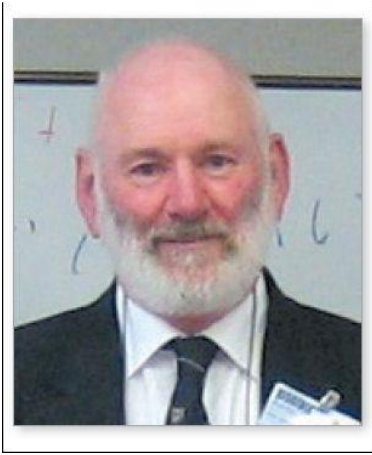




Dr Martin Reeve

Medical Officer

Communicable Diseases

Auckland Regional Public Health Service

Saturday, August 10, 2019

(Plenary)

15:20 - 15:40 TB in NZ - The Battle Rages On

TB IN NEW ZEALAND

Dr Martin Reeve Auckland Regional Public Health
Service

Auckland Regional Public Health Service

Rātonga Hauora ā Iwi o Tamaki Makaurau



Waitemata
District Health Board
Best Care for Everyone



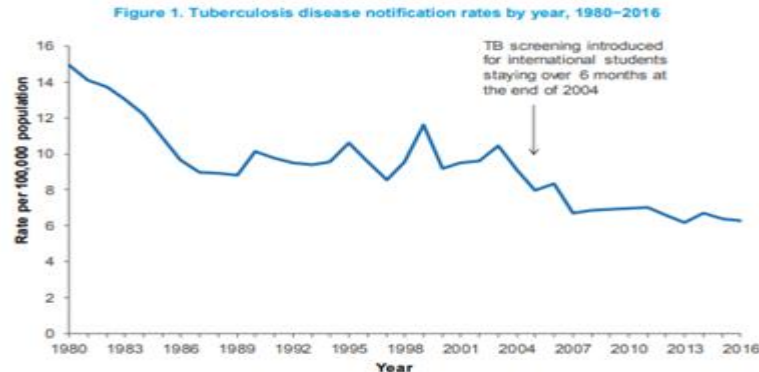
Working with the people of Auckland, Waitemata and Counties Manukau

TREATMENT OF ACTIVE TB

- Highly specialised
- Strict international guidelines
- Minimum 6 month course for simple TB, minimum two year course for multi drug resistant TB
- Start with 4 drugs, two months, then four months with two
- Isoniazid, Rifampicin, Pyrazinamide, Ethambutol
- P – DOT, E- DOT

HOW MUCH TB IN NEW ZEALAND

- About 300 new cases each year
- Rate 6/100, 000, comparable to other countries
- Rate dropping, but slowing
- Many of you will never see a case, but 85% found by GP's



Note: Census population data was used as the denominator to calculate rates before 1991 and the Statistics New Zealand mid-year population estimates were used from 1991 onwards.

WHO GETS TB?

Table 3. Numbers and rates of tuberculosis notifications (new case) by age group and sex, 2016

Age group (years)	Male		Female		Total	
	Cases	Rate ^a	Cases	Rate ^a	Cases	Rate ^a
<5	4	2.6	2	1.4	6	2.0
5–14	4	1.3	4	1.3	8	1.3
15–39	86	10.8	74	9.4	160	10.1
40–59	21	3.5	32	5.0	53	4.3
≥60	33	7.3	22	4.3	55	5.8
Total	148	6.4	134	5.6	282	6.0

^a Rate per 100,000 based on 2016 mid-year population estimates; caution as rates shown for counts with less than 5 cases

Table 5. Risk factors reported for tuberculosis (new case) notifications, 2016

Risk factor	Cases ^a	Total ^b	%
Born outside New Zealand	224	282	79.4
Current/recent residence with person born outside New Zealand	183	255	71.8
Contact with confirmed case	72	248	29.0
Has immunosuppressive illness	40	269	14.9
Exposure in a healthcare setting	22	264	8.3
On immunosuppressive medication	15	274	5.5
Current/recent residence in an institution	5	273	1.8

^a Number of cases with 'yes' recorded for the risk factor.

