



Mr Ben Harris

Medical Laboratory Scientist

Honorary Lecturer, University of Otago

FULLY BOOKED

Saturday, August 13, 2016

(Room 2)

16:30 - 17:30 WS #134: The Challenge of Rest Home Environments

The Challenge of Rest Home LTCF Environments



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45 Country Death Comparisons

of 16 million deaths in 45 countries
older people dying in LTCF:

average 18%

NZ 31%

median LTCF age now > 85yrs

Dying in LTCF NZ Stats

40,000 elderly resident in LTCF (NZACA 2013)

In NZ 20% die $\leq$ 6 months

But 36% if admitted ex acute hospital

i.e. 'de facto hospice'

so palliative care very important



Need multidisciplinary holistic care

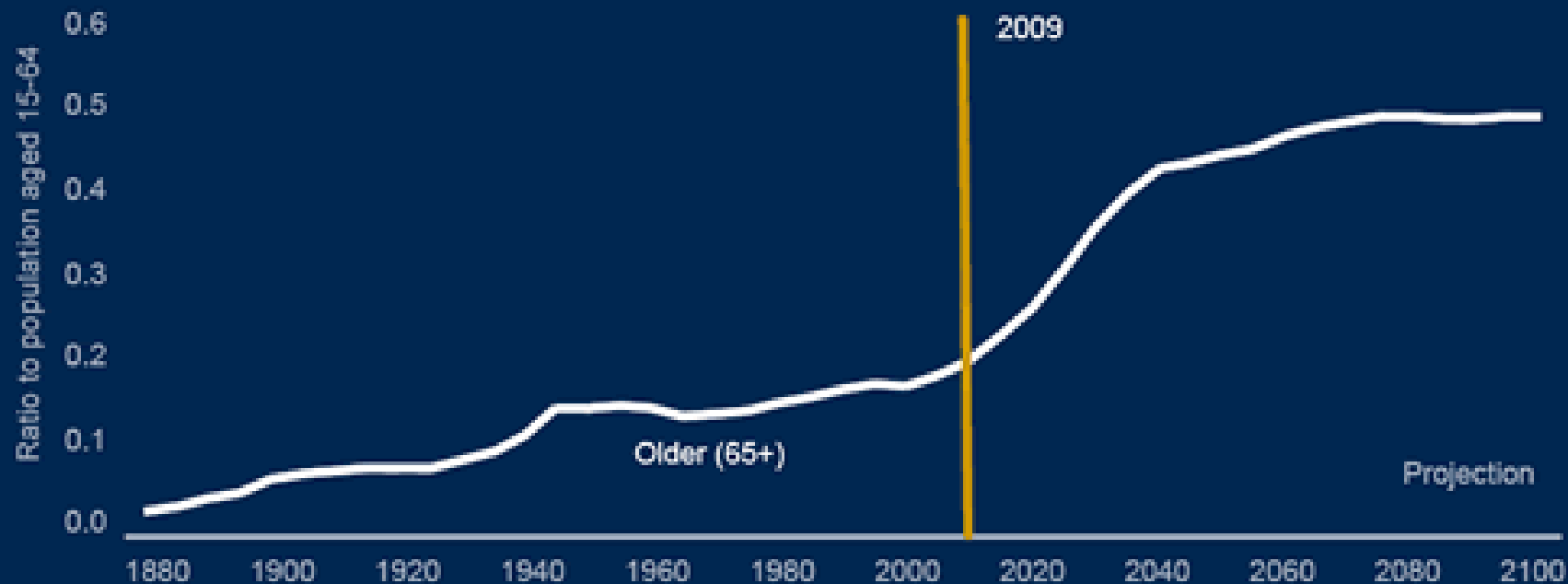
for mind, body, spirit

given by Cleaners (mainly),

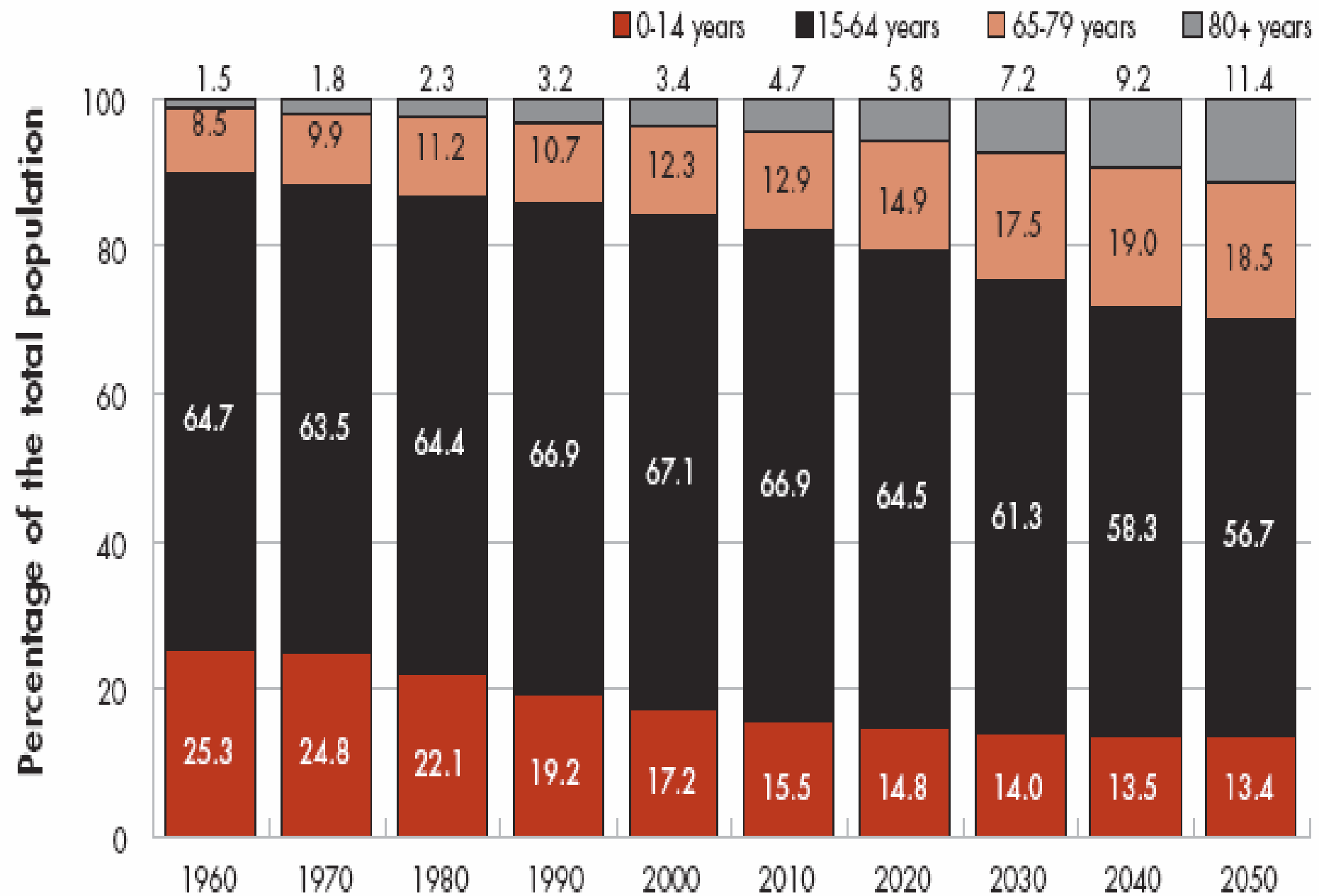
+ GP, RN, HCA

Demographics

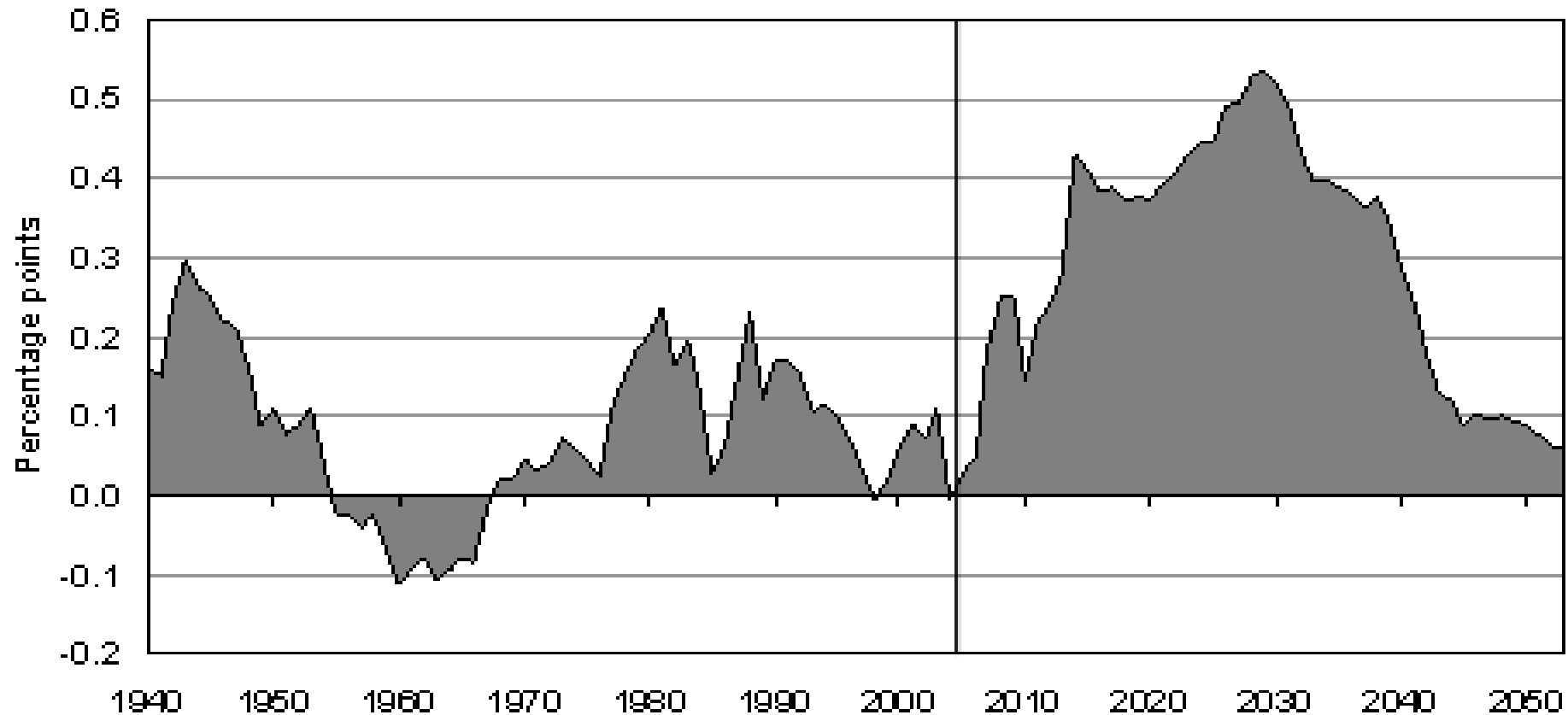
Ratio of those aged 65+ to 15-64



Source: Statistics New Zealand



NZ Yearly Change in Share of People 65 and Older



In 1900 there were 15 people working for every person over 65
Now 5 people working for every person 65 and over
by mid-century there are projected to be only two

