

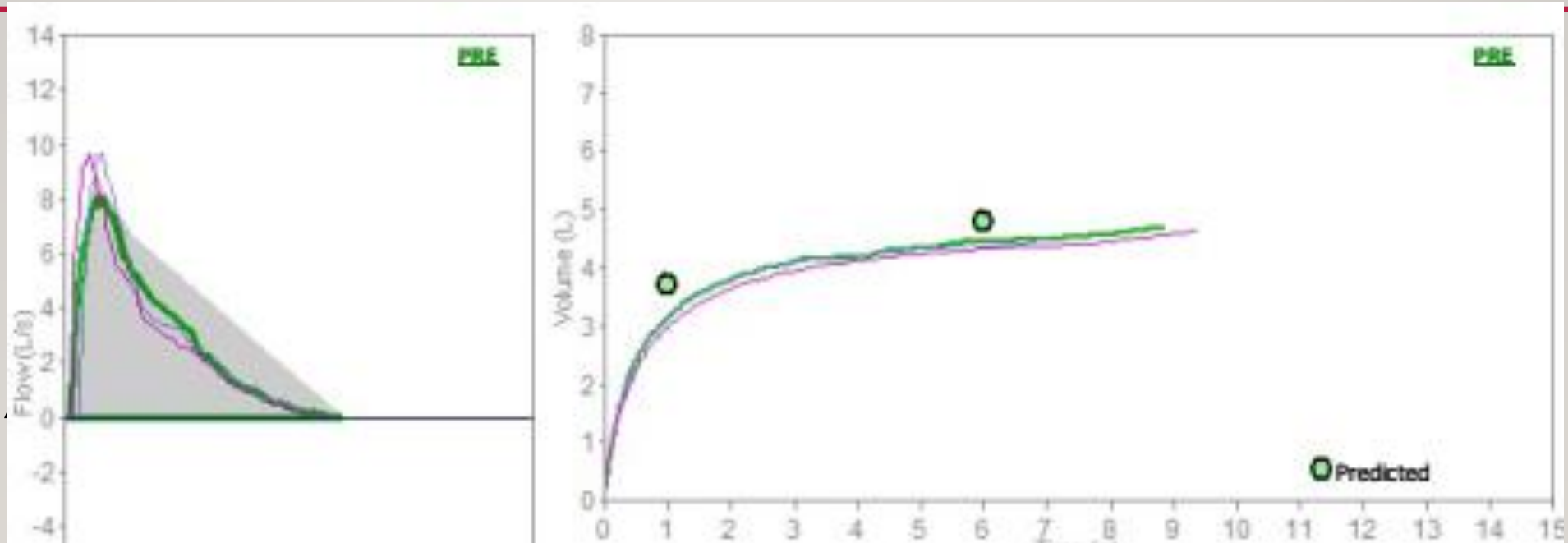
# RESPIRATORY CASES

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25<sup>TH</sup> JULY 2022

DONNY WONG

# 60 YEAR MALE OLD EX SMOKER (30 PACK)



- FEV1/FVC of 3.69/4.72 (85/99%) ratio 67%

**What's the diagnosis?**

# FOLLOW UP TESTS

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- HRCT (chest): Diffuse bronchial wall thickening and functional small airways disease
- Bloods, eosinophilia 0.7, IgE 110, Aspergillus serology unremarkable. No skin prick test
- Formal lung functions, FEV1/FVC of 3.23/4.83 (86/100%), ratio 67%. TLC 7.6 (103%), DLCO 103%, FeNO 53/54
- CT sinus: Variable pansinusitis, maximal through right maxillary ethmoid and frontal sinus regions. Extensive polyposis about the middle turbinates. Left mastoid opacification medially and bilateral earwax.
- **?Asthma with fixed airflow limitation**
  - So changed to Qvar with higher dose ICS 100 mcg 3 puffs bd.
- Referred to ENT after script for maximal sinusitis treatment

# FOLLOW UP

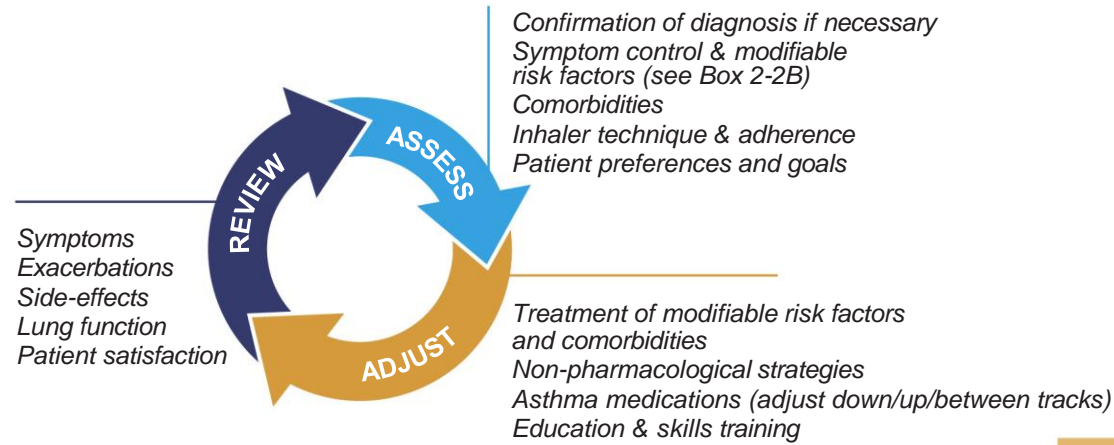
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- 4/12 later sinus surgery
- Improved post surgery only 1x prn Qvar a month
- Reviewed 2/12 post surgery spirometry normal
  - FEV1/FVC is 4.60/6.02 (125/126%) with ratio 76%. Peak flow 650L/min.
- Changed to Symbicort Smart

