

GEMCITABINE with concurrent RT

INDICATION (ICD10) C25

1. Locally advanced non-metastatic pancreatic cancer unable to receive capecitabine (unlicensed)

REGIMEN

Days 1, 8, 15, 22, 29 and 36

GEMCITABINE 300mg/m² infusion in #ml sodium chloride 0.9% IV infusion over 30 minutes

diluent and diluent volume for dose prescribed as per national standardised product specification or licensed dose

CYCLE FREQUENCY AND NUMBER OF CYCLES

1 cycle (5.5 weeks) with radiotherapy

ADMINISTRATION

Day 1 to be prescribed to start on day 2 of radiotherapy (total planning target volume should be restricted to smaller than 800cm³).

Gemcitabine should be given prior to RT when pt is due both.

ANTI-EMETICS

Low risk days 1, 8, 15, 22, 29 and 36

CONCURRENT MEDICATION REQUIRED

| | |
|-------------|---------------|
| Gemcitabine | None required |
|-------------|---------------|

EXTRAVASATION AND TYPE OF LINE / FILTERS

Gemcitabine – neutral

No filters required

Central or peripheral line

INVESTIGATIONS

Blood results required before SACT administration

FBC every dose, U&E, LFTs and creatinine every cycle

Neutrophils x 10⁹/L ≥1.5

Platelets x 10⁹/L ≥100

Baseline weight and every cycle

MAIN TOXICITIES AND ADVERSE REACTIONS

| | |
|-------------|-----------------------------------------------------------------------------------------------------------------------------------|
| Gemcitabine | Diarrhoea – see dose modifications, treat with loperamide or codeine Mucositis – see dose modifications, use routine mouthcare |
|-------------|-----------------------------------------------------------------------------------------------------------------------------------|

DOSE MODIFICATIONS

Haematological

Gemcitabine

| | |
|-------------------------------------------------------------------------|-----------------------------------------------------|
| Neutrophils $>1.5 \times 10^9/L$ and platelets $>100 \times 10^9/L$ | give 100% dose |
| Neutrophils $1.0-1.5 \times 10^9/L$ or platelets $75-100 \times 10^9/L$ | Discuss with consultant |
| Neutrophils $<1.0 \times 10^9/L$ or platelets $<100 \times 10^9/L$ | delay treatment (day 1) or otherwise omit treatment |

Non-haematological

Gemcitabine

| | |
|---------------------------------------------|--------------------------------------------------------|
| Diarrhoea and/or mucositis grade 2 toxicity | omit until toxicity resolved then restart at 100% dose |
| Diarrhoea and/or mucositis grade 3 | omit until toxicity resolved then restart at 75% dose |
| Diarrhoea and/or mucositis grade 4 | omit until toxicity resolved then restart at 50% dose |

Hepatic impairment

Gemcitabine

| | |
|---------------------------------|----------------------------------|
| Bilirubin $>27 \mu\text{mol/L}$ | initiate treatment with 80% dose |
|---------------------------------|----------------------------------|

Renal impairment

Gemcitabine

No need for dose adjustment

REFERENCES

1. International archives of Medicine 2009 2:7-13
2. Huang et al IJROBP (2009) 73 (1): 159-165