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Does this document meet the requirements of the Equality Act 2010 in relation to Race, Religion and Belief, Age, Disability, Gender, Sexual Orientation, Gender Identity, Pregnancy & Maternity, Marriage and Civil Partnership, Carers, Human Rights and Social Economic Deprivation discrimination? Initial Assessment		

26/01/2018 – TEMPORARY NOTICE: Due to a shortage of IV clindamycin, please see the notices <u>attached here</u>.

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1 PURPOSE

This guideline has been produced to provide evidence based recommendations to optimise the benefits of using prophylactic antibiotics:

The goals of prophylactic administration of antibiotics to surgical patients are to:

- Reduce the incidence of surgical site infection.
- Use antibiotics in a manner that is supported by evidence of effectiveness.
- Minimise the effect of antibiotics on the patient's normal bacterial flora including driving healthcare associated infections.
- Minimise adverse effects.
- Cause minimal changes to the patient's host defences.

2 SCOPE

This guideline applies to all adult patients in Blackpool Teaching Hospitals NHS Foundation Trust whom are undergoing a surgical procedure.

This guideline must be used by all authorised prescribers involved in the patients care.

Currently this guideline covers surgical prophylaxis to be used in Cardiology, General Surgery, Orthopaedics, Urology and Obstetrics and Gynaecology.

Other areas should continue to use their existing in-house surgical prophylaxis protocols.

3 PROCEDURE

3.1 Benefits and risks of antibiotic prophylaxis

The final decision regarding the benefits and risks of prophylaxis for an individual patient will depend on:

- The patient's risk of surgical site infection (SSI).
- The potential severity of the consequence of SSI.
- The effectiveness of prophylaxis in that operation.
- The consequences of prophylaxis for that patient (e.g. increased risk of colitis).

3.2 Antibiotic choice and dosing

- The antibiotics selected for prophylaxis must cover the expected pathogens for that operative site.
- The choice of antibiotic should take into account local resistance patterns.
- A single standard therapeutic dose of antibiotic is sufficient for prophylaxis under most circumstances.

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3.3 Classification of type of surgery

Surgery may be classified as clean, clean-contaminated, contaminated or dirty.

Class	Definition
Clean	Non-traumatic No inflammation Respiratory, alimentary or genitourinary tracts are NOT entered No break in aseptic technique
Clean- contaminated	Operations in which the respiratory, alimentary or genitourinary tracts are entered but without significant spillage
Contaminated	Operations where acute inflammation (without pus) is encountered, or where there is visible contamination of the wound.
Dirty	Operations in the presence of pus, where there is a previously perforated hollow viscous or compound / open injuries more than four hours old.

Antibiotic prophylaxis is not routinely required for clean non-prosthetic uncomplicated surgery. Antibiotic prophylaxis is required for clean surgery involving the placement of a prosthesis or implant, clean-contaminated surgery and contaminated and dirty surgery.

3.4 Duration of surgery

Prophylactic antibiotic should be limited to an evidence based single dose except in special circumstances (e.g. prolonged surgery, major blood loss or as indicated).

For operations lasting more than 4 hours, re-dosing may be necessary depending on the half-life [T½] antibiotics used. Subsequent doses may be used at the following intervals:

- Cefuroxime 8-hourly
- Metronidazole 8-hourly
- Gentamicin No further doses necessary
- Teicoplanin No further doses necessary

Gentamicin is ALWAYS given as a single dose, and NO subsequent doses should be required, except in exceptional circumstances. Discuss with Microbiology if this situation arises.

In the event of major intra-operative blood loss in adults (>1500mL) additional dosage of antibiotic should be considered after fluid replacement.

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3.5 Timing of administration

The aim of prophylaxis is to have maximum tissue levels at the time of first incision (the only exception is where microbiological specimens are to be taken, in which case prophylaxis should be given immediately after specimens have been obtained).

For this reason, oral and intramuscular prophylaxis is usually administered 1 hour pre-op, whereas intravenous antibiotics are given so that the infusion or dose has just been completed at the time of incision.

3.6 Prophylaxis in patients known to be colonised with Methicillin-Resistant Staphylococcus Aureus (MRSA), or at high risk of MRSA

ALL patients undergoing elective surgery should be screened for MRSA pre-operatively and managed accordingly. In elective patients known to be colonised with MRSA, emergency patients known to be colonised with MRSA and those admitted from Nursing Homes and Care Homes (at higher risk of MRSA carriage) undergoing prosthetic/implant surgery, the antibiotic choice should cover MRSA (i.e. add Teicoplanin, unless already included as part of regimen).

For ease and practicality, Teicoplanin is the antibiotic of choice for prophylaxis against MRSA. Vancomycin remains the glycopeptide of choice for the treatment of MRSA infection (see Antimicrobial Formulary)

Teicoplanin should be used in the following dose regimen:

Teicoplanin 600mg IV as a single dose at induction.

3.7 MRSA Carriage, Cardiac and Orthopaedic Surgery:

Please contact MRSA Decolonisation Failure Clinic if patients with MRSA carriage have failed primary MRSA decolonisation regime before elective surgery.

