

# SPE3 Feedback session

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Barts & The London

# User notes

- This is a formative exam
- **Please do not share outside of your cohort**
  - This will lessen the use for future years

*If not sure about something based on the feedback or anything else PSA related, feel free to email me at [v.kapil@qmul.ac.uk](mailto:v.kapil@qmul.ac.uk) with your query and if we can't figure out by email, we can Teams*

# User notes

- Please use this for your learning around the topics and familiarising yourself with the BNF(s)
- The PWS questions will be hand-marked
- Any truly disputable questions/answers will not have made it into a final PSA exam
- PWS (10-point prescribing) mark schemes potentially will vary in the real exam, but this is a reasonable indicative guide
- There is no reasonable way to reproduce the full “look” of the PSA exam with the Rogo/SPE system

# Summary statistics

- Internal Anghoff method pass mark 61%
  - Fail (<61%)
  - Borderline pass (61-65%)
  - Good pass (>65%)
- As a rule of thumb, you should do the PWS and REV questions with at least half your overall time left
- If you think you will spend too long on a calculation, I suggest you move on first
- Be familiar with the BNF
- Work through any screenshots I have provided

**Case presentation**

A 46-year-old woman has been unable to take her usual medications for Addison's disease due to viral gastroenteritis causing vomiting. She is admitted into hospital in a collapsed state **PMH**. Addison's disease. **DH**. Hydrocortisone orally 10mg in morning, 5mg in afternoon, 5mg in evening; fludrocortisone 100 micrograms orally daily.

She is diagnosed with an Addisonian crisis with hypotension and hypoglycaemia. She has been given appropriate intravenous fluid and glucose resuscitation.

**Examination (after initial fluid resuscitation and glucose treatment)**

BP 118/68 mmHg, pulse 96 regular  
Sats 98% OA, RR 12, chest clear  
Blood capillary glucose 8.2 mmol/l  
GCS 14/15

ECG sinus rhythm

**Prescribing request**

Write a prescription for ONE drug that will help to treat her condition  
(use the hospital 'once-only medicines' prescription chart provided)

ONCE ONLY MEDICINES							
Date DD/MM/YYYY	Time HH:MM	Medicine (Approved Name)	Dose	Route	Prescriber Signature <i>including surname</i>	Time Given	Given By

# Learning point – select the dose for the right indication!

## Indications and dose

### For HYDROCORTISONE

#### Thyrotoxic crisis (thyroid storm)

By intravenous injection

##### For Adult

100mg every 6 hours, to be administered as sodium succinate.

#### Adrenocortical insufficiency resulting from septic shock

By intravenous injection

##### For Adult

50mg every 6 hours, given in combination with fludrocortisone.

#### Acute hypersensitivity reactions such as angioedema of the upper respiratory tract and anaphylaxis (adjunct to adrenaline)

By intravenous injection

##### For Adult

100–300mg, to be administered as sodium succinate.

#### Corticosteroid replacement, in patients who have taken more than 10mg prednisolone daily (or equivalent) within 3 months of minor surgery under general anaesthesia

By intravenous injection, or by intravenous infusion

##### For Adult

Initially 25–50 mg, to be administered at induction of surgery; the patient's usual oral corticosteroid dose is recommenced after surgery.

### Adrenal crisis

#### Adrenal crisis

Initially by intramuscular injection, or by intravenous injection

##### For Adult

Initially 100 mg, then (by continuous intravenous infusion) 200 mg every 24 hours, diluted in Glucose 5%, alternatively (by intramuscular injection or by intravenous injection) 50 mg every 6 hours, dose increased to 100 mg every 6 hours in patients who are severely obese.

A. Drug choice				Score	Feedback/justification	
1	Hydrocortisone	5	Hydrocortisone is the initial management of Addisonian crises			

**Case presentation**

A 38-year-old woman in her 2<sup>nd</sup> trimester is attends the antenatal ward with a 3-day history of foul-smelling urine, dysuria and fever. **PMH.** None. **DH.** No regular medicines. **SH.** She is a non-smoker.

**On examination**

Appears well

Temperature 37.8°C.

There is no renal angle or abdominal tenderness.

Pulse 92/min, blood pressure 112/84 mmHg

**Investigations**

Urinalysis reveals nitrites +++, leucocytes +++, no protein.

She is clinically well, and is due to be discharged with medications.

**Prescribing request**

Write a prescription for ONE drug that will be continued as an outpatient to treat her condition  
(use the hospital 'regular medicines' prescription chart provided)

		Date →					
		Time ↕					
Medicine (Approved name)		6	<input type="checkbox"/>				
<input type="text"/>		8	<input type="checkbox"/>				
Dose	Route	12	<input type="checkbox"/>				
<input type="text"/>	<input type="text"/>						
Prescriber Signature including surname	Start date (DD/MM/YYYY)	14	<input type="checkbox"/>				
<input type="text"/>	<input type="text"/>	18	<input type="checkbox"/>				
Notes	Pharmacy	22	<input type="checkbox"/>				



NICE National Institute for Health and Care Excellence

Evidence search BNF BNFC CKS Journals and data

urinary tract infection

Drugs | Interactions | Treatment Summaries | What's Changed?

Home > Search: urinary tract infection

Showing 1-10 of 84 results for "urinary tract infection"

**Urinary-tract infections** *Treatment summary*

of age; in patients with suspected upper **urinary-tract infection**; complicated **infection**; or recurrent **infection**; if resistant organisms are suspected; if

**Urinary retention** *Treatment summary*

acute **urinary retention**, haematuria, renal failure, bladder calculi or recurrent **urinary-tract infection**. Related drugs Other drugs used for **urinary retention**:

NICE National Institute for Health and Care Excellence

Evidence search BNF BNFC CKS Journals and data

Search...

Drugs | Interactions | Treatment Summaries | What's Changed?

Home > Treatment summary > Urinary-tract infections

## Urinary-tract infections

### Overview

Urinary-tract infection is more common in women than in men; when it occurs in men there is frequently an underlying abnormality of the renal tract. Recurrent episodes of infection are an indication for radiological investigation especially in children in whom untreated pyelonephritis may lead to permanent kidney damage.

*Escherichia coli* is the most common cause of urinary-tract infection; *Staphylococcus saprophyticus* is also common in sexually active young women. Less common causes include *Proteus* and *Klebsiella* spp. *Pseudomonas aeruginosa* infections usually occur in the hospital setting and may be associated with functional or anatomical abnormalities of the renal tract. *Staphylococcus epidermidis* and *Enterococcus faecalis* infection may complicate catheterisation or instrumentation.

A specimen of urine should be collected for culture and sensitivity testing before starting antibacterial therapy.

- In men;
- In pregnant women;
- In children under 3 years of age;

**Related drugs**

- AMOXICILLIN
- AMPICILLIN
- CEFALOXIN
- CEFUROXIME
- CIPROFLOXACIN
- CO-AMOXICLAV
- FOSFOMYCIN
- NITROFURANTOIN
- NITROFURANTOIN HYDROCHLORIDE

**Scroll down or "Ctrl-F" Pregnancy**

**Pregnant women**

An immediate antibacterial prescription should be given and a midstream urine sample obtained before treatment is taken and sent for culture and susceptibility testing.

### Choice of antibacterial therapy

- **Oral first line:**
  - Nitrofurantoin.
- **Oral second line** (if no improvement after at least 48 hours, or first line not suitable):
  - Amoxicillin (only if culture susceptible), or cefalexin.
- **Alternative second line:**
  - Consult local microbiologist.
- **Asymptomatic bacteriuria:**
  - Amoxicillin, cefalexin, or nitrofurantoin.

**Prostatitis, acute**

# Pregnancy antibiotics for UTIs - principles

	First trimester	Second trimester	Third trimester
Nitrofurantoin	Suitable in first and second trimesters (but may be second line to penicillins)		<b>should be avoided at term (may produce neonatal haemolysis)</b>
Penicillins and cephalosporins	Suitable for treating urinary-tract infection during pregnancy, BUT second line in BNF <b>(if not allergic)</b>		
Sulfonamides (e.g. sulfamethoxazole – in folate synthesis pathway) and quinolones (e.g. ciprofloxacin / levofloxacin)	<b>Avoided during pregnancy</b>		
Trimethoprim	<b>Avoided particularly in the first trimester</b>	<b>Manufacturers recommend avoiding throughout</b>	

## A. Drug choice

		Score	Feedback/justification
1	Nitrofurantoin	5	First line in BNF (but in other trusts amox may be first line)
2	Amoxicillin Cefalexin	4	Beta lactams or orally active cephalosporins are appropriate in pregnancy May be second line
3	Cephalosporins with only IV formulations	1	These will work but are reserved for those unable to tolerate oral medications
4	Trimethoprim	0	Should be avoided if possible, especially in first trimester
5	Sulphonamides	0	Contraindicated in pregnancy

## B. Dose, route, frequency

	Score	Feedback/justification
50 mg orally 6-hrly (or 100mg orally 12hrly for MODIFIED RELEASE)	5	This is the optimum dose
100 mg orally 6 hrly	3	
500 mg orally 8-hrly 500 mg orally 12-hrly	4	This is the optimal dosage and should be given for up to 7 days No clear indication for intravenous antibiotics
Any above but either too low or high oral dose	2	Too high dose
Appropriate iv dose	1	Intravenous route is reserved for those unable to take orally.
Appropriate iv dose	1	
All doses	0	
All doses	0	

**Case presentation**

A 50-year-old man presents to the Emergency Department with complete dysphagia due to a food bolus. He is made “nil by mouth” pending endoscopy tomorrow. **PMH** Nil. **DH.** Nil.

**On examination**

Temperature 37.1°C, HR 72/min and regular, BP 122/70 mmHg, O<sub>2</sub> sat 99% on RA. Weight 70kg

He has received already:

1L 5% dextrose/0.15% KCl over 12h

**Prescribing request**

Write a prescription for ONE intravenous fluid that would be most appropriate for the patient to receive next  
(use the hospital fluid prescription chart provided)

Date DD/MM/YYYY	Start Time HH:MM	Infusion Solution	Volume	Duration	Infusion Rate [mL/min]	Signature including surname
				please select ▼		

[Home](#) > [Treatment summary](#) > [Fluids and electrolytes](#)

You are viewing BNF. If you require BNF for Children, use [BNFC](#).

# Fluids and electrolytes

## Electrolyte replacement therapy

The electrolyte concentrations (intravenous fluid) table and the electrolyte content (gastro-intestinal secretions) table may be helpful in planning replacement electrolyte therapy. Faeces, vomit, or aspiration should be saved and analysed where possible if abnormal losses are suspected.

## Oral preparations for fluid and electrolyte imbalance

Sodium and potassium salts, may be given by mouth to prevent deficiencies or to treat established deficiencies of mild or moderate degree.

## Oral potassium

Compensation for potassium loss is especially necessary:

- in those taking digoxin or anti-arrhythmic drugs, where potassium depletion may induce arrhythmias.

### Related drugs

- [ALBUMIN SOLUTION](#)
- [CALCIUM GLUCONATE](#)
- [FUROSEMIDE](#)
- [GELATIN](#)
- [GLUCOSE](#)

[SODIUM](#)

**This section is not helpful for PSA**

- [SODIUM CHLORIDE WITH GLUCOSE](#)

### Related Treatment Summaries

- [Diarrhoea \(acute\)](#)

## Fluids prescribing question

Is this a question about emergency resuscitation?  
(Examples include acute volume loss, hypotensive, tachycardic etc.)

