PANCO¹ CASE STUDY: Stage III T4NOMO 24cc Tumour

• 83-year-old male

PRESENTATION

- Pre-existing asthma and hypertension
- ECOG 1 performance status

- LAPC diagnosed 50 days prior to enrolment in the PanCO study:
- Stage III T4NOMO
- 5.4cm longest diameter*/24cc tumour volume*



biliary stent and encasing the proximal superior mesenteric artery (blue arrow)



FDG-PET/CT: There is a focal area of increased FDG uptake (SUVmax 6.4*) in the uncinate process/head of the pancreas, adjacent to the biliary stent, in keeping with the known primary tumour (yellow arrow).

Dr Harpreet Wasan

Imperial College and NHS Foundation Trust, London, UK

on behalf of the PanCO Study investigators

Pancreatic duct is dilated proximal to the lesion (yellow arrow). Portal vein is patent. Aerobila are observed.

Abbreviations used in this case study

AE:	Adverse event
CT:	Computed tomography
FDG-PET:	Fluorodeoxyglucose-positron emission tomography
LAPC:	Locally advanced pancreatic cancer
LD:	Longest diameter
SUVmax:	Maximum standardised uptake value
TEAE:	Treatment-emergent adverse event
TV:	Tumour volume

Reference: 1. The PanCO study: ClinicalTrials.gov Identifier: NCTO3003078. Data on file. * By Central Image Reader

INTENDED USE/INDICATIONS FOR USE: OncoSII" is intended for intratumoural implantation into a pancreatic tumour via injection under endoscopic ultrasound guidance. OncoSil" is indicated for the treatment of patients with locally advanced unresectable pancreatic cancer, in combination with gemcitabine-based chemotherapy The OncoSil[™] System is supplied sterile and is intended for single-patient, single-use.

This information is intended for healthcare professionals only. All medical treatments carry benefits and risks. For safety related information, please refer to the OncoSil" System Instructions for Use.





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• The post-implant TEAEs occurred at 6 months (high CRP levels), 21 months (anaemia) and 24.6 months (anaemia) from implant, all of which were possibly related to chemotherapy

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TUMOUR RESPONSE



Bremsstrahlung

4 hours post OncoSil[™] 7 days implantation implan

T days post OncoSil™ implantation

On post-implantation Bremsstrahlung planar and SPECT/CT images, the highest activity is localised to the head of the pancreas, anterior to the biliary stent (yellow arrow). There is contiguous activity in the uncinate process medially. There is much lower activity in the proximal small bowel loops.





4 hours post OncoSil[™] implantation

7 days post OncoSil™ implantation



4 hours post OncoSil[™] implantation

7 days post OncoSil™ implantation

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37% reduction in the FDG avidity of the primary pancreatic tumour by SUVmax* [6.4 to 4.07] (yellow arrow), indicating a favourable response to treatment.



-44% LD reduction

-86% TV reduction (3cc vs 24cc at baseline)



CT Imaging at 80 weeks

Notes

- CA 19-9 response not applicable patient had CA 19-9 <ULN at baseline
- Patient unsuitable for surgical resection due to age, co-morbidities etc



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TUMOUR RESPONSE