

# QUEEN



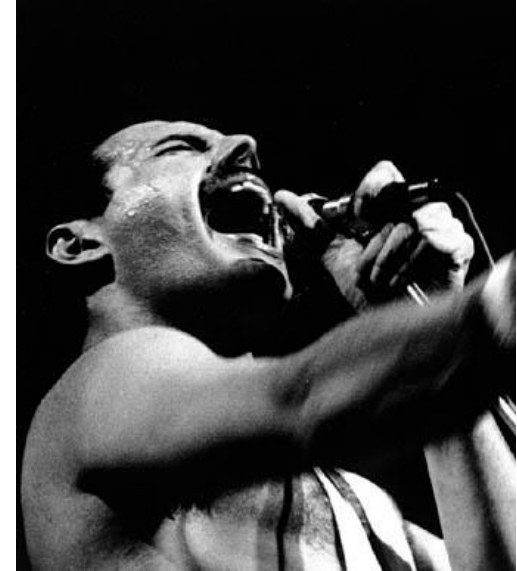
**DON'T STOP ME NOW**

Pathogenesis, progress and prevention in  
management of diabetes foot disease

Brian Kennon, Southern General, Glasgow

# Why the interest in foot disease?





**In which year did Freddie Mercury die?**

**1991**

**1993**

# Outline



- Introduction
- Why is diabetes so bad for feet?
  - Neuropathy - Infection - Ischaemia
- Whose at risk of diabetes foot disease?
- What's the outcome of diabetes foot disease?
- How can we improve the outcome?
- Summary



# Foot Facts I

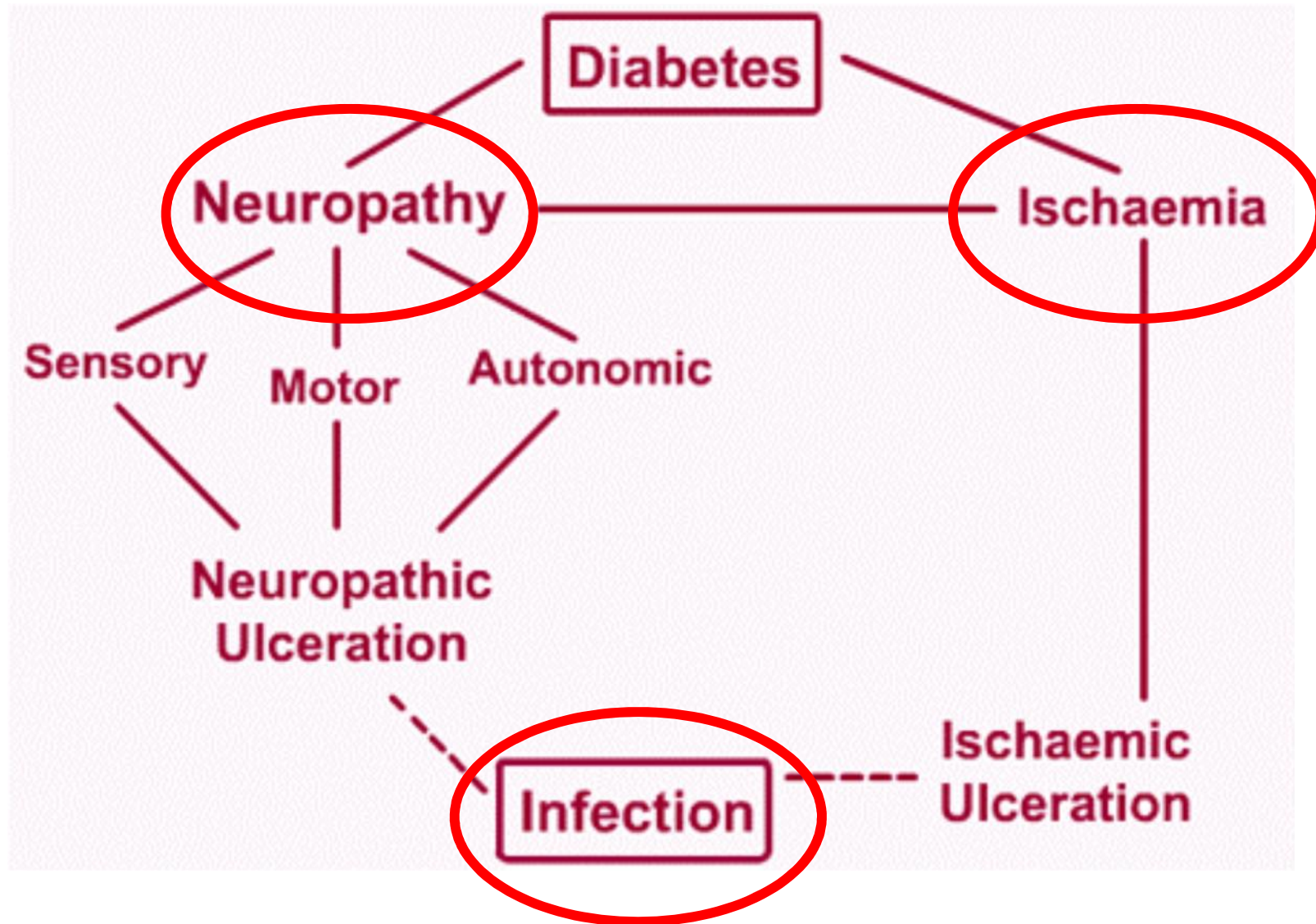
- Every 30 seconds a leg is lost to diabetes somewhere in the world
- In developed countries, up to 5% of people with diabetes have a foot problem
- 47-50% of diabetes-related admissions
- Average length of admission is 6 weeks
- Estimated that diabetic foot disease costs £662M annually in England & Wales



# Foot Facts II

- 85% amputations preceded by a foot ulcer
- One in every six people with diabetes will have a foot ulcer during their lifetime
- 60 % of minor and 43% of major lower limb amputations are in individuals with diabetes
- Well-organised diabetic foot care teams, good diabetes control and well-informed self care can significantly reduce amputation rates

# Why is Diabetes so bad for feet?



# Is there neuropathy?



## Sensory

- 10 g monofilament
- Don't test over callus
- Test over 10 sites



## Motor

- Neuropathic foot with high medial arch and prominent metatarsal heads



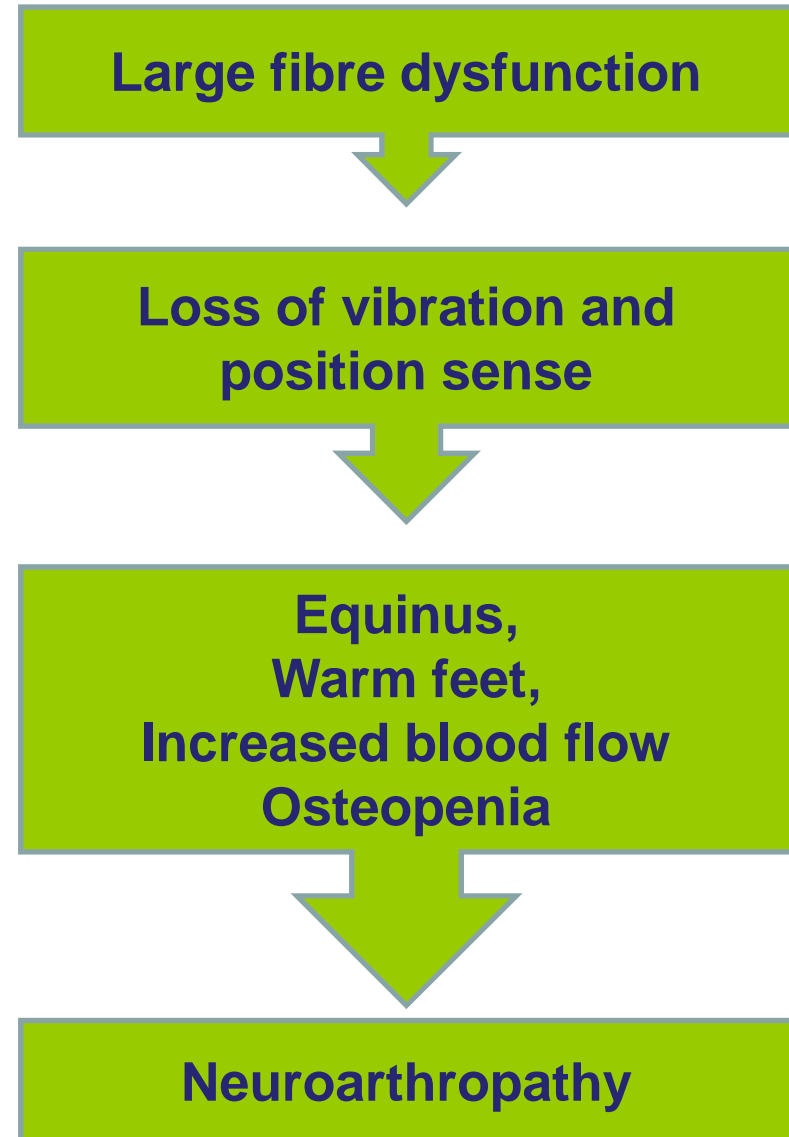
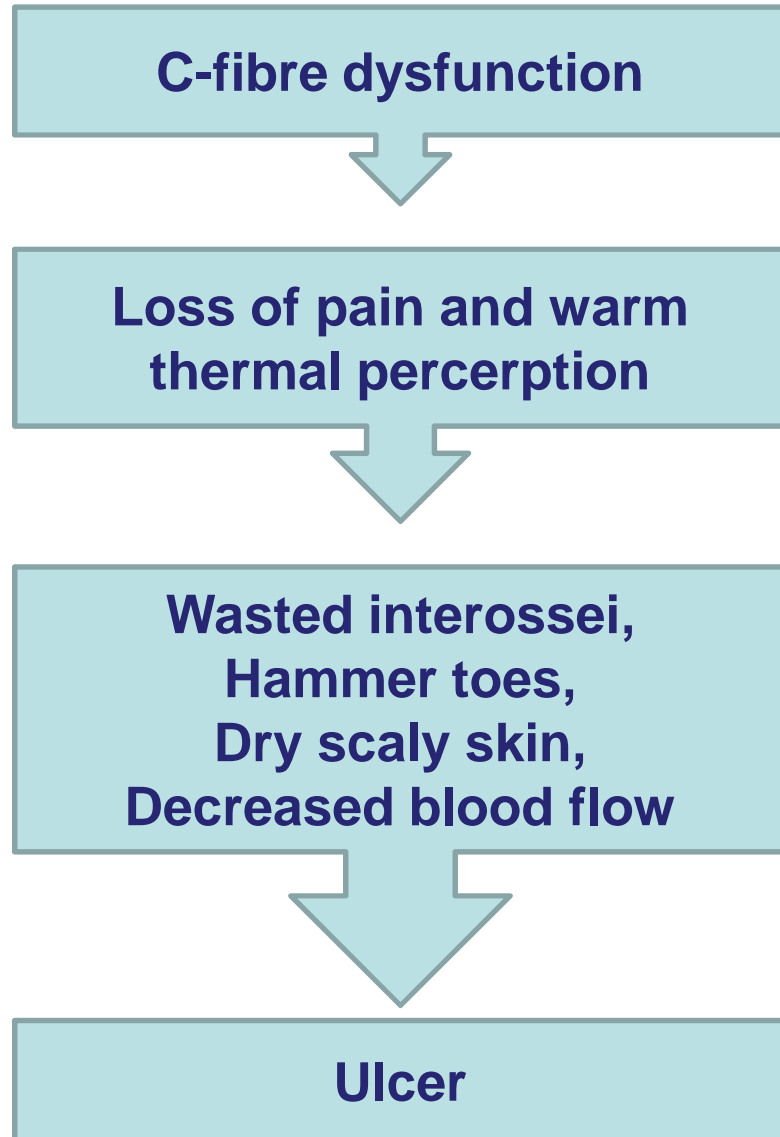
## Autonomic

- Dry skin with fissuring
- Distended veins on dorsum of foot





# Neuropathy & Pathology



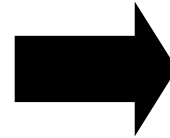
# Neuropathic Ulceration



- 40 years old man
  - Type 1 DM for 22 years
  - Chronic poor control
  - Retinopathy, Nephropathy & Neuropathy
  - Presents with an ulcer of 2 weeks duration
- 
- Classic neuropathic ulcer over a pressure area
  - Mainstay of treatment is pressure relief

