

## How to Reach Me





#### Clinical Scenario History

- \* diagnosed with MS 5 years ago
- has had several acute flairs, including 2 episodes of optic neuritis treated with pulse corticosteroids
- long-standing urinary retention, left arm tremor
- intermittent self-catheterization
- \* has had recurrent urinary tract infections

Hamilton Health Sciences morriand@hhsc.ca

History

**Clinical Scenario** 

- \* no other medical problems
- NKDA
- has been taking cephalexin x 21 days for an infected heel ulcer with mild leg cellulitis
- scheduled to undergo elective lumbar surgery (discectomy)
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#### Clinical Scenario Physical Examination

- \* looks generally well and medically stable
- \* H&N, resp, cardiac and abdominal exams normal
- left CN VI palsy
- \* lower limb spasticity; power 3/5 on left and 4/5 on right
- diminshed vibration and joint position sensation on left
- \* cerebellar left arm tremor
- mild coccygeal skin erythema from decubitus pressure, but no overt skin breakdown
- \* no evidence of active cellulitis (i.e. clinically healed)

Hamilton Health Sciences morriand@hhsc.ca





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## Co-factors of SSIs Patient factors - Microbial virulence

- \* all organisms have the potential to produce SSI
- not all organisms have the same potential to produce SSI
  - the presence of S. aureus in nares is associated with increased risk of SSI
  - eradication of S. aureus in nares reduces the risk of SSI in cardiac surgery
- relatively non-pathogenic organisms (e.g. CNSt, diphtheroids) are especially pathogenic to prosthetic devices

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Co-factors of SSIs
Patient factors - Host resistance
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- poor glycæmic control (both pre-operative and postoperative) has been associated with risk of SSI
- poor vascular supply to the surgical site increases risk of SSI
  - this has been best shown in sternotomy infections following bilateral internal mammary artery grafting of coronary arteries
- there is no good data confirming steroid use as a risk factor for SSI
- malnutrition has been shown to be an independent risk factor for SSI
  - trial evidence has repeatedly failed to show a benefit of TPN in preventing SSI
- Hamilton Health Sciences morriand@hhsc.ca





#### Institutional risk of drug resistance The case of vancomycin

- this has long been an argument against vancomycin as surgical prophylaxis
- vancomycin was also much more expensive than cefazolin prior to coming off-patent in the late 1990s
- \* a recent decision analysis looking at CABG gave the following results
- cefazolin → 7% more SSIs, 1% more deaths, US \$117/procedure
   placebo → 42% more SSIs, 3.6% more deaths, US \$888/procedure
   there has been no sufficient decision model accounting for the risk of developing drug resistance

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Low Risk	Controversial
<ul> <li>varicose vein surgery</li> </ul>	<ul> <li>breast surgery</li> </ul>
<ul> <li>cardiac catheterization</li> </ul>	<ul> <li>herniorraphy</li> </ul>
<ul> <li>most dermatologic and plastic surgery</li> </ul>	<ul> <li>other clean surgery</li> </ul>
<ul> <li>arterial puncture</li> </ul>	
<ul> <li>thoracentesis</li> </ul>	
<ul> <li>paracentesis</li> </ul>	
<ul> <li>lumbar puncture</li> </ul>	

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S. aureus, CNSt ↔ all grafts, implants,	S. aureus, CNSt, GN bacilli	GN bacilli, anaerobes
prostheses	<ul> <li>orthopædic surgery</li> </ul>	<ul> <li>appendectomy</li> </ul>
<ul> <li>neurosurgery</li> </ul>	<ul> <li>cardiac surgery</li> </ul>	<ul> <li>biliary tract</li> </ul>
<ul> <li>breast surgery</li> <li>vascular surgery</li> </ul>	<ul> <li>thoracic surgery</li> </ul>	<ul> <li>gastroduodenal (also has streptococci)</li> </ul>
		<ul> <li>colorectal</li> </ul>
		<ul> <li>♦ OB/GYN (also has Group B strep, enterococci)</li> </ul>
	ciences morriand@hhsc.ca	



