

What is a Clinical Pathway?



An evidence-based guideline that decreases unnecessary variation and helps promote safe, effective and consistent patient care.

Objectives of Pathway



- Improve recognition of shunt malfunction on presentation to ED
- Initiate appropriate care for patient with suspected shunt malfunction
- Prevent delay in treatment and management
- Improve patient and family satisfaction
- Improve standard of care

Why do we need this pathway?



- To change practice for these select group of patients with early recognition of potential shunt malfunction and early appropriate imaging and care
- To guide care for these children
- To ensure standard of care is successfully implemented for the safety of the patient

Background Info



- Ventriculoperitoneal (VP) shunt insertion remains the mainstay of treatment for hydrocephalus despite a high rate of complications
- In the United States alone, more than 30,000 procedures to relieve hydrocephalus are performed every year
- The 1-year failure rate for VP shunts had been reported at around 40-50% for pediatric patients
- VP shunt malfunction remains the most frequent reason for shunt revisions and one of the most frequent complication
- Early recognition and treatment improves patient outcomes and decreases hospital stays

The Shunt Malfunction pathway has 2 areas of care: Emergency Department and Inpatient.

We will be reviewing each component in the following slides.

CLINICAL PATHWAY:

Suspected Neurosurgical Shunt Malfunction

Inclusion Criteria: A child that presents with a pre-existing shunt (VP/VA/Vpleural) AND has symptoms associated with malfunction (see below)

- Infants: Enlargement of head, full and tense fontanelle while positioned upright and calm, prominent scalp veins, swelling along the shunt tract, vomiting, irritability, sleepiness, downward deviation of the eyes
- Toddlers: enlargement of head, vomiting, headache, irritability, sleepiness, loss of previous abilities (sensory or motor function)
- Children and adults: vomiting, headache, vision problems, photophobia, irritability, sleepiness, personality change, difficulty in waking up or staying

Exclusion Criteria: Concern for neurosurgical shunt infection (see Suspected Neurosurgical Shunt Infection Clinical Pathway), identification of alternate source for symptoms, or symptoms not related to shunt malfunction as defined

ED Evaluation

- Vitals: BP, HR, O2 sat, RR, temperature
- Head circumference (if age <2 years)
- Place on continuous cardiac and respiratory monitoring
- Notify Neurosurgery attending immediately if bradycardia, hypertension, depressed level of consciousness (LOC)

Initial evaluation:

Obtain a detailed history and initial exam (see Appendix A)

Initial Management

CBC, CRP, BMP

- Shunt tap by Neurosurgery (at the discretion of Neurosurgery attending) c If tapped, send STAT cerebrospinal fluid culture and gram stain

Head ultrasound if fontanelle is open or

- Reduced shunt protocol MRI brain without contrast is preferred imaging modality if can confirm patient has a non-programmable shunt (if not documented in chart, may confirm via skull x-ray; Appendix B: Radiographic Appearance of Shunt Valves)
- c If programmable shunt is present: prior to ordering MRI, please ensure a Neurosurgery provider is able to reprogram the shunt within 24 hours of imaging. Make MRI aware that patient has a
- c If MRI not available: CT head without contrast
- Abdominal ultrasound if abdominal symptoms are present
- Order VP Shunt series at the discretion of the neurosurgery attending

FEN/GI:

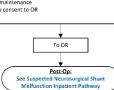
IVF D5 NS with 20 mEq KCI/L at maintenance rate

- Ondansetron 0.1 mg/kg/dose q8hr PRN nausea (max 4 mg/dose)
- Acetaminophen 15 mg/kg/dose q6hr PRN pain/headache (max 75 mg/kg/day or 4,000 mg/day)

Notify Neurosurgery attending via Intellidesk

Admit to Neurosurgery service on the floor if stable, or to the PICU if unstable

- OR case request for shunt revision to be completed by Neurosurgery attending or APP
- Continuous CR monitoring (close monitoring for bradycardia)
- NPO and IVF at maintenance
- Neurosurgery to consent to OR







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We will start with reviewing the Emergency Department pathway.

