# PRESCRIBING ANTIBIOTICS

Jos WM van der Meer Emeritus professor of Medicine Radboud umc Nijmegen NL

# **Conflicts of interest**

none



- Arrives on Saturday night at the ER because of *high fever and cough*
- Previous history: Hypertension (R/ACE inh)
- Yearly Flu vaccine, recent Pneumovax and COVID Pfizer vaccine booster March 2022
- Smokes 10 cigars per day
- 6 units alcohol per day
- No previous antibiotic exposure
- No exposure to Livestock/animals

- No problems until this morning
- After breakfast chills and rigors, temperature rise >39.9°
- Cough, no sputum
- Slight R-sided chest pain
- Feels ill
- Took paracetamol 500 mg
- Covid-19 antigen self test negative.
- Further history not remarkable

- Physical exam:
  - Dyspnea & tachypnea (32/min)
  - BP 88/58, pulse rate 100
  - Temp 39.5°
  - Small herpes labialis lesion
  - Dull percussion over right lower part of thorax
  - Ausc: Bronchial breathing with crepitations



Lab: **CRP 80** Leukocytes 13.8 Hb 8 mmol/l creat 100 µmol/l BUN 6 mmol/l **Bilirubin slightly** elevated Sodium 129 mmol/l pH 7.2; PO<sub>2</sub> 8.1 kP Radboudumc

What is your differential diagnosis now?



- What is your differential diagnosis now?
  - CAP
    - Lobar pneumonia probably pneumococcal
    - Other causes?
      - Legionella (male, smoker, alcohol, hotel)
      - Aspiration pneumonia (male, smoker, alcohol)
      - Other causes of pneumonia unlikely (Staph/Gramnegative/Mycoplasma/Qfever/Chlamydia psittaci or C. pneumoniae/ viral\*)

\* SARS CoV2?  $\rightarrow$  antigen test and PCR negative

What is your assessment of severity?



# CURB-65

- Confusion
- BUN >7 mmol/l
- Resp rate  $\geq$  30
- Syst BP < 90 or</li>
   Diast BP ≤ 60
- Age ≥ 65

- No: 0 Yes: +1

# CURB-65

- Confusion
- BUN >7 mmol/l
- Resp rate  $\geq$  30
- Syst BP < 90 or</li>
   Diast BP ≤ 60
- Age ≥ 65

- No: 0 Yes: +1

# CURB-65

- Confusion
- BUN >7 mmol/l
- Resp rate  $\geq$  30
- Syst BP < 90 or</li>
   Diast BP ≤ 60
- Age ≥ 65

- No: 0 Yes: +1

Radboudumc

3 points: 14% 30-day mortality  $\rightarrow$  ward admission

• What about antibiotics?



- What about antibiotics?
  - Follow the guidelines...
    - Which guidelines?
  - Think!
  - (or do both!)

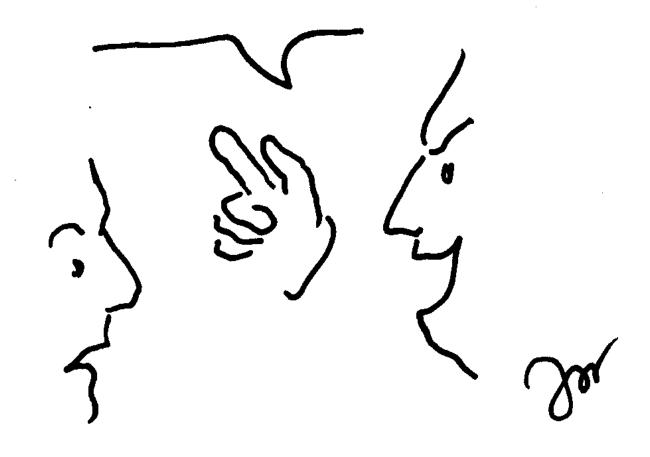


#### This is not what I mean:



This is not what I mean either:

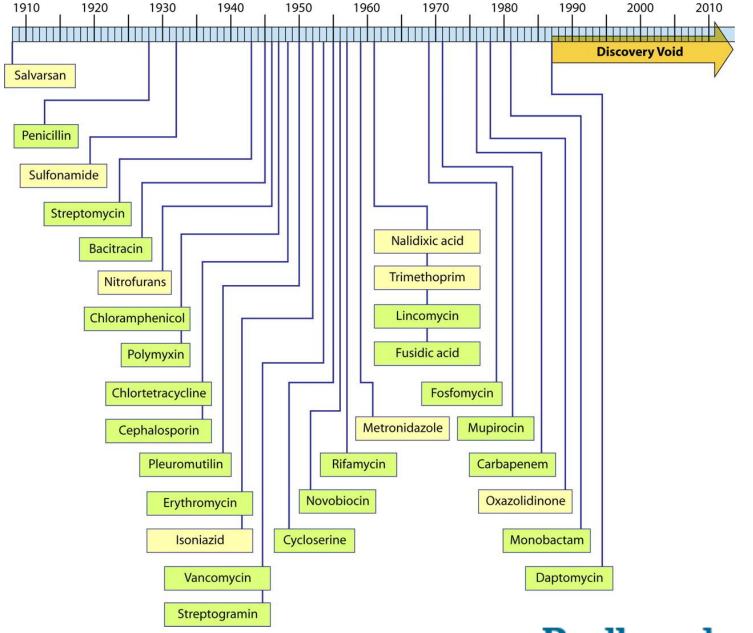
### Guidelines are for beginners!



# **Prescribing antibiotics**

- Is there an infection?
- If so, is it likely to be bacterial?
- If so, which bacterium?
- If so, is antibiotic treatment beneficial?\*

\* e.g., acute bronchitis, acute otitis media, sinusitis → antibiotics rarely indicated



#### The preferred antibiotic:

- Is effective against the (presumed) causative bacterium
- Attains effective concentrations at the site of infection
- Does lead to the least development of antimicrobial resistance
  - Of the causative micro-organism
  - Of the colonizing microflora
  - Of the micro-organisms in the environment
- Is (relatively) non toxic
- Can be administered in the most preferable way
- Is not more expensive than an equivalent

The preferred antibiotic:

• Is effective against the (presumed) causative bacterium

How do we know this?



#### The preferred antibiotic:

• Is effective against the (presumed) causative bacterium

#### How do we know this?

- 1. In-vitro susceptibility studies
  - MIC
  - Time-kill curves
- 2. In-vivo studies
  - In experimental animals
  - In patients

# In-vitro susceptibility studies

Minimal inhibitory concentration

- Is a better indicator for resistance than for susceptibility
- performs better to compare the effect of one antibiotic on different bacteria (pneumococci are more sensitive than enterococci to benzylpenicillin) than

the effect of different antibiotics to the same bacterium (pneumococci are more sensitive to benzylpenicillin than to amoxicillin)

 The translation to the in-vivo situation is difficult, because:

# In-vitro susceptibility studies

Minimal inhibitory concentration

- The in-vitro conditions differ greatly from in vivo:
  - Culture medium does not resemble the conditions in tissues and body fluids
  - The bacterium is exposed to a constant antibiotic concentration in vitro
  - Overnight incubation is not relevant in vivo
  - Protein binding is often not taken into account
  - Judgement of the in-vitro effect is visible growth
- Direct quantitive translation of MIC to in-vivo concentrations is unscientific.

#### The preferred antibiotic:

• Is effective against the (presumed) causative bacterium

#### How do we know this?

- 1. In-vitro susceptibility studies
  - MIC
  - Time-kill curves
- 2. In-vivo studies
  - In experimental animals
  - In patients

**Teach us about pharmacokinetics:** 

- Time dependent killing
- Concentration-dependent killing

The preferred antibiotic:

• Is effective against the (presumed) causative bacterium

So we need good RCTs to know the real effectiveness of antibiotics!

Note of caution:

The older antibiotics have not been tested in modern-type RCTs! Most RCTs have been performed with quinolones...

So there is a strong bias with regard to type A evidence!

#### The preferred antibiotic:

- Is effective against the (presumed) causative bacterium
- Attains effective concentrations at the site of infection
- Does lead to the least development of antimicrobial resistance
  - Of the causative micro-organism
  - Of the colonizing microflora
  - Of the micro-organisms in the environment
- Is (relatively) non toxic
- Can be administered in the most preferable way
- Is not more expensive than an equivalent

The preferred antibiotic:

- Is effective against the (presumed) causative bacterium
- Attains effective concentrations at the site of infection For most infections, this is not a problem. It is a problem for sites such as:
  - Prostatitis
  - Eye
  - Meningeal space and brain and for
  - abscesses
  - foreign bodies (such as prostheses)
  - empyema
  - bone sequesters

Here again time-dependent versus dose dependent killing is important!

