

# PROLONGED SEIZURE

**MODULE: NEUROLOGY**

**TARGET: ALL PAEDIATRIC TRAINEES; NURSING STAFF**

## **BACKGROUND:**

Prolonged seizure is the most common neurological medical emergency in children. It continues to be associated with significant morbidity and mortality.

The Royal College of Paediatrics and Child Health (RCPCH) has set standards for training; by the completion of level one training, all trainees are expected to be able to initiate therapy in a child presenting with prolonged seizure.

## INFORMATION FOR FACULTY

### LEARNING OBJECTIVES

At the end of the session, participants should:

1. Recognise role of intraosseus access when IV access unsuccessful
2. Be able to initiate and continue anticonvulsant treatment for acute status epilepticus
3. Understand principles of anticonvulsant treatment
4. Have knowledge of common causes of seizures in babies and children
5. Form differential diagnosis for status epilepticus
6. Refer to intensive care team whilst maintaining patient safety until they take over.
7. Engage in multidisciplinary team management

### SCENE SETTING

Location: Emergency Department

Expected duration of scenario: 15 mins      Expected duration of debriefing: 30 mins

### EQUIPMENT AND CONSUMABLES

Mannequin (baby)  
 Monitoring  
 O<sub>2</sub> facemask  
 Bag valve mask  
 Laryngoscope  
 Size 3.5, 4.0 ETT  
 IV cannula plus fixation stickers  
 EZ-IO device  
 0.9% saline  
 10 % dextrose  
 Simulated Drugs:  
   IV Lorazepam  
   Buccal Midazolam/PR Diazepam  
   PR Paraldehyde  
   IV Phenytoin  
   IV Phenobarbitone  
 Syringes (enteral, 5ml, 10ml, 50ml)  
 Drug chart  
 Obs chart  
 Blood gas results  
 SORT Emergency drug chart (if requested – see appendix)

### PERSONNEL-IN-SCENARIO

ST1-3 trainee *and/or*  
 ST4-6 trainee  
 ED or paediatric nurse  
 Mollie's mother

### ADDITIONAL INFORMATION

Differential diagnosis for floppy/seizing neonate:

- Hypoglycaemia
- Sepsis
- Non-accidental injury
- Electrolyte disturbance
- Metabolic syndrome

(NB In this scenario, unable to get initial IV access so intraosseus route should be used).

**PARTICIPANT BRIEFING**

2-month-old Mollie has been brought to the Emergency Department by her mum who is worried because Mollie has been difficulty to settle and not feeding properly.

The triage nurse has asked you to review Mollie urgently.

**FACULTY BRIEFING****'VOICE OF THE MANIKIN' BRIEFING**

2-month old baby; very quiet and drowsy.