MRSA Decolonisation Failure Clinic contacts see Appendix 2.

3.8 Surgical prophylaxis guidance

See Appendix 1.

4 ATTACHMENTS	3
Appendix Number	Title
Appendix 1	Surgical prophylaxis guidance

5	ELECTRONIC AND MANUAL RECORDING OF INFORMATION
Ele	ectronic Database for Procedural Documents
He	eld by Policy Co-ordinators/Archive Office

6 LOCATIONS THIS DOCUMENT ISSUED TO		
Copy No Location Date Issued		
1	Intranet	18/06/2015
2	Wards, Departments and Service	18/06/2015

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7 OTHER RELEVANT/ASSOCIATED DOCUMENTS	
Unique Identifier Title and web links from the document library	

8 SUPPORTING REFERENCES/EVIDENCE BASED DOCUMENTS

References In Full

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American urological association 2014

https://www.auanet.org/common/pdf/education/clinical-guidance/Antimicrobial-

Prophylaxis-PocketTable.pdf <accessed 5.2.15>

9 CONSULTATION WITH STAFF AND PATIENTS

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10 DEFINITIONS/GLOSSARY OF TERMS	
MRSA	Methicillin-Resistant Staphylococcus Aureus
SSI	surgical site infection

11 AUTHOR/DIVISIONAL/DIRECTORATE MANAGER APPROVAL			
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Appendix 1: Antibiotic prophylaxis for surgical procedures		
Antibiotic prophylaxis for Cardiovascular	Procedures	
Surgery / Procedure	Regimen	
PERMANENT PACEMAKER AND CARDIOVERTER DEFIBRILLATOR IMPLANTATION	Flucloxacillin 1g IV as a single dose at induction	
Hypersensitivity to penicillins or cephalosporins	Clindamycin 600mg IV as a single dose at induction	
In patients known to be colonised with MRSA or at high risk of MRSA	Teicoplanin 600mg IV as a single dose at induction	
Subsequent doses are NOT usually required		
TEMPORARY PACING WIRE IMPLANTATION	Teicoplanin 600mg IV as a single dose at induction, followed by two further doses of 600mg at 12-hours followed by 600mg 24-hourly until permanent pacing wire is situated or temporary pacing wire is removed	
TRANSCATHETER AORTIC VALVE REPLACEMENT	Single dose Teicoplanin 600mg IV at induction [preferably 20min before incision] Plus Single dose Gentamicin 160mg IV [240mg if over 90Kg] at induction (ensure patient does not have any renal impairment]	
Patients with renal impairment (CrCl < 50mL/min) Lower doses may be required in patients with severe renal impairment CrCl < 10mL/min	Cefuroxime 1.5g plus 2 further doses at 8 and 16 hours post-op	

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Appendix 1: Antibiotic prophylaxis for surgical procedures **Antibiotic prophylaxis for Cardiovascular Procedures Surgery / Procedure** Regimen **CARDIO-THORACIC SURGERY** Gentamicin IV 3mg/kg as a single dose at induction plus Flucloxacillin IV 1g as a single dose at induction, followed by 3 further doses of flucloxacillin 1g at 6, 12 and 18hours post op Hypersensitivity to penicillins / In patients Gentamicin IV 3mg/kg as a single dose at known to be colonised with MRSA induction plus Teicoplanin IV 800mg as a single dose at induction **patients with severe renal impairment CrCl Discussed with microbiologist for an <10mL/min individualised plan microbiologist **Patients for thoracic surgery with existing Discussed with an active infection individualised plan

Consultant cardiac surgeons may choose to use a 2nd dose of gentamicin on day 2 post op. However this should be a clinical decision and following obtaining a gentamicin trough level <1mg/l (sample collected between 18-24hours after 1st dose) before administering the 2nd dose.

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Prophylaxis against infective endocarditis

In March 2008, NICE issued guidance on antimicrobial prophylaxis against infective endocarditis in adults and children undergoing interventional procedures.

Antibiotic prophylaxis has not been proven to be effective and there is no clear association between episodes of infective endocarditis and interventional procedures. Any benefits from prophylaxis need to be weighed against the risks of adverse effects for the patient and of antibiotic resistance developing. As a result, **NICE have recommended that antibiotic prophylaxis is no longer offered routinely for defined interventional procedures.**

When to offer prophylaxis

Do **NOT** offer antibiotic prophylaxis against infective endocarditis:

- To people undergoing dental procedures
- To people undergoing non-dental procedures at the following sites:
- Lower and upper gastrointestinal tract
- Genitourinary tract; this includes urological, gynaecological and obstetric procedures and childbirth
- Upper and lower respiratory tract; this includes ear, nose and throat procedures and bronchoscopy.

Do **NOT** offer chlorhexidine mouthwash as prophylaxis against infective endocarditis to people at risk undergoing dental procedures.

Whilst these procedures can cause bacteraemia, there is no clear association with the development of infective endocarditis. Prophylaxis may expose patients to the adverse effects of antimicrobials when the evidence of benefit has not been proven.

