Clinical Pathways

Delirium Clinical Pathway Emergency Department and Inpatient Care

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Objectives of the Pathway



- Define delirium and understand the causes of delirium in pediatric patients
- Describe strategies to prevent delirium
- Demonstrate how to use and interpret the Cornell Assessment of Pediatric Delirium (CAPD) to screen for delirium
- Review important components of the new ED Delirium Clinical Pathway and order set
- Review important components of the new Inpatient Delirium Evaluation, Workup and Management Clinical Pathway and order sets

What is Delirium?



- Acute-onset neuropsychiatric syndrome characterized by disturbances of cognition, attention, consciousness or perception that is potentially life-threatening
 - Secondary to a medical etiology (not an isolated psychiatric condition)
 - o Can occur as a result of underlying illness, hospitalization, medications or trauma
- Treatment requires inter-professional collaboration between primary physicians, specialists, nursing, and family
 - Early recognition and treatment may prevent adverse outcomes

Why do we care?



- Delirium is a high risk diagnosis, serving as a sign of acute brain dysfunction and a marker for potential significant clinical decompensation
- All hospitalized patients are at risk of developing delirium
- Often under-recognized in children:
 - O Affects 10-44% of hospitalized children and up to 30% of PICU patients (Bettencourt 2017, Traube 2014, Traube 2017, Smith 2013)
 - Signs may be very subtle

Clinical Presentation of Delirium



- Acute onset (hours-days)
- Waxing/waning course with lucid intervals
- Sleep/wake cycle disruption (often reversed)
- Disturbed consciousness
- Neurocognitive deficits
- Perception, hyper/hypoactivity, mood/affect
- Direct physiological consequence of medical/organic etiology

Clinical Presentation of Psychiatric Illness



Psychosis:

o Presence of hallucinations, delusional thoughts

Mania:

 Elated mood, increased energy, rapid speech, grandiosity, decreased need for sleep, impulsivity, flight of ideas, distractibility

Depression:

o Depressed mood, anhedonia, change in sleep/energy/concentration, guilt, suicidal ideation

Some symptoms overlap with delirium, but....

Delirium vs. Psychiatric Illness



Primary psychiatric illness does **not** have...

- Acute onset
- Fluctuating course
- Disorientation
- Disturbed consciousness
- Memory/Language/Visuospatial impairment
- Confusion, Inattention

Types of Delirium



Delirium in children can present as hypoactive, hyperactive or mixed type

Table 2 Types of delirium						
Туре	Signs	Patient example				
Hypoactive	Child looks apathetic and seems uninterested ²	Toddler who lies quietly in the bed and does not make eye contact or reach for toys or family members				
Hyperactive	Child is irritable despite adequate pain medication and may be thrashing ²	School-aged child receiving mechanical ventilation who is constantly moving around in bed despite adequate pain medication Patient is difficult to sedate				
Mixed	Child fluctuates between a hypoactive and a hyperactive state ²	Teenager who vacillates between yelling at staff and thrashing in the bed to being calm and staring off into the distance with no interactions with staff at different times of the day				

Potential Causes of Delirium



- Infection (intracranial or systemic)
 - o Fever
 - Sepsis
- Drug intoxication
- Drug withdrawal
- Medications
 - Opioids, Benzodiazepines
 - Anti-histamines
 - Corticosteroids
- Metabolic/Endocrine disturbance
 - o Electrolyte abnormality
 - Hypoglycemia

- Traumatic Brain Injury
- Seizures
- Hypoxia
- Neoplasm
- Cerebrovascular event
- Autoimmune encephalitis
- Organ dysfunction/Insufficiency
- Hospitalization (Environment)
 - Sleep/wake cycle disruption
 - Prolonged immobilization
 - Unfamiliar surroundings, sensory loss
 - Unmanaged painful stimuli

Independent Risk Factors for Developing Delirium



- Age < 2yo
- Developmental delay
- Illness severity
- Prior coma
- Mechanical ventilation
- Receiving benzodiazepines or anticholinergics

Traube 2017, Silver 2015

Complications from Delirium



- Increased:
 - Length of stay
 - Safety events (i.e. pulling lines, falls)
 - Morbidity and mortality
 - Cost of hospitalization
 - Use of restraints and sedatives

(Traube 2017, Traube 2016, Turkel 2017)

 Reported long term neuro-developmental and behavioral consequences, including development of PTSD following hospitalization (Brummel 2014)

Delirium Clinical Pathways



- Pathway can help guide appropriate medical evaluation and management for patients with recognized delirium
 - o There is a high clinical suspicion for delirium if a patient has <u>any one</u> of the following features:
 - Acute mental status change
 - Acute onset hallucinations/delusions
 - Confusion or impaired memory
 - Alteration in attention or arousal
 - New catatonic features

Delirium Clinical Pathways



There are 2 Delirium Clinical Pathways:

1. Delirium Emergency Department Care

o This pathway is focused on identifying delirium and initiating work-up prior to admission

2. Delirium Inpatient Care

- O This pathway has three main aims:
 - Prevent and identify delirium in the inpatient setting
 - Guide work-up
 - Manage symptoms

Pathway Overview

The first page is a general overview of the ED and inpatient pathway.

Note that phases of care and scoring tools are easily accessible.

Note that all patients admitted to the med/surg floors are screened for delirium, in order to identify patients early.

CLINICAL PATHWAY:

Delirium Emergency Department and Inpatient

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

Inclusion Criteria: Any patient in the Emergency Department or Inpatient Med/Surg Units with any of the following:

- Acute mental status change, acute onset hallucinations or delusions, confusion, impaired memory, alteration of attention or arousal, acute catatonia; <u>OR</u>
- Clinical suspicion of delirium based on Vanderbilt Assessment for Delirium in Infants and Children (VADIC)
 Assessment Tool or Cornell Assessment of Pediatric Delirium (CAPD) Score
- All patients admitted to Medical/Surgical floors will be screened for delirium

Exclusion Criteria: Patient located in the NICU, ambulatory and perioperative areas, infusion patients, PICU. If in PICU, follow PICU protocol for screening and prevention.

Etiologies to consider:

CNS infection, fever, sepsis/end organ dysfunction (see Sepsis Pathway), Multi-system Inflammatory Syndrome in Children (see MIS-C Pathway), hypoxemia, hypoglycemia, electrolyte abnormality, CNS abnormality, intoxication, autoimmune encephalitis, SLE, vasculitis, drug withdrawal, metabolic disease, neoplasm

Phase of Care - Navigation Links

Emergency Department

Inpatient and ED (Zone C) Management

Inpatient Prevention and Screening

Inpatient Evaluation and Work Up

Scoring Tools - Navigation Links

Appendix A: Vanderbilt Assessment for Delirium in Infants and Children (VADIC) Assessment Tool

Appendix B: Cornell Assessment of Pediatric Delirium (CAPD) Score

Appendix C: Developmental Anchors

NEXT PAGE



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- 1. Evaluation and Work Up
- 2. Management

It is important to identify potential etiologies of delirium first, and disease specific management should occur.

If a specific etiology for delirium is not identified on initial assessment, further lab and imaging studies are recommended.

