

Seizure

SESLHDPR/497

<p>Aim:</p> <ul style="list-style-type: none"> • Early identification and treatment of a patient following a seizure. • Early initiation of treatment / clinical care and symptom management within benchmark time. 																			
<p>Assessment Criteria: On assessment the patient may have one or more of the following presenting symptoms:</p> <table border="0"> <tr> <td>☒ Tonic - clonic seizure activity</td> <td>☒ Partial / Focal seizure (limited to one side of the body or extremity)</td> <td>☒ Alteration to mental status</td> </tr> <tr> <td>☒ Absent seizure activity</td> <td>☒ Atonic seizure activity</td> <td>☒ Tonic seizure activity</td> </tr> <tr> <td></td> <td></td> <td>☒ Persistent eye deviation</td> </tr> </table>		☒ Tonic - clonic seizure activity	☒ Partial / Focal seizure (limited to one side of the body or extremity)	☒ Alteration to mental status	☒ Absent seizure activity	☒ Atonic seizure activity	☒ Tonic seizure activity			☒ Persistent eye deviation									
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<p>Escalation Criteria: Immediate life-threatening presentations that require escalation and referral to a Senior Medical Officer (SMO):</p> <table border="0"> <tr> <td>☒ Status Epileptics (> 30 min continuous seizure activity)</td> <td>☒ Hypotension</td> <td>☒ Pregnancy</td> </tr> <tr> <td>☒ Traumatic head injury / fall</td> <td>☒ History of brain cancer</td> <td>☒ History of drug and alcohol abuse / overdose</td> </tr> <tr> <td>☒ Suspected Stroke / TIA</td> <td>☒ Apnea</td> <td>☒ Preceding severe headache</td> </tr> </table>		☒ Status Epileptics (> 30 min continuous seizure activity)	☒ Hypotension	☒ Pregnancy	☒ Traumatic head injury / fall	☒ History of brain cancer	☒ History of drug and alcohol abuse / overdose	☒ Suspected Stroke / TIA	☒ Apnea	☒ Preceding severe headache									
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<p>History:</p> <ul style="list-style-type: none"> • Presenting complaint • Allergies • Medications: Recent non-compliance with medications, Anticoagulant Therapy, Anti-hypertensive medications, Diabetic medications, Analgesics, Inhalers, Chemotherapy, Non-prescription medications, any recent change to medications. • Past medical past surgical history relevant: <ul style="list-style-type: none"> ○ History of seizures / epilepsy, cancer, infections, CVA / TIA, metabolic disorders, ingestion of toxins, drug and alcohol use, stress, lack of sleep ; recent trauma or fall or head injury; recent overseas travel or immigration; fevers; pregnancy • Last ate / drank and last menstrual period (LMP) • Events and environment leading to presentation i.e. Red flags – History of central nervous system (CNS) pathology (stroke, neoplasms, recent surgery) • Pain Assessment / Score: PQRST (Palliating / provoking factors, Quality, Region / radiation, Severity, Time onset) 																			
<p>Systems Assessment:</p> <ul style="list-style-type: none"> • Focused neurological assessment: <i>Inspection / Palpation / Auscultation (listen)</i> <ul style="list-style-type: none"> ○ Inspect - Level of consciousness, restlessness, pupil size and reaction, abnormal posturing / behaviour, tongue biting, incontinence ○ Listen- Patient complaints; headache, nausea or vomiting ○ Palpate- Equal limb strength, signs of injury 																			
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Nursing Interventions / Management Plan:		
<p>Resuscitation / Stabilisation:</p> <ul style="list-style-type: none"> Oxygen therapy and cardiac monitoring [as indicated] IV Cannulation (consider large bore i.e. 16-18 gauge) <p>Management of an active seizure:</p> <ul style="list-style-type: none"> Airway maneuver Administer oxygen Roll to recovery position Full set of vital signs including BSL Administer medications as per local protocol 	<p>Symptomatic Treatment:</p> <ul style="list-style-type: none"> Antiemetic: as per district standing order Analgesia: as per district standing order IV Fluids: as per district standing order 	