Predisposing sepsis factors in elderly patients

Risk Factors for Severe Infection

Dementia, delirium,
excess injury, aspiration

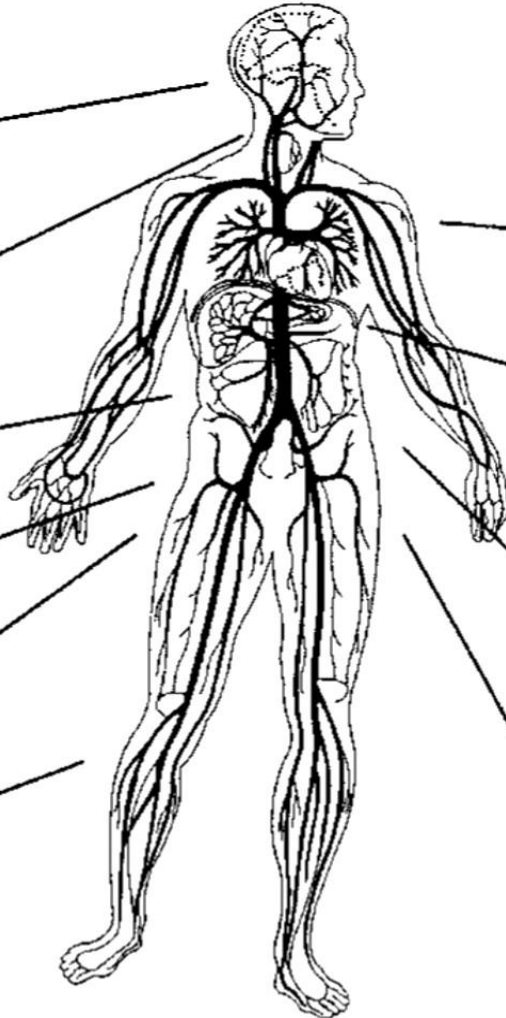
Decreased gag and
cough reflex

Endocrine deficiency
(adrenal, gonads, thyroid)

Poor nutrition

Immunosenescence
T and B cells

Immobility, skin
breakdown



Risk Factors for Severe Sepsis and Mortality

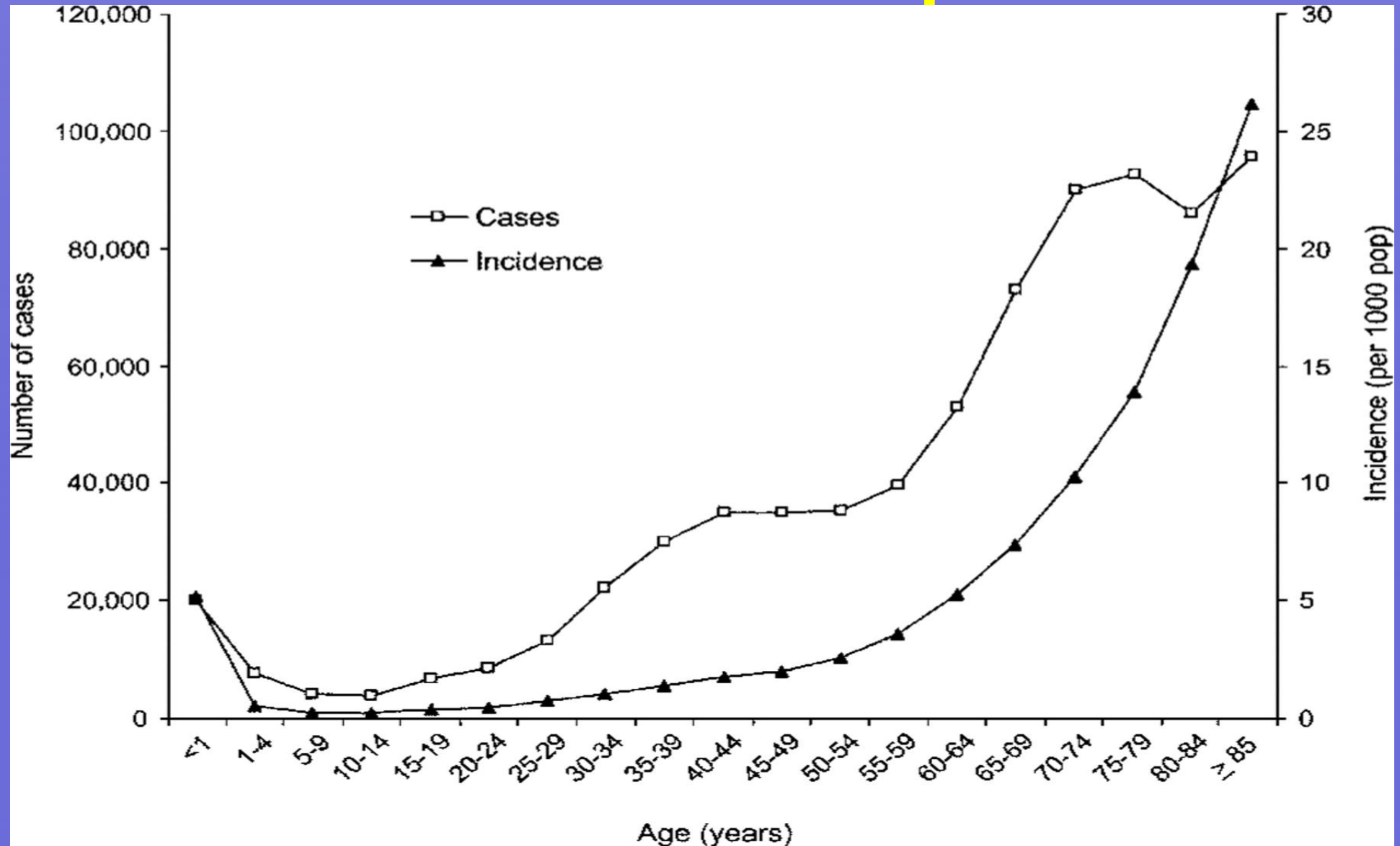
Concomitant medical
diseases

Diminished
cardiopulmonary reserve

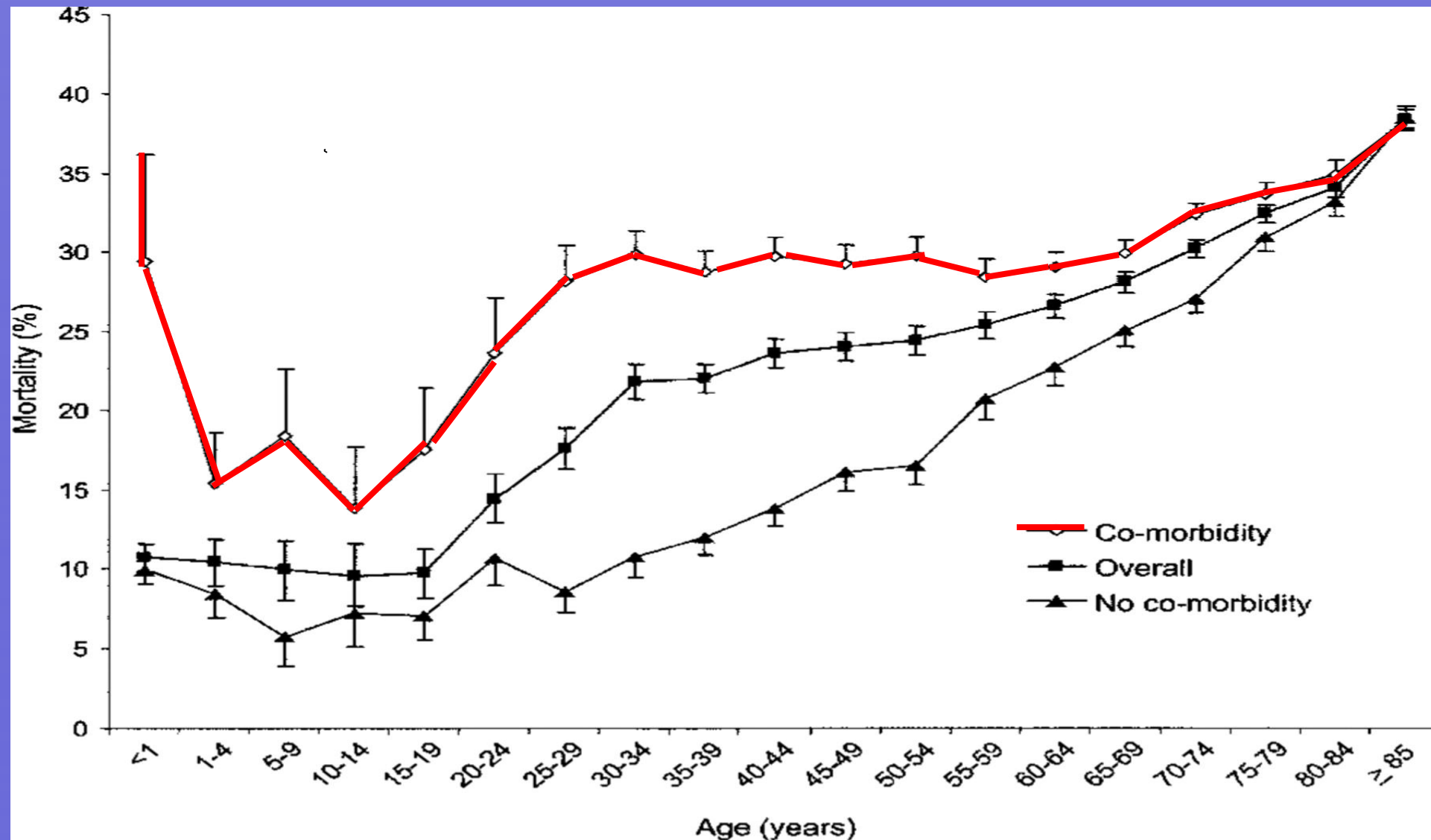
Age-related decrease in
organ function

Intact, or even enhanced,
innate immune responses
and cytokine production

Age-specific number and incidence cases of severe sepsis



Age-specific mortality rates for all severe sepsis with or without associated underlying comorbidity



NZ Growth Projections

Statistics NZ

4 million today to 5 million in 2050
25% increase in 45 years

The 'Old' - **over 65 year olds** projected growth
three-fold by 2050 **i.e. 300%**↑

The 'Oldest of the Old' - **85 year olds and over**
grow six-fold by 2050 **i.e. 600%**↑↑

Immigration Solution?