The goal of the Emergency Department Pathway is to rapidly identify and diagnose patients with shunt malfunction so they can be prepared for surgery as soon as possible.

CLINICAL PATHWAY:

Suspected Neurosurgical Shunt Malfunction

THIS PATHWAY SERVES AS A GUID AND DOES NOT REPLACE CLINICAL

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 <u>Children and adults</u>: vomiting, headache, vision problems, photophobia, irritability, sleepiness, personality change, difficulty in waking up or staying awake

Exclusion Criteria: Concern for neurosurgical shunt infection (see Suspected Neurosurgical Shunt Infection Clinical Pathway), identification of alternate source for symptoms, or symptoms not related to shunt malfunction as defined

ED Evaluation

Triage:

- Vitals: BP, HR, O2 sat, RR, temperature
- Weight
- Head circumference (if age <2 years)
- Pain so
- Place on continuous cardiac and respiratory monitoring
- Notify Neurosurgery attending immediately if bradycardia, hypertension, depressed level of consciousness (LOC)

Initial evaluation:

Obtain a detailed history and initial exam (see Appendix A)

Initial Management

CBC, CRP, BMP

Shunt tap by Neurosurgery (at the discretion of Neurosurgery attending)

c If tapped, send STAT cerebrospinal fluid culture and gram stain

Imaging

Head ultrasound if fontanelle is open or

- Reduced shunt protocol MRI brain without contrast is preferred imaging modality if can confirm patient
 has a nan programmable shunt (if not documented in chart, may confirm via skull x ray; Appendix B:
 Radiographic Appearance of Shunt Valves)
- If programmable shunt is present: prior to ordering MRI, please ensure a Neurosurgery provider is able to reprogram the shunt within 24 hours of imaging. Make MRI aware that patient has a programmable shunt.
- c If MRI not available: CT head without contrast
- Abdominal ultrasound if abdominal symptoms are present
- Order VP Shunt series at the discretion of the neurosurgery attending

FEN/GI:

IVF D5 NS with 20 mEg KCI/L at maintenance rate

Medications:

Ondansetron 0.1 mg/kg/dose q8hr PRN nausea (max 4 mg/dose)

Acetaminophen 15 mg/kg/dose q6hr PRN pain/headache (max 75 mg/kg/day or 4,000 mg/day)

Notify Neurosurgery attending via Intellidesk Pre-Op: Admit to Neurosurgery service on the floor is stable, or to the PICU if unstable OR case request for shunt revision to be completed by Neurosurgery attending or APP Continuous CR monitoring (close monitoring for bradycardia) NPO and IVF at maintenance Neurosurgery to consent to OR To OR

NEXT PAGE

Post-Op:
See Suspected Neurosurgical Shun
Malfunction Inpatient Pathway



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CLINICAL PATHWAY:
Suspected Neurosurgical Shunt Malfunction

THIS PATHWAY SERVES AS A GUIDE AND DOES NOT REPLACE CLINICAL JUDGMENT.

Inclusion Criteria: A child that presents with a pre-existing shunt (VP/VA/Vpleural) AND has symptoms associated with malfunction (see below

If there is concern for shunt infection, please follow the Shunt Infection Clinical Pathway.

Inclusion Criteria: A child that presents with a pre-existing shunt (VP/VA/Vpleural) AND has symptoms associated with malfunction (see below)

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Exclusion Criteria: Concern for neurosurgical shunt infection (see Suspected Neurosurgical Shunt Infection Clinical Pathway), identification of alternate source for symptoms, or symptoms not related to shunt malfunction as defined

ED Evaluation

Note: Any bradycardia, hypertension, and depressed level of consciousness (LOC) are signs of increased intracranial pressure (ICP) and should prompt immediate notification of the Neurosurgery attending



Children may present with different symptoms based on their age.