**IN-SCENARIO PERSONNEL BRIEFING (MUM)**

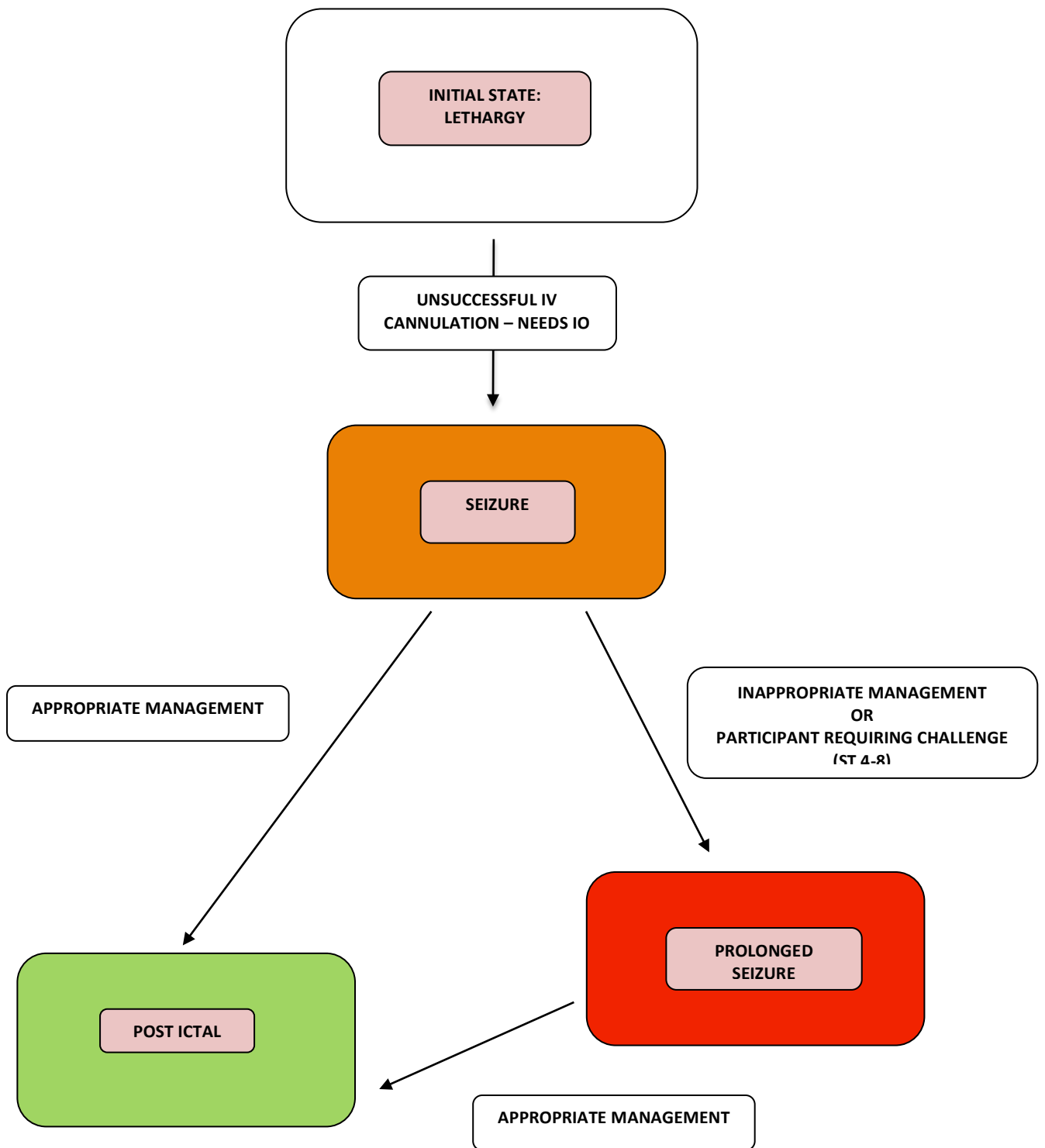
Mollie is your two-month old baby and you brought her to ED because you are worried she is getting floppier. She has been unsettled for the past day and has not been feeding well from her bottle.

If asked: Mollie was born at term by normal vaginal delivery and did not require special care after birth. Mollie has been well so far, with no medical concerns and no regular medications. She is due her first set of immunisations next week. You live with Mollie's father (your boyfriend) and Mollie. You have no other children.

**IN-SCENARIO PERSONNEL BRIEFING (NURSE)**

You have just triaged Mollie who was brought into ED via taxi with her mum. You are very concerned about her because she is very quiet and floppy with a bulging soft spot.

**CONDUCT OF SCENARIO**



**INITIAL STATE: LETHARGY**

VITAL SIGNS					
Rhythm	SR	HR	130	BP	68/34
Resp rate	30	SaO <sub>2</sub>	95%	ETCO <sub>2</sub>	
Temp	35.7	AVPU	V	Pupils	3 ERL
Other	Wt = 5kg	Mannequin dressed in vest and babygro			
ASSESSMENT					
Pulses	Normal	Cap refill	2-3 sec	Skin	No rashes
Airway	Maintained	Breathing	Normal	Breath sounds	Normal
Work of breathing	Normal	Recession	None	Neuro	Bulging fontanelle
Other	Baby quiet; hypotonic				
EXPECTED OUTCOMES					
<b>Participants should:</b>	<p>Obtain brief history. Key features:</p> <ul style="list-style-type: none"> <li>poor feeding</li> <li>difficult to settle</li> </ul> <p>Full examination, exposing skin. Key features:</p> <ul style="list-style-type: none"> <li>Very quiet</li> <li>Bulging fontanelle</li> <li>Hypotonic</li> </ul> <p>Apply monitoring</p> <ul style="list-style-type: none"> <li>Note capillary refill 2-3 seconds but HR not tachycardic for baby of this age</li> </ul> <p>Attempt IV access</p> <ul style="list-style-type: none"> <li>Unsuccessful – should move to IO access.</li> </ul> <p>Consider fluid bolus once IO/IV access obtained</p> <ul style="list-style-type: none"> <li>Note floppy child with prolonged capillary refill</li> </ul>				
<b>Facilitators should:</b>	After five minutes move to state <b>'Seizure'</b>				

