

OVERVIEW

- 3 clinical cases in Starship between 2000-2010
- Worldwide & New Zealand Epidemiology
- Diagnosis & Management of Tetanus
- The preventable cost of tetanus

CLINICAL CASE: LB

- 4 yr old unimmunised Caucasian boy from Auckland
- · Presented to ED with 1 day history of jaw stiffness
- History of trauma:
- Stood on a plastic toy 3 wks ago with puncture wound to heel
 Stood on a thorn 2 days prior with a small scratch
- · Rapid progression to generalised body spasms
- Clinically noted to have generalised ¹tone with opisthotonic spasms occurring every 10 minutes with variable heart rate and reduced mouth opening
- In ED, he was given TIG & metronidazole
- Admitted to PICU for management of spasms & ventilatory support

CLINICAL CASE: LB

• In PICU:

- Ventilated & sedated with midazolam infusion initially then required active cooling & paralysis for high temperatures
- Autonomic instability including hypertension
- Received nasogastric nutrition

CLINICAL CASE: LB

• In PICU:

- Complications:
 - Failed extubation after 2 weeks and required insertion of tracheostomy
 - * Pneumonia requiring prolonged antibiotic treatment
 - × PICC line wound infection with MRSA
 - \times Significant pain issues managed with methadone and diazepam
- Transferred to ward with tracheostomy after 3 weeks admission in PICU

Treatments included:

- × Intragam P
- \times 2 weeks treatment with Metronidazole
- × Orthopaedic debridement of foot wound
- Extensive discussions regarding immunisation followed by his first immunisation with Infanrix-Hexa
 Clonidine for hypertension
- × 15 days of Magnesium infusion
- × High dose oral vitamin C supplementation due to parental request

CLINICAL CASE: LB

• On the ward:

- SLT assessment improvement in voice quality & gradual increase in oral intake with subsequent removal of NGT
- Decannulation of tracheostomy just under 1 month from admission
- 2nd immunisation of Infanrix-Hexa given
 Ongoing PT and OT input with improvement in core strength & balance
- Ongoing PT and OT input with improvement in core strength & balance

Discharged with physiotherapy plan

• Follow-up in Paed ID clinic 1 month from discharge:

- Observed to run into clinic
- Riding bicycle with trainer wheels removed
- e ?altered hearing awaiting hearing assessment
- ← Due for further DTaP, IPV, Hep B & MMR
- Parental reports of uptake of ongoing immunisation including all older siblings

CLINICAL CASE: MP

- 16m old unimmunised Maori girl from Northland
- Presented to Whangarei Hospital with 4 day history of increasing respiratory effort (?infective exacerbation of asthma)
- Ongoing deterioration and subsequently intubated
- Transferred to Starship PICU 2 days following presentation
- In PICU:
- Developed episodic breath holding attacks with associated stiffness and mouth clenching
- Treated with IV Cefotaxime, Vancomycin and Acyclovir.
- e Normal CSF including viral PCR and cultures
- Normal CT and MRI brain
- Normal EEG

CLINICAL CASE: MP

• In PICU:

- Given TIG in view of opisthotonic posturing to cover for potential tetanus and penicillin was commenced.
- Extubated on day 3 and transferred to ward
- Readmitted on day 9 due to increasing spasms and requirement for IV diazepam
- c Commenced on morphine infusion
- e Penicillin was changed to metronidazole on day 10
- Re-examination found two skin cracks between her great & 2nd toe which was debrided
- c Transferred back to ward on day 15 with PICC line and NJT for feeding.