 All children under 2 years of age should have a head circumference documented

Providers should complete a thorough history and physical exam

See Appendix A

Triage:

- Vitals: BP, HR, O2 sat, RR, temperature
- Weight
- Head circumference (if age <2 years)
- Pain score
- Place on continuous cardiac and respiratory monitoring
- Notify Neurosurgery attending immediately if bradycardia, hypertension, depressed level of consciousness (LOC)

Initial evaluation:

Obtain a detailed history and initial exam (see Appendix A)

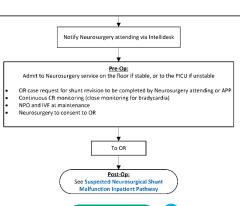
NPO
 IVF D5

IVF D5 NS with 20 mEq KCI/L at maintenance rate

Medications:

Ondansetron 0.1 mg/kg/dose q8hr PRN nausea (max 4 mg/dose)

Acetaminophen 15 mg/kg/dose q6hr PRN pain/headache (max 75 mg/kg/day or 4,000 mg/day)



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CLINICAL PATHWAY: Suspected Neurosurgical Shunt Malfunction Appendix A: Obtaining a Detailed History

Appendix A: Obtaining a detailed history and physical

Important factors to include:

- Shunt history, including:
 - Location of shunt (ventricular-atrial shunt, ventricular-pleural shunt, ventricularperitoneal shunt)
 - Date of shunt placement
 - Date of last shunt revision
 - Signs/symptoms present at presentation/last revision
- Headache history, including:
 - Quality
 - Duration
 - Location
 - Past treatment
- Vomiting history, including:
 - Timing
 - Any precipitating events
- Neurological symptoms, including:
 - Change in LOC
 - Increased irritability

 - Increased lethargy
- Abdominal symptoms, including:
 - Significant increase in abdominal girth
 - Pain Tenderness

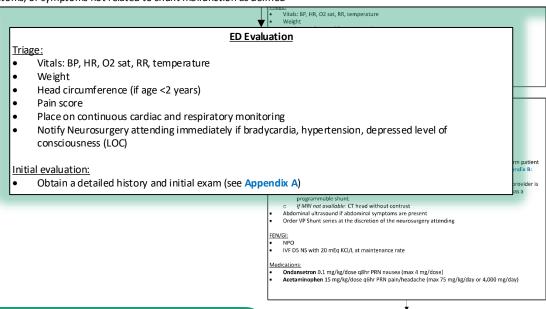
- Trauma history
- - Head circumference
 - Decreased breath sounds for pleural shunt

CLINICAL PATHWAY: **Suspected Neurosurgical Shunt Malfunction**

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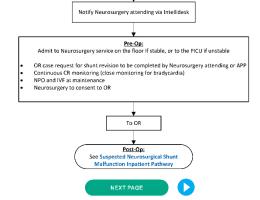
Appendix A provides guidelines for pertinent history and physical exam factors which will be important for correct diagnosis.











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Initial management includes obtaining imaging, sending screening lab work, and making the patients NPO in prep for surgery.

> The neurosurgery attending should be notified after imaging is completed and with ANY signs of increased ICP

CLINICAL PATHWAY:

Suspected Neurosurgical Shunt Malfunction

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Initial Management

- CBC, CRP, BMP
- Shunt tap by Neurosurgery (at the discretion of Neurosurgery attending)
 - If tapped, send STAT cerebrospinal fluid culture and gram stain

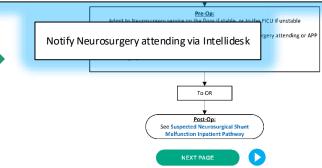
Imaging:

- Head ultrasound if fontanelle is open or
- Reduced shunt protocol MRI brain without contrast is preferred imaging modality if can confirm patient has a non-programmable shunt (if not documented in chart, may confirm via skull x-ray; Appendix B: Radiographic Appearance of Shunt Valves)
 - If programmable shunt is present: prior to ordering MRI, please ensure a Neurosurgery provider is able to reprogram the shunt within 24 hours of imaging. Make MRI aware that patient has a programmable shunt.
 - If MRI not available: CT head without contrast
- Abdominal ultrasound if abdominal symptoms are present
- Order VP Shunt series at the discretion of the neurosurgery attending

FEN/GI:

- IVF D5 NS with 20 mEq KCI/L at maintenance rate

- Ondansetron 0.1 mg/kg/dose q8hr PRN nausea (max 4 mg/dose)
- Acetaminophen 15 mg/kg/dose q6hr PRN pain/headache (max 75 mg/kg/day or 4,000 mg/day)



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Before MRI, confirm that the patient has a non-programmable shunt.

