

國軍左營總醫院放射腫瘤科

2026 年口腔癌放射治療指引

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口腔癌放射治療指引與監測修正對照表

| 2025 | 2026 | 說明 |
|------|--|--------------------------------------|
| | 根治性放射治療技術之 二、無手術之放射治療新增 Reduce field boost 可考慮再做 CT scan simulation 評估療效，並做為 treatment plan 參考。 | Adaptive radiotherapy for H&N cancer |
| | 更新 references | 補充 |

放射治療適應症

一、根治性目的(curative intent)

1. T1-2N0 Definitive RT, T1-2N1: CCRT
2. 手術後之輔助性放射治療(adjuvant radiotherapy)
 - A. Inadequate margin
 - B. ECS (Extracapsular nodal spread)
 - C. LN(+) (N1 in level IV/V or N2, N3)
 - D. PNI (Perineural invasion)
 - E. LVP(Lymphovascular permeation)
 - F. T3 or T4
3. 可切除但有內科問題或開刀危險不適合手術(Resectable, but with poor medical problem or surgical risk)
4. 未產生遠端轉移之局部復發

二、緩解性目的(palliative intent)

1. 無法手術切除:T4b or Unresectable nodal disease
2. 有遠端轉移病灶
3. 併有遠端轉移且產生症狀之局部復發

根治性放射治療必要流程

一、治療計劃前完整的臨床評估

1. 確認期別、手術紀錄及病理報告，包括組織型態、腫瘤大小、惡性級數、開刀邊緣、有無神經旁侵犯、有無淋巴血管浸潤、

有無淋巴結轉移(包括數目/區域)、有無 ECS 等等。

2. 安排必要檢驗以排除有全身轉移之可能。
3. 經團隊會議討論及相關科別照會。
4. 必要時會做放射治療前的牙科會診及牙齒處置。

二、治療體位設定

1. 病人採仰臥，以頭頸模具固定，治療標記設定於模具及身體上。

三、模擬攝影

1. 病人依設定體位躺上電腦斷層攝影床，以金屬線進行必要標記(如手術疤痕、腫大之頸部淋巴、可疑腫塊)，視腫瘤位置及病患狀況需要時口中可含 cork 固定，並配合模具固定身體位置。
2. 通常電腦斷層掃描每切面為 2.5mm，掃描範圍應包含整個口腔腫瘤及頸部淋巴區域，通常至少包括從眼眶到鎖骨下緣。
3. 掃描後應以油性水洗不掉簽字筆作好標記供治療辨認。

四、治療計劃(treatment planning)

1. 臨床腫瘤體積(CTV: clinical target volume)

- A. 手術後之輔助性放射治療：CTV 包括原發腫瘤區(primary tumor bed)、侵犯淋巴部位/淋巴區以及潛在風險淋巴區。
- B. 無手術之放射治療：CTV 包括原發腫瘤部位、侵犯淋巴部位/淋巴區以及潛在風險淋巴區。
- C. 局部復發病人：CTV 包括復發部位。

2. 採用強度調控放射治療(IMRT)為治療方式，以減少危急器官放射劑量。

3. 治療計劃標靶體積(PTV: planning target volume)：PTV 依 CTV 增加 0.3 至 0.5 公分。鄰近腦幹處，考慮器官忍受劑量可為 0.1 公分。

4. 劑量評估參數：至少包括腦幹、脊髓、腮腺、顛下頷關節，當腫瘤位置較高時，尚要包括眼睛、視神經，視交叉等劑量。

五、放射治療前評估紀錄：包括病理報告、期別、核磁共振或電腦斷層攝影影像報告、病人簡史、理學檢查、重要檢查結果、診斷評估、體能狀態及治療計劃。

六、首次治療前應使用定位照相驗證片以確保照射範圍正確性，並由主治醫師確認簽章後才能進行。

根治性放射治療技術

一、手術後之輔助性放射治療

1. 原發腫瘤區(primary tumor bed) 至少給予 59.4Gy,侵犯淋巴部位(involved nodes)給予 60-66Gy ,未侵犯但有潛在風險淋巴區給予 45-63 Gy。每天一次，每週五次。每次 1.8-2Gy。
2. 輔助性放射治療建議於手術後三至四週後開始實施。
3. 建議分高，低危險區域做二階段不同劑量治療。