# Adults & adolescents 12+ years

## Personalized asthma management

Assess, Adjust, Review  
for individual patient needs



**CONTROLLER** and **PREFERRED RELIEVER** (Track 1). Using ICS-formoterol as reliever reduces the risk of exacerbations compared with using a SABA reliever

<b>STEPS 1 – 2</b> As-needed low dose ICS-formoterol	<b>STEP 3</b> Low dose maintenance ICS-formoterol	<b>STEP 4</b> Medium dose maintenance ICS-formoterol	<b>STEP 5</b> Add-on LAMA Refer for assessment of phenotype. Consider high dose maintenance ICS-formoterol, ± anti-IgE, anti-IL5/5R, anti-IL4R, anti-TSLP
RELIEVER: As-needed low-dose ICS-formoterol			

See GINA severe asthma guide

**CONTROLLER** and **ALTERNATIVE RELIEVER** (Track 2). Before considering a regimen with SABA reliever, check if the patient is likely to be adherent with daily controller

<b>STEP 1</b> Take ICS whenever SABA taken	<b>STEP 2</b> Low dose maintenance ICS	<b>STEP 3</b> Low dose maintenance ICS-LABA	<b>STEP 4</b> Medium/high dose maintenance ICS-LABA	<b>STEP 5</b> Add-on LAMA Refer for assessment of phenotype. Consider high dose maintenance ICS-LABA, ± anti-IgE, anti-IL5/5R, anti-IL4R, anti-TSLP
RELIEVER: As-needed short-acting beta <sub>2</sub> -agonist				

Other controller options for either track (limited indications, or less evidence for efficacy or safety)

	Low dose ICS whenever SABA taken, or daily LTRA, or add HDM SLIT	Medium dose ICS, or add LTRA, or add HDM SLIT	Add LAMA or LTRA or HDM SLIT, or switch to high dose ICS	Add azithromycin (adults) or LTRA. As last resort consider adding low dose OCS but consider side-effects
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# MILD ASTHMA

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- Even apparently mild asthma are still at risk:
  - adults with acute asthma: 30–37%
  - near-fatal asthma: 16%
  - adults dying of asthma: 15–27%
- Triggers are unpredictable
- 4–5 steroids courses can increase risk of osteoporosis, DM, cataract

# Investigate and manage difficult-to-treat asthma in adults and adolescents

Consider referring to specialist or severe asthma clinic at any stage

DIAGNOSIS:  
"Difficult-to-treat asthma"

**1 Confirm the diagnosis**  
(asthma/differential diagnoses)

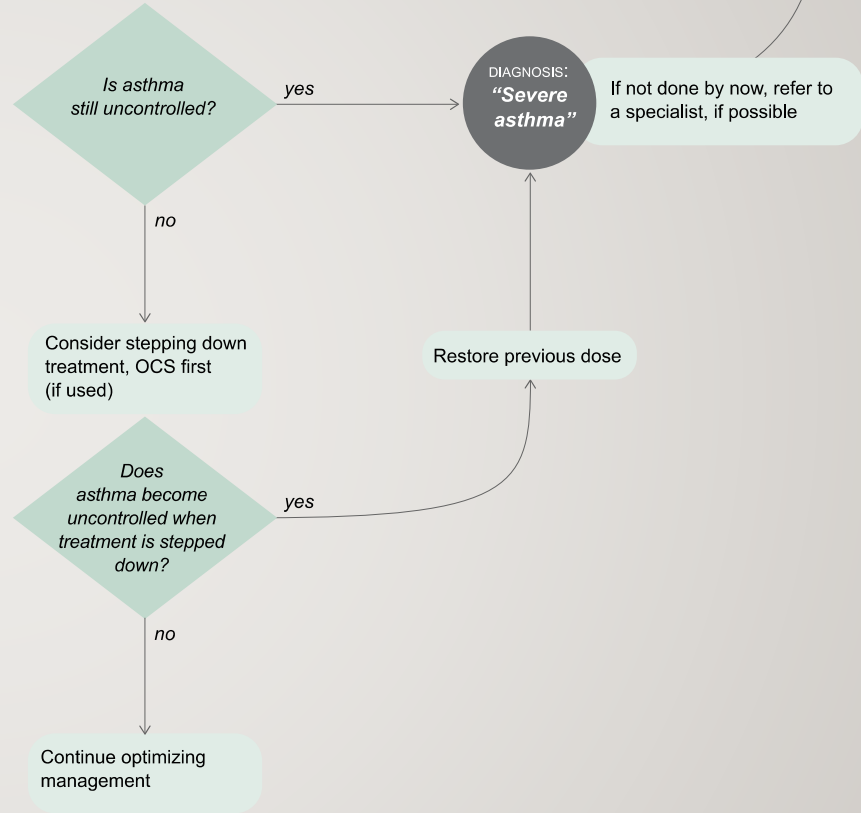
**2 Look for factors contributing to symptoms, exacerbations and poor quality of life:**