One way of keeping the aged-dependency ratio under 20% (where it was in 2005) all the way out to 2050 would require 300,000 net immigrants each year from 2020

i.e. Immigration not a solution

LTCF Challenges

- ↑ demographic, ↑ ↑ > 80 year olds
- ↑ complex cases
- ↑ ↑ dementia
- Earlier public hospital discharge
means ↑ LTCF hospital care cases
- Later community entry → ↑ care needs
- More underlying illnesses

LTCF Challenges (2)

- ↓ Carer pool
- ↑ Gen Y demand more equality or leave
(e.g. shift 'equality')
- ↑ turnover
- ↓ continuity
- ↓ institutional knowledge
- ↓ IPC knowledge
- ↓ average formal training time

Staff Turnover LTCF (USA)

STAFF	TURNOVER	VACANCY
Administrator	43%	Sharp declines in new administrators
Registered nurse	49%	18%
Licensed practical nurse	50%	14%
Direct care workers	76%	12%

LTCF Challenges (3)

↑ ↑ Diagnostic challenges

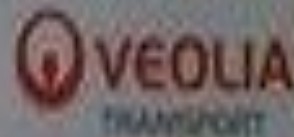
- clinical
- subtle signs & symptoms
- lack readily available onsite tests
e.g. radiology, laboratory
- good quality samples difficult to
obtain (e.g. resp, urine)
- no routine onsite Dr or Specialist

LTCF Challenges (4)

- ↑ MDRO's e.g. MRSA, ESBL, CRE, VRE
- ↑ Outbreaks e.g. *Norovirus, influenza, scabies*
(*C. difficile*)
- Financial constraints
- Prescribing often remote, by phone, etc



**DO NOT Pick
your nose and
wipe it on the
walls or doors**





Two Infection Sources

A. Endogenous Source

- We 'catch' the infection from ourselves
- The microbes emerge from our normal microbial protective flora when a compromised site or situation

B. Exogenous Source

- We catch the microbe (infection) from others who are infected (or carriers e.g. MDRO)

Infections

Endogenous vs Exogenous

Endogenous

- Staph aureus
- E.coli, Klebsiella coliforms, etc
- Strep
- Anaerobes

i.e. Most non viral infections

Exogenous

- Influenza, Norovirus
- Scabies
- Hepatitis, HIV
- Bordetella pertussis
- TB
- Campylobacter
- Salmonella, Shigella

Endogenous Infections

MOUTH

Strep.viridans
anaerobes
Candida

SKIN & NASAL

Staph. aureus
MSSA or MRSA

SKIN

Staph.epidermidis

BOWEL

Anaerobes
E.coli, ESBL, CRE VRE

VAGINA

Lactobacilli
Gardnerella
Candida



Microbial Sharing

Freshly washed skin 1,000 – 10,000 bacteria sq cm

We each carry 90 trillion bacteria normally

'my bugs are your bugs'

Newly arrived skin microbes



normal flora from

others skin (touch)

