

MDT Discussion

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Disclosures



Dr David Woolf: I have received honoraria for travel expenses from Roche and Elekta, for consultancy services from Genesis Care and for my participation in the 2024 Think Again Lung Summit from AstraZeneca.

Mr Alessandro Brunelli: I have received honoraria for my participation in advisory boards from AstraZeneca, Bristol Myers Squibb, Merck Sharp & Dohme, Roche, Medtronic, Medela and Ethicon, and for my participation in the 2024 Think Again Lung Summit from AstraZeneca.

Dr Craig Dyer: I have received honoraria from Roche, Lilly, and AstraZeneca, and for my participation in the 2024 Think Again Lung Summit from AstraZeneca.

Dr Kevin Franks: I have received honoraria for consultancy and my participation in advisory boards from AstraZeneca, Amgen, BioNTech, Boehringer-Ingelheim, Bristol Myers Squibb, Lilly, Roche, and Takeda, and for my participation in the 2024 Think Again Lung Summit from AstraZeneca.

Dr Spyros Gennatas: I have received honoraria for my participation in the 2024 Think Again Lung Summit from AstraZeneca.

Ms Emma Halkyard: I have received honoraria from Takeda, AstraZeneca, Roche, Daiichi Sankyo, and Janssen, and for my participation in the 2024 Think Again Lung Summit from AstraZeneca.

Dr Anna Sharman: I have received honoraria for my participation in the 2024 Think Again Lung Summit from AstraZeneca.

MDT Discussion

Case study 1

Mr Alessandro Brunelli, Dr Kevin Franks and Dr Spyros Gennatas

Based on a real patient case study, with certain features adapted.



63-year-old female



- Presented to the GP with a cough
- CXR showed a large mass and went on to have a CT and a biopsy

CT, computed tomography; CXR, chest radiography; GP, general practitioner.

PMH



- **Asthma**
- **Type 2 DM**
- **Hypertension**
- **Previous left MCA aneurysm clip**
- **Ex-smoker: 15 pack-years**
- **ECOG PS 1**

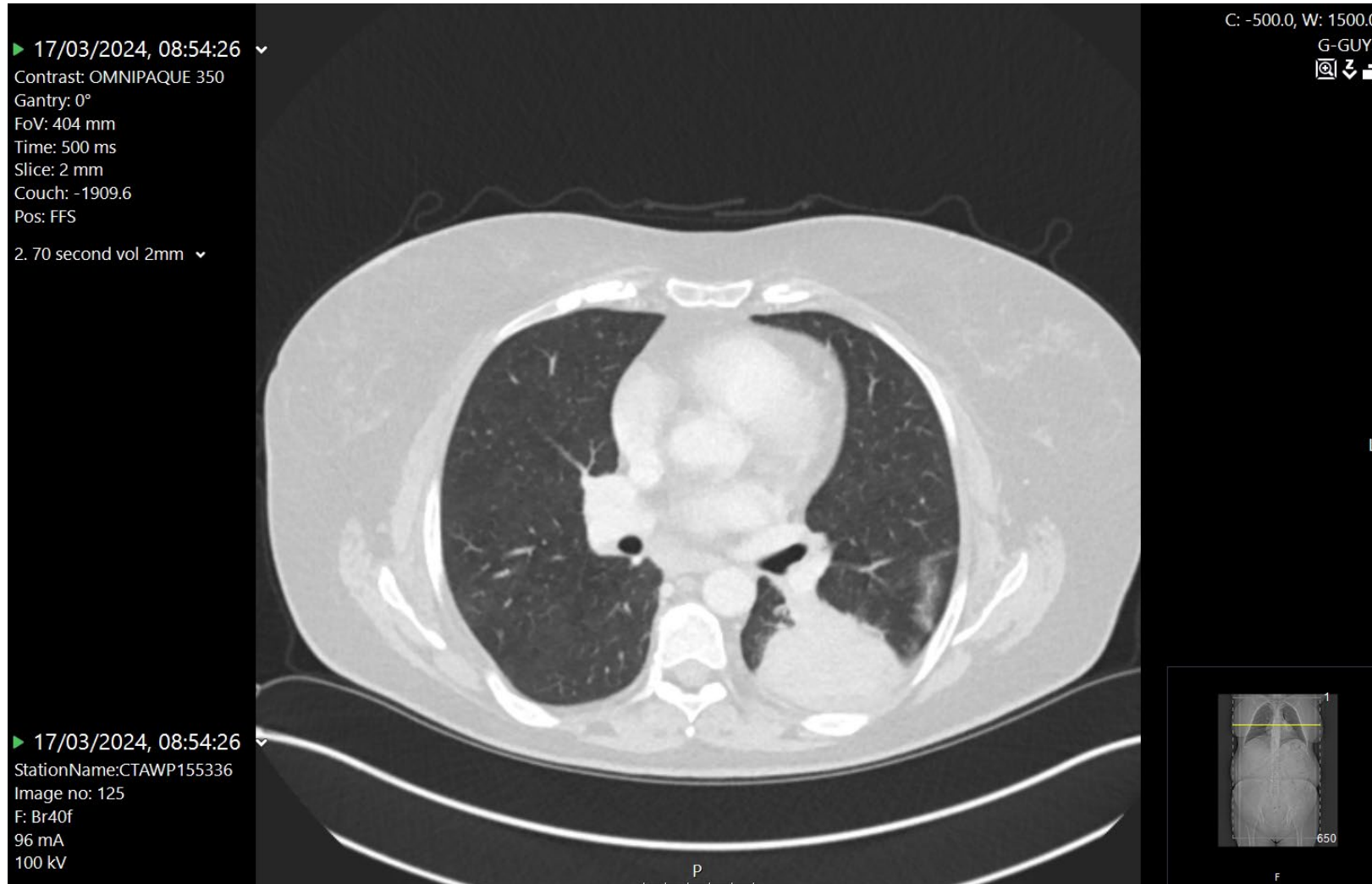
DM, diabetes mellitus; ECOG PS, Eastern Cooperative Oncology Group Performance Status; MCA, middle cerebral artery; PMH, past medical history.

- **Metformin**
- **Losartan**
- **Salbutamol**
- **Escitalopram**
- **Lansoprazole**
- **Gliclazide**
- **Seretide**
- **Diazepam**

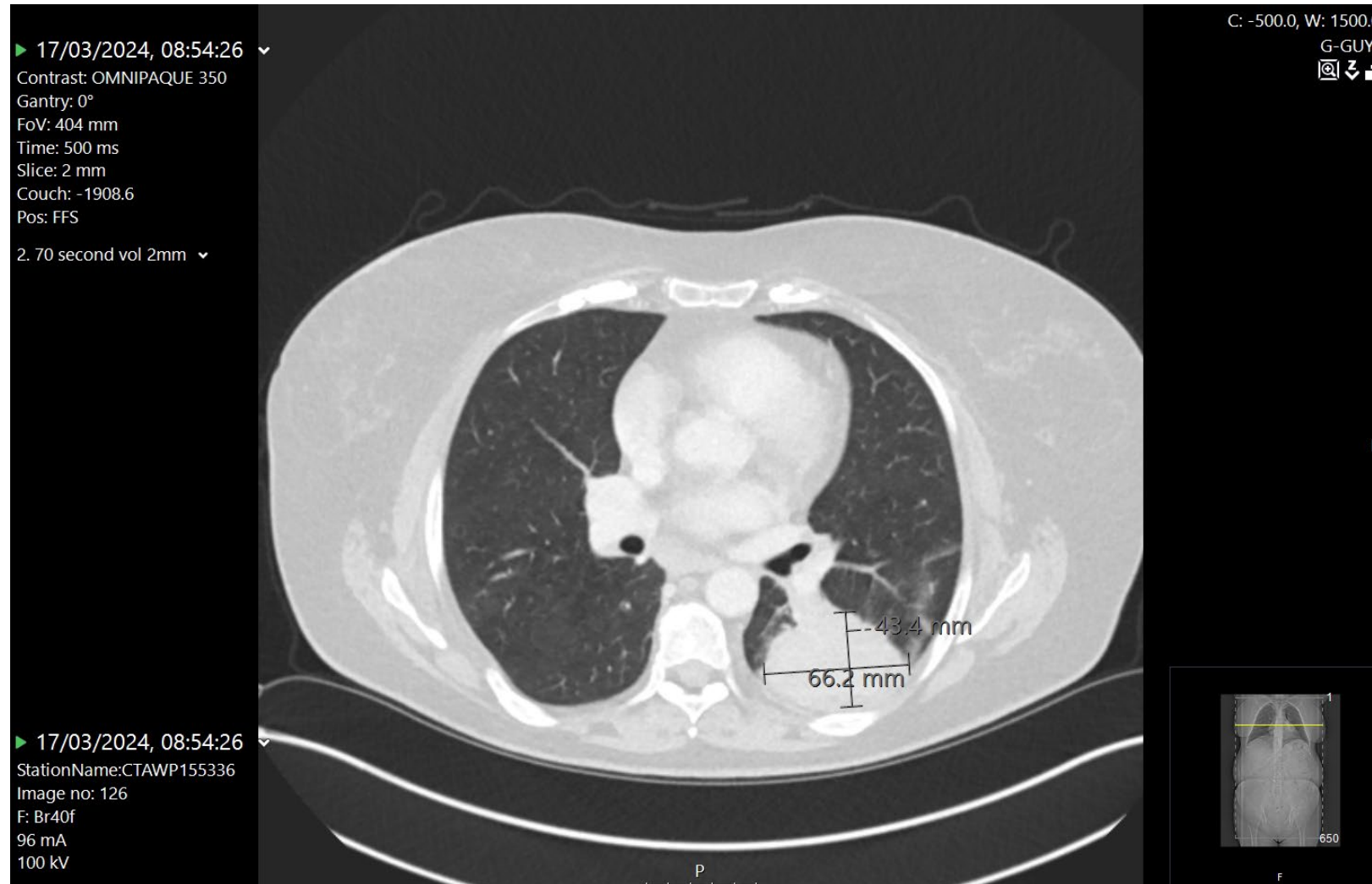
Please refer to the individual SmPCs for each product before prescribing, to review detailed information on indications, posology, special warnings, adverse events and their management.