CLINICAL PATHWAY:

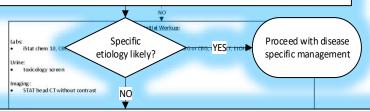
Delirium - Emergency Department Care



Etiologies to consider:

Etiologies to conside

CNS infection, fever, sepsis/end organ dysfunction (see Sepsis Pathway), Multi-system Inflammatory Syndrome in Children (see MIS-C Pathway), hypoxemia, hypoglycemia, electrolyte abnormality, CNS abnormality, intoxication, autoimmune encephalitis. SLE. vasculitis. drug withdrawal, metabolic disease, neoplasm



Initial Workup:

lahs.

iStat chem 10, CBC, CRP, ESR, ammonia, PT/PTT/INR, TSH, free T4, VBG or CBG, AST, ALT, EtOH level, ANA

Urine:

toxicology screen

Imaging:

STAT head CT without contrast

If febrile:

- Blood and urine cultures
- Strongly consider LP: cell count with differential, protein, glucose, gram stain and culture, HSV PCR, enterovirus PCR, opening pressure. Ask lab to hold 3 mL CSF for further studies.
- Begin all empiric IV antimicrobials listed below:
 - Ceftriaxone IV 100 mg/kg/day q12hr (max 2,000 mg/dose) x48 hours AND
 - Vancomycin IV x48 hours:
 - <52 weeks PMA[†]/about <3 mo old: 15 mg/kg q8hr or as determined by pharmacy based on estimated AUC</p>
 - ≥52 weeks PMA[†]/about ≥3 months old 11 years old: 70 mg/kg/day div q6hr
 - ≥12 yrs old: 60 mg/kg/day div q8hr AND
 - Acyclovir 20 mg/kg/dose IV q8hr until HSV studies negative

[‡]PMA (Post-Menstrual Age) = gestational age + postnatal age

infectious proces







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- 1. Evaluation and Work Up
- 2. Management

If patient is **febrile**, blood and urine cultures should be obtained, and an LP is strongly recommended.

When performing the LP, please send as much CSF possible to the lab to be saved for potential future studies.

- * Minimum of 3 ml of CSF should be saved, but as much as 6 ml may be needed for some panels.
- * Please call the lab to confirm CSF is being held.

CLINICAL PATHWAY:

Delirium - Emergency Department Care



Etiologies to consider:

Etiologies to conside

CNS infection, fever, sepsis/end organ dysfunction (see Sepsis Pathway), Multi-system Inflammatory Syndrome in Children (see MIS-C Pathway), hypoxemia, hypoglycemia, electrolyte abnormality, CNS abnormality, intoxication, autoimmune encephalitis. SLE. vasculitis. drug withdrawal, metabolic disease, neoplasm



Initial Workup:

Labs:

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- 1. Evaluation and Work Up
- 2. Management

If the patient is febrile with delirium, empiric broad spectrum antimicrobial coverage should be initiated.

Note: the pharmacy's vancomycin protocol was updated in Feb 2021.

- All patients who have vancomycin IV ordered will be followed by the clinical pharmacist to help determine appropriate dosing parameters.
- Providers will order initial doses per pathway/order set and provide indication within the order.
- IV vancomycin dosing and recommended labs will be managed by pharmacy in conjunction with primary teams.

CLINICAL PATHWAY:

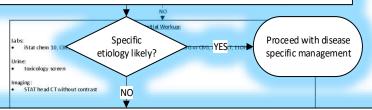
Delirium - Emergency Department Care

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Etiologies to consider:

Etiologies to conside

CNS infection, fever, sepsis/end organ dysfunction (see Sepsis Pathway), Multi-system Inflammatory Syndrome in Children (see MIS-C Pathway), hypoxemia, hypoglycemia, electrolyte abnormality, CNS abnormality, intoxication, autoimmune encephalitis. SLE. vasculitis. drug withdrawal, metabolic disease, neoplasm



Initial Workup:

Labs:

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- 1. Evaluation and Work Up
- 2. Management

Consider ED Social Work and/or Psychiatric consult to help determine and support behavioral health needs and establish follow up plan.

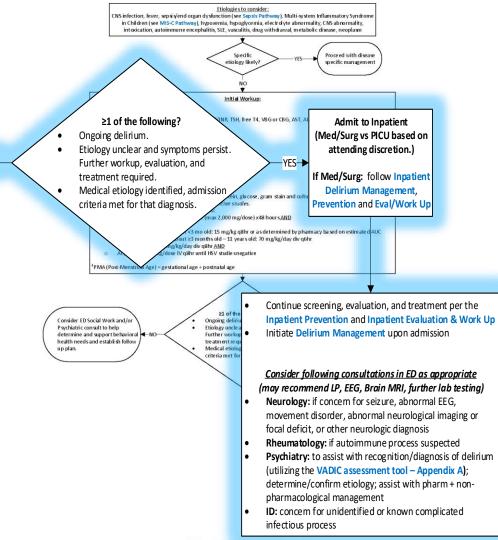
Based on initial testing and continual evaluation, disposition can be determined.

Specific criteria warrant inpatient admission, including ongoing delirium, or ongoing symptoms with need of further interventions.

Considerations for additional consults are outlined.

CLINICAL PATHWAY: Delirium - Emergency Department Care

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





RETURN TO THE BEGINNING



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- 1. Evaluation and Work Up
- 2. Management

Etiologies to consider: NS infection, fever, sepsis/end organ dysfunction (see Sepsis Pathway), Multi-system Inflammatory Syndrome in Children (see MIS-C Pathway), hypoxemia, hypoglycemia, electrolyte abnormality, CNS abnormality, intoxication, autoimmune encephalitis, SLE, vasculitis, drug withdrawal, metabolic disease, neoplasm Proceed with disease etiology likely? Initial Workup: ≥1 of the following? TSH, free T4, VBG or CBG, AST, A Admit to Inpatient Consider ED Social Work and/or Ongoing delirium. (Med/Surg vs PICU based on Psychiatric consult to help Etiology unclear and symptoms persist. attending discretion.) determine and support behavioral NO--YES→ Further workup, evaluation, and health needs and establish follow If Med/Surg: follow Inpatient treatment required. up plan. Medical etiology identified, admission Delirium Management, n, glucose, gram stain and cul criteria met for that diagnosis. Prevention and Eval/Work Up mo old: 15 mg/kg q8hr or as determined by pharmacy based on estimated A t ≥3 months old - 11 years old: 70 mg/kg/day div q6hr kg/day div q8hr AND tal Age) = gestational age + postnatal age Continue screening, evaluation, and treatment per the Inpatient Prevention and Inpatient Evaluation & Work Up Consider ED Social Work and/or Ongoing deli P sychiatric consult to help Etiology uncle Initiate **Delirium Management** upon admission determine and support behaviora health needs and establish follow tre atment re Medical etic criteria met fo Consider following consultations in ED as appropriate (may recommend LP, EEG, Brain MRI, further lab testing) Neurology: if concern for seizure, abnormal EEG, movement disorder, abnormal neurological imaging or focal deficit, or other neurologic diagnosis Rheumatology: if autoimmune process suspected **Psychiatry:** to assist with recognition/diagnosis of delirium (utilizing the VADIC assessment tool - Appendix A); determine/confirm etiology; assist with pharm + nonpharmacological management ID: concern for unidentified or known complicated infectious process

Delirium - Emergency Department Care

Appendix A: The Vanderbilt Assessment for Delirium in Infants and Children (VADIC)

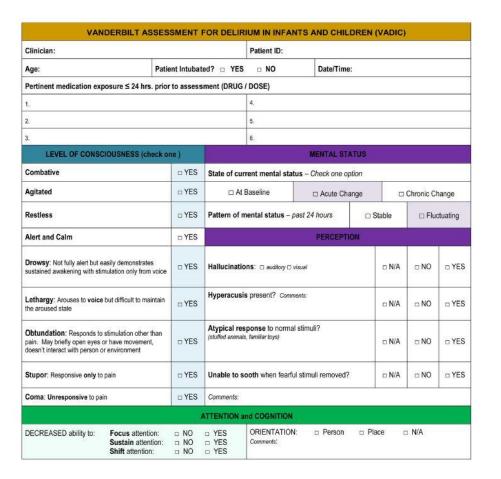


CLINICAL PATHWAY:

- 1. Evaluation and Work Up
- 2. Management

Appendix A: The Vanderbilt Assessment for Delirium in Infants and Children (VADIC)

This is tool provides a comprehensive framework to standardize pediatric delirium assessment by psychiatrists.



