Clinical Pathways

Diabetes Insipidus (DI) Post-operative Neurosurgical Management

Cem Demirci, MD
Rebecca Riba-Wolman, MD
David Hersh, MD
Jonathan Martin, MD
Elliot Melendez, MD









What is a Clinical Pathway?



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

Background



- Diabetes insipidus (DI) refers to the passage of large volumes of dilute urine and may result from decreased secretion of antidiuretic hormone (ADH) by the posterior pituitary gland.
- Patients undergoing surgery in the sellar or parasellar region are at risk for postoperative DI, which may be transient, triphasic [DI > SIADH > DI], or permanent.
- Patients without an intact thirst mechanism (adipisic central DI) are a particular challenge, as they may not drink enough to replace their urine losses, resulting in severe hypernatremia.

Why is the DI Pathway Necessary?



- Uncontrolled hypernatremia has adverse effects, including an increased risk of neurological sequela and venothromboembolism
- Provider variability and inconsistent care delivery/monitoring are barriers to establish diagnosis and deliver timely and effective care in the absence of a standardized protocol

Objectives of the DI Pathway



- Standardize the management of postoperative patients at risk for developing DI
 - Initial PICU monitoring for development of DI
 - Initial PICU management if DI develops
 - Standardized clearance for patient's transfer to med/surg floors
- Standardize the management of post-operative patients with confirmed DI in the PICU and on the floors
 - Minimize fluctuations in sodium level and volume status
 - Expedite the development of an outpatient plan in order to facilitate a safe discharge to home

Pathway Overview



- This is the Diabetes
 Insipidus (DI) Post-operative
 Neurosurgical Management
 Clinical Pathway.
- There are 3 portions of the pathway:
 - PICU Post-operative Monitoring for DI
 - PICU Management of DI
 - Med/Surg Management of DI
- We will be reviewing each component in the following slides.

Page 1

CLINICAL PATHWAY:

Diabetes Insipidus (DI) - Post-operative Neurosurgical ManagementPICU Post-operative Monitoring for DI

Page 2

CLINICAL PATHWAY:

Diabetes Insipidus (DI) - Post-operative Neurosurgical ManagementPICU Management of DI

Page 3

CLINICAL PATHWAY:

Diabetes Insipidus (DI) - Post-operative Neurosurgical ManagementMed/Surg Management of DI

CLINICAL PATHWAY:
Diabetes Insipidus (DI) Post-operative Neurosurgical Management
PICU Post-operative Monitoring for DI

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

Pre-Operative
Risk Factors for
DI:
Hx of nocturia,
enuresis

Indusion Criter

Indusion Criter

Admit

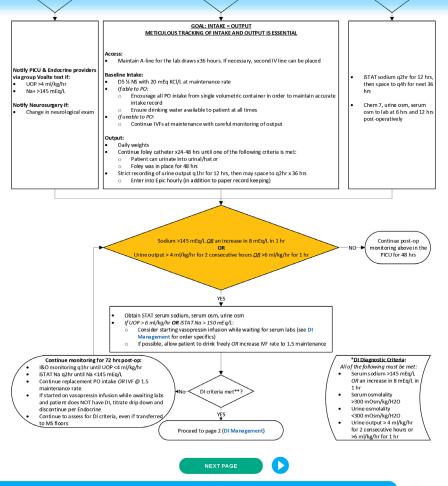
Indusion Criteria: Patients ≥ 1 year old undergoing surgery involving the sellar or parasellar regions of the brain Exclusion Criteria: Age < 1 year, acute kidney injury or chronic kidney disease

Record fluid management intra-operatively

Because any surgical procedure that involves the sellar or parasellar regions of the brain can increase the risk of DI development, any child that is ≥1 year of age that has such a procedure will be monitored for the development of DI post-operatively.

Of note, those with any acute kidney injury or chronic kidney disease are excluded from the pathway.

Inclusion Criteria: Patients ≥ 1 year old undergoing surgery involving the sellar or parasellar regions of the brain Exclusion Criteria: Age < 1 year, acute kidney injury or chronic kidney disease



CLINICAL PATHWAY:
Diabetes Insipidus (DI) Post-operative Neurosurgical Management
PICU Post-operative Monitoring for DI

ndusion Criteria: Patients ≥ 1 year old undergoing surgery involving the sellar or parasellar regions of the brain

Exclusion Criteria: Age < 1 year, acute kidney injury or chronic kidney disease

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

All patients with sellar/parasellar surgery will be admitted to the PICU for monitoring of DI development.

If DI is noted immediately post-op, DI management should be followed.