- Incorrect inhaler technique
- Suboptimal adherence
- Comorbidities including obesity, GERD, chronic rhinosinusitis, OSA
- Modifiable risk factors and triggers at home or work, including smoking, environmental exposures, allergen exposure (if sensitized); medications such as beta-blockers and NSAIDs
- Overuse of SABA relievers
- Medication side effects
- Anxiety, depression and social difficulties

**3 Optimize management, including:**

- Asthma education
- Optimize treatment (e.g. check and correct inhaler technique and adherence; switch to ICS-formoterol maintenance and reliever therapy, if available)
- Consider non-pharmacological interventions (e.g. smoking cessation, exercise, weight loss, mucus clearance, influenza and COVID-19 vaccination)
- Treat comorbidities and modifiable risk factors
- Consider non-biologic add-on therapy (e.g. LABA, LAMA, LM/LTRA, if not used)
- Consider trial of high dose ICS-LABA, if not used

**4 Review response after ~3-6 months**



**Key**

- decision, filters
- intervention, treatment
- diagnosis, confirmation

For adolescents and adults with symptoms and/or exacerbations despite medium or high dose ICS-LABA, or taking maintenance OCS

# KEY MESSAGES

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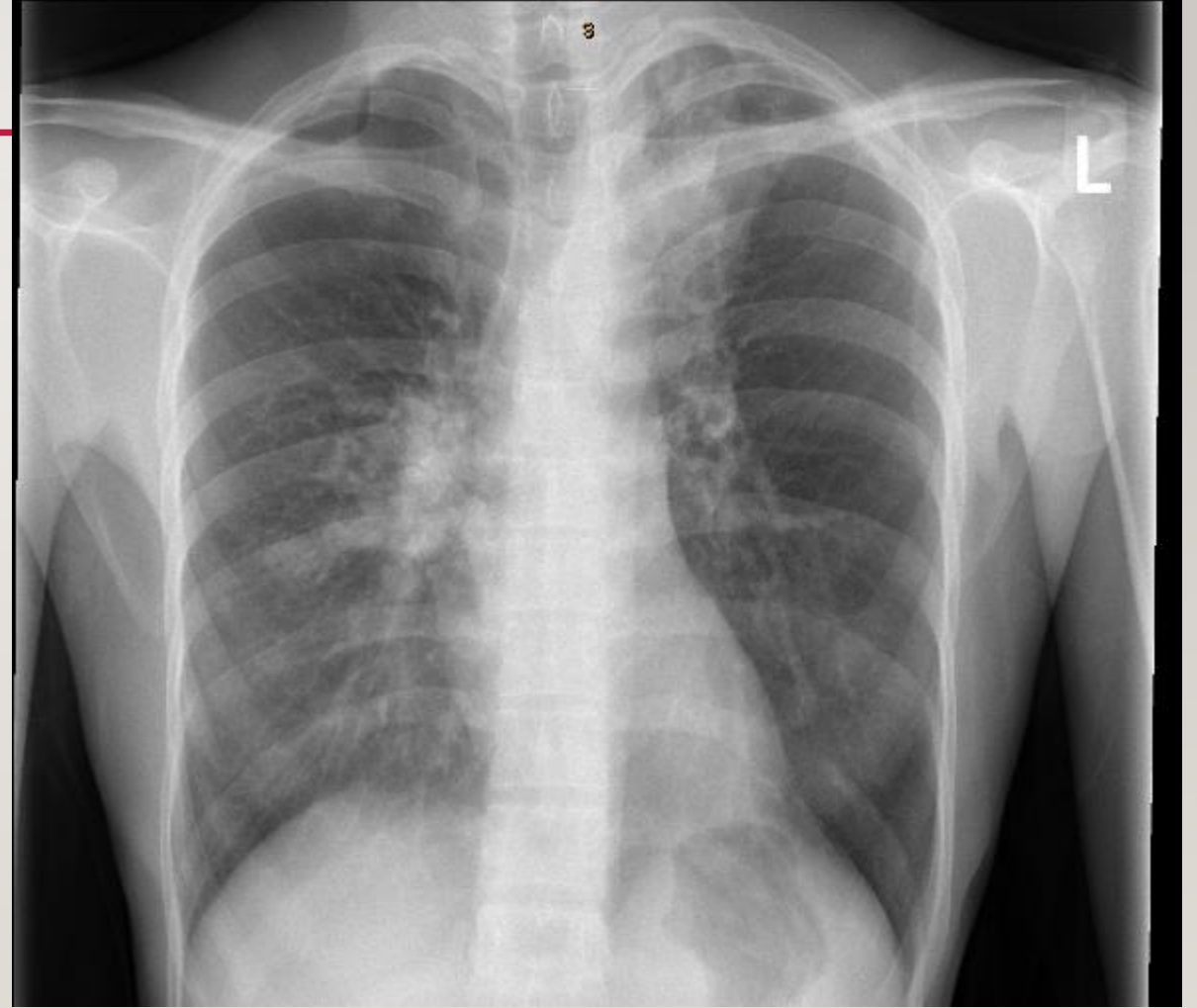
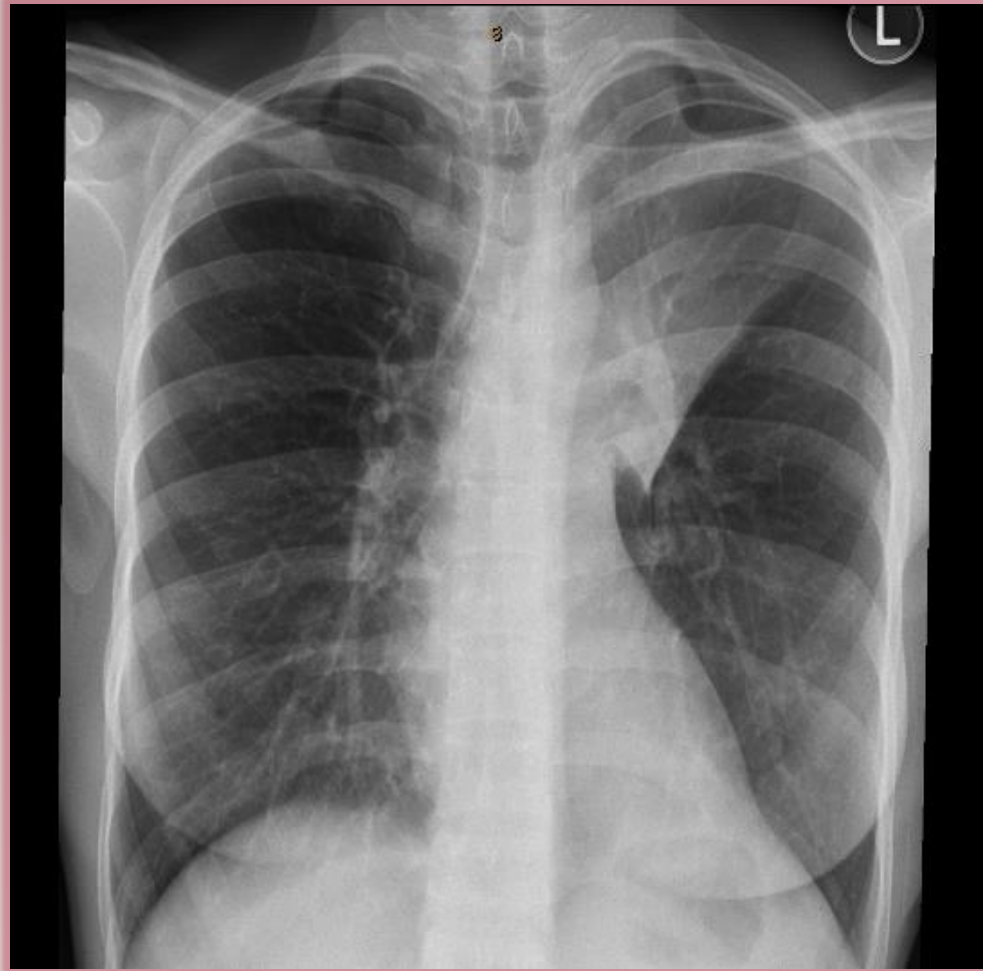
- Symbicort SMART is preferable
  - Same reduction of exacerbation as daily maintenance ICS
  - Less hospitalisations and ED visits cf. daily maintenance
- Mild asthma can be bad
- GINA would like spirometry prior diagnosis, NZ guidelines more pragmatic



# 31 YEAR YO FEMALE

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- Presented to me after routine CXR.
- Asymptomatic and never smoker
- 2019, LUL collapse ?endobronchial lesion, bronch normal and no Tb found
- On hindsight previous bloods showed, eosinophilia up to 2.25