YES

0.9% Sodium chloride, 500 ml in 10 minutes  
(NO potassium)

NO

Is this a question about emergency hypoglycaemia?  
(If symptomatic severe hypoglycaemia in the fluids questions)

YES

10% Glucose, 150 ml in 15 minutes  
(NO potassium)

NO

Is this a question about symptomatic severe hypercalcaemia?  
(unless complicated by below)

YES

0.9% Sodium chloride, 1000 ml in 2-4 hours  
(NO potassium)

NO

Is there only a need to maintain input without  
fluid/electrolyte deficit/ongoing losses?  
(Maintenance alone – e.g. nil by mouth for surgery/stroke  
without prior losses such as diarrhoea / vomiting)

YES

For routine maintenance alone  
25–30 ml/kg/day of water  
~ 1 mmol/kg/day of K<sup>+</sup>, Na<sup>+</sup>, Cl<sup>-</sup>  
~ 50–100 g/day of glucose to limit starvation ketosis  
Likely to be penalised if use small volumes if overall daily prescription ≥3L

Example FIRST prescription:  
0.9% sodium chloride with 0.3% potassium chloride, 1000ml over 8-12 hours  
5% glucose with 0.3% potassium chloride, 1000ml over 8-12 hours

**This  
question**

BUT: Sometimes the question already provides first prescription, and you need to calculate what is missing for second prescription

NO

Is there also a fluid deficit?  
(Fluid replacement ± maintenance)

YES

Try to estimate deficit for **BOTH** volume and electrolytes (looks at U+Es, and high Na/K loss with GI losses)  
– aim to replace in 1-2 days  
Add on replacing ongoing losses

### Volume

- Likely >3L / day, so likely 1L over ≤6 hours
- But be careful of too giving fluids too fast (unlikely to be >6L/day, so unlikely to be 1L <4 hours)

### Electrolytes

- Likely to need 0.9% sodium chloride (due to electrolyte losses)
- If hypokalaemia (especially with active loss) may require initial fluids with maximal rate of KCl (10mmol/hr):
  - for example → 1L 0.9% sodium chloride with 0.3% potassium chloride in 4 hrs

# IV fluids – this question

Daily requirements are [NICE guidelines]:		Previous / ongoing losses	This scenario	Suggested prescription
25–30 ml/kg/d water	= 1750-2100mL water/day	None	Already had 1L in 12hrs	Therefore consider <b><u>1000mL in 12 hours</u></b>
1 mmol/kg/day sodium, chloride	= 70 mmol/day	None	None in previous 12hr	Therefore other fluids should contain Na <sup>+</sup> and Cl <sup>-</sup> <ul style="list-style-type: none"> <li>Consider <b><u>0.9% sodium chloride (+- potassium, see later)</u></b> or other balanced crystalloids (cannot adjust potassium)</li> </ul>
1 mmol/kg/day potassium	= 70 mmol/day	None	(Has had 1L with 0.15% KCl) Therefore already had 20 mmol/L in 12hrs	Therefore requires at least 40 mmol KCl <ul style="list-style-type: none"> <li>Available in <b><u>0.9% sodium chloride with 0.3% KCl (note: 0.3% KCl = 40mmol/L KCl)</u></b></li> <li>I would suggest that balanced crystalloids (with only 3-5 mmol/L KCl) may be insufficient in context of earlier</li> </ul>
50–100 g/day glucose (NB. glucose 5% contains 5 g/100ml)		None	Already had 50g in 12hrs	No need to have glucose, so should still be based on above

**Case presentation**

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(use the hospital fluid prescription chart provided)

Date DD/MM/YYYY	Start Time HH:MM	Infusion Solution	Volume	Duration	Infusion Rate [mL/min]	Signature including surname
				please select ▼		



A. Drug choice		Score	Feedback/justification		B. Dose and route		Score	Feedback/justification
1	0.9% sodium chloride with 0.3% KCl	5	See previous slide for feedback		1L in 12 hours (accept 10 hours)	5	See previous slide for feedback	
					1L in 4-8 hours OR 1L in 16-24 hours	3	Either too fast or too slow	
					Smaller volumes	1	Inappropriate to prescribe smaller volumes if patient stable	
2	0.9% sodium chloride with 0.15% KCl	4	Insufficient potassium		1L in 12 hours	4	See previous slide for feedback	
					1L in 4-8 hours OR 1L in 16-24 hours	2	Either too fast or too slow	
3	Hartmann's, Ringer's, Plasmalyte (or other balanced solutions) 0.9% sodium chloride (without KCl)	3	insufficient potassium		1L in 12 hours	3	See previous slide for feedback	
					1L in 4-8 hours OR 1L in 16-24 hours	1	Either too fast or too slow	
4	5% glucose with 0.3% KCl	3	Insufficient to meet sodium requirements		1L in 12 hours	3	See previous slide for feedback	
					1L in 4-8 hours OR 1L in 16-24 hours	1	Either too fast or too slow	
5	5% glucose (without KCl)	1	Insufficient sodium and potassium		1L in 12 hours	1	See previous slide for feedback	
6	Other fluids	0	Likely to score very low					

# **Be familiar with the PSA exam interface**

- Use the demonstration paper and click through

ID : UK-01-1208

Prescribing

Hospital Fluid

10 marks



☐ Mark for review

A A



#### Case presentation

A 74-year-old man is admitted to hospital 7 hours after an acute stroke. He had been unwell for 2 days and had been eating and drinking less than usual.

**PMH.** Hypertension. **DH.** Ramipril 10 mg PO daily.

#### On examination

Temperature 36.4°C, HR 88/min and regular, BP 135/76 mmHg. Alert, dysphasic and has a right hemiparesis. He is unable to swallow and does not tolerate insertion of a nasogastric tube.

#### Investigations

Na<sup>+</sup> 144 mmol/L (137–144), K<sup>+</sup> 3.9 mmol/L (3.5–4.9), U 7.5 mmol/L (2.5–7.0), Cr 85 µmol/L (60–110), random plasma glucose 7.2 mmol/L.  
CXR shows no evidence of cardiac failure.  
ECG shows no changes suggestive of recent myocardial infarction.

#### Prescribing request

Write a prescription for ONE IV fluid that is *most appropriate* for the patient at this stage.

(use the 'hospital IV fluid' prescription form provided)

#### PRESCRIPTION FORM

##### Infusion Fluid

normal  
sodium chloride 0.45% solution  
sodium chloride 0.9% solution

##### Duration

Enter characters to search

##### Infusion Rate (mL/min) ⓘ

-

##### Prescriber

Fu Ng

##### Date

14/11/2019

##### Time

15:50

NOTE: Not “normal saline”

**Case presentation**

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**Prescribing request****PRESCRIPTION FORM****Infusion Fluid**

sodium

sodium bicarbonate 1.26% solution

sodium bicarbonate 8.4% solution

sodium chloride 0.18%/glucose 4% solution

sodium chloride 0.18%/glucose 4%/potassium chloride 0.15% solution

sodium chloride 0.18%/glucose 4%/potassium chloride 0.3% solution

sodium chloride 0.45% solution

sodium chloride 0.45%/glucose 2.5% solution

sodium chloride 0.45%/glucose 5% solution

**Scroll down****Case presentation**

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**Prescribing request**

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(use the 'hospital IV fluid' prescription form provided)

**PRESCRIPTION FORM****Infusion Fluid**

sodium

sodium chloride 0.45%/glucose 5%/potassium chloride 0.15% solution

sodium chloride 0.9% solution

sodium chloride 0.9%/glucose 5% solution

sodium chloride 0.9%/potassium chloride 0.15% solution

sodium chloride 0.9%/potassium chloride 0.3% solution

dextran '70' 6% /sodium chloride 7.5% solution

disodium hydrogen phosphate 5.75 g/L, potassium dihydrogen phosphate 1.295 g/L

ID : UK-01-1208

Prescribing

Hospital Fluid

10 marks

☐ Mark for review

A A

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**Prescribing request**

Write a prescription for ONE IV fluid that is *most appropriate* for the patient at this stage.

(use the 'hospital IV fluid' prescription form provided)

**PRESCRIPTION FORM****Infusion Fluid**

hart

Hartmann's solution (Na<sup>+</sup> 131/K<sup>+</sup> 5/Ca<sup>2+</sup> 2/HCO<sub>3</sub><sup>-</sup> 29/Cl<sup>-</sup> 111 mmol/L)

Enter characters to search

IV

**Duration**

Enter characters to search

**Infusion Rate (mL/min) ⓘ**

-

**Prescriber**

Fu Ng

**Date**

14/11/2019

**Time**

15:50

ID : UK-01-1208

Prescribing

Hospital Fluid

10 marks

☐ Mark for review

A A

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**Prescribing request**

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(use the 'hospital IV fluid' prescription form provided)

**PRESCRIPTION FORM****Infusion Fluid**

plas

Plasma-Lyte® 1:1:8 (glucose 5%) solution (Na+ 140/K+ 5/Mg2+ 1.5/Cl- 98 mmol/L)

Plasma-Lyte® 1:1:8 (water) solution (Na+ 140/K+ 5/Mg2+ 1.5/Cl- 98 mmol/L)

fresh frozen plasma

**Prescriber**

Fu Ng

**Date**

14/11/2019

**Time**

15:50

**NOTE: Not “dextrose”**

**ID : UK-01-1208** **Prescribing** **Hospital Fluid** **10 marks** ☐ Mark for review A A

**Case presentation**  
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**Prescribing request**  
Write a prescription for ONE IV fluid that is *most appropriate* for the patient at this stage.  
(use the 'hospital IV fluid' prescription form provided)

**PRESCRIPTION FORM**  
**Infusion Fluid**

glucose 10% solution
glucose 20% solution
glucose 5% solution
glucose 5%/potassium chloride 0.15% solution
glucose 5%/potassium chloride 0.3% solution
glucose 50% solution
Plasma-Lyte® 148 (glucose 5%) solution (Na <sup>+</sup> 140/K <sup>+</sup> 5/Mg2 <sup>+</sup> 1.5/Cl <sup>-</sup> 98 mmol/L)
sodium chloride 0.18%/glucose 4% solution

**NOTE:**

0.15% potassium chloride = 20mmol/L

0.3% potassium chloride = 40mmol/L

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**Prescribing request**

Write a prescription for ONE IV fluid that is *most appropriate* for the patient at this stage.

(use the 'hospital IV fluid' prescription form provided)

**PRESCRIPTION FORM****Infusion Fluid****Volume**

500 mL

50 mL

5 mL

**Prescriber**

Fu Ng

**Route**

IV

**Infusion Rate (mL/min)**

-

**Date**

17/11/2019

**Time**

23:21

**NOTE:**

- Fixed volumes

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(use the 'hospital IV fluid' prescription form provided)

**PRESCRIPTION FORM****Infusion Fluid****Volume**

1000 mL

100 mL

10 mL

150 mL

1 unit (blood product)

**Route**

IV

**Infusion Rate (mL/min)**

-

**Date**

17/11/2019

**Time**

23:21



ID : UK-01-1208   Prescribing   Hospital Fluid   10 marks  

**Case presentation**  
A 74-year-old man is admitted to hospital 7 hours after an acute stroke. He had been unwell for 2 days and had been eating and drinking less than usual.  
**PMH.** Hypertension. **DH.** Ramipril 10 mg PO daily.

**On examination**  
Temperature 36.4°C, HR 88/min and regular, BP 135/76 mmHg. Alert, dysphasic and has a right hemiparesis. He is unable to swallow and does not tolerate insertion of a nasogastric tube.

**Investigations**  
Na<sup>+</sup> 144 mmol/L (137–144), K<sup>+</sup> 3.9 mmol/L (3.5–4.9), U 7.5 mmol/L (2.5–7.0), Cr 85 μmol/L (60–110), random plasma glucose 7.2 mmol/L.  
CXR shows no evidence of cardiac failure.  
ECG shows no changes suggestive of recent myocardial infarction.