^b Number of cases for which information was recorded for the risk factor. Cases can have multiple risk factors.

WHERE IN THE BODY DOES IT OCCUR?

- Born in New Zealand – 75% pulmonary, 25 % not
- Not born in New Zealand – 50% pulmonary, 50% not
- Extrapulmonary. Lymphatic, pleural, intra-abdominal not renal

WHAT CAUSES TB?

- 97% Mycobacterium TB, 3% M Bovis
- M Bovis first herd in New Zealand, now spread to possums
- Environmental mycobacteria AKA non-tuberculous mycobacteria AKA atypical mycobacteria occasionally cause TB like diseases, but they are not infectious.

HOW IS TB SPREAD?

The three I's

- Inhalation – most important by far
 - Ingestion – unpasteurised milk
 - Inoculation – meat workers, possum trappers, BCG vaccination
-
- Extrapulmonary TB is not considered infectious – needs no contact tracing.

TWO PHASES OF TB – BIPHASIC ILLNESS

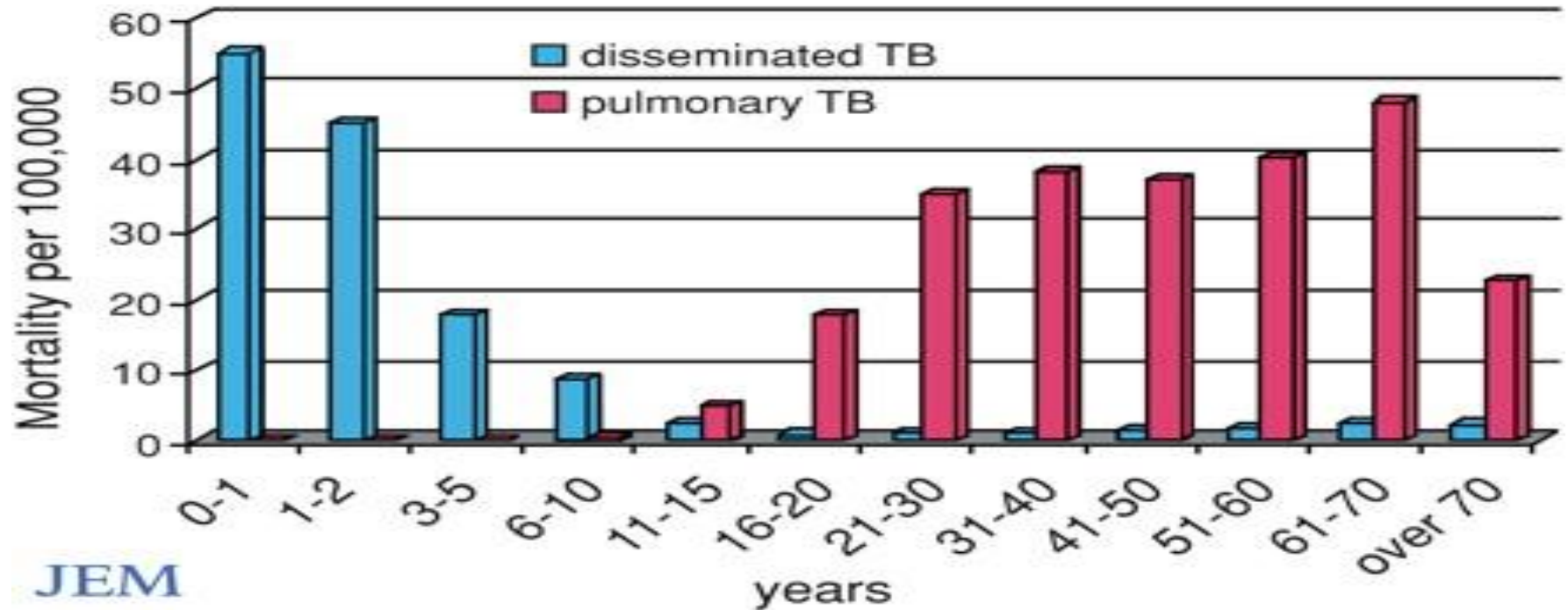
Primary – childhood

- Local – small lesion
- Regional – large nodes
- Systemic – spreads easily

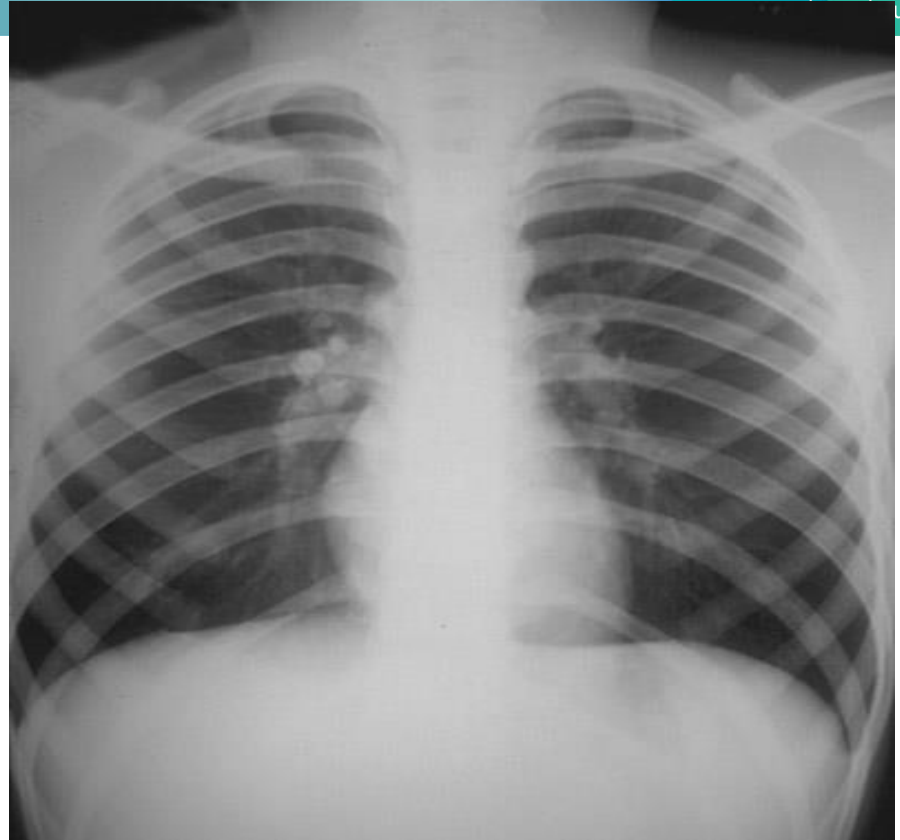
Secondary – adult

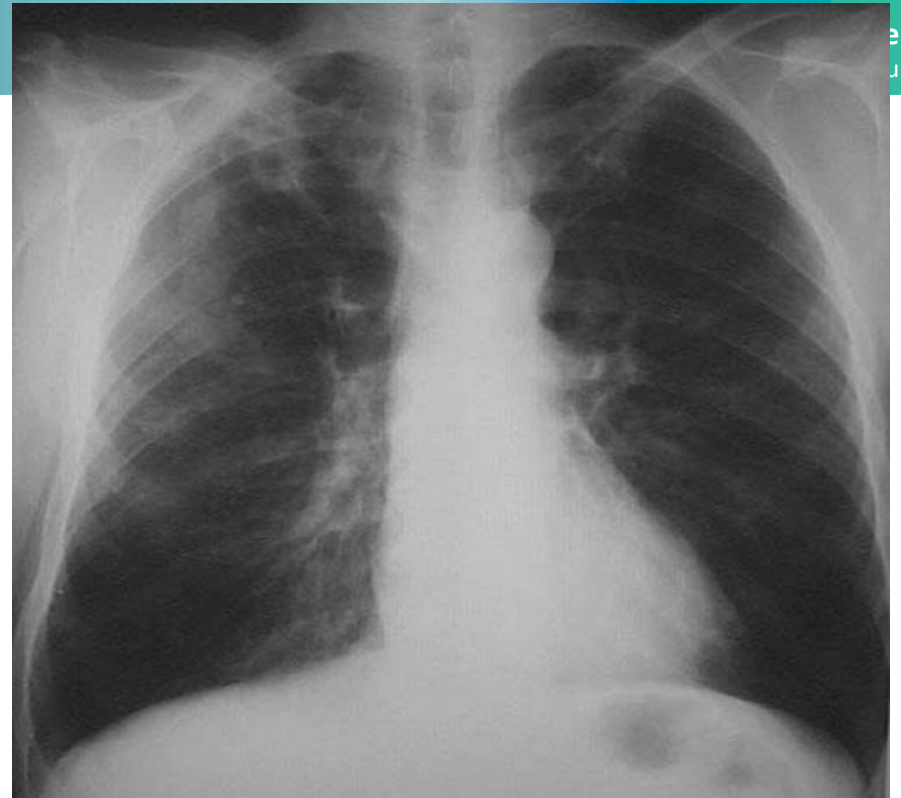
- Local – large lesion
- Regional – small nodes
- Systemic – does not spread easily