Managing infection

Any infection in patients at risk of endocarditis_should be **investigated promptly** and **treated appropriately** to reduce the risk of endocarditis.

If patients at risk of endocarditis are undergoing a gastro-intestinal or genitourinary tract procedure at a site where infection is suspected, they should receive appropriate antibacterial therapy that includes cover against organisms that cause endocarditis.

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Advice

Offer people at risk of infective endocarditis clear and consistent information about prevention including:

- The benefits and risks of antibiotic prophylaxis, and an explanation of why antibiotic prophylaxis is no longer recommended
- The importance of maintaining good oral health
- Symptoms that may indicate infective endocarditis and when to seek expert advice
- The risks of undergoing invasive procedures, including non-medical procedures such as body piercing or tattooing

The following cardiac conditions are at risk of developing infective endocarditis:

- Acquired valvular heart disease with stenosis or regurgitation
- Valve replacement
- Patients with a prosthetic cardiac valve
- Structural congenital heart disease including:
 - Unrepaired cyanotic CHD, including palliative shunts and conduits
 - Completely repaired congenital heart defect with prosthetic material or device, whether placed by surgery or by catheter intervention, during the first 6 months after the procedure (i.e. pre-endothelialisation)
 - Repaired CHD with residual defects at the site or adjacent to the site of a prosthetic patch or prosthetic device (which inhibit endothelialisation)
- Hypertrophic cardiomyopathy
- Patients with previous infective endocarditis
- Cardiac transplantation recipients who develop cardiac valvulopathy

If prophylaxis is considered appropriate for individual patients, please choose from one of the regimens below:

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Appendix 1: Antibiotic prophylaxis for sur	gical procedures
Surgery / Procedure	Regimen
PROPHYLAXIS AGAINST INFECTIVE ENDOCARDITIS	Amoxicillin 1g IV as a single dose at induction PLUS Gentamicin 3mg/kg IV as a single dose at induction
Serious allergy to penicillins/allergy to cephalosporins	Teicoplanin 600mg IV as a single dose at induction PLUS Gentamicin 3mg/kg IV as a single dose at induction
Patients with renal impairment (CrCl < 50mL/min)	Cefuroxime 1.5g IV as a single dose at induction
In patients known to be colonised with MRSA or at high risk of MRSA	Teicoplanin 600mg IV as a single dose at induction PLUS Gentamicin 3mg/kg IV as a single dose at induction
In patients known to be colonised with MRSA or at high risk of MRSA with renal impairment (CrCl < 50mL/min)	Teicoplanin 600mg IV as a single dose at induction PLUS Cefuroxime 1.5g IV as a single dose at induction

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Antibiotic prophylaxis for Urological Procedures

MUST check previous sensitivity if available and if resistance to suggested regimen discuss with microbiologist

Surgery / Procedure	Regimen
ROUTINE SHORT-TERM CATHETERISATION Acute retention and peri-operatively Catheter insertion: If febrile (>38°C) or urine dipstick positive for nitrites, send urine for culture and treat	Antibiotic prophylaxis is NOT required
appropriately i.e. symptomatic Catheter removal:	Antibiotic prophylaxis is NOT required
LONG-TERM CATHETERISATION (including intermittent self-catheterisation)	No antibiotic prophylaxis required for routine change, even if colonised with bacteria. Prophylaxis is ONLY indicated if there has been a history of sepsis with previous catheter changes. The antibiotic chosen in these cases should be based on sensitivity of urine isolate.
TRANSURETHRAL RESECTION OF PROSTATE (TURP)	If systemic signs of infection, treat appropriately Gentamicin 3mg/kg IV as a single dose at induction
Patients with renal impairment (CrCl < 50mL/min) TRANSURETHRAL RESECTION OF	Cefuroxime 1.5g IV as a single dose at induction Antibiotic prophylaxis is NOT required
BLADDER TUMOUR (TURBT)	If prophylaxis indicated (*high risk – see patient related factors on next page and large tumours), use Gentamicin 3mg/kg IV as a single dose at induction
Patients with renal impairment (CrCl < 50mL/min) TRANSRECTAL PROSTATE BIOPSY	Cefuroxime 1.5g IV as a single dose at induction Ciprofloxacin 500mg PO (1 hour before) induction followed by one further dose of 500mg PO 12 hours later or Gentamicin 3mg/kg IV as a single dose at induction if no renal impairment

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Surgery / Procedure	Regimen
URO-DYNAMICS SHOCK WAVE LITHOTRIPSY	Ciprofloxacin 500mg PO (1 hour before) induction followed by one further dose of 500mg PO 12 hours later or Gentamicin 3mg/kg IV as a single dose at induction if no renal impairment
Prophylaxis is NOT required unless: Known infected stone, bacteriuria, indwelling catheter, stent, nephrostomy tube Patients with renal impairment (CrCl < 50mL/min)	Gentamicin 3mg/kg IV as a single dose at induction Cefuroxime 1.5g IV as a single dose at induction
CYSTOSCOPY (include hydrodistension, urethral dilation)	Antibiotic prophylaxis is NOT required unless high risk of UTI: patients with bacteriuria, indwelling catheters, history of a urogenital infection (recurrent or recent infection in the last month) or other patient related risk factors* Gentamicin 3mg/kg IV as a single dose at induction or according to MSII.
Patients with renal impairment (CrCl < 50mL/min) In patients known to be colonised with MRSA or at high risk of MRSA	Cefuroxime 1.5g IV as a single dose at induction or according to MSU Teicoplanin 600mg IV as a single dose at induction PLUS gentamicin 3mg/kg IV as a single dose at induction (if MSU resistant to gentamicin — replace gentamicin with alternative sensitive antimicrobial)