**Prescribing request**  
Write a prescription for ONE IV fluid that is *most appropriate* for the patient at this stage.  
(use the 'hospital IV fluid' prescription form provided)

**PRESCRIPTION FORM**  
**Infusion Fluid**  
sodium chloride 0.9% solution x

**Volume**  
x

**Duration**  
2m  
5m  
10m  
15m  
20m  
30m  
60m  
90m  
2h

**Route**  
IV

**Infusion Rate (mL/min) ⓘ**

**Date**  
17/11/2019

**Time**  
23:23

**NOTE:**

Resus Bolus of 500ml “less than 15 minute”

**NOTE:**

- Fixed time durations
- No “stat” prescription
- unlikely able to give bolus 500ml in 2 minutes

**Case presentation**

A 72 year old man presents to his General Practitioner after developing severe pain on his torso, 2 weeks after an attack of shingles. **PMH.** Hypertension, type II diabetes, gastric ulcer secondary to NSAIDs. **DH.** Amlodipine 5 mg orally daily, Metformin 850mg orally twice a day, paracetamol 1 grams four times a day, omeprazole 20mg orally daily.

Pharmacy Stamp	Age	
Please don't stamp over age box	D.o.B.	
Number of days' treatment N.B. Ensure dose is stated		
Endorsements	Drug Name	
	Dose	
	Route	
	Frequency	
	Signature of Prescriber including surname	Date DD/MM/YYYY
For Dispenser No. of Prescns. on form	Sunshine Health Centre Dr Tom Baker Mill Lane SZ9 9FF, London Tel: 123456789	

**Prescribing request**

Write the initial prescription for ONE drug that will help to improve his symptoms.

*(use the general practice prescription form provided)*

# Learning Point - What is being treated?

- **KEY MESSAGE** – Read the question on which is the condition/symptom is being treated.
  - The correct answer to this question was to treat the pain
  - If you were asked to treat the condition, you would consider treating the infection
    - (e.g. while the infection is active within 72 hours of the rash)

A. Drug choice				B. Dose, route, freq.			
		Score	Feedback/justification		Score	Feedback/justification	
1	Pregabalin	5	Part of first-line choices for neuropathic pain	Orally, Initially 150 mg daily in 2–3 divided doses, then increased if necessary to 300 mg daily in 2–3 divided doses, dose to be increased after 3–7 days, then increased if necessary up to 600 mg daily in 2–3 divided doses, dose to be increased after 7 days.	5	Choose initial script not titration	
				Higher than starting dose	2		
2	Amitriptyline or Nortriptyline	5	Part of first-line choices for neuropathic pain [Unlicensed indication]	Orally. Initially 10-25 mg once daily for A or 10mg once daily for N	5		
				Higher than starting dose	2		
3	Gabapentin	5	Part of first-line choices for neuropathic pain	Orally. Initially 300 mg once daily on day 1, then 300 mg twice daily on day 2, then 300 mg 3 times a day on day 3, alternatively initially 300 mg 3 times a day on day 1, then increased in steps of 300 mg every 2–3 days in 3 divided doses, adjusted according to response; maximum 3.6 g per day	5	Choose initial script not titration	
				Higher than starting dose	2		
4	Tramadol	2	Opioid analgesia may be effective, but not first line	Appropriate dosing	2		
5	Codeine/Lidocaine	1	Opioid analgesia may be effective, but not first line	Appropriate dosing	1		
6	Morphine/Capsaicin	0	Although may be effective, only to be started in specialist care (not in GP)				
7	Co-codamol	0	Already on full dose				

### Case presentation

A 79-year-old woman presents to Accident and Emergency with a painful left hip. She reports that she became dizzy two hours after taking her tablets. Her friend tested her capillary blood glucose minutes after the fall – 9.5 mmol/L. **PMH.** Hypercholesterolaemia, hypertension, type 2 diabetes mellitus, neuropathic pain, giant cell arteritis. **DH.** Her current regular medicines are listed (right). **SH.** Lives alone, independent.

### Investigations

Blood Glucose: 8.2 mmol/L

Full blood count and U+Es - normal

X-ray of left hip: fractured neck of femur.

### Question A

Select TWO medications that are most likely to have raised her risk of fractures.

(mark them with a tick in column A)

### Question B

Select TWO prescriptions that are *most likely* to be contributing to her fall.

(mark them with a tick in column B)

### CURRENT PRESCRIPTIONS

Drug name	Dose	Route	Freq.	A	B
Amitriptilline	75 mg	ORAL	Nightly		✓
Aspirin	75 mg	ORAL	Daily		
Atorvastatin	20 mg	ORAL	Daily		
Gaviscon	10 ml	ORAL	PRN		
Gliclazide MR	30mg	ORAL	DAILY		
Metformin	1000 mg	ORAL	Twice daily		
Omeprazole	20mg	ORAL	Daily	✓	
Prednisolone	15 mg	ORAL	Daily	✓	
Ramipril	10 mg	ORAL	Daily		✓
Senna	15 mg	ORAL	Nightly		

### Answer box

Question A Marks per correct tick [2]

Prednisolone in this dose is common in giant cell arteritis. However, steroids in these doses taken long-term cause major problems including osteoporosis and diabetes

Proton-pump inhibitors can increase the risk of fractures (particularly when used at high doses for over a year in the elderly) (in adults)

Question B Marks per correct tick [1]

Ramipril-associated hypotension can make the patient feel dizzy and lead to falls. For ramipril daily doses of 5mg and above, ideally split into two doses given 12-hourly. A hypoglycaemic episode from the sulfonylurea is unlikely given glucose level. Amitriptilline, particularly at higher doses, increases the risk of falls and postural hypotension. (GI bleed is a possibility taking steroid and aspirin without PPI protection but urea and haemoglobin are reported in 'normal range').

# Question (A) – Risk of medicines-related osteoporosis

- Steroids
- PPIs at high doses can increase risk of fractures, especially in elderly over long courses
- Long-term androgen suppression (e.g. LHRH agonists such as buserelin, goserelin for prostate cancer)
- LMWHs (long term)
- There are other rarer causes

# Learning point: Consider other subheadings

- Side effects can also be under:
  - Cautions
  - Important safety information etc.

treatment of SCLL only if there are no signs of remission after a few weeks or months.

## Cautions

### For all PROTON PUMP INHIBITORS

Can increase the risk of fractures (particularly when used at high doses for over a year in the elderly) (in adults); may increase the risk of gastro-intestinal infections (including *Clostridioides difficile* infection); may mask the symptoms of gastric cancer (in adults); patients at risk of osteoporosis

### Cautions, further information

#### Risk of osteoporosis

Patients at risk of osteoporosis should maintain an adequate intake of calcium and vitamin D, and if necessary, receive other preventative therapy.

#### Gastric cancer

##### In adults

Particular care is required in those presenting with 'alarm features', in such cases gastric malignancy should be ruled out before treatment.

## Interactions

# Question (B) – Risk of medicines-related falls

- Drugs that increase the overall risk of falls (and the baseline risk has many other factors)
  - Benzodiazepines, Z-drugs
  - Antidepressants (especially TCAs and SNRIs, less so SSRIs)
  - Monoamine oxidase inhibitors
  - Most antipsychotics
  - Opiates
  - Most antihypertensives (especially alpha-blockers, diuretics, centrally acting antihypertensives)
  - Some anti-Parkinson's medications (e.g. selegiline, ropinirole)
  - (Less commonly) some antiepileptics
  - In theory, those that cause hypoglycaemia, bleeding but need some evidence of those to answer in PSA...



**Case presentation**

A 57-year-old man attends his GP following discharge from hospital after being treated for an infective exacerbation of chronic obstructive pulmonary disease. He is reporting muscle aches and headaches. **PMH.** Chronic back pain, chronic obstructive pulmonary disease and hypertension. **DH.** In addition to simvastatin 80 mg orally daily, current regular medicines are listed (right).

**Examination**

BP 188/102 mmHg, pulse 84/min regular

**Question A**

Select the TWO prescriptions that are most likely to interact with simvastatin to increase the risk of myopathies.  
(mark them with a tick)

**Question B**

Select TWO prescriptions that are *most likely* to be contributing to his poorly controlled hypertension.  
(mark it with a tick)

**CURRENT PRESCRIPTIONS**

Drug name	Dose	Route	Freq.	A	B
Amiloride	5 mg	ORAL	daily		
Amlodipine	10mg	ORAL	Daily	✓	
Amoxicillin	500mg	ORAL	8-hrly		
Beclomethasone	200 micrograms	INH	12-hrly		
Clarithromycin	500mg	ORAL	12-hrly	✓	
Enoxaparin	40mg	SC	daily		
Ipratropium	40 micrograms	INH	8-hrly		
Naproxen	500 mg	ORAL	12-hrly		✓
Omeprazole	20mg	ORAL	daily		
Prednisolone	40 mg	ORAL	daily		✓

**Answer box**

**Question A** Marks per correct tick 1

Clarithromycin inhibits statin metabolism and therefore puts the patient at risk of statin-induced rhabdomyolysis.

**Question B** Marks per correct tick 1

both glucocorticoids and non-steroidal anti-inflammatory drugs can raise blood pressure through salt and water retention

## **Question (B) – Risk of medicines-related hypertension**

- Inhaled beclomethasone is not associated with hypertension due to limited systemic effects
- NSAIDs are known to contribute to higher blood pressure

### Case presentation

A 33-year-old woman is reviewed in the pre-conception clinic with a view to getting pregnant soon. **PMH.** Atrial fibrillation, mitral stenosis from previous rheumatic fever, hypertension, rheumatoid arthritis, asthma. **DH.** Listed on table.

### Question A

Select the TWO prescriptions that should not be prescribed in someone actively considering pregnancy.  
(mark them with a tick in column A)

### Question B

Select the ONE prescription that contain a dosing error.  
(mark them with a tick in column B)

### CURRENT PRESCRIPTIONS

Drug name	Dose	Route	Freq.
Apixaban	5 mg	ORAL	12-hrly
Digoxin	125 mg	ORAL	Daily
Folic acid	5 mg	ORAL	Weekly
Paracetamol	1 grams	ORAL	6-hrly
Methotrexate	12.5 mg	ORAL	Weekly
Methyldopa	250 mg	ORAL	6-hrly
Symbicort® 100/6	Two puffs	INH.	12-hrly
Terbutaline	500 micrograms	INH.	6-hrly

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### Answer box

Question A and B	Marks per correct tick	1 (for A) 2 (for B)
------------------	------------------------	------------------------

Methotrexate is teratogenic.	D
Apixaban has not been studied in pregnancy, and is currently not recommended. LMWH is recommended anticoagulant during pregnancy.	rl
Digoxin is prescribed in micrograms.	3

# Learning Points Question (A) - Anticoagulation in pregnancy

- Low molecular weight heparins are used because they do not cross the placenta
  - LMWH have a lower risk of osteoporosis and of heparin-induced thrombocytopenia compared to unfractionated heparin
- **Monitoring**
  - anti-Factor Xa activity may be necessary in pregnancy
  - NOTE: will also need to monitor platelets (for heparin-induced thrombocytopenia)
- Dosed according to **renal function**
- Other anticoagulants are not recommended / contraindicated in pregnancy
  - Warfarin and direct oral anticoagulants (e.g. apixaban, dabigatran, edoxaban, rivaroxaban)

# Learning Points Question (A) - DMARDs in pregnancy

- With pregnancy, **most** DMARDs are recommended to be **avoided**:
  - e.g. methotrexate, penicillamine, hydroxychloroquine, azathioprine, ciclosporin, leflunomide, monoclonal antibodies
- If you see a DMARD with pregnancy, there is a strong chance that this is the medication that is contraindicated.
  - It may very rarely be used in highly specialised cases, but that is never as a F1 doctor (and therefore not the PSA)

# Learning Points Question (B) – Unit errors and 10x (or other factor) errors

- Be wary of medications that are typically in **micrograms** or **grams** ranges (most medications are in the mg ranges)
- A **non-exhaustive** list:

<b>Micrograms</b>	Tamsulosin, fludrocortisone (not hydrocortisone), levothyroxine, digoxin, naloxone, inhalers, ipratropium nebs
<b>Milligrams (1 to low 100s)</b>	Most medications
<b>100s mg – grams</b>	Some antibiotics, metformin, some anti-epileptics
<b>Grams</b>	Paracetamol, calcium carbonate, N-acetylcysteine

- 10x errors can be more difficult to spot and depends on your experience

### Case presentation

A 63-year-old woman is admitted to hospital for an elective knee replacement. Her post operative recovery is complicated by a mild surgical site infection. **PMH.** Atrial fibrillation, chronic obstructive pulmonary disease, osteoarthritis. **DH.** In addition to apixaban 5 mg orally 12hrly, her other medications are listed on table. Allergy to penicillin.