If there is a programmable shunt present, check with the Neurosurgery team and ensure they are able to reprogram the shunt within 24 hours of imaging.

CLINICAL PATHWAY:

Suspected Neurosurgical Shunt Malfunction

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AND DOES NOT
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Initial Management

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Imaging:

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- Order VP Shunt series at the discretion of the neurosurgery attending

FEN/GI:

- NPO
- IVF D5 NS with 20 mEg KCI/L at maintenance rate

Medications:

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Pre-Op:
Admit to Neurosurgery service on the floor if stable, or to the PICU if unstable

Notify Neurosurgery attending via Intellides k

To OR

Post-Op:
See Suspected Neurosurgical Shunt
Malfunction Inpatient Pathway

NEXT PAGE

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CLINICAL PATHWAY: Suspected Neurosurgical Shunt Malfunction Appendix B: Radiographic Appearance of Shunt Valves

When evaluating the radiographic markings of any implanted device, it is important to recognize that the veracity of your interpretation depends on the quality of the radiographic images. For the best results, x-rays should be taken orthogonally to the plane of the shunt valve. The positioning of the valve relative to the skull base may also obscure the valve markings, as overlapping radiodensities along the skull base can blur valve markings. In more difficult cases, fluoroscopy or 3D CT reconstruction may be used to properly identify the radio-opaque markings on a shunt valve.

It is important to realize that an exhaustive list of all shunt valve radiographic markings is beyond the scope of this appendix. For additional information regarding common shunt valve markings found in North American neurosurgical patients, you may also reference the ISPN's website on the same topic.

Please see the next several pages for examples of radiographic images of nonprogrammable and programmable shunts. The sources of these images are:

- http://www.kinderneurochirurgie-leipzig.de/therapeuticfocus/hydrocephalus/radiologicidentification-of-vp-shunt-valves-and-adjustment/
- https://www.ispn.guide/
- https://www.medtronic.com/us-en/index.html
- https://radiopaedia.org/

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Initial Management

Labs:

- CBC, CRP, BMP
- Shunt tap by Neurosurgery (at the discretion of Neurosurgery attending)
 - If tapped, send STAT cerebrospinal fluid culture and gram stain

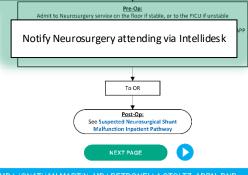
Imaging:

- Head ultrasound if fontanelle is open or
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Appendix B outlines radiographic considerations when evaluating a shunt, with imaging examples provided.

urgery attending

- Ondansetron 0.1 mg/kg/dose q8hr PRN nausea (max 4 mg/dose)
- Acetaminophen 15 mg/kg/dose q6hr PRN pain/headache (max 75 mg/kg/day or 4,000 mg/day)



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Once a patient is identified as having a shunt malfunction, they will be admitted (to the Med/Surg unit or PICU depending on their clinical stability) or taken to the OR.

Post-operatively, the inpatient portion of the pathway will be launched.

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ED Evaluation

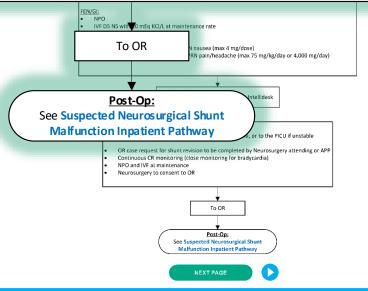
- Vitals: BP, HR, O2 sat, RR, temperature
- Head circumference (if age <2 years)
- Place on continuous cardiac and respiratory monitoring
- Notify Neurosurgery attending immediately if bradycardia, hypertension, depressed level of

Obtain a detailed history and initial exam (see Appendix A)

Pre-Op:

Admit to Neurosurgery service on the floor if stable, or to the PICU if unstable

- OR case request for shunt revision to be completed by Neurosurgery attending or APP
- Continuous CR monitoring (close monitoring for bradycardia)
- NPO and IVF at maintenance
- Neurosurgery to consent to OR