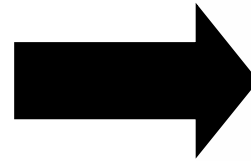


# Neuropathic Ulcersion



- Neuropathy affects up to 30% of patients with diabetes
- Increased with:
  - age
  - duration of diabetes
  - smoking
  - gender
  - sub-optimal glycaemic control

# Recurrent Ulceration



## Aetiology?

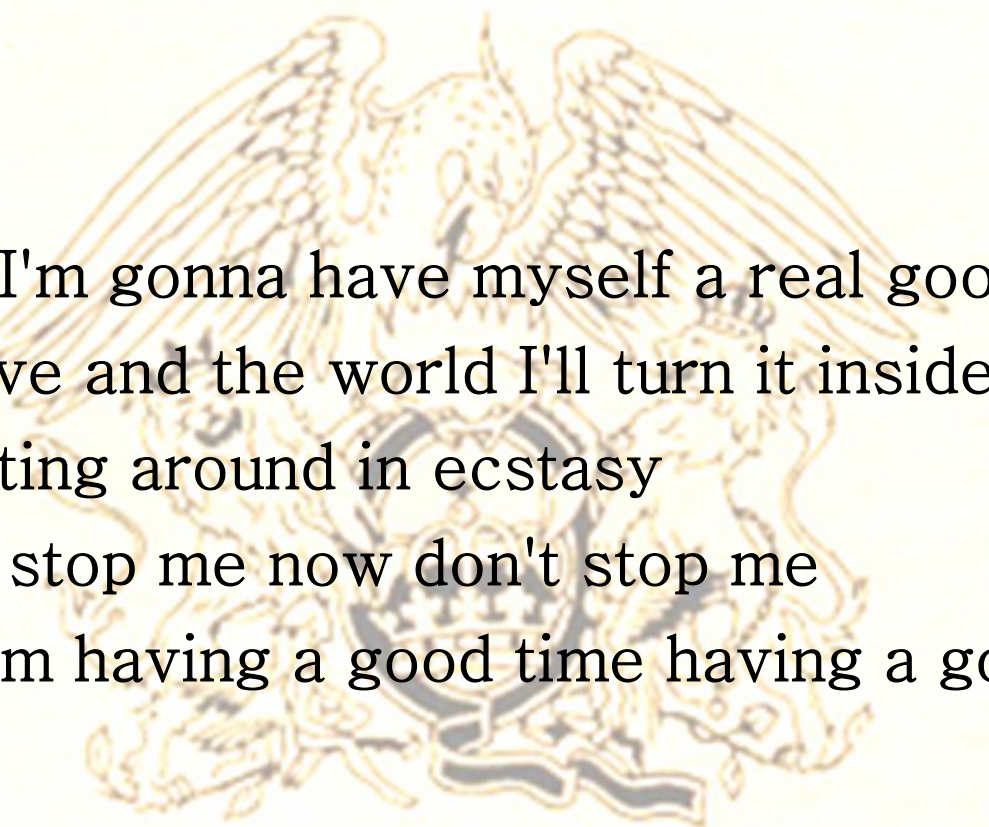


Why did the patient develop an ulcer?

How do we prevent it from recurring?

1 year 34%

5 years 70%

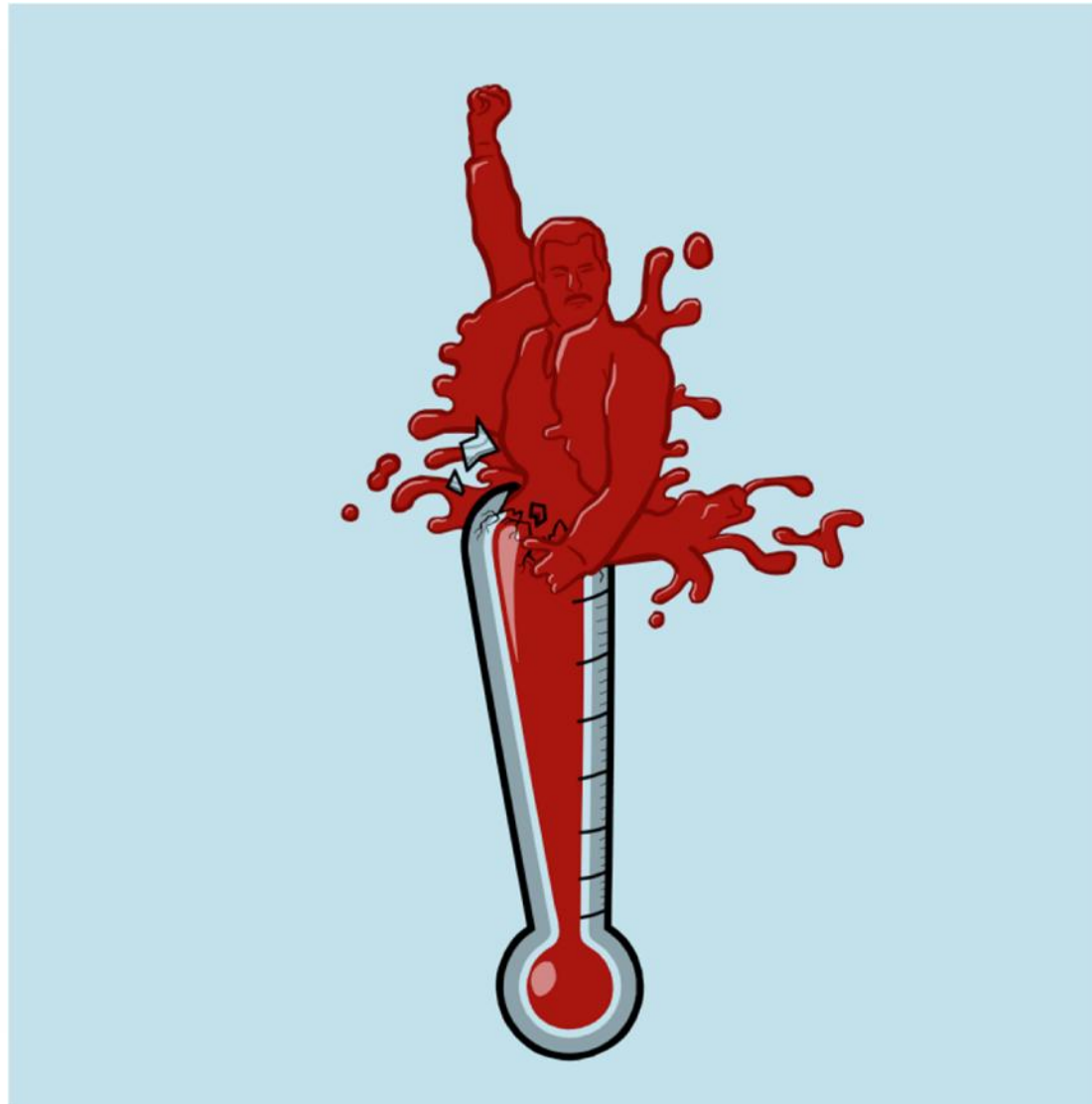


Tonight I'm gonna have myself a real good time  
I feel alive and the world I'll turn it inside out - yeah  
And floating around in ecstasy  
So don't stop me now don't stop me  
'Cause I'm having a good time having a good time

DON'T STOP ME NOW

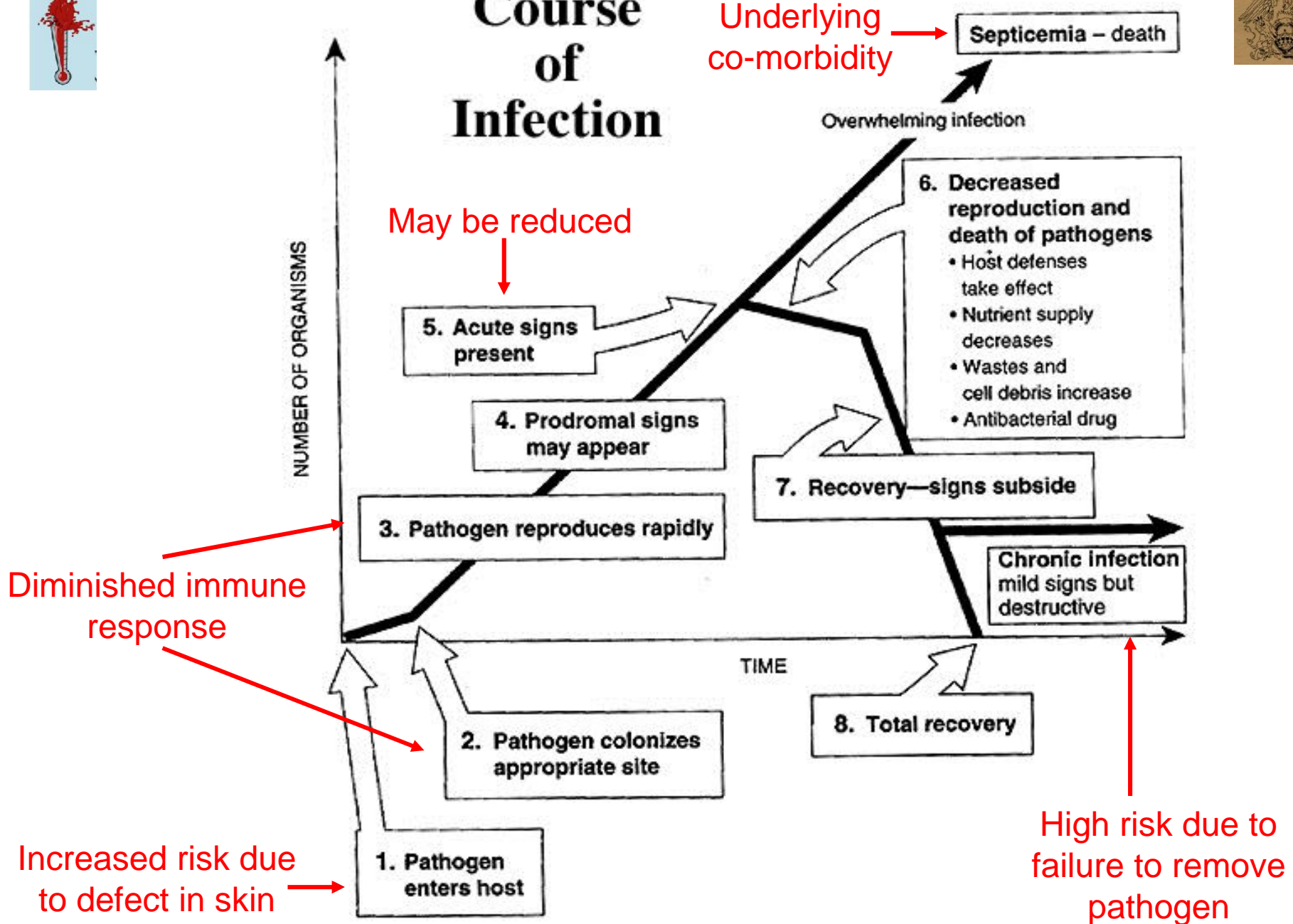
TIP: Footwear is often the aetiology of ulceration

Is there evidence of infection?





# Course of Infection





**What % of diabetic foot ulcers are infected at initial presentation?**

**58%**

**78%**





# High prevalence of ischaemia, infection and serious comorbidity in patients with diabetic foot disease in Europe. Baseline results from the Eurodiale study

10 European countries, 14 centres, 1229 consecutive ulcers 2003-4

Stage	Definition	Number of patients	Percentage of study population
A	PAD -, infection -	270	24
B	PAD -, infection +	305	27
C	PAD +, infection -	205	18
D	PAD +, infection +	347	31

← 58% ulcers are infected

Infection was diagnosed if two or more were present: frank purulence, local warmth, erythema, lymphangitis, oedema, pain, fever and foul smell.

82% subjects hospitalised had infection      Rate higher if PAD at 63%v53%



# How to assess the severity...

Clinical Manifestation of Infection	PEDIS Grade	IDSA Infection Severity
No symptoms or signs of infection	1	Uninfected
Infection present, as defined by the presence of at least 2 of the following items:		
<ul style="list-style-type: none"> <li>• Local swelling or induration</li> <li>• Erythema</li> <li>• Local tenderness or pain</li> <li>• Local warmth</li> <li>• Purulent discharge (thick, opaque to white or sanguineous secretion)</li> </ul>		
Local infection involving only the skin and the subcutaneous tissue (without involvement of deeper tissues and without systemic signs as described below). If erythema, must be >0.5 cm to ≤2 cm around the ulcer. Exclude other causes of an inflammatory response of the skin (eg, trauma, gout, acute Charcot neuro-osteoarthropathy, fracture, thrombosis, venous stasis).	2	Mild
Local infection (as described above) with erythema > 2 cm, or involving structures deeper than skin and subcutaneous tissues (eg, abscess, osteomyelitis, septic arthritis, fasciitis), <b>and</b> No systemic inflammatory response signs (as described below)	3	Moderate
Local infection (as described above) with the signs of SIRS, as manifested by ≥2 of the following: <ul style="list-style-type: none"> <li>• Temperature &gt;38°C or &lt;36°C</li> <li>• Heart rate &gt;90 beats/min</li> <li>• Respiratory rate &gt;20 breaths/min or PaCO<sub>2</sub> &lt;32 mm Hg</li> <li>• White blood cell count &gt;12 000 or &lt;4000 cells/μL or ≥10% immature (band) forms</li> </ul>	4	Severe <sup>a</sup>

How have your blood sugars been?