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Appendix 1: Antibiotic prophylaxis for surgical procedures	
Surgery / Procedure	Regimen
URETEROSCOPY	Antibiotic prophylaxis is NOT required unless high risk of UTI: patients with bacteriuria, indwelling catheters, history of a urogenital infection (recurrent or recent infection in the last month) or other patient related risk factors*
Mild allergy to penicillins / patients with renal impairment (CrCl < 50mL/min)	Amoxicillin 1g IV as a single dose at induction PLUS gentamicin 3mg/kg IV as a single dose at induction or according to MSU
Serious allergy to penicillins/allergy to cephalosporins/ In patients known to be	Cefuroxime 1.5g IV as a single dose at induction or according to MSU
colonised with MRSA or at high risk of MRSA	Teicoplanin 600mg IV as a single dose at induction PLUS gentamicin 3mg/kg IV as a single dose at induction (if MSU resistant to gentamicin — replace gentamicin with alternative sensitive antimicrobial)
In patients known to be colonised with MRSA or at high risk of MRSA with renal impairment (CrCl < 50mL/min)	Teicoplanin 600mg IV as a single dose at induction PLUS Cefuroxime 1.5g IV as a single dose at induction

*Other patient related factors for infectious complications:

General risk factors	Special risk factors associated with an increased bacterial load
Older age	Long preoperative hospital stay or recent hospitalisation
Deficient nutritional status	History of recurrent urogenital infections
Impaired immune response	Surgery involving bowel segment
Diabetes mellitis	Colonisation with microorganisms
Smoking	Long term drainage
Extreme weight	Urinary obstruction or stone
Co-existing infection at a remote site	

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Appendix 1: Antibiotic prophylaxis for surgical procedures		
Surgery / Procedure	Regimen	
BLADDER LITHOPAXY	Antibiotic prophylaxis is NOT required unless high risk of UTI: patients with bacteriuria, indwelling catheters, history of a urogenital infection (recurrent or recent infection in the last month) or other patient related risk factors* Gentamicin 3mg/kg IV as a single dose at	
	induction or according to MSU	
Patients with renal impairment (CrCl < 50mL/min)	Cefuroxime 1.5g IV as a single dose at induction or according to MSU	
In patients known to be colonised with MRSA or at high risk of MRSA	Teicoplanin 600mg IV as a single dose at induction PLUS gentamicin 3mg/kg IV as a single dose at induction (if MSU resistant to gentamicin — replace gentamicin with alternative sensitive antimicrobial)	
In patients known to be colonised with MRSA or at high risk of MRSA with renal impairment (CrCl < 50mL/min)	Teicoplanin 600mg IV as a single dose at induction PLUS Cefuroxime 1.5g IV as a single dose at induction	
Renal stent change/removal	Gentamicin 3mg/kg IV as a single dose at induction or according to MSU	
Patients with renal impairment (CrCl < 50mL/min)	Cefuroxime 1.5g IV as a single dose at induction or according to MSU	
*Other patient related factors for infectious complications:		
General risk factors	Special risk factors associated with an increased bacterial load	
Older age	Long preoperative hospital stay or recent hospitalisation	
Deficient nutritional status	History of recurrent urogenital infections	
Impaired immune response	Surgery involving bowel segment	
Diabetes mellitis	Colonisation with microorganisms	
Smoking	Long term drainage	
Extreme weight	Urinary obstruction or stone	
Co-existing infection at a remote site		