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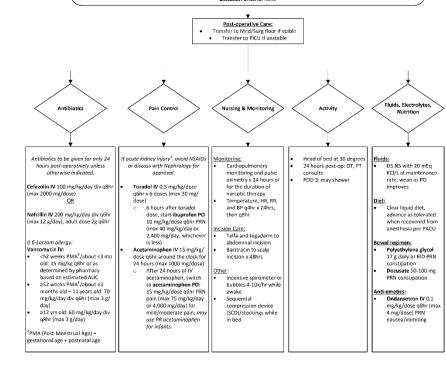
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The goal of the Inpatient pathway is to guide postoperative care of patients who underwent surgical correction of a shunt malfunction.

CLINICAL PATHWAY:

Suspected Neurosurgical Shunt Malfunction

Inclusion Criteria: post-operative care for any patient diagnosed by Neurosurgery to have shunt malfunction requiring surgical correction Exclusion Criteria: none



Discharge Criteria:

- Baseline neurological examination
- Pain well-controlled on oral medication
- Afebrile x 24 hours
- Taking adequate fluid and nutrition orally Cleared by PT & OT

Discharge Medications:

- Ibuprofen PO 10 mg/kg/dose q6hr PRN (max 40 mg/kg/day or 2,400 mg/day, whichever is less) for mild/moderate pair
- Acetaminophen PO: 15 mg/kg/dose q6hr PRN pain (max 75 mg/kg/day or 4,000 mg/day) for mild/ moderate pain
- Polyethylene glycol PO and/or docusate to prevent constination

Discharge Instructions

Call Neurosurgery for fever >101.5, vomiting >3x in 12 hr period, excessive irritability o

Follow up outpatient 2-3 weeks after discharge

Definition of Acute Kidney Injury (It should be noted that this

definition does not apply to children <1 year of age)

- AKI is defined by having either At least a 50% increase in Scr above baseline* and new Scr ≥0.5 mg/dL OR
- An increase by 0.3 mg/dL from baseline*, and new Scr ≥0.5

*If a baseline creatinine is unknown estimate baseline Cr using the Schwartz Calculation (buseline creatinine = (0.413 + height cm)/ 120 GFR). For patients with Chronic Kidney Disease (CKD), use the CKID





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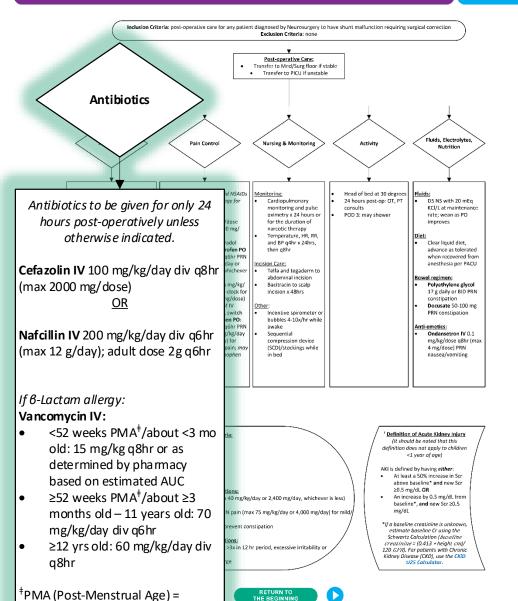
Antibiotics are only given for the first 24 hours post-operatively, unless otherwise indicated.

CLINICAL PATHWAY:

Suspected Neurosurgical Shunt Malfunction

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Connecticut



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gestational age + postnatal age

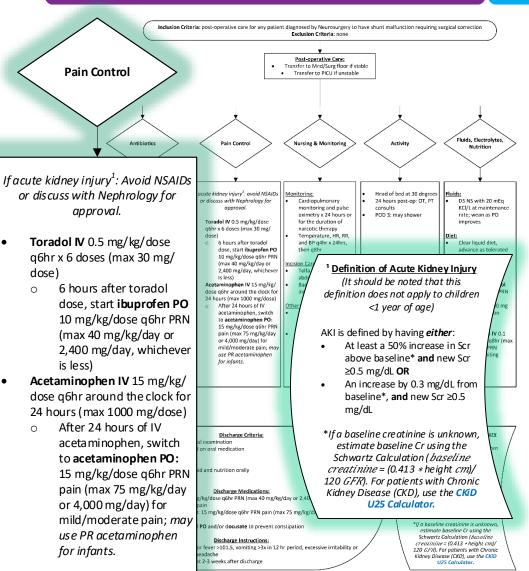
Pain can typically be managed by toradol/ibuprofen and acetaminophen.