+ exogenous microbes from

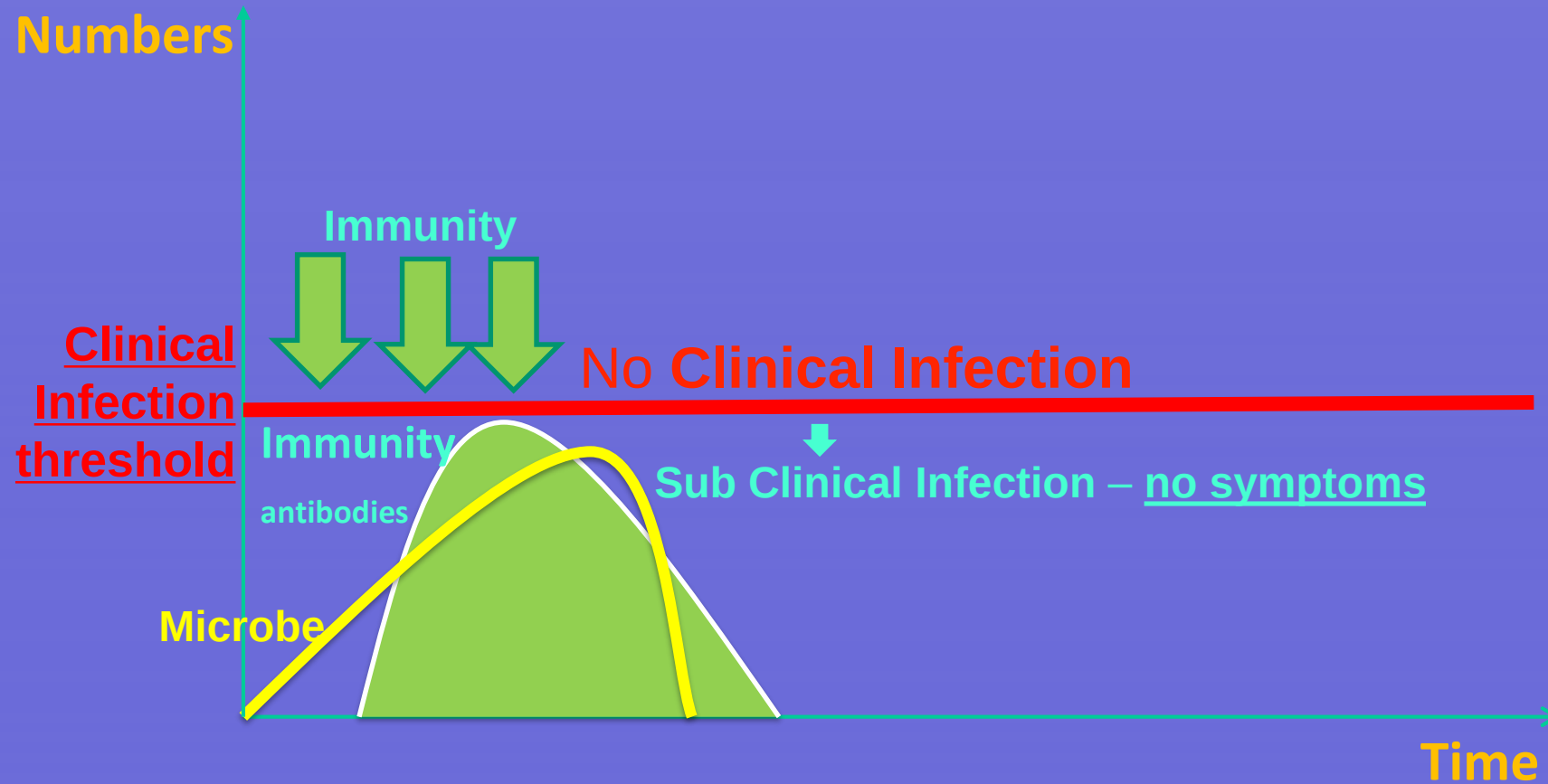
someone else (infection) contact

can always be washed off with good technique

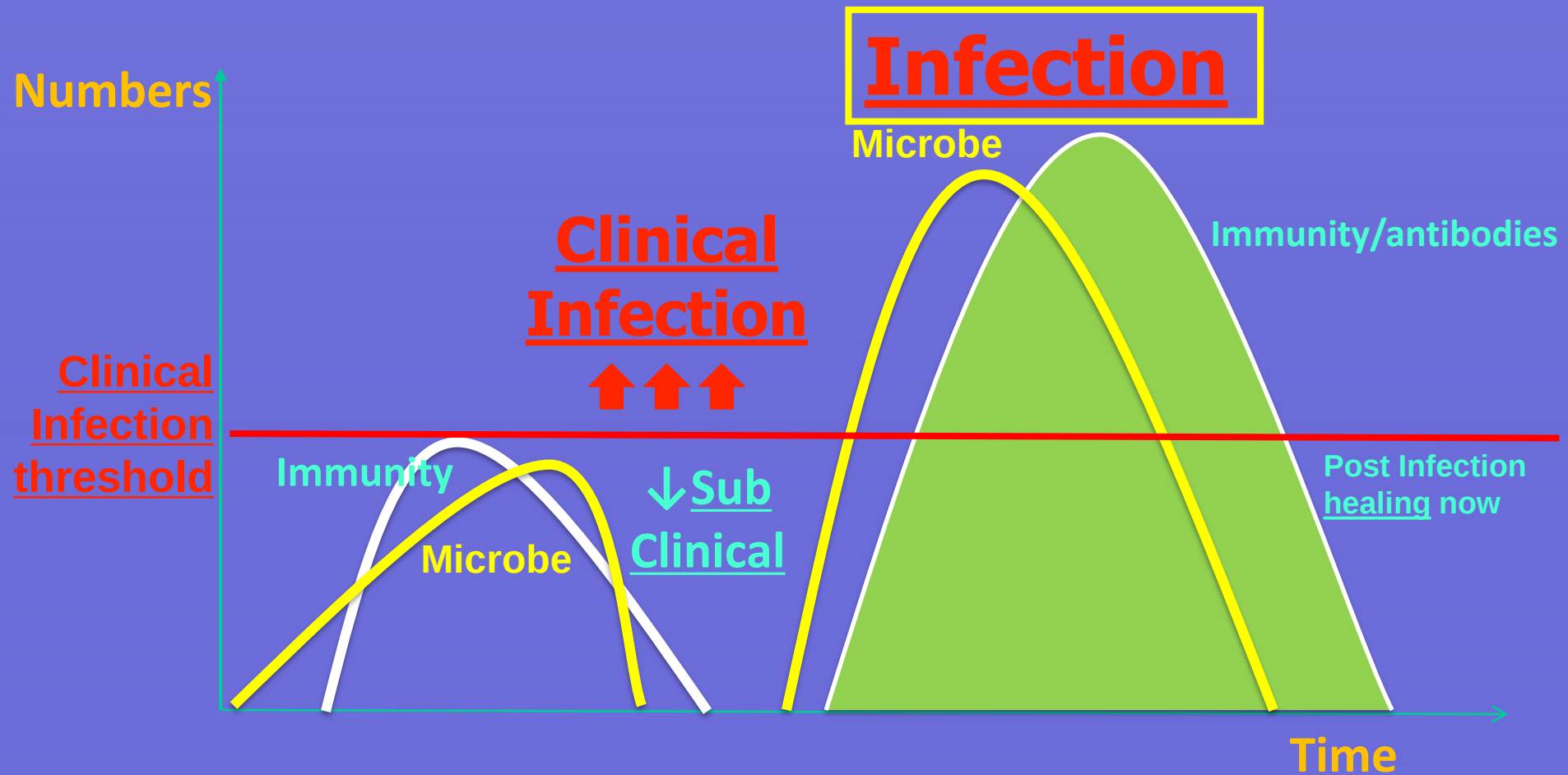
no matter how resistant or virulent

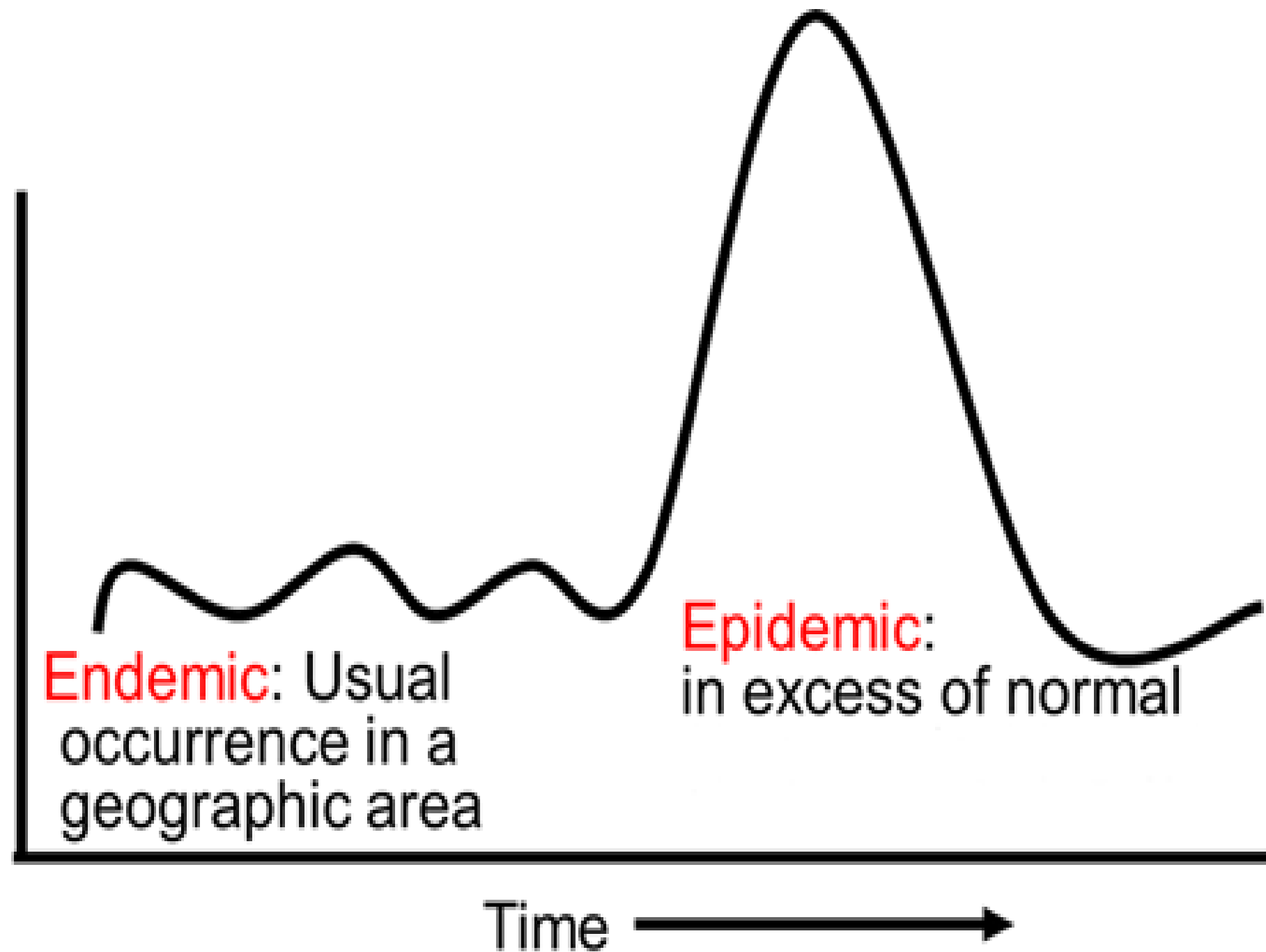


Infection vs Immunity



Infection vs Immunity





Gastroenteritis

Bacteria

- Campylobacter
- Salmonella
- Shigella
- Clostridium difficile
- E. coli (EPEC, EIEC, UTEC O157, etc)
- 'travellers diarrhoea'

Virus

- Norovirus
- Rotavirus
- Sapovirus

Parasites

- Giardia
- Cryptosporidium

NOROVIRUS



YOU DON'T WANT IT



DO NOT USE

**Restrooms for
VOMITING! Vomit goes
OVER RAIL ONLY!**



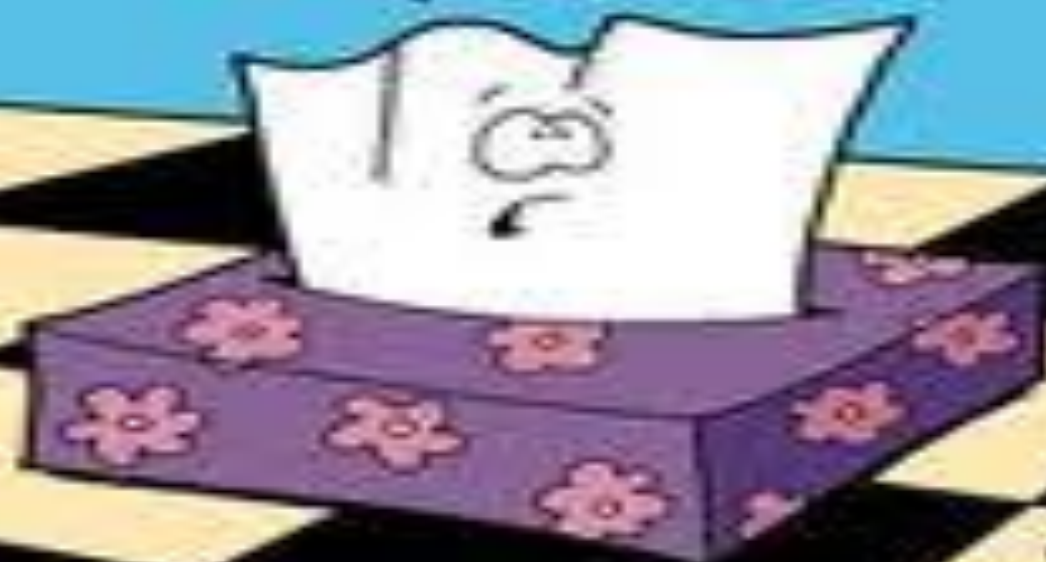
**Thank You,
Your Crew**

Gastroenteritis

use liquid soap water (Norovirus, *C. difficile*)



Yup, that's
my job... I
can't think of
anything more
humiliating...
I'm so ashamed.
By the way,
What do you do?

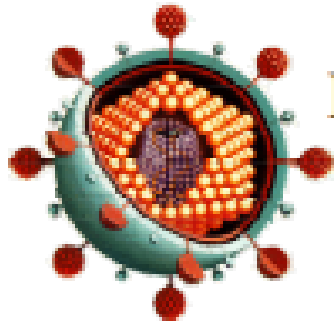




Influenza Surveillance Criteria



INFLUENZA-LIKE ILLNESS **CASE DEFINITION**



Influenza-like Illness (ILI) =

fever ($\geq 100^{\circ}$ F)*

AND

cough and/or sore throat

**(in the absence of a known cause
other than influenza)**

**Temperature can be measured in the office or at home*

Please report all patients that meet the ILI case definition above unless diagnostic tests confirm a cause other than influenza. For example, a patient with fever, cough, and

ILI Frequency 207 Patients Hospitalised then found to have Influenza Infection 3 years retrospective evaluation

Ref: Infection Control & Hospital Epidemiology vol 27 No 3 pp266-270

TABLE 3. Symptoms and Signs Reported by 207 Patients Before Influenza Diagnosis, by Age

Symptom	No. (%) of Patients, by Age		P	Overall
	Aged Less Than 65 Years (n = 116)	Aged 65 Years or More (n = 91)		
Cough	104 (90)	82 (90)	.85	186 (90)
Subjective fever	84 (72)	53 (58)	.04	137 (66)
Fatigue	52 (45)	56 (61)	.02	108 (52)
Chills and/or rigors	48 (41)	29 (32)	.13	77 (37)
Nausea and/or vomiting	51 (44)	25 (28)	.02	76 (37)
Myalgias	41 (35)	26 (29)	0.11	67 (32)
Coryza	42 (36)	21 (23)	.05	63 (30)
Headache	35 (30)	19 (21)	.12	54 (26)
Sore throat	35 (30)	9 (10)	<.01	44 (21)
Diarrhea	20 (17)	16 (17)	.98	36 (17)

NOTE. Data in bold type are statistically significant.

ORIGINAL ARTICLE

Is Influenza an Influenza-Like Illness?

Clinical Presentation of Influenza in Hospitalized Patients

Hilary M. Babcock, MD; Liana R. Merz, MPH; Victoria J. Fraser, MD

BACKGROUND. Early recognition of influenza virus

OBJECTIVE. To determine the clinical presentation of

DESIGN. Case series. Data were collected retrospectively from medical records, clinical symptoms and signs, microbiologic test results

SETTING. A 1,400-bed teaching hospital.

PATIENTS. A total of 207 inpatients who received a

RESULTS. Over the course of 3 seasons, 207 patients were infected with influenza B virus). The most common clinical presentation was fever (100%); 124 patients (60%) had a documented cough (66%); 124 patients (60%) had a documented sore throat (66%). Centers for Disease Control and Prevention (CDC) criteria for influenza-like illness (ILI)—temperature $\geq 38.0^{\circ}\text{C}$ or greater and either cough or sore throat—were met by 107 patients (51%). There were no differences in the proportion of patients who met ILI criteria with respect to age, sex, season, influenza virus type, or time to diagnosis in the hospital. Most patients (150 [72%]) received acetaminophen. Only 41 patients (20%) had positive results of clinical cultures; 178 patients (86%) received antibiotic therapy. Fifty-six patients (27%) had pneumonia; 36 (17%) required admission to the ICU, and 25 (12%) required ventilatory support. Patients with pulmonary disease were more likely to require ventilatory support (12 [26%] vs 13 [8%]; $P = .003$).

CONCLUSIONS. Only half of hospitalized patients with influenza met CDC criteria for ILI. These criteria may be more appropriate in outpatient settings. A high index of suspicion is needed to recognize influenza in hospitalized patients.