CLINICAL PATHWAY:

Delirium - Emergency Department Care

THIS PATHWAY
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AND DOES NOT
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CNS infection, fever, sepsi s/end organ dy sfunction (see Sepsis Pathway), Multi-system Inflammatory Syndr in Children (see MIS-C Pathway), hypoxemia, hypoglycemia, electrolyte abnormality, CNS abnormality.	
in Children (see \$400 C Dethrond) has a coming to produce only all outside the constitution of the constit	ome
intoxication, autoimmune encephalitis, SLE, vasculitis, drug withdrawal, metabolic disease, neoplasm	

DECREASED ability to screen out ext DECREASED ability to interact with to DECREASED social smile in response Object permanence present? (interacts	ys/objects appropriately? (No inte to toys or stuffed animals?		oriately)		
SLEEP-WAKI	ECYCLE	AF	FECT		
Normal Nap Patterns (02-4h infants, 08s preschod): Nocturnal Disturbance: (initial, midd insonnia, phase shift) Day-Night Reversal (more difficult to recog	e, terminal ONO OYES	Irritability or anger DO YES Inconsolability PES			
Comments:					
3	LANGUAGE	and THOUGHT			
	YES YES YES YES Abated)	Describe baseline speech and language per parent/nurse if available: Appropriate Decreased smount Decreased sontaneity Increased latency Change from baseline Circumstantial Tangential Obstructed due to disease or device			
	IS ACUTE DEL	RIUM PRESENT?			
UTA When LOC severely depressed	, unable to directly clinically assess p	atient AND prior clinical assessment not a	ovallable.		
■ NO If NO consider → Subsynd	romal delirium(SS) (Delirium pro	bable but NOT all criteria met): 🛚	NO - YES		
YES If YES then choose type →	HYPOACTIVE HYPERACT	IVE II MIXED Drug Withdi	rawai? N/A NO YES		
24-HOUR assessment → IS DELIF	RIUM PRESENT? PRESEN	T - ABSENT - SUBSYN	IDROMAL - UTA		
□ 1. Acute change Mental Status	a 3. Inattention present	□ 5. Change in Cognition	□ 7. Change in Affect		
p 2. Fluctuating Course	p 4. Inconsolability	6. Change in Lan- guage/Thought	a 8. Change in Sleep/Wake Cycle		

DELIRIUM = 1+2+3+5+7 AND 4 OR 6 OR 8



1. Management

Management of Delirium in the Inpatient or Zone C setting encompasses 4 key categories:

- Treat the suspected etiology
- Medications & Assessment
- Nursing Care
- Optimize Environment

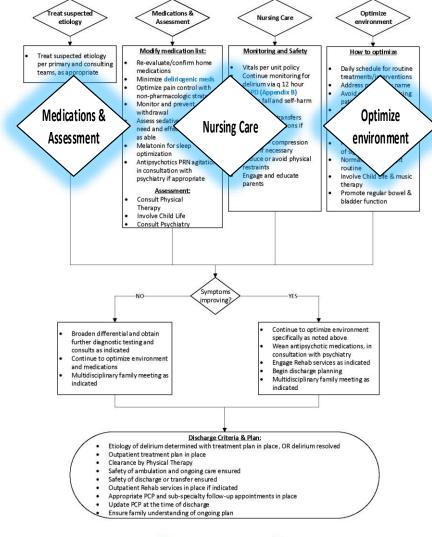
We will discuss these strategies more in depth in later slides.



CLINICAL PATHWAY:

Delirium - Inpatient and Zone C Management

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1. Management

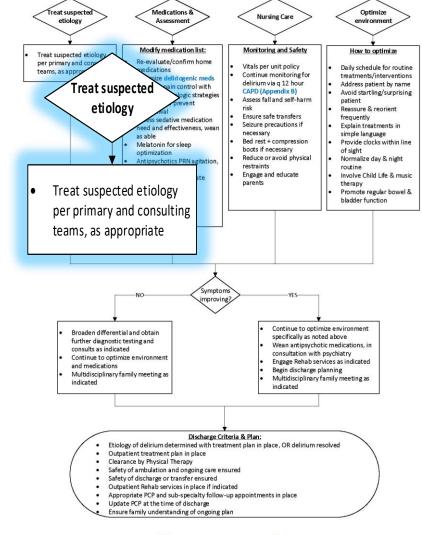
It is always important to assess for the most likely etiology of delirium.

Be sure to involve any consulting teams as appropriate.

CLINICAL PATHWAY:

Delirium - Inpatient and Zone C Management











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AST UPDATED: 02:02:21



1. Management

Because certain medications can contribute to delirium, it is important to re-evaluate medications, and minimize any deliriogenic medications the patient is on.

Clicking on "deliriogenic meds" will bring you to a list of medications listed in the Inpatient Prevention portion.

Psychiatry may assist with treatment of agitation.

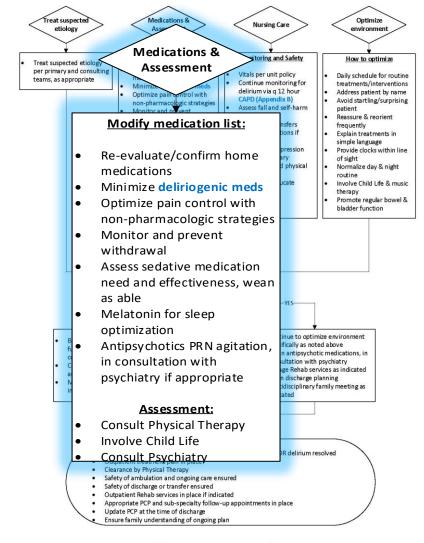
Physical therapy and Child life should become involved as early as it is safe to do so.

Child life is helpful for creating a functional plan to help normalize day time and night time routines.

CLINICAL PATHWAY:

Delirium - Inpatient and Zone C Management











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1. Management

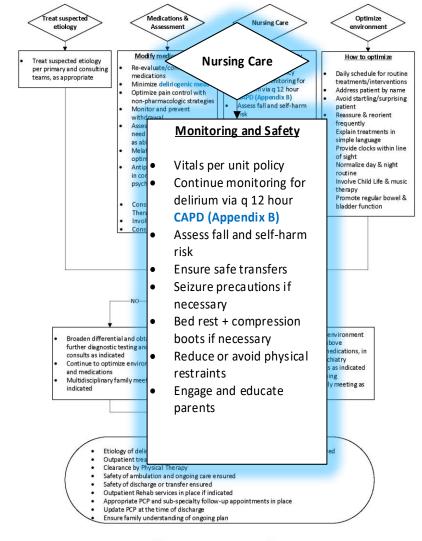
Continued and regular assessment of delirium is very important to assess for improvement or worsening.

Modified nursing care and safety monitoring are a vital part of the management plan.