<p>Supportive Treatment:</p> <ul style="list-style-type: none"> Nil By Mouth (NBM) Monitor vital signs as clinically indicated (BP, HR, T, RR, SpO₂) Monitor neurological status - GCS 30 minutely then hourly (as per monitoring recommendation above) 	<ul style="list-style-type: none"> Fluid Balance Chart (FBC) Monitor pain assessment / score 	
<p>Practice Tips / Hints:</p> <ul style="list-style-type: none"> Monitor and assess vital signs including GCS, pupil response and limb strength every 30 min for the first hour and then hourly during the post ictal phase During a generalised seizure the patient may experience a period of transient apnea and hypoxia. In a physiologic effort to maintain cerebral oxygenation, the patient may become hypertensive Hyperthermia, hyperglycemia and lactic acidosis are common following seizures with vigorous muscle activity. These symptoms usually resolve within one hour. A seizure is caused by a sudden and disorderly discharge of cerebral neurons resulting in a change to behavior, sensory perception or motor activity. Seizures are common: approximately 10% of the population will have a seizure within their lifetime and more than 50 million people worldwide are diagnosed with Epilepsy. (Craft, Gordon and Tiziani, 2011; WHO, 2015) It is important to assess the pathophysiology of the seizure to ensure early recognition of life threatening causes and timely treatment is commenced (Craft, Gordon and Tiziani, 2011; Pillow, 2015) There are more than 40 different types of seizure which can be grouped into three classifications: Partial, Generalised and Unclassified seizures (Craft, Gordon and Tiziani, 2011). <ul style="list-style-type: none"> Partial or focal seizures: usually involve one hemisphere of the brain. The area of epileptic neuronal activity will dictate the seizure symptoms. Partial seizures can also be subdivided into simple (no loss of consciousness or awareness) or complex seizures (lowered level of consciousness or awareness) (Craft, Gordon and Tiziani, 2011) Generalised seizures can be subdivided into: Absence, Myoclonic, Tonic-Clonic, Tonic and Atonic seizures (Craft, Gordon and Tiziani, 2011) Unclassified seizures involve seizure activity which does not follow the pattern of either partial or generalised seizures (Craft, Gordon and Tiziani, 2011). 		
<p>Further Reading / References:</p> <ol style="list-style-type: none"> Craft, J., Gordon, C., and Tiziani, A. (2011). Understanding pathophysiology. Mosby, Sydney. Emergency Care Institute (2021) Seizures https://aci.health.nsw.gov.au/networks/eci/clinical/clinical-resources/clinical-tools/neurology/seizures Pillow, T. (2015). Seizure assessment in the Emergency Department. http://emedicine.medscape.com/article/1609294-overview SESLHDPR/283 Deteriorating Patient – Clinical Emergency Response System for the Management of Adult and Maternity inpatients The World Health Organisation (2015). Epilepsy fact sheet. http://www.who.int/mediacentre/factsheets/fs999/en/ 		
<p>Acknowledgements: <i>SESLHD Adult Emergency Nurse Protocols were developed and adapted with permission from:</i></p> <ul style="list-style-type: none"> Murphy, M (2007) Emergency Department Toolkits. Westmead Hospital, SWAHS Hodge, A (2011) Emergency Department, Clinical Pathways. Prince of Wales Hospital SESLHD. 		
Revision and Approval History		
Date	Revision No.	Author and Approval
June 2018	0	Kylie Howes, Nurse Educator. Prince of Wales Hospital Emergency Department.
July 2015	1	Kelly Wright, Clinical Nurse Consultant. The Sutherland Hospital Emergency Department.
September	2	Edited by Alana Clements, CNC St George Hospital Emergency Stream CNC/NE Working Group SESLHD.
October	3	Endorsed by: SESLHD Emergency Clinical Stream Committee
December 2015	4	Endorsed by: SESLHD Drug and Quality Use Medicines Committee
13 April 2016	5	Endorsed by: SESLHD District Clinical and Quality Council on 13 April 2016
June 2018	6	Minor review endorsed by Executive Sponsor.

Adult Emergency Nurse Protocol



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July 2018	6	Endorsed by SESLHD Quality Use of Medicine Committee.
April 2021	7	Revised by Kelly Wright, Clinical Nurse Consultant. The Sutherland Hospital.
May 2021	7	Approved by Executive Sponsor.
June 2021	7	Endorsed by: SESLHD Quality Use of Medicine Committee