“CONCLUSIONS:

Only half of hospitalised patients with influenza met CDC criteria for ILI
A high index of suspicion needed to recognise influenza in hospitalised patients”

Surveillance vs Clinical criteria

ILI Frequency 207 Patients Hospitalised then found to have
Influenza Infection 3 years retrospective evaluation

Infection Control & Hospital Epidemiology 27 No 3 pp266-270

<u>ILI Symptom</u>	%
• Temperature* $\geq 37.8^{\circ}\text{C}$	60
• Temperature $\geq 37.8^{\circ}\text{C}$ + cough	51
• Temperature $\geq 37.8^{\circ}\text{C}$ + sore throat	12
• Temperature $\geq 37.8^{\circ}\text{C}$ + either cough or sore throat	51

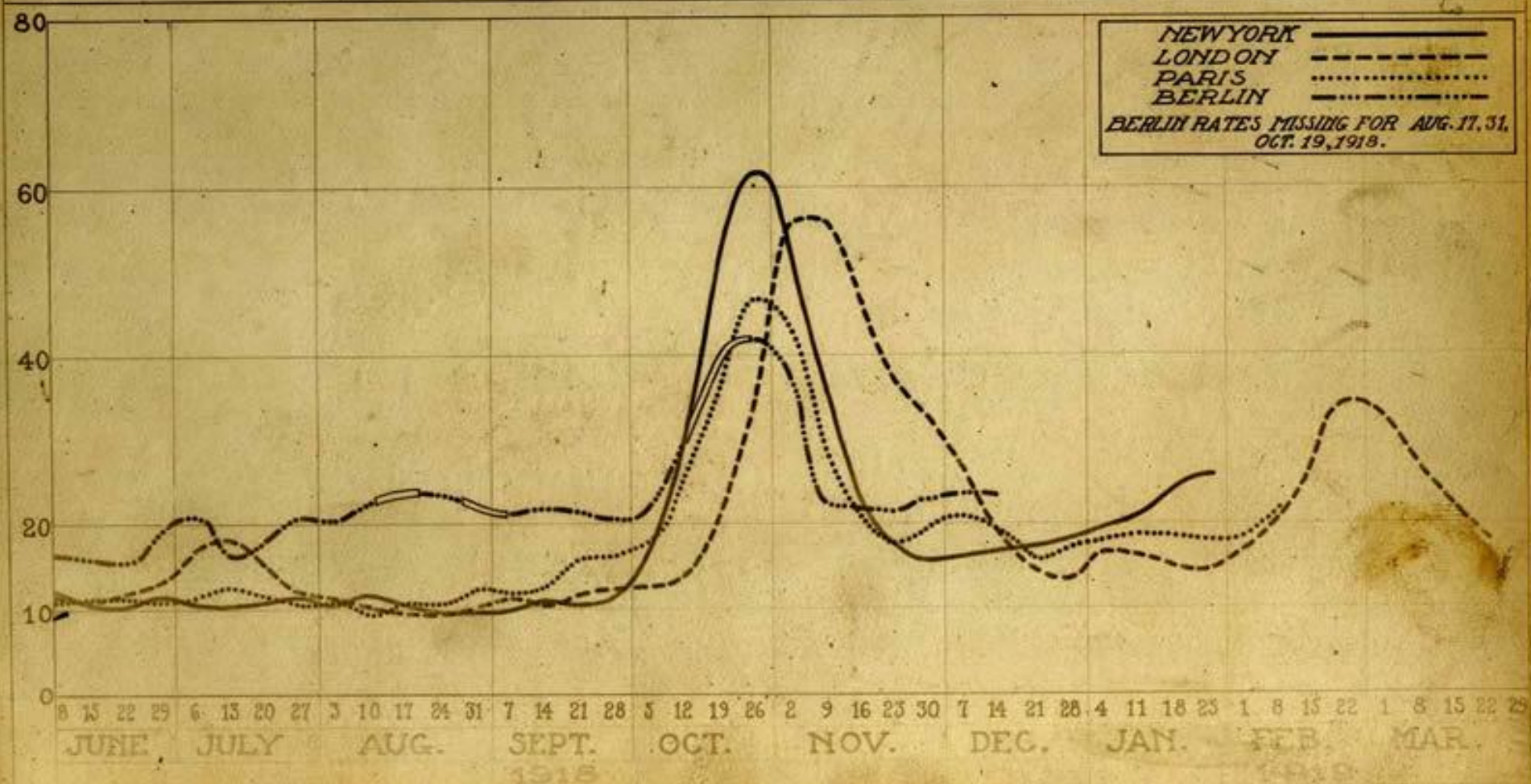
* Measured temperature on admission



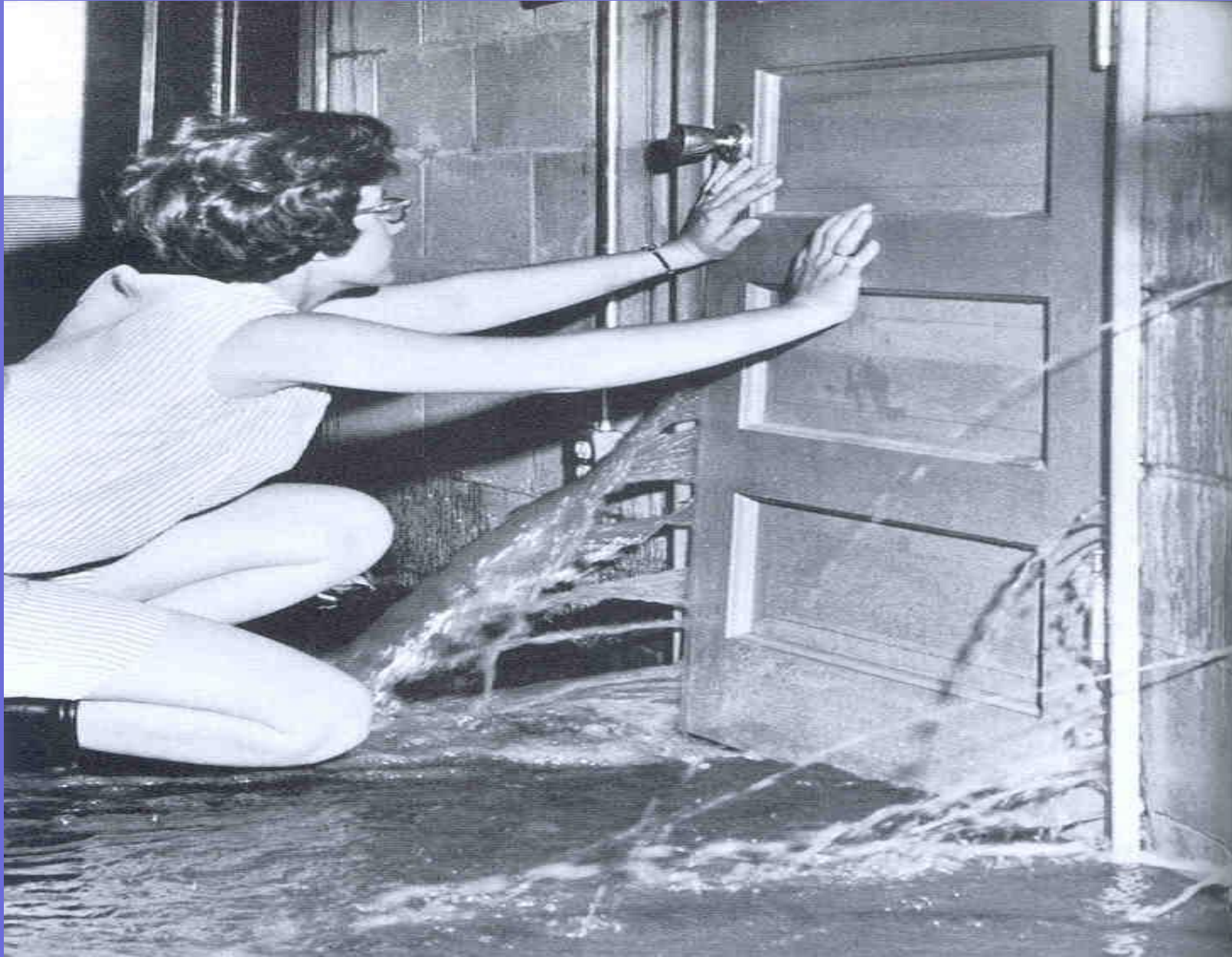
INFLUENZA PANDEMIC

MORTALITY IN AMERICA AND EUROPE DURING 1918 AND 1919

DEATHS FROM ALL CAUSES EACH WEEK
EXPRESSED AS AN ANNUAL RATE PER 1000



UTI in LTCF's



HAEMATOLOGY		BIOCHEMISTRY		MICROBIOLOGY		CYTOLOGY	
☒ CBC		☐ Urea	☒ Creatinine	SWABS Site _____ Culture ☐		☐ Cx Smear	☐ One ☐ Two slides
☐ ESR	☐ Reticulocytes	☐ Electrolytes	☐ Uric Acid	Site _____ Chlamydia ☐		☐ Vag Smear	☐ Cervex Broom
☐ Iron Studies	☐ Ferritin	☐ Calcium	☐ Amylase	Site _____ Herpes ☐		LMP ☐	☐ Sputula
☐ Red Cell Folate	☐ B12/Folate	☐ PO ₄	☐ Lipid Profile	URINE MSU/CSU/OTHER (add details below)		☐ Post Menopause	☒ Other 3
☐ INR (Prothrombin)	☐ Coag. Screen	☐ PSA	☐ GTT (75g)	☒ Culture	☐ Tb Culture x3	☐ Urine Cytology x3	
IMMUNOLOGY		☐ Glucose	☐ Polycose Screen	SPUTUM			
☐ First Antenatal Screen E.D.D.		☐ Urine Microalb	☐ HbA1c	☐ Culture	☐ Tb Culture x3	☐ Sputum Cytology x3	
☐ Later Antenatal Screen ☐		☐ Myoc. Enzymes	☐ Prot & Elec	FAECES		☐ FNA* Add diagram & site details below	
☐ Hep BsAg	☐ Hep A IgG		☐ Immunoglobulins	☐ Culture	☐ Occult Blood	HISTOLOGY	
☐ Hep BsAb	☐ Hep A IgM	☐ Liver Function		☐ Ova & Parasites x2	☐ Giardia Antigen	☐ Site (details below)	
☐ Hep B Core Ab	☐ Hep C Ab	ENDOCRINE		SKIN SCRAPING/FUNGI (add details below)		SPECIMEN COLLECTION (Blood/Urine)	
☐ Thyroid Antibodies		☐ FT4	☐ FSH	Site(s) _____		☐ Fasting ☐ Non Fasting	
☐ HIV	☐ Treponema	☐ TSH	☐ LH	SEMINAL FLUID		Time _____ am/pm Date _____	
☐ ANA	☐ Rubella Status	☐ FT3	☐ Progesterone	☐ Fertility	☐ Post Vasectomy	URGENT	
☐ dsDNA	☐ Infect. Mono	☐ Prolactin	☐ Quant HCG	ASPIRATE Site _____		☐ Phone _____	
☐ Tissue Autoabs	☐ EBV	☐ Oestradiol		☐ Culture	☐ Crystals	☐ Fax _____	
☐ Rheumatoid Factors	☐ Toxoplasma	☐ Cortisol am/pm		THERAPEUTIC DRUGS Dose _____		Mark appropriate tests with "Ph" or "Fax"	
☐ C. Reactive Protein	☐ CMV	Specify _____ Last dose at _____		☐ Blood Culture x2	☐ Mantoux	*Tests requiring an appointment	
☐ Streptococcal Abs	☐ Brucella			☐ Skin Allergy*		☐ Tick if not eligible for health benefit fee	
Additional Tests:		Clinical Details:		I certify that this patient and these tests are eligible for publicly funded laboratory services.			
PSA		Discomfort with full bladder ? cause					
		Medication:					