二、無手術之放射治療

1. 原發腫瘤部位(primary tumor) 及侵犯淋巴部位(involved nodes)給予 66-72Gy ,未侵犯但有潛在風險淋巴區給予 45-63Gy。每天一次，每週五次。每次 1.8-2Gy。合併化學治療時可考慮向下調整治療劑量。
2. 建議分高，中，低危險區域做三階段不同劑量治療。Reduce field boost 可考慮再做 CT scan simulation 評估療效，並做為 treatment plan 參考。

三、Example:

A. Postoperative RT:

Sequential: Two phases: deliver the initial phase (week 4-5) followed by high-dose boost volume phase (weeks 6-7) using 2 separate dose plans

CTV-H: primary tumor bed and involved nodal area:

59.4 Gy/33fr-68.4Gy/38fr

CTV-L: uninvolved nodal area :

41.4 Gy/23fr-50.4Gy/28fr

CTV-H: primary tumor bed with 1.0-1.5 cm margin /

involved nodal bed with 0.5-1.0 cm margin or involved nodal level

CTV-L: uninvolved nodal level

B. Definitive RT:

Sequential IMRT technique: deliver the initial phase (week 4-5) followed by moderate-dose boost volume phase (week 5-6.5), then gross tumor boost to high dose (week 6.5 to 8), usually using 3 separate dose plans

CTV-H: primary tumor with involved nodal area:

66.6Gy/37fx – 75.6Gy/42fx

CTV-M: primary tumor with 1.5-2.0 cm margin /

involved node with 1.0-1.5 cm margin or involved nodal level :

57.6 Gy/ 32 fx– 61.2 Gy/ 34 fx

CTV-L: uninvolved nodal area :

41.4Gy/23fx-50.4Gy/28fx

重要器官劑量評估參數

NOTE: All dose constraints below should be met whether the patient undergoes 3D-CRT or IMRT techniques.

Critical Normal Structures

Dose constraints are given below:

| Structure | true structure constraint | PRV constraint |
|----------------------|---|---------------------------------|
| Brainstem | 54 Gy max dose | no more than 1% to exceed 60 Gy |
| Spinal Cord | 45 Gy max dose | no more than 1% to exceed 50 Gy |
| Optic Nerves, Chiasm | 54Gy max dose | 54 Gy max dose |
| Mandible, TM joint | 70 Gy, if not possible then nomore than 1cc to exceed 75 Gy | |

Parotid glands: Mean dose <26 Gy (optimal) or 30 Gy(acceptable), should be achieved in at least one gland; or at least 50% of one gland will receive < 33 Gy (optimal) or 35Gy(acceptable) (should be achieved in at least one gland).

Other normal structurescan be considered:

| | |
|--------------------------------|--|
| Each cochlea | No more than 5% receives 55 Gy or more |
| Eyes | Max dose less than 45Gy |
| Lens | Max dose less than 10Gy |
| Glottic Larynx | Mean dose less than 40Gy |
| Esophagus, Postericoid pharynx | Mean dose less than 45 Gy |

根治性口腔癌放射治療常見之副作用及程度分級：

CTCAE v5.0

| Adverse event | Grade | | | | |
|---|---|---|---|--|-------|
| | 1 | 2 | 3 | 4 | 5 |
| Oral pain Definition: A disorder characterized by a sensation of marked discomfort in the mouth, tongue or lips | Mild pain | Moderate pain; limiting Instrumental ADL | Severe pain; limiting self care ADL | - | - |
| Mucositis oral Definition: A disorder characterized by inflammation of the oral mucosal | Asymptomatic or mild symptoms; intervention nor indicated | Moderate pain; not interfering with oral intake; modified diet indicated | Severe pain; interfering with oral intake | Life-threatening consequences; urgent intervention indicated | death |
| Dry mouth Definition: A disorder characterized by reduced salivary flow in the oral cavity | Symptomatic(e.g., dry or thick saliva) without significant dietary alteration; unstimulated saliva flow>0.2ml/min | Moderate symptoms; oral intake alteration(e.g., copious water, other lubricants, diet limited to purees and/or soft, moist foods); unstimulated saliva 0.1-0.2 ml/min | Inability to adequately aliment orally; tube feeding or TPN indicated; unstimulated saliva <0.1 ml/min | - | - |
| Dysphagia Definition: A disorder characterized by difficulty in swallowing | Symptomatic, able to eat regular diet | Symptomatic and altered eating/swallowing | Severely altered eating/swallowing; tube feeding or TPN or hospitalization indicated | Life-Threatening consequences; urgent intervention indicated | Death |
| Dermatitis radiation Definition: A finding of inflammatory reaction occurring as a result of exposure to biologically effective levels of ionizing radiation | Faint erythema or dry desquamation | Moderate to brisk erythema; patchy moist desquamation, mostly confined to skin folds and creases; moderate edema | Moist desquamation in areas other than skin folds and creases; bleeding induced by minor trauma or abrasion | Life-threatening consequences; skin necrosis or ulceration of full thickness dermis; spontaneous bleeding from involved site; skin graft indicated | Death |

