LUNG CANCER UPDATE

# The Respiratory Perspective

Dr Emma Ball Respiratory Physician Royal Hobart Hospital



Basics

Referral to clinic

Topics

Diagnosis

Staging

Lung Cancer MDT

## **History and Examination**

Symptoms Cough/SOB Weight loss Haemoptysis Night sweats Chest pain Smoking history

Occupational history

#### Ex:

Clubbing Lymphadenopathy Collapse/Effusion Bony tenderness





# Initial Investigations

CXR CT Chest and upper abdomen with contrast Liver/adrenals/lymphadenopathy for staging Bloods: FBC U+E LFT Ca Coagulation

# Case Study

59 yo male Cough and SOB for 4 months 40 pack yr history Examination: NAD CXR: ?opacity RLL CT Chest: 10mm nodule in RLL WHAT NEXT?

#### Fleischner 2017 Guideline

Solid	Size	Follow up		
	< 6 mm (<100mm <sup>3</sup> )	Single	Low risk High risk	No routine follow Optional CT at 12 months
		Multiple	Low risk High risk	No routine follow Optional CT at 12 months
	6-8 mm (100-250mm <sup>3</sup> )	Single	Low risk High risk	CT at 6-12 mo, then consider CT at 18-24 CT at 6-12 mo, then CT at 18-24
		Multiple	Low risk High risk	CT at 3-6 mo, then consider CT at 18-24 CT at 3-6 mo, then CT at 18-24
	> 8 mm (> 250mm <sup>3</sup> )	Single	All	Consider CT at 3 mo, PET/CT or Biopsy
		Multiple	Low risk High risk	CT at 3-6 mo, then consider CT at 18-24 CT at 3-6 mo, then CT at 18-24

# Referral for work up

Important points for triaging: Age Main symptoms/examination findings E.g. Pain, hypoxia Performance status (ECOG 0-4) Co morbidities Copy of bloods and imaging If concerned please contact Respiratory on call to discuss

# ECOG Performance status

#### 0 Fully active

- 1 Restricted in physical activity but able to carry out work of a light sedentary nature
- 2 Ambulatory and capable of all self care but unable to carry out work activities; up and about for more than 50% of waking hours
- 3 Limited self-care; confined to bed or chair for more than 50% of waking hours
- 4 Completely disabled and unable to carry out any self care

# Clinic

Urgent Seen within two weeks Clinic (consultant or advanced trainee) Repeat history and examination Discuss imaging Discuss diagnostic pathway Lung Function

# **Diagnostic Pathway**

Patient, imaging findings and is a case-by-case decision

## THE THREE HOOPS



## THE THREE HOOPS



# WHAT IS IT: Options for diagnosis?

Bronchoscopy **CT** Guided Biopsy Endobronchial Ultrasound **Pleural aspirate** USS guided Supraclavicular lymph node biopsy USS/Aspirate CT guided biopsy of bone/soft tissue lesions Surgical biopsy Pulmonary nodules increasing in size, PET positive but difficult position to biopsy. On table frozen section

# Which one to choose?

Location of the lesion Central versus peripheral Lung function: FEV1 >1L Co-existing medical disease Location of metastatic disease

# Bronchoscopy



Endobronchial lesions Biopsy, brushings, washings Risk of bleeding Local sedation Determine distance from carina





# Specific Procedures

Biopsy Small pair of forceps Endobronchial lesion

Brushings:

Thin brush on the end of a wire Abnormal mucosa

Endobronchial lesion seen

Bronchial washings

# Bronchoscopy

Day procedure

Fasting

Withhold anticoagulants

Risks

Bleeding

Infection

Cough/Fever

# CT- Guided FNA

High yield(> 80%) if close to pleura Lower yield if more central Not possible in all cases (extensive emphysema) FNA and Core Biopsies Molecular testing Pneumothorax rate up to 20%





# Endobronchial Ultrasound and Biopsy

Bronchoscope with ultrasound

Visualise enlarged lymph nodes

Trans bronchial biopsy

Doppler

Diagnosis and staging of malignancy

?N2 disease especially if surgical candidate





#### Passed sequentially into sub-segmental bronchi until lesion located





## THE THREE HOOPS







#### **Superior Mediastinal Nodes 1** Highest Mediastinal 2 Upper Paratracheal **4** Lower Paratracheal (including Azygos Nodes) N<sub>2</sub>=single digit, ipsilateral **Aortic Nodes**

- 5 Subaortic (A-P window)
- 6 Para-aortic (ascending aorta or phrenic)

#### **Inferior Mediastinal Nodes**

- 7 Subcarinal
- 8 Paraesophageal  $\bigcirc$ (below carina)
  - 9 Pulmonary Ligament

#### N<sub>1</sub> Nodes

- O 10 Hilar
- 11 Interlobar
- 12 Lobar
- 13 Segmental
- 14 Subsegmental

WHERE IS IT: Staging CT PET **EBUS** EUS Mediastinoscopy Pleural tap Biopsy other CT Brain/MRI

- 3 Pre-vascular and Retrotracheal
- N3=single digit, contralateral or supraclavicular

# <u>CT and</u> <u>PET: Nodal</u> disease

# CT Nodal staging:

- Sensitivity 60–65% Specificity 60–70%
- Incorrect staging in 40%

PET:

- PPV 79% NPV 97%
- False positive results are seen in ~ 15% therefore node sampling is still necessary
  - Roberts et al AnnThoracSurg 2000, Toloza Chest 2003