SYNAECOLGY

4.43 H.V./Cx Swab
 7.30 Chlamydia
 7.30 Herpes
 4.98 Smear ☐ Cx ☐ Vag
☐ Spatula ☐ One Slide
☐ Cervex Broom ☐ Two Slides
 MP: ___/___/___ ☐ Brush

HISTOLOGY FNA

31.03 134.64
 Site(s) _____

Seminal fluid
 22.65 Fertility
 13.47 Post-vasectomy

Aspirate
 14.51 Culture
☐ Crystals
 Site _____

MYCOLOGY

13.41
 Site(s) _____

8.07 Streptococcal abs
 6.32 C. Reactive Protein
 6.20 Rheumatoid Factors
 6.90 ANA
 10.00 ds DNA
 10.00 Tissue Autoabs
 2.65 Skin Allergy*
 5.13 Tuberculin

URGENT

☐ Ph _____
☐ Fax _____
 Mark appropriate test(s) with Ph or Fax

RF-C-V/2

tests.

Collection Date: ___/___/___ Time: _____

Clinical Details:
 Medication: **Yellow Urine**

I confirm that the patient and these tests will stay confidential. Further information on services. Date _____



Urine Dipstick



30-50% LTCF residents have bacteria in
their urine without symptoms

i.e. asymptomatic bacteriuria

The urine will likely be turbid, smelly
WBCs present, often nitrites positive

i.e. **DIPSTICK POSITIVE**

DO NOT TREAT

DO NOT SEND URINE TO LAB !!

Urine Dipstick



Half of the rest will have WBCs in their
urine without bacteria

i.e. DIPSTICK POSITIVE

DO NOT TREAT

DO NOT SEND URINE TO LAB !!

Urine Dipstick



Only 20% average of all residents will have
a negative dipstick

Every dipstick use promotes significant
antibiotic overuse to the detriment of all

i.e. a simple, misleading, 'dangerous', test

↑↑ Facility wide MDRO making & sharing
plus significant microbiome damage

Urine Dipstick LTCF



A positive dipstick is of no diagnostic value

A positive culture is not helpful
without Genitourinary Symptoms

The Lab report can NEVER make the
Diagnosis of UTI

UTI LTCF

- Residents with Symptoms from any cause will likely have a 'positive' urine culture
- Acute confusion can be a UTI indicator but most acute confusion has other causes – e.g. respiratory infection
- In absence of fever, UTI an unlikely cause of a non specific decline in clinical status (non catheterised)

UTI LTCF

- Mortality is rarely associated with UTI, unless febrile and/or catheterised then this can be much more serious
- Only about 10% of fever episodes without localising GU Sx have a urine source (if not catheterised)
- But if catheterised this increases to about one third (30%)

Elderly, urine

Asymptomatic

Untreated bacteriuria

has no effect on mortality

But treating will promote needless

Antibiotic resistance, shared by all

E.coli in Midstream Urine

Ampicillin	- resistant
Augmentin	- resistant
Cefotaxime	- intermediate
Ciprofloxacin	- resistant
Gentamicin	- resistant
Co-Trimoxazole	- resistant
Cephalothin	- resistant
Chloramphenicol	- resistant
Nitrofurantoin	- intermediate
Tetracycline	- resistant
Imipenem	- sensitive BUT how much? longer

PROBIOTIC





Myoc. Enzymes

Immunoglobulins

Liver Function

CRINE

T4 FSH

SH LH

T3 Progesterone

Prolactin Quant HCG

Destradiol

Cortisol am/pm

APPEUTIC DRUGS Dose

Last dose at

FAECES

Culture Occult Blood

Ova & Parasites x2 Giardia Antigen

SKIN SCRAPING/FUNGI (add details below)

Site(s)

SEMINAL FLUID

Fertility Post Vasectomy

ASPIRATE Site

Culture Crystals

Blood Culture x2

Skin Allergy* Mantoux

FNA* Add diagram & site details below

HISTOLOGY

Site (details below)

SPECIMEN COLLECTION (Blood/Urine)

☐ Fasting ☐ Non Fasting

Time am/pm Date

URGENT

☐ Phone

☐ Fax

Mark appropriate tests with "Ph" or "Fax"

*Tests requiring an appointment

☐ Tick if not eligible for health benefit fee

I certify that this patient and these tests are eligible for publicly funded laboratory services.

Clinical Details:

Medication:

Dark, smelly urine

? UTI

Reluctant to drink

Date: / /

Signature:

TSH FT3 Progesterone Prolactin Quant HCG Oestradiol Cortisol am/pm	Fertility Post Vasectomy ASPIRATE Site Culture Crystals Blood Culture x2 Skin Allergy* Mantoux	Fasting Non Fasting Time _____ am/pm Date _____ URGENT Phone _____ Fax _____ Mark appropriate tests with "Ph" or "Fax" *Tests requiring an appointment Tick if not eligible for health benefit fee I certify that this patient and these tests are eligible for publicly funded laboratory services.
THERAPEUTIC DRUGS Dose _____ Specify _____ Last dose at _____		
Clinical Details: Last urine at night Medication:		
Date: ____/____/____		Signature:

Pneumonia LTCF



Bedside Criteria ? Pneumonia

≥2 following symptoms or signs :

- New cough onset with or without sputum
- Fever (rectal temperature ≥ 37.8 C)
- Shortness of breath
- Respiratory rate ≥ 25 breaths per minute
- Heart rate ≥ 100 beats per minute
- Hypoxemia (oxygen saturation $< 94\%$ breathing room air)
- Acute change in cognitive or functional status
- Localised congestion (rales/ronchi)

Diagnostic Studies ? Pneumonia LTCF

- Pulse oximetry
- Chest radiograph (??!)
- Sputum, if obtainable, gram stain & culture
- Complete blood count with differential
- Blood urea nitrogen to assess hydration status





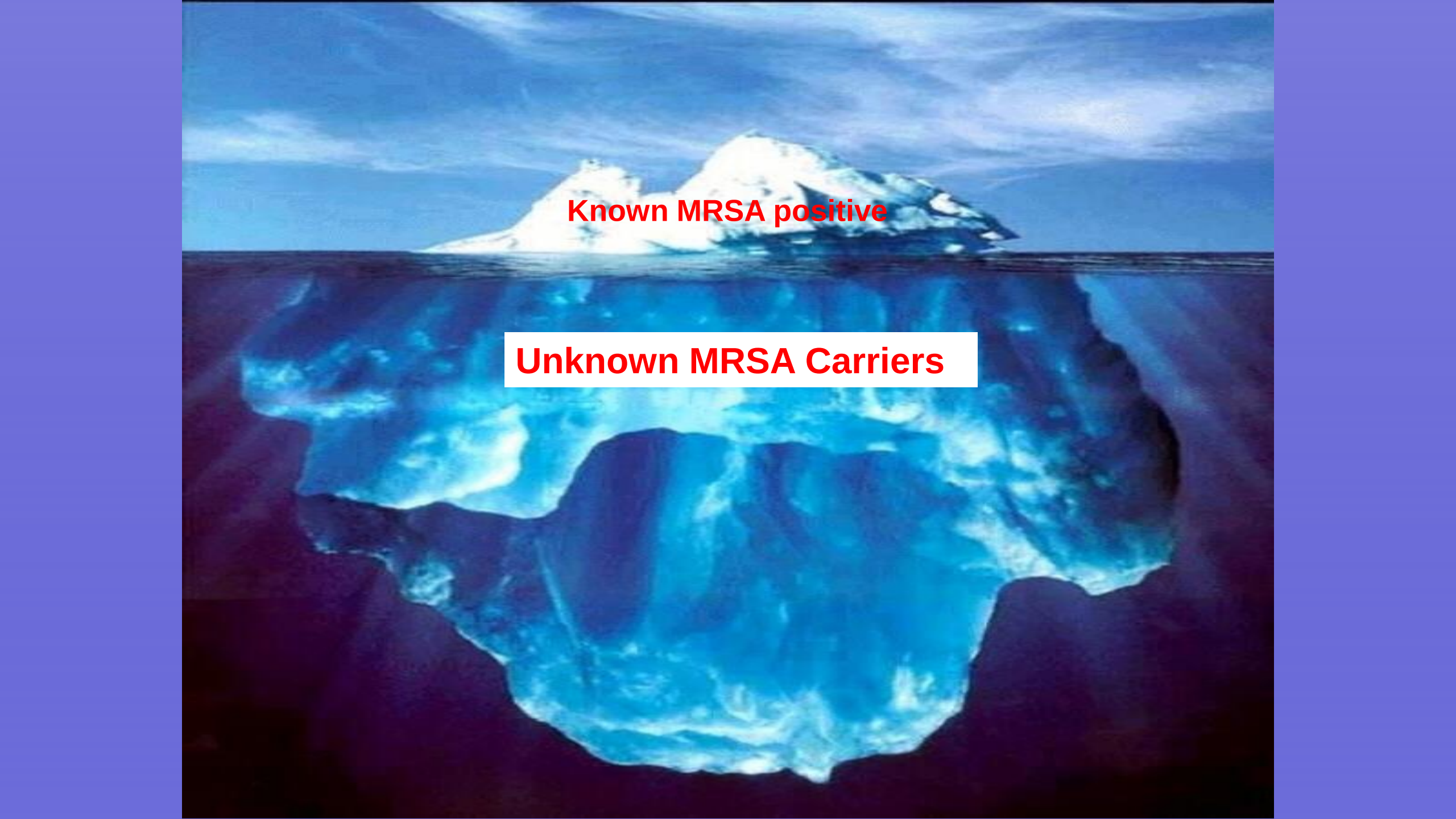




Skin & Soft Tissue





An iceberg floating in a dark blue ocean under a cloudy sky. The visible tip of the iceberg is small and white, while the submerged part is much larger and a deep blue color. The image is used as a metaphor for MRSA carriers, with the visible tip representing known positive cases and the submerged part representing unknown carriers.