## STATE: SEIZURE / PROLONGED SEIZURE

VITAL SIGNS					
Rhythm	SR	HR	170	BP	Unable to pick up (seizing)
Resp rate	30	SaO <sub>2</sub>	Poor signal	ETCO <sub>2</sub>	
Temp	35.7	AVPU	U	Pupils	3 ERL
Other	Mannequin in 'seizure' state				
ASSESSMENT					
Pulses	Normal	Cap refill	2-3 sec	Skin	No rashes
Airway	Maintained	Breathing	Erratic pattern	Breath sounds	Reduced
Work of breathing	Normal	Recession	None	Neuro	Unresponsive
Other	Seizure is generalised; not focal. Still has bulging fontanelle.				
EXPECTED OUTCOMES					
Participants should:	<ul style="list-style-type: none"> <li>• Gain IO access if not done already</li> <li>• Once blood gas results available, give bolus 10% dextrose</li> <li>• Give rectal diazepam 2.5mg or buccal midazolam if no IV/IO access</li> <li>• Give intravenous/IO lorazepam 0.1mg/kg</li> <li>• Give PR paraldehyde whilst IV phenytoin being prepared</li> <li>• Could also give 0.9% saline bolus via IO and reattempt IV access</li> </ul>				
Facilitators should:	<p>- If suboptimal management (e.g. hypoglycaemia not addressed), or if senior trainee requiring challenge, baby continues to have seizure.</p> <p>- If no progression after 10 minutes, use 'pause and perfect' principle to help participant understand current state and treatment options, before restarting scenario and allowing them to instigate appropriate management.</p> <p>- If appropriate management, seizure stops before phenytoin given. Progress to state '<b>Post-Ictal</b>'.</p>				

## STATE: POST-ICTAL

VITAL SIGNS					
Rhythm	SR	HR	130	BP	74/38
Resp rate	30	SaO <sub>2</sub>	95%	ETCO <sub>2</sub>	
Temp	36	AVPU	P	Pupils	3 ERL
Other	Seizure terminated – no abnormal movements				
ASSESSMENT					
Pulses	Normal	Cap refill	2-3 sec	Skin	No rashes
Airway	Maintained	Breathing	Normal	Breath sounds	Normal
Work of breathing	Normal	Recession	None	Neuro	Responsive to pain
Other	Seizure terminated. Baby now floppy. Cries in response to pain.				
EXPECTED OUTCOMES					
Participants should:	<ol style="list-style-type: none"> <li>Recognise post-ictal state</li> <li>Suggest bloods: <ul style="list-style-type: none"> <li>Ongoing BM/blood gas monitoring</li> <li>U+E, FBC, Coagulation screen,</li> <li>Serum amino acids, ammonia</li> <li>Blood cultures</li> </ul> </li> <li>Suggest urine tests: <ul style="list-style-type: none"> <li>MC+S</li> <li>Ketones</li> <li>Amino and organic acids</li> </ul> </li> <li>Suggest radiology: <ul style="list-style-type: none"> <li>Cranial USS</li> <li>CT head</li> <li>?Chest X-ray</li> <li>?Skeletal survey</li> </ul> </li> <li>Arrange transfer to ward/HDU</li> </ol>				
	<ul style="list-style-type: none"> <li>- If suboptimal management (e.g. hypoglycaemia not addressed), or if senior trainee requiring challenge, baby continues to have seizure.</li> <li>- If no progression after 10 minutes, use 'pause and perfect' principle to help participant understand current state and treatment options, before restarting scenario and allowing them to instigate appropriate management.</li> <li>- If appropriate management, seizure stops. Progress to state 'Post-ictal'.</li> </ul>				







**APPENDIX 3 – EMERGENCY DRUG CALCULATOR**

Date Dec 20, 2012

**Southampton  
Oxford  
Retrieval  
Team**

**DRUG CALCULATOR**

**WEIGHT** 5 Kg

*Enter weight and click calculate*

Calculate
Print

**Emergency**

Adrenaline 1:10,000	0.5 ml (0.1 ml/kg)
Atropine 600mcg/ml	0.17 ml (20mcg/kg, min 100mcg)
Atropine 100mcg/ml	1 ml (20mcg/kg min 100mcg)
Sodium Bicarbonate 8.4%	5 ml (1 ml/kg)
Calcium Gluconate 10%	2.5 ml (0.5 ml/kg)

