診療指引 | Ver.1 修訂於2022.03.02 Page 8 (Ref. 5-8)

Regimens of Chemotherapy

Induction or adjuvant, 建議2-3cycles

Regimen 1 : q3w G^{註5} P

- Gemcitabine (1000mg/ m²) D1, 8
- Cisplatin (80mg/ m²) D1

Regimen 2 : q3w G^{註5} Carboplatin

- Gemcitabine (1000mg/ m²) D1, 8
- Carboplatin (AUC x 5mg) D1

Regimen 3 : q3-4 weeks T + P ± F ± weekly Cetuximab^{註5}

- Taxotere(60 mg/ m²) D1 註5
- Cisplatin(60-75 mg/ m²) D1
- Fluorouracil (5-FU)(600-750mg/ m²) D2-D5
- Cetuximab (400mg/ m²) loading dose first week, then Cetuximab (250mg/ m²) maintain dose

Regimen 4 : q3-4 weeks T + Carboplatin ± F ± weekly Cetuximab^{註5}

- Taxotere(60 mg/ m²) D1 註5
- Carboplatin (AUC x 5mg) D1
- Fluorouracil (5-FU)(600-750mg/ m²) D2-D5
- Cetuximab (400mg/ m²) loading dose first week, then Cetuximab (250mg/ m²) maintain dose

Carcinoma of Nasopharynx

註3

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Regimens of Chemotherapy

Induction or adjuvant, 建議2-3cycles

Regimen 5: q3-4 weeks CDDP ± F ± weekly Cetuximab 註5

- Cisplatin(80-100mg/ m²) D1 or Cisplatin (20mg/ m²) D1-D5
- Fluorouracil (5-FU) (1000mg/ m²) D1-D5
- Cetuximab(400mg/ m²) loading dose first week, then weekly Cetuximab (250mg/ m²)

Regimen 6: q3-4 weeks Carboplatin ± F ± weekly Cetuximab 註5

- Carboplatin (AUC x 5mg) D1
- Fluorouracil (5-FU) (1000mg/ m²) D2-D5
- Cetuximab(400mg/ m²) loading dose first wk, then weekly Cetuximab (250mg/ m²)

Regimen 7: oral Fluorouracil

- Ufur cap (tegafur 100mg+uracil 224mg) 2# BID-TID

(作為取代 IV form 5-FU之替代藥物)

Carcinoma of Nasopharynx

註4

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Regimens of Chemotherapy

Recurrent or metastatic disease

Regimen 1 (First line): q3w G^{註5} ± P

- Gemcitabine (1000mg/ m²) D1, 8
- Cisplatin (80mg/ m²) D1

Regimen 2: q4w GGG^{註5} ± P

- Gemcitabine (1000mg/ m²) D1, 8, 15
- Cisplatin (50-60mg/ m²) D22

Regimen 3: q3w G^{註5} ± Carboplatin

- Gemcitabine (1000mg/ m²) D1, 8
- Carboplatin (AUC x 5mg) D1

Regimen 4: q3-4 weeks P ± F

- Cisplatin(80-100mg/ m²) D1 or Cisplatin (20mg/ m²) D1-D5
- Fluorouracil (5-FU) (600-1000 mg/m²) D2-D5

Carcinoma of Nasopharynx

註4

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Regimens of Chemotherapy

Recurrent or metastatic disease

Regimen 5: q3-4 weeks Carboplatin ± F

- Carboplatin (AUC x 5mg) D1
- Fluorouracil (5-FU) (1000mg/ m²) D2-D5

Regimen 6: q3-4 weeks T ± P

- Taxotere(60 mg/ m²) D1 註5
- Cisplatin(60-75 mg/ m²) D1

Regimen 7: q3-4 weeks T ± Carboplatin

- Taxotere(60 mg/ m²) D1 註5
- Carboplatin (AUC x 5mg) D1

Regimen 8: q3-4 weeks Carboplatin ± weekly Cetuximab 註5

- Carboplatin (AUC x 5mg) D1
- Cetuximab(400mg/ m²) loading dose first week, then weekly Cetuximab (250mg/ m²)

Carcinoma of Nasopharynx

註4

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Regimens of Chemotherapy

Recurrent or metastatic disease

Regimen 9: q3w G ± P + Pembrolizumab / Nivolumab(q2w) ^{31, 32, 註5}

- Gemcitabine (1000mg/ m²) D1, 8
- Cisplatin (80mg/ m²) D1
- Pembrolizumab(200mg) D1 / Nivolumab(3mg/kg) D1

Regimen 10: weekly Methotrexate

- Methotrexate (40-60mg/ m²)