CLINICAL PATHWAY:

Delirium - Inpatient and Zone C Management











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1. Management

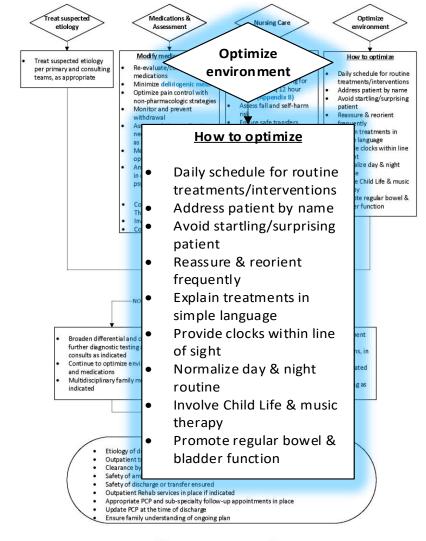
Optimizing the environment to help re-orient the child to their surroundings can help improve delirium.

Having a daily schedule, providing clocks, and decreasing potential stressors are all examples.

CLINICAL PATHWAY:

Delirium - Inpatient and Zone C Management











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1. Management

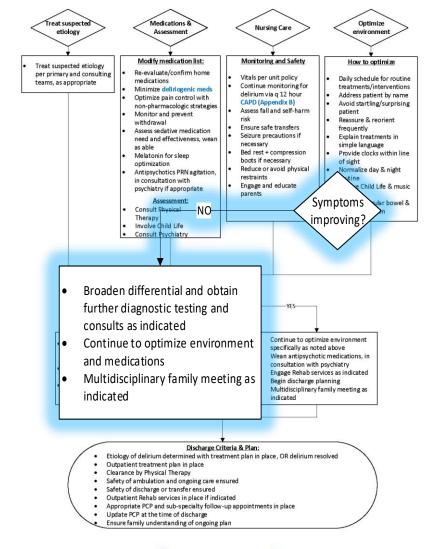
If symptoms are not improving, the differential should be broadened to further assess for a potential etiology.

Optimization of the environment and the patient's medication should be ongoing during this time.

CLINICAL PATHWAY:

Delirium - Inpatient and Zone C Management











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1. Management

If symptoms of delirium are improving, management strategies to continue while planning towards discharge.

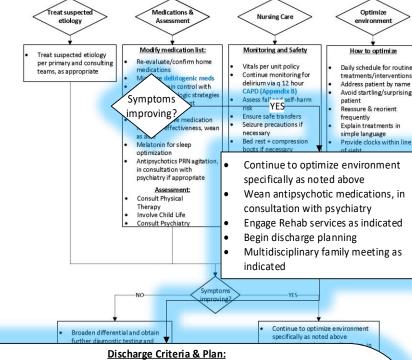
> Depending on the circumstance, a multidisciplinary family meeting may be necessary.

The patient must have specific criteria met in order to be discharged – specifically, delirium should have resolved (or a treatment plan is in place for etiologies that have been determined).

CLINICAL PATHWAY:

Delirium - Inpatient and Zone C Management





- Etiology of delirium determined with treatment plan in place, OR delirium resolved
- Outpatient treatment plan in place
- Clearance by Physical Therapy
- Safety of ambulation and ongoing care ensured
- Safety of discharge or transfer ensured
- Outpatient Rehab services in place if indicated
- Appropriate PCP and sub-specialty follow-up appointments in place
- Update PCP at the time of discharge
- Ensure family understanding of ongoing plan
 - Ensure family understanding of ongoing plan







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- 1. Prevention and Identification
- 2. Evaluation and Work up

The Delirium – Inpatient Prevention and Screening algorithm is meant for all patients admitted on the Med/Surg units at CT Children's, not just for those with suspected or known delirium.

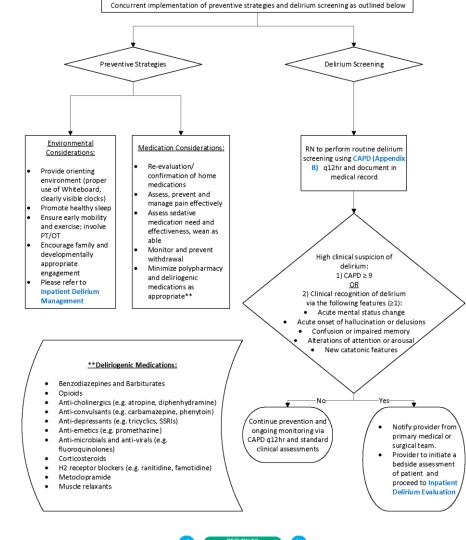
Of note:

PICU patients should be excluded from this pathway. They are being screened with the CAPD score, but providers and nurses should follow the specific protocols for screening, prevention and treatment for PICU patients.

CLINICAL PATHWAY:

Delirium - Inpatient Prevention and Screening

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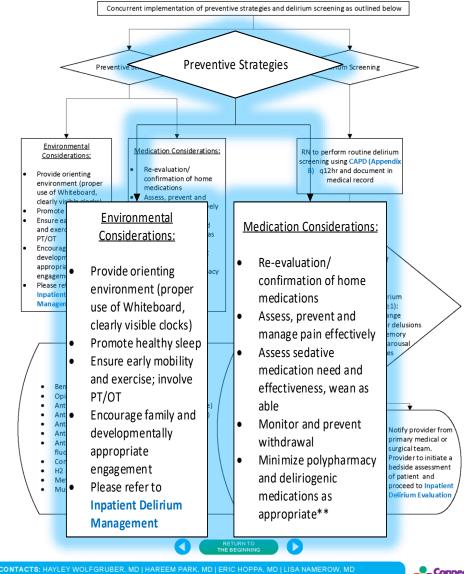
- 1. Prevention and Identification
- 2. Evaluation and Work up

Prevention is key!

Proactive measures to prevent delirium include both environmental and medication considerations.

CLINICAL PATHWAY:

Delirium - Inpatient Prevention and Screening





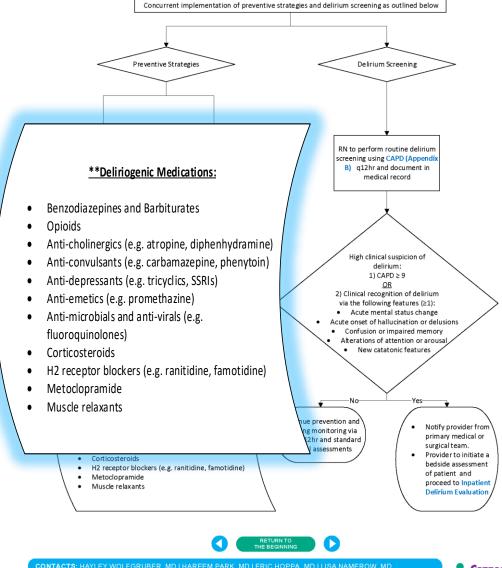
- 1. Prevention and Identification
- 2. Evaluation and Work up

Deliriogenic medications are listed and should be reviewed for every patient presenting with delirium.

CLINICAL PATHWAY:

Delirium - Inpatient Prevention and Screening

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- 1. Prevention and Identification
- 2. Evaluation and Work up

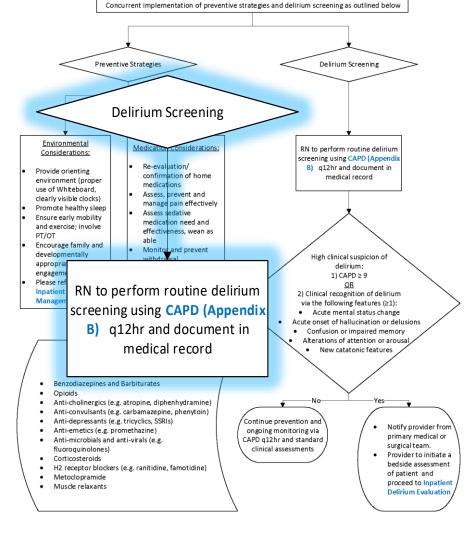
All med/surg patients will be screened with the CAPD tool.