**Respiratory**

Magnesium Sulphate	200 mg (40 mg/kg over 20 minutes)
Salbutamol load	75 mcg (15 mcg/kg over 10 minutes)
Hydrocortisone	20 mg (4 mg/kg, max 100mg)
Aminophylline load	25 mg (5 mg/kg over 20 minutes)
Adrenaline 1:1000 Nebulised	2.5 ml (0.5 ml/kg, max 5 mls) Make up to 5 ml with saline

**Cardiac**

Cardioversion (sync)	5 Joules (1J/kg) (use 2J/kg if fails)
Shockable rhythm (async)	20 Joules (4J/kg)
Adenosine	500 mcg (100 mcg/kg)
Amlodarone Load	25 mg (5 mg/kg over 30 minutes to 4hrs)

**Anaesthesia**

Ketamine	10 mg (2mg/kg)
Thiopentone	5 to 25 mg (1-5mg/kg)
Fentanyl	10 to 25 mcg (2-5mcg/kg)
Morphine	0.5 mg (0.1 mg/kg)
Rocuronium	5 mg (1mg/kg)
Atracurium	2.5 mg (0.5mg/kg)
Vecuronium	0.5 mg (0.1mg/kg)
Suxamethonium	7.5 mg (1.5mg/kg)

**Neuro**

Lorazepam	0.5 mg (0.1 mg/kg)
Midazolam Buccal	0.5 mg (0.1 mg/kg)
Phenytoin	100 mg (20 mg/kg over 20 minutes)
Phenobarbitone	100 mg (20 mg/kg)
Paraldehyde PR	2 ml (0.4 ml/kg, mix 1:1 with oil)
3% Saline	15 ml (3ml/kg)
Mannitol 10%	25 ml (5ml/kg, eqivalent to 0.5g/kg)

**Anaphylaxis**

Adrenaline IM	0.15 ml of 1:1000
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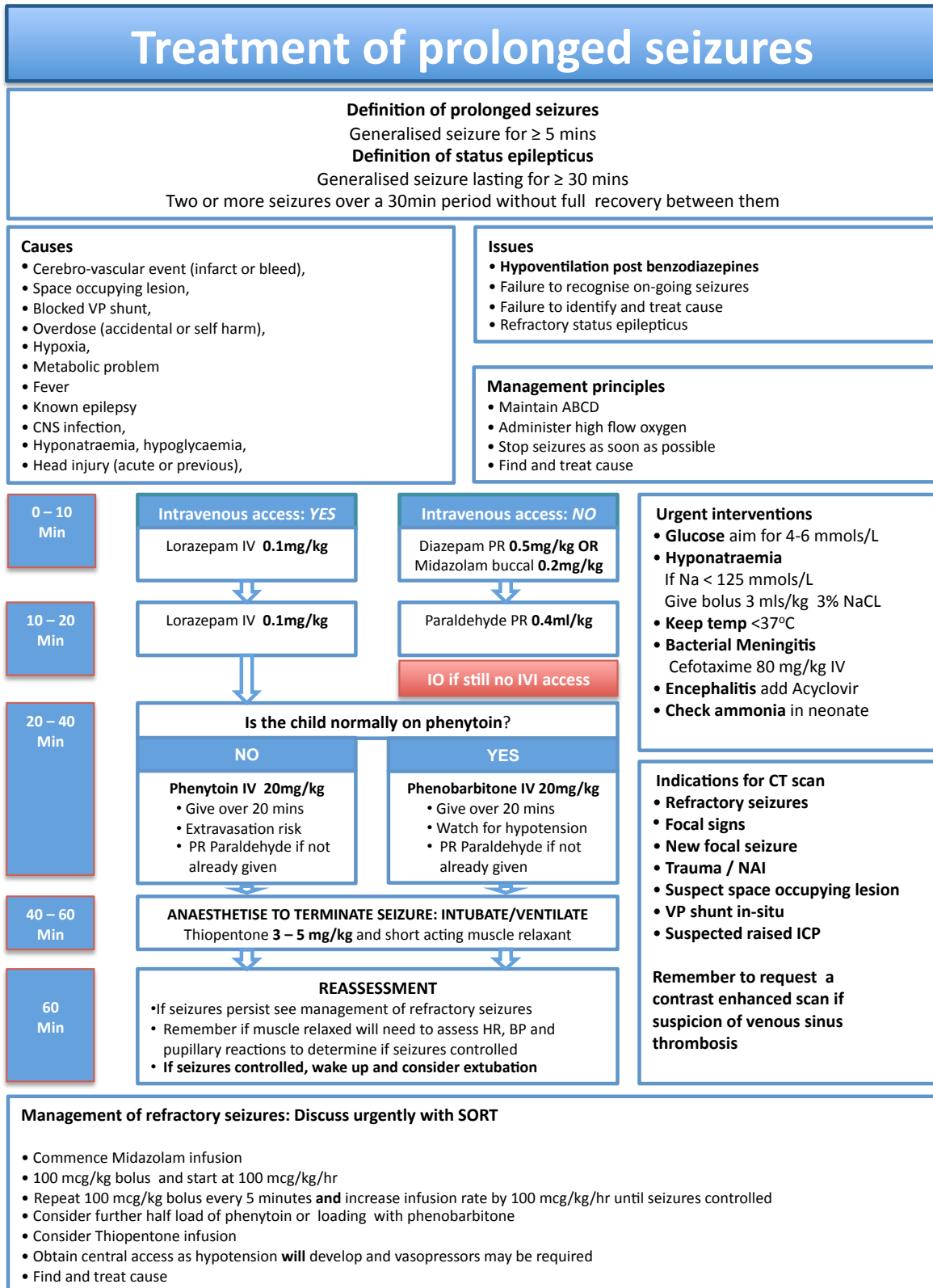
**Infusions**

