

# 高雄榮民總醫院

## 胰臟癌癌症診療指引

2023年03月22日 第一版

胰臟癌醫療團隊擬訂

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。

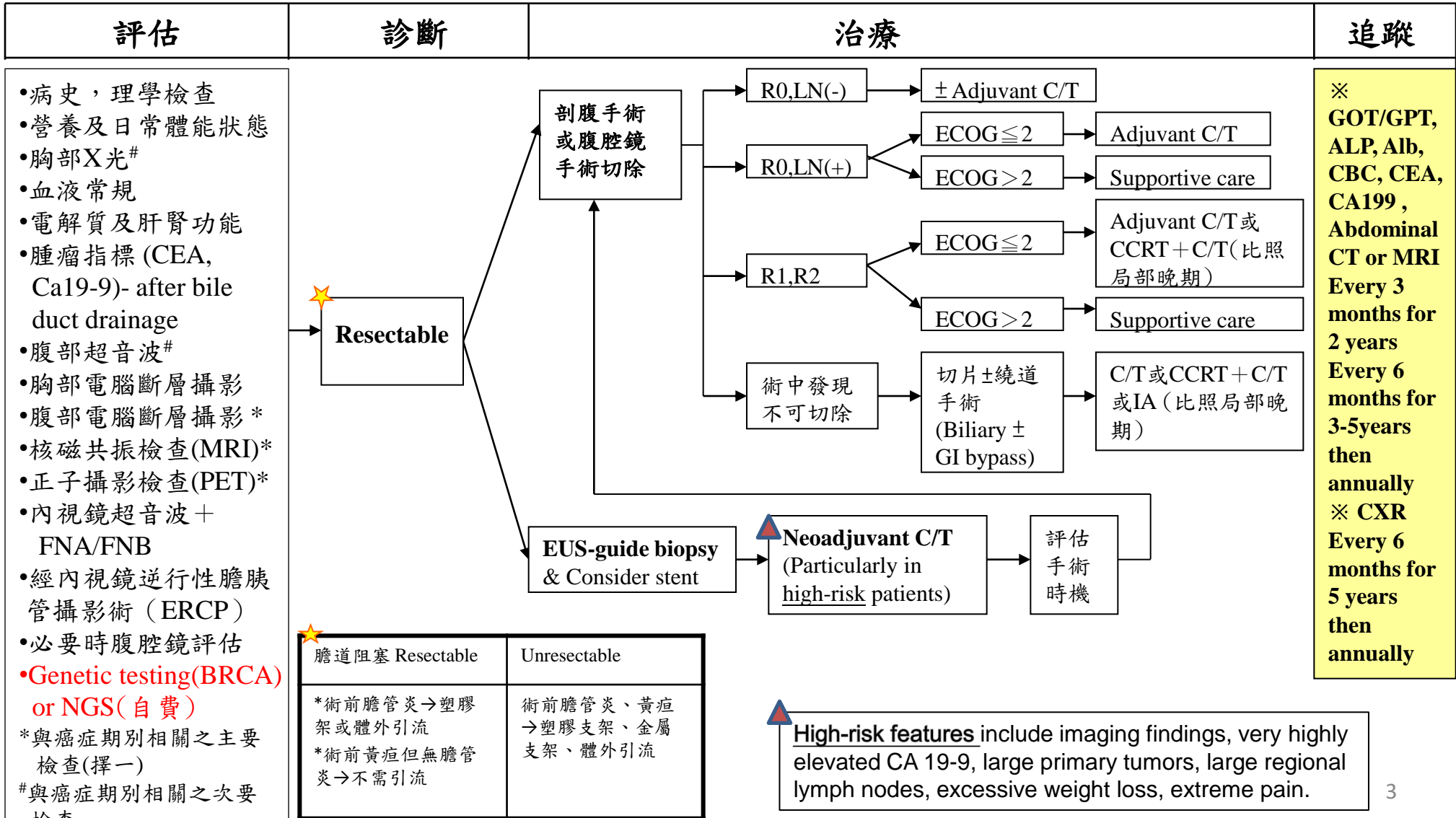
# 會議討論

上次會議：2022/03/22(第一版)