Known MRSA positive

Unknown MRSA Carriers

MDRO Approach

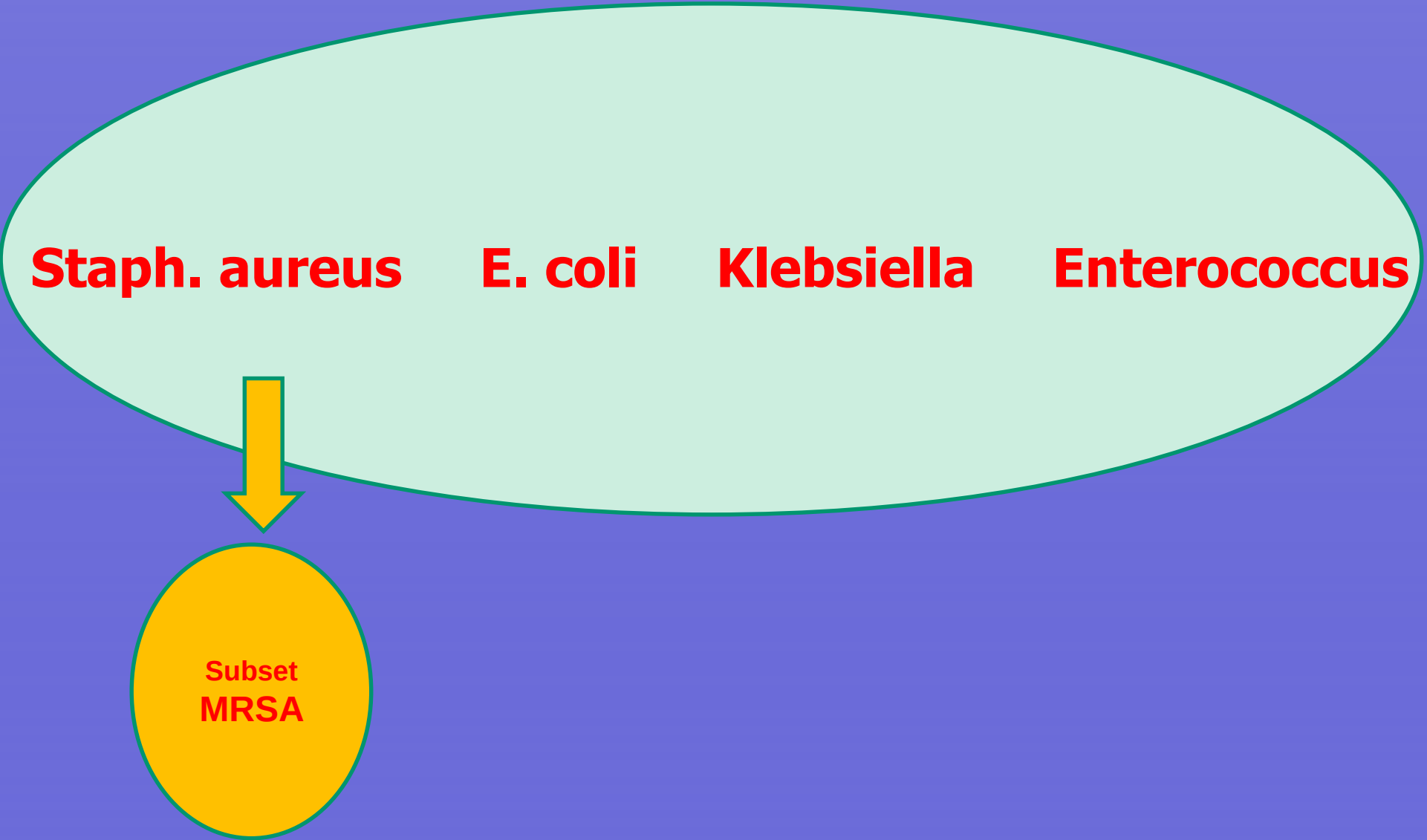
Silo or Horizontal Required ?

Staph. aureus

E. coli

Klebsiella

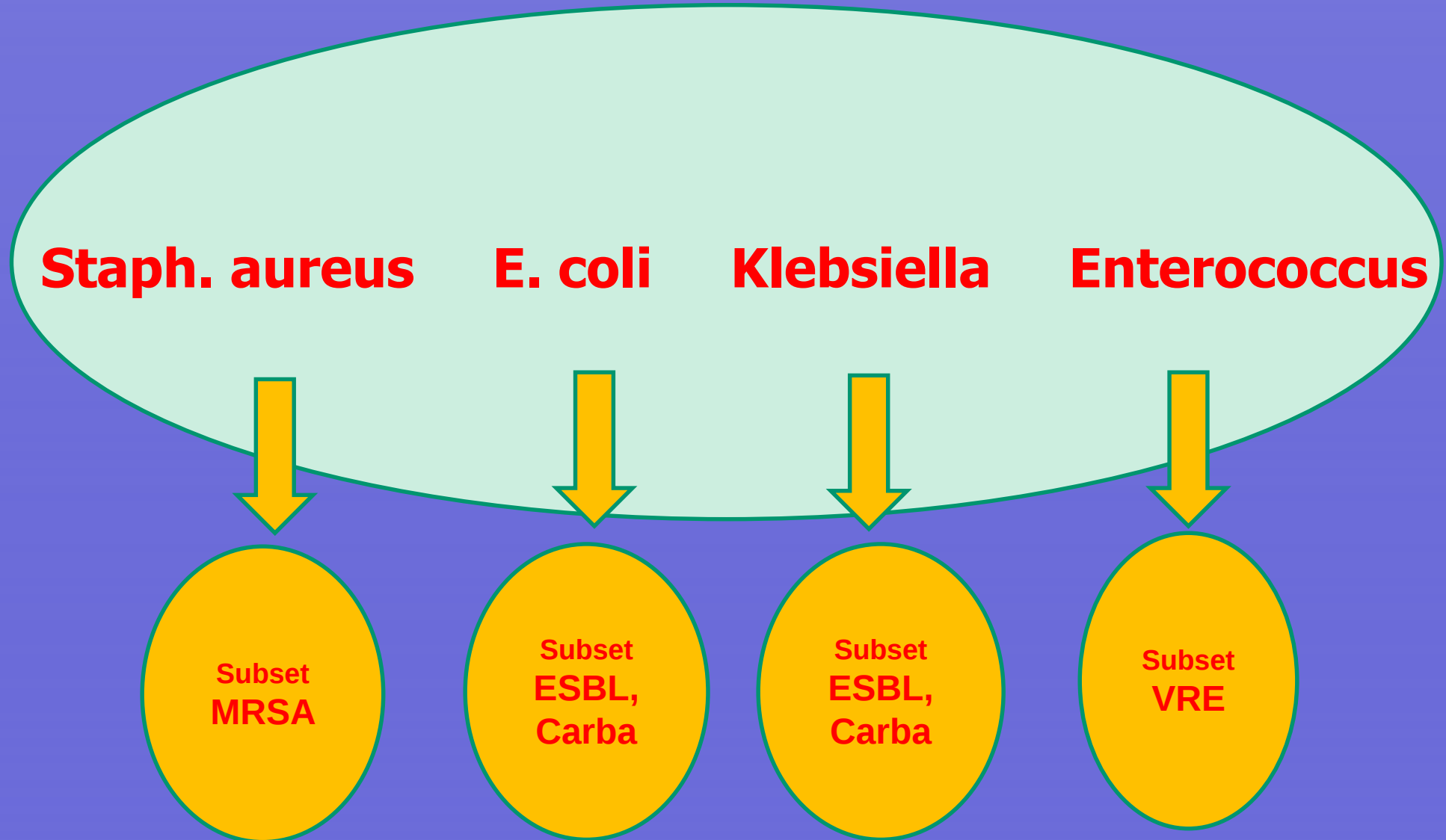
Enterococcus



**Subset
MRSA**

MDRO Approach

Silo or Horizontal Required ?



Largest Rest Home MRSA Study

Std Precautions vs 'Search & Destroy'

104 LTCF's 6,036 residents Vaud Switzerland

Largest Randomised Controlled one-year MRSA study

A. Standard Precautions only e.g. hand hygiene
(51 LTCF's control group)

versus

B. Universal MRSA screening followed by
decolonisation of all MRSA carriers
(53 LTCF's intervention group)

MRSA

Std Precautions vs 'Search & Destroy'

- 51 LTCF Control: Standard Precautions ONLY
anonymised MRSA screening at study entry
- 53 LTCF Intervention group:
all residents + new admissions MRSA screened
all MRSA carriers 5-day topical decolonisation
chlorhexidine skin + pharynx + Nasal mupirocin
plus environmental disinfection
(daily change clothes, sheets, bedding,
+ alcohol disinfection of surrounds)

Decolonisation process was repeated when failure

MRSA

Std Precautions vs 'Search & Destroy'

Results:

- Mean MRSA prevalence at baseline
8.9% in both groups (range 0- 44%)
declined to 6.6% (control group)
and 5.8% (intervention group)

60% of carriers in the intervention group were successfully decolonised, and 47% remained negative at study end
but others became MRSA carriers

MRSA Std Precs vs 'Search & Destroy' i.e. decolonisation

Conclusions:

Universal MRSA screening followed by
decolonisation of carriers had

no significant additional impact in reducing
prevalence of MRSA carriage rate

at one year compared to only Standard Precautions

5% of 500 Infections
which ones are MRSA positive???



5% + 5% of 500 People
? MRSA &/or ESBL positive???