DH, drug history; SmPCs, Summary of Product Characteristics.

Imaging



T4N0M0



Biopsy



- **TTF-1** positive adenocarcinoma
- **PD-L1** 10%, **ALK/ROS1** negative on IHC
- **EGFR, KRAS, BRAF, MET, ERBB2** WT

ALK, anaplastic lymphoma kinase; BRAF, v-raf murine sarcoma viral oncogene homolog B1; EGFR, epidermal growth factor receptor; ERBB2, v-erb-b2 avian erythroblastic leukaemia viral oncogene homolog 2; IHC, immunohistochemistry; KRAS, Kirsten rat sarcoma viral oncogene homolog; MET, mesenchymal epithelial transition factor receptor; PD-L1, programmed death ligand 1; ROS1, ROS proto-oncogene 1; TTF-1, thyroid transcription factor 1; WT, wild type.

Pulmonary function tests



- **FEV1:** 1.88 (86%)
- **FEV1/FEV:** 72%
- **FVC:** 2.54 (98%)
- **KCO:** 1.90 (122%)

FEV, forced expiratory volume; FEV1, forced expiratory volume in the first second; FVC, forced vital capacity; KCO, carbon monoxide transfer coefficient.

What would you do next?



Surgery



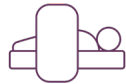
Chemotherapy



Chemo-immunotherapy



SABR/radiotherapy



PET



EBUS

EBUS, endobronchial ultrasound; PET, positron emission tomography; SABR, stereotactic ablative radiotherapy.

What would you do next?



1. Surgery
2. Chemotherapy
3. Chemo-immunotherapy
4. SABR/Radiotherapy
5. PET → No metastatic disease
6. EBUS

EBUS, endobronchial ultrasound; PET, positron emission tomography; SABR, stereotactic ablative radiotherapy.

Let's remind ourselves



- **T4N0M0 LLL adenocarcinoma**
- **PD-L1 10%**
- **Molecular diagnostics all negative**
- **Very good lung function**

LLL, left lower lobe; PD-L1, programmed death ligand 1.

What treatment would you offer next?



Bi-lobectomy



Pneumonectomy



Chemotherapy



Chemo-immunotherapy



SABR/radiotherapy

EBUS, endobronchial ultrasound; PET, positron emission tomography; SABR, stereotactic ablative radiotherapy.

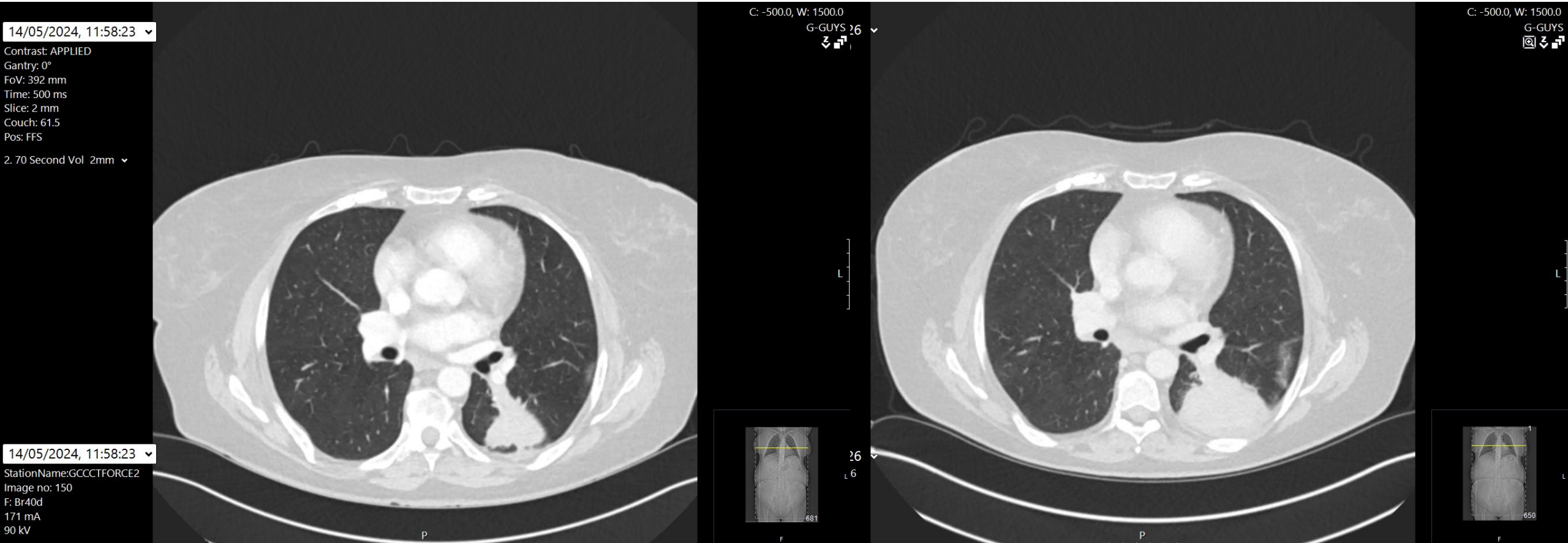
This was supposed to be straight-forward



1. Bi-lobectomy
2. Pneumonectomy
3. Chemotherapy
4. **Chemo-immunotherapy** → **Neoadjuvant chemo-immunotherapy**
5. SABR/Radiotherapy

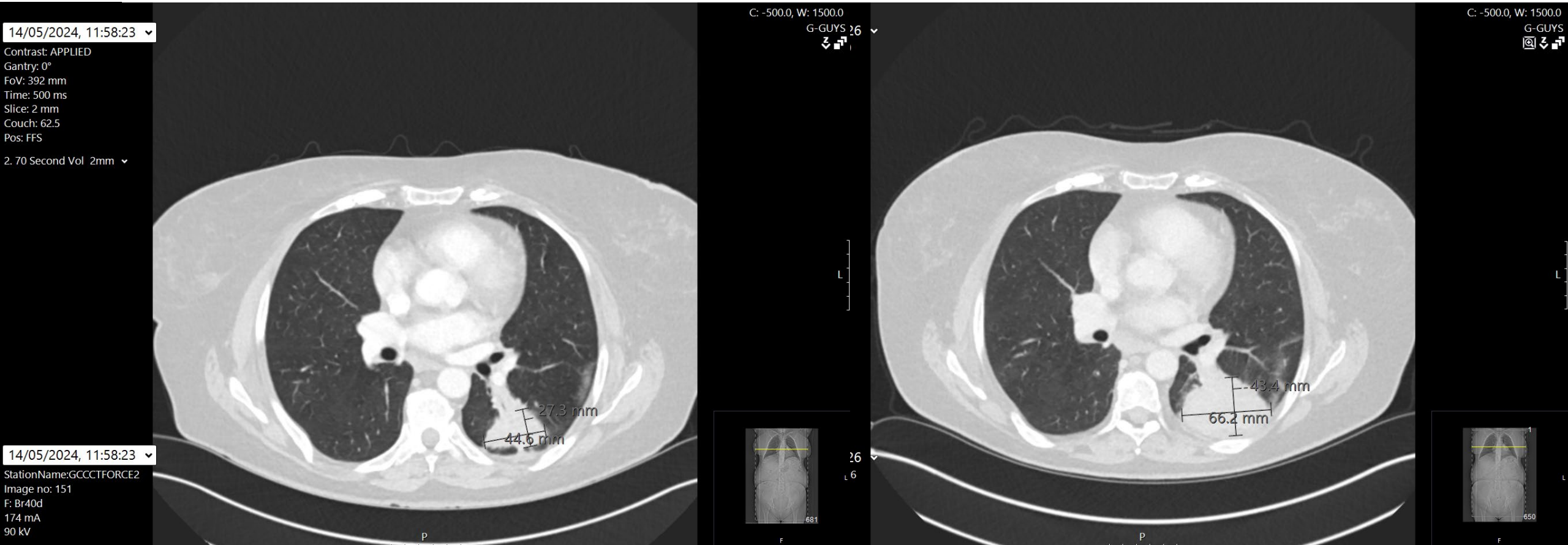
SABR, stereotactic ablative radiotherapy.