本共識與上一版的差異

上一版	新版
<p>1. Resectable: 術後 ECOG &gt; 2 分 「支持性治療」改「Supportive care」。(P.3)</p> <p>2. Metastatic disease:</p> <p>2-1. 「支持性治療」改「Palliative care」。(P.5)</p> <p>2-2. Metastatic disease: ECOG &gt; 2 分新增「Hospice」(P.5)</p> <p>3. 新增轉移癌的維持藥物: Olaparib(Lynpraza) (P.12)</p>	<p>1. 評估加上「Genetic testing(BRCA) or NGS(自費)」。(ppt.3-5)</p> <p>2. Borderline Resectable 修改診斷切片順序及 Neoadjuvant therapy 前如有黃疸症狀之說明。(ppt.4)</p> <p>3. Unresectable/Locally advanced 跟 Metastatic Disease (ppt.5)</p> <p>3-1. ECOG <math>\leq</math> 0-2 選項後修改診斷順序並新增「±繞道手術(Biliary ± GI bypass)」。</p> <p>3-2. 修改治療措辭。</p> <p>4. 新輔助化療處方新增「SLOG」。(ppt.8)</p> <p>5. 二線化療處方新增「GAS」。(ppt.14)</p> <p>6. 轉移癌維持處方修改項目名稱。(ppt.15)</p> <p>7. 放射治療 (ppt.17)</p> <p>7-1. 修改 indication 條件。</p> <p>7-2. 新增 CRT regimen: Capecitabine。</p>

# 胰臟腺癌



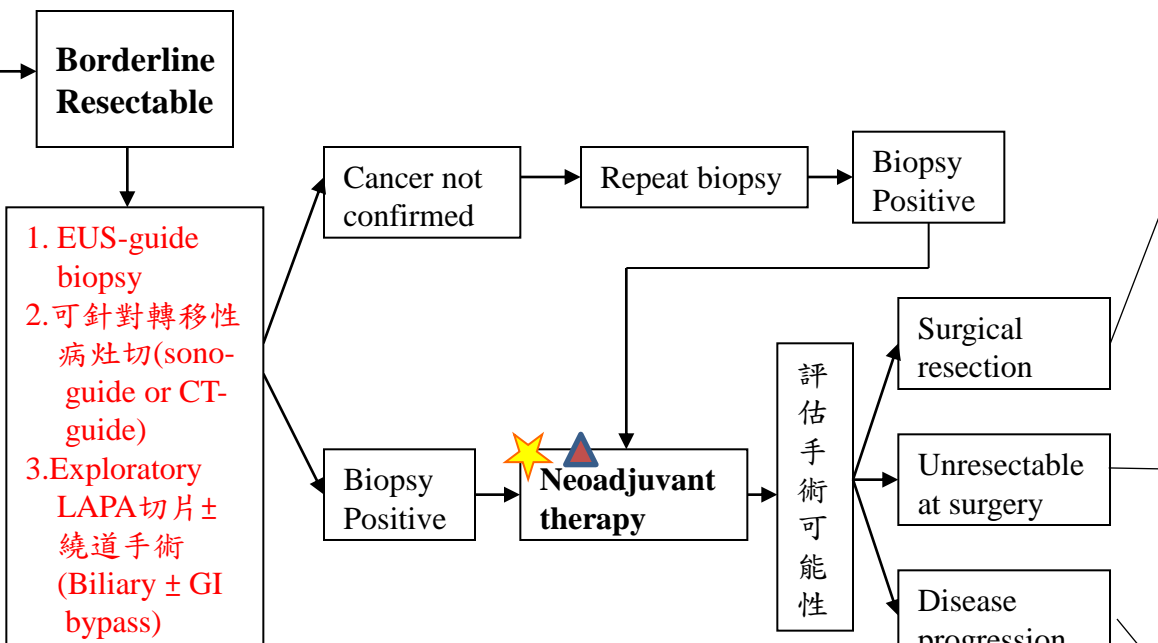
# 胰臟腺癌

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癌症診療指引

2023年第一版

評估	診斷	治療	追蹤
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- 病史，理學檢查
  - 營養及日常體能狀態
  - 胸部X光<sup>#</sup>
  - 血液常規
  - 電解質及肝腎功能
  - 腫瘤指標 (CEA, Ca19-9)- after bile duct drainage
  - 腹部超音波<sup>#</sup>
  - 胸部電腦斷層攝影
  - 腹部電腦斷層攝影\*
  - 核磁共振檢查(MRI)\*
  - 正子攝影檢查(PET)\*
  - 內視鏡超音波 + FNA/FNB
  - 經內視鏡逆行性膽胰管攝影術 (ERCP)
  - 必要時腹腔鏡評估
  - Genetic testing(BRCA) or NGS (自費)
- \*與癌症期別相關之主要檢查(擇一)
- #與癌症期別相關之次要檢查