Regimen 11: q3 weeks Pembrolizumab

- Pembrolizumab(200mg) D1

Regimen 12: q2 weeks Nivolumab

- Nivolumab(3mg/kg) D1

Regimen 13: weekly Cetuximab ^{註5}

- Cetuximab (400mg/ m²) loading dose first week, then Cetuximab (250mg/m²) maintain dose

Carcinoma of Nasopharynx

註4

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Regimens of Chemotherapy

Recurrent or metastatic disease

Regimen 14: oral Fluorouracil

- Ufur cap (tegafur 100mg+uracil 224mg) 2# BID-TID
(作為取代 IV form 5-FU之替代藥物)

Regimen 15: FL

- Leucovorin (250 mg/ m²) D1
- Fluorouracil (5-FU) (2500 mg/ m²) D1

Regimen 16: P-FL

- Cisplatin (60mg/ m²) week 1, 3, 5, 7
- Fluorouracil (5-FU)(2500mg/ m²) + Leucovorin (250mg/ m²) mixed week 2, 4, 6, 8

Carcinoma of Nasopharynx

註5 高雄榮民總醫院 臨床診療指引 Ver.1 修訂於2022.03.02 Page 14

特殊用藥健保給付規定

Taxotere

- 頭頸部癌，限局部晚期且無遠端轉移之頭頸部鱗狀細胞癌且無法手術切除者。
- 與Cisplatin 及5-FU 併用，作為放射治療前的引導治療，限使用四個療程。

Cetuximab

- 限與放射線療法合併使用於局部晚期之口咽癌、下咽癌及喉癌患者，使用總療程以接受8 次輸注為上限。需經事前審查核准後使用。

符合下列條件之一：

- 1.年齡 \geq 70 歲
- 2.Ccr < 50ml/min
- 3.聽力障礙者 (聽力障礙定義為500Hz、1000Hz、2000Hz 平均聽力損失大於25 分貝)
- 4.無法耐受platinum-based 化學治療。

- 限無法接受局部治療之復發及/或轉移性頭頸部鱗狀細胞癌，且未曾申報 cetuximab 之病患使用。需經事前審查核准後使用，使用總療程以18週為限，每9週申請一次，需無疾病惡化情形方得繼續使用。(106/4/1)

Carboplatin

- 限腎功能不佳 (CCr < 60) 或曾作單側或以上腎切除之惡性腫瘤患者使用。

Carcinoma of Nasopharynx

註5

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特殊用藥健保給付規定

Pembrolizumab、Nivolumab

- 先前已使用過 platinum 類化學治療失敗後，又有疾病惡化的復發或轉移性頭頸部鱗狀細胞癌成人患者。本類藥品與 cetuximab 僅能擇一使用，且治療失敗時不可互換。
- 符合下列條件：
 - 病人身體狀況良好(ECOG≤1)
 - NYHA (the New York Heart Association) Functional Class I 或 II
 - GOT<60U/L及GPT<60U/L，且 T-bilirubin<1.5mg/dL；Creatinine<1.5mg/dL，且 eGFR>60mL/min/1.73m²
 - PD-L1 表現量 TPS≥50%
- 初次申請以12週為限，申請時需檢附以下資料：病理或細胞檢查報告、生物標記(PD-L1)表現量檢測報告、病人身體狀況良好(ECOG≤1)及心肺與肝腎功能之評估資料、符合 i-RECIST 定義之影像檢查及報告(上述影像檢查之給付範圍不包括PET)、先前已接受過之治療與完整用藥資料、使用免疫檢查點抑制劑之治療計畫(treatment protocol)。
- 用藥後每 12 週評估一次，以 i-RECIST 或 mRECIST 標準評定反應，依下列原則給付：
 - 有療效反應者(PR 及 CR)得繼續使用；
 - 出現疾病惡化(PD)或出現中、重度或危及生命之藥物不良反應時，應停止使用；
 - 疾病呈穩定狀態者(SD)，可持續再用藥 4 週，並於 4 週後再次評估，經再次評估若為 PR、CR 者，得再繼續使用 12 週。若仍為 SD 或已 PD 者，應停止使用。

Carcinoma of Nasopharynx

註5

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特殊用藥健保給付規定

Gemcitabine

限用於：

- 1.晚期或無法手術切除之非小細胞肺癌及胰臟癌病患。
- 2.晚期膀胱癌病患。
- 3.Gemcitabine與paclitaxel併用，可使用於曾經使用過anthracycline之局部復發且無法手術切除或轉移性之乳癌病患。
- 4.用於曾經使用含鉑類藥物(platinum-based) 治療後復發且間隔至少6個月之卵巢癌，作為第二線治療。
- 5.無法手術切除或晚期或復發之膽道癌(含肝內膽管)病患。

備註：頭頸癌與鼻咽癌目前無健保給付

Carcinoma of Nasopharynx

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