

### Generalized Convulsive **Status Epilepticus (GCSE)**

Emergency Department (ED)

### **Center for Clinical Excellence**

Generalized **Convulsive Status** 

**Epilepticus:** 

Bilateral tonic-clonic

(BTC) seizures lasting ≥ 5 minutes

≥ 2 discrete BTC

seizures between

which there is

incomplete recovery of

consciousness

Prepare next medication

while waiting for current

medication to take effect

#### **Inclusion Criteria:**

Patients ≥ 28 days in generalized convulsive status epilepticus (GCSE)

#### **Exclusion Criteria:**

- Focal status epilepticus
- Psychogenic nonepileptic events (PNEE)
- Presenting history incudes trauma prior to seizure activity (follow trauma activation protocols)
- Hyponatremic etiology for seizures
- Known toxicological ingestion

Patient on Chronic **Antiseizure Medications** 

**GCSE versus PNEE** 

**Neuroimaging** 

**Concern for CNS** Infection

#### Initial management

- Cardiorespiratory monitoring
- Suction secretions
- Supplemental oxygen for O2 sats < 90%
- Place Intravenous (IV) line

#### Labs

- Point of care Glucose
- Consider blood gas if clinical concern for electrolyte disturbance
- Complete Metabolic Panel
- Complete Blood Count
- Serum pregnancy for females ≥ 10 years
- Urine drugs of abuse screen, if indicated
- Anticonvulsant levels, if indicated

Do not delay medication administration in order to obtain

labs

Lorazepam IV 0.1 mg/kg (max 4 mg/dose) over 1 minute

- Midazolam IN 0.3 mg/kg (max 10 mg/dose), split between nostrils
- If glucose < 60 mg/dL give 5 mL/kg D10% IV bolus (max 250 mL)

Seizure continues for 5 minutes

#### Repeat

Lorazepam IV 0.1 mg/kg (max 4 mg/dose) over 2 minutes

Midazolam IN 0.3 mg/kg (max 10 mg/dose), split between nostrils

Seizure continues for 5 minutes

Levetiracetam IV

60 mg/kg (max 4500 mg/dose) infuse over 5 minutes

Seizure continues for 10 minutes

#### Fosphenytoin IV

20 mg PE/kg (max 1500 mg PE/dose) infuse over 10 minutes

If patient has known allergy to fosphenytoin/phenytoin, consider Phenobarbital or Valproic Acid

Seizure continues for 10 minutes

Additional medical management in consultation with Neurology

Consult neurology if GCSE not responsive to Fosphenytoin or for any other concerns

> **Admission criteria** Neurology (HP6)

versus **PICU** 

**Discharge Criteria** Planning

CPP-ED Generalized Convulsive Status Epilepticus Clinical Pathway Published: 6/19/2023; Revised: 6/19/2023

# Generalized Convulsive Status Epilepticus (GCSE) versus Psychogenic Non-Epileptic Events (PNEE)

Distinguishing GCSE and PNEE can be challenging, and it is important to recognize that exceptions can occur.

Below is generalized guidance to distinguish GCSE and PNEE

#### Signs Favoring PNEE **Indeterminate Signs** Signs Favoring Epileptic Seizures Long duration **Gradual Onset** Fluctuating Course Occurrence from Non-Stereotyped Physiologic Sleep Events Asynchronous Postictal Confusion Flailing or Thrashing Movements\* Movements Stertorous Breathing Pelvic Thrusting\* Side-to-side Head or Opisthotonos Body Movements\*\* **Tongue Biting** Forced Eye Closure **Urinary Incontinence Ictal Crying** Memory Recall

\*May not reliably differentiate between PNEE and frontal lobe partial epileptic seizures

\*\* May only be helpful in distinguishing convulsive PNEE and epileptic seizures

#### *Adapted from:*

Avbersek A, Sisodiya S. Does the primary literature provide support for clinical signs used to distinguish psychogenic nonepileptic seizures from epileptic seizures? J Neurol Neurosurg Psychiatry. 2010; 81:719–725.