However, those with renal disease or impairment should avoid the use of NSAIDs.

Note: the definition of AKI has been updated and is available as a key.

CLINICAL PATHWAY: Suspected Neurosurgical Shunt Malfunction

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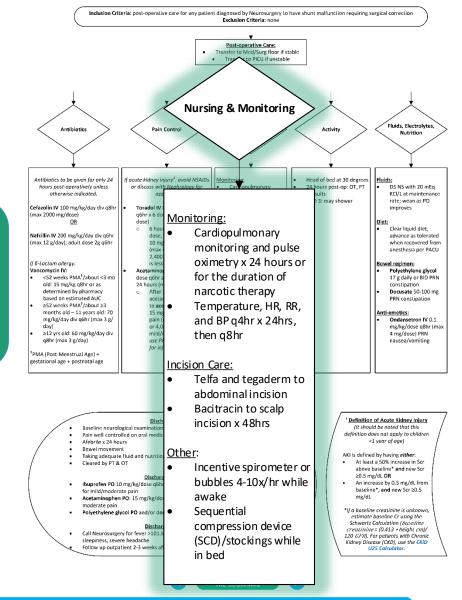


Patients will need typical post anesthesia nursing care but with close observation of the surgical sites for leakage.

CLINICAL PATHWAY:

Suspected Neurosurgical Shunt Malfunction

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PT and OT are initiated on post operative day 1 to encourage early movement.

CLINICAL PATHWAY:

Suspected Neurosurgical Shunt Malfunction

THIS PATHWAY
SERVES AS A GUIDE
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REPLACE CLINICAL
JUDGMENT:

Inclusion Criteria: post-operative care for any patient diagnosed by Neurosurgery to have shunt malfunction requiring surgical correction Exclusion Criteria: none Post-operative Care: Transfer to Med/Surg floor if stable
Transfer to PICU if unstable Antibiotics Pain Control **Activity** Antibiotics to be given for only 24 acute kidney injury¹: avoid NSAIDs hours post-operatively unless or discuss with Nephrology for D5 NS with 20 mEq otherwise indicated. monitoring and pulse KCI/L at maintenance aximetry x 24 hours or rate; wean as PO Cefazolin IV 100 mg/kg/day diy o8hr Toradol IV 0.5 mg/kg/dose for the duration of improves (max 2000 mg/dose) q6hr x 6 doses (max 30 mg/ narcotic therapy Temperature, HR, RR 6 hours after toradol and BP q4hr x 24hrs, Clear liquid diet, Nafcillin IV 200 mg/kg/day div q6hr dose, start ibuprofen PO (max 12 g/day); adult dose 2g q6hr 10 mg/kg/dose q6hr PRN (max 40 mg/kg/day or 2,400 mg/day, whichever If 6-Lactam allergy: is less) e glycol BID PRN Vancomycin IV: Acetaminophen IV 15 mg/kg/ <52 weeks PMA[†]/about <3 mo dose q6hr around the clock for Head of bed at 30 old: 15 mg/kg o8hr or as 24 hours (max 1000 mg/dose) After 24 hours of IV 0-100 mg determined by pharmacy based on estimated AUC acetaminophen, switch degrees ≥52 weeks PMA[†]/about ≥3 to acetaminophen PO: months old - 11 years old: 70 15 mg/kg/dose q6hr PRN mg/kg/day div q6hr (max 3 g/ pain (max 75 mg/kg/day 24 hours post-op: OT, or 4,000 mg/day) for ≥12 yrs old: 60 mg/kg/day div mild/moderate pain; may **PT consults** q8hr (max 3 g/day) use PR acetaminophen for infants. PMA (Post-Menstrual Age) = POD 3: may shower gestational age + postnatal age Discharge Criteria: Baseline neurological examination (It should be noted that this Pain well-controlled on oral medication definition does not apply to children Afebrile x 24 hours <1 year of age) AKI is defined by having either: Taking adequate fluid and nutrition orally At least a 50% increase in Scr Cleared by PT & OT above baseline* and new Scr ≥0.5 mg/dL OR Discharge Medications: An increase by 0.3 mg/dL from Ibuprofen PO 10 mg/kg/dose q6hr PRN (max 40 mg/kg/day or 2,400 mg/day, whichever is less) baseline*, and new Scr ≥0.5 for mild/moderate pair Acetaminophen PO: 15 mg/kg/dose q6hr PRN pain (max 75 mg/kg/day or 4,000 mg/day) for mild/j moderate pain *If a baseline creatinine is unknown Polyethylene glycol PO and/or docusate to prevent constination estimate baseline Cr using the Schwartz Calculation (buseline Discharge Instructions: creatinine = (0.413 + height cm)/ Call Neurosurgery for fever >101.5, vomiting >3x in 12 hr period, excessive irritability or 120 GFR). For patients with Chronic Kidney Disease (CKD), use the CKID U25 Calculator. Follow up outpatient 2-3 weeks after discharge