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Appendix 1: Antibiotic prophylaxis for surgical procedures	
Surgery / Procedure	Regimen
CORRECTION OF HYDROCELE	Antibiotic prophylaxis is NOT required unless high risk of infections e.g. immunocompromised - discuss with microbiologist
CIRCUMCISION	Antibiotic prophylaxis is NOT required for routine elective procedures
FRENULOPLASTY	Flucloxacillin 1g IV as a single dose at induction PLUS gentamicin 3mg/kg IV as a single dose at induction
Mild allergy to penicillins/ Patients with renal impairment (CrCl < 50mL/min)	Cefuroxime 1.5g IV as a single dose at induction
SERIOUS ALLERGY TO PENICILLINS OR MRSA COLONISED OR SUSPECTED	Teicoplanin 600mg IV as a single dose at induction PLUS gentamicin 3mg/kg IV as a single dose at induction
In patients known to be colonised with MRSA or at high risk of MRSA with renal impairment (CrCl < 50mL/min)	Teicoplanin 600mg IV as a single dose at induction PLUS Cefuroxime 1.5g IV as a single dose at induction
Excision of epidymal cyst	Antibiotic prophylaxis is NOT required
Penile biopsy	Antibiotic prophylaxis is NOT required
Optical urethrotomy	Antibiotic prophylaxis is NOT required unless high risk of UTI: patients with bacteriuria, indwelling catheters, history of a urogenital infection (recurrent or recent infection in the last month) or other patient related risk factors*
	Amoxicillin 1g IV as a single dose at induction PLUS gentamicin 3mg/kg IV as a single dose at induction
Mild allergy to penicillins / patients with renal impairment (CrCl < 50mL/min)	Cefuroxime 1.5g IV as a single dose at induction
SERIOUS ALLERGY TO PENICILLINS/ In patients known to be colonised with MRSA or at high risk of MRSA	Teicoplanin 600mg IV as a single dose at induction PLUS gentamicin 3mg/kg IV as a single dose at induction
In patients known to be colonised with MRSA or at high risk of MRSA with renal impairment (CrCl < 50mL/min)	Teicoplanin 600mg IV as a single dose at induction PLUS Cefuroxime 1.5g IV as a single dose at induction

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Appendix 1: Antibiotic prophylaxis for surgical procedures **Antibiotic prophylaxis for Upper Gastro-Intestinal Surgery** Regimen **Surgery / Procedure OESOPHAGEAL SURGERY** Gentamicin 3mg/kg IV as a single dose at **GASTRO DUODENAL SURGERY** induction **GASTRIC BYPASS SURGERY BARIATRIC SURGERY** Patients with renal impairment **Cefuroxime** 1.5g IV as a single dose at (CrCl < 50mL/min)induction In patients known to be colonised with Teicoplanin 600mg IV as a single dose at MRSA or at high risk of MRSA induction PLUS Gentamicin 3mg/kg IV as a single dose at induction In patients known to be colonised with Teicoplanin 600mg IV as a single dose at MRSA or at high risk of MRSA with renal induction PLUS Cefuroxime 1.5g IV as a impairment (CrCl < 50mL/min) single dose at induction

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Appendix 1: Antibiotic prophylaxis for surgical procedures		
Antibiotic prophylaxis for Hepato-biliary Surgery		
Surgery / Procedure	Regimen	
BILIARY SURGERY – OPEN PANCREATIC SURGERY LIVER SURGERY GALL BLADDER SURGERY (OPEN) GALL BLADDER SURGERY (LAPAROSCOPIC)	Gentamicin 3mg/kg IV as a single dose at induction PLUS Metronidazole 500mg IV as a single dose at induction	
Patients with renal impairment (CrCl < 50mL/min)	Cefuroxime 1.5g IV as a single dose at induction PLUS Metronidazole 500mg IV as a single dose at induction	
In patients known to be colonised with MRSA or at high risk of MRSA	Teicoplanin 600mg IV as a single dose at induction PLUS Metronidazole 500mg IV as a single dose at induction PLUS Gentamicin 3mg/kg IV as a single dose at induction	
In patients known to be colonised with MRSA or at high risk of MRSA with renal impairment (CrCl < 50mL/min)	Teicoplanin 600mg IV as a single dose at induction PLUS Metronidazole 500mg IV as a single dose at induction PLUS Cefuroxime 1.5g IV as a single dose at induction	
LAPAROSCOPIC CHOLECYSTECTOMY – UNCOMPLICATED	Antibiotic prophylaxis is NOT required	
ERCP	Gentamicin IV 3 mg/Kg [preferred] or Ciprofloxacin 500mg PO as a single dose at induction	

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Appendix 1: Antibiotic prophylaxis for surgical procedures **Antibiotic prophylaxis for Lower Gastro-Intestinal Surgery** Regimen **Surgery / Procedure APPENDECTOMY** Gentamicin 3mg/kg IV as a single dose at induction + Metronidazole 500mg IV as a COLORECTAL SURGERY SMALL BOWEL SURGERY single dose at induction Patients with renal impairment Cefuroxime 1.5g IV as a single dose at induction +/- Metronidazole 500mg IV as a (CrCl < 50mL/min)Lower doses may be required in patients single dose at induction with severe renal impairment CrCl < 10mL/min In patients known to be colonised with Teicoplanin 600mg IV as a single dose at induction PLUS Metronidazole 500mg IV MRSA or at high risk of MRSA as a single dose at induction PLUS Gentamicin 3mg/kg IV as a single dose at induction In patients known to be colonised with Teicoplanin 600mg IV as a single dose at MRSA or at high risk of MRSA with renal induction PLUS Metronidazole 500mg IV impairment (CrCl < 50mL/min) as a single dose at induction PLUS Cefuroxime 1.5g IV as a single dose at induction