Bonus content – CT post 3 cycles of chemo-immunotherapy



CT, computed tomography.

Bonus content – CT post 3 cycles of chemo-immunotherapy



CT, computed tomography.

MDT Discussion

Case study 2

Mr Alessandro Brunelli, Dr Craig Dyer and Dr Kevin Franks

Based on a real patient case study, with certain features adapted.



Persistent complaints

3/52

- Cough, non-productive
- Weight loss (4-5kg)

6/12

- Progressive SOB
- Exertional fatigue – tolerance 500m (flat)



SOB, shortness of breath.

67/Male Tyre Fitter



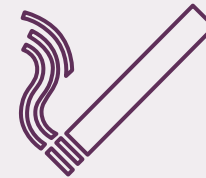
Past medical history

- Performance Status - 1
- Hypercholesterolaemia
- Type 2 Diabetes Mellitus



Smoking history

- Smoker – 60 pack year history
- Possible occupational asbestos exposure
- Lives alone – partner died last year



Bloods



Results				
Hb	105 (L)		Na	143
Plt	421 (H)		K	4.6
MCV	70 (L)		Urea	10.1 (H)
WCC	12.0 (H)		Creat	107
Neut	10.4			
Lymph	0.6		HbA1c	78 (H)
B12	371		LFT	Normal
Folate	23.2		Bone	Normal
Ferr	10 (L)			
T-Sat	5 (L)			

B12, vitamin B12; Creat, creatinine; Ferr, ferritin; Hb, hemoglobin; HbA1c, hemoglobin A1c; K, potassium; LFT, liver function tests; Lymph, lymphocyte count; MCV, mean corpuscular volume; Na, sodium; Neut, neutrophil count; Plt, platelet count; T-Sat, transferrin saturation; Urea, urea nitrogen; WCC, white cell count.

Modifiable risk factors?



- **Prehabilitation**
- **Dietetics**
- **Diabetic review**
- **Airways management**
- **Smoking cessation**
- **Iron replacement**
- **Social support / finance support / psychology input**

Radiology



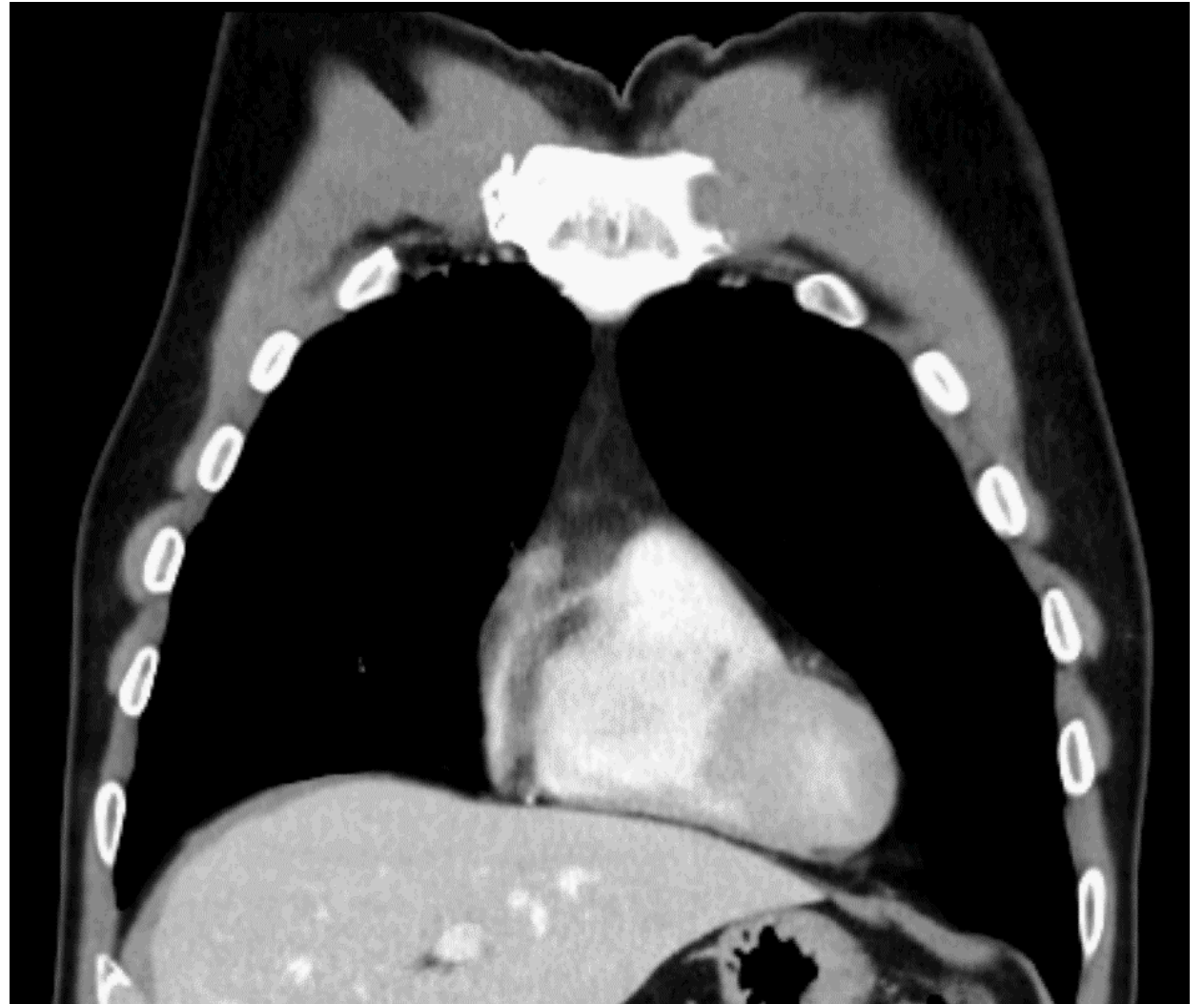
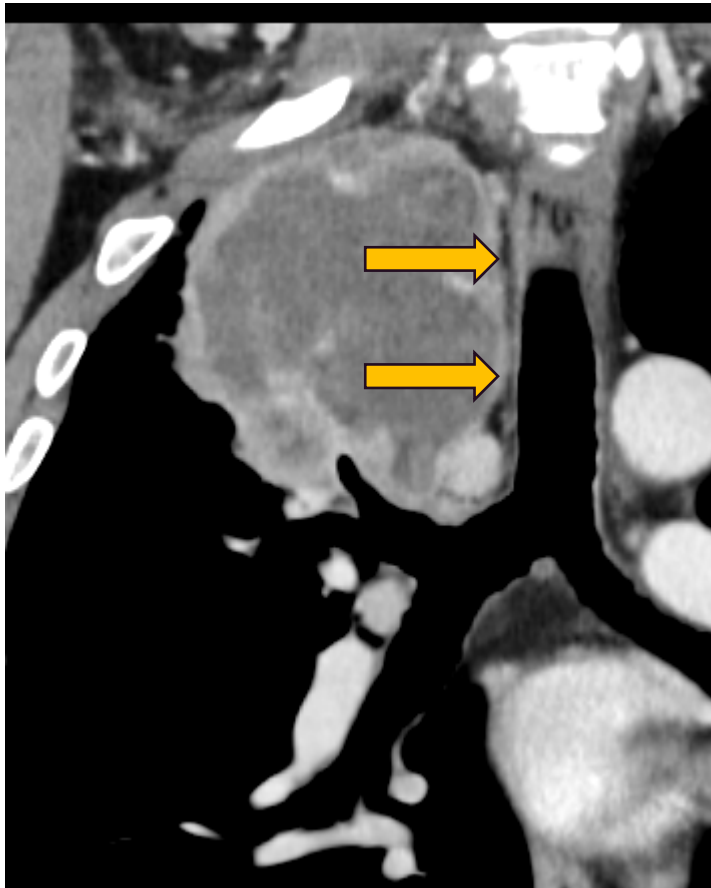
Radiological summary