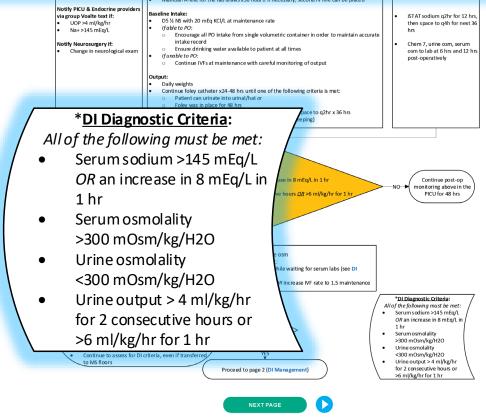
DI criteria are noted here.

Note that all of the criteria must be met, which includes serum sodium, serum osmolality, urine osmolality, and urine output. These parameters will be assessed frequently.

- Record fluid management intra-operatively
- Admit to PICU post-operatively and follow care below for 48 hrs

Risk Factors for DI: Hx of nocturia,

- If DI criteria** present immediately post-operatively, proceed to page 2 (DI management)
- Neurosurgery to stress dose hydrocortisone post-operatively; wean per endocrine until cosyntropin stim test performed







The most important aspect of DI is to ensure intake = output. Meticulous tracking of I&Os is essential.

Intake:

- All should be started on maintenance IVF.
- In order to maintain accurate intake, PO from one container is encouraged, and patient should have water available at all times.

Output:

- Foley catheter is maintained for 24-48
 hours until the patient can urinate or the
 foley has been in for 48 hours.
- Strict recording of output is needed every 1 hour for 12 hours, and then every 2 hours for 36 hours thereafter.



GOAL: INTAKE = OUTPUT METICULOUS TRACKING OF INTAKE AND OUTPUT IS ESSENTIAL

Access:

• Maintain A-line for the lab draws x36 hours. If necessary, second IV line can be placed

Baseline Intake:

- D5 ½ NS with 20 mEq KCI/L at maintenance rate
- If able to PO:
 - Encourage all PO intake from single volumetric container in order to maintain accurate intake record
 - Ensure drinking water available to patient at all times
- If unable to PO:
 - o Continue IVFs at maintenance with careful monitoring of output

Output:

- Daily weights
- Continue foley catheter x24-48 hrs until one of the following criteria is met:
 - Patient can urinate into urinal/hat or
 - o Foley was in place for 48 hrs
- Strict recording of urine output q1hr for 12 hrs, then may space to q2hr x 36 hrs
 - Enter into Epic hourly (in addition to paper record keeping)





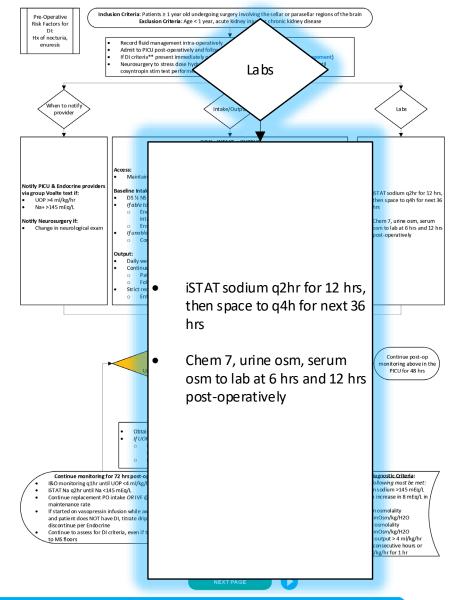


Laboratory monitoring is also essential to ensure that DI has not developed.

Recommendations are listed here.

CLINICAL PATHWAY: Diabetes Insipidus (DI) Post-operative Neurosurgical Management PICU Post-operative Monitoring for DI

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

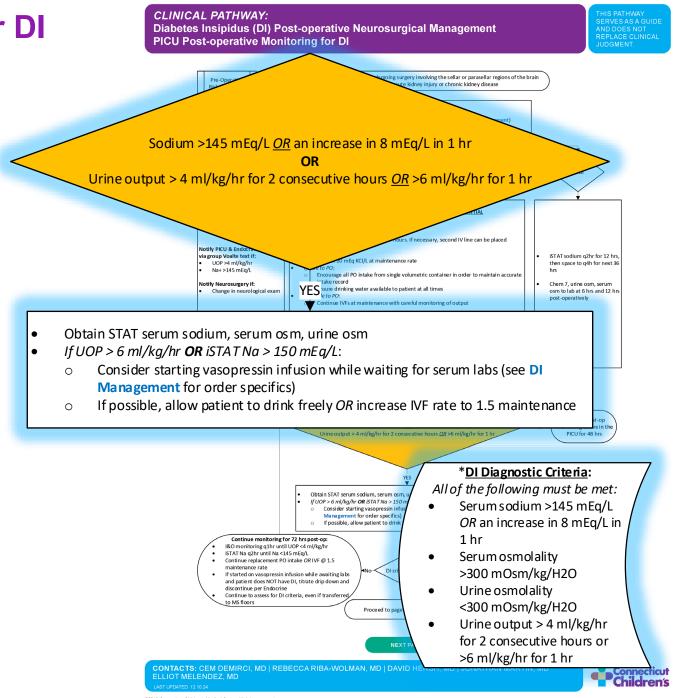


CONTACTS: CEM DEMIRCI, MD | REBECCA RIBA-WOLMAN, MD | DAVID HERSH, MD | JONATHAN MARTIN, MD ELLIOT MELENDEZ, MD



©2019 Connecticut Children's Medical Center. All rights reserved.