RETURN TO THE BEGINNING



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There is NO routine blood work required post operatively.

Diet is advanced as tolerated.

Bowel regimen is essential and should be started as soon as possible post procedure.

CLINICAL PATHWAY:

Suspected Neurosurgical Shunt Malfunction

FHIS PATHWAY SERVES AS A GUIDE AND DOES NOT REPLACE CLINICAL JUDGMENT.

nausea/vomiting

Inclusion Criteria: post-operative care for any patient diagnosed by Neurosurgery to have shunt malfunction requiring surgical correction Exclusion Criteria: none Post-operative Care: Transfer to Med/Surg floor if stable Transfer to PICU if unstable Fluids, Electrolytes, Nutrition Antibiotics Pain Control Nursing & Monitoring Antibiotics to be given for only 24 cute kidney injury¹: avoid NSAIDs Head of bed at 30 degre hours post-operatively unless or discuss with Nephrology for Cardiopulmonary otherwise indicated. monitoring and pulse aximetry x 24 hours or Cefazolin IV 100 mg/kg/day diy o8h Toradol IV 0.5 mg/kg/dose for the duration of (max 2000 mg/dose) q6hr x 6 doses (max 30 mg/ narcotic therapy Temperature, HR, RF Fluids: 6 hours after toradol and BP q4hr x 24hrs, Nafcillin IV 200 mg/kg/day div q6hr dose, start ibuprofen PO D5 NS with 20 mEq (max 12 g/day); adult dose 2g q6h 10 mg/kg/dose q6hr PRN (max 40 mg/kg/day or 2,400 mg/day, whicheve Telfa and tegaderm to KCI/L at maintenance If 6-Lactam allergy. is loss) abdominal incision /ancomycin IV: Acetaminophen IV 15 mg/kg/ Bacitracin to scalp rate; wean as PO <52 weeks PMA[†]/about <3 mg dose q6hr around the clock for incision x 48hrs old: 15 mg/kg o8hr or as 24 hours (max 1000 mg/dose) improves determined by pharmacy After 24 hours of IV based on estimated AUC acetaminophen, switch Incentive spirometer or ≥52 weeks PMA[†]/about ≥3 to acetaminophen PO: bubbles 4-10x/hr while months old - 11 years old: 70 15 mg/kg/dose q6hr PRN mg/kg/day div q6hr (max 3 g/ pain (max 75 mg/kg/day Sequential Diet: or 4,000 mg/day) for compression device ≥12 yrs old: 60 mg/kg/day div mild/moderate pain; mo Clear liquid diet. q8hr (max 3 g/day) use PR acetaminophen for infants. advance as tolerated PMA (Post-Menstrual Age) = estational age + postnatal age when recovered from anesthesia per PACU Bowel regimen: Discharge Criteria: Baseline neurological examination Polyethylene glycol Pain well-controlled on oral medication Afebrile x 24 hours 17 g daily or BID PRN Bowel movemen Taking adequate fluid and nutrition orally constipation Cleared by PT & OT Discharge Medications: Docusate 50-100 mg Ibuprofen PO 10 mg/kg/dose q6hr PRN (max 40 mg/kg/day or 2,400 mg/day, whichever is less) for mild/moderate pair PRN constipation Acetaminophen PO: 15 mg/kg/dose q6hr PRN pain (max 75 mg/kg/day or 4,000 mg/day) for mild/ moderate pain Polyethylene glycol PO and/or docusate to prevent constination Anti-emetics: Discharge Instructions Call Neurosurgery for fever >101.5, vomiting >3x in 12 hr period, excessive irritability of Ondansetron IV 0.1 Follow up outpatient 2-3 weeks after discharg mg/kg/dose q8hr (max 4 mg/dose) PRN