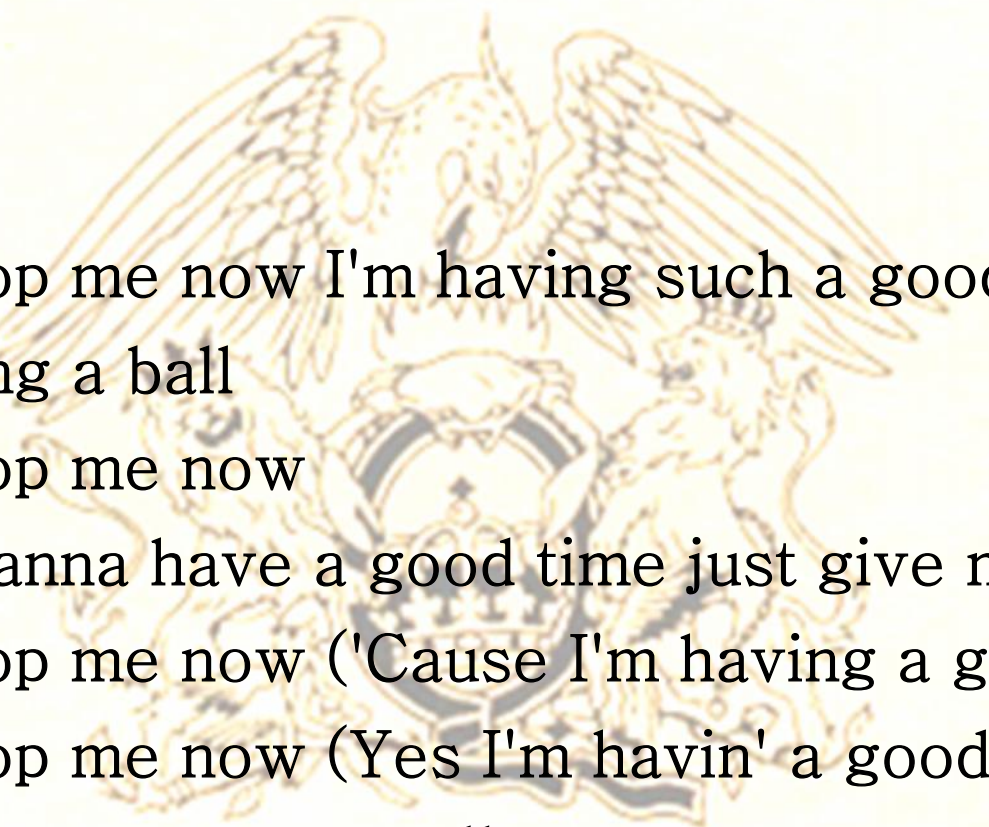
3 things to evaluate: **P**erson, **L**imb and **W**ound



# Infection



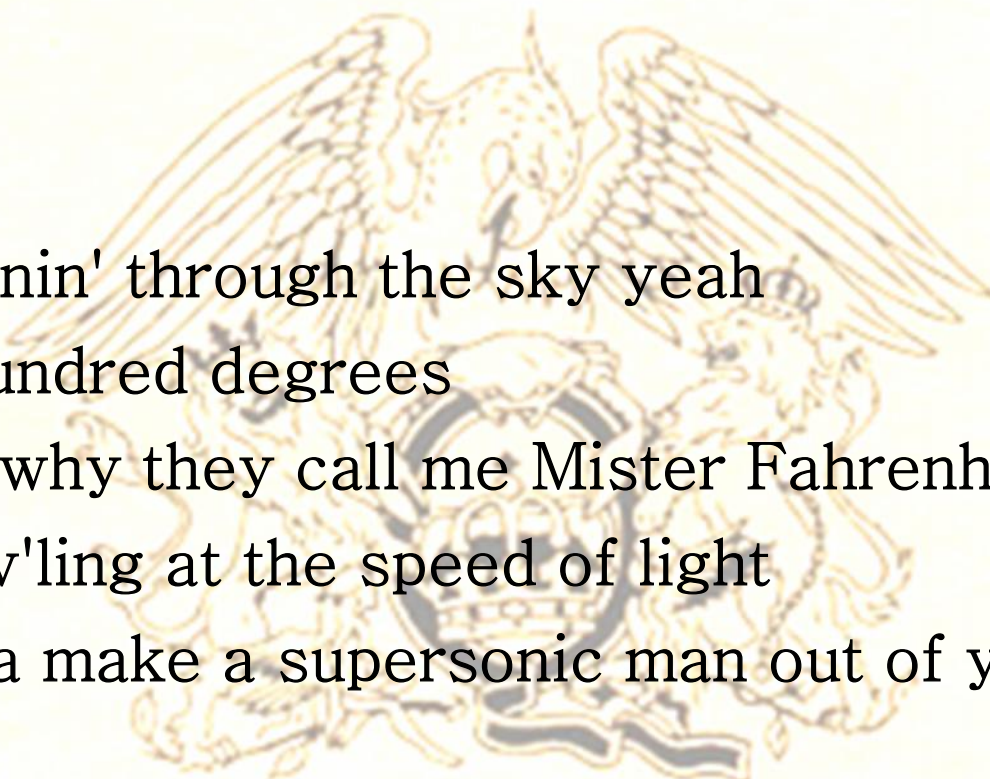
- 43 years old man
- Type 2 DM for 3 years
- PHx Obesity
- Neuropathy
- Returned from holiday with a swollen hot red foot
- Pain in his foot
- Systemically unwell



Don't stop me now I'm having such a good time  
I'm having a ball  
Don't stop me now  
If you wanna have a good time just give me a call  
Don't stop me now ('Cause I'm having a good time)  
Don't stop me now (Yes I'm havin' a good time)  
I don't want to stop at all

DON'T STOP ME NOW

TIP: Pain in a neuropathic foot consider  
urgent surgical assessment



I'm burnin' through the sky yeah  
Two hundred degrees  
That's why they call me Mister Fahrenheit  
I'm trav'ling at the speed of light  
I wanna make a supersonic man out of you

DON'T STOP ME NOW

TIP: Treat infection aggressively

TIP: Some people find a prosthesis a  
positive experience



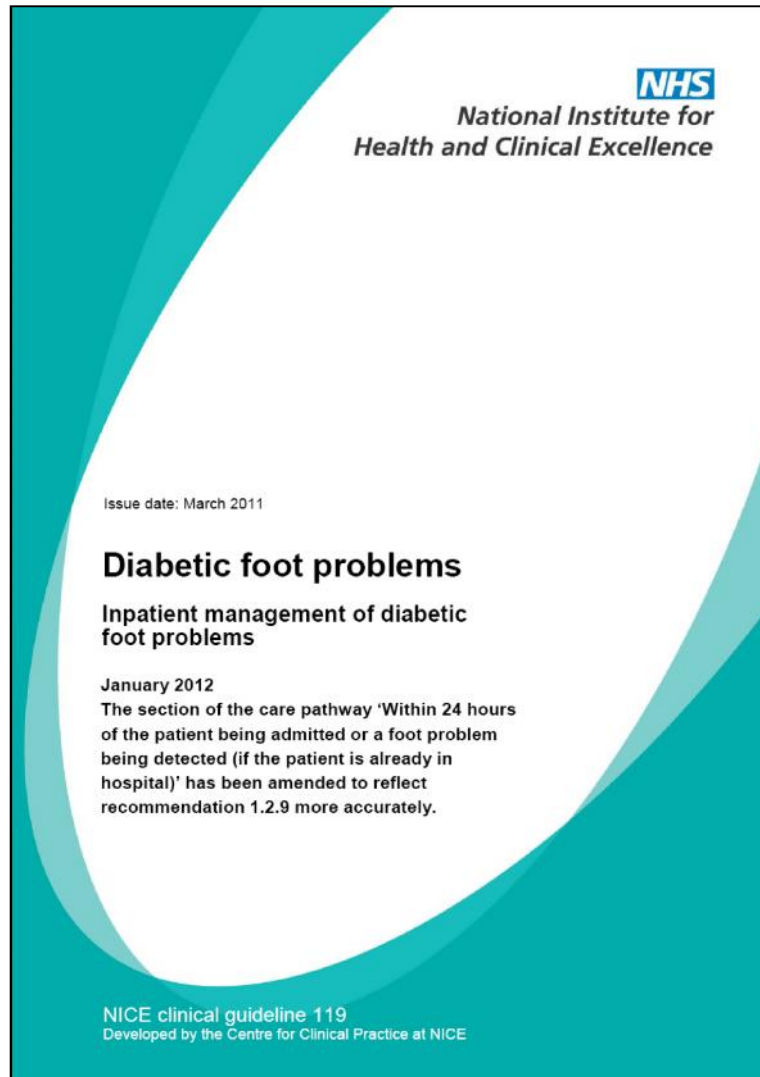
# Therapeutic Options

- Treat infection
  - IV antibiotics, drainage & VAC device
- Maximise pressure relief
  - Bed rest & orthosis
- Maximise blood supply
  - Good blood supply
- Control blood glucose
  - Very erratic due to illness
- Manage CV risk factors
  - On appropriate treatment
- Surgical intervention
  - Amputation?





# What antibiotics should you use?



## What is the clinical effectiveness of different antibiotic regimens and antimicrobial therapies for diabetic foot infections (with or without osteomyelitis)?

Systematic search: 9817 studies.

13 studies were included:

- all were head-to-head trials
- no 2 studies with the same pair-wise comparisons.

### Quality of the evidence

**Evidence was inconclusive**

**Not possible to make recommendations on individual antibiotics**



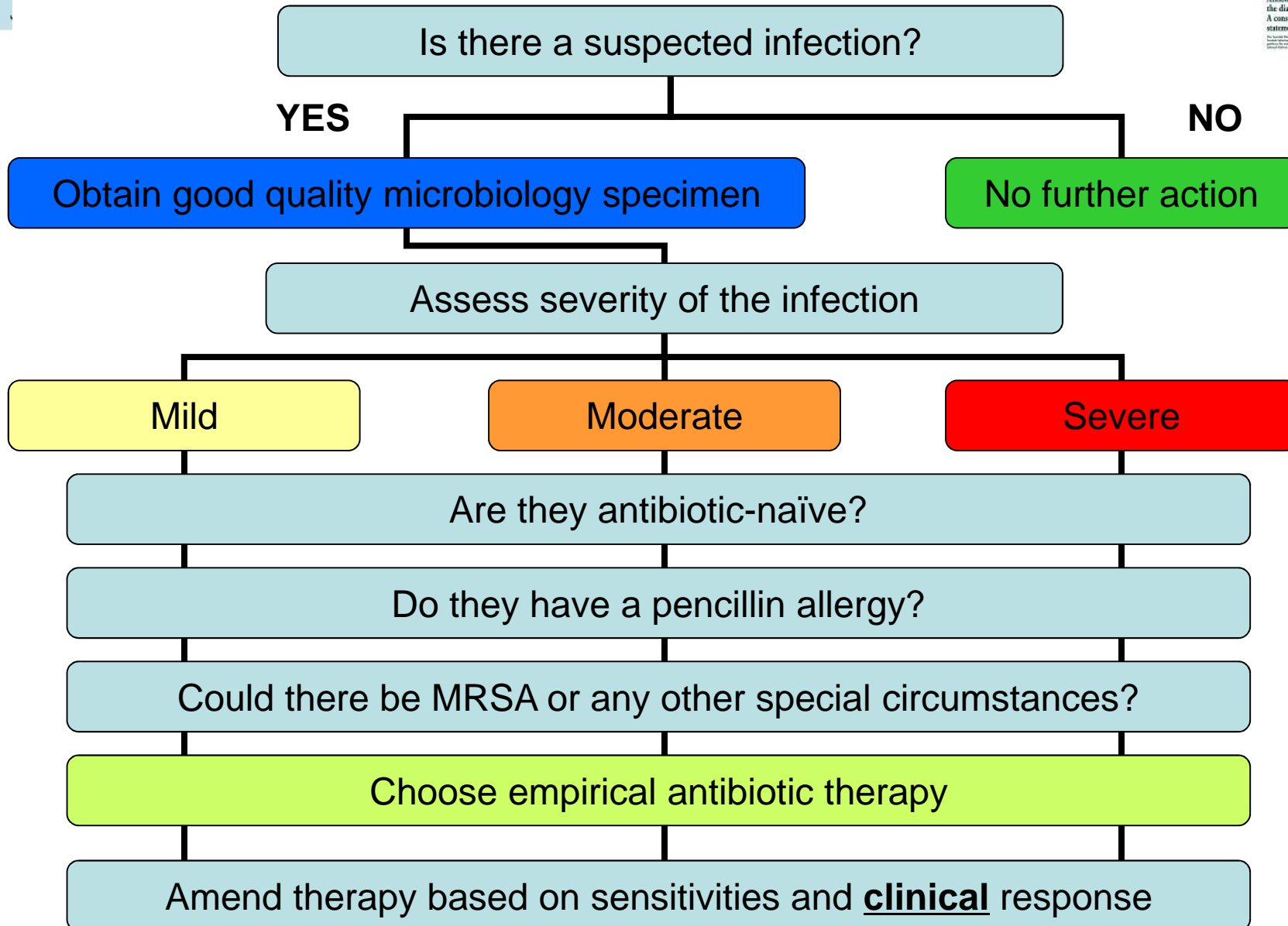
# Recommendations

- Each hospital should have antibiotic guidelines
- Therapy for suspected osteomyelitis should not be delayed pending MRI
- Empirical therapy should be started based on severity of infection
- Definitive regimen is informed by microbiology results
- Lowest cost drugs appropriate to clinical setting should be used