Staging CT



Mediastinal fat plane



Next steps

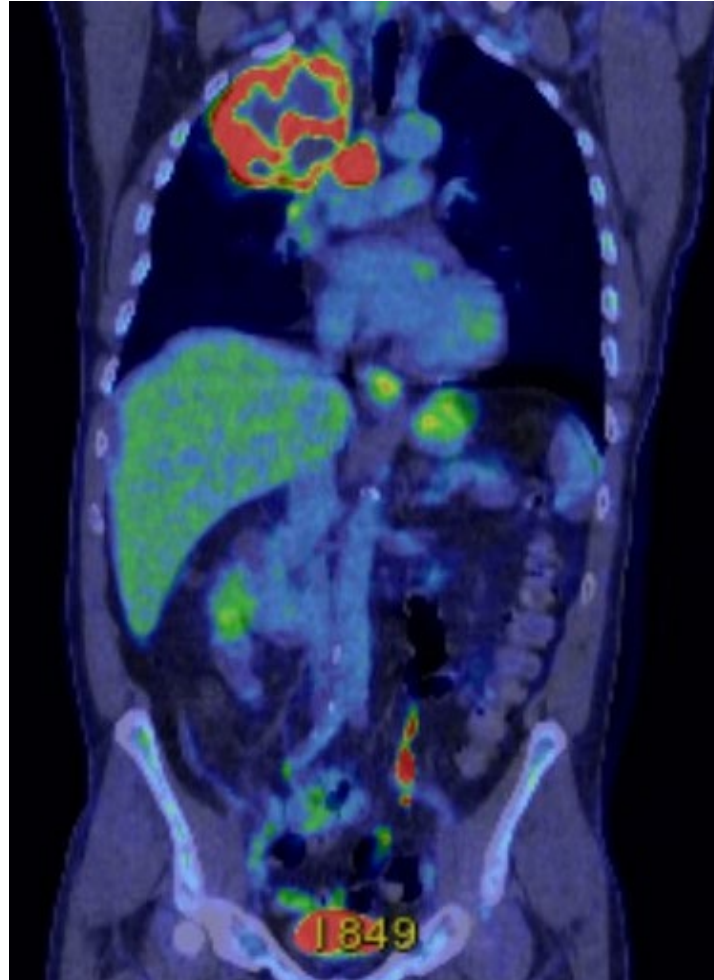


- **CT PET**
- **Bronchoscopy**
- **EBUS**
- **CT Guided Biopsy**
- **CT-DNA**

CT, computed tomography; CT-DNA, circulating tumor DNA; EBUS, endobronchial ultrasound; PET, positron emission tomography.

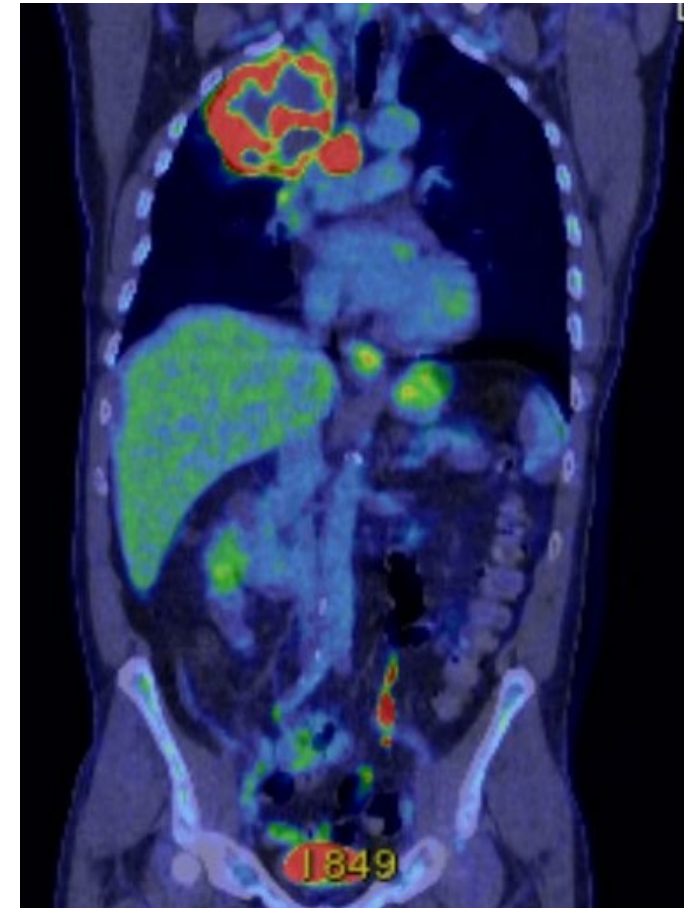
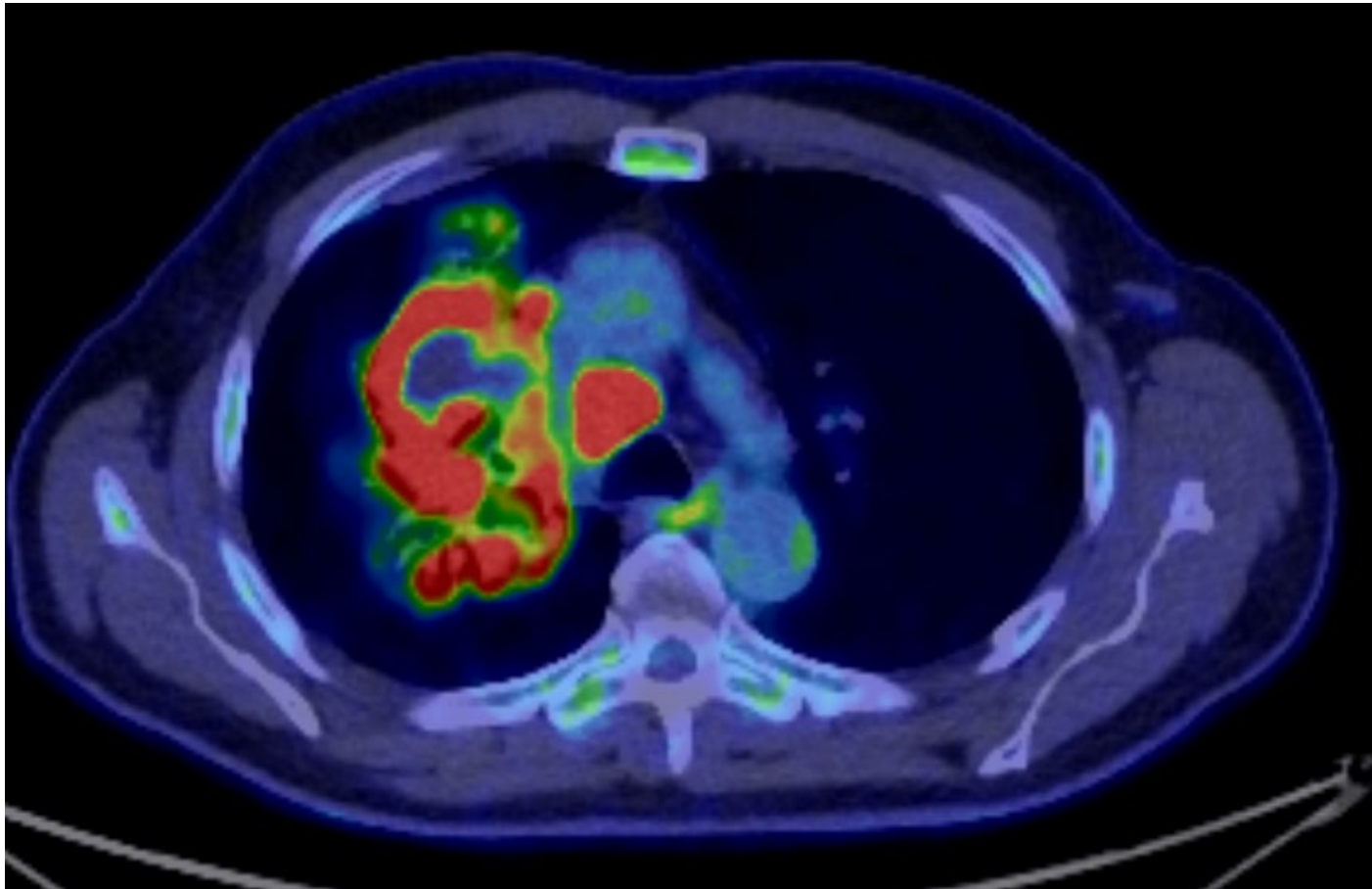
CT-PET