#### Don't bother giving post-operative antibiotics

- \* because infection is usually introduced from peri-incisional flora, there is little theoretical benefit from post-operative antibiotics
- \* most published studies included 1-2 post-op doses of antibiotic
- \* a meta-analysis looking at appendectomies showed no additional benefit from multi-dose antibiotics over single-dose pre-op prophylaxis

Cochrane Database Systematic Reviews 2001; 2

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Don't bother arguing with surgeons about postoperative antibiotics (e.g. cardiac surgery)

	Overall results	British Columbia	Alberta	Sask	Manitoba	Ontario	Quebec	NB	Nova Scotia	Nfld
Postoperative	97%	100%	100%	100%	100%	100%	100%	100%	100%	0%
prophylaxis	(32/33)	(4/4)	(2/2)	(2/2)	(2/2)	(9/9)	(11/11)	(1/1)	(1/1)	(0/1)
Regimen (n)*										
Ampicillin 1 g every 6 h Cefazolin	1	-	-	-	-	-	1	-	-	-
Total	33	4	2	2	3	10	9	2	1	-
0.5 g every 6 h	1	-	-	-	-	1	-	-	-	-
1.0 g every 6 h	1	-	-	-	-	-	1	-	-	-
1.0 g every 8 h	23	2	1	21	2	8‡	6	19	1	-
2.0 g every 8 h	6	-	1	-	1	12	2	19	-	-
2.0 g every 12 h	1	1	-	-	-	-	-	-	-	-
3.0 g every 6 h	1	1	-	-	-	-	-	-	-	-
Cloxacillin										
Total	2	-	-	-	-	-	2	-	-	-
1.0 g every 4 h	1	-	-	-	-	-	1	-	-	-
1.0 g every 6 h	1	-	-	-	-	-	1	-	-	-
Vancomycin										
Total	2	-	-	-	-	-	2	-	-	-
0.5 g every 8 h	1	-	-	-	-	-		-	-	-
1.0 g every 12 h	1	-	-	-	-	-	11	-	-	-
Some institutions reported m 1 g for body mass index less I every 8 h for patients weighing	ore than one , han 30, celsu g 100 kg or g	ocaloperative pr olin 2 g for body eater; *Patients	ophydaxia reg r maaa index with valve re	imen; †Celiur 30 or greater placements	olin 2 g admin ; FCetuzolin 1	istered onc g every 8 t	e patient wa I for patients	e off cardio weighing h	ouimonery by ass then 100	osas: ‡Celszolin kg. celszolin 2 g
n duration c	of pro	phyla>	kis is	36h		Can	J In	fect	Dis 2	2002; 1
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#### Generally recommended antimicrobials for preoperative prophylaxis

<ul> <li>neurosurgery</li> <li>breast surgery</li> <li>cefazolin</li> <li>cefazolin</li> <li>cefazolin</li> <li>cefazolin</li> <li>cefazolin</li> <li>cefazolin</li> <li>vancomycin</li> <li>deficient</li> <li>deficient</li> <li>cefazolin</li> </ul>	<ul> <li>all grafts, implants, prostheses</li> <li>neurosurgery</li> <li>breast surgery</li> <li>vascular surgery</li> <li>cefazolin</li> <li>vancomycin</li> </ul>	<ul> <li>orthopædic surgery</li> <li>cardiac surgery</li> <li>thoracic surgery</li> <li>cefazolin</li> <li>cefuroxime</li> <li>vancomycin</li> </ul>	<ul> <li>appendectomy</li> <li>biliary tract</li> <li>gastroduodena</li> <li>has streptococ</li> <li>colorectal</li> <li>OB/GYN (also Group B strep, enterococci)</li> <li>cefotetan</li> <li>cefotetan</li> <li>cefazolin + metronidazo</li> </ul>
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morriand@hhsc.ca

#### **Procedures/Surgeries and Prosthetic Joints**

- \* prosthetic joints are almost always infected at the time of surgery
- rarely, systemic bacteræmia can result in seeding a prosthetic joint
- oral organisms rarely cause prosthetic joint infection
- despite lack of evidence, the ADA and American Academy of Orthopaedic Surgeons recommend giving antibiotics for
  - long procedures

  - surgery in an infected area
    other procedures with a high risk of bacteræmia
- there is generally no need to worry about infected prosthetic joints during dental, GI or GU procedures

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