# CURRENT WORK UP

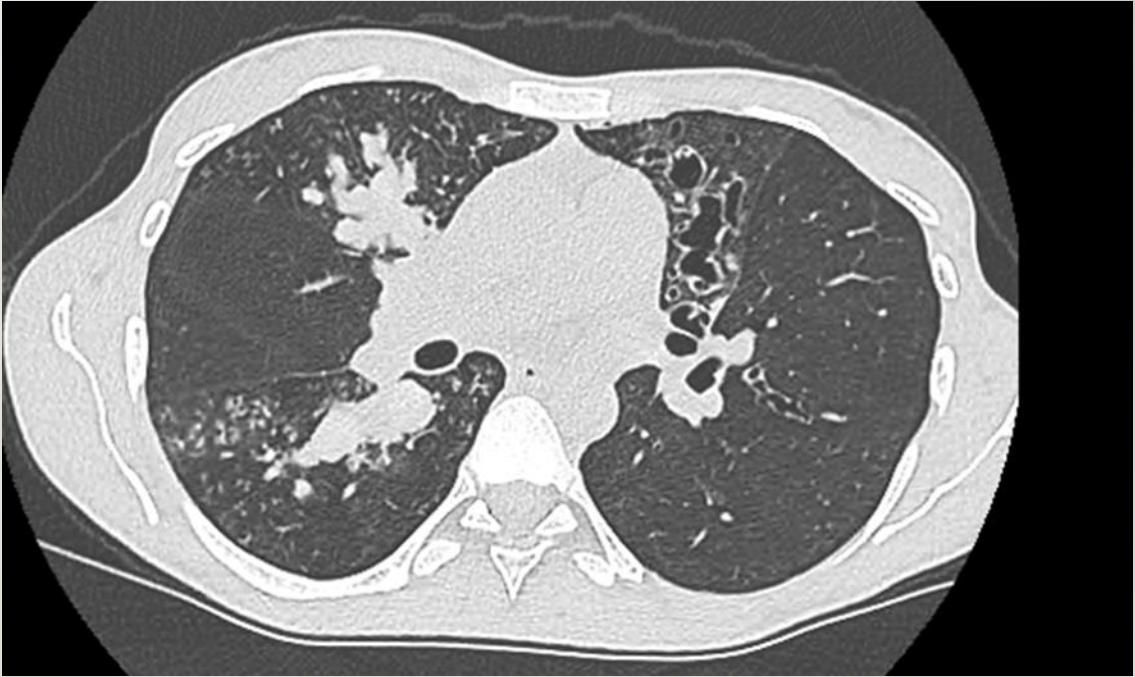
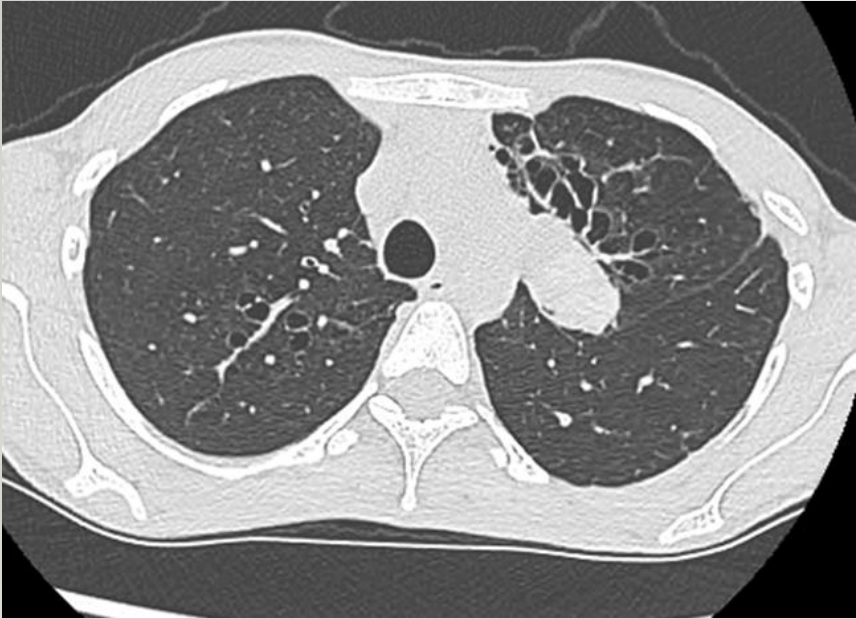
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- IgE >2500, Eosinophils
- SPT only positive to aspergillus
- Aspergillus serology RAST 4+, Precipitins >200
- Spirometry not done.
- CT done, bronchiectasis with mucus plugs



# CT SCAN

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# ALLERGIC BRONCHOPULMONARY ASPERGILLOSIS (ABPA)

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## DIAGNOSIS

- ?Asthma
- IgE
- Aspergillus RAST/SPT
- Aspergillus precipitins
- CT findings
- Eosinophilia

## TREATMENT

- Prednisone mainstay for months
- No antifungal role first
- Monitoring IgE and eosinophils
- ICS maybe enough.