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Appendix 1: Antibiotic prophylaxis for surgical procedures		
Antibiotic prophylaxis for Abdominal Surgery		
Surgery / Procedure	Regimen	
HERNIA REPAIR – WITHOUT MESH	Antibiotic prophylaxis is NOT usually required	
HERNIA REPAIR – WITH MESH	Gentamicin 3mg/kg IV as a single dose at induction	
Patients with renal impairment (CrCl < 50mL/min) Lower doses may be required in patients with severe renal impairment CrCl < 10mL/min	Cefuroxime 1.5g IV as a single dose at induction	
In patients known to be colonised with MRSA or at high risk of MRSA	Teicoplanin 600mg IV as a single dose at induction PLUS Gentamicin 3mg/kg IV as a single dose at induction	
In patients known to be colonised with MRSA or at high risk of MRSA with renal impairment (CrCl < 50mL/min)	Teicoplanin 600mg IV as a single dose at induction PLUS Cefuroxime 1.5g IV as a single dose at induction	

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Appendix 1: Antibiotic prophylaxis for surgical procedures **Antibiotic Prophylaxis in Vascular Surgery Surgery / Procedure** Regimen Flucioxacillin 1g IV as a single dose at VASCULAR PROCEDURES: includes induction, followed by three further doses of lower limb amputations and all grafts 1g at 6, 12, and 18 hours whether vein or prosthetic Metronidazole 500mg IV as a single dose at induction, followed by two further doses of 500mg at 8 and 16 hours PLUS Gentamicin 3mg/kg IV as a single dose at induction Mild allergy to penicillins / patients with Cefuroxime 1.5g IV as a single dose at renal impairment (CrCl < 50mL/min) induction, followed by two further doses of Lower doses may be required in patients at 8 and 16 **PLUS** 750mg hours with severe renal impairment CrCl < Metronidazole 500mg IV as a single dose 10mL/min at induction, followed by two further doses of 500mg at 8 and 16 hours Serious allergy to penicillins/allergy Teicoplanin 600mg IV as a single dose at cephalosporins/ colonised induction, followed by one further dose of MRSA or suspected 600mg at 12 hours PLUS Metronidazole 500mg IV as a single dose at induction, followed by two further doses of 500mg at 8 and 16 hours PLUS Gentamicin 3mg/kg IV as a single dose at induction In patients known to be colonised with Teicoplanin 600mg IV as a single dose at induction, followed by one further dose of MRSA or at high risk of MRSA with renal 600mg at 12 hours PLUS Metronidazole impairment (CrCl < 50mL/min) 500mg IV as a single dose at induction, two further doses of 500mg at 8 and 16 hours PLUS Cefuroxime 1.5q IV as a single dose at induction, followed by two further doses of 750mg at 8 and 16 hours

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Appendix 1: Antibiotic prophylaxis for surgical procedures		
Antibiotic Prophylaxis for Orthopaedic Procedures		
Surgery / Procedure	Regimen	
ARTHROSCOPY	Antibiotic prophylaxis is NOT required	
ORTHOPAEDIC SURGERY WITHOUT PROSTHESIS (except spinal surgery, open traumatic wound)	Antibiotic prophylaxis is NOT required	
MINOR METALWORK INSERTION (e.g. K-wires, screws, small orthopaedic plates)	Flucloxacillin 1g IV +/- Gentamicin 3mg/kg IV at induction (single dose)	
MINOR METALWORK REMOVAL	If removing uninfected metal work – Antibiotic prophylaxis is NOT required If removing infected metal work – empiric antibiotic prophylaxis or guided by positive bacteriology would be needed	
Mild allergy to penicillins / patients with renal impairment (CrCl < 50mL/min) Lower doses may be required in patients with severe renal impairment CrCl < 10mL/min	Cefuroxime 1.5g IV as a single dose at induction, followed by two further doses of 750mg at 8 and 16 hours	
Serious allergy to penicillins/allergy to cephalosporins/ MRSA colonised or suspected	Teicoplanin 600mg IV as a single dose at induction PLUS Gentamicin 3mg/kg IV as a single dose at induction	
In patients known to be colonised with MRSA or at high risk of MRSA with renal impairment (CrCl < 50mL/min)	Teicoplanin 600mg IV as a single dose at induction PLUS Cefuroxime 1.5g IV as a single dose at induction	

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Appendix 1: Antibiotic prophylaxis for surgical procedures Flucloxacillin 1g IV as a single dose at SURGERY ORTHOPAEDIC PROSTHESIS (joint replacement surgery, induction, followed by three further doses of 1g at 6, 12, and 18 hours hip fracture repair, spinal surgery, insertion of prosthetic device, internal fixation of Gentamicin 3mg/kg IV as a single dose at closed fracture, clean open fracture) induction Mild allergy to penicillins / patients with Cefuroxime 1.5g IV as a single dose at renal impairment (CrCl < 50mL/min) induction, followed by two further doses of Lower doses may be required in patients 750mg at 8 and 16 hours with severe renal impairment CrCl < 10mL/min Teicoplanin 600mg IV as a single dose at Serious allergy to penicillins/allergy induction, followed by one further dose of to 600mg at 12 hours PLUS Gentamicin cephalosporins/ MRSA colonised or 3mg/kg IV as a single dose at induction suspected Teicoplanin 600mg IV as a single dose at induction PLUS Cefuroxime 1.5g IV as a In patients known to be colonised with single dose at induction MRSA or at high risk of MRSA with renal impairment (CrCl < 50mL/min) **DIRTY FRACTURE (CONTAMINATED)** Add Metronidazole 500mg IV as a single dose at induction, followed by two further doses of 500mg at 8 and 16 hours NB - consider tetanus immunity for contaminated wound