Perez DL, LaFrance WC. Nonepileptic seizures: an updated review. CNS Spectr. 2016 June;21(3:) 239-246.

## Patient on Chronic Antiseizure Medications

### Patient on chronic antiseizure medications:

- Follow GCSE pathway
- Consult neurology
- Check medication level if on any of the following as results may impact care in the acute setting:
  - o Carbamazepine
  - Phenobarbital
  - o Phenytoin
  - o Valproic acid

May consider checking levels for other anti-seizure medications – discuss with neurology

### **Neuroimaging**

### Consider neuroimaging when clinically stable if any of the following:

- Acute focal deficit (see Sudden Neurologic Deficit Pathway)
- Risk for bleeding
- Presence of VP shunt

Fast MRI preferred

### **Concern for CNS Infection**

### **Complete:**

- CSF studies
- Other tests as indicated
- Antibiotics/Acyclovir per Febrile Infant 29-60-day order set

**Do not delay** antibiotic or acyclovir to wait for completion of lumbar puncture

### **Medications**

- Glucose < 60 mg/dL: 5 ml/kg D10% IV bolus (max dose 250 mL).</li>
- Lorazepam IV: 0.1 mg/kg (max 4 mg/dose) over 2 minutes.
- Midazolam IN: 0.3 mg/kg (max 10 mg/dose). Split dose between nostrils.
- Levetiracetam IV loading dose: 60 mg/kg (max 4500 mg/dose) infuse over 10 minutes.
- Fosphenytoin IV loading dose: 20 mg PE/kg (max 1500 mg PE/dose) infuse over 10 minutes.
- Known allergy to fosphenytoin/phenytoin, consider:
  - Phenobarbital IV loading dose: 20 mg/kg (max 1000 mg/dose) infuse over 15-20 minutes (monitor for respiratory depression and potential need for intubation).

or

- Valproic acid IV loading dose: 30 mg/kg (maximum 3000 mg/dose) infuse over 10 minutes.
  - Contraindications:
    - < 2 years old</p>
    - Known mitochondrial disorder
  - Relative Contraindications:
    - Pregnancy
    - Hepatic dysfunction

### **Admission Criteria**

### Neurology (HP6)

- Status epilepticus resolved
- No airway compromise
- Improving mental status

### **Pediatric Intensive Care Unit**

- Concern for persistent GCSE
- Concern for subclinical status epilepticus
- Requiring continuous IV medications for seizure control
- Airway compromise (requiring nasal trumpet, BiPAP, endotracheal intubation)

### Discharge Criteria & Planning

### Discharge home from the Emergency Department may be considered if all of the following criteria are met:

- GCSE that resolves spontaneously or after 1 dose of benzodiazepine
- Patient mental status has returned to baseline
- Family comfortable with discharge plan and follow-up

### If criteria met:

- Provide prescription for seizure rescue medication
- Follow Up:
  - PCP within 2-3 days if Neurology follow-up is not already established and / or
  - o Neurology follow-up as recommended

### References

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Origination Date: *June, 2023* Last Revision Date: *June, 2026* 

### **Clinical Pathway Development**

This clinical pathway was developed using the process described in the NCH Clinical Pathway Development Manual Version 6, 2022. Clinical Pathways at Nationwide Children's Hospital (NCH) are standards which provide general guidance to clinicians. Patient choice, clinician judgment, and other relevant factors in diagnosing and treating patients remain central to the selection of diagnostic tests and therapy. The ordering provider assumes all risks associates with care decisions. NCH assumes no responsibility for any adverse consequences, errors, or omissions that may arise from the use or reliance on these guidelines. NCH's clinical pathways are reviewed periodically for consistency with new evidence; however, new developments may not be represented, and NCH makes no guarantees, representations, or warranties with respect to the information provided in this clinical pathway.

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### **Metrics**

### **Pathway Goal**

Timely treatment and evaluation of Generalized Convulsive Status Epilepticus (GCSE)

### **Quality Measures**

### **Outcome Metrics**

- Time to first antiepileptic drug administration
- Time to IV access

#### **Process Metrics**

- Pathway Visualization
- Pathway Order Set Utilization