★ Neoadjuvant 前如有阻塞性黃疸，需適當引流，包括金屬支架、塑膠支架、體外引流等

▲ High-risk features include imaging findings, very highly elevated CA 19-9, large primary tumors, large regional lymph nodes, excessive weight loss, extreme pain.

- ※ GOT/GPT, ALP, Alb, CBC, CEA, CA199, Every 6 months
- Abdominal CT or MRI Every 3 months for 2 years then annually
- ※ CXR Every 6 months for 5 years then annually

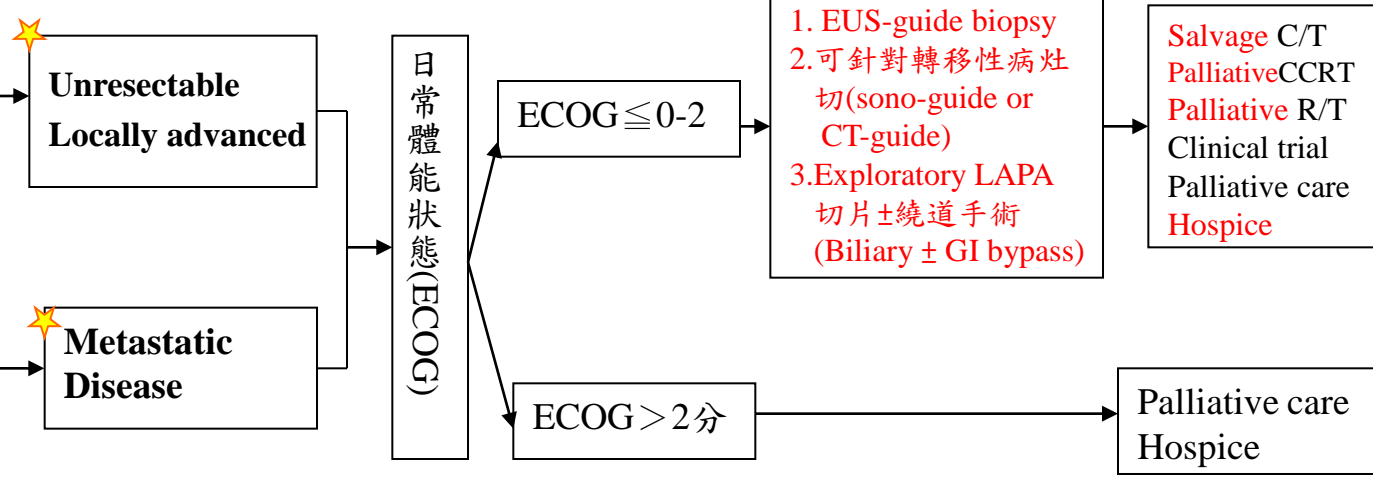
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- #與癌症期別相關之次要檢查



- ★ 如有黃疸症狀
  - 金屬支架
  - 塑膠支架
  - 體外引流

※ GOT/GPT, ALP, Alb, CBC, CEA, CA199, Abdominal CT or MRI  
 Every 3 months for 2 years  
 Every 6 months for 3-5 years then annually  
 ※ CXR  
 Every 6 months for 5 years then annually

## Criteria defining resectability status at diagnosis

Reference (No): 1

\*可手術切除 (MD-CT or MRI) :

- ① 無遠處轉移
- ② 上腸繫膜靜脈(SMV)或肝門靜脈(PV)完好
- ③ 腹腔動脈幹(celiac trunk)、肝動脈(HA)、上腸繫膜動脈(SMA)完好

\* **Borderline**可切除 :