#### The preferred antibiotic:

- Is effective against the (presumed) causative bacterium
- Attains effective concentrations at the site of infection
- Does lead to the least development of antimicrobial resistance
  - Of the causative micro-organism
  - Of the colonizing microflora
  - Of the micro-organisms in the environment
- Is (relatively) non toxic
- Can be administered in the most preferable way
- Is not more expensive than an equivalent

#### **Development of antimicrobial resistance**

- Various mechanisms
- Antibiotics vary in their capacity to induce resistance
- Microorganisms vary in their capacity to become resistant

- Of the causative micro-organism → rare
- Of the colonizing microflora → dependent on the effect of the antibiotic on commensals
- Of the micro-organisms in the environment

   → dependent on (urinary) excretion of the drug

and its stability

#### The preferred antibiotic:

- Is effective against the (presumed) causative bacterium
- Attains effective concentrations at the site of infection
- Does lead to the least development of antimicrobial resistance
  - Of the causative micro-organism
  - Of the colonizing microflora
  - Of the micro-organisms in the environment
- Is (relatively) non toxic
- Can be administered in the most preferable way
- Is not more expensive than an equivalent

# Toxicity

- Allergy: especially penicillins
- Dose-dependent toxicity:
  - Aminoglycosides
  - Polymyxins
  - Quinolones



#### The preferred antibiotic:

- Is effective against the (president of the section of the sec
- Attains effective concentration
- Does lead to the least develop
  - Of the causative micro-organisn
  - Of the colonizing microflora
  - Of the micro-organisms in the er
- Is (relatively) non toxic
- Can be administered in the most preferable way
- Is not more expensive than an equivalent



- What is your differential diagnosis now?
  - CAP
    - Lobar pneumonia probably pneumococcal
    - Other causes?
      - Legionella (male, smoker, alcohol, hotel)
      - Aspiration pneumonia (male, smoker, alcohol)
      - Other causes of pneumonia unlikely (Staph/Gramnegative/Mycoplasma/Qfever/Chlamydia psittaci or C. pneumoniae/ viral)
- Classification

Curb-65 score +3

# What do guidelines say?

#### Dutch SWAB guideline

#### - For CURB-65 score +3:

For patients with risk category III (severe CAP – ward admission; CURB-65: 3-5; PSI: 5; hospitalized on non-ICU ward) therapy should be started with a 2nd or 3rd generation cephalosporin. No empiric coverage for atypical microorganisms is given. A Legionella and pneumococcal urinary antigen test should be carried out as a routine procedure within 12-24 hours of admission.

So let us choose for ceftriaxon...

#### The preferred antibiotic: $\rightarrow$ ceftriaxon

Assuming pneumococcus/aspiration pneumonia

- Is effective against the (presumed) causative bacteria  $\checkmark$
- Attains effective concentrations at the site of infection  $\sqrt{}$
- Does lead to the least development of antimicrobial resistance
  - Of the causative micro-organism  $\sqrt{}$
  - Of the colonizing microflora
  - Of the micro-organisms in the environment  $\checkmark$
- Is (relatively) non toxic  $\sqrt{}$

- What is your differential diagnosis now?
  - CAP
    - Lobar pneumonia probably pneumococcal
    - Other causes?
      - Legionella (male, smoker, alcohol, hotel)
      - Aspiration pneumonia (male, smoker, alcohol)
      - Other causes of pneumonia unlikely (Staph/Gramnegative/Mycoplasma/Qfever/Chlamydia psittaci or C. pneumoniae/ viral)
- Classification

Curb-65 score +3

- What is your differential diagnosis now?
  - CAP
    - Lobar pneumonia probably pneumococcal
    - Other causes?
      - Legionella (male, smoke
      - Aspiration pneumonia (r
      - Other causes of pneumoniae/ negative/Mycoplasma/Q<sup>i</sup>
         pneumoniae/ viral)
- Classification

Curb-65 score +3

But this is classical Pneumococcal pneumonia!!



- We decide to start treatment with ceftriaxon and ciprofloxacin
- The next day the pneumococcal antigen test in urine is positive, the legionella antigen test is negative.
- The temperature has normalised

So what is the next step?



# Streamline!



- We decide to start treatment with ceftriaxon and ciprofloxacin
- The next day the pneumococcal antigen test in urine is positive, the legionella antigen test is negative.
- The temperature has normalised

#### So what is the next step?

#### Stop ciprofloxacin

Switch to oral amoxicillin (5 days)

Culture result sputum (day 2): penicillin-sensitive
 Pneumococcus