

Paracetamol poisoning

Version 96

Proforma to guide ED management of ORAL ingestions in adults**Includes overdoses due to therapeutic excess****Manage and document any co-ingestions separately**

Disclaimer:
This is a clinical template; clinicians should always use judgment when managing individual patients

June 2020

Patient details

Full name

DoB

Unit number

(use sticker if available)

① Sources of further advice

- Go to toxbase.org or toxbasebackup.org for online management advice regarding Paracetamol poisoning, including IV and other routes (see ED INsite guidance page 'Drugs & Fluids' for login details)
- National Poisons Information Service (NPIS)** is available anytime if remaining uncertainties after advice from ED senior
☎ **0344 892 0111**
- Liver unit** referrals should be made to the 'liver unit medical registrar' at the **Queen Elizabeth Hospital Birmingham** (see box 7 for criteria)
☎ **0121 627 2000**

② Significant ingestion?

Work out ingested dose in mg/kg

Total Dose	<input type="text"/>	mg	=	<input type="text"/>	mg/kg
Patient weight	<input type="text"/>	kg			

Disregard any additional kilos in excess of 110kg
If pregnant, enter pre-pregnancy not actual weight

Yes, as one of the below

Ingested dose >75 mg/kg/24h ☐
Reported dose unreliable ☐

No, as none of the above**③ NAC treatment needed?****YES, as one or more of the below**

4-15h after single ingestion, level on or above treatment line ☐
>15h after single ingestion, paracetamol is still detectable ☐
>4h after last tablets of a staggered ingestion taken, paracetamol is detectable ☐
>4h after an ingestion of uncertain timing, paracetamol is detectable ☐
INR >1.3 * ☐
ALT >53 IU/L * ☐

NO, as none of the above

* Call NPIS for advice (see box 1 for details) if INR or ALT are known to be chronically elevated due to other causes (e.g. warfarin or chronic liver disease) AND paracetamol cannot be detected or concentration is below the treatment line on the nomogram; patient **MAY** not need NAC

④ Single ingestion >24h ago

Obtain INR, venous gas, U&E, LFT, Paracetamol level and FBC

If jaundice or liver tenderness

→ Start NAC immediately (do not wait for blood results) and admit to AMU.
NB: check if referral to a liver unit is required (see box 6 for criteria).

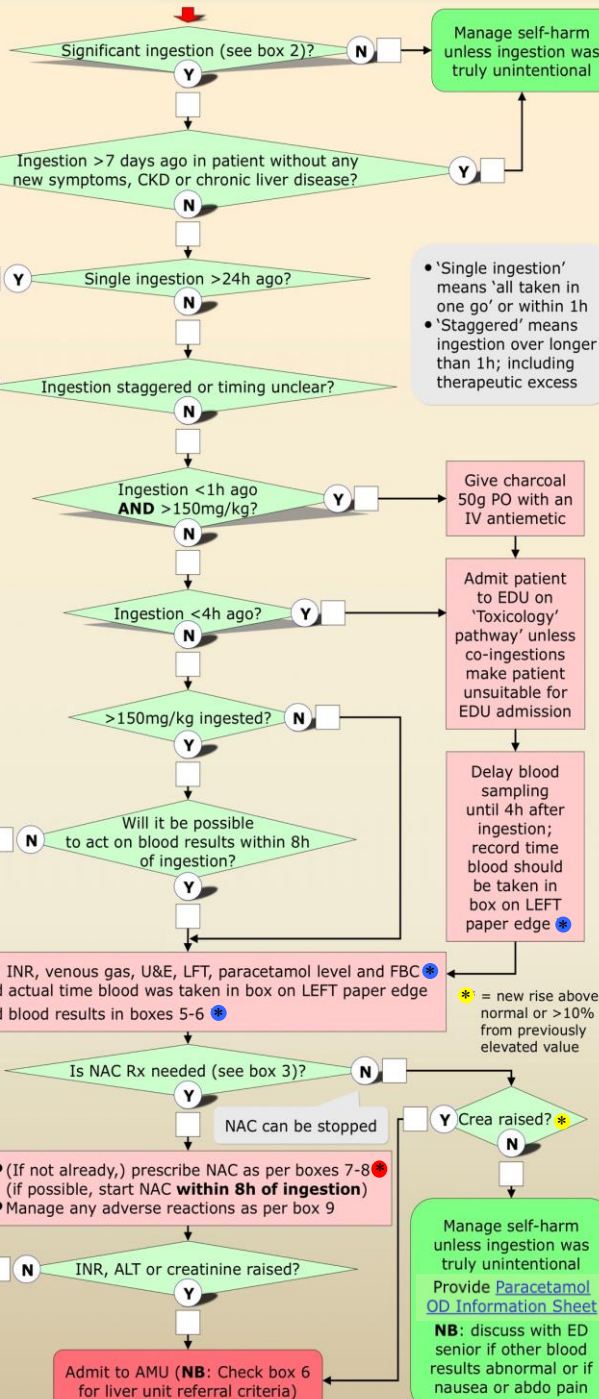
Otherwise await blood results

If NAC treatment needed (see box 3)

→ Start NAC and admit to AMU
NB: check if referral to a liver unit is required (see box 6 for criteria)

If serum creatinine is abnormal
(i.e. new rise above upper limit of normal or >10% from previous abnormal value)
→ Admit to AMU

If none of the above
Patient is not a risk of liver toxicity



Note times & tasks in the boxes below

DD/MM/YY	Current date
HH:MM	Current time
DD/MM/YY	Date of ingestion
Time of ingestion (24h clock)	
<input type="checkbox"/> Single ingestion; all tablets at	
<input type="checkbox"/> Staggered; last tablets taken at	
HH:MM	hours passed since
HH:MM	
<input type="checkbox"/> Timing unclear	
HH:MM	Sample needed at
Blood sampling delegated to	
HH:MM	Sample taken at
Result checking delegated to	
HH:MM	Start NAC before
NAC administration delegated to	
HH:MM	NAC started at

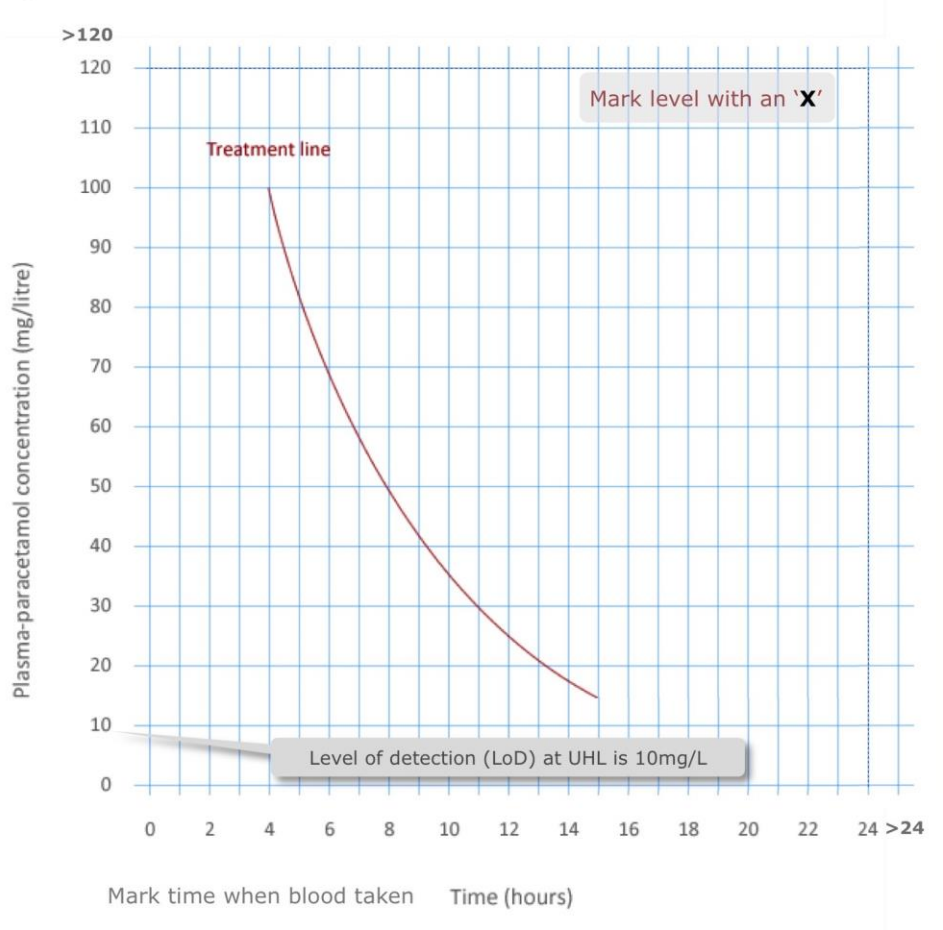
This patient was managed by

Print name

Signature

Role

⑤ Paracetamol blood level



- ### ⑦ NAC regimen
- N-Acetylcysteine (NAC) ampoules contain 2G NAC in 10mL (200mg/mL)
 - Regimen consists of 2 infusions given consecutively over 12h
 - Tick applicable weight range (in pregnancy, here: **ACTUAL** weight)
 - Prescribe NAC on fluid page of drug chart as per example in box 8

Patient weight (kg)	First infusion			Second infusion		
	Add required amount of NAC to a 200 mL bag of Glucose 5%			Add required amount of NAC to a 1000mL bag of Glucose 5%		
	NAC 100mg/kg		Rate	NAC 200mg/kg		Rate
	Dose	Volume		Dose	Volume	
	mg	mL	mL/h	mg	mL	mL/h
40-49	4600	23	112	9000	45	105
50-59	5600	28	114	11000	55	106
60-69	6600	33	117	13000	65	107
70-79	7600	38	119	15000	75	108
80-89	8600	43	122	17000	85	109
90-99	9600	48	124	19000	95	110
100-109	10600	53	127	21000	105	111
>109	11000	55	128	22000	110	111
Run time	2 hours			10 hours		

⑧ NAC example prescription for 62kg patient as per table in box 7

Date	Infusion fluid		Additions to infusion		IV or SC	Line	Start Time	Time to run or mL/hr	Fluid Batch No.	Prescriber
	Type/strength	Volume	Drug	Dose						
DD/MM/YY	Glucose 5%	200mL	N-Acetylcysteine	6600mg = 33mL	IV		HH:MM	117 mL/h (i.e. runs over 2h)		Dr.'s Name
DD/MM/YY	Glucose 5%	1000mL	N-Acetylcysteine	13000mg = 65 mL	IV		HH:MM	107 mL/h (i.e. runs over 10h)		Dr.'s Name

⑥ Blood results

initially | post-NAC

Time

liver unit referral criteria (NB: also include hepatic encephalopathy >grade II)

pH <7.3

pCO₂

Bicarb

Lactate >3.5*

Glucose

* >3 after fluid resuscitation/24h post-ingestion

Paracetamol

NB: Patients with paracetamol levels >700mg/L who are also in coma with a high lactate may require haemodialysis alongside NAC; d/w NPIS

Na

K

Urea

Crea >300

Bili

ALT

Alb

AP

WBC

Hb

Platelets

INR

Prothrombin time >20

⑨ NAC adverse reactions

NAC can cause nausea, vomiting, flushing, urticarial rash, angioedema, tachycardia, bronchospasm and, rarely, shock.

Reactions are more likely in women, asthmatics, those with a family history of allergies and patients with low paracetamol levels. They are usually seen during infusion of the 1st bag (large amounts being given rapidly).

Reactions can often be controlled by simply stopping the infusion temporarily; consider giving chlorphenamine 10mg IV if not. Add salbutamol 5mg neb if bronchospasm.

If unsuccessful use anaphylaxis pathway.

NB: Once reaction settled, restart the infusion at the normal rate.

Previous reaction is **NO** contraindication to NAC. If patient reports previous reactions consider pre-treatment with chlorphenamine 10mg and ranitidine 50mg IV. Pre-treat with salbutamol if history of bronchospasm.