- ① 無遠處轉移
- ② 上腸繫膜靜脈(SMV)或肝門靜脈(PV)可能被侵犯，但可手術切除部份血管並清除腫瘤
- ③ 胃十二指腸動脈(GDA)或肝動脈(HA)被侵犯，但可手術切除部份血管並清除腫瘤
- ④ 上腸繫膜動脈(SMA)完好，但未超過180°

\* 不可切除 :

胰臟頭部腫瘤

- ① 有遠處轉移
- ② 上腸繫膜動脈(SMA)被侵犯>180°，或celiac trunk被侵犯
- ③ 上腸繫膜靜脈(SMV)或肝門靜脈(PV)不可切除(無法重建血管) ④ 主動脈或下腔靜脈被侵犯

胰臟體部腫瘤

- ① 有遠處轉移
- ② 上腸繫膜動脈(SMA)被侵犯>180°
- ③ 上腸繫膜靜脈(SMV)或肝門靜脈(PV)不可切除(無法重建血管) ④ 主動脈被侵犯

胰臟尾部腫瘤

- ① 有遠處轉移 ② 上腸繫膜動脈(SMA)被侵犯>180° ③ 淋巴結轉移至切除範圍外

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## 化學治療處方建議表：新輔助化療-1

Chemotherapy for Neo-adjuvant (ECOG grade $\leq 2$ )	Schedule	Reference (No)/ strength of Evidence
<b>FOLFIRINOX</b> Oxaliplatin 85 mg/m <sup>2</sup> ,IV,2hrs Leukovorin 400 mg/m <sup>2</sup> ,IV,2hrs Irinotecan 180 mg/m <sup>2</sup> ,IV,90mins 5-FU 400 mg/m <sup>2</sup> ,IV bolus 5-FU 2400 mg/m <sup>2</sup> ,IV,46hrs	Q2W	NO.08/Level V
Cisplatin 50 mg/m <sup>2</sup> , IV,D1, D15 Gemcitabine 1000 mg/m <sup>2</sup> , IV,D1,D15	Q28 d	NO.17/Level V、 NO.22/Level V
<b>mFOLFIRINOX</b> Oxaliplatin 85 mg/m <sup>2</sup> ,IV,2hrs Leukovorin 400 mg/m <sup>2</sup> ,IV,2hrs Irinotecan 150 mg/m <sup>2</sup> ,IV,90mins 5-FU 2400 mg/m <sup>2</sup> ,IV,46hrs	Q2W	NO.01/Level I

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## 化學治療處方建議表:新輔助化療-2

<b>Chemotherapy for Neo-adjuvant (ECOG grade <math>\leq 2</math>)</b>	<b>Schedule</b>	<b>Reference (No)/ strength of Evidence</b>
<b>SLOG</b> Gemcitabine 800 mg/m <sup>2</sup> , IV, D1 Oxaliplatin 85 mg/m <sup>2</sup> ,IV,2hrs, D1 TS-1 35mg/m <sup>2</sup> /daily, BIDPC (Max daily dose 120mg), D1-D7 Calcium Folate Folic acid(15mg/tab) 20mg/m <sup>2</sup> /daily, BID, D1-D7	Q2W/cycle	NO.20 /Level V



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## 化學治療處方建議表：輔助化療

Adjuvant chemotherapy (R0切除) (ECOG grade $\leq 2$ )	Schedule	Reference (No)/ strength of Evidence
<b>TS-1*</b> 80-120mg/day (PO 4 weeks on, 2 weeks off or PO 2 weeks on, 1 weeks off) BSA $\geq 1.5\text{m}^2$ : 120mg /day, 1.25 $\text{m}^2$ - 1.5 $\text{m}^2$ : 100mg/day, <1.25 $\text{m}^2$ : 80mg/day	Q42 d /cycle x 4	NO.04/Level IB
<b>Gemcitabine*</b> 1000 mg/m <sup>2</sup> , IV,D1,D8,D15	Q28 d /cycle x 6	NO.05/Level IB NO.06 /Level IB
<b>5-FU/LV</b> Leucovorin 20mg/m <sup>2</sup> , IV bolus, and then 5-FU 425mg/m <sup>2</sup> , IV bolus, total 5 days	Q28 d/cycle x 6	NO.07/Level IB