Nursing will complete the screening about every 12 hours. The screen will occur towards the end of the shift to capture the "overall assessment" or average behavior. This is NOT a "moment in time" assessment.

CLINICAL PATHWAY:

Delirium - Inpatient Prevention and Screening

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- 1. Prevention and Identification
- 2. Evaluation and Work up

CAPD screening tool (Cornell Assessment of Pediatric Delirium)

- Validated for patients 0-21 yrs
- Easy to use
- Can trend over time
- Based on developmental anchor points for patients
 years old or developmentally delayed
- Detects hypoactive and hyperactive forms of delirium

In developmentally normal children, CAPD sensitivity 92% and specificity 86.5% In developmentally delayed children, CAPD sensitivity 96% and specificity 51%

CLINICAL PATHWAY:

Delirium - Inpatient Prevention and Screening

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Concurrent implementation of preventive strategies and delirium screening as outlined below

Figure 1. Cornell Assessment of Pediatric Delirium (CAPD) revised							
RASS Score (if -4 or -5 do not proceed)							
Please answer the following questions based on your interactions with the patient over the coryour shift:						ırse of	
	Never	Rarely	Sometimes	Often	Always	Score	
	4	3	2	1	0		
1. Does the child make eye contact with the caregiver?							
2. Are the child's actions purposeful?							
3. Is the child aware of his/her surroundings?							
4. Does the child communicate needs and wants?							
	Never	Rarely	Sometimes	Often	Always		
	0	1	2	3	4		
5. Is the child restless?							
6. Is the child inconsolable?							
7. Is the child underactive—very little movement while awake?							
8. Does it take the child a long time to respond to interactions?							
					TOTAL		

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- 1. Prevention and Identification
- 2. Evaluation and Work up

CAPD uses <u>Developmental</u> <u>Anchor Points</u>

- Anchor points are a reference for normative behaviors based on age/developmental level
- Used for patients < 2 years of age (and/or of that developmental level)
- Observable behaviors as they would be seen in hospital setting
- Adjusted for alterations by "sick behavior," pain, anxiety, and developmental delay

CLINICAL PATHWAY: Delirium - Inpatient Prevention and Screening

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

2 years

C		and and balance
Concurrent implementation of	preventive strategies and delirium screening	as outlined below

	NB	4 weeks	o weeks	o weeks	Zo weeks	1 year	Z years
I. Does the child make eye contact with the caregiver?	Fixates on face	Holds gaze briefly Follows 90 degrees	Holds gaze	Follows moving object/caregiver past midline, regards examiner's hand holding object, focused attention	Holds gaze. Prefers primary parent. Looks at speaker.	Holds gaze. Prefers primary parent. Looks at speaker.	Holds gaze. Prefers primary parent. Looks at speaker
2. Are the child's actions purposeful?	Moves head to side, dominated by primitive reflexes	Reaches (with some discoordination)	Reaches	Symmetric movements, will passively grasp handed object	Reaches with coordinated smooth movement	Reaches and manipulates objects, tries to change position, if mobile may try to get up.	Reaches and manipulates objects, tries to change position, if mobile may try to get up and walk
3. Is the child aware of his/her surroundings?	Calm awake time	Awake alert time Turns to primary caretaker's voice May turn to smell of primary care taker	Increasing awake alert time Turns to primary caretaker's voice May turn to smell of primary care taker	Facial brightening or smile in response to nodding head, frown to bell, coos	Strongly prefers mother, then other familiars. Differentiates between novel and familiar objects	Prefers primary parent, then other familiars, upset when separated from preferred care takers. Comforted by familiar objects especially favorite blanket or stuffed animal	Prefers primary parent, then other familiars, upset when separated from preferred care takers. Comforted by familiar objects, especially favorite blanket or stuffed animal
4. Does the child communicate needs and wants?	Cries when hungry or uncomfortable	Cries when hungry or uncomfortable	Cries when hungry or uncomfortable	Cries when hungry or uncomfortable	Vocalizes /indicates about needs, e.g., hunger, discomfort, curiosity in objects, or surroundings	Uses single words or signs	3 to 4 word sentences, or signs. May indicate toilet needs, calls self or me
5. Is the child restless?	No sustained awake alert state	No sustained calm state	No sustained calm state	No sustained awake alert state	No sustained calm state	No sustained calm state	No sustained calm state
6. Is the child inconsolable?	Not soothed by parental rocking, singing, feeding, comforting actions	Not soothed by parental rocking, singing, feeding, comforting actions	Not soothed by parental rocking, singing, feeding, comforting actions	Not soothed by parental rocking, singing, comforting actions	Not soothed by usual methods, e.g., singing, holding, talking	Not soothed by usual methods, e.g., singing, holding, talking, reading	Not soothed by usual methods, e.g., singing, holding, talking, reading (may tantrum, but can organize)
7. Is the child underactive—very little movement while awake?	Little if any flexed and then relaxed state with primitive reflexes (Child should be sleeping comfortably most of the time)	Little if any reaching, kicking, grasping (still may be somewhat discoordinated)	Little if any reaching, kicking, grasping (may begin to be more coordinated)	Little if any purposive grasping, control of head and arm movements, such as pushing things that are noxious away	Little if any reaching, grasping, moving around in bed, pushing things away	Little if any play, efforts to sit up, pull up, and if mobile crawl or walk around	Little if any more elaborate play, efforts to sit up and move around, and if able to stand, walk, or jump
8. Does it take the child a long time to respond to interactions?	Not making sounds or reflexes active as expected (grasp, suck, moro)	Not making sounds or reflexes active as expected (grasp, suck, moro)	Not kicking or crying with noxious stimuli	Not cooing, smiling, or focusing gaze in response to interactions	Not babbling or smiling/laughing in social interactions (or even actively rejecting an interaction)	Not following simple directions. If verbal, not engaging in simple dialogue with words or jargon	Not following I-2 step simple commands. If verbal, not engaging in more complex dialogue

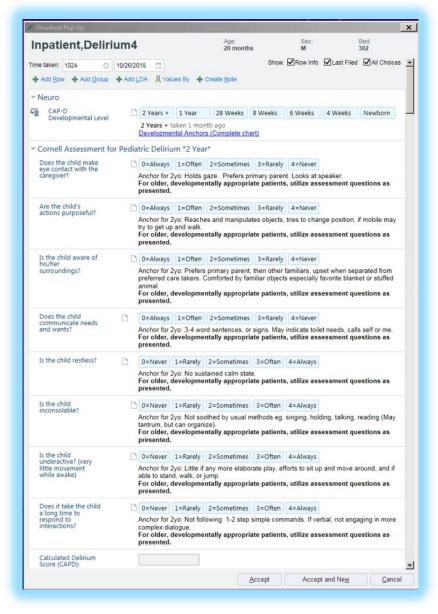
8 weeks

4 weeks

The CAPD: Documenting it in the chart

- CAPD Screening Tool is a screening tool based on these 8 questions, answered based on observed patient behaviors over the course of the shift and reflective of their current developmental level
- Scoring will be completed by nursing twice daily, ideally towards the end of their shift
- Providers may be asked by nursing to help answer some questions in the tool that they are having trouble evaluating (Can be completed in a team approach for a patient that is difficult to assess)
- Parents may also be a resource to help answer these questions based on parents observation, comparing to baseline behaviors





The CAPD: Where is it in the Chart?