# ABPA

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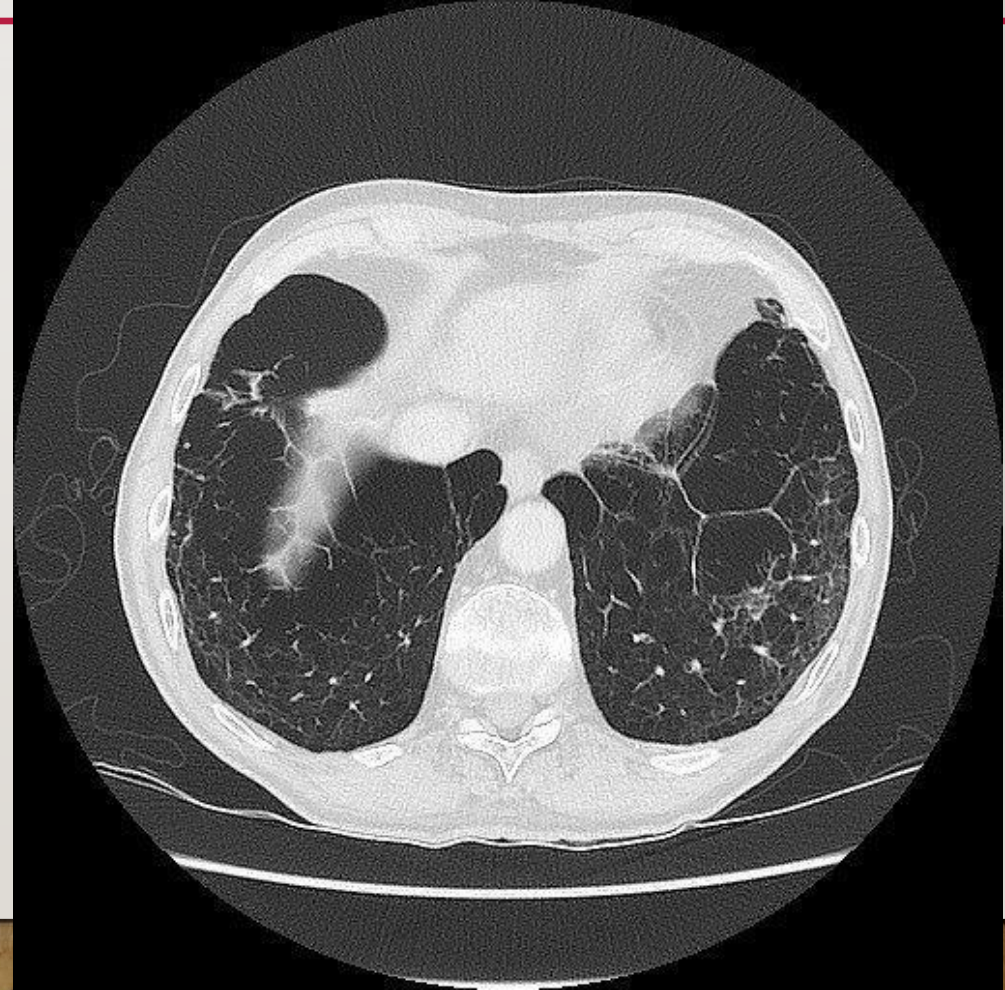
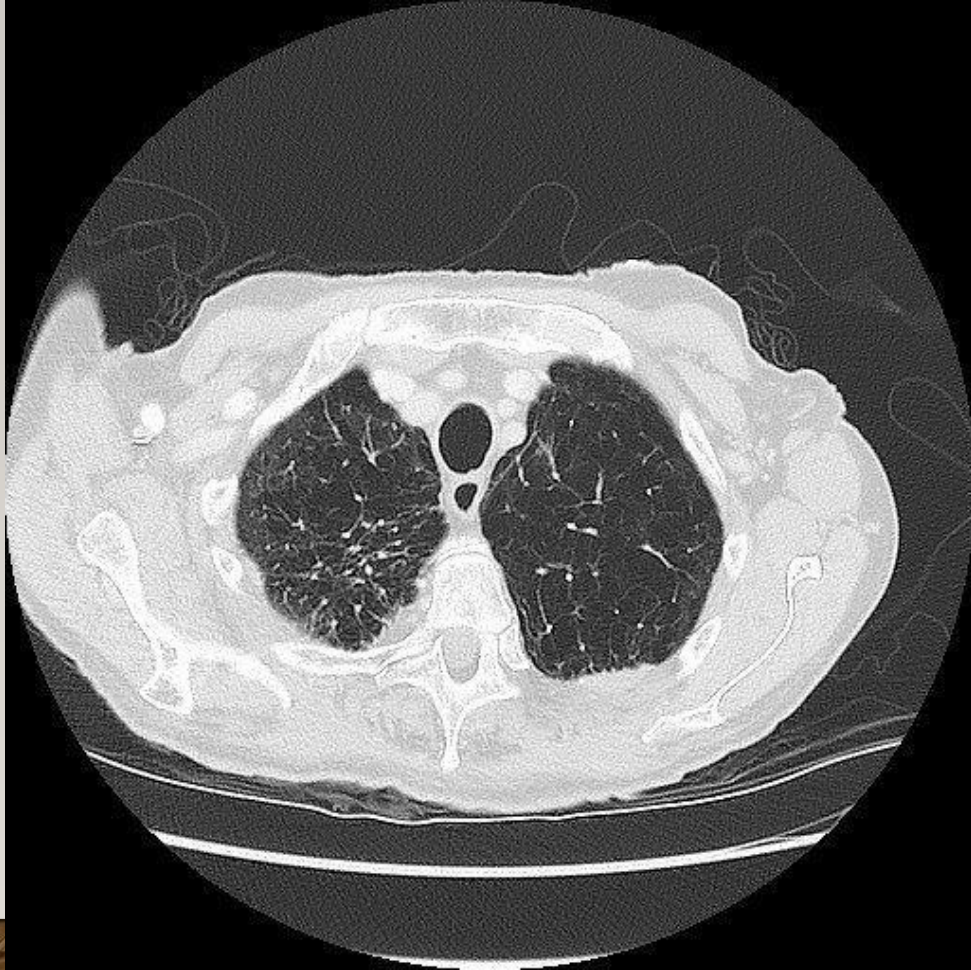
- Bad case of extreme allergy to an allergen in this case aspergillus
- Aspergillus is an ubiquitous fungi
- In a spectrum of aspergillus disease but not an infection per se.
- Monitored by IgE and Eosinophils