\*健保用藥9.4.1：Gemcitabine限用於晚期或無法手術切除之非小細胞肺癌及胰臟癌病患。

\*健保用藥9.46：TS-1治療局部晚期無法手術切除或轉移性胰臟癌病人。

a. 若淋巴結陽性，符合「晚期」。可以開立健保給付之Gemcitabine與TS-1。

b. 若淋巴結陰性，不符合「晚期」。Gemcitabine與TS-1需用自費開立；或使用5-FU/LV則無給付之疑慮，但證據強度較Gemcitabine低。

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## 化學治療處方建議表:輔助化療

Adjuvant chemotherapy (R0切除) (ECOG grade $\leq 2$ )	Schedule	Reference (No)/ strength of Evidence
<b>mFOLFIRINOX</b> <b>Oxaliplatin</b> 85 mg/m <sup>2</sup> ,IV,2hrs <b>Leukovorin</b> 400 mg/m <sup>2</sup> ,IV,2hrs <b>Irinotecan</b> 150 mg/m <sup>2</sup> ,IV,90mins <b>5-FU</b> 2400 mg/m <sup>2</sup> ,IV,46hrs	Q2W /cycle x 12	NO.26/Level I

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## 化學治療處方建議表:局部晚期、轉移癌化療-1

Chemotherapy for unresectable、metastasis (ECOG grade $\leq 2$ )	Schedule	Reference (No)/ strength of Evidence
<b>FOLFIRINOX*</b> <b>Oxaliplatin</b> 85 mg/m <sup>2</sup> ,IV,2hrs <b>Leukovorin</b> 400 mg/m <sup>2</sup> ,IV,2hrs <b>Irinotecan</b> 180 mg/m <sup>2</sup> ,IV,90mins <b>5-FU</b> 400 mg/m <sup>2</sup> ,IV bolus <b>5-FU</b> 2400 mg/m <sup>2</sup> ,IV,46hrs	Q2W	NO.08/Level IB
<b>Gemcitabine</b> 1000 mg/m <sup>2</sup> , IV,D1,D8,D15	Q28 d	NO.09/Level IA
<b>Gemcitabine</b> 1000 mg/m <sup>2</sup> , IV,D1,D8 + <b>TS-1</b> 60-100mg/day BSA $\geq 1.5\text{m}^2$ : 100mg /day, 1.25m <sup>2</sup> - 1.5m <sup>2</sup> : 80mg/day, <1.25m <sup>2</sup> : 60mg/day,PO,D1-14	Q21 d	NO.10 /Level IB NO.15 /Level III
<b>TS-1</b> 80-120mg/day (PO 4 weeks on, 2 weeks off/ or PO 2 weeks on, 1 weeks off) BSA $\geq 1.5\text{m}^2$ : 120mg /day, 1.25m <sup>2</sup> - 1.5m <sup>2</sup> : 100mg/day, <1.25m <sup>2</sup> : 80mg/day	Q21d~ Q42 d /cycle	NO.10 /Level IB

\*健保用藥9.10: 3.與 5-fluorouracil、leucovorin 及 irinotecan 併用(FOLFIRINOX)，作為轉移性胰臟癌之第一線治療(限用 Oxalip、Opatin、Eloxatin、Folep)。(自110年5月1日生效)

# 胰臟腺癌

## 化學治療處方建議表:局部晚期、轉移癌化療-2

Chemotherapy for unresectable 、metastasis ( ECOG grade $\leq 2$ )	Schedule	Reference (No)/ strength of Evidence
<b>SLOG</b> Gemcitabine 800 mg/m <sup>2</sup> , IV, D1 Oxaliplatin 85 mg/m <sup>2</sup> ,IV,2hrs, D1 TS-1 35mg/m <sup>2</sup> /daily, BIDPC (Max daily dose 120mg), D1-D7 Calcium Folate Folic acid(15mg/tab) 20mg/m <sup>2</sup> /daily, BID, D1-D7	Q2W/cycle	NO.20 /Level V
<b>FIRINOX</b> Oxaliplatin 85 mg/m <sup>2</sup> ,IV,2hrs Irinotecan 150 mg/m <sup>2</sup> ,IV,90mins 5-FU 2400 mg/m <sup>2</sup> ,IV,46hrs	Q2W/ cycle x 4	NO.24/Level V
<b>Nab-paclitaxel (Abraxane)*</b> 125 mg/m <sup>2</sup> , IV, D1, D8, D15 Gemcitabine 1000 mg/m <sup>2</sup> , IV, D1, D8, D15	Q4W/cycle	NO.21 /Level I