根治性口腔癌放射治療可能副作用與處置：

一、急性副作用：

1. 口腔黏膜炎：常以溫開水漱口，嚴重時可請醫師處方漱口劑及藥膏。
2. 嗅覺味覺遲鈍：需配合調節食物口味，在治療後將漸漸恢復。
3. 皮膚炎：減少磨擦，嚴重時可請醫師處方藥膏。
4. 口乾：隨身攜帶水壺漱口或飲用。
5. 下巴、頸部淋巴水腫：嚴重時可請醫師處理或開處方藥物。
6. 短暫性脊髓病變：在治療後將漸漸恢復。
7. 在經濟條件許可下，治療期間可建議病患服用 Glutamine 降低黏膜副作用。

二、慢性副作用：

1. 口乾：隨身攜帶水壺漱口或飲用，嚴重時可請醫師處方藥物。
2. 蛀牙：保持口腔清潔，定期牙科門診防治。
3. 牙關緊閉：練習張口運動。
4. 頸部僵硬：經常做頸部柔軟運動及手部按摩。
5. 中耳炎及聽力減退：定期耳鼻喉科門診追蹤。
6. 少數零星個案且較嚴重的副作用，如腦組織壞死、視神經及視網膜病變、腦幹病變、腦下垂體功能低下、永久性脊髓病變、骨頭壞死、白內障、吞嚥困難、大量出血及中風等等：定期門診追蹤，嚴重時可考慮介入處置。

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2026 年口腔癌放射治療品質監測指標

1. 根治性口腔癌病人接受放射治療前，主治醫師對該療程進行確認及簽章比率:閾值:95%

分子定義：監測期間內，因口腔癌進行根治性放射治療，於接受放射治療前，主治醫師對病患療程進行確認及簽章之人數

分母定義：監測期間內，因口腔癌進行根治性放射治療總人數

2. 根治性口腔癌病人接受放射治療前，使用定位照相以確保照射範圍正確性之比率:閾值:95%

分子定義：監測期間內，因口腔癌進行根治性放射治療，於接受放射治療前，使用定位照相或影像導引以確保照射範圍正確性之人數

分母定義：監測期間內，因口腔癌進行根治性放射治療總人數

3. 根治性口腔癌病人接受放射治療時，劑量符合標準政策之比率:閾值:90%

分子定義：監測期間內，因口腔癌進行根治性放射治療，於療程完成時總劑量與標準劑量誤差為正負(含)10%以內之人數

分母定義：監測期間內，因口腔癌進行根治性放射治療總人數

4. 根治性口腔癌病人接受放射治療時，治療時間符合標準政策之比率:閾值:90%

分子定義：監測期間內，因口腔癌進行根治性放射治療，於療程完成時，總治療時間與標準治療時間誤差為正負(含)兩週以內之人數

分母定義：監測期間內，因口腔癌進行根治性放射治療總人數

5. 根治性口腔癌病人接受放射治療時，治療次數符合標準政策之比率:閾值:90%以上

分子定義：監測期間內，因口腔癌進行根治性放射治療，於療程完成時，實際次數與標準次數誤差為正負(含)10%以內之人數

分母定義：監測期間內，因口腔癌進行根治性放射治療總人數

6. 根治性口腔癌病人接受放射治療時，急性期非血液副作用出現第三級或以上之反應的比率:閾值:30%

分子定義：監測期間內，因口腔癌進行根治性放射治療；於療程完成時，急性期副作用出現第三級或以上之反應之人數

分母定義：監測期間內，因口腔癌進行根治性放射治療總人數