On the Vital Signs screen listed under the vitals signs

Cornell Assessment of Pediatric Delirium CAP-D Score CAP-D Score

Neuro

You can also add a column for the CAPD to "My List" for easy viewing when looking at your patient list

Additional Documentation Under the flowsheet "Pedi A&I"

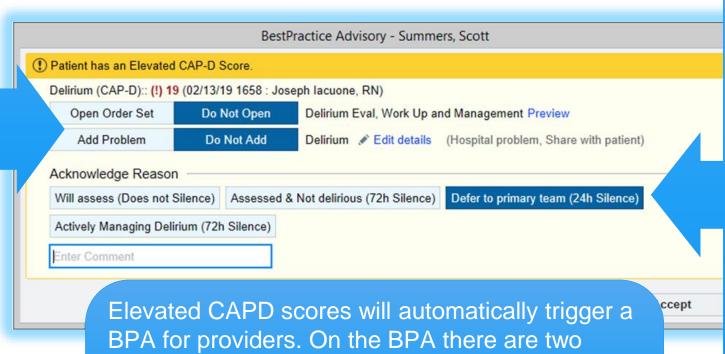
Screening Assessments CAP-D Developmental Level State Behavioral Scale (PICU) Cornell Assessment for Pediatric Delirium Does the child make eye contact with Are the child's actions purposeful? Is the child aware of his/her Does the child communicate needs and Is the child restless? Is the child inconsolable? Is the child underactive? (very little Does it take the child a long time to Calculated Delirium Score (CAPD):

The CAPD: Best Practice Alerts (BPA)

sections.

By selecting to open the order set or add the problem, you are saying that you are performing the appropriate actions.

This means that you DO NOT have to select an acknowledgement reason below



On the top you can Open the Order Set and

If you do neither you will need to chose a

reason why on the bottom "Acknowledge

Add Delirium as a Problem

reason section"



The Acknowledge Reason section should be used when you do not want to perform one of the above two actions.

- Actively Managing
 Delirium could be used
 when you have already
 placed orders and
 added the problem but it
 has been 72 hours and
 the patient is still getting
 an elevated score.
- When you select one of the acknowledge reasons the top two actions will automatically change to "Do Not...".

- 1. Prevention and Identification
- 2. Evaluation and Work up

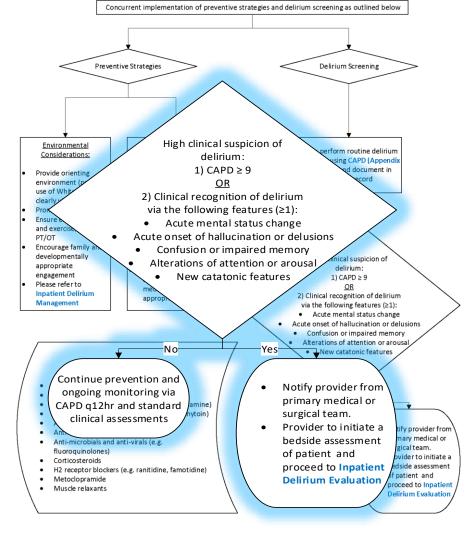
If there is a high clinical suspicion of delirium, proceed to the Inpatient Delirium Evaluation and Work Up.

If not, continual assessment, and optimization of environment/medications should occur.

CLINICAL PATHWAY:

Delirium - Inpatient Prevention and Screening

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









CONTACTS: HAYLEY WOLFGRUBER, MD | HAREEM PARK, MD | ERIC HOPPA, MD | LISA NAMEROW, MD

LAST UPDATED: 02.02.21

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- 1. Prevention and Identification
- 2. Evaluation and Work up

Once a patient has been identified as having delirium due to clinical presentation and/ or elevated CAPD score, the primary provider should perform a bedside assessment of the patient.

Notify the primary attending if a patient is confirmed to have delirium based on the bedside evaluation.

Inpatient delirium management should occur simultaneously as the work up. Management is the same as Zone C management – as previously discussed.

CLINICAL PATHWAY:

Delirium – Inpatient Evaluation and Work Up

THIS PATHWAY
SERVES AS A GUID!
AND DOES NOT
REPLACE CLINICAL

Provider bedside evaluation of patient.
 Initiation of inpatient Delirium Management while evaluation and workup is occurring simultaneously.

- Provider bedside evaluation of patient.
- Initiation of Inpatient Delirium Management while evaluation and workup is occurring simultaneously.

Labo:

Primary Work, Up

Labo:

Stat chem 10, CBC, CRP, ESR, ammonia, PT/PTT/INR, TSH, free T4, VBG or CBG, AS
ALT, ECH level, AVA

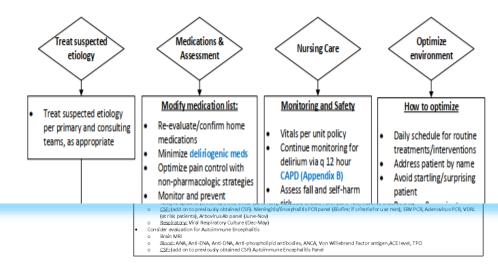
Urine:

Toxicology screen imaging:

CLINICAL PATHWAY:

Delirium - Inpatient and Zone C Management

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









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- 1. Prevention and Identification
- 2. Evaluation and Work up

If etiology not clear, work up should follow a tiered evaluation including:

- Lab testing
- Imaging
- Consult services

Overall evaluation and escalation of work up should involve a *multidisciplinary team* approach

Non pharmacologic interventions should start as soon as delirium identified

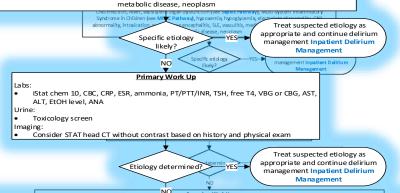
CLINICAL PATHWAY:

Delirium – Inpatient Evaluation and Work Up

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

tiologies to consider:

CNS infection, fewer, sepsis/end organ dysfunction (see Sepsis Pathway), Multi-system Inflammatory Syndrome in Children (see MIS-C Pathway), hypoxemia, hypoglycemia, electrolyte abnormality, CNS abnormality, intoxication, autoimmune encephalitis, SLE, vasculitis, medication effect, drug withdrawal,



Secondary Work Up

If febrile:

- Blood and urine cultures
- Strongly consider LP: cell count with differential, protein, glucose, gram stain and culture, HSV PCR, enterovirus PCR, opening pressure. Ask lab to hold 3 mL CSF for further studies.
- Begin all empiric IV antimicrobials listed below:
 - Ceftriaxone IV 100 mg/kg/day q12hr (max 2,000 mg/dose) x48 hours AND
 - Vancomycin IV x48 hours:
 - <52 weeks PMA[‡]/about <3 mo old: 15 mg/kg q8hr or as determined by pharmacy based on estimated AUC</p>
 - ≥52 weeks PMA[‡]/about ≥3 months old 11 years old: 70 mg/kg/day div q6hr
 - ≥12 yrs old: 60 mg/kg/day div q8hr AND
 - Acyclovir 20 mg/kg/dose IV q8hr until HSV studies negative

Consider following consultations (who may recommend further work up)

- Neurology (if concern for seizure, abnormal EEG, movement disorder, abnormal neurological imaging or focal deficit, or other neurologic diagnosis)
- Rheumatology (if autoimmune process suspected)
- <u>Psychiatry</u> (to assist with recognition/diagnosis of delirium utilizing the Vanderbilt Assessment for Delirium in Infants and Children (VADIC)
 assessment tool Appendix A; determine/confirm etiology; assist with pharm + non-pharmacological management; help with ongoing monitoring/
 response to therapies; for ongoing co-management)
- If diagnosis or treatment plan involves multidisciplinary approach, strongly consider family meeting.