\*健保用藥9.5.2：Albumin-based paclitaxel (如Abraxane):(108/11/01)限併用gemcitabine，作為轉移性胰臟癌患者之第一線治療。(自108年11月1日生效)

## 化學治療二線處方建議表-1

Chemotherapy for unresectable/recurrent disease ( ECOG grade $\leq 2$ )	Schedule	Reference (No)/ strength of Evidence
<b>Liposomal irinotecan and fluorouracil</b> <b>Onivyde</b> *60-80 mg/m <sup>2</sup> ,IV, keep 90mins <b>Leucovorin</b> 400 mg/m <sup>2</sup> ,IV, over 30mins <b>5-FU</b> 2400 mg/m <sup>2</sup> , IV, for 46hrs	Q2W/cycle Until progression	NO.16/Level IB
<b>FOLFIRI</b> <b>Irinotecan</b> 180 mg/m <sup>2</sup> ,IV, D1 <b>Leucovorin</b> 400 mg/m <sup>2</sup> ,IV, 2hrs <b>5-FU</b> 400 mg/m <sup>2</sup> , IV bolus <b>5-FU</b> 2400 mg/m <sup>2</sup> ,IV,46hrs	Q2W/cycle Until progression	NO.23/Level I

\*健保用藥9.12.2：Irinotecan微脂體注射劑(如Onivyde):(自107年8月1日生效)

- 1.與5-FU及leucovorin合併使用於曾接受過gemcitabine治療後復發或惡化之轉移性胰臟腺癌。
- 2.需經事前審查核准後使用。

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## 化學治療二線處方建議表-2

Chemotherapy for unresectable/recurrent disease ( ECOG grade $\leq 2$ )	Schedule	Reference (No)/ strength of Evidence
<b>FIRINOX</b> <b>Oxaliplatin</b> 85 mg/m <sup>2</sup> ,IV,2hrs <b>Irinotecan</b> 150 mg/m <sup>2</sup> ,IV,90mins	Q2W/cycle Until progression	NO.25/Level V
<b>SOXIRI</b> <b>Oxaliplatin</b> 85 mg/m <sup>2</sup> ,IV,2hrs <b>Irinotecan</b> 150 mg/m <sup>2</sup> ,IV,90mins <b>TS-1</b> 80mg/m <sup>2</sup> , BID	Q2W/cycle Until progression	NO.25/Level V
<b>GAS</b> <b>Nab-paclitaxel (Abraxane)*</b> 80~100 mg/m <sup>2</sup> , IV <b>Gemcitabine</b> 600~1000 mg/m <sup>2</sup> , IV <b>TS-1*</b> 60-100mg/m <sup>2</sup> /day(by BSA) , PO, D1-D7 BSA $\geq 1.5\text{m}^2$ : 100mg /m <sup>2</sup> /day, 1.25m <sup>2</sup> - 1.5m <sup>2</sup> :60-100mg/m <sup>2</sup> /day, <1.25m <sup>2</sup> : 60mg/m <sup>2</sup> /day	Q2W/cycle Until progression	NO.28、29 /Level V

\*健保用藥9.46：TS-1治療局部晚期無法手術切除或轉移性胰臟癌病人。

\*健保用藥9.5.2：Albumin-based paclitaxel (如Abraxane):(108/11/01)限併用gemcitabine，作為轉移性胰臟癌患者之第一線治療。

(自108年11月1日生效)

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## 標靶治療處方建議表：未惡化之轉移癌維持治療

Chemotherapy for metastasis (maintenance therapy)	Schedule	Reference (No)/ strength of Evidence
<b>Olaparib</b> (Lynparza) 300-600mg/day, PO, BID	QD Until progression	NO.27/Level IB

\*Olaparib(Lynparza)單一療法之維持治療，可用於遺傳性BRCA突變且經第一線含鉑化療至少16週後疾病未惡化之轉移性胰臟腺癌成年病人。

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## 動脈內化學放射治療處方建議表

Indications:

1. Post-operative liver metastasis from pancreatic cancer

<b>1. Intra-arterial Chemoradiotherapy for post-operative liver metastasis(術後肝轉移，ECOG grade <math>\leq 2</math>)</b>	<b>Schedule</b>	<b>Reference (No)/ strength of Evidence</b>
IA Chemotherapy 5-FU D1~D5 and IA Gemcitabine, D1~D5 Gemcitabine 400mg/m <sup>2</sup> /d, IA, over 30mins 5-FU 250mg/m <sup>2</sup> /d, IA, over 24hrs from day1 to 5	Q 2~ 4 W	NO.13/Level IIB NO.18/Level IV



## 放射治療處方建議表

### Indication :

Reference (No)/ strength of Evidence N0.14、30

- (1)Adjuvant CCRT for R1 resection and R2 resection
- (2)For medically fit patients but **marginal resectable**/unresectable cancer without distant metastasis
- (3)For medically unfit patients without distant metastasis
- (4)Following CCRT, additional maintenance chemotherapy is suggested
- (5)Recurrence**

### CCRT:

#### (1)Radiation therapy:

Target volume: tumor bed, adjacent LN and surgical anastomosis (for post OP adjuvant CCRT)  
Dose: 45-54 Gy (1.8-2 Gy/day)

#### (2)Chemotherapy regimen:

- Capecitabine(1000 ~1500mg/m<sup>2</sup>/day )/day in two divided doses, PO**
- Gemcitabine (600 mg/m<sup>2</sup> ) beginning the first day of RT (before RT), then weekly thereafter during RT



### PRINCIPLES OF SYSTEMIC THERAPY

#### Chemoradiation

##### Preferred Regimens

- Capecitabine + concurrent RT
- Continuous infusion 5-FU + concurrent RT

##### Other Recommended Regimens

- Gemcitabine + concurrent RT<sup>30</sup>

##### Useful in Certain Circumstances

- None

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## 癌症藥物停藥準則

影像學檢查，腫瘤有變大或轉移變多，應停止或改變治療方式。

## AJCC 8<sup>th</sup> 胰臟癌分期

Reference (No): 1

**Table 1. Definitions for T, N, M**  
American Joint Committee on Cancer (AJCC) TNM Staging of Pancreatic Cancer (8th ed., 2017)

<b>T</b>	<b>Primary Tumor</b>	<b>N</b>	<b>Regional Lymph Nodes</b>
<b>TX</b>	Primary tumor cannot be assessed	<b>NX</b>	Regional lymph nodes cannot be assessed
<b>T0</b>	No evidence of primary tumor	<b>N0</b>	No regional lymph node metastases
<b>Tis</b>	Carcinoma <i>in situ</i> This includes high-grade pancreatic intraepithelial neoplasia (PanIn-3), intraductal papillary mucinous neoplasm with high-grade dysplasia, intraductal tubulopapillary neoplasm with high-grade dysplasia, and mucinous cystic neoplasm with high-grade dysplasia	<b>N1</b>	Metastasis in one to three regional lymph nodes
<b>T1</b>	Tumor ≤2 cm in greatest dimension	<b>N2</b>	Metastasis in four or more regional lymph nodes
<b>T1a</b>	Tumor ≤0.5 cm in greatest dimension	<b>M</b>	<b>Distant Metastasis</b>
<b>T1b</b>	Tumor >0.5 cm and <1 cm in greatest dimension	<b>M0</b>	No distant metastasis
<b>T1c</b>	Tumor 1–2 cm in greatest dimension	<b>M1</b>	Distant metastasis
<b>T2</b>	Tumor >2 cm and ≤4 cm in greatest dimension		
<b>T3</b>	Tumor >4 cm in greatest dimension		
<b>T4</b>	Tumor involves the celiac axis, superior mesenteric artery, and/or common hepatic artery, regardless of size		

**Table 2. AJCC Prognostic Groups**

	<b>T</b>	<b>N</b>	<b>M</b>
<b>Stage 0</b>	Tis	N0	M0
<b>Stage IA</b>	T1	N0	M0
<b>Stage IB</b>	T2	N0	M0
<b>Stage IIA</b>	T3	N0	M0
<b>Stage IIB</b>	T1, T2, T3	N1	M0
<b>Stage III</b>	T1, T2, T3	N2	M0
	T4	Any N	M0
<b>Stage IV</b>	Any T	Any N	M1

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