Intra-Abdominal Peritonitis in Adults, Lower Gastrointestinal Tract Origin – Microbiology Full Clinical Guideline

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Introduction

- The lower gastrointestinal tract consists of the small intestine (distal duodenum, jejunum, and ileum) and large intestine (caecum, colon [ascending, transverse, descending, and sigmoid], rectum, and anal canal).
- Intra-abdominal infection of lower gastrointestinal tract origin can be caused by multiple pathogens, i.e. polymicrobial infectious disease.
- Gram negatives (e.g. *Escherichia coli, Klebsiella* spp, and *Proteus* spp), Gram positives (e.g. *Streptococcus* spp and *Enterococcus* spp), and anaerobes (e.g. *Bacteroides fragilis* and *Clostridium* spp) are commonly identified bacterial causes.
- Mechanisms of transmission include mucosal breach, enabling inoculation of gastrointestinal tract flora. Breaches in the mucosa can be secondary to:
 - Perforated viscera.
 - Surgical anastomotic breakdowns.
- Other mechanisms of transmission include contiguous inoculation. Another focus of intra-abdominal viscera infection (e.g. appendicitis or diverticulitis) disseminates locally and invades the abdominal cavity.
- One of the outcomes of:
 - Microbial invasion from the lower gastrointestinal tract into the abdominal cavity; and
 - The subsequent inflammatory response

is peritonitis.

- Manifestations include abdominal pain and tenderness.
- Temperatures > 38°C or < 36°C, a respiratory rate > 20 breaths/minute, a heart rate > 90 beats/minute, and hypotension can denote progression of localised infectious disease into <u>sepsis</u> or septic shock.

Investigation

Radiology

- First line: in general, computed tomography (CT) abdomen pelvis.
- Second line: discuss with the surgical senior and collaborate with the consultant radiologist.

Microbiology

- With the range of bacterial pathogens, variations in resistance and susceptibility profiles, contraindications, and side-effects, microbiological investigation enables best antibiotic practice:
 - Before starting antibiotics: blood cultures × 2, drawn approximately 1-15 minutes apart, from 2 locations/venepunctures.
 - If surgery intervenes: fluid, pus, or tissue for microscopy, culture, and susceptibility (MC&S).

Blood sciences

• Full blood count (FBC), C reactive protein (CRP), lactate, urea and electrolytes (U&Es), and liver function tests (LFTs).



Treatment

Surgical opinion ± intervention

- Intra-abdominal peritonitis can progress from localised infectious disease into <u>sepsis</u> or septic shock.
- Intra-abdominal peritonitis can be secondary to perforated viscera, anastomotic breakdown, or another focus of intra-abdominal infection (e.g. appendicitis or diverticulitis). Therefore, early discussion with the lower gastrointestinal tract registrar/consultant on call is recommended.
- Surgical intervention could enable: (i) elimination of the origin(s) of the infectious episode; (ii) reduction of the microbial inoculum; (iii) identification of the causative agent(s); and, (iv) restoration of host physiological function.

Empiric, intravenous antibiotics

 Community acquired (symptoms, signs, and/or radiological findings of intraabdominal peritonitis within 48 hours of hospital admission):
If clinically stable If clinically unstable (haemodynamic)

	If clinically stable	If clinically unstable (haemodynamic
		instability, sepsis, or septic shock)
First line	Co-amoxiclav 1.2	Piperacillin tazobactam 4.5 g 8 hourly
	g 8 hourly	
Second line, if non-	Ceftriaxone 2 g	Ceftazidime 1 g 8 hourly and
immediate without	24 hourly and	Vancomycin or teicoplanin, dose as per
systemic involvement	Metronidazole	hospital guidelines, vancomycin target
penicillin allergy	500 mg 8 hourly	pre dose level 15-20 mg/l, teicoplanin
		target pre dose level 15-30 mg/l and
		Metronidazole 500 mg 8 hourly
Third line, <u>if</u>	Co-trimoxazole	Ciprofloxacin 400 mg 12 hourly and
immediate rapidly	960 mg 12 hourly	Vancomycin or teicoplanin, dose as per
evolving or non-	and	hospital guidelines, vancomycin target
immediate with	Metronidazole	pre dose level 15-20 mg/l, teicoplanin
systemic involvement	500 mg 8 hourly	target pre dose level 15-30 mg/l and
penicillin allergy	_	Metronidazole 500 mg 8 hourly