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Antibiotic Prophylaxis for Obstetric Procedures

Surgery / Procedure	Regimen
CAESAREAN SECTION All patients undergoing caesarean section must receive prophylactic antibiotics. Antibiotics should be administered wherever possible prior to knife incision, or as soon as possible during the procedure (e.g. in the case of Grade 1 Caesarean section).	Cefuroxime 1.5g IV as a single dose at induction
History of immediate rash, angioedema or anaphylaxis to penicillin	Clindamycin IV 600mg as a single dose at induction PLUS (if emergency caesarean section) Gentamicin 3mg/kg IV as a single dose calculated according to ideal (pre-pregnancy) body weight after the cord is clamped
If renal impairment and concern with the use of gentamicin	Discuss with microbiologist Single dose prophylaxis is usually sufficient. Some patients with prolonged ruptured membranes, a long labour, raised CRP or raised white cell count may require further doses of antibiotic therapy.
MANUAL REMOVAL OF PLACENTA	Gentamicin 3mg/kg IV as a single dose calculated according to ideal (prepregnancy) body weight and metronidazole 500mg IV as single dose at induction
Patients with renal impairment (CrCl < 50mL/min) Lower doses may be required in patients with severe renal impairment CrCl < 10mL/min	Cefuroxime 1.5g IV as a single dose at induction PLUS Metronidazole 500mg IV as a single dose at induction

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Appendix 1: Antibiotic prophylaxis for surgical procedures		
THIRD OR FOURTH DEGREE TEARS	Flucloxacillin IV 1g as a single dose plus gentamicin 3mg/kg* IV calculated according to ideal (pre-pregnancy) body weight as a single dose plus metronidazole 500mg IV as a single dose at induction	
Allergy to penicillins	Clindamycin 600mg IV as a single dose plus gentamicin 3mg/kg* IV calculated according to ideal (pre-pregnancy) body weight as a single dose at induction	
If renal impairment and concern with the use of gentamicin	Discuss with microbiologist	

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Antibiotic Prophylaxis for Gynaecological Procedures

Surgery / Procedure	Regimen
CONSERVATIVE MANAGEMENT OF RUPTURED MEMBRANES	Use O&G Directorate Policy
KNOWN GROUP B HAEMOLYTIC STREPTOCOCCUS +VE PATIENTS	Use O&G Directorate Policy for antibiotic prophylaxis during labour
Prophylaxis for any gynaecological procedure involving opening of the peritoneum i.e. HYSTERECTOMY OOPHORECTOMY Mild allergy to penicillins Lower doses may be required in patients with severe renal impairment CrCl < 10mL/min Antibiotic prophylaxis to be part of WHO check list during time-out.	Gentamicin 3mg/kg IV as a single dose calculated according to ideal (prepregnancy) body weight and metronidazole 500mg IV as single dose at induction or if gentamicin is contraindicated: Co-amoxiclav 1.2g IV as a single dose at induction Cefuroxime 1.5g IV as a single dose at induction PLUS Metronidazole 500mg IV as a single dose at induction
Endoscopic procedures e.g. LAPAROSCOPY CYSTOSCOPY	Antibiotic prophylaxis is not usually required.
COLPOSCOPY	Consider prophylaxis for some cardiac patients, see "Prophylaxis against infective endocarditis".
Mild allergy to penicillins / patients with renal impairment (CrCl < 50mL/min) Lower doses may be required in patients with severe renal impairment CrCl < 10mL/min	Gentamicin 3mg/kg IV as a single dose calculated according to ideal (prepregnancy) body weight and metronidazole 500mg IV as single dose at induction or if gentamicin is contraindicated: Co-amoxiclav 1.2g IV as a single dose at induction Cefuroxime 1.5g IV as a single dose at induction PLUS Metronidazole 500mg IV as a single dose at induction

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Appendix 1: Antibiotic prophylaxis for surgical procedures				
VAGINAL REPAIR PROCEDURES		Antibiotic prophylaxis is not usually required provided the abdominal peritoneum is not opened.		
TRANSURETHRAL	TAPE	OR	Discuss with Consultant Gynaecologist	
TRANSOBTURATOR		TAPE	-	
PROCEDURES				

Antibiotic Prophylaxis for Gynaecological Procedures Continued

Surgery / Procedure	Regimen
SUCTION TERMINATION OF PREGNANCY	Appropriate antibiotic treatment for patients who have screened Chlamydia positive.
	If genital Chlamydial infection cannot be ruled out, post-operative Azithromycin 1g PO as a single oral dose should be given.
SURGICAL EVACUATION OF RETAINED PRODUCTS	As per Suction Termination
PERFORATION OF UTERUS DURING HYSTEROSCOPY, D&C OR ABLATION TREATMENT	Co-amoxiclav 1.2g IV as a single dose at induction
Mild allergy to penicillins / patients with renal impairment (CrCl < 50mL/min) Lower doses may be required in patients with severe renal impairment CrCl < 10mL/min	Cefuroxime 1.5g IV as a single dose at induction PLUS Metronidazole 500mg IV as a single dose at induction
BOWEL PERFORATION DURING OPEN OR ENDOSCOPIC PROCEDURE	Follow hospital prophylaxis for clean-contaminated surgery.
	Provided prophylaxis has not already been given:
	Gentamicin 3mg/kg IV as a single dose PLUS Metronidazole 500mg IV as a single dose at the time of recognising perforation.
	If prior prophylaxis, consider the addition of Gentamicin dependent on the degree of perforation and after discussion with a Consultant Microbiologist.