*PMA (Post-Menstrual Age) = gestational age + postnatal age



Tertiary Work Up:

- Consult Infectious Disease
- Infectious Encephalitis Panel
- Blood: Mycoplasma IgM/igG, bartonella IgM/igG, lyme IgM/igG, West Nile IgM/igG (June-Nov), Anaplasma Phagocytophilium IgG/igM (June-Nov), Anaplasma (Ehrlichia) blood smear (June-Nov), Rickettsial Disease Panel (June-Nov, travel to endemic area)
- CSE: (add on to previously obtained CSF). Meningitis/Encephalitis PCR panel (Biofire; if criteria for use met), EBV PCR, Adenovirus PCR, VDRL (at risk patients), Arbovirus Ab panel (June-Nov)
- Respiratory: Viral Respiratory Culture (Dec-May)
- Consider evaluation for Autoimmune Encephalitis
 - Brain N
 - Blood: ANA, Anti-ENA, Anti-DNA, Anti-phospholipid antibodies, ANCA, Von Willebrand Factor antigen, ACE level, TPO
 - o CSE: (add on to previously obtained CSF) Autoimmune Encephalitis Panel



- 1. Prevention and Identification
- 2. Evaluation and Work up

Primary work up is intended to screen for easily identifiable sources of delirium.

As soon as an etiology is positively identified, it should be treated as appropriate, while continuing to manage delirium.

CLINICAL PATHWAY:

Delirium – Inpatient Evaluation and Work Up

THIS PATHWAY SERVES AS A GUID! AND DOES NOT REPLACE CLINICAL IUDGMENT

Provider bedside evaluation of patient.
Initiation of Inpatient Delirium Management while evaluation and workup is occurring simultaneously.

Etiologies to consider:

CNS infection, fever, sepsis/end organ dysfunction (see Sepsis Pathway), Multi-system Inflammatory Syndrome in Children (see MIS-C Pathway), hypoxemia, hypoglycemia, electrolyte abnormality, CNS abnormality, intoxication, autoimmune encephalitis, SLE, vasculitis, medication effect, drug withdrawal, metabolic disease, neoplasm



<u>Primary Work Up</u>

Labs:

• iStat chem 10, CBC, CRP, ESR, ammonia, PT/PTT/INR, TSH, free T4, VBG or CBG, AST, ALT, EtOH level, ANA

Urine:

Toxicology screen

Imaging:

Consider STAT head CT without contrast based on history and physical exam





Consult Infectious Disease Infectious Encephalitis Panel:

- Blood: Mycoplasma ig M/igG, bartonella ig M/igG, lyme ig M/igG, West Nile ig M/igG (June-Nov), Anaplasma Phag ocytophilium ig G/igM (June-Nov), Anaplasma (Ehrlichia) blood smear (June-Nov), Rickettsial Disease Panel (June-Nov, travel to endemic area)
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- 1. Prevention and Identification
- 2. Evaluation and Work up

If febrile, further evaluation (including an LP) and empiric antimicrobials is warranted. Specialists may be consulted depending on specific concerns.

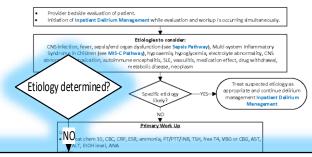
When performing the LP, please send as much CSF possible to the lab to be saved for potential future studies.

- * Minimum of 3ml of CSF should be saved, but as much as 6ml may be needed for some panels.
- * Please call the lab to confirm CSF is being held

CLINICAL PATHWAY:

Delirium – Inpatient Evaluation and Work Up

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



Secondary Work Up

If febrile:

- Blood and urine cultures
- Strongly consider LP: cell count with differential, protein, glucose, gram stain and culture, HSV PCR, enterovirus PCR, opening pressure. Ask lab to hold 3 mL CSF for further studies.
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- If diagnosis or treatment plan involves multidisciplinary approach, strongly consider family meeting.

[†]PMA (Post-Menstrual Age) = gestational age + postnatal age



RETURN TO





- 1. Prevention and Identification
- 2. Evaluation and Work up

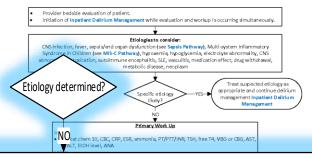
The pharmacy's vancomycin protocol was updated in Feb 2021.

- All patients who have vancomycin IV ordered will be followed by the clinical pharmacist to help determine appropriate dosing parameters.
- Providers will order initial doses per pathway/order set and provide indication within the order.
- IV vancomycin dosing and recommended labs will be managed by pharmacy in conjunction with primary teams.

CLINICAL PATHWAY:

Delirium – Inpatient Evaluation and Work Up

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



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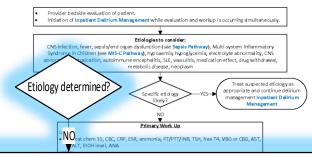
- 1. Prevention and Identification
- 2. Evaluation and Work up

Note that the VADIC assessment tool will again be used by Psychiatry to provide consistent standardized assessment of patients with concern for Delirium

CLINICAL PATHWAY:

Delirium – Inpatient Evaluation and Work Up

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RETURN TO THE BEGINNING





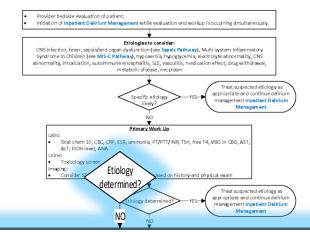


- 1. Prevention and Identification
- 2. Evaluation and Work up

CLINICAL PATHWAY:

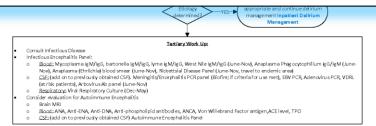
Delirium – Inpatient Evaluation and Work Up

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



Tertiary Work Up:

- Consult Infectious Disease
- Infectious Encephalitis Panel:
 - Blood: Mycoplasma IgM/IgG, bartonella IgM/IgG, lyme IgM/IgG, West Nile IgM/IgG (June-Nov), Anaplasma Phagocytophilium IgG/IgM (June-Nov), Anaplasma (Ehrlichia) blood smear (June-Nov), Rickettsial Disease Panel (June-Nov, travel to endemic area)
 - <u>CSF</u>: (add on to previously obtained CSF). Meningitis/Encephalitis PCR panel (Biofire; if criteria for use met), EBV PCR, Adenovirus PCR, VDRL (at risk patients), Arbovirus Ab panel (June-Nov)
 - Respiratory: Viral Respiratory Culture (Dec-May)
- Consider evaluation for Autoimmune Encephalitis
 - Brain MRI
 - Blood: ANA, Anti-ENA, Anti-DNA, Anti-phospholipid antibodies, ANCA, Von Willebrand Factor antigen, ACE level, TPO
 - CSE: (add on to previously obtained CSF) Autoimmune Encephalitis Panel