• Hospital acquired (symptoms, signs, and/or radiological findings of intra-abdominal peritonitis > 48 hours after hospital admission):

First line	Piperacillin tazobactam 4.5 g 6 hourly
Second line, if non-	Ceftazidime 2 g 8 hourly and
immediate without	Vancomycin or teicoplanin, dose as per hospital guidelines,
systemic involvement	vancomycin target pre dose level 15-20 mg/l, teicoplanin
penicillin allergy	target pre dose level 15-30 mg/l and
	Metronidazole 500 mg 8 hourly
Third line, if immediate	Ciprofloxacin 400 mg 8 hourly and
rapidly evolving or non-	Vancomycin or teicoplanin, dose as per hospital guidelines,
immediate with	vancomycin target pre dose level 15-20 mg/l, teicoplanin
systemic involvement	target pre dose level 15-30 mg/l and
penicillin allergy	Metronidazole 500 mg 8 hourly

• NB Empiric anti-fungals can be considered in specific patients, including recurrent intra-abdominal peritonitis (for example, post-operative recurrence or after completion of anti-bacterials) or history of immunocompromise. However, in general, anti-fungals are reserved for patients with cultures of *Candida* species from blood or intra-operative fluid, pus, or tissue.



Directed, intravenous antibiotics (with susceptibilities)

• Reflecting the polymicrobial nature of intra-abdominal peritonitis, microbiologists commonly recommend antibiotics (both for empiric and directed antimicrobial chemotherapy) with Gram negative, Gram positive, and anaerobic spectrums:

If the pre-operative blood and/or intra- operative fluid, pus, or tissue cultures: Gram negatives (e.g. <i>Escherichia coli</i> , <i>Klebsiella</i> spp,	First line Narrowest spectrum of co- amoxiclav or	Second line, <u>if</u> <u>non-immediate</u> <u>without systemic</u> <u>involvement</u> <u>penicillin allerqy</u> Ceftriaxone 2 g 24 hourly and Metronidazole	Third line, <u>if immediate</u> rapidly evolving or non- immediate with systemic involvement penicillin allergy <u>Co-trimoxazole</u> 960 mg 12 hourly and Metronidazole 500 mg 8
Proteus spp)	piperacillin tazobactam <u>standard dosage</u>	500 mg 8 hourly	hourly
<i>Streptococcus</i> species	Co-amoxiclav 1.2 g 8 hourly	Ceftriaxone 2 g 24 hourly and Metronidazole 500 mg 8 hourly	Vancomycin or teicoplanin, <u>dose as per</u> <u>hospital guidelines</u> , vancomycin target pre dose level 15-20 mg/l, teicoplanin target pre dose level 15-30 mg/l and <u>Co-trimoxazole</u> 960 mg 12 hourly and Metronidazole 500 mg 8 hourly
Enterococcus species	Co-amoxiclav 1.2 g 8 hourly	Vancomycin or teicoplanin, <u>dose</u> <u>as per hospital</u> <u>guidelines</u> , vancomycin target pre dose level 15-20 mg/l, teicoplanin target pre dose level 15-30 mg/l and Ceftriaxone 2 g 24 hourly and Metronidazole 500 mg 8 hourly	Vancomycin or teicoplanin, <u>dose as per</u> <u>hospital guidelines</u> , vancomycin target pre dose level 15-20 mg/l, teicoplanin target pre dose level 15-30 mg/l and <u>Co-trimoxazole</u> 960 mg 12 hourly and Metronidazole 500 mg 8 hourly
Anaerobes (e.g. Bacteroides fragilis, Clostridium spp)	Co-amoxiclav 1.2 g 8 hourly	Ceftriaxone 2 g 24 hourly and Metronidazole 500 mg 8 hourly	<u>Co-trimoxazole</u> 960 mg 12 hourly and Metronidazole 500 mg 8 hourly

 NB Please note, directed antimicrobial chemotherapy relates to pre-operative blood cultures and/or intra-operative fluid, pus, or tissue sterile site MC&S. Postoperative wounds and chronic drains can become colonised with single or multiple microorganisms. With the administration of pre- and post-operative broad spectrum anti-bacterials, non-sterile site investigations may isolate multi-drug resistant, colonising flora only.