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Antibiotic Prophylaxis for Gynaecological Procedures Continued

Surgery / Procedure	Regimen
PROCEDURES FOR SUSPECTED TUBO- OVARIAN ABSCESS	Cefuroxime 1.5g IV as a single dose at induction PLUS Metronidazole 500mg IV as a single dose at induction PLUS Gentamicin 3mg/kg IV as a single dose at induction. Post-operative therapy to continue with advice from the Consultant Microbiologist as required.
REMOVAL OF TRANSLOCATED IUCD	Co-amoxiclav 1.2g IV as a single dose at induction
Mild allergy to penicillins / patients with renal impairment (CrCl < 50mL/min) Lower doses may be required in patients with severe renal impairment CrCl < 10mL/min	Cefuroxime 1.5g IV as a single dose at induction PLUS Metronidazole 500mg IV as a single dose at induction

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Appendix 2: MRSA Decolonisation Failure Clinic Contacts

- Jan Tipping [Orthopaedics]
 Bernadette McAlea [Cardio-thoracic]
- Dr Guleri [Consultant Microbiologist].

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Appendix 3: Equality Impact Assessment Form

 Department
 Organisation Wide
 Service or Policy
 Guideline
 Date Completed:
 July 2015

GROUPS TO BE CONSIDERED

Deprived communities, homeless, substance misusers, people who have a disability, learning disability, older people, children and families, young people, Lesbian Gay Bi-sexual or Transgender, minority ethnic communities, Gypsy/Roma/Travellers, women/men, parents, carers, staff, wider community, offenders.

EQUALITY PROTECTED CHARACTERISTICS TO BE CONSIDERED

Age, gender, disability, race, sexual orientation, gender identity (or reassignment), religion and belief, carers, Human Rights and socio economic/deprivation.

socio economic/deprivation.				
QUESTION	RESPONSE		IMPACT	
	Issue	Action	Positive	Negative
What is the service, leaflet or policy development? What are its aims, who are the target audience?	The Procedural Document is to ensure that all members of staff have clear guidance on processes to be followed. The target audience is all staff across the Organisation who undertakes this process.	Raise awareness of the Organisations format and processes involved in relation to the procedural document.	Yes – Clear processes identified	
Does the service, leaflet or policy/ development impact on community safety Crime Community cohesion	Not applicable to community safety or crime	N/A	N/A	
Is there any evidence that groups who should benefit do not? i.e. equal opportunity monitoring of service users and/or staff. If none/insufficient local or national data available consider what information you need.	No	N/A	N/A	
Does the service, leaflet or development/ policy have a negative impact on any geographical or sub group of the population?	No	N/A	N/A	
How does the service, leaflet or policy/ development promote equality and diversity?	Ensures a cohesive approach across the Organisation in relation to the procedural document.	All policies and procedural documents include an EA to identify any positive or negative impacts.		
Does the service, leaflet or policy/ development explicitly include a commitment to equality and diversity and meeting needs? How does it demonstrate its impact?	The Procedure includes a completed EA which provides the opportunity to highlight any potential for a negative / adverse impact.			
Does the Organisation or service workforce reflect the local population? Do we employ people from disadvantaged groups	Our workforce is reflective of the local population.			
Will the service, leaflet or policy/ development i. Improve economic social conditions in deprived areas ii. Use brown field sites Improve public spaces including creation of green spaces?	N/A			
Does the service, leaflet or policy/ development promote equity of lifelong learning?	N/A			
Does the service, leaflet or policy/ development encourage healthy lifestyles and reduce risks to health?				
Does the service, leaflet or policy/ development impact on transport? What are the implications of this?	N/A			
Does the service, leaflet or policy/development impact on housing, housing needs, homelessness, or a person's ability to remain at home?	N/A			
Are there any groups for whom this policy/ service/leaflet would have an impact? Is it an adverse/negative impact? Does it or could it (or is the perception that it could exclude disadvantaged or marginalised groups?	None identified			

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Appendix 3: Equa	lity Impact Assessment Form			
	ACTION:			
Please identify if you are now	required to carry out a Full Equality Analysis	No	(Please delete appropriate)	as
Name of Author: Signature of Author:	Michelle Wong		Date Signed:	
Name of Lead Person: Signature of Lead Person:	Rashmi Sharma		Date Signed:	
Name of Manager: Signature of Manager	Alastair Gibosn		Date Signed:	
	·			