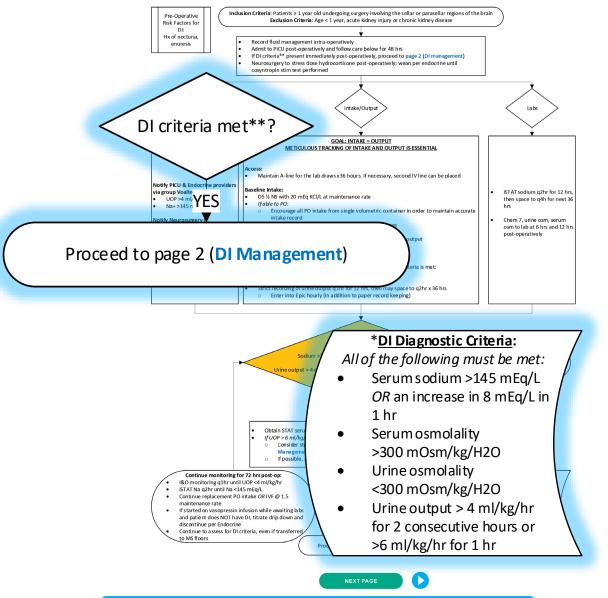
- If sodium is >145 mEq/L (or an increase by 8 mEq/L in 1 hr)
 OR there is urine output that is >4 ml/kg/hr for 2 consecutive hours (or >6 ml/kg/hr for 1 hr), you MUST obtain stat labs to evaluate if DI is present.
- If UOP or Na is concerning, you can consider starting vasopressin right away while waiting for the other confirmatory labs.
- While waiting, it is advisable to increase IVF to 1.5 M to help combat losses, or allow the patient to drink freely.



 If DI criteria is met after those STAT labs are obtained, then you will proceed to DI management on page 2.

CLINICAL PATHWAY: Diabetes Insipidus (DI) Post-operative Neurosurgical Management PICU Post-operative Monitoring for DI

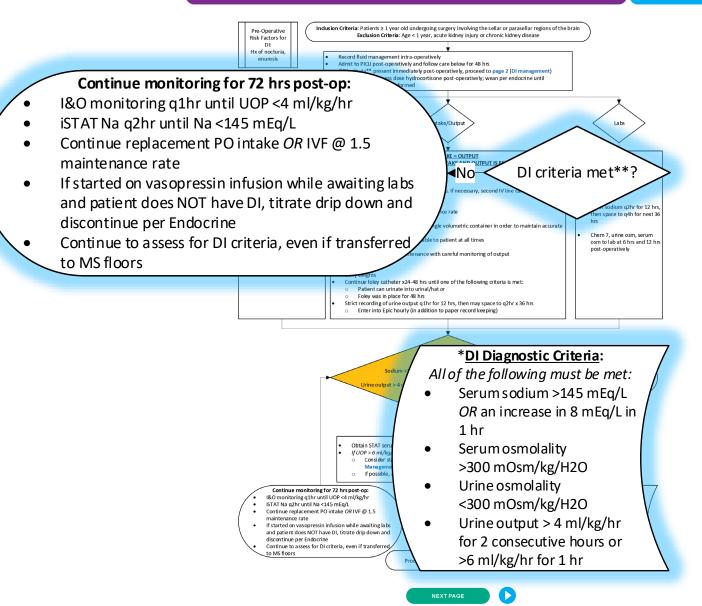
THIS PATHWAY
SERVES AS A GUIDE
AND DOES NOT
REPLACE CLINICAL
JUDGMENT.