Directed, per oral antibiotics (with susceptibilities)

• Reflecting the polymicrobial nature of intra-abdominal peritonitis, microbiologists commonly recommend antibiotics (both for empiric and directed antimicrobial chemotherapy) with Gram negative, Gram positive, and anaerobic spectrums:

If the pre-operative blood and/or intra-operative fluid, pus, or tissue cultures:	First line	Second line	Third line
Gram negatives (e.g. <i>Escherichia coli, Klebsiella</i> spp, <i>Proteus</i> spp)	Co-amoxiclav 625 mg 8 hourly plus amoxicillin 500 mg 8 hourly	Co-trimoxazole 960 mg 12 hourly and Metronidazole 400 mg 8 hourly	Ciprofloxacin 500 mg 12 hourly and Metronidazole 400 mg 8 hourly
Streptococcus species	Co-amoxiclav 625 mg 8 hourly plus amoxicillin 500 mg 8 hourly	Clindamycin 300 mg 6 hourly and <u>Co-trimoxazole</u> 960 mg 12 hourly	Linezolid 600 mg 12 hourly and <u>Co-trimoxazole</u> 960 mg 12 hourly and Metronidazole 400 mg 8 hourly
Enterococcus species	Co-amoxiclav 625 mg 8 hourly plus amoxicillin 500 mg 8 hourly	Linezolid 600 mg per oral 12 hourly and <u>Co-trimoxazole</u> 960 mg 12 hourly and Metronidazole 400 mg 8 hourly	Linezolid 600 mg per oral 12 hourly and Ciprofloxacin 500 mg 12 hourly and Metronidazole 400 mg 8 hourly
Anaerobes (e.g. Bacteroides fragilis, Clostridium spp)	Co-amoxiclav 625 mg 8 hourly plus amoxicillin 500 mg 8 hourly	Co-trimoxazole 960 mg 12 hourly and Metronidazole 400 mg 8 hourly	Ciprofloxacin 500 mg 12 hourly and Metronidazole 400 mg 8 hourly

Duration of antibiotics

- In general, 4 days from surgical intervention and source control.
- NB If the episode of intra-abdominal peritonitis has been complicated (e.g. suboptimal source control or surgical drain in situ or bloodstream infection), collaborate with the microbiology consultant responsible for sterile site investigations.



Management

Clinical concerns re intra-abdominal peritonitis (manifesting symptoms and signs include abdominal pain and tenderness) Investigation Radiology: • First line: in general, CT abdomen pelvis o Second line: discuss with the surgical senior and collaborate with the consultant radiologist Microbiology: • Before starting antibiotics: blood cultures x 2, drawn approximately 1-15 minutes apart, from 2 locations/venepunctures Blood sciences: • FBC, CRP, lactate, U&Es, and LFTs Treatment • Surgical opinion ± intervention: o Consult with the lower gastrointestinal tract registrar/consultant on call • Empiric, intravenous antibiotics (please note, page 2) • NB Empiric anti-fungals can be considered in specific patients, including recurrent intra-abdominal peritonitis (for example, post-operative recurrence or after completion of anti-bacterials) or history of immunocompromise. However, in general, anti-fungals are reserved for patients with cultures of Candida species from blood or intra-operative fluid, pus, or tissue Investigation (if surgery intervenes): Microbiology: • Fluid, pus, or tissue for MC&S

Treatment

• Directed, intravenous antibiotics (please note, pages 3-4)

• In general, 4 days from surgical intervention and source control

References

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