THINK
AGAIN



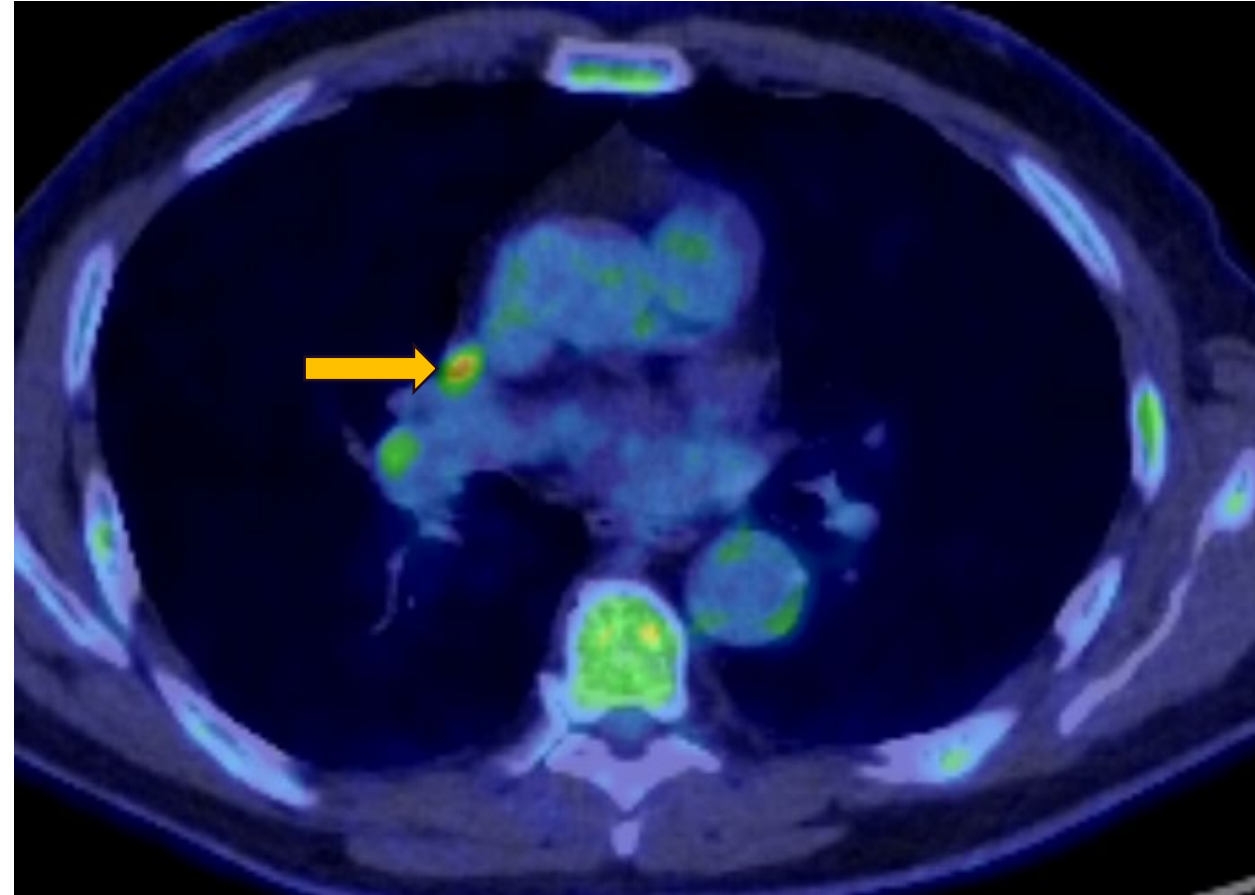
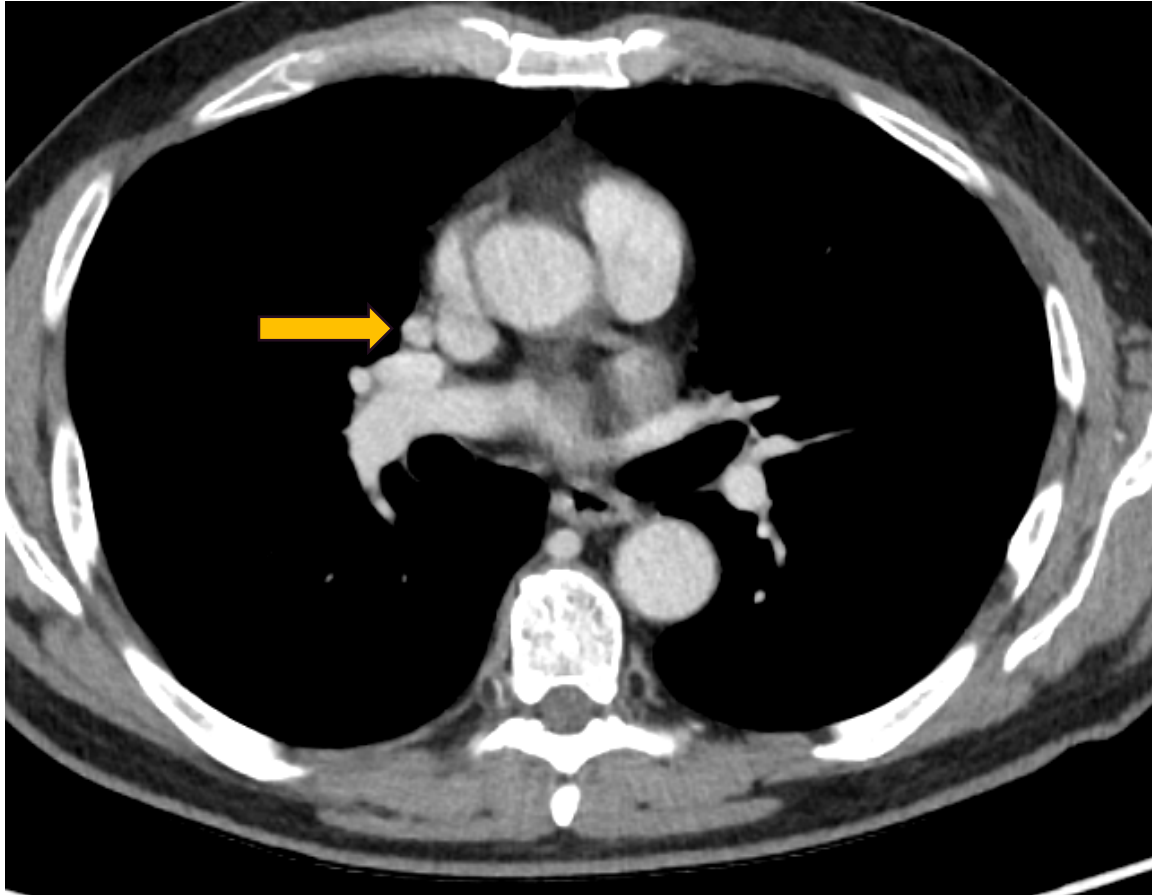
CT-PET=computed tomography-positron emission tomography.