**3 of this 500 have infected wound
should the MRSA be isolated??**













Mary Saybolt
Port Richmond

*The only thing golden
about my golden years is
my urine.*



Dolores
Port Richmond

*The best
thing old
more t*



We're not
ALL BAD!!



ben.harris@sclabs.co.nz

www.canterburyscl.co.nz

Infection Control tab



NZ Hand Hygiene Compliance Rates

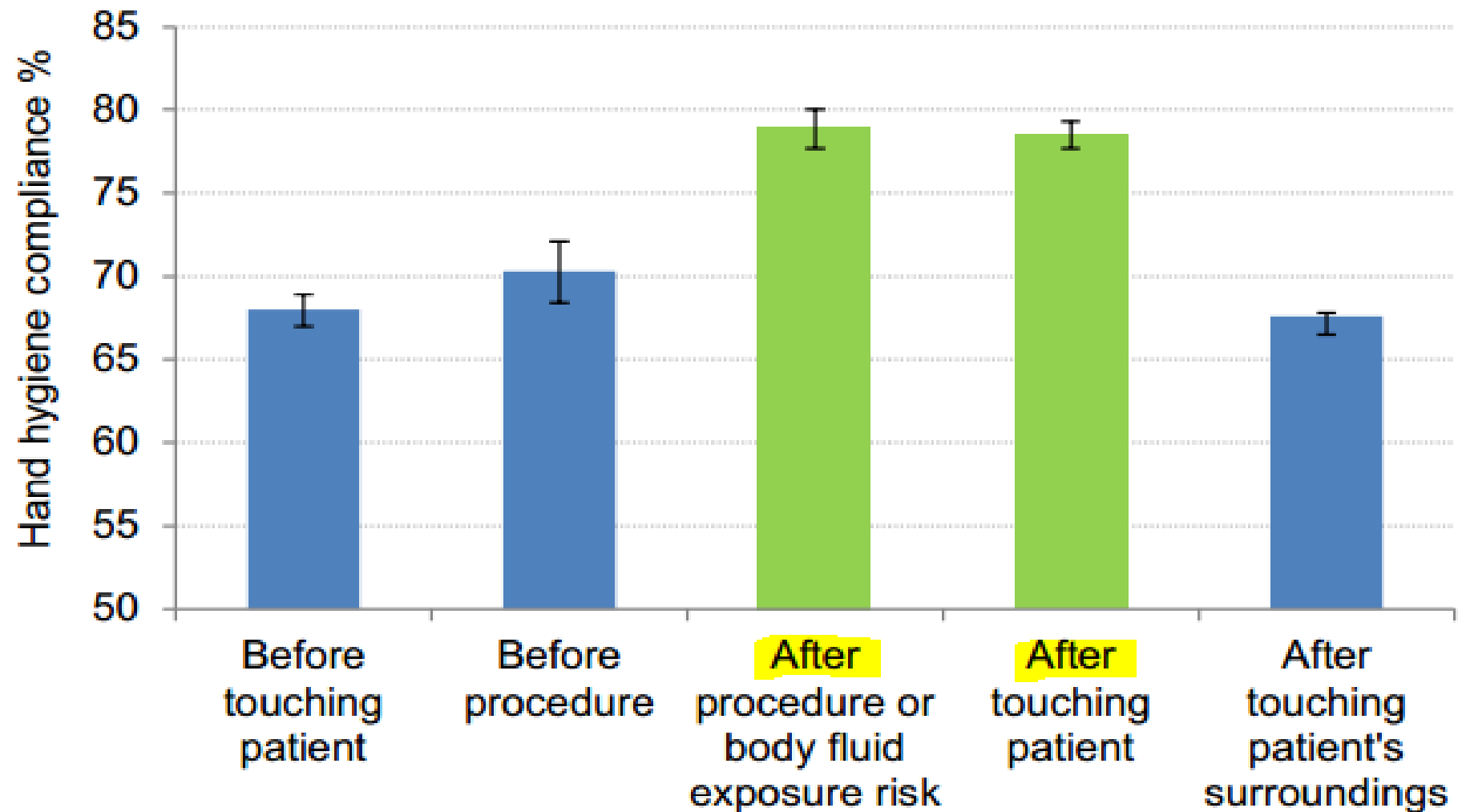
80% average

X20 DHBs

1 April – 30 June 2015

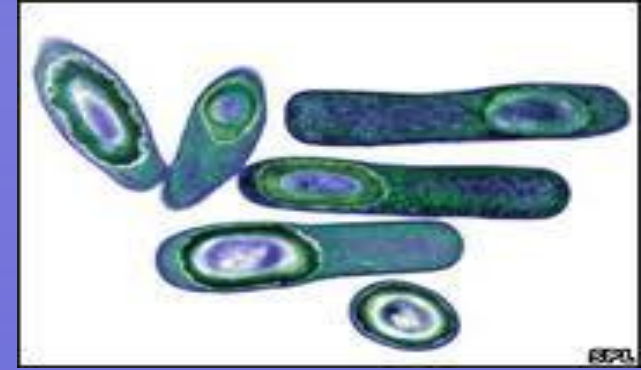
NZ DHB Hand Hygiene

Hand hygiene compliance by moment, all New Zealand Mar 2014



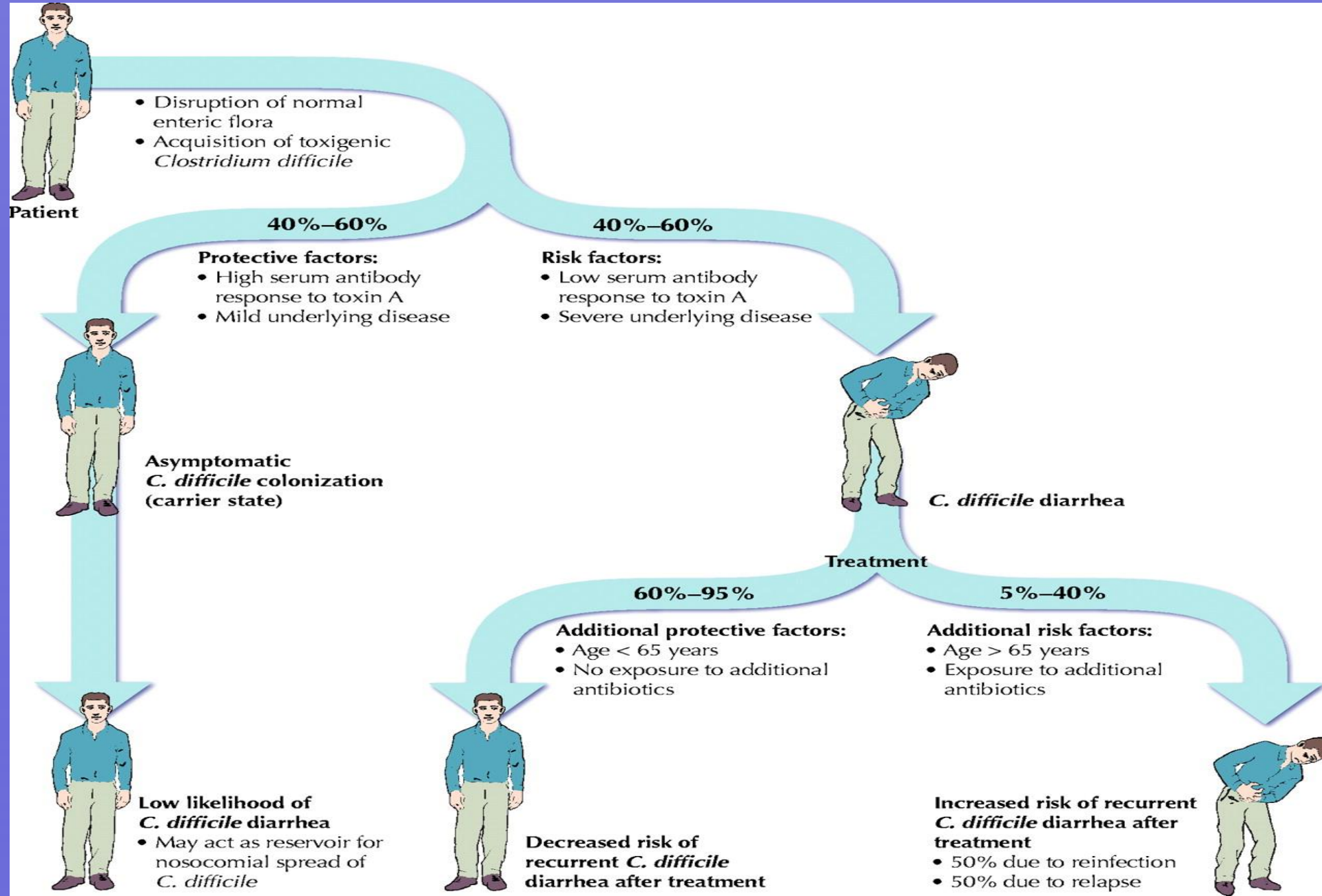
Clostridium difficile

Brat to Tyrant



- 50% infants colonised with CD
- 3% adults colonised, but greater in health institutions
- Only an issue when toxin A, B or hypervirulent CD binary toxin strain BI/NAP1/027 produced
- Mortality in the northern hemisphere has more than doubled from 3.6% to 8% since year 2000

Clostridium difficile



Clostridium difficile

Issues

- **Antibiotics** (use, type and stewardship)
- **Hand Hygiene** (spores alcohol resistant)
- **Environmental** cleaning (scrupulous,
chlorine, how to monitor)
- Binary toxin ↑ mortality (8%)
- Recurrence - discharge to LTCF
implications
- Faecal transplants

? UTI Empiric ABs when

A. No indwelling urinary catheter:

1. Acute dysuria (pain on urination) *alone, or*
2. Fever $\geq 37.9^{\circ}\text{C}$ or 1.5°C above baseline **plus at least one** of the following:
 - (a) New or worsening urgency
 - (b) Frequency
 - (c) Suprapubic pain
 - (d) Gross haematuria (new)
 - (e) Costovertebral angle tenderness
 - (f) Urinary incontinence

? UTI Empiric ABs when

B. Indwelling urinary catheter:

At least one of the following

- (a) Fever $\geq 37.9^{\circ}\text{C}$ or 1.5°C above baseline
- (b) New costovertebral angle tenderness
- (c) Rigors
- (d) New onset of delirium

UTI LTCF

Pyuria, Dipstick, WBC

- 90% colonised urines have pyuria (WBC)
- 25 – 50% residents will have positive white cells/leucocytes in the urine without UTI
- A negative dipstick or culture effectively rules out UTI (only ? 20% of all residents)

? Respiratory Infection

Empiric Antibiotics

1. Fever > 38.9°C plus at least one of the following:
 - (a) Respiratory rate > 25 breaths per minute
 - (b) Productive cough

2. Fever > 37.9°C but ≤ 38.9°C plus cough plus at least one of the following:
 - (a) Pulse > 100 beats per minute
 - (b) Delirium
 - (c) Rigors
 - (d) Respiratory rate > 25 breaths per minute

? Respiratory Infection

Empiric Antibiotics

3. Afebrile and high risk COPD (age $\geq$ 65 years)
 - New or worsening cough **and** purulent sputum

4. Afebrile without COPD
 - New cough, purulent sputum plus at least one of the following:
 - Respiratory rate $>$ 25 breaths per minute
 - Delirium

Pneumonia & Pulse Oximetry

Pulse oximetry good if fully aware of its limitations

A normal pulse oximetry is difficult to interpret in LTCF residents with baseline functional and mental deficits

But a saturation of less than 90% oxygen
+ *respiratory rate >25 is suggestive*
impending respiratory failure

Then consider some urgent interventions
e.g. aggressive suctioning, chest physical therapy & bronchodilators to improve the residents condition. Nursing staff in a LTCF would need to determine their own comfort level based on their knowledge of the resident's medical problems

Pneumonia

Pulse Oximetry

A single oxygen saturation of $<94\%$ is 80% sensitive and 91% specific for the diagnosis of pneumonia with a PPV of 95%

A decrease in oxygen saturation of 3% from baseline is less sensitive (73%) but more specific (100%) for pneumonia with a PPV of 100%