She reports that her tongue has a change in colour.

### Examination

Oral candidiasis

### Question A

Select the ONE prescriptions that should not be prescribed together with apixaban.  
(mark them with a tick in column A)

### Question B

Select the TWO prescriptions that *most likely* to increase the risk of oral candidiasis.  
(mark them with a tick in column B)

### CURRENT PRESCRIPTIONS

Drug name	Dose	Route	Freq.
Digoxin	125 micrograms	ORAL	Daily
Doxycycline	200 mg	ORAL	Daily
Enoxaparin	20 mg	S.C.	Daily
Ibuprofen	200 mg	ORAL	PRN
Morphine sulfate	10 mg	ORAL	PRN
Paracetamol	1 g	ORAL	6-hrly
Simvastatin	20 mg	ORAL	Daily
Symbicort® 100/6	Two puffs	INH.	12-hrly
Tiotropium	18 micrograms	INH.	Daily

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### Answer box

Question A Marks per correct tick 2

The patient is already fully anti-coagulated on apixaban [a non VKA anticoagulant] and therefore there is an increased risk of bleeding with dual therapy and the LMWH should be stopped.

Question B Marks per correct tick 1

Symbicort contains an inhaled corticosteroid, which is appropriate mouthcare is not instituted, can increase the risk of oral candidiasis  
Broad spectrum antibiotics also increase the risk of oral candidiasis

# Question (B) – Medications increasing the risk of fungal infections

- Candidiasis may be more prevalent in patients receiving:
  - Corticosteroids
    - Inhaled → oral candidiasis
    - Systemic → vaginal (more than oral) candida
  - Broad-spectrum antibiotics
  - Cytotoxics / immunosuppression
- For oral candidiasis
  - Treatment
    - First line - Oral nystatin (100,000 units, 4x/day)
    - Second line oral fluconazole, IF:
      - Unresponsive infections
      - Topical antifungal drug cannot be used or if the patient has dry mouth
      - Immunocompromised patients



### Case presentation

A 54-year-old man is an in-patient for management of infective endocarditis. He feels light-headed and sweaty after taking his regular insulin but is then taken for an echocardiogram before getting a chance to eat his breakfast. **PMH.** Type I diabetes mellitus, hypertension. **DH.** Flucloxacillin 2g intravenously 4-hrly; Mixtard 30 Insulin 28 units twice daily with meals.

### Examination

Blood pressure 128/72 mmHg, pulse 88/minute regular  
Glasgow Coma Scale 15/15  
Capillary blood glucose 2.2 mmol/L

### Question

Select the *most appropriate* management option at this stage.  
(mark it with a tick)

### MANAGEMENT OPTIONS

- |   |                                 |
|---|---------------------------------|
| A | 50% glucose 30 mL intravenously |
| B | Glucagon 1 mg intramuscularly   |
| C | Liquid glucose 15 grams orally  |
| D | Reduce morning insulin          |
| E | Withhold evening insulin        |

### Answer box

Option A Justification

Option B Justification

Glucagon IM is option is obtunded/seizures

Option C Justification

BNF states: initially [glucose](#) 15–20 g is given by mouth either in fast acting carb form. As awake and GCS normal, oral route preferable to invasive treatment

Option D Justification

Doesn't treat current hypoglycaemia, and the cause of the hypoglycaemia is missing a meal (rather than dose being too high)

Option E Justification

Doesn't treat current hypoglycaemia, and the cause of the hypoglycaemia is missing a meal (rather than dose being too high)

# Hypoglycaemia management

- Dependent on:
  - Alert vs. unconscious
  - IV access vs. no access

	Management
<b>Alert</b>	Glucose 15-20 g orally in liquid form followed by sustained carbohydrate
<b>Reduced consciousness</b>	<ol style="list-style-type: none"><li>1. Glucagon (1mg subcutaneously or intramuscularly)</li><li>2. 10% glucose (150-200ml) intravenously<ul style="list-style-type: none"><li>• NOT 50% (too concentrated and risk of extravasation injury)</li></ul></li></ol>
<b>Unconscious Seizures Aggressive</b>	<ol style="list-style-type: none"><li>1. Glucagon (1mg subcutaneously or intramuscularly)</li><li>2. 10% glucose (150-200ml) intravenously or 20% (75mL-100) iv delivering 15-20g<ul style="list-style-type: none"><li>• NOT 50% (too concentrated and risk of extravasation injury)</li></ul></li></ol>

### Case presentation

A 21-year old woman developed severe morning sickness requiring treatment in the first trimester of her first pregnancy.

**PMH.** None. **DH.** Folic acid 400 micrograms orally daily.

### On examination

Temperature 37°C, HR 88/min and regular, BP 124/74 mmHg, RR 14/min, O<sub>2</sub> sat 99% on air, HS normal, chest sounds clear.

### Question

Select the *most appropriate* management option at this stage.  
(mark it with a tick)

## MANAGEMENT OPTIONS

- |          |                           |
|----------|---------------------------|
| <b>A</b> | Dexamethasone 8 mg orally |
| <b>B</b> | Domperidone 10 mg orally  |
| <b>C</b> | Haloperidol 1.5 mg orally |
| <b>D</b> | Ondansetron 80 mg orally  |
| <b>E</b> | Promethazine 25 mg orally |

x

## Answer box

### Option A Justification

Dexamethasone is typically used for nausea/vomiting related to cancer. It is not typically used in pregnancy

### Option B Justification

Domperidone is not typically used in pregnancy

### Option C Justification

Haloperidol is used in nausea / vomiting relating to palliative care, but not pregnancy

### Option D Justification

Ondansetron is an option but wrong dose

### Option E Justification

Promethazine is a first-line treatment for nausea and vomiting in pregnancy.

# Nausea and labyrinth disorders

## Drug treatment

Scroll  
down



Antiemetics should be prescribed only when the cause of vomiting is known because otherwise they may delay diagnosis, particularly in children. Antiemetics are unnecessary and sometimes harmful when the cause can be treated, such as in diabetic ketoacidosis, or in [digoxin](#) or antiepileptic overdose.

If antiemetic drug treatment is indicated, the drug is chosen according to the aetiology of vomiting

**Antihistamines** are effective against nausea and vomiting resulting from many underlying conditions. ~~There is no evidence that any one antihistamine is superior to another but their duration of action and inci~~

Nausea and vomiting in the first trimester of pregnancy is common and will usually resolve spontaneously within 16 to 20 weeks. For women who have nausea and vomiting, offer appropriate self-care advice (such as rest, oral hydration and dietary changes), and inform them about other available support (e.g. self-help information and support groups) and when to seek urgent medical advice. Take into consideration that a number of interventions may have already been tried. Antiemetics should be considered for women with persistent symptoms where self-care measures have been ineffective. If a non-pharmacological option is preferred, ginger may be helpful for mild to moderate nausea.

For women who choose pharmacological treatment, offer an antiemetic considering the advantages and disadvantages of each option, as well as patient preference, and their experience with treatments in previous pregnancies. Although few drug options are specifically licensed for nausea and vomiting associated with pregnancy, their use is established practice. Antiemetic options include: [chlorpromazine hydrochloride](#), [cyclizine](#), [doxylamine with pyridoxine](#), [metoclopramide hydrochloride](#), [prochlorperazine](#), [promethazine hydrochloride](#), [promethazine teoclate](#), and [ondansetron](#). For further information on antiemetic options, see NICE guideline: **Antenatal care** (available at: [www.nice.org.uk/guidance/hg201](http://www.nice.org.uk/guidance/hg201)). Assess the response to treatment after 24 hours; if the response is inadequate, switch to an antiemetic from a different therapeutic class. Reassess after 24 hours and if symptoms have not settled, specialist opinion should be sought. For women who have moderate to severe nausea and vomiting, consider intravenous fluids and adjunctive treatment with acupuncture.

# Feedback Session Discussion Point

- If there are more than one plausible options, the way to differentiate between them could be:
  - A treatment summary which states which is first line (see previous slide in this case)
  - Other PMH (including pregnancy) which makes some options less appropriate
  - Other investigations (e.g. renal function) which makes some options (or the proposed dose) less appropriate

### Case presentation

A 56 year old woman attends the Emergency Department with increasing shortness of breath worsening over two days. She is wheezy, but has not had an increase in sputum production or fevers. **PMH.** Chronic obstructive pulmonary disease ( $FEV_1 < 50\%$  predicted). **DH.** Fostair 100/6 (100 micrograms beclomethasone dipropionate / 6 micrograms formoterol) one puff inhaled 12hrly, Spiriva (tiotropium) 18 micrograms inhaled once daily, salbutamol 200 micrograms inhaled as required.

### On examination

Temperature 36°C, HR 88/min and regular, BP 124/74 mmHg, RR 26/min, O<sub>2</sub> sat 85% on air, HS normal, chest sounds wheezy throughout.

### Investigations:

Arterial blood gases on air - pH 7.37 (7.35–7.45), pO<sub>2</sub> 6.8 kPa (11.3–12.6), pCO<sub>2</sub> 9.0 kPa (4.7–6.0), HCO<sub>3</sub><sup>-</sup> 36.2 mmol/L (21–29), base excess 3.2 mmol/L (±2).

CXR hyperinflated. No consolidation, effusions, oedema or pneumothorax

She is also being treated with nebulised bronchodilators and oral steroids.

### Question

Select the *most appropriate* oxygen therapy at this stage.  
(mark it with a tick)

### MANAGEMENT OPTIONS

A	15L/min oxygen via non-rebreath mask	
B	3L/minute oxygen via nasal cannulae	
C	28% oxygen via Venturi mask	x
D	Non-invasive ventilation, titrating oxygen to achieve oxygen saturations of 88-92%	
E	No supplementary oxygen	

# Oxygen therapy – a reminder of theory (may differ in clinical practice)

**Low-concentration (controlled) oxygen therapy** for patients at risk of hypercapnic respiratory failure, more likely in those with:

- chronic obstructive pulmonary disease;
- advanced cystic fibrosis;
- severe non-cystic fibrosis bronchiectasis;
- severe kyphoscoliosis or severe ankylosing spondylitis;
- severe lung scarring caused by tuberculosis;
- musculoskeletal disorders with respiratory weakness.

Initial oxygen should be given using a controlled concentration of 24 or 28%, titrated towards a target oxygen saturation of 88–92%.

- The controlled concentration is usually achieved with a venturi valve/mask
- Repeated arterial blood gasses may be required

**High concentration** oxygen therapy

- safe in uncomplicated cases of conditions such as pneumonia, pulmonary thromboembolism, pulmonary fibrosis, shock, severe trauma, sepsis, or anaphylaxis.