CT-PET

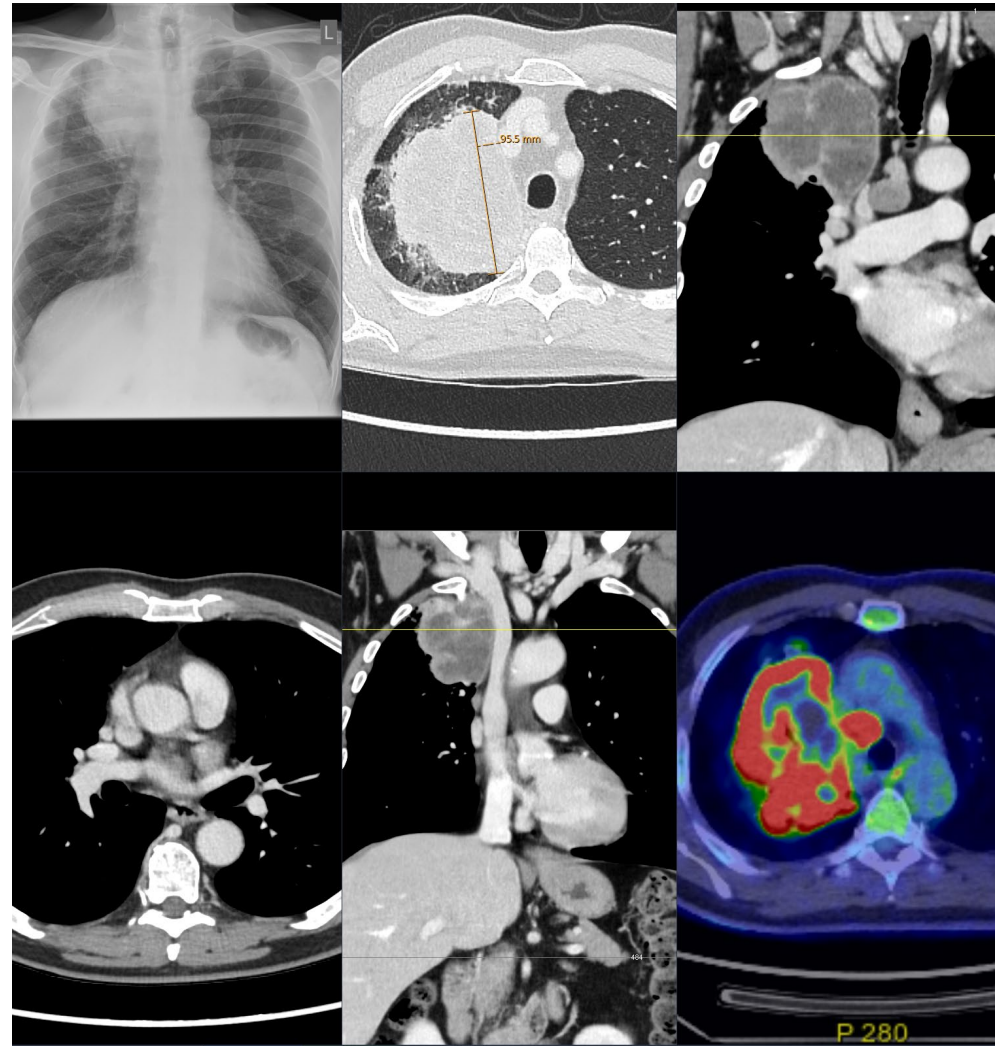


CT-PET=computed tomography-positron emission tomography.

Subcentimetre but avid node



Radiological summary – RUL cancer: T4 N2 M0



Physiology



Measurement	Test	Predicted
FEV1	2.5	91%
FVC	4.00	113%
FEV1/FVC	63%	
TLCO	5.51	74%
KCO	1.12	79%

ECHO:

- Ejection Fraction – 55%
- No valvular abnormality

MRI Brain:

- Normal

ECHO, echocardiogram; FEV1, forced expiratory volume in one second; FVC, forced vital capacity; KCO, transfer coefficient of the lung for carbon monoxide; MRI, magnetic resonance imaging; TLCO, transfer factor of the lung for carbon monoxide.

Pathology results



Staging EBUS Results	
Station 11L	No malignant cells
Station 10L	No malignant cells
Station 4L	No malignant cells
Station 7	No malignant cells
Station 4R	Adenocarcinoma – lung primary
Station 10 R	Adenocarcinoma – lung primary
Station 11R	Adenocarcinoma – lung primary

MDT – T4N2M0 Adenocarcinoma – PD-L1/NGS sent

EBUS, endobronchial ultrasound; MDT, multidisciplinary team; NGS, next-generation sequencing; PDL1, programmed death-ligand 1.

MDT discussion – day 24



Confirmed

- T4N2M0 Adenocarcinoma
- PD-L1 80%
- NGS – *KRAS* G12C positive (*EGFR/ALK* negative)
- Performance status 1
- Prehab – doing well – stable/improving, 6MWT 650m

6MWT, six-minute walk test; ALK, anaplastic lymphoma kinase; EGFR, epidermal growth factor receptor; MDT, multidisciplinary team; NGS, next-generation sequencing; PDL-1, programmed death-ligand 1.

What treatment would you offer next?



Surgery?



Radical radiotherapy?



Chemo-radiotherapy?



Neo-adjuvant chemo-immunotherapy?



Targeted therapies?

MDT, multidisciplinary team.

What would you do next?



1. Surgery
2. Radical radiotherapy
3. Chemo-radiotherapy
- 4. Neo-adjuvant chemo-immunotherapy**
5. Targeted therapies

MDT Discussion

Case study 3

Dr Anna Sharman and Dr David Woolf

Based on a real patient case study, with certain features adapted.



LUNG
SUMMIT

67-year-old female



- **MEDSURV patient**
- **Left VATS upper lobectomy T1c N1 adenosquamous 2022**
- ***EGFR* negative – seen by medical oncology but patient opted against chemotherapy**
- **Referral to MDT for abnormal RUL**

EGFR, epidermal growth factor receptor; MDT, multidisciplinary team; MEDSURV, medical survivorship programme; RUL, right upper lobe; VATS, video-assisted thoracoscopic surgery.