# 77 YEAR OLD MALE

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- First saw me in 2019 with exacerbation of COPD with RSV
- FEV1 in hospital showed 0.47 L, 13% predicted, FVC 1.93 (39%)
- LUL lesion ?cancer, CT diffuse emphysema and bullous disease.
- Opted for review in clinic with repeat spirometry but declined FU.
- Further yearly exacerbations, CT chest showed resolution
- AF anticoagulated

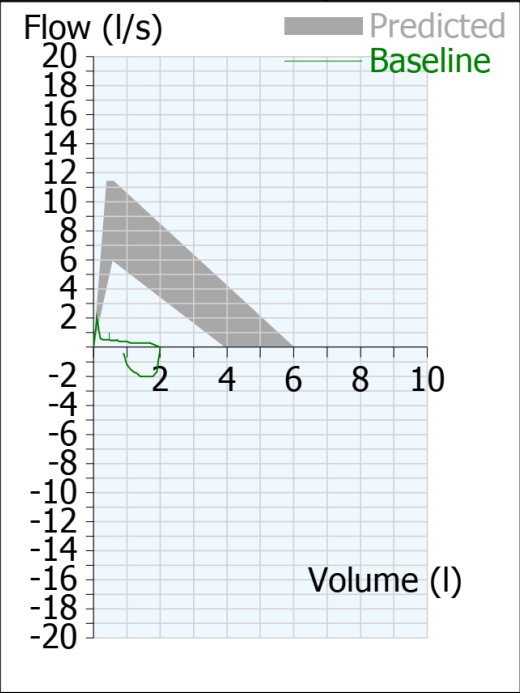
CT



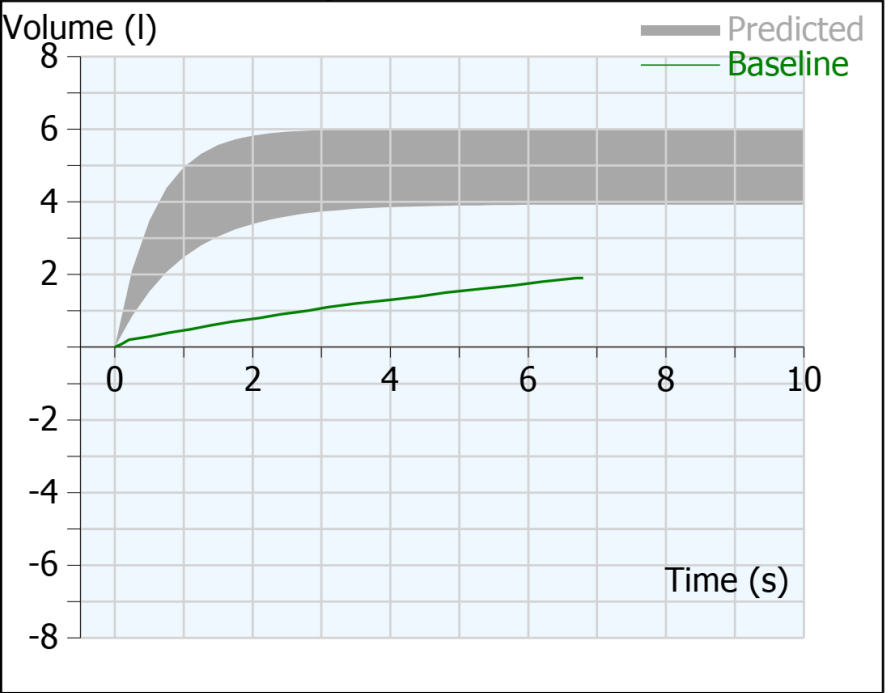


# SPIRO

Best Flow Volume Graph



Best Volume Time Graph



# CURRENT STATUS

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- Latest Nov 21 exacerbation discharge on home short term oxygen.
- Declined LTOT assessment clinic 6 weeks post discharge opted for telehealth
- Doing okay independent ADLS though ?help required
- No exacerbations
- ?Mild leg swelling
  