# Stepwise Approach to Antibiotic Choice





## Mild infection (IDSA) or PEDIS 2: Antibiotic-naïve

Signs of infection with cellulitis <2cm, No systemic illness

Likely pathogens : *Staph. aureus* or beta-haemolytic streptococci.



### Antibiotics

*Primary:* Oral FLUCLOXACILLIN 1 g qds for 5-7 days.

*Oral alternatives:* DOXYCYCLINE 100 mg bd  
or CLINDAMYCIN 300–450 mg qds

**\*\*Second course is rarely effective if first appropriate & unsuccessful\*\***



## Moderate infection (IDSA) or PEDIS 3: Antibiotic-naïve

Either: (a) lymphatic streaking, deep tissue infection or abscess or (b) Cellulitis >2 cm. No systemic illness.



Likely pathogens : *S aureus*, streptococci.and/or anaerobes

### Antibiotics

#### Primary

Oral or IV Flucloxacillin 1 g qds

#### Oral alternatives

Co-trimoxazole 960 mg bd or

Co-amoxiclav 625 mg tds

## Moderate infection (IDSA)/PEDIS 3: Not antibiotic-naïve

Likely pathogens : polymicrobial



### Antibiotics

*Primary* : IV co-amoxiclav 1.2 g tds

*Oral switch* : Co-amoxiclav 625 mg tds, or Co-trimoxazole 960 mg bd.



*Allergic to penicillins*: IV ciprofloxacin 400 mg tds and IV metronidazole 500 mg tds, or IV gentamicin and IV metronidazole 500 mg tds.

– Add vancomycin if MRSA infection is suspected.



## Severe infection (IDSA)/PEDIS 4: Antibiotic-naïve

Any infection accompanied by systemic toxicity. The presence of critical ischaemia of the involved limb may make the infection severe.

**\*\*Generally advised admission to hospital\*\***

Likely pathogens

*S. aureus* or beta-haemolytic streptococci.

Anaerobes, enterobacteriaceae

*Pseudomonas aeruginosa* may also need to be treated.

*P. aeruginosa* is usually a coloniser.



Antibiotics

*Primary*

IV co-amoxiclav 1.2 g tds +/- add gentamicin

*Allergic to penicillins or concern about renal function*

IV ciprofloxacin 400 mg bd and IV metronidazole 500 mg tds.

– Add vancomycin if MRSA infection is suspected.



## Severe infection (IDSA)/PEDIS 4: Specific circumstances

- Recent antibiotic therapy (i.e. preceding 90 days).
- Proven drug-resistant
- Extended-spectrum beta-lactamase-producing (ESBL) *Escherichia coli* or *Klebsiella* spp. : seek specialist advice

### Antibiotics

*Primary* : IV piperacillin/tazobactam 4.5 g tds,  
– Add vancomycin if MRSA infection is suspected

### *Penicillin allergy*

IV ciprofloxacin 400 mg bd and IV metronidazole 500 mg tds.



## Consider empirical treatment of MRSA in the following:

- Hx of previous MRSA infection/colonization within the past year.
- Local prevalence of MRSA (ie, % of all *S.aureus* clinical isolates) is high
- The infection is sufficiently severe that failing to empirically cover MRSA while awaiting definitive cultures would pose an unacceptable risk of treatment failure.





# Osteomyelitis

## Diagnosis

GOLD STANDARD: Bone biopsy

Clinical assessment:

PROBE-TO-BONE Test:

- a. Infected wound and PTB +ve high likelihood
- b. Non-infected wound and PTB –ve very unlikely

## Investigations

Clinical suspicion: initial IXS is plain X-ray

**\*\*May take 2 weeks before any changes of acute osteomyelitis on plain radiograph and thus serial X-rays may be required\*\***

Secondary investigations :

1. MRI
2. Isotope white cell scan
3. Triple phase bone scan (highly sensitive but not specific)





# Osteomyelitis

## **Management:** Surgery v Conservative Therapy

- Local surgery can increase healing time and decrease need for antibiotics
- Lone antibiotic therapy can eliminate infection in 80% of cases of osteomyelitis
- No evidence of best antibiotic regimen
- Recommended duration of therapy 4-6 weeks (if no surgery) may be longer depending on clinical response
- No evidence IV therapy superior to Oral
- Limited evidence if MRSA: add Rifampicin 660mg bd or Fusidic Acid 500mg tds

# Diabetes & PAD



## Diagnosis and assessment of peripheral arterial disease in the diabetic foot

- UKPDS each 1% increase in HbA1c associated 28% increased risk PAD
- Diffuse disease which is often distal
- Less adaptive mechanisms to ischaemia and decreased collaterals
- AV shunting with capillary hypoperfusion
- Often asymptomatic as co-existent neuropathy
- 49% ulcers co-existing PAD
- EURODIALE study only 56% with severe PAD (ABPI <0.5) had vascular imaging and of those only 43% revascularised



# Is there evidence of ischaemia?

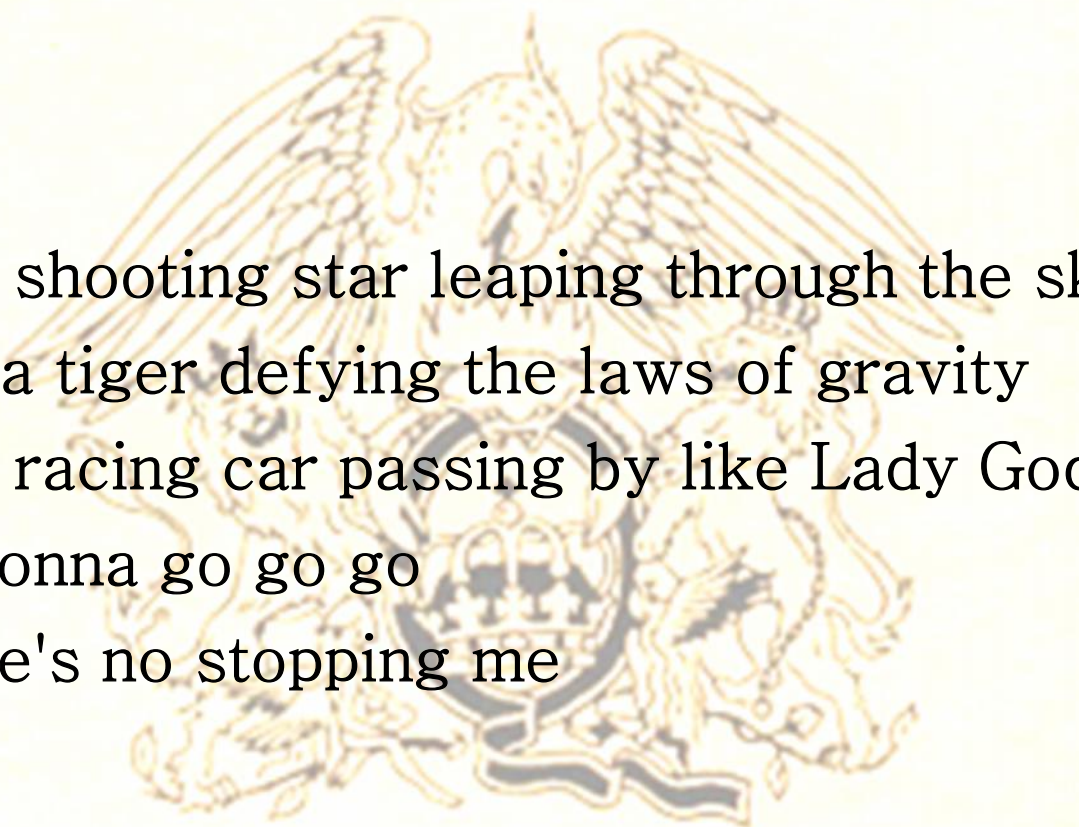


## Clinical assessment

- Assess lower limb pulses & capillary return
  - (30% subjects deficient DP pulse)
- Lift limb above neutral position and note if pallor and then hyperperfusion once dependent
- Doppler examination: normal triphasic. In severe PAD absent or monophasic (often inaccurate)
- Femoral bruit can indicate sig PAD
- ABPI  $<0.9$  or toe: brachial  $<0.7$  indicates probable PAD
- ABPI  $<0.6$  poor wound healing potential



If in doubt or a non-healing ulcer then image



I'm a shooting star leaping through the sky  
Like a tiger defying the laws of gravity  
I'm a racing car passing by like Lady Godiva  
I'm gonna go go go  
There's no stopping me

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TIP: Elevate the leg to assess the severity of PAD

TIP: Do NOT drive wearing orthosis

# Assessing the vascular system?



## Doppler ultrasonography

- Pros: safe, inexpensive, good –ve predictor
- Cons: operator dependent, less reliable for infra-popliteal

## CT angiography

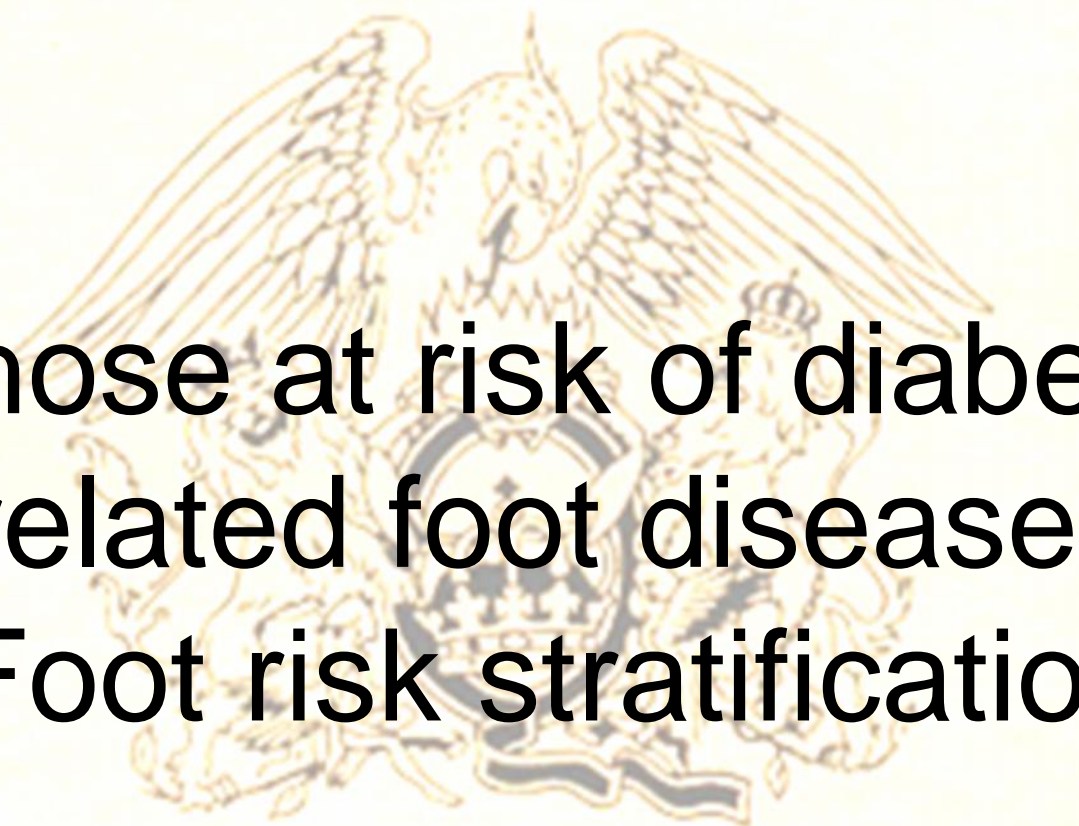
- Pros: 90-95% sens & 92-96% spec detecting >50% stenosis
- Cons: risk of contrast, interference