Calculations based on Southampton PICU infusions guidelines (2011)

Dopamine (central)	75 mg In 50ml of 0.9% Saline or 5% Glucose	1 ml / hr =	5 mcg/kg/min
Dopamine (peripheral)	7.5 mg In 50ml of 0.9% Saline or 5% Glucose	1 ml / hr =	0.5 mcg/kg/min
Adrenaline	1.5 mg In 50ml of 0.9% Saline or 5% Glucose	1 ml / hr =	0.1 mcg/kg/min
Noradrenaline	1.5 mg In 50ml of 0.9% Saline or 5% Glucose	1 ml / hr =	0.1 mcg/kg/min
Milrinone	10 mg In 50ml of 0.9% Saline or 5% Glucose	0.75 ml / hr =	0.5 mcg/kg/min
Dinoprostone (Prostin E2)	50 mcg In 50ml of 0.9% Saline or 5% Glucose	1.5 ml / hr =	5 ng/kg/min
Morphine	5 mg In 50ml of 0.9% Saline or 5% Glucose	1 ml / hr =	20 mcg/kg/hr
Midazolam	5 mg In 50ml of 0.9% Saline or 5% Glucose	1 ml / hr =	20 mcg/kg/hr
Salbutamol	10 mg In 50ml of 0.9% Saline or 5% Glucose	1.5 ml / hr =	1 mcg/kg/min
Aminophylline	250 mg In 250ml of 0.9% Saline or 5% Glucose	5 ml / hr =	1 mg/kg/hr

**It is the prescribers responsibility to ensure the correct dose is prescribed**      Compiled by Tom Bennett - May 2012

**APPENDIX 4 – SORT GUIDELINE – PROLONGED SEIZURE**



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**APPENDIX 5 – INTRAOSSEOUS NEEDLE GUIDANCE (SORT)**

**Intraosseous needle (IO) observation chart and guidance**

**Patient Label**

Time/Date			
Sites inserted		Sites attempted	
R Tibial		R Tibial	
L Tibial		L Tibial	
Other		Other	


**Guidance**  
 There should only be one attempt at insertion in each bone  
 The IO needle should be secured without bandaging the limb so that the limb can be seen and monitored  
 One nurse must be solely responsible for monitoring the IO needle at all times

**Observation Chart – to be completed every 15 minutes**

Time							
Capillary refill time in distal limb (secs)							
Colour of limb *							
Visible swelling of limb Y/N **							
Limb feels firm Y/N **							

\*  
 Colour of limb  
 Pink = 1  
 Pale = 2  
 Blue/White = 3

\*\*  
 Always compare with other limb

 **Suspect extravasation if CRT > 4secs / colour is 2 or 3 / limb swelling / limb firmness**

**Actions if suspected infiltration**  
 Stop infusion (s) immediately  
 Remove IO needle  
 Call senior orthopaedic doctor to consider immediate fasciotomy  
 Orthopaedic registrar at Southampton or Oxford available for advice if none available locally