CLINICAL CASE: MP

• On the ward:

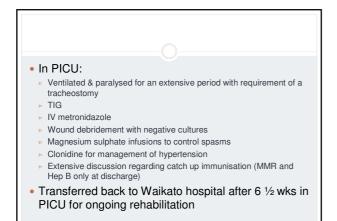
- e Brief spasms treated with intermittent PR diazepam
- Congoing SLT, PT and OT input
- Discharged home on day 23 following immunisations with DTPa, Hib/HepB and Polio.
- $\ensuremath{\scriptscriptstyle \subset}$ Ongoing community paediatric $% \ensuremath{\mathsf{and}}$ and neurodevelopmental follow-up

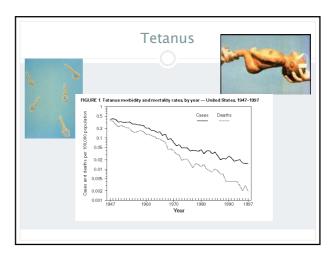
Her immunisations:

- For catch up immunisations of HepB and DTPa with GP
- MMR deferred for 6 months due to use of TIG
- For booster tetanus 12 months later and again at 4 years of age

CLINICAL CASE: LA

- 9 yr old unimmunised Caucasian girl from Waikato
- Presented to GP with trismus and spasms following an infection of a leg wound
- Admitted to Waikato ICU with clinical tetanus & transferred to PICU Starship



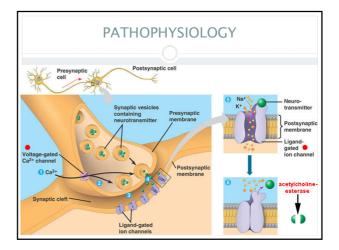


CLOSTRIDIUM TETANI

- Obligate anaerobic bacillus
- 2 toxins: tetanospasmin & tetanolysin
- Mature organisms lose their flagellae & develop a terminal spore
 Spores are extremely stable and retain ability to germinate
- therefore can cause disease indefinitely

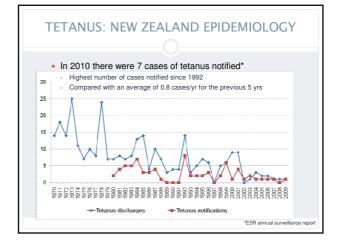
 Spores are found worldwide as constituents of soil & in GI tracts
- Spores are found worldwide as constituents of soil & in Gi tracts of animals

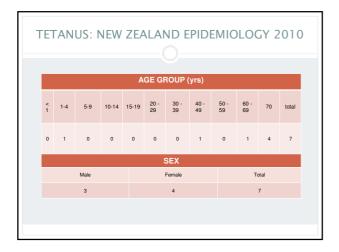




TETANUS: EPIDEMIOLOGY

- Tetanus is an important cause of hospital admission and death in parts of the world with limited vaccination programmes
- At least 1 million cases require hospital treatment worldwide every year
- ~400,000 deaths from tetanus each year
- Neonatal tetanus accounts for ~ half of the tetanus deaths in developing nations
- In developed countries, injuries account for ~70% of cases
 Evenly divided between punctures and lacerations





Tetanus Immunisation

- Universal childhood immunisation with tetanus toxoid containing vaccines in NZ since 1958
- Still elderly cohort in NZ never fully immunised
- Tetanus in NZ usually follows minor injury
- Vaccine is prepared from cell free toxin treated with formaldehyde to create toxoid
- Stimulates the production of antitoxin, providing immunity against the effects of the toxin. It does not prevent *C.tetani* from growing in a contaminated wound.

Tetanus Immunisation....cont.

- Highly immunogenic vaccine
 - 100% effective when given to pregnant women to prevent neonatal tetanus
- Studies show 100% infants have protective levels after 3 doses at least 4 weeks apart
- Antibody decay studies predict 5 year protection by 3 infant doses: booster at 5 years should give at least further 21 years protection
- Convenience and administrative issues determine frequency of boosters (10 or 20 yearly?)
- × NZ: at 11, 45 and 65 years

TETANUS: CLINICAL MANIFESTATIONS

- ← Incubation period is usually 3-21 days (median 8 days)
- Generalised type is most common
- ⇐ Trismus, risus sardonicus
- +/- abdominal rigidity
- e Opisthotonic posturing
- upper airway obstruction & respiratory compromise
- e Autonomic dysfunction with severe sustained/labile BP & arrhythmias
- progression for ~2wks, approx 10% mortality, or recovery is usually complete
- Recurrent tetanus may occur if patient does not receive active immunisation