CONTACTS: MARKUS BOOKLAND, MD | JONATHAN MARTIN, MD | PETRONELLA STOLTZ, APRI

Certain criteria must be met prior to discharge, including adequate pain control and bowel movements.

Medications focus on pain management and maintaining adequate bowel movements.

Education regarding when to call neurosurgery post discharge is very important to ensure no complications exist post operatively. Early recognition is important.

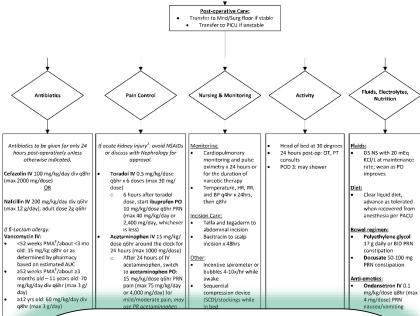
CLINICAL PATHWAY:

Suspected Neurosurgical Shunt Malfunction

THIS PATHWAY SERVES AS A GUIDI AND DOES NOT REPLACE CLINICAL JUDGMENT.

Inclusion Criteria: post-operative care for any patient diagnosed by Neurosurgery to have shunt malfunction requiring surgical correction

Exclusion Criteria: none



MA (Post-Mensestational

Discharge Criteria:

- Baseline neurological examination
- Pain well-controlled on oral medication
- Afebrile x 24 hours
- Bowel movement
- Taking adequate fluid and nutrition orally
- Cleared by PT & OT

Discharge Medications:

- Ibuprofen PO 10 mg/kg/dose q6hr PRN (max 40 mg/kg/day or 2,400 mg/day, whichever is less) for mild/moderate pain
- Acetaminophen PO: 15 mg/kg/dose q6hr PRN pain (max 75 mg/kg/day or 4,000 mg/day) for mild, moderate pain
- Polyethylene glycol PO and/or Docusate to prevent constipation

Discharge Instructions:

- Call Neurosurgery for fever >101.5, vomiting >3x in 12 hr period, excessive irritability or sleepiness, severe headache
- Follow up outpatient 2-3 weeks after discharge

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Review of Key Points



- Appropriate imaging to rule out shunt malfunction is imperative to determine need for surgical intervention
- Timely pre operative care helps facilitate timely transfer to OR
- Standardized post-operative care assists in management, discharge planning and follow up

Quality Metrics



- Percent of patients with pathway order set usage
- Percent of patients with deep wound infections
- Percent of patients with superficial wound infections
- Number of patients with organ space infection within 30 days of principal operative procedure
- Number of patients with shunt malfunction within 90 days of principal operative procedure
- Percentage of patients with cerebrospinal fluid leak
- Number of readmissions within 30 days
- Number of patients with return to the OR within 30 days

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Thank You!



About Connecticut Children's Pathways Program

Clinical pathways guide the management of patients to optimize consistent use of evidence-based practice. Clinical pathways have been shown to improve guideline adherence and quality outcomes, while decreasing length of stay and cost. Here at Connecticut Children's, our Clinical Pathways Program aims to deliver evidence-based, high value care to the greatest number of children in a diversity of patient settings. These pathways serve as a guide for providers and do not replace clinical judgment.