The screenshot shows a web page with a navigation bar at the top containing 'Home', 'Treatment summary', and 'Oxygen'. The 'Oxygen' link is highlighted with a red box. Below the navigation bar, the page title 'Oxygen' is displayed in a large, bold font. Underneath the title, the section 'Overview' is shown. The main content area contains several paragraphs of text. The first paragraph states that oxygen should be regarded as a drug and is prescribed for hypoxaemic patients to increase alveolar oxygen tension and decrease the work of breathing. The second paragraph explains that oxygen is the most common drug used in medical emergencies and should be prescribed initially to achieve a normal or near-normal oxygen saturation. The third paragraph discusses high concentration oxygen therapy, noting it is safe in uncomplicated cases of conditions such as pneumonia, pulmonary thromboembolism, pulmonary fibrosis, shock, severe trauma, sepsis, or anaphylaxis. The fourth paragraph mentions that in acute severe asthma, the arterial carbon dioxide ( $P_aCO_2$ ) is usually subnormal but as asthma deteriorates it may rise steeply. The fifth paragraph states that low concentration oxygen therapy (controlled oxygen therapy) is reserved for patients at risk of hypercapnic respiratory failure, which is more likely in those with: chronic obstructive pulmonary disease; advanced cystic fibrosis; severe non-cystic fibrosis bronchiectasis; severe kyphoscoliosis or severe ankylosing spondylitis; severe lung scarring caused by tuberculosis; musculoskeletal disorders with respiratory weakness, especially if on home ventilation; and an overdose of opioids, benzodiazepines, or other drugs causing respiratory depression. The final paragraph notes that until blood gases can be measured, initial oxygen should be given using a controlled concentration of 28% or less, titrated towards a target oxygen saturation of 88–92%.

Home Treatment summary > Oxygen

## Oxygen

### Overview

Oxygen should be regarded as a drug. It is prescribed for hypoxaemic patients to increase alveolar oxygen tension and decrease the work of breathing. The concentration of oxygen required depends on the condition being treated; the administration of an inappropriate concentration of oxygen can have serious or even fatal consequences.

Oxygen is probably the most common drug used in medical emergencies. It should be prescribed initially to achieve a normal or near-normal oxygen saturation; in most acutely ill patients with a normal or low arterial carbon dioxide ( $P_aCO_2$ ), oxygen saturation should be 94–98% oxygen saturation. However, in some clinical situations such as cardiac arrest and carbon monoxide poisoning it is more appropriate to aim for the highest possible oxygen saturation until the patient is stable. A lower target of 88–92% oxygen saturation is indicated for patients at risk of hypercapnic respiratory failure.

High concentration oxygen therapy is safe in uncomplicated cases of conditions such as pneumonia, pulmonary thromboembolism, pulmonary fibrosis, shock, severe trauma, sepsis, or anaphylaxis. In such conditions low arterial oxygen ( $P_aO_2$ ) is usually associated with low or normal arterial carbon dioxide ( $P_aCO_2$ ), and therefore there is little risk of hypoventilation and carbon dioxide retention.

In acute severe asthma, the arterial carbon dioxide ( $P_aCO_2$ ) is usually subnormal but as asthma deteriorates it may rise steeply (particularly in children). These patients usually require high concentrations of oxygen and if the arterial carbon dioxide ( $P_aCO_2$ ) remains high despite other treatment, intermittent positive-pressure ventilation needs to be considered urgently.

Low concentration oxygen therapy (controlled oxygen therapy) is reserved for patients at risk of hypercapnic respiratory failure, which is more likely in those with:

- chronic obstructive pulmonary disease;
- advanced cystic fibrosis;
- severe non-cystic fibrosis bronchiectasis;
- severe kyphoscoliosis or severe ankylosing spondylitis;
- severe lung scarring caused by tuberculosis;
- musculoskeletal disorders with respiratory weakness, especially if on home ventilation;
- an overdose of opioids, benzodiazepines, or other drugs causing respiratory depression.

Until blood gases can be measured, initial oxygen should be given using a controlled concentration of 28% or less, titrated towards a target oxygen saturation of 88–92%. The aim is to provide the patient with enough oxygen to achieve an acceptable arterial oxygen tension without worsening carbon dioxide retention and respiratory acidosis. Patients may carry an *oxygen alert card*.

### Case presentation

A 78 year-old woman with end-stage lung cancer has been an inpatient, and has her analgesia uptitrated. For the last three days her opiate analgesia has been:

- Regular morphine sulfate MR 60 mg orally twice daily, and
- As required Oramorph oral solution (morphine sulphate 10mg/5ml) 7.5 ml, 4 doses in total

It is decided to transfer her on to Durogesic Dtrans (fentanyl) patches.

### Question

Which one of the following dosage regimens would provide an equivalent dose for current opiate use?  
(mark it with a tick)

### MANAGEMENT OPTIONS

A	'100' fentanyl patch transdermally replace every 3 days	
B	'100' fentanyl patch transdermally replace every 5 days	
C	'50' fentanyl patch transdermally replace every 3 days	
D	'75' fentanyl patch transdermally replace every 3 days	X
E	'75' fentanyl patch transdermally replace every 5 days	

### Answer box

#### REGULAR dose

60mg x2/day = 120mg/24hr

#### AS REQUIRED dose

7.5mls of morphine solution 10mg/5ml = 15mg/dose

Four doses in 24hours = 60mg morphine/24hrs.

Therefore total 180 mg morphine /24 hours

"Prescribing in Palliative" care section of BNF shows approximations for interchanging morphine / fentanyl. Fentanyl '75' patch is most appropriate. Opioid Analgesic section of BNF shows fentanyl patches release 75microg/hour of fentanyl for 72hours. Hence new prescription is: one fentanyl '75' patch every 72 hours.

NOTE: different patches have different durations of action



# Converting opiate formulations - requested topic

- Please be familiar with “Prescribing in palliative care” section of your BNF
- Opiate equivalence is in the “Transdermal route” subsection
- Conversions typically happen when pain is controlled on stable doses
- Calculate the equivalent daily oral morphine dose
  - Regular **AND** “prn” medications
- Be wary of:
  - Concentration of oral morphine
  - Parenteral morphine requires further conversion
  - Other “non-morphine” preparations that requires further calculations
    - e.g. codeine, diamorphine, oxycodone, hydromorphone, tramadol
    - In “Equivalent doses of opioid analgesics” subsection

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## Prescribing in palliative care

### Overview

Palliative care is an approach that improves the quality of life of patients with threatening illness, through the prevention and relief of suffering by means of

**Scroll down**

**Buprenorphine patches are *approximately* equivalent to the following 24-hour doses of oral morphine**

morphine salt 12 mg daily	≡ buprenorphine '5' patch
morphine salt 24 mg daily	≡ buprenorphine '10' patch
morphine salt 36 mg daily	≡ buprenorphine '15' patch
morphine salt 48 mg daily	≡ buprenorphine '20' patch
morphine salt 84 mg daily	≡ buprenorphine '35' patch
morphine salt 126 mg daily	≡ buprenorphine '52.5' patch
morphine salt 168 mg daily	≡ buprenorphine '70' patch

Formulations of transdermal patches are available as 72-hourly, 96-hourly and 7-day patches, for further information see [buprenorphine](#). Conversion ratios vary and these figures are a guide only. Morphine equivalences for transdermal opioid preparations have been approximated to allow comparison with available preparations of oral morphine.

**72-hour Fentanyl patches are *approximately* equivalent to the following 24-hour doses of oral morphine**

morphine salt 30 mg daily	≡ fentanyl '12' patch
morphine salt 60 mg daily	≡ fentanyl '25' patch
morphine salt 120 mg daily	≡ fentanyl '50' patch
morphine salt 180 mg daily	≡ fentanyl '75' patch
morphine salt 240 mg daily	≡ fentanyl '100' patch

**Case presentation**

A 15 year old man is about to be discharged from hospital with a new diagnosis of Type I diabetes mellitus. He has been issued a prescription for long- and short-acting insulin.

**Question**

Select the *ONE most appropriate* information option that should be communicated to the patient.  
(mark it with a tick)

**INFORMATION OPTIONS**

A	He can miss his insulin dose once a week without consequence	<input type="checkbox"/>
B	He should ensure the timing of taking the short acting insulin should remain constant	<input type="checkbox"/>
C	He should seek medical advice if he is unable to drink fluids due to nausea and vomiting	<input checked="" type="checkbox"/>
D	He will need to have blood tests for HbA1c levels	<input type="checkbox"/>
E	He will need to monitor his capillary blood glucose six times daily	<input type="checkbox"/>

**Answer box****Option A** Justification

Adherence to an insulin regime is of the utmost importance, and young males, especially teenagers, find this difficult to adapt to at first.

**Option B** Justification

The timing of insulin should be adjusted according to meal times.

**Option C** Justification

If he is unable to eat/drink, he may require hospitalisation / intravenous fluids

**Option D** Justification

Although he will need HbA1c levels, this is not the most important information to be discussed

**Option E** Justification

Although he needs to monitor his capillary glucose, this is too frequent

# Sick day rules - Type I diabetes mellitus

- Never omit insulin (may need increased – local guidance usually provided)
- Maintain adequate (sugar-free) fluid intake
- Maintain regular carbohydrate intake – if unable to take solids, in liquid carbohydrate format
- Consider anti-emetic if nauseated
- Consider oral electrolyte replacement in diarrhoea
- If prolonged inability to keep down fluids (e.g. >4hrs), then likely needs hospital admission
- Increased blood glucose monitoring (e.g. 4hr-ly, and even more frequently if >moderate ketones)
- Ketone testing 2-4 hrly
  - If persistently elevated, or elevated while low blood glucose – may need hospital admission
- Diabetic specialist nurse should provide individualised plan

# Sick day rules - Type II diabetes mellitus

Medications	Advice
Patients on oral medication only	<ul style="list-style-type: none"><li>• Stop metformin, SGLT2i, GLP-1 analogues (and potentially some CV meds like ACE-I and diuretics) when unwell and restart when well</li></ul>
Patients taking <u>insulin</u> therapy	<ul style="list-style-type: none"><li>• Never omit insulin (if regular prescription)</li><li>• Emphasis on the importance of regular carbohydrate intake</li><li>• Minimum twice-daily self-blood glucose monitoring</li><li>• Seek advice if blood glucose persistently elevated (e.g. &gt; 17)</li><li>• Diabetic specialist nurse should provide individualised plan</li></ul>

**Case presentation**

An 64 year old lady has *C Difficile* colitis and has been started on oral vancomycin.

**Question**

Select the *most appropriate* information option that should be communicated to the patient.  
(mark it with a tick)

**INFORMATION OPTIONS**

A	If vancomycin fails to clear the infection, then she may require anti-TNF therapies	
B	Intravenous vancomycin is as effective as oral vancomycin for C Difficile colitis	
C	Oral vancomycin is given 4 times daily	x
D	She will be cleared of any skin colonisation of MRSA	
E	She will require regular blood tests to check liver function	

**Answer box****Option A** Justification

Immunosuppression would be likely life threatening in the case of severe C Difficile infection

**Option B** Justification

IV vancomycin would not be effective as would not reach bowel lumen.  
IV metronidazole is secreted into bowel which is why it is an option in combination with other therapies

**Option C** Justification

Correct

**Option D** Justification

Oral vancomycin does not clear MRSA colonization (IV vancomycin may be appropriate for MRSA associated infections)

**Option E** Justification

Medication works mainly in bowel, but manufacturer advises monitoring serum-vancomycin concentration in inflammatory intestinal disorders.

# Clostridium difficile

Higher risk of developing <i>C. difficile</i> colitis when also taking:	
Antibiotics	Clindamycin and second- and third-generation cephalosporins (especially in older people)
	Quinolones, carbapenems (for example, imipenem and meropenem)
	Prolonged courses of aminopenicillins (for example, ampicillin and amoxicillin).
Proton pump inhibitors	

# Clostridium difficile

		Treatment
First line (mild/moderate – first episode)		<u>Oral</u> vancomycin
Second line		<u>Oral</u> fidaxomicin
	(or with recurrence)	

- Note: Oral route is preferable for local site of action in GI tract
- Oral vancomycin not well absorbed, but manufacturer advises monitoring serum-vancomycin concentration in inflammatory intestinal disorders.
- Metronidazole iv (+oral vanc) reserved for life threatening infection under specialist supervision





**Case presentation**

A 54-year-old man with a new diagnosis of atrial fibrillation was assessed for his risk of thromboembolic disease, and the patient-doctor decision was to start warfarin.