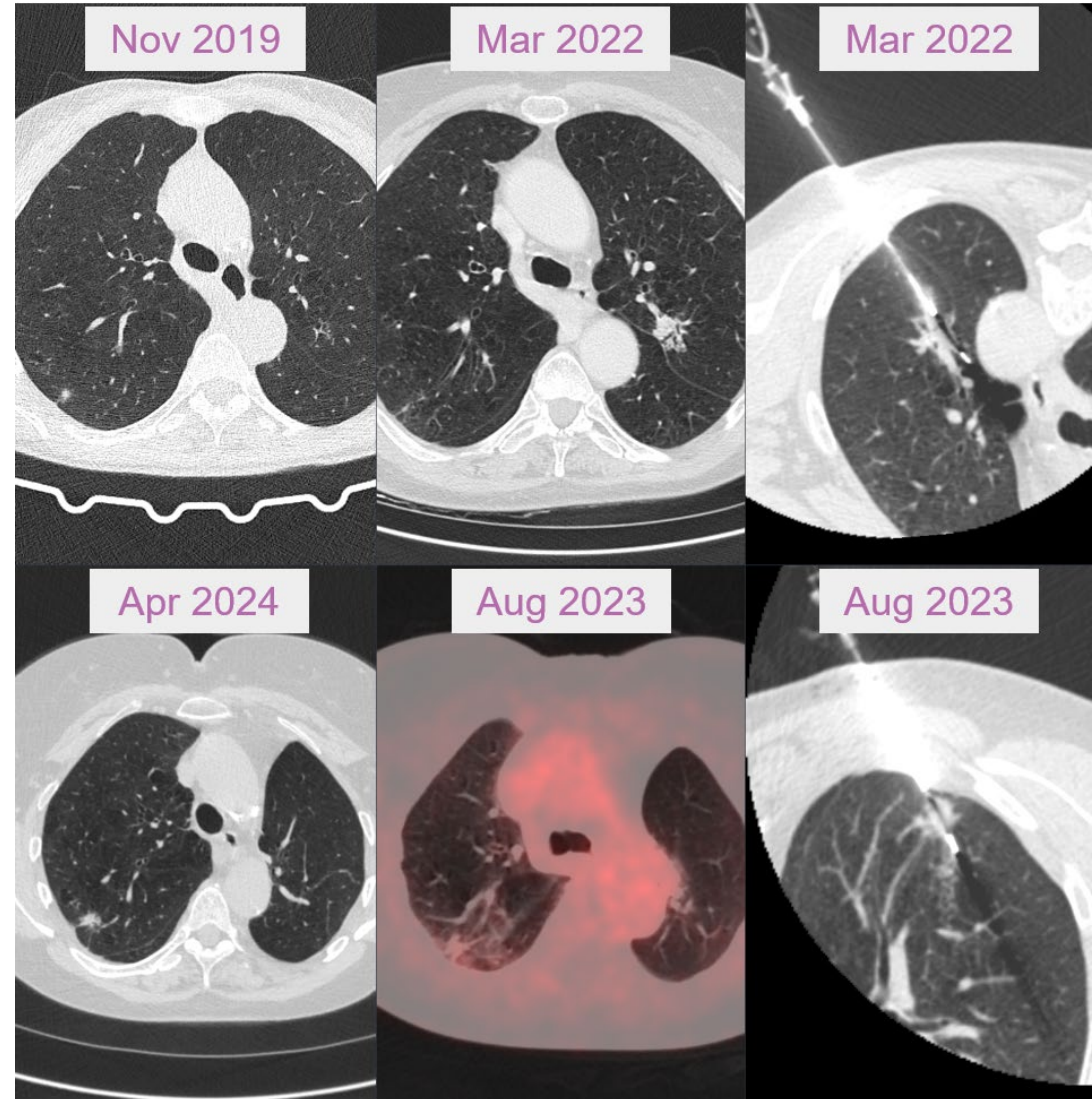
Past medical history



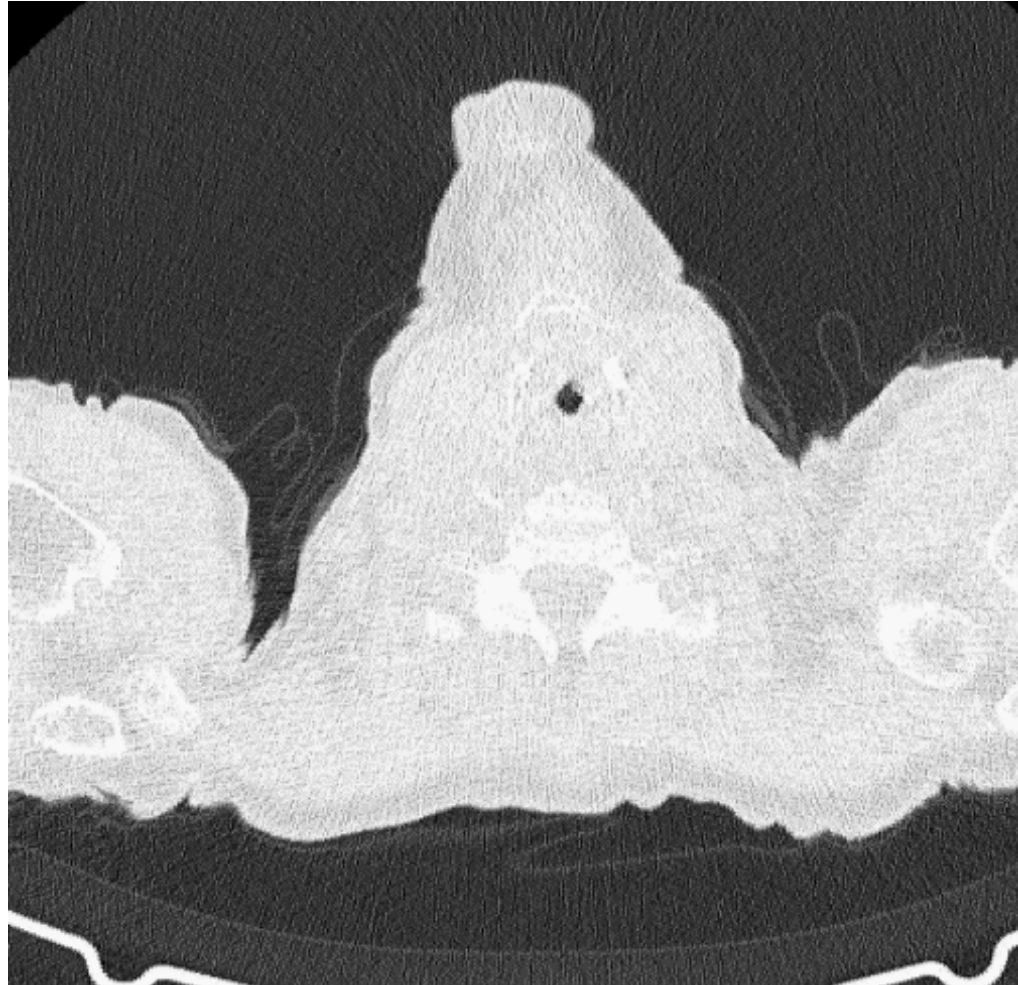
- **Ex-smoker**
- **COPD**
- **Temporal arteritis**
- **Thoracic aortic aneurysm under surveillance (4.2cm)**
- **PS 1**

COPD, chronic obstructive pulmonary disease; PS, performance status.

Radiological summary from 2019–2024



Back to the beginning – screening CT Nov 2019



CT, computed tomography.

2019 – multiple abnormal areas



RUL

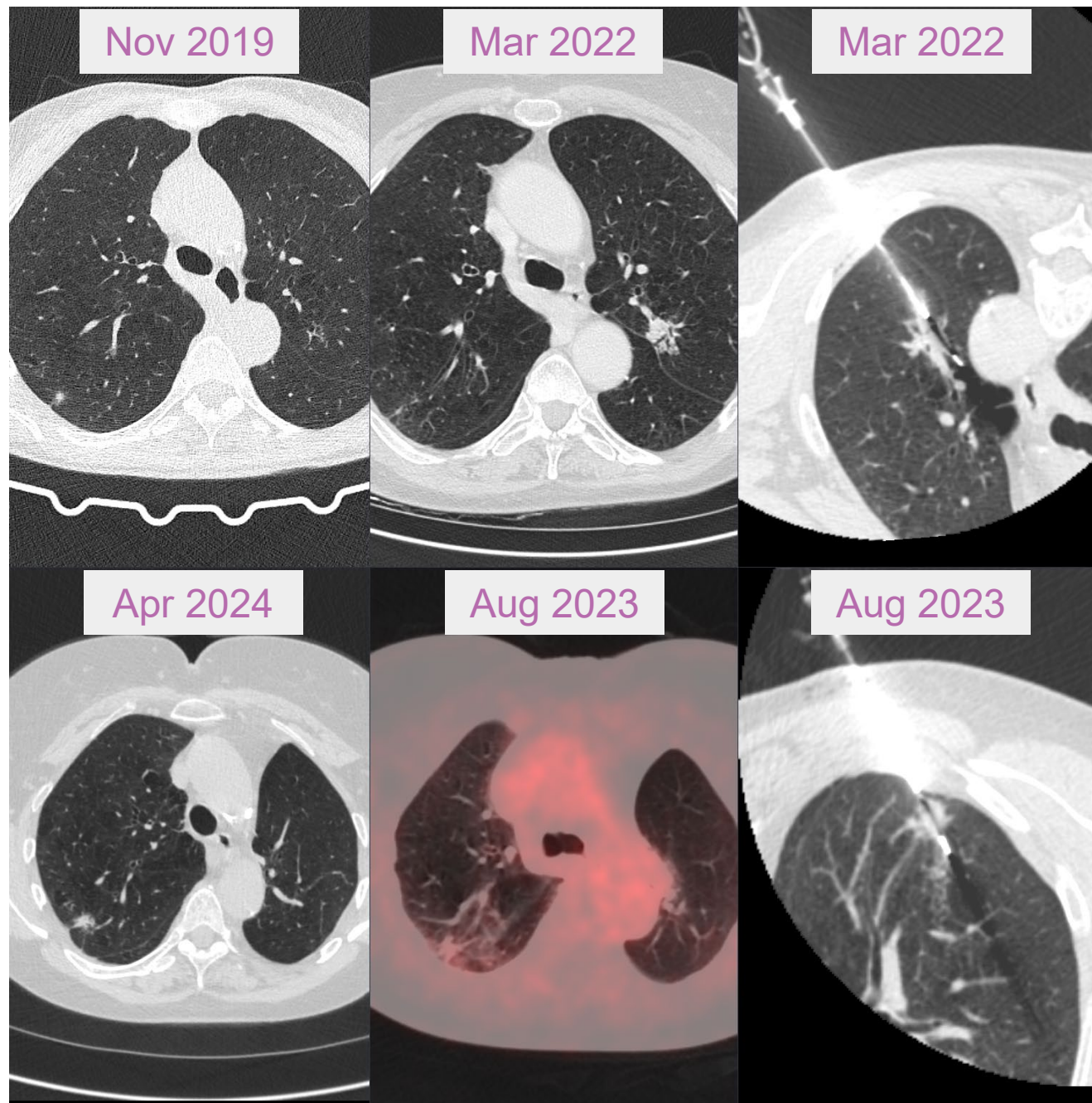
RML

LUL

LLL



LLL, left lower lobe; LUL, left upper lobe; RML, right middle lobe; RUL, right upper lobe.