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## DEBRIEFING

### POINTS FOR FURTHER DISCUSSION

- Hypoglycaemia and another underlying cause for seizures may co-exist (e.g. NAI, meningitis) so treatment of hypoglycaemia may not be enough to stop seizure.
- Important to recognise Human Factors in high-stress, high-noise environment when a child begins to have seizure.
- Is everyone on team comfortable with giving PR diazepam, buccal midazolam, PR paraldehyde? Often the team member nearest the child (e.g. doctor) will be handed the syringe – important to know how to give it to prevent under-dosing.
- Important to remember benzodiazepines may have been given by parents or ambulance staff. If so, **do** count this dose in algorithm and progress onto paraldehyde/phenytoin after 2<sup>nd</sup> benzodiazepine (including pre-hospital dose(s)). Risk of significant respiratory depression if you do not take these pre-hospital doses into account.
- In baby having seizure with hypoglycaemia and difficult IV access, the priority should be to correct hypoglycaemia. Investigation of hypoglycaemia is less crucial at that stage, especially if obtaining blood samples is difficult.

### DEBRIEFING RESOURCES

SORT (Southampton Oxford Retrieval Service) Emergency Drug Calculator

<http://www.sort.nhs.uk/Media/Guidelines/Drugcalculator.pdf>

SORT guideline and obs chart for IO needles

[http://www.sort.nhs.uk/Media/Guidelines/Intraosseousneedle\(IO\)observationchartandguidance.pdf](http://www.sort.nhs.uk/Media/Guidelines/Intraosseousneedle(IO)observationchartandguidance.pdf)

**PROLONGED SEIZURE - HANDOUT****INFORMATION FOR PARTICIPANTS****KEY POINTS**

- Hypoglycaemia and another underlying cause for seizures may co-exist (e.g. NAI, meningitis) so treatment of hypoglycaemia may not be enough to stop seizure.
- Important to remember benzodiazepines may have been given by parents or ambulance staff. If so, **do** count this dose in algorithm and progress onto paraldehyde/phenytoin after 2<sup>nd</sup> benzodiazepine (including pre-hospital dose(s)). Risk of significant respiratory depression if you do not take these pre-hospital doses into account.
- Unlike with IV infusions, PR or buccal medications are often administered by the person closest to the patient – which may be you! Make sure you are happy with how to give buccal and PR medications safely and effectively.

**FURTHER RESOURCES**

SORT (Southampton Oxford Retrieval Service) Emergency Drug Calculator  
<http://www.sort.nhs.uk/Media/Guidelines/Drugcalculator.pdf>

SORT guideline and obs chart for IO needles  
[http://www.sort.nhs.uk/Media/Guidelines/Intraosseousneedle\(IO\)observationchartandguidance.pdf](http://www.sort.nhs.uk/Media/Guidelines/Intraosseousneedle(IO)observationchartandguidance.pdf)

## RELEVANT AREAS OF THE CURRICULUM

### Level One

L1_GEN_STA_02	Effective responses to challenge, complexity and stress in paediatrics
L1_GEN_STA_03	Advanced neonatal and paediatric life support skills
L1_GEN_STA_05	Effective skills in paediatric assessment
L1_GEN_STA_06	Skills in formulating an appropriate differential diagnosis in paediatrics
L1_GEN_STA_07	Effective initial management of ill-health and clinical conditions in paediatrics seeking additional advice and opinion as appropriate
L1_GEN_STA_09	Safe practical skills in paediatrics
L1_GEN_STA_15	Knowledge of common and serious paediatric conditions and their management
L1_GEN_STA_29	Effective communication and interpersonal skills with colleagues
L1_GEN_STA_30	Professional respect for the contribution of colleagues in a range of roles in paediatric practice
L1_GEN_STA_32	Effective handover, referral and discharge procedures in paediatrics
L1_GEN_STA_34	Ethical personal and professional practice in providing safe clinical care
L1_GEN_STA_35	Reliability and responsibility in ensuring their accessibility to colleagues and patients and their families
PAED_L1_ENDO_ACU_HYPO_01	Know the causes, complications and treatment of hypoglycaemia in the neonatal period and beyond
PAED_L1_ENDO_ACU_HYPO_02	Know that blood glucose is an urgent investigation in patients with impaired conscious level
PAED_L1_NEURO_GEN_10	Understand the life-threatening nature of acute neurological deterioration and when to call for help
PAED_L1_NEURO_GEN_11	Be able to recognise, initiate diagnostic tests and outline the management of common (neuro) disorders
PAED_L1_NEURO_SEIZ_01	Know the common causes of seizures in newborn babies and children
PAED_L1_NEURO_SEIZ_07	Be able to initiate treatment for acute continuing seizures
PAED_L1_NEURO_SEIZ_08	Be able to form a differential diagnosis in continuing seizures
PAED_L1_NEURO_SEIZ_09	Work effectively with the multidisciplinary team (in continuing seizures)