## Contrast enhanced MRI

- Pros: 96% sens & 97% spec detecting >50% stenosis
- Cons: risk of contrast, limits of MRI

## Digital subtraction angiography

- Pros: supposed gold standard, allows intervention
- Cons: invasive, complication rate 2%, risk of contrast



**Whose at risk of diabetes  
related foot disease?  
Foot risk stratification**

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# Foot screening and risk stratification has been shown to reduce ulcer rate:



Don't know

No



In specific groups

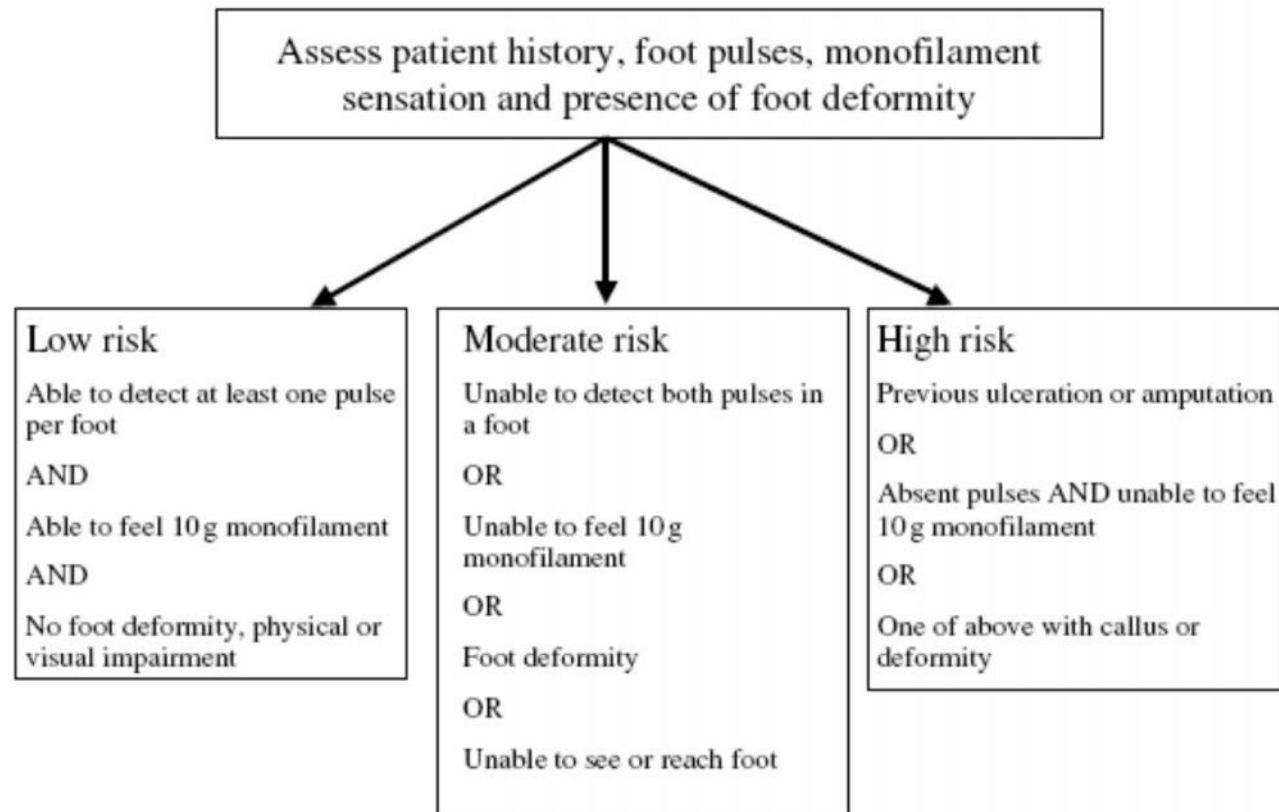
Yes



# What is the evidence for foot screening?



Stratification of foot ulcer risk in patients with diabetes:  
a population-based study



# Ulcer rate depending on risk stratification

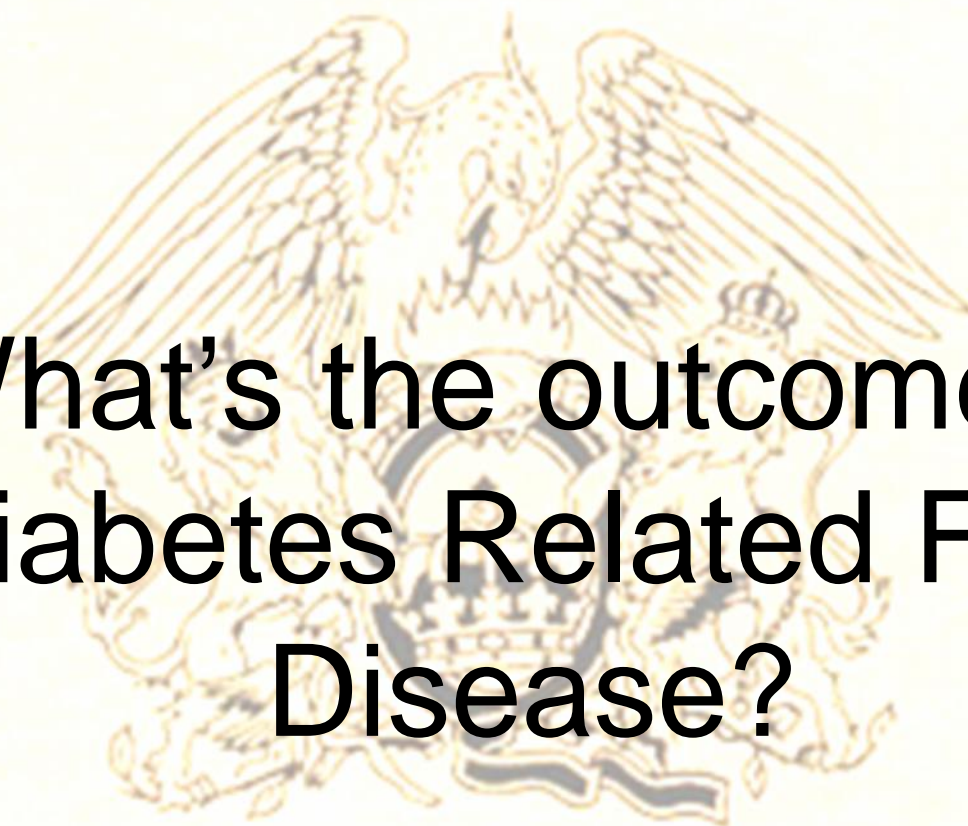


**Table 1** Ulcer outcome for patients who underwent foot risk stratification over 1.7-year follow-up

<i>Overall results</i>	<i>Developed ulcer during follow-up</i>	<i>Did not develop ulcer during follow-up</i>	<i>Total</i>
<b>Risk score</b>			
'High risk'	140 (29.4%)	337 (70.6%)	477
'Moderate risk'	18 (2.3%)	778 (97.7%)	796
'Low risk'	8 (0.36%)	2245 (99.6%)	2253
Total	166 (4.7%)	3360 (95.3%)	3526
	<i>Sensitivity %</i>	<i>Specificity %</i>	<i>Positive predictive value %</i>
High-risk group	84.3 (83.1–85.7)	90.0 (89.0–90.9)	29.4 (27.9–30.9)
High- and moderate-risk group	95.2 (94.5–95.9)	66.8 (65.3–68.4)	
			<i>Negative predictive value %</i>
Low-risk group			99.6 (99.5–99.7)

**High risk v Low risk = x 83 risk ulceration**

**Mod risk v Low risk = x 6 risk ulceration**



# What's the outcome of Diabetes Related Foot Disease?

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**One year after developing a diabetic foot ulcer more patients will have:**

**Died**

**Amputated**

# **Assessing the Outcome of the Management of Diabetic Foot Ulcers Using Ulcer-Related and Person-Related Measures**

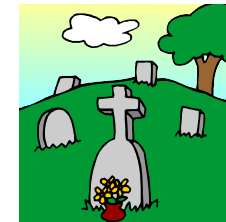


- **Ulcer related outcomes**
  - **Healing, Amputation, Death, Persistent ulcer**
- **Patient related outcomes**
  - **Survival, Ulcer free, Amputation**
- **449 patients recruited over a 3 year period**
  - **352 (78%) superficial ulcers**
  - **183 (41%) infected**
  - **216 (48%) PVD**

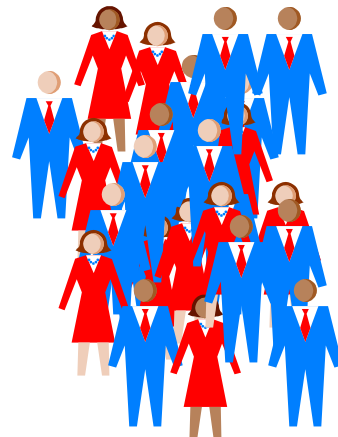
# Assessing the Outcome of the Management of Diabetic Foot Ulcers Using Ulcer-Related and Person-Related Measures



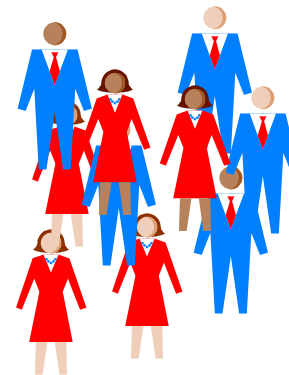
## Person related outcomes



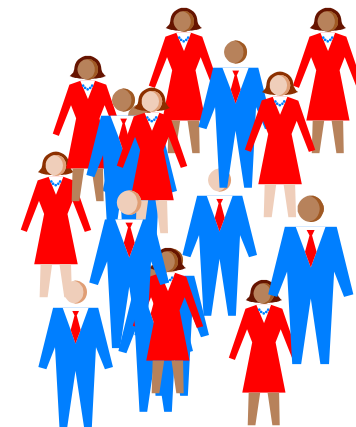
45%



25%



11%



17%



**What factors influence  
outcome?**

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**Which factor(s) presents the greatest risk of an ulcer progressing to amputation?**

**Ulcer down to bone**

**Infection & ischaemia**

# University of Texas wound classification system



- Grade 0            Pre or post ulceration
- Grade 1            Superficial ulcer
- Grade 2            Probing tendon or capsule
- Grade 3            Probing to bone
  
- Stage A            No infection, No ischaemia
- Stage B            Infection
- Stage C            Ischaemia
- Stage D            Infection + Ischaemia