CT scan



- LUL lobectomy
- Multiple nodules in RUL progressing from 2019
- Enlarging RML cyst



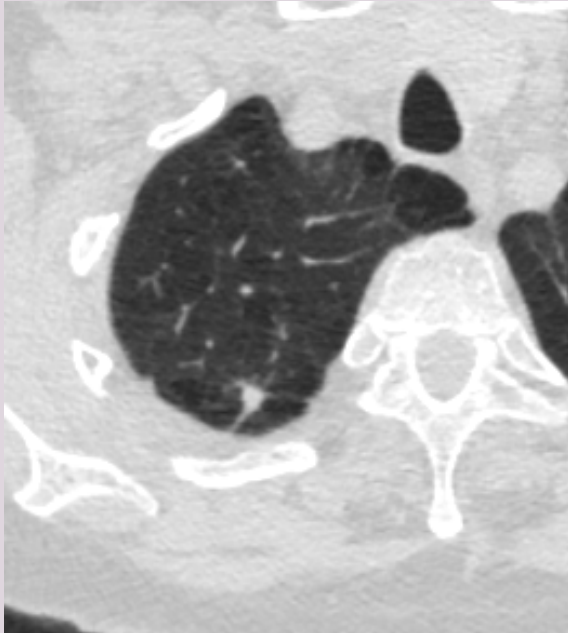
LUL, left upper lobe; RML, right middle lobe; RUL, right upper lobe.

PET done

THINK
AGAIN

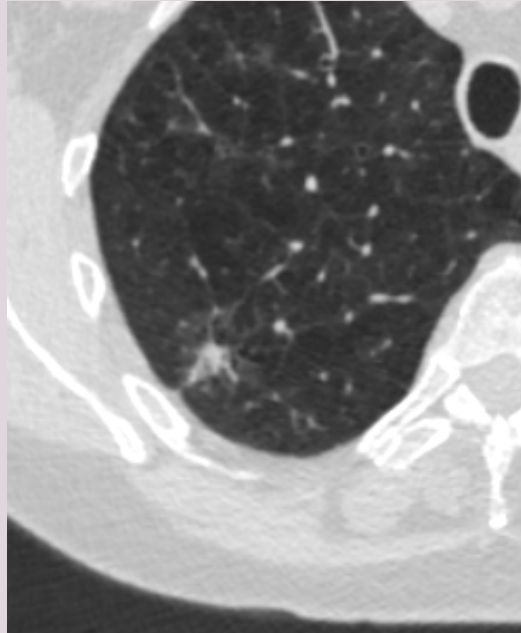
SUV max 2.6, Right apical 0.6 x 0.6 cm nodule (3.51). This lesion has slightly increased since 2022, was 0.5 x 0.4 cm and barely perceivable in 2019.

If NSCLC; radiological stage T1a N0



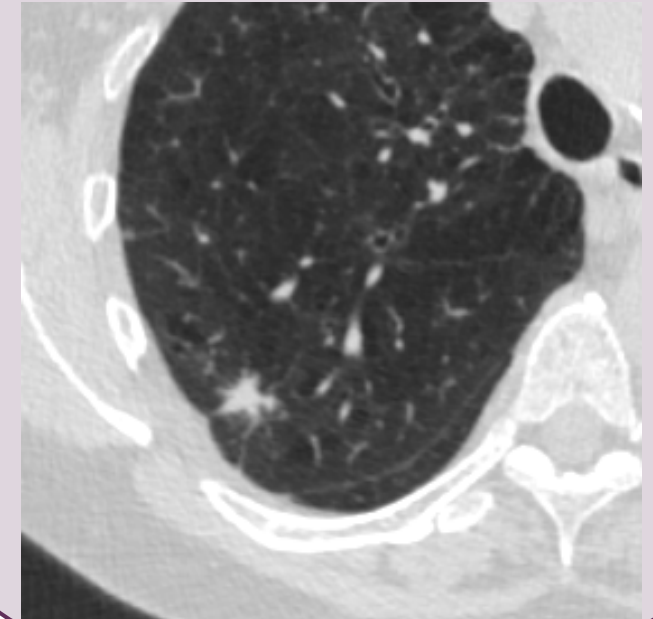
SUV max 1.5, RUL 1 x 0.7 cm nodule (3.59). Stable from 2019.

If NSCLC; radiological stage T1a N0



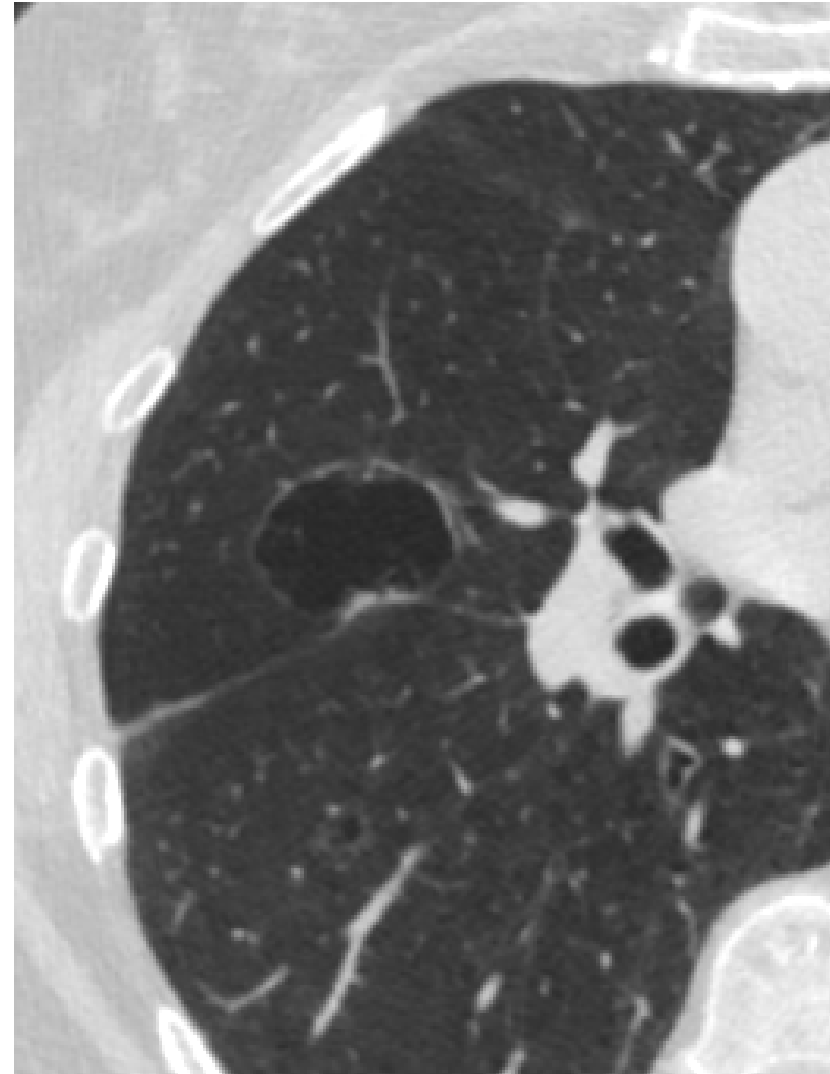
SUV max 2.9, RUL 1.1 x 0.8 cm nodule (3.64). Increased from 0.5 cm in Nov 2019 - this has been biopsied.

If NSCLC radiological stage T1b N0



NSCLC, non-small cell lung cancer; SUV max, maximum standardised uptake value.

- Right middle lobe 3cm cyst
- No measurable thickening
- **No focal increased metabolic activity**



Radiology conclusion



- **Complex imaging with multifocal abnormalities assumed to represent synchronous primary lung adenocarcinomas**
- **The dominant lesion is the most caudal nodule within the right upper lobe posteriorly which has been targeted for biopsy**
 - T1b (m) N0
- **The middle lobe cyst will require ongoing surveillance**

Pathology from RUL nodule



- **Adenocarcinoma**

RUL, right upper lobe.

Physiology



- **Echo: mild LVH, EF > 55%, mild AR/MR, dilated aortic root and AA, dilated LA**
- **FEV1 2.29L (105% predicted)**
- **DLCO % predicted: 46%**

AA, ascending aorta; AR, aortic regurgitation; DLCO, diffusing capacity for carbon monoxide; Echo, echocardiogram; FEV1, forced expiratory volume in 1 second; LA, left atrium; LVH, left ventricular hypertrophy; MR, mitral regurgitation.

What treatment would you offer next?



CT surveillance



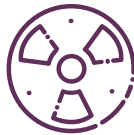
Surgery with RUL lobectomy



Wedge resection RUL of biopsied lesion



Chemotherapy



SABR

CT, computerised tomography; RUL, right upper lobe; SABR, stereotactic ablative radiotherapy.

What would you do next?



1. CT surveillance
- 2. Surgery with RUL lobectomy**
3. Wedge resection RUL of biopsied lesion
4. Chemotherapy
5. SABR

CT, computerised tomography; RUL, right upper lobe; SABR, stereotactic ablative radiotherapy.