Cause of death in untreated cases – Bavaria 1905



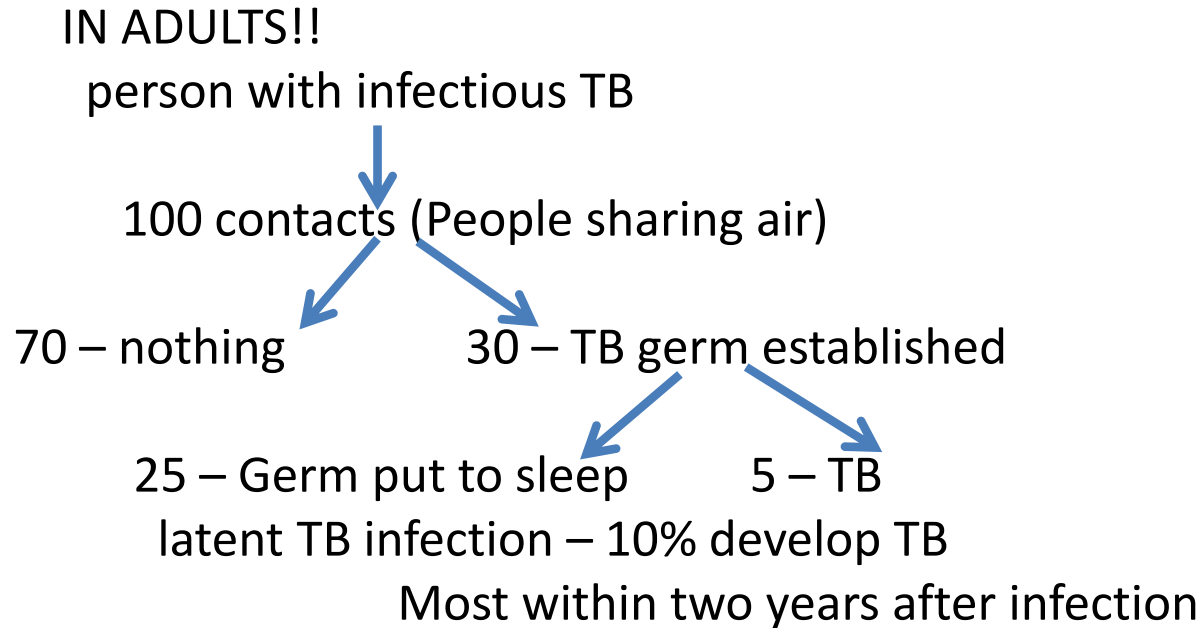
JEM



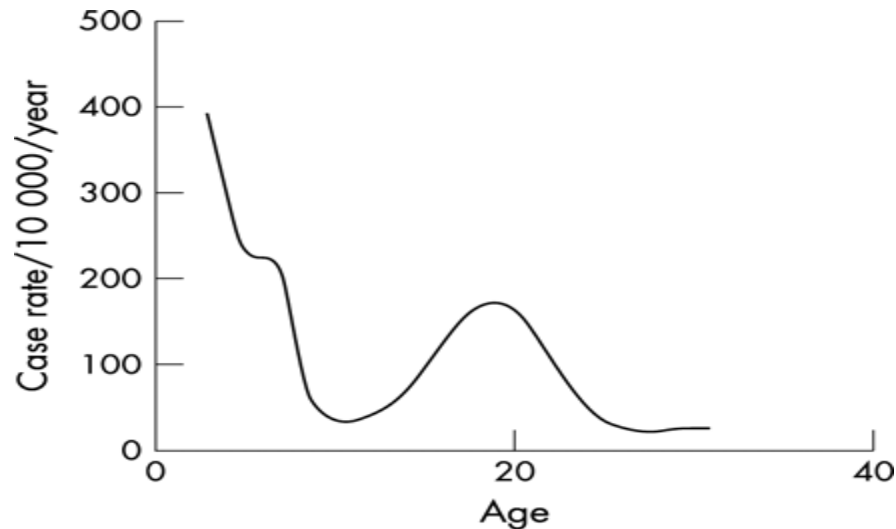


What happens if someone inhales TB germ?

Not very infectious – shared air for minimum of 8 hours



Likelihood of latent TB infection becoming active TB by age BCG reduces chance in under 5's



Lubeck disaster – 1929 – 251 neonates given oral BCG contaminated with live TB bacteria

- 100% infected
- 85% developed TB disease
- 30% died

How is it controlled?

Contact tracing, looking for

- Origin of disease
- Those infected – treated for LTBI, but number needed to treat may be large.
- Those vulnerable, particularly young children – started on meds

Immigrant screening – CXR

Contact tracing testing

- Mantoux test – skin test
- Antigen testing – Quantiferon blood test
- Cannot distinguish between TB disease and LTBI
- Usually two tests done 8 weeks apart as it takes that long for body's immune system to react
- Mantoux probably more sensitive in young children

Contact tracing

- Extent based on infectiousness of case
- Extra-pulmonary – confined to asking if anyone with symptoms in family
- Pulmonary microscopy –ve, (smear –ve), culture +ve, family only
- Pulmonary microscopy +ve, culture +ve, all those who have shared air for more than 8 hours since the person started coughing. Concept of concentric circles

BCG vaccination

- All strains developed from M Bovis strain which was attenuated in 1929 in Germany
- Protection when given in those older than 5 years varies – probably depends on background rate of non-TB mycobacteria.
- Given routinely only to neonates or children <5 at increased risk of TB – see Immunisation Handbook
- Needs only to be given correctly, no need for mantoux or blistering.
- Lesion appears after weeks and lasts weeks, up to 1cm is acceptable
- If concerned, leave it alone and consult specialists.

IF YOU THINK SOMEONE HAS TB

- Don't do mantoux or quantiferon. Person with acute illness and +ve mantoux does not have TB
- Chest xray, sputum, isolate, consult

IF YOUR PATIENT IS A CONTACT OF TB

- Public health should follow up – may take some time, eg culture +ve TB
- Not all contacts need follow up
- Don't do testing yourself, if you think person has been overlooked, contact public health

OLD TB ON CHEST XRAY

- Infectiousness of non-acute TB cannot be determined by CXR
- Only by induced sputum, bronchoscopy
- Any suggestion of old TB, no matter how minor, consult with Chest clinic

THE END