### Level Two (as above plus):

L2_GEN_STA_02	Increasing credibility and independence in response to challenge and stress in paediatrics
L2_GEN_STA_03	Leadership skills in advanced neonatal and paediatric life support
L2_GEN_STA_04	Responsibility for conducting effective paediatric assessments and interpreting findings appropriately

L2_GEN_STA_06	Improving skills in formulating an appropriate differential diagnosis in paediatrics
L2_GEN_STA_09	Effective skills in performing and supervising practical procedures in paediatrics ensuring patient safety
L2_GEN_STA_15	Extended knowledge of common and serious paediatric conditions and their management
L2_GEN_STA_29	Skill in ensuring effective relationships between colleagues
L2_GEN_STA_32	Effective skills in ensuring handover, referral and discharge procedures in paediatrics
L2_GEN_STA_34	Sound ethical, personal and professional practice in providing safe clinical care
L2_GEN_STA_35	Continued responsibility and accessibility to colleagues, patients and their families
PAED_L2_ENDO_ACU_HYPO_01	Know when to consider rare causes of hypoglycaemia and what investigations to perform during the hypoglycaemic episode
PAED_L2_ENDO_ACU_HYPO_02	Be able to treat hypoglycaemia safely and effectively with intravenous glucose or glucagon where appropriate
PAED_L2_NEURO_SEIZ_04	Be able to refer to intensive care teams appropriately and maintain patient safety until that team takes over (acute continuing seizures)
PAED_L2_NEURO_SEIZ_05	Be able to decide initial and continuing anticonvulsant therapy in babies and children

### Level Three (as above plus):

L3_GEN_STA_02	Responsibility for an effective response to complex challenges and stress in paediatrics
L2_GEN_STA_03	Leadership skills in advanced neonatal and paediatric life support
L3_GEN_STA_06	Effective skills in making safe decisions about the most likely diagnoses in paediatrics
L3_GEN_STA_07	Leadership skills in the management of common and complex conditions in general paediatrics and paediatric subspecialties seeking additional advice and opinion as appropriate
L3_GEN_STA_09	Expertise in a range of practical procedures in paediatrics specific to general and sub-specialist training
L3_GEN_STA_15	Detailed knowledge of common and serious paediatric conditions and their management in General Paediatrics or in a paediatric sub-specialty
L3_GEN_STA_29	Positive and constructive relationships form a wide range of professional contexts
L3_GEN_STA_32	Effective leadership skills in the organisation of paediatric team-working and effective handover
L3_GEN_STA_34	Exemplary professional conduct so as to act as a role model to others in providing safe clinical care
L3_GEN_STA_35	Responsibility for ensuring their own reliability and accessibility and that of others in their team
PAED_L3_NEURO_SEIZ_01	Work effectively with the multidisciplinary team and lead the care maintaining patient safety until that team takes over (acute continuing seizures)