**Question**

Select the *most appropriate* information option that should be communicated to the patient.  
(mark it with a tick)

**INFORMATION OPTIONS**

A	He must not drink alcohol on warfarin
B	He must not eat green leafy vegetables
C	He must use barrier contraception during sexual intercourse
D	If he has haematuria he should seek medical attention
E	If he requires antibiotics, he will need to stop warfarin

**Answer box****Option A** Justification

Should try to abstain but can drink alcohol as long as similar amounts most days as acute binges can be cause enzyme inhibition of the enzymes of warfarin metabolism

**Option B** Justification

Recommended to eat similar amounts per day as contains vitamin K

**Option C** Justification

Warfarin is teratogenic for women of childbearing age, but unlike methotrexate, men taking warfarin does not influence fetal development

**Option D** Justification

Unexpected bleeding must be investigated, at least with an INR check

**Option E** Justification

Antibiotics may be enzyme inducers/inhibitors of warfarin metabolism so will need careful INR monitoring, but not necessarily means stopping warfarin

# Warfarin COMS

Good info from: <https://www.nhs.uk/conditions/warfarin/> (not accessible during PSA)

- Missed doses
  - Take if on same day, but do not double dose the next day
- Seek medical attention if you:
  - Pass melaena, haematemesis, haemoptysis, haematuria, PR bleeding, increased or unexpected PV bleeding
  - Prolonged (>10 minutes) epistaxis, bleeding gums
  - Unusual headaches
  - Have a fall or an accident, or significant head injury
- Rashes and hair loss is common
- Be wary of drug-drug interactions (including OTC medications)
- Be wary of green leafy vegetables, vegetable oils and cereal grains which may contain large amounts of Vitamin K.
- Do not binge drink
- May need to be stopped prior to surgery or dental work
- Body piercings are not recommended (risk of bleeding / infection)
- Can have IM injection (e.g. vaccines) if INR is within range, but may also consider SC injection
- Avoid martial arts and kickboxing (and be wary of other contact sports)



### Case presentation

A 54-year old woman is seen in clinic for an autoimmune condition. The plan is for her to wean off the oral steroids slowly. She is currently prescribed prednisolone 30 mg orally daily. She is to continue on this dose for 5 days, and to step down by 5 mg every subsequent 5 days until she completely stops the prednisolone

Prednisolone is to be dispensed as 5mg tablets for her.

### Calculation

How many 5mg prednisolone tablets should be dispensed to allow her to finish her tapering course?  
(Write your answer in the box below)

Answer

105

tablets

### Answer box

#### Correct answer

105 tablets

#### Working

30mg = 6 tablets for 5 days → 30 tablets

25mg = 5 tablets for 5 days → 25 tablets

20mg = 4 tablets for 5 days → 20 tablets

15mg = 3 tablets for 5 days → 15 tablets

10mg = 2 tablets for 5 days → 10 tablets

5mg = 1 tablet for 5 days → 5 tablets

Therefore total number of tablets required =  
30+25+20+15+10+5 = 105 tablets



### Case presentation

A 28-year old woman is in Obstetrics High Dependency Unit with a diagnosis of pre-eclampsia. In addition to her other management, she has been prescribed intravenous labetalol infusion at a rate of 500 micrograms/minute.

Labetalol is available as 100mg/20mL.

### Calculation

What rate of labetalol infusion should be started?  
(Write your answer in the box below)

Answer

6

mL/hr

### Answer box

#### Correct answer

6mL/hr

#### Working

Labetalol concentration =  $100\text{mg}/20\text{ml} = 5\text{mg}/\text{ml}$

Dose = 500 micrograms/minute.  
=  $0.5\text{mg}/\text{minute}$   
=  $0.5 \times 60 \text{ mg}/\text{hour}$   
=  $30 \text{ mg}/\text{hr}$

Rate required = dose/concentration  
=  $30\text{mg}/\text{hr} / 5\text{mg}/\text{ml}$   
=  $6 \text{ ml}/\text{hr}$



### Case presentation

A 34 year old man is in the Emergency Department being treated for diabetic ketoacidosis. To prepare his fixed rate intravenous insulin infusion, a final concentration of 50 units soluble insulin in 50 mL 0.9% sodium chloride is required.

Soluble insulin is presented as a 10mL vial containing 100 units/mL.

### Calculation

What volume (mL) of stock soluble insulin (100 units/mL) should be used to make the infusion?

*(Write your answer in the box below)*

Answer

0.5

mL

### Answer box

#### Correct answer

0.5 mL

#### Working

100 units of insulin in 1mL

Therefore 50 units in 0.5mL.

The information regarding 50ml of 0.9% sodium chloride is not required.

Total volume of the vial (10ml) is also not required.



### Case presentation

A 6-month-old child (Body surface area =  $0.4 \text{ m}^2$ ) has been admitted to the paediatric ward. He requires  $500 \text{ mg/m}^2$  acyclovir intravenously 8hrly for 10 days.

Aciclovir is available in 250 mg vials, and must be used immediately after reconstitution. Any left over drug must be disposed and not kept for the next dose.

### Calculation

How many 250 mg vials of aciclovir will he require over the period of his treatment?

(Write your answer in the box below)

Answer

30

vials

### Answer box

#### Correct answer

30 vials

#### Working

Each dose =  $500 \text{ mg} \times 0.4 \text{ m}^2 = 200 \text{ mg}$

Each vial must be used immediately therefore need 1 vial for each dose

One vial, for three doses for 10 days

=  $1 \times 3 \times 10$  vials

= 30 vials

**Case presentation**

A 56-year-old man with Parkinson's Disease is having his medications uptitrated. His dose has been increased to 'Sinemet' 37.5/150 (37.5 mg carbidopa and 150 mg levodopa) one tablet orally three times a day

**Question**

Select the adverse effect that is *most likely* to be caused by this treatment.  
(mark it with a tick)

**MANAGEMENT OPTIONS**

- |          |              |
|----------|--------------|
| <b>A</b> | Dyskinesia   |
| <b>B</b> | Falls        |
| <b>C</b> | Fevers       |
| <b>D</b> | Palpitations |
| <b>E</b> | Weight gain  |

<input checked="" type="checkbox"/>
<input type="checkbox"/>
<input type="checkbox"/>
<input type="checkbox"/>
<input type="checkbox"/>

**Answer box**

**Option A** Justification

Dyskinesias (together with nausea) is one of the most common side effects

**Option B** Justification

While falls can occur (with multiple contributing factors), it is less common compared to dyskinesias

**Option C** Justification

Not a known side effect, and while abrupt discontinuation may result in neuroleptic malignant syndrome, this is uncommon

**Option D** Justification

Palpitations can occur, but not as common as dyskinesias

**Option E** Justification

Weight changes (both increase and loss) can occur, but not as common as dyskinesias



# Feedback Session Discussion Points

- The issue is that almost all the side effects listed in BNF is “Frequency not known”.
  - This is an ALPHABETICAL ORDERED list (i.e. not a list in priority/frequency order)
  - This group encompasses side effects that can range from “very common” to “very rare”.
  - This is typical for older drugs where reporting of side effects from clinicians are poor – so they do not have exact values to provide a frequency
- Be wary that many dopaminergic drugs for Parkinson’s Disease have the important (but less common) issues with impulse control disorders

### Case presentation

A 43-year-old woman develops muscle rigidity, high temperatures, delirium and sweating 2 weeks after being commenced on several medications following an in-patient psychiatric stay for depressive psychosis. **PMH.** Type II diabetes mellitus, gastroesophageal reflux disease, depression, chronic back pain. **DH.** See right

### Question

Select the medication that is *most likely* to have caused this adverse effect.  
(mark it with a tick)

### ADVERSE EFFECT OPTIONS

A	Amitriptiline 25 mg orally daily
B	Metformin 850 mg orally twice daily
C	Morphine MR 20 mg orally 12hrly
D	Olanzapine 10 mg orally daily
E	Ranitidine 150 mg orally 12hrly

<input type="checkbox"/>
<input type="checkbox"/>
<input type="checkbox"/>
<input checked="" type="checkbox"/>
<input type="checkbox"/>

### Answer box

Option A Justification

TCA's do not cause neuroleptic malignant syndrome

Option B Justification

No issues

Option C Justification

Morphine does not cause neuroleptic malignant syndrome

Option D Justification

She has neuroleptic malignant syndrome, very similar features to serotonin syndrom]. 4-14 days usually after exposure to anti-psychotics.

Option E Justification

No issues

# Neuroleptic malignant syndrome

- Mainly caused by antipsychotic drugs (“typical” higher risk than “atypical”)
- Features
  - Hyperthermia, fluctuating level of consciousness, muscle rigidity, and autonomic dysfunction with pallor, tachycardia, labile blood pressure, sweating, and urinary incontinence
  - Complications include rhabdomyolysis, acute renal failure, hyperkalaemia, seizures
- Management
  - Discontinuation of the antipsychotic drug is essential because there is no proven effective treatment
  - But bromocriptine and dantrolene have been used
- Easily confused with (apart from offending drug):
  - Serotonin syndrome
  - Malignant hyperthermia
  - Anticholinergic toxicity

**Case presentation**

An 18-year-old woman attends her General Practitioner. She reports that her depression persists despite non-pharmacological management. They jointly decide to start paroxetine 20 mg orally daily. **PMH.** Depression, irritable bowel syndrome, migraines and polycystic ovarian syndrome. **DH.** In addition to the paroxetine, her medications are listed (right).

**Question**

Select the prescription that is *most likely* to be interact with paroxetine to cause adverse effects.  
(mark it with a tick)

**PRESCRIPTION OPTIONS**

<b>A</b>	Cerazette [desogestrel] 75 micrograms daily	<input type="checkbox"/>
<b>B</b>	Mebeverine 150mg, 20min before meals	<input type="checkbox"/>
<b>C</b>	Metformin 500mg twice daily	<input type="checkbox"/>
<b>D</b>	Omeprazole 20mg daily	<input type="checkbox"/>
<b>E</b>	Sumatriptan 100 mg orally as required (maximum 300 mg total daily dose)	<input checked="" type="checkbox"/>

**Answer box**

Option A	Justification
There is no significant interaction with desogestrel	
Option B Justification	
Drug used for irritable bowel syndrome	
Option C Justification	
There is no significant interaction with metformin	
Option D Justification	
There is no significant interaction with omeprazole	
Option E Justification	
Sumatriptan has serotonin receptor agonist activity, and in combination with paroxetine increases the risk of serotonin syndrome	

# Serotonin syndrome (page 1/2)

- What is it?
  - Collection of symptoms, which varies and ranges in severity
    - Fever, agitation, **increased reflexes (and clonus)**, tremor, sweating, dilated pupils, and diarrhoea
    - Occasionally seizures and rhabdomyolysis
- When does it occur?
  - Typically due to the use of two or more serotonergic medications (or overdose of one serotonergic agent)
  - SSRIs (e.g. citalopram, fluoxetine, paroxetine), SNRIs (e.g. duloxetine), TCAs (e.g. amitriptyline), MAOi (e.g. selegiline), buspirone
  - Pethidine, tramadol, codeine, linezolid, triptans
  - Amphetamines (pharmaceutical or recreational), ecstasy/MDMA
  - Antiemetics affecting serotonin - metoclopramide, ondansetron
  - Components of certain over the counter drugs
    - Dextromethorphan, St. John's wort, L-tryptophan, 5-hydroxytryptophan

# Serotonin syndrome (page 2/2)

- How to treat?
  - Get advice from National Poisons Information Service
  - Stopping offending medications, supportive therapy, active cooling (not antipyretic drugs)
  - Possibly cyproheptadine (ITU setting)
  - +/- benzodiazepines
- Easily confused with (apart from offending drug):
  - Neuroleptic malignant syndrome
  - Malignant hyperthermia
  - Anticholinergic toxicity

**Case presentation**

A 30-year-old man attends a Gastroenterology clinic as follow up after a recent flare of his Crohn's disease. He has been taking prednisolone 10 mg orally daily, and it is expected that he will be continuing for at least 3 to 6 months. **PMH.** Crohn's disease. **DH.** In addition to the prednisolone, he is also taking azathioprine 125mg orally daily.