# Lung Function

Calculating predicted post operative pulmonary function using FEV1 and DLCO

>60% low risk

<30% high risk

FEV1 >2L generally tolerate a pneumonectomy

FEV1>1.5L generally tolerate a lobectomy

TNM (Tumour, node, metastases) 8th Edition Lung Cancer

<u>T – P</u>	T – Primary Tumour				
ΤX		Primary tumour cannot be assessed, or tumour proven by the presence of malignant cells in sputum or bronchial washings but not visualized by imaging or bronchoscopy			
T0		No evidence of primary tumour			
Tis		Carcinoma in situ			
T1		Tumour 3 cm or less in greatest dimension, surrounded by lung or visceral pleura, without bronchoscopic evidence of invasion more proximal than the lobar bronchus (i.e., not in the main bronchus) <sup>1</sup>			
	T1mi	Minimally invasive adenocarcinoma <sup>2</sup>			
	T1a	Tumour 1 cm or less in greatest dimension <sup>1</sup>			
	T1b	Tumour more than 1 cm but not more than 2 cm in greatest dimension <sup>1</sup>			
	T1c	Tumour more than 2 cm but not more than 3 cm in greatest dimension <sup>1</sup>			
T2		<ul> <li>Tumour more than 3 cm but not more than 5 cm; or tumour with any of the following features<sup>3</sup></li> <li>Involves main bronchus regardless of distance to the carina, but without involving the carina</li> <li>Invades visceral pleura</li> <li>Associated with atelectasis or obstructive pneumonitis that extends to the hilar region, either involving part of the lung or the entire lung</li> </ul>			
	T2a	Tumour more than 3 cm but not more than 4 cm in greatest dimension			
	T2b	Tumour more than 4 cm but not more than 5 cm in greatest dimension			
T3		Tumour more than 5 cm but not more than 7 cm in greatest dimension or one that directly invades any of the following: chest wall (including superior sulcus tumours), phrenic nerve, parietal pericardium; or associated separate tumour nodule(s) in the same lobe as the primary			
T4		Tumours more than 7 cm or one that invades any of the following: diaphragm, mediastinum, heart, great vessels, trachea, recurrent laryngeal nerve, oesophagus, vertebral body, carina; separate tumour nodule(s) in a different ipsilateral lobe to that of the primary			

#### N – Regional Lymph Nodes

NX	Regional lymph nodes cannot be assessed
NO	No regional lymph node metastasis
N1	Metastasis in ipsilateral peribronchial and/or ipsilateral hilar lymph nodes and intrapulmonary nodes, including involvement by direct extension
N2	Metastasis in ipsilateral mediastinal and/or subcarinal lymph node(s)
N3	Metastasis in contralateral mediastinal, contralateral hilar, ipsilateral or contralateral scalene or supraclavicular lymph node(s)

No regional lymph node metastases



Metastasis in ipsilateral intrapulmonary/ peribronchial/ hilar lymph node(s), including nodal involvement by direct extension

N2

Metastasis in ipsilateral mediastinal and/or subcarinal lymph node(s), including "skip" metastasis without N1 involvement



Metastasis in ipsilateral mediastinal and/or subcarinal lymph node(s) associated with N1 disease

#### M- Distant Metastasis

MO		No distant metastasis
M1		Distant metastasis
	M1a	Separate tumour nodule(s) in a contralateral lobe; tumour with pleural or pericardial nodules or malignant pleural or pericardial effusion <sup>4</sup>
	M1b	Single extrathoracic metastasis in a single organ ⁵
	M1c	Multiple extrathoracic metastases in one or several
		organs





## THE THREE HOOPS



WHAT CAN WE DO: Lung Cancer MDT

Respiratory, Medical Oncology, Radiation Oncology, Surgeons, Pathology, Radiology including Nuclear Medicine and Palliative care plus supporting nursing staff.

Weekly meeting

New cases, surgical pathology specimens, review imaging of patients undergoing treatment, recurrence.

All patients, regardless of their disease stage, have access to all relevant treatment and supportive care options

Patients are then fully informed of their treatment choices

# New Patients

History Radiology Pathology Confirm Stage ECOG/PFTs

Management Discussion

Follow up in Respiratory clinic with the results of the MDT and then referral on to the appropriate specialist for their ongoing management.



**Clinical Details:** 69yo LUL nodule picked up on skeletal survey by haem for MGUS. Ex-smoker 30 pack year. FEVI 1.97L (68%), DLCO 62%. X2 left sided pneumothoraces requiring ICC insertion 2019. ECOG 0.

Please review:

Radiology: CT Chest 7/1/21 Rad Tas, CTB pending, PET RHH

Pathology: Bronchoscopy biopsy and brushings 12/2/21

 $\label{eq:Radiology: CT chest scan at Rad Tas on 7^{th} January shows 21mm spiculated LUL nodule. Background centrilobular and paraseptal COPD changes and a 10mm 11L lymph node.$ 

PET scan at RHH on 23<sup>rd</sup> February shows LUL lesion and 11L lymph node are both intensely avid. Bilateral hilar lymph node mild uptake appears inflammatory.

Pathology: Bronchoscopy biopsy is consistent with adenocarcinoma, TTF-1 positive, ROS and ALK negative. There is insufficient tissue for PDL1 and NGS studies. Stage: cT1cN1

ECOG: 0

Management Suggestions:

1. Referral to cardiothoracics for consideration of LUL lobectomy.

Please note this is a summary for your records and no further action is required. If there are any queries about the above please discuss with the lead consultant.

Yours sincerely

Edited but not signed

Chair, TOMM

# Questions?