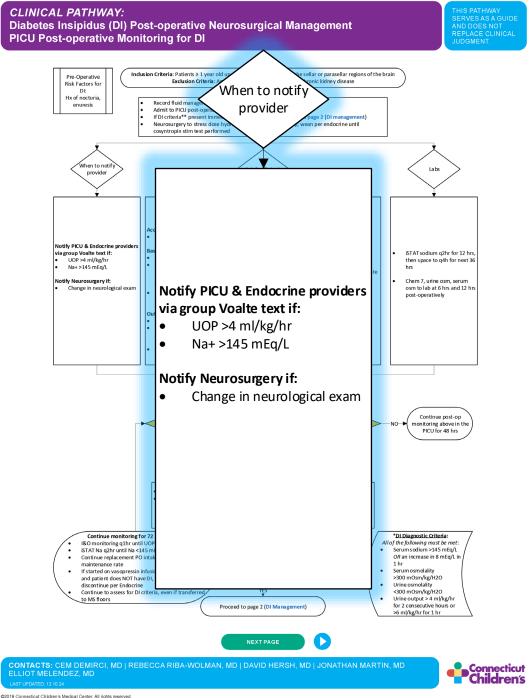


If DI criteria is not met:

- Continue closer monitoring until patient is more stable
- Increase input by increasing IVF to 1.5 maintenance or replace PO to meet UOP
- Titrate vasopressin if it was started
- Continue to closely monitor for DI as previously reviewed



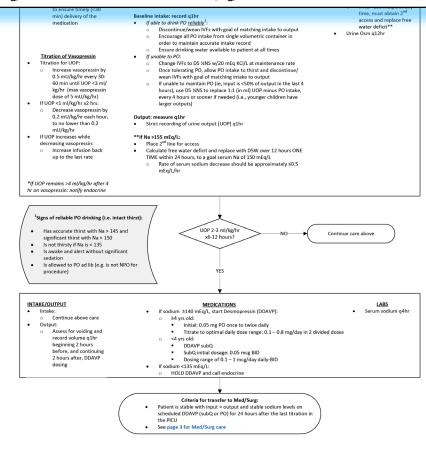
- Note that PICU/Endocrine providers should be notified immediately if UOP >4 ml/kg/hr or sodium is >145 mEq/L. This is to allow for immediate intervention and closer monitoring.
- Neurosurgery should be notified if there is any change in neurological examination.



- If DI criteria is met, follow the 2nd page of the algorithm for PICU management of DI
- Diagnostic criteria are listed here. Remember that all criteria must be met.

Diabetes Insipidus diagnosed if all of the following are met:

- Serum sodium >145 mEg/L OR an increase in 8 mEg/L in 1 hr
- Serum osmolality >300 mOsm/kg/H2O
- Urine osmolality <300 mOsm/kg/H2O
- Urine output > 4 ml/kg/hr for 2 consecutive hours or >6 ml/kg/hr for 1 hr











- Part of the initial management of DI is vasopressin.
- One role of vasopressin is to stimulate arginine vasopressin receptors (aka, antidiuretic hormone, or ADH).
- This results in decreased urine output and increased osmolality.
- Vasopressin is titrated based on UOP.
- If UOP stays >4 ml/kg/hr despite 4 hrs of vasopressin therapy, endocrine should be notified.

CLINICAL PATHWAY:
Diabetes Insipidus (DI)
PICU Management of DI

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

Serum sodium q2hr in the firs 24 hrs after diagnosis; then

> If Na >155 mEq/L at any time, must obtain 2nd

access and replace free

LABS

can space out to q4hr

water deficit**

Urine Osm a12hr

MEDICATION

- Order STAT Vasopressin IV infusion at 0.5 mU/kg/hr (max vasopressin dose of 5 mU/kg/ hr)
 - call pharmacy in order to ensure timely (<30 min) delivery of the medication

Titration of Vasopressin

- Titration for UOP:
 - Increase vasopressin by
 0.5 mU/kg/hr every 30 60 min until UOP <3 ml/kg/hr (max vasopressin dose of 5 mU/kg/hr)
- If UOP <1 ml/kg/hr x2 hrs:
 - Decrease vas opressin by 0.2 mU/kg/hr each hour, to no lower than 0.2 mU/kg/hr
- If UOP increases while decreasing vasopressin:
 - Increase infusion back up to the last rate

*If UOP remains >4 ml/kg/hr after 4 hr on vasopressin: notify endocrine

Diabetes Insipidus diagnosed if all of the following are met: Serum sodium >145 mEq/L OR an increase in 8 mEq/L in 1 hr Serum sondality >300 mOsm/kg/H20 Ufine osmolality <300 mOsm/kg/H20 Ufine osmolality <300 mOsm/kg/H20 Ufine output > 4 m/kg/hr for 2 consecutive hours or >6 m/kg/hr for 1 hr

INTAKE/OUTPUT GOAL: INTAKE = OUTPUT

METICULOUS TRACKING OF INTAKE AND OUTPUT IS ESSENTIAL

Maintain A-line and foley catheter as long as patient is on vasopres: Baseline intake: record altr

- If able to drink PO reliably²:
 Discontinue/wean IVFs with goal of matching intake to outpue
 Encourage all PO intake from single volumetric container in order to maintain accurate intake record
- Ensure