**Question**

Select the *most appropriate* option to prevent complications of the long-term prednisolone prescription.  
(mark it with a tick)

**MANAGEMENT OPTIONS**

A	Adcal-D3® (1.5 g calcium carbonate, 400 units colecalciferol) one tablet 12hrly	
B	Alendronic acid 70 mg orally once weekly	x
C	Lansoprazole 30 mg orally daily	
D	None required	
E	Sando-K® (12 mmol K <sup>+</sup> / 8 mmol Cl <sup>-</sup> ) three tablets 12hrly	

**Answer box****Option A** Justification

AdcalD3 is used for Vitamin D / Calcium deficiency (and not for bone prophylaxis in long-term steroid use)

**Option B** Justification

CORRECT - Taking long term steroids requires bisphosphonate therapy to reduce the risk of osteoporosis

**Option C** Justification

PPI for the prevention of gastritis for those with long-term therapy are used in selected cases (see next slide) that this man does not clearly fit. It might be considered, but it will not be as high priority as a bisphosphonate

**Option D** Justification

This man requires bone protection in context of his long-term steroids

**Option E** Justification

There is no evidence of hypokalaemia, and there is no need for routine potassium supplementation in long-term steroid therapy

# Long-term steroids – when to consider bisphosphonates / PPI?

Consider...	If...
Consider a bisphosphonate (alendronate or risedronate)	Taking high doses of oral corticosteroids ( $\geq$ prednisolone 7.5 mg daily for 3 months or longer)
Consider a proton pump inhibitor	At high risk of gastrointestinal bleeding or dyspepsia: <ul style="list-style-type: none"><li>• History of gastroduodenal ulcer, gastrointestinal bleeding, or gastroduodenal perforation.</li><li>• Older age.</li><li>• Use of drugs that are known to increase the risk of gastrointestinal bleeding, such as NSAIDs / anticoagulants</li><li>• Serious comorbidity, such as advanced cancer</li></ul>
Potassium supplementation <u>not routinely</u> prescribed (but U+Es may be monitored)	



**Case presentation**

A 86-year-old woman with asthma and atrial fibrillation was admitted to hospital with a fractured neck of femur. She was normally maintained successfully on digoxin 125 micrograms orally daily. This was stopped for 3 days peri-operatively and re-commenced at 8am today. **PMH.** Atrial fibrillation, severe asthma.

**DH.** Warfarin (target INR 2-3), Fostair® 200/6 (beclomethasone dipropionate 200 micrograms / formoterol 6 micrograms) two inhaled 12hly, tiotropium 5 micrograms inhaled daily, salbutamol inhaled as required.

**Examination**

Fully conscious and comfortable.

Pulse 122-136/minute, irregular

BP 152/74 mmHg.

**Investigations**

Plasma digoxin level 0.6 nmol/L (range 1-2) when tested 6h after dose.

**Question**

Based on the information provided, select the *most appropriate* decision option with regard to optimising her atrial fibrillation rate control  
(mark it with a tick)

**DECISION OPTIONS**

A	Digoxin loading dose 250 micrograms intravenously	<input checked="" type="checkbox"/>
B	Digoxin-Fab fragments intravenously	<input type="checkbox"/>
C	Increase daily digoxin dose to 250 micrograms orally daily	<input type="checkbox"/>
D	Reduce daily digoxin dose to 62.5 micrograms orally daily	<input type="checkbox"/>
E	Stop digoxin and start bisoprolol 5mg orally daily	<input type="checkbox"/>

**Answer box****Option A** Justification

This represents an appropriate loading dose to increase her serum level acutely whilst the oral dose is restarted & reaches steady state

**Option B** Justification

There are no symptoms or signs of digoxin toxicity.

**Option C** Justification

This represents a very large increase in usual daily dose which is inappropriate as the reason her level is low is due to her medication having been stopped for 3 days

**Option D** Justification

This represents an inappropriate decrease in dose which requires increasing as her plasma level is low

**Option E** Justification

She may not tolerate a  $\beta$ -blockade due to asthma, and it would be preferred to treat her atrial fibrillation with a drug that she is known to tolerate

# Feedback Session Discussion Point

- (Digoxin should not have been stopped for surgery)

she:

- Is not digitoxic (so answers B+D are wrong)
- Is currently not haemodynamically compromised (so she has time to get the dosing right)
- Is known to do well with digoxin, and there is potential contraindication to beta-blockers (so answer E is wrong)
- Was previously on the right dose prior to temporarily stopping digoxin (so increasing the dose in C is wrong)

**Case presentation**

A 24-year-old woman presents to the Emergency Department 4h post overdose with an unknown number of aspirin tablets. **PMH.** depression. She is asymptomatic **DH.** St John's Wort

**Investigations**

Plasma aspirin level:	632 mg/L	
Plasma potassium: 4.0 mmol/L	(3.8-5.1)	
Venous pH:	7.3	(7.35-7.45)

**Question**

Select the *most appropriate* decision option with regard to the management of her aspirin overdose  
(mark it with a tick)

**DECISION OPTIONS**

<b>A</b>	0.9% sodium chloride 1L over 4 hours
<b>B</b>	Activated charcoal 50 grams orally
<b>C</b>	Haemodialysis
<b>D</b>	Repeat pH monitoring in 4h
<b>E</b>	Sodium bicarbonate 225 mmol intravenously

✓

**Answer box****Option A** Justification

Supportive care will form part of managing aspirin overdose, but this treatment of intravenous fluids will not increase salicylate excretion

**Option B** Justification

May be useful if within <1h post overdose

**Option C** Justification

When levels >700mg/kg, may consider straight to dialysis (senior decision)

**Option D** Justification

There can be slow absorption of aspirin so more regular monitoring is used with clinical assessment to guide prognosis and treatment

**Option E** Justification

Used in the treatment of aspirin overdose (those which do not meet criteria for haemodialysis)

# Aspirin overdose

- Features
  - Initial respiratory alkalosis, followed by severe metabolic acidosis
  - Nausea, vomiting, hypoglycaemia, hyperpyrexia, non-cardiogenic pulmonary oedema, coma
- Management
  - Consider activated charcoal if
    - <1hr of ingestion,
    - Toxic dose (>125mg/kg),
    - and protected airway (see next slide)
  - If metabolic acidosis, give sodium bicarbonate (50-100 mmol); aim urine pH 7.5-8.5, blood pH<7.55
  - If aspirin levels >500mg/L, give intravenous sodium bicarbonate 225mmol
  - If aspirin levels >700mg/L or adverse features (coma, seizures, pulmonary oedema, renal failure), consider haemodialysis

# Feedback Session Discussion Point

- Is pH 7.30 severe acidosis and therefore haemodialysis is the right answer?
- There is no regularly agreed cut-offs, but most would say:
  - 7.25-7.35      Mild
  - 7.15-7.25      Moderate
  - <7.15          Severe
- So haemodialysis is not the right option for now

# Activated charcoal for selected poisoning cases

- Prevent absorption
  - Activated charcoal within 1h of ingestion
  - 50 mg orally in conscious patient
  - 50 mg by NG tube if obtunded (but needs protected airway)
  - Also caution in those with reduced gastrointestinal motility (risk of obstruction)
- NOT to be used with substances that do not bind to charcoal
  - Ethylene glycol, Iron, Lithium, Methanol, Strong acids and alkalis

# Haemodialysis for overdose for limited poisoning cases

- Haemodialysis can be consider for severe overdoses with drugs with low volume of distribution:
  - E.g. Aspirin, lithium, valproate, ethylene glycol, methanol and phenobarbital
- This is a specialist decision

NICE National Institute for Health and Care Excellence

Evidence search BNF BNF CKS Journals and databases

poisoning

Home > Search: poisoning

Showing 1-10 of 150 results for "poisoning"

**Poisoning, emergency treatment** Treatment summary

blockers poisoning Features of calcium-channel blocker poisoning include nausea, vomiting, dizziness, agitation, confusion, and coma in severe poisoning. Metabolic

Guidance on prescribing Medicines guidance

be given to the possibility of allergy, fire, explosion, radiation, or poisoning. Substances such as corticosteroids, some antimicrobials, phenothiazines

SODIUM NITRITE | Indications and dose Drug

Indications and dose Poisoning with cyanides (used in conjunction with sodium thiosulfate) By intravenous injection For Child Consult the National Poisons

NICE National Institute for Health and Care Excellence

Evidence search BNF BNF CKS Journals and databases

Search...

Home > Treatment summary > Poisoning, emergency treatment

**Poisoning, emergency treatment**

Overview

These notes provide only an overview of the treatment of poisoning, and it is strongly recommended that either TOXBASE or the UK National Poisons Information Service be consulted when there is doubt about the degree of risk or about management.

Hospital admission

Patients who have features of poisoning should generally be admitted to hospital. Patients who have taken poisons with delayed action should also be admitted, even if they appear well. Delayed-action poisons include:

- ACETYLCHOLINE
- ADRENALINE/EPINEPHRINE
- ASPIRIN
- ATROPINE SULFATE
- CALCIUM CHLORIDE
- CONATE
- FINE
- ACTIVATED
- LOPE
- DESFERRIOXAMINE MESILATE
- DIAZEPAM

Be familiar with the contents of this (very long) page!

Scroll down of Ctrl-F "Aspirin"

Aspirin poisoning

The main features of salicylate poisoning are hyperventilation, tinnitus, deafness, vasodilatation, and sweating. Coma is uncommon but indicates very severe poisoning. The associated acid-base disturbances are complex.

Treatment must be in hospital, where plasma salicylate, pH, and electrolytes can be measured; absorption of aspirin may be slow and the plasma-salicylate concentration may continue to rise for several hours, requiring repeated measurement. Plasma-salicylate concentration may not correlate with clinical severity in the young and the elderly, and clinical and biochemical assessment is necessary. Generally, the clinical severity of poisoning is less below a plasma-salicylate concentration of 500mg/litre (3.6 mmol/litre), unless there is evidence of metabolic acidosis. Activated charcoal can be given within 1 hour of ingesting more than 125 mg/kg of aspirin. Fluid losses should be replaced and intravenous sodium bicarbonate may be given (ensuring plasma-potassium concentration is within the reference range) to enhance urinary salicylate excretion (optimum urinary pH 7.5–8.5).

Plasma-potassium concentration should be corrected before giving sodium bicarbonate as hypokalaemia may complicate alkalinisation of the urine.

Haemodialysis is the treatment of choice for severe salicylate poisoning and should be considered when the plasma-salicylate concentration exceeds 700mg/litre (5.1 mmol/litre) or in the presence of severe metabolic acidosis.

Opioid poisoning



**Case presentation**

A 14-year-old girl is being treated with tobramycin 150mg intravenously 8-hrly for a pseudomonal chest infection. **PMH.** Cystic fibrosis, secondary diabetes. **DH.** Insulin, salbutamol, carbocysteine

**Investigations**

Serum tobramycin level:

Pre dose                    3 mg/L (range <2 mg/L)

Post dose                10mg/L (range 8-12 mg/L)

**Question**

Select the *most appropriate* decision option with regard to the management of her tobramycin prescription  
(*mark it with a tick*)

**DECISION OPTIONS**

<b>A</b>	Change to tobramycin 150mg inhaled 8-hrly
<b>B</b>	Change to tobramycin 100mg intravenously 8-hrly
<b>C</b>	Change to tobramycin 150mg intravenously 12-hrly
<b>D</b>	Change to tobramycin 200mg intravenously 12-hrly
<b>E</b>	No change to treatment

<input type="checkbox"/>
<input checked="" type="checkbox"/>
<input type="checkbox"/>
<input type="checkbox"/>

**Answer box**

Option A	Justification
Inhaled therapy is not appropriate in this setting	
Option B	Justification
Will reduce peak (post-dose) level required for antibacterial activity and potentially result in reduce antimicrobial activity	
Option C	Justification
Correct - Trough too high, need to increase length of time between doses to allow to drop down to below trough level required	
Option D	Justification
Increasing dose will potentially increase the post-dose (peak) levels to toxic (above range) levels. Additionally, there is an unpredictable effects on trough levels (pre-dose) when changing 2 parameters	
Option E	Justification
Pre dose too high: risk of toxicity	

# Antibiotics therapeutic drug monitoring

- For antibiotics that require therapeutic drug monitoring, BNF does provide the usual target levels
- This however should already be in the PSA question
- In “real-life” you should also consult local microbiology guidance

## In adults

Serum aminoglycoside concentrations **must** be determined in the elderly.