- **What to Do?**



# MANAGEMENT

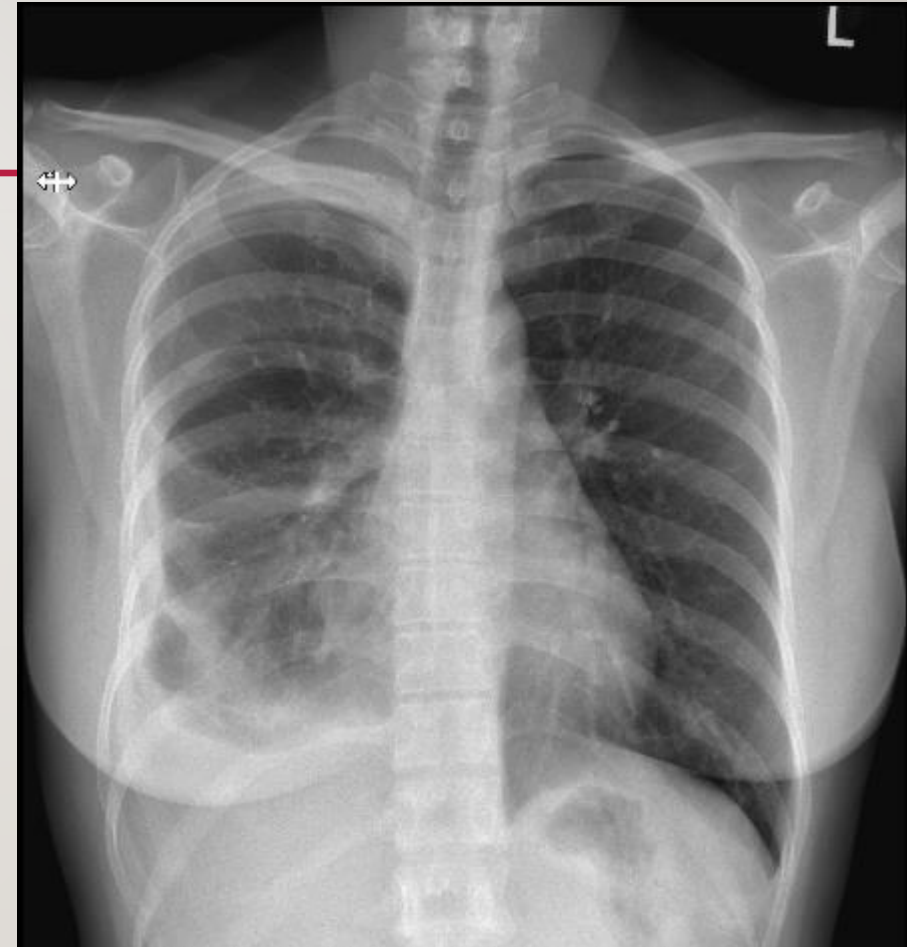
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- Already been referred to Pulmonary rehab, declined to attend
- PT already referred to hospice
- Offered NASC referral, also decline
- Ideally in person to assess for LTOT and spirometry
- Assess for LTOT with cor pulmonale/polycythemia , ie when sats 92% or less
  - Criteria PaO<sub>2</sub> <8.0 vs less than 7.3
- Already on triple inhaler
  - assess eosinophilia (0.3) ?stop ICS.

# 34 YO FEMALE

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- 3-4 kg weight loss with cough over a month
- CXR shows moderate R) effusion
- Thoraconcentesis completed
  - Adenosine deaminase 37, Lymphocytic exudate
  
- Any risk of infection?



# PROGRESS

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- TB PCR positive on pleural fluid, but culture even after 8 weeks negative.
- 3 Induced sputa done, all smear positive and 1x culture positive at 4 weeks
- Started RHEZ and progressing appropriately

# TB FACTS FROM LATEST 2018

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- 307 cases a year, 80% are migrants; 80.5% with symptoms to doctor, 10.8% immigration screening, 4.7% contact tracing
  - Counties Manukau (11.8 per 100,000), Capital & Coast (10.1 per 100,000) and Waitemata (8.3 per 100,000) DHBs
  - Asian ethnic group (26.6 per 100,000), followed by MELAA (18.2 per 100,000) and Pacific peoples (11.8 per 100,000)
- Pulmonary 56.2% and over the years (2014-2018) 16.2% pleural