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AST LIDDATED: 02.02.21

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Order Set

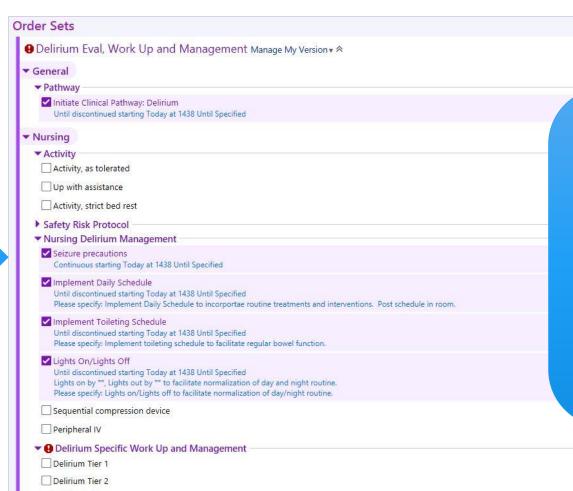


Nursing orders are prefilled out and preselected for ease of ordering

Notice the orders are broken down into Tiers

Delirium Tier 3

Medications



There are two order sets for inpatient use:

- 1. Admit to MS Delirium And
- 2. Delirium Evaluation, Work up, and Management

Either can be used at any time, but the second is meant for patients already admitted.

Review of Key Points



- Pediatric delirium is an under-recognized and high-risk diagnosis in pediatric patients that can lead to several complications
- Delirium is a condition caused by a medical etiology, it is not a psychiatric illness
- Many factors contribute to the development of delirium, including underlying illness, medications and disruption of normal routine
- CAPD screening tool can help earlier identify patients with delirium in the inpatient and ICU setting
- New Clinical Pathways for Pediatric Delirium Evaluation, Work-up and Management provides a consistent approach to preventing, screening, evaluating, and managing delirium

Quality Metrics



- Percentage of patients on medical surgical units who were not screened with the CAPD
- Percent of patients who were screened with CAPD tool twice daily
- Percent of patients with CAPD score ≥ 9 with delirium pathway order set usage
- Average time from CAPD score ≥ 9 to the initiation of the delirium pathway order set
- Number of PICU transfers following CAPD score ≥ 9
- Number of MET activations following CAPD score ≥ 9
- Percent of patients with CAPD score ≥ 9 who have delirium ICD-10 codes applied
- Percent of patients with CAPD score ≥ 9 who have a psychiatry evaluation
- Percent of patients with CAPD score ≥ 9 who have a CT scan
- ALOS for patients with a CAPD score ≥ 9 (days)

Pathway Contacts



- Hayley Wolfgruber, MD
 - o Pediatric Hospital Medicine
- Hareem Park, MD
 - Pediatric Hospital Medicine
- Eric Hoppa, MD
 - Pediatric Emergency Medicine
- Jennifer Downs, MD
 - Child Psychiatry

References



- Schieveld JNM, Janssen NJJF. "Delirium in the Pediatric Patient: On the Growing Awareness of Its Clinical Interdisciplinary Importance." JAMA Pediatrics 2014; 168 (7): 595-596
- Moldonado JR. "Delirium pathophysiology: An updated hypothesis of the etiology of acute brain failure." Int J Geriatr Psychiatry 2017; doi: 10.1002/gps.4823
- Cerejeira J, Firmino H, Vaz-Serra A et al. "The neuroinflammatory hypothesis of delirium." Acta Neuropathol 2010; 119: 737-754
- American Psychiatric Association. Diagnostic and Statistical Manual of Mental Disorders. Fifth Edition. Washington DC: American Psychiatric Publishing 2013
- Bettencourt, A., and Mullen, J. "Delirium in children: Identification, Prevention, and Management." Critical Care Nurse. 2017; 37(3).
- Traube C, Silver G, Reeder R, et al. "Delirium in critically ill children: An international point prevalence study." Critical Care Medicine 2017; 45: 584-590
- Traube C, Silver G, Gerber LM, et al. "Delirium and Mortality in Critically III Children: Epidemiology and Outcomes of Pediatric Delirium." Critical Care Medicine 2017; 45(5): 891-898
- Smith HAB, Berutti T, Brink E, Strohler B, Fuchs DC, Ely EW, et al. "Pediatric Critical Care Perceptions on Analgesia, Sedation and Delirium." Semin Respir Crit Care Med 2013; 34:244-261
- Silver G, Kearney J, Traube C, et al. "Pediatric delirium: Evaluating the gold standard." Palliat Support Care 2015; 13(3): 513-51
- Silver G, Traube C, Gerber LM, et al. "Pediatric Delirium and Associated Risk Factors: A Single-Center Prospective Observational Study." Pediatr Crit Care Med 2015; 16(4): 303-309
- Traube C, Silver G, Kearney J, et al. "Cornell Assessment of Pediatric Delirium: A Valid, Rapid, Observational Tool for Screening Delirium in the PICU." Pediatr Crit Care Med 2014; 42(3): 656-663
- Traube C, Mauer EA, Gerber LM, et al. "Cost Associated with Pediatric Delirium in the ICU." Crit Care Med 2016; 44(12): e1175-e1179
- Turkel SB. "Pediatric Delirium: Recognition, Management, and Outcome." Curr Psychiatry Rep 2017; 19(12): 101. doi: 10.1007/s11920-017-0851-1
- Smith HA, Boyd J, Fuchs DC, et al. "Diagnosing delirium in critically ill children: Validity and reliability of the Pediatric Confusion Assessment Method for the Intensive Care Unit". Critical Care Medicine 2011; 39(1): 150
- Smith HAB, Gangopadhyay M, Goben CM, et al. "The Preschool Confusion Assessment Method for the ICU: Valid and Reliable Delirium Monitoring for Critically III Infants and Children." Pediatr Crit Care Med 2016; 44(3): 592-600
- Silver G, Kearney J, Traube C, et al. "Delirium screening anchored in child development: The Cornell Assessment for Pediatric Delirium." Palliat Support Care 2015; 13(4): 1005-1011
- Silver GH, Kearney JA, Bora S, et al. A Clinical Pathway to Standardize Care of Children with Delirium in Pediatric Inpatient Settings. Hosp Pediatr. 2019 Nov;9(11):909-916.
- Malas N, Brahmbhatt K, McDermott C, et al. "Pediatric Delirium: Evaluation, Management and Special Considerations." Curr Psychiatry Rep 2017; 19(9): 65
- Foster, J, et al. "Melatonin and melatonin agonists to prevent and treat delirium in critical illness: a systematic review protocol." Systematic Reviews 2016. 5:199
- Hunter A, Johnson L, Coustasse A. "Reduction of intensive care unit length of stay: The case of early mobilization." The Health Care Manager 2014; 33(2):

Thank You!



About Connecticut Children's Clinical Pathways Program

The Clinical Pathways Program at Connecticut Children's aims to improve the quality of care our patients receive, across both ambulatory and acute care settings. We have implemented a standardized process for clinical pathway development and maintenance to ensure meaningful improvements to patient care as well as systematic continual improvement. Development of a clinical pathway includes a multidisciplinary team, which may include doctors, advanced practitioners, nurses, pharmacists, other specialists, and even patients/families. Each clinical pathway has a flow algorithm, an educational module for end-user education, associated order set(s) in the electronic medical record, and quality metrics that are evaluated regularly to measure the pathway's effectiveness. Additionally, clinical pathways are reviewed annually and updated to ensure alignment with the most up to date evidence. These pathways serve as a guide for providers and do not replace clinical judgment.