In patients with normal renal function, aminoglycoside concentrations should be measured after 3 or 4 doses of a multiple daily dose regimen and after a dose change.

For multiple daily dose regimens, blood samples should be taken approximately 1 hour after intramuscular or intravenous administration ('peak' concentration) and also just before the next dose ('trough' concentration). If the pre-dose ('trough') concentration is high, the interval between doses must be increased. If the post-dose ('peak') concentration is high, the dose must be decreased.

For once daily dose regimens, consult local guidelines on serum concentration monitoring.

## In children

In children with normal renal function, aminoglycoside concentrations should be measured after 3 or 4 doses of a multiple daily dose regimen.

Blood samples should be taken just before the next dose is administered ('trough' concentration). If the pre-dose ('trough') concentration is high, the interval between doses must be increased. For multiple daily dose regimens, blood samples should also be taken approximately 1 hour after intramuscular or intravenous administration ('peak' concentration). If the post-dose ('peak') concentration is high, the dose must be decreased.

# Feedback Session Discussion Point

- a short “syllabus” for DAT questions
  - Be familiar with how to work out your next management step based on results of these investigations
  - Be familiar with how to find guides in the BNF (if available)

Data interpretation – please know how to manage (learning points of your previous sessions):		
Paracetamol overdose	Know/find management for all different timepoints, and staggered	[Treatment summaries → Poisoning, emergency treatment]
Warfarin and INR	Know/find what to do with high INR <ul style="list-style-type: none"> <li>Need to know INR and severity of any bleeding</li> </ul>	[Treatment summaries → Oral anticoagulant]
Other kinds of anticoagulation monitoring	Know what monitoring options for other anticoagulants	Not easily found in BNF
Antibiotic sensitivity / MC+S result	Know how to work through <ul style="list-style-type: none"> <li>e.g. sensitivities, contraindications/allergies, severity, renal dysfunction with adjusted doses</li> </ul>	Not in BNF
Antibiotic therapeutic drug monitoring	Know how to interpret peak / trough levels	In the “monitoring requirements” for <u>some</u> antibiotics (e.g. gentamicin)
Adjusting insulin doses	Know how to adjust insulin doses based on blood glucose diary	Not in BNF
Statins and CK / LFTs / cholesterol	Know/find when to adjust/stop statins	In the “monitoring requirements” for <u>most</u> statins [ <u>BUT NOT</u> for cholesterol targets – that you will need to know]
Oxygen therapy	Know the different co-morbidities with different oxygen therapy strategies <ul style="list-style-type: none"> <li>Know the different targets for these comorbidities</li> <li>Know how to interpret arterial blood gases</li> </ul>	Not very useful, should know clinically [Treatment summaries → Oxygen]
Converting opiates	Know how to calculate <i>total daily</i> opiate requirements Know how to find equivalent doses	Needs practice!!! [Treatment summaries → Prescribing in Palliative Care]
TFT monitoring	Know how to interpret and adjust medications	Not in BNF
Lithium levels	Know what if toxic, or if to change doses	Monitoring not explained in BNF (you need to know) Treating lithium toxicity in [Treatment summaries → Poisoning, emergency treatment]
Adjusting dose for renal dysfunction	Be able to recognise when patient has renal dysfunction Be familiar with how to find dosing adjustments (if required)	Be familiar with how to change doses based on “Renal Impairment” information with each drug

# Monitoring requirements (and other tabs) are (sometimes) your friends

Home > Drugs > ISOTRETINOIN

## ISOTRETINOIN

Indications and dose	Important safety information
Contra-indications	Cautions
Interactions	Side-effects
Conception and contraception	Pregnancy
Breast feeding	Hepatic impairment
Renal impairment	Monitoring requirements
Prescribing and dispensing information	Patient and carer advice
Medicinal forms	

Indications and dose

## Important safety information

MHRA/CHM advice: Isotretinoin (Roaccutane®): rare reports of erectile dysfunction and decreased libido (October 2017)

### With oral use

An EU-wide review has concluded that on rare occasions, oral isotretinoin, indicated for severe acne, may cause sexual side-effects, including erectile dysfunction and decreased libido.

## Monitoring requirements

### Monitoring of patient parameters

#### With oral use

Measure hepatic function and serum lipids before treatment, 1 month after starting and then every 3 months (reduce dose or discontinue if transaminase or serum lipids persistently raised).

## Patient and carer advice

### With oral use

Warn patient to avoid wax epilation (risk of epidermal stripping), dermabrasion, and laser skin treatments (risk of scarring) during treatment and for at least 6 months after stopping; patient should avoid exposure to UV light (including sunlight) and use sunscreen and emollient (including lip balm) preparations from the start of treatment.

### With topical use

Patients should be warned that some redness and skin peeling can occur initially but settles with time. If undue irritation occurs, the frequency of application should be reduced or treatment suspended until the reaction subsides; if irritation persists, discontinue treatment. Several months of treatment may be needed to achieve an optimal response and the treatment should be continued until no new lesions develop.

If sun exposure is unavoidable, an appropriate sunscreen or protective clothing should be used.

### With oral use

Patients and carers should be told how to recognise signs and symptoms of psychiatric disorders such as depression, anxiety, and rarely suicidal thoughts.

**Case presentation**

A 71-year-old woman is admitted with severe community-acquired pneumonia. She has had a cough productive of thick green sputum for 2 days.

**Examination**

Temperature 38.4C, RR 24/min

Coarse crepitations at the right lung base

**Investigations**

Chest X-ray shows right lower lobe pneumonia.

Treatment with co-amoxiclav 1.2 g IV 8-hrly is initiated.

**Question**

Select the *most appropriate* monitoring option to assess the beneficial effects of this prescription in the *first three days of treatment*.  
(mark them with a tick)

**Case presentation**

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Select the *most appropriate* monitoring option to assess the **beneficial** effects of this prescription in the *first three days of treatment*.  
(mark them with a tick)

# Feedback Session Discussion Point

- BNF usually does not inform you how to monitor for beneficial effects



**Case presentation**

A 71-year-old woman is admitted with severe community-acquired pneumonia. She has had a cough productive of thick green sputum for 2 days.

**Examination**

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**Investigations**

Chest X-ray shows right lower lobe pneumonia.

Treatment with co-amoxiclav 1.2 g IV 8-hrly is initiated.

**Question**

Select the *most appropriate* monitoring option to assess the beneficial effects of this prescription in the *first three days of treatment*.  
(mark them with a tick)

**MONITORING OPTIONS**

A	Blood pressure	<input type="checkbox"/>
B	Chest X-ray appearance	<input type="checkbox"/>
C	Disappearance of basal crackles	<input type="checkbox"/>
D	Respiratory rate	<input checked="" type="checkbox"/>
E	Sputum colour	<input type="checkbox"/>

**Answer box**

Option A	Justification
Blood pressure may be low in event of severe sepsis, but in the short term a better reflection of fluid resuscitation (or inotropic support) rather than antibiotics	
Option B	Justification
The chest X-ray appearance is unlikely to resolve in the early stages of treatment	
Option C	Justification
The auscultatory finding of basal crackles at the base of the lung will take several days to resolve.	
Option D	Justification
Successful treatment of the pneumonia will improve gas exchange and any hypoxia and reduce the respiratory rate.	
Option E	Justification
Sputum colour is a poor guide to the success of treatment for pneumonia.	

**Case presentation**

A 25-year-old woman attends her diabetes review. She has been asymptomatic, and her capillary blood glucose diary was reviewed. **PMH.** Type I diabetes mellitus. **DH.** Insulin aspart 6 units with meals, insulin glargine 8 units at night.

Her night-time insulin glargine dose has been increased to 10 units.

**Question**

Select the *most important* monitoring option that is required for this change in dosing.  
(mark it with a tick)

**MONITORING OPTIONS**

A	HbA1c	
B	Pre-afternoon meal capillary blood glucose	
C	Pre-bedtime capillary blood glucose	
D	Pre-breakfast capillary blood glucose	x
E	Pre-evening meal capillary blood glucose	

**Answer box**

Option A Justification

Option B Justification

Option C Justification

Option D Justification

Option E Justification

**Case presentation**

A 10-year-old boy attends paediatric rheumatology clinic. In addition to his methotrexate and folic acid, he has been prescribed adjunctive treatment with oral corticosteroids (30 mg prednisolone orally daily).

**Question**

Select the *most appropriate* monitoring option that is required for his prednisolone  
(mark it with a tick)

**MONITORING OPTIONS**

- |   |               |
|---|---------------|
| A | Visual fields |
| B | Serum calcium |
| C | Growth chart  |
| D | cholesterol   |
| E | Serum sodium  |

✓

**Answer box**

Option A	Justification
Option B	Justification
Option C	Justification
To prevent unexpected growth retardation	
Option D	Justification
Option E	Justification

**Case presentation**

A 68-year old man attends the Emergency Department with shortness of breath and swelling of both legs of gradual onset for the last 4 days. **PMH.** Severe heart failure, myocardial infarction and coronary artery bypass grafting 5 years ago. **DH.** Aspirin 75mg orally daily, ramipril 10mg orally daily, bisoprolol 10mg orally daily, furosemide 40mg orally twice a day, spironolactone 25mg orally daily.

**Examination**

BP 136/88 mmHg, pulse 68/min regular  
ECG sinus rhythm, no ischaemia  
JVP raised at 4cm, peripheral oedema to knees  
Weight 86kg  
Urinary catheter output in last hour 80 ml

**Investigations**

Na<sup>+</sup> 139 mmol/L (137–144), K<sup>+</sup> 4.6 mmol/L (3.5–4.9), eGFR 78 mL/min (previously >90)  
CXR – pulmonary oedema

He has been given furosemide 80mg intravenously, and has been started on an intravenous infusion of GTN, initially at 1mg/hr.

**Question**

Select the *most appropriate* monitoring option to guide the rate of GTN infusion  
(mark it with a tick)

**MONITORING OPTIONS**

<b>A</b>	Blood pressure	<input checked="" type="checkbox"/>
<b>B</b>	eGFR	<input type="checkbox"/>
<b>C</b>	Serum potassium	<input type="checkbox"/>
<b>D</b>	Urine output	<input type="checkbox"/>
<b>E</b>	Weight	<input type="checkbox"/>

**Answer box**

Option A	Justification
	Blood pressure (and pulse) is an important guide to titrating IV GTN minute-to-minute/hour-to-hour
Option B	Justification
	eGFR will be important longer term, particularly with ongoing intravenous diuretic treatment
Option C	Justification
	Serum potassium will be important longer term, particularly with ongoing intravenous diuretic treatment
Option D	Justification
	Urine output will be important to the overall assessment of changes of fluid balance, but does not change immediate GTN dosing.
Option E	Justification
	Daily weights will be important to the overall assessment of changes of fluid balance day-to-day

# Resources summary

1. Online practice papers (<https://prescribing-safety-assessment.ac.uk>)
  2. All lectures, SPEs
  3. Workshops 1-4
  4. Fluids and calculations workbooks on QM+
- 
1. Email me queries as needed (v.kapil@qmul.ac.uk)