TETANUS: CLINICAL MANIFESTATIONS

· Localised tetanus

- Involves rigidity of muscles at site of spore inoculation
- May be mild & can resolve spontaneously
- More commonly is a prodrome for generalised tetanus

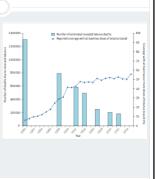
Cephalic tetanus

- Special form of localised disease
- Almost always follow an apparent head injury & causes a facial paresis

TETANUS: CLINICAL MANIFESATIONS

Neonatal tetanus

- Follows an infection of the umbilical stump when the mother is inadequately immunised
- Newborns present with generalised weakness, feeding difficulties, rigidity & spasms
- Mortality rate exceeds 90%
- Apnoea is leading cause of death in 1st wk of life & sepsis in 2nd wk.



TETANUS: DIAGNOSIS

- · Diagnosis is made strictly on clinical grounds
- · Laboratory testing cannot confirm or exclude the diagnosis C.tetani cultures from wound are not useful
- Negligible serum tetanus antibody concentrations can support the diagnosis
- · Strychnine poisoning is the only condition that truly mimics tetanus
- · Other differential diagnosis include: dental infections, tonsillitis, parotitis, TMJ disease and dystonic reactions to medications

TETANUS: MANAGEMENT

DIAGNOSIS & STABILISATION (first hr after presentation)

- Assess airway & ventilation: consider ETT
- Routine bloods including antitoxin level, strychnine, CK & urinary myoglobin
- Determine portal of entry, incubation period, period of onset and immunisation history
- Administer benztropine or diphenhydramine to rule out dystonic reaction
- Administer benztropine or benzodiazepine to control spasm & decrease rigidity
- Transfer to a dark quiet area with reduced stimulation

TETANUS: TREATMENT

EARLY MANAGEMENT PHASE (first 24hrs)

- Tetanus immunoglobulin IM or IVIG
- Active immunisation
- Metronidazole IV Q6H for 7-10 days
- Consider tracheostomy
- Wound debridement as indicated
- Nutrition: NGT or TPN
- Benzodiazepines to control spasms & provide sedation
- « Consider neuromuscular blockade if unable to achieve adequate spasm control

TETANUS: TREATMENT

• INTERMEDIATE MANAGEMENT PHASE (NEXT 2-3WKS)

- C Treat sympathetic hyperactivity with beta blockers or morphine
- Sustained bradycardia usually requires a pacemaker
- Prophylactic heparin
- e Pressure cares
- Maintain benzodiazepines until neuromuscular blockade has been terminated & then taper over 2 to 3 weeks as spasms diminish

Rehabilitation planning

- CONVALESCENT PHASE (2-6 WKS) Physiotherapy once spasms have completely resolved
- Repeat dose of tetanus vaccine
- Schedule for 3rd dose of toxoid to be given 4wks after the 2nd dose

TETANUS: Prevention Summary

- · Tetanus is a vaccine preventable disease:
- · passive immunisation and wound management is a poor second
- · 3 injections at intervals of 4-8 weeks recommended
- Immunity is established with 2 doses and 3rd dose prolongs its duration
- Boosters are needed in adult life
- · Full series of maternal immunisations would be ideal but even one dose confers substantial protection against neonatal tetanus
- · Previous tetanus does not confer immunity
- No herd immunity possible

WHAT HAPPENED TO THESE CHILDREN?

- Parental reports of compliance with ongoing immunisation in follow-up I.D. clinic
- No further immunisations given following hospital discharge Siblings only received 2 sets of immunisations during the period LB was admitted

- Lives with large extended whanau in a rural setting
- · Limited access to primary healthcare
- No further immunisations given following hospital discharge according to NIR

- Family strongly against immunisationNo tetanus immunisations given in hospital
- No further immunisations in the community following discharge home

