

PATIENT GROUP DIRECTION EEAST 2022-25 Administration of Tranexamic Acid Injection

Auministration of Tranexamic Acid Injection					
Clinical Condition					
Indications	Patient with significant internal or external haemorrhage suspected				
Inclusion criteria	 Injured patients with traumatic injury where serious bleeding (significant internal/external haemorrhage) is suspected, within 3 hours of injury Patients aged over 18 years old with head injury and a GCS of 12 or less, within 3 hours of injury Post-Partum Haemorrhage within 3 hours of delivery or onset and bleeding is estimated to be >500mls and continuing Haemorrhage due to disorders of obstetric origin e.g, antepartum hours of placental placental obstetric origin e.g. 				
Exclusion criteria	haemorrhage, suspected placental abruption Critical intervention required i.e., if critical interventions leave insufficient time for administration of tranexamic acid.				
	Bleeding now stopped				
	Known allergy to tranexamic acid				
	Haemorrhage suspected of occurring more than three hours prior to treatment				
Cautions/Need for further advice	Rapid intravenous injection might rarely cause hypotension.				
	There is good data that this treatment is safe and effective (giving a 9% reduction in the number of deaths in patients in the CRASH2 trial).				
	Manufacturer's list of clinical precautions:				
	https://www.medicines.org.uk/emc/product/1220/smpc#CLINICAL PRECAUTIONS				
Pregnancy and breast feeding	No evidence of harm in pregnancy. However, manufacturer advises use only if potential benefit outweighs risk—crosses the placenta.				
	Manufacturer's information:				
	https://www.medicines.org.uk/emc/product/1220/smpc#PREGNANCY				
Action to be taken if patient is excluded	Ensure the medical details and all advice given and the actions of the patient (including guardian or relative) are handed over to the receiving facility.				
Action to be taken if notion t	Seek advice from a medical practitioner				
Action to be taken if patient declines treatment	Ensure medical records detail the advice given and the actions of the patient (including guardian or relative)				
	Seek advice from a medical practitioner if present				



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Drug Detail	S												
Name, form & strength of	Tranexamic Acid 500mg in 5ml ampoule (100 mg/ml).												
Legal Status	Prescription Only Medicine [POM]												
Licensed or Unlicensed	Licensed												
Route/Method	Intravenous or Intraosseous (off-licence use)												
	Must not be administered IM												
Dosage	Adults and children ≥12 years							Volume					
	Age		Initial Dose	Repeat Dose	Dose Interval			e Max Dose					
	≥12 years-A	dult	1gram	NONE	N/A	100mg/ml		10mls	1g				
	administration and monitor blood pressure carefully for possible hypotensive response giving appropriate fluid challenges where necessary to maintain perfusion pressure.												
	If required to aid slow IV administration draw up dose in a syringe with Sodium Chloride 0.9% to a total of 10ml. (e.g., 6-year-old, draw up 3ml of TXA, add 7ml of NaCl to make 10ml total volume)												
	Age	Initial Dose		Dose Interval	Concen	tration	Dose Vo	olume	Max Dose				
	11 years	500m	ng NONE	N/A	100mg/i	nl	5ml		500 mg				
	10 years	500m	ng NONE	N/A	100mg/i	ml	5ml		500 mg				
	9 years	450m	ng NONE	N/A	100mg/ml 100mg/ml 100mg/ml 100mg/ml 100mg/ml		4.5mls		450 mg				
	8 years	400m	ng NONE	N/A			4ml 4		400 mg				
	7 years	350m	ng NONE	N/A			3.5ml 3		350 mg				
	6 years	300m	ng NONE	N/A			3ml 3		300 mg				
	5 years	300m	ng NONE	N/A			3ml 3		300 mg				
	4 years	250m	ng NONE	N/A	100mg/i	nl	2.5ml		250 mg				
	3 years	200m	•	N/A	100mg/i		2ml		200 mg				
	2 years	200m	•	N/A	100mg/i		2ml		200 mg				
	18 months	150m	•	N/A	100mg/i		1.5ml		150 mg				
	12 months	150m	•	N/A	100mg/i		1.5ml		150 mg				
	9 months	150m	•	N/A	100mg/i		1.5ml		150 mg				
	6 months	100m	•	N/A	100mg/i		1ml		100 mg				
	3 months	100m	•	N/A	100mg/i		1ml		100 mg				
	1 month	50mg	-	N/A	100mg/i		0.5ml		50 mg				
	Birth	50mg) NONE	N/A	100mg/i	nl	0.5ml		50 mg				



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Drug Details	
Frequency	Single dose
Duration of treatment	Single dose
Maximum or minimum treatment period	Single episode
Quantity to administer	As above
Adverse effects	Rapid injection might rarely cause hypotension. Nausea, vomiting and diarrhoea
	Possible visual disturbances including visual impairment, vision blurred, impaired colour vision
	Manufacturer's listing:
	https://www.medicines.org.uk/emc/product/1220/smpc#UNDESIRABLE EFFECTS
	If any untoward reactions occur these must be reported using the Yellow Card system to the MHRA http://yellowcard.mhra.gov.uk/
	If such a report is made an internal Trust Incident Report form must be completed
Advice to patient/carer	The effects and side effects of the procedure must be explained to the patient and consent gained before administration.
Follow up	All patients who have received Tranexamic Acid must be transported to hospital and handed over to the care of a suitably qualified doctor or nurse.
Key References	BNF: JRCALC:
	https://bnf.nice.org.uk/drug/tranexamic-acid.html
	BNFc:
	https://bnfc.nice.org.uk/drug/tranexamic-acid.html
	eMC (Accord Healthcare):
	https://www.medicines.org.uk/emc/product/8953/smpc
	JRCALC:
	https://jrcalc-web.netlify.app/#/tab/dash/drug/D0360



Authorisation by EEAST		
Medical Advisor	Name: Dr Tom Davis Position: Medical Director GMC: 6077328 Signature: Date: 9 th June 2021	
Trust Pharmacist	Name: Dr Brian Wells Position: Pharmacist (Wells Offshore) GPhC: 2016621 Signature: BUCC Date: 19 th May 2021	