# Treatment of prolonged seizures

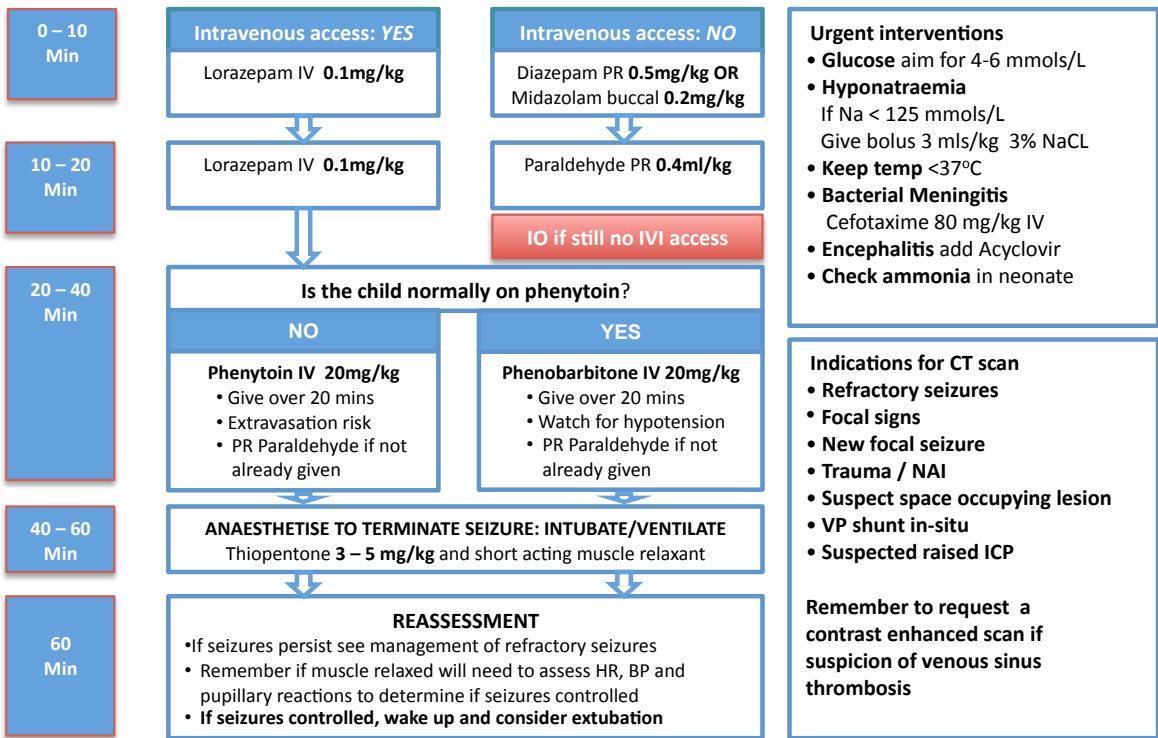
**Definition of prolonged seizures**  
Generalised seizure for  $\geq 5$  mins

**Definition of status epilepticus**  
Generalised seizure lasting for  $\geq 30$  mins  
Two or more seizures over a 30min period without full recovery between them

- Causes**
- Cerebro-vascular event (infarct or bleed),
  - Space occupying lesion,
  - Blocked VP shunt,
  - Overdose (accidental or self harm),
  - Hypoxia,
  - Metabolic problem
  - Fever
  - Known epilepsy
  - CNS infection,
  - Hyponatraemia, hypoglycaemia,
  - Head injury (acute or previous),

- Issues**
- Hypoventilation post benzodiazepines
  - Failure to recognise on-going seizures
  - Failure to identify and treat cause
  - Refractory status epilepticus

- Management principles**
- Maintain ABCD
  - Administer high flow oxygen
  - Stop seizures as soon as possible
  - Find and treat cause



- Management of refractory seizures: Discuss urgently with SORT**
- Commence Midazolam infusion
  - 100 mcg/kg bolus and start at 100 mcg/kg/hr
  - Repeat 100 mcg/kg bolus every 5 minutes **and** increase infusion rate by 100 mcg/kg/hr until seizures controlled
  - Consider further half load of phenytoin or loading with phenobarbitone
  - Consider Thiopentone infusion
  - Obtain central access as hypotension **will** develop and vasopressors may be required
  - Find and treat cause

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## PARTICIPANT REFLECTION

What have you learned from this experience? (Please try and list 3 things)

How will your practice now change?

What other actions will you now take to meet any identified learning needs?

**PARTICIPANT FEEDBACK**

Date of training session:.....  
 ...

Profession and grade:.....  
 .....

What role(s) did you play in the scenario? (Please tick)

Primary/Initial Participant	<input type="checkbox"/>
Secondary Participant (e.g. 'Call for Help' responder)	<input type="checkbox"/>
Other health care professional (e.g. nurse/ODP)	<input type="checkbox"/>
Other role (please specify): ..... .....	<input type="checkbox"/>
Observer	<input type="checkbox"/>

	Strongly Agree	Agree	Neither agree nor disagree	Disagree	Strongly Disagree
I found this scenario useful					
I understand more about the scenario subject					
I have more confidence to deal with this scenario					
The material covered was relevant to me					



Please write down one thing you have learned today, and that you will use in your clinical practice.

How could this scenario be improved for future participants? This is especially important if you have ticked anything in the disagree/strongly disagree box.

**FACULTY DEBRIEF – TO BE COMPLETED BY FACULTY TEAM**

What went particularly well during this scenario?

What did not go well, or as well as planned?

Why didn't it go well?

How could the scenario be improved for future participants?