# Validation of a Diabetic Wound Classification System



## Amputation & UoT Class

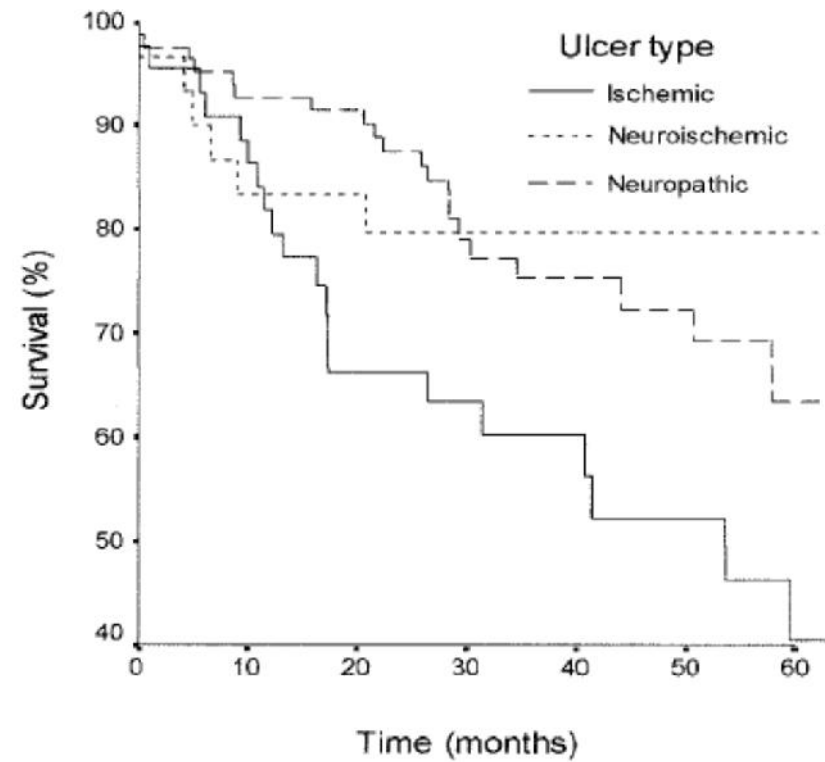
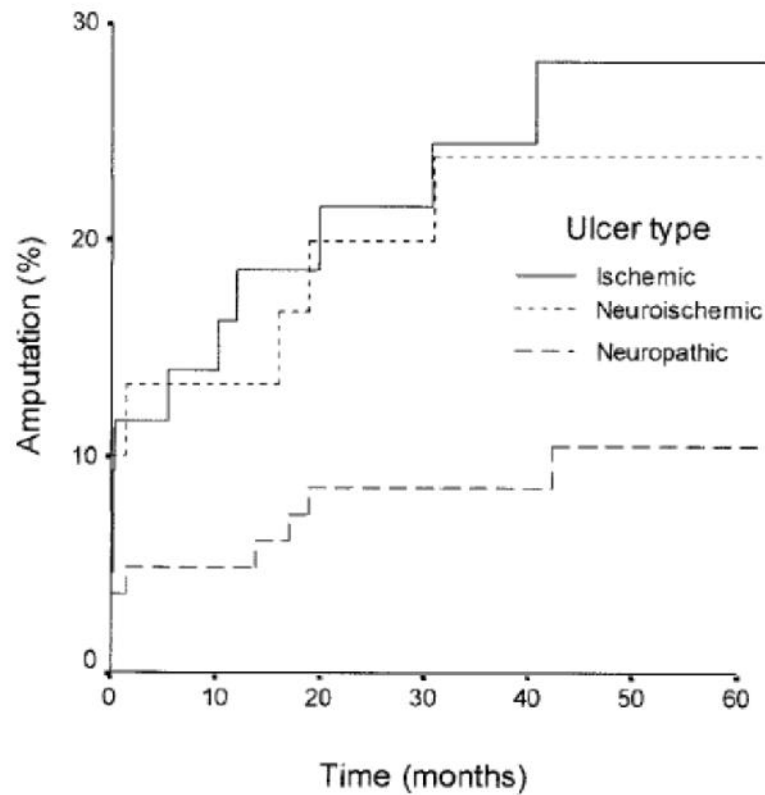
		Grade			
		0 = healed	1 = superficial	2 = tendon or capsule	3 = bone or joint
Stage	A: no infection & no ischaemia	0%	0%	0%	0%
	B: infection	13%	9%	29%	92%
	C: ischaemia	25%	21%	25%	100%
	D: infection & ischaemia	50%	50%	100%	100%

X 11

X 90

Armstrong et al, Diabetes Care 21:855-9 1998

# Ulcer type: amputation & mortality rate





The impact of foot ulceration and amputation on mortality in diabetic patients. I: From ulceration to death, a systematic review

## **Key Messages**

- lower limb ulceration in diabetic patients is associated with amputation and high mortality
- in this systematic review, we quantify the role new-onset ulceration plays in mortality in diabetic patients
- five-year mortality rates after ulceration were around 40%
- risk factors for death commonly identified were increased age, male gender, peripheral vascular disease and renal disease

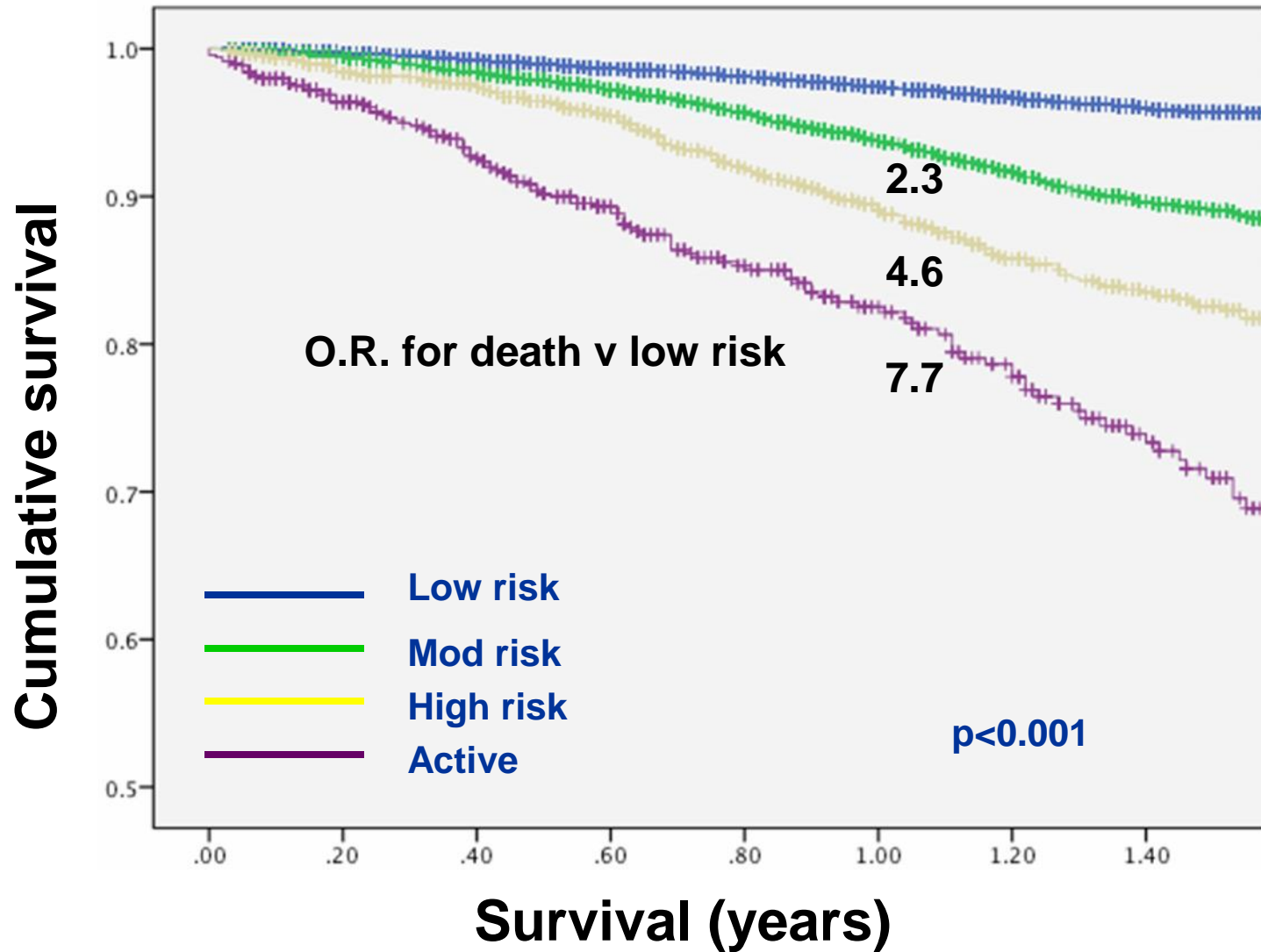
# It's not just ulcerated subjects!

## Foot Risk Stratification & One Year Mortality

- A cohort study of 33,268 subjects in Greater Glasgow and Clyde, Scotland who had diabetes foot risk stratification performed from the 1st January 2009 onwards.
- 1 year mortality was calculated from the time of screening according to foot risk score

<b>Foot risk score</b>	<b>Total number of subjects (%)</b>	<b>Dead within 1 year</b>	<b>1 year survival</b>	<b>O.R. for death v low risk</b>
<b>Low</b>	21151(63.7%)	347	98.4%	1.0
<b>Moderate</b>	9274 (27.8%)	343	96.3%	2.3
<b>High</b>	2145 (6.4%)	138	93.6%	4.6
<b>Active</b>	698 (2.1%)	86	87.7%	7.7
<b>Overall</b>	33268	914	97.3%	

# Diabetes Foot Risk Stratification & One Year Mortality



The image features the Royal Coat of Arms of the United Kingdom, rendered in a light gold or yellow color. It depicts a central shield supported by a lion on the left and a unicorn on the right. Above the shield is a crown, and a large eagle with spread wings is positioned at the top. The entire emblem is set against a light yellow background.

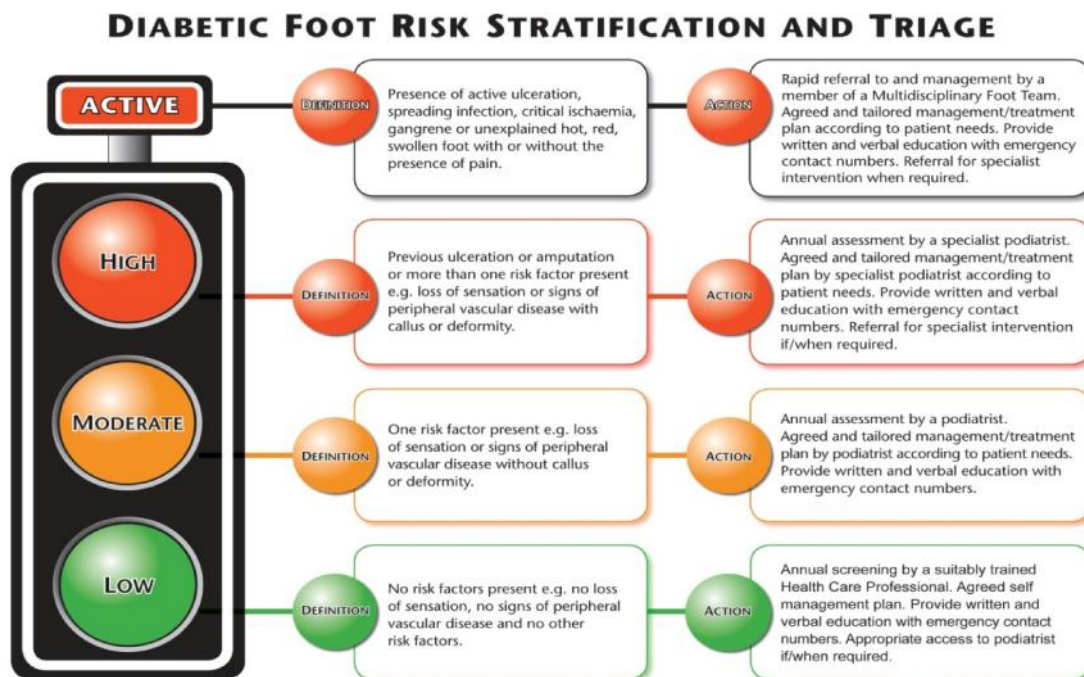
**What can we do to improve  
outcomes?**

DON'T STOP ME NOW



# Improving Outcomes

- Identify those individuals 'at risk'
  - Foot risk stratification
  - Attempt to modify risk status
  - Unclear if intervention prevents ulceration



Produced by the Scottish Diabetes Group - Foot Action Group

These risk categories relate to the use of the SCI-DC foot risk stratification tool



# Ensure appropriately skilled HCP



**Online, interactive, e-learning resource using animations and case scenarios to ensure whichever healthcare professional carrying out foot screening, has the competence and confidence to do so**

## Foot Risk Awareness and Management Education

# **F R A M E**

[www.diabetesframe.org](http://www.diabetesframe.org)



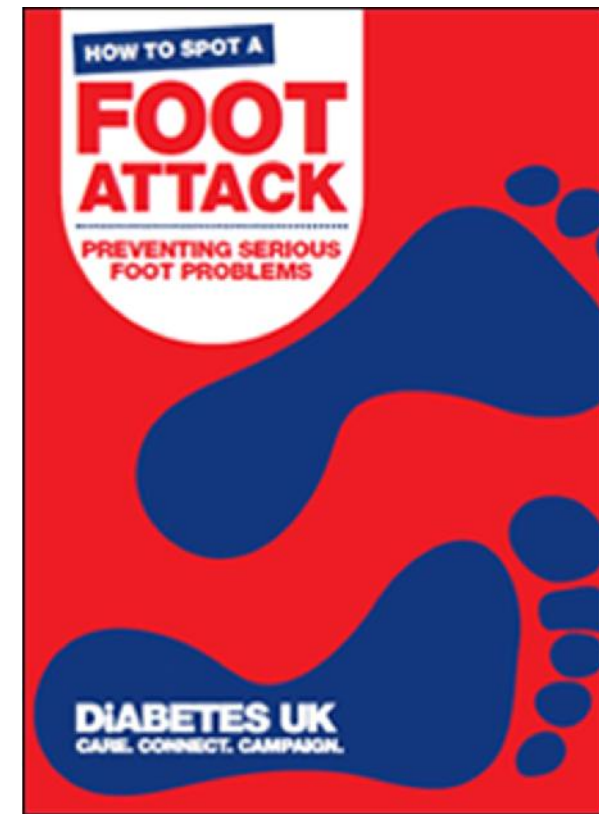
# Person Centred Care



## FOCUS ON FEET



- Pilot aimed at developing a structured education resource for those with active foot disease or 'high risk'
- Concentrates on why foot ulcers develop, what you can do to get them healed and how to reduce the risk of recurrence



# Specialist Assessment & Intervention



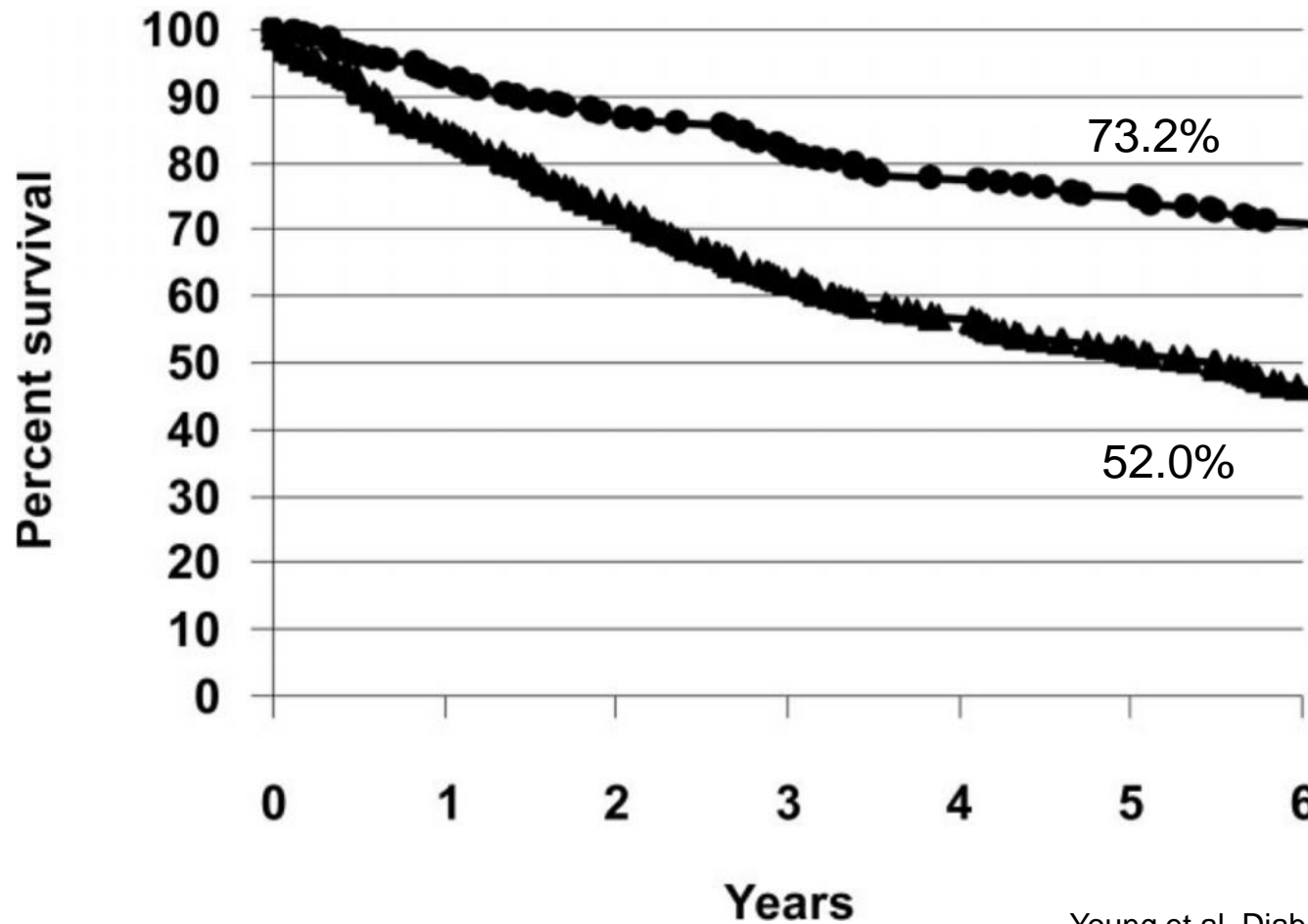
- Early identification of acute problems and immediate referral to specialist foot teams
  - MDFC have been shown to decrease amputation rates
- Immediate assessment focusing on 4 key areas:
  - Infection    - Ischaemia    - Pressure    - Glycaemia
- Early referral for possible surgical intervention
  - Delays in revascularisation and drainage of infection increases the risk of amputation and potentially death
- Develop care pathways similar to stroke
  - Timely imaging and revascularisation



# Aggressive CV risk factor management



## Improved Survival of Diabetic Foot Ulcer Patients 1995–2008



# Reduce Iatrogenic Harm



The Scottish Inpatient Diabetic Foot Audit in November 2013 revealed that:

- 2.4% of in patients with diabetes developed a new foot lesion whilst in hospital
- 57% of in patients had not had their feet checked
- 60% who were discovered to be at risk of developing a foot ulcer did not have any pressure relief in place

*(Scottish Diabetes Foot Action Group 2013)*

Have your patients with diabetes had:  
**CPR for their Feet?**

**C**  
**Check**



**C**heck both feet:

- Is there an ulcer or gangrene?
- Is neuropathy present?
- Is action required?

**P**  
**Protect**



**P**rotect feet if at risk due to:

- Neuropathy
- Previous ulcer or amputation
- Bed bound or fragile skin

**R**  
**Refer**



**R**efer all patients with a foot ulcer, gangrene or other major concern to the podiatry department or diabetes team.

**Ext** .....

# Diabetic foot problems: prevention and management of foot problems in people with diabetes



Care within 24 hours of a person with diabetic foot problems being admitted to hospital, or the detection of diabetic foot problems (if the person is already in hospital)

- Each hospital should have a care pathway for people with diabetic foot problems who need inpatient care. **[2011] [1.1.1]**

## Assessing the risk of developing a diabetic foot problem

- For adults with diabetes, assess their risk of developing a diabetic foot problem at the following times: when diabetes is diagnosed, at least annually thereafter (see recommendation 1.3.11), if problems arise, and on any admission to hospital. **[1.3.3]**

# Diabetic foot problems: prevention and management of foot problems in people with diabetes



## Assessing the risk of developing a diabetic foot problem

- Refer people with an active diabetic foot problem to the foot protection service or multidisciplinary foot care service within 24 hours for appropriate triage according to local protocols. [1.4.1]

1.2.3 The multidisciplinary foot care service should be led by a named healthcare professional, and consist of specialists with skills in the following areas:

- Diabetology.
- Podiatry.
- Diabetes specialist nursing.
- Vascular surgery.
- Microbiology
- Orthopaedic surgery.
- Orthotics and/or biomechanics.
- Interventional radiology.
- Casting.
- Tissue viability.

# Charcot Neuroarthropathy



## Investigation

- 1.7.1 Be aware that if a person with diabetes fractures their foot or ankle, it may progress to Charcot arthropathy.
- 1.7.2 Suspect acute Charcot arthropathy if there is redness, warmth, swelling or deformity (in particular, when the skin is intact), especially in the presence of peripheral neuropathy or renal failure. Think about acute Charcot arthropathy even when deformity is not present or pain is not reported.
- 1.7.3 Refer the person urgently (within 24 hours) to the multidisciplinary foot care service to confirm the diagnosis, and offer non-weight-bearing treatment until definitive treatment can be started.



# Charcot Neuroarthropathy

- 1.7.4 If acute Charcot arthropathy is suspected, X-ray the affected foot. Consider an MRI if the X-ray is normal but clinical suspicion still remains.

## Treatment

- 1.7.5 If the multidisciplinary foot care service suspects acute Charcot arthropathy, offer treatment with a non-removable off-loading device. Only consider treatment with a removable off-loading device if a non-removable device is not advisable because of the clinical or the person's circumstances.
- 1.7.6 Do not offer bisphosphonates to treat acute Charcot arthropathy, unless as part of a clinical trial.



# Charcot Neuroarthropathy

- 1.7.7 Monitor the treatment of acute Charcot arthropathy using clinical assessment. This should include measuring foot–skin temperature difference and taking serial X-rays until the acute Charcot arthropathy resolves. Acute Charcot arthropathy is likely to resolve when there is a sustained temperature difference of less than 2 degrees between both feet and when X-ray changes show no further progression.
- 1.7.8 People who have a foot deformity that may be the result of a previous Charcot arthropathy are at high risk of ulceration and should be cared for by the foot protection service.

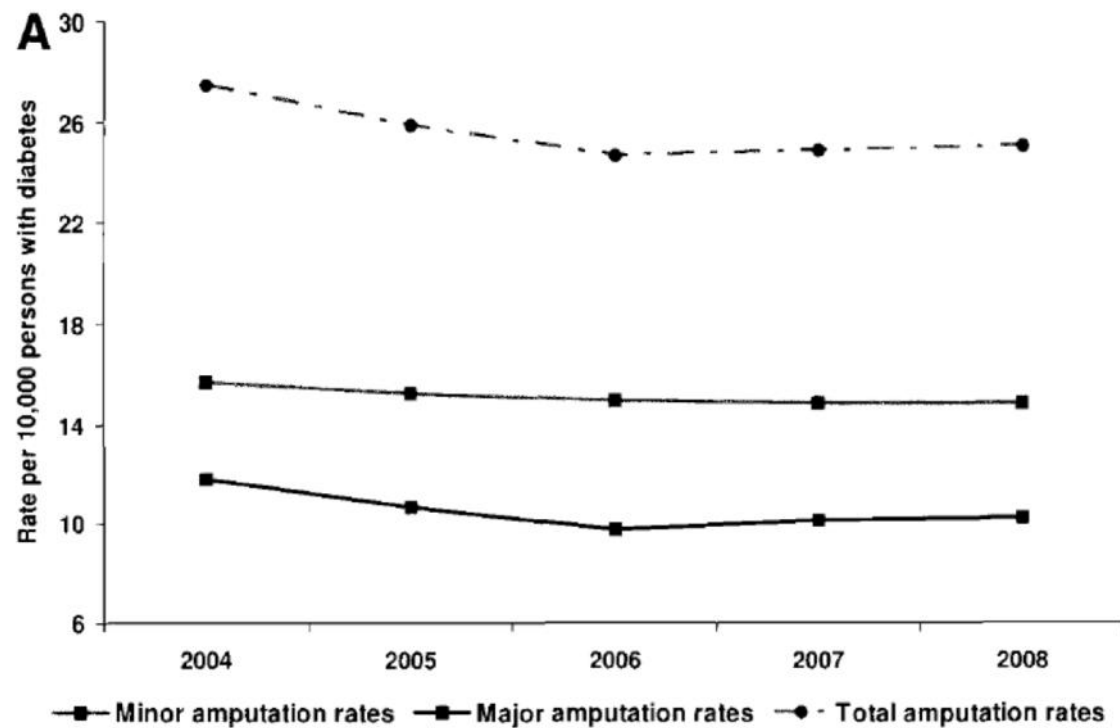


**Are we improving  
outcomes?**

DON'T STOP ME NOW

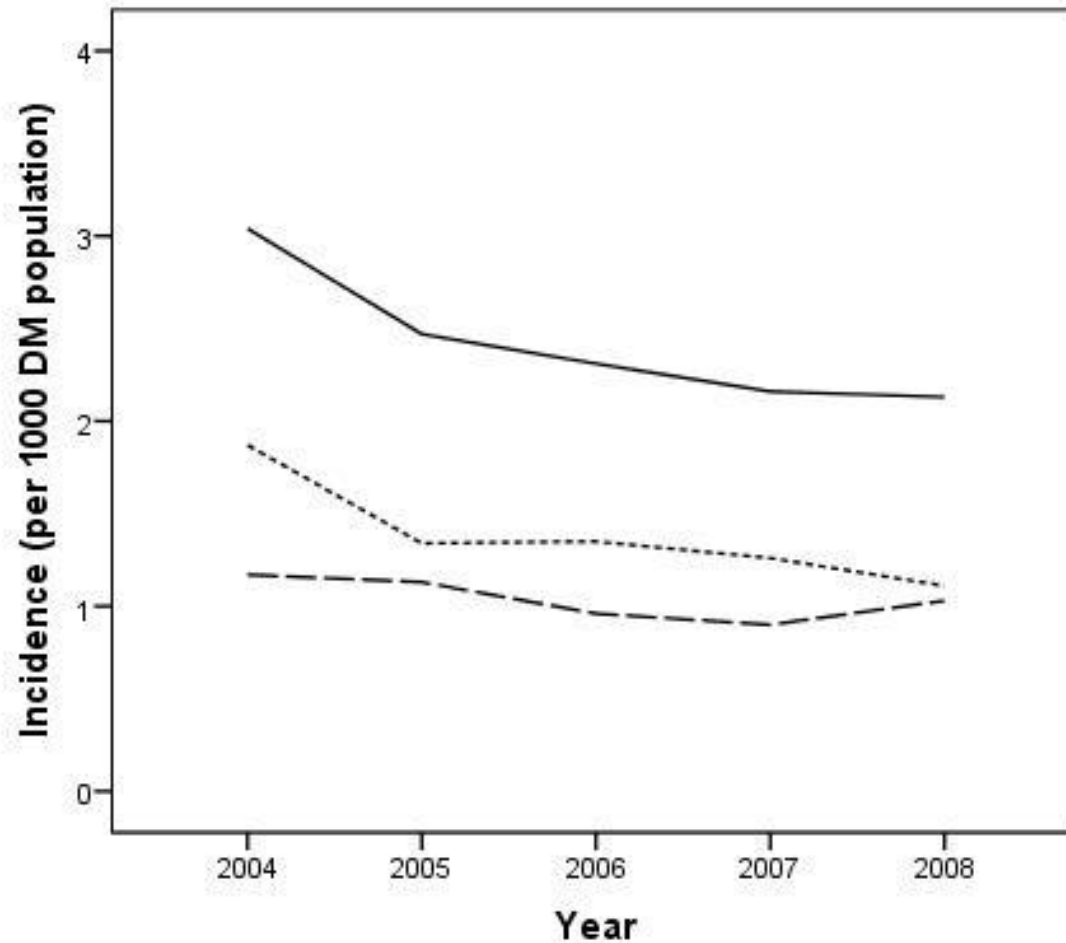


# Lower Extremity Amputation Rates in England



Incidence of major & minor  
LEA rate: unchanged

# Lower Extremity Amputation Rates in Scotland



**Class**  
-- Minor  
..... Major  
— Total

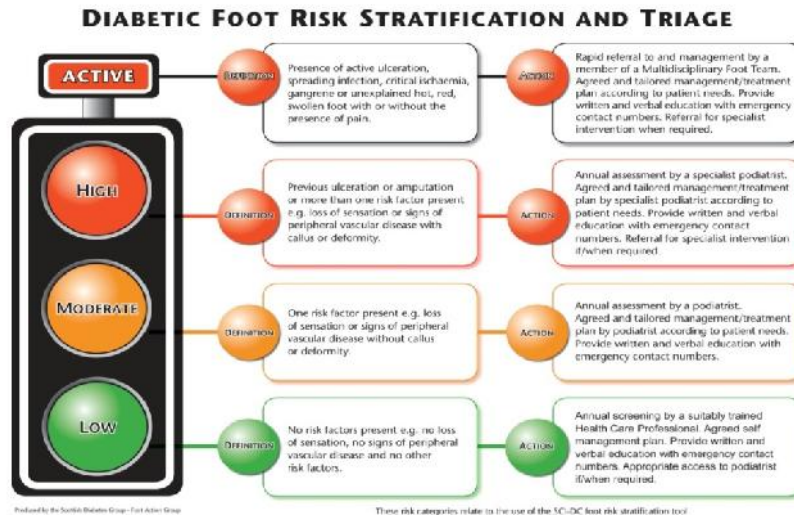
29.8% fall in any LEA  $p < 0.001$

40.7% fall in major LEA  $p < 0.001$

# Reasons for the discrepancy?




- Unified IT system in SCI-diabetes
- Standardised approach to risk stratification and management
- Patient information
- ? Less regional variation
- ? High baseline rate to begin with



# Summary

- Diabetes increases the risk of foot ulceration & amputation
- Risk stratification identifies those 'at risk'
- Immediate referral and review by specialist teams improves outcomes
- Multi-disciplinary foot teams can reduce major amputation rates (& mortality?)
- Acute surgical intervention can save limbs/lives
- Reducing iatrogenic harm for in-patients should be a priority for acute services

DON'T STOP ME NOW



Freddie highlights the importance of  
holistic care.....

Yeah, I'm a rocket ship on my way to Mars  
On a collision course  
I am a satellite I'm out of control  
I am a sex machine ready to reload  
Like an atom bomb about to  
Oh oh oh oh oh explode

DON'T STOP ME NOW

TIP: Podiatrists are well placed to assess  
for erectile dysfunction

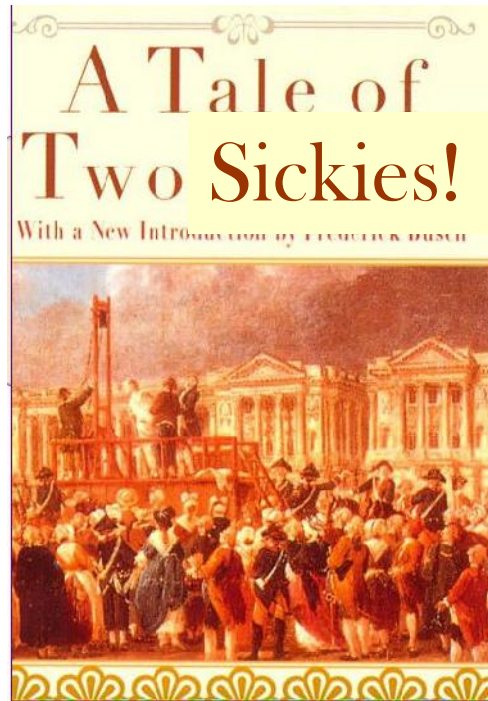
# Acknowledgements



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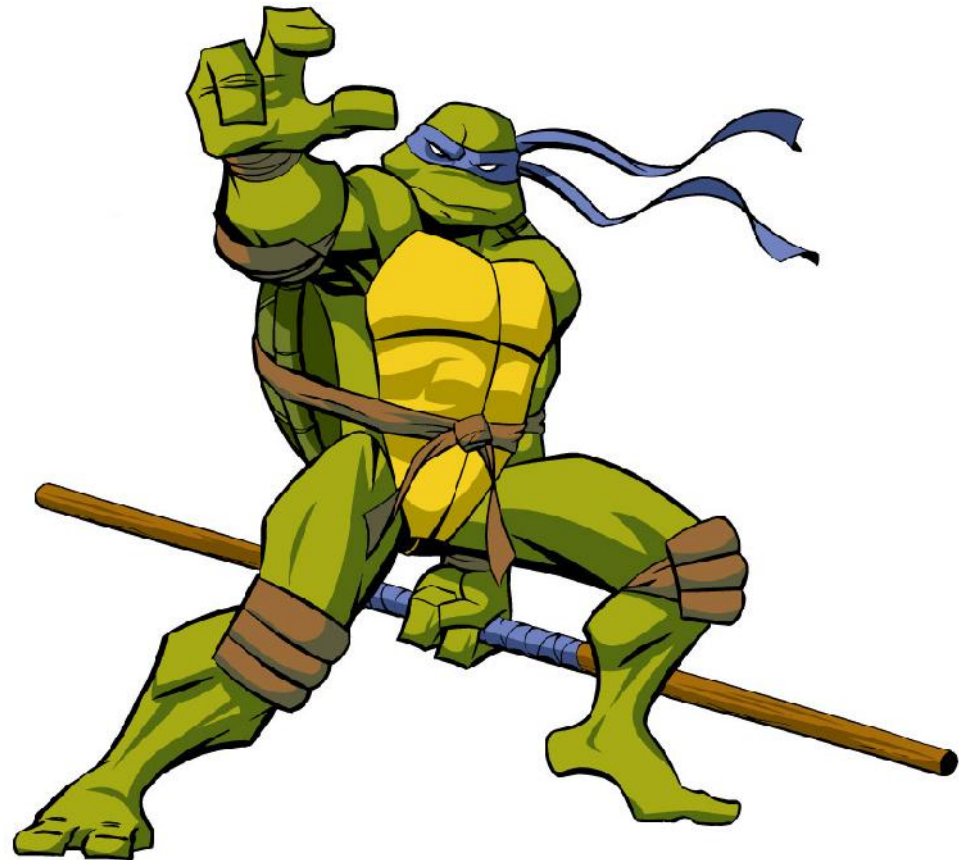
Can elective amputation be  
the best therapeutic option?

# Case Studies





# Case 1



Elective amputation?

# Therapeutic Options

- INFECTION - IV antibiotics
- PRESSURE - Bed rest / Aircast boot
- VASCULAR - PHx angioplasty no further options
- GLYCAEMIA - Very erratic due to illness
- CVS RISK FACTORS – on appropriate treatment
- SURGERY - would mean AKA

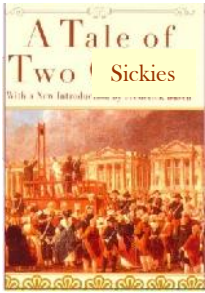
## Risk v Benefit

Anaesthetic  
Immobility  
Quality of Life  
Stump problems



Definitive therapy  
? Improve BG  
Quality of Life

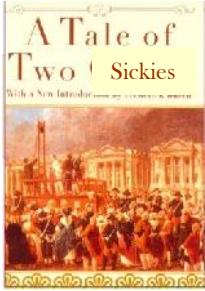
**Elective amputation?**



## Case 2



- 60 years old lady
- Type 1 DM for 30 years
- PHx Left BKA 1994, PVD, Nephropathy, IHD, Retinopathy
- Presents with a hot red swollen foot
- Absent pulses
- Recurrent neuroischaemic ulcer UoT 3D



## Case 2



- **Recurrent severe sepsis**
- **Erratic blood glucose**
- **Poor appetite**

**Elective amputation?**

# Therapeutic Options

- INFECTION - IV antibiotics (venous access very poor)
- PRESSURE - Bed rest & orthosis
- VASCULAR - Severe distal vessel disease
- GLYCAEMIA - Very erratic due to illness
- CVS RISK FACTORS - On appropriate treatment
- SURGERY - Only option BKA (bilateral amputee)

## Risk v Benefit

Anaesthetic  
Immobility  
Quality of Life  
Stump problems



Definitive therapy  
Improve BG  
Quality of Life  
Avoid recurrence

**Elective amputation?**

# Factors determining outcome post amputation

- Age
- Activity level pre-op
- Co-morbidities
- Limb length
- Stump problems
- Planning
- Patient motivation

Elective	Emergency
++	+
+	+++
++	+
++	+++
++++	+
+++	+

# Case 1



- Father would NOT move house
- Refused an amputation
- Readmitted with overwhelming sepsis
- Developed multi-organ failure
- Died aged 49

Would amputation have improved his mortality?

Statistically a success!!

# Case 2



- Right transtibial amputation
- Bilateral amputee
- Independently mobile
- Good diabetes control
- 'Feels great'
- Just back from Canada.....

**A definite statistical failure!**





# Larvae Therapy



## Antimicrobial Activity

- Active removal of bacteria through ingestion and digestion
- Create alkaline wound (pH 8.5) which is an unfavourable environment for bacteria
- Increased irrigation in the wound due to larvae exudate flushes bacteria from site
- Broad spectrum activity: MRSA, Strep. pyogenes and Strep. Pneumoniae, Candida Albicans, Pseudomonas aeruginosa, Enterobacter cloacae, Escherichia coli, varios bacillus strains
- MRSA colonisation was eliminated from 12 of the 13 ulcers (92%)
- Larvae therapy and antibiotics work synergistically

# Improving Performance



## National Diabetes Foot care Audit (NDFA)

This will assess the:

- Structure
- Process
- Outcomes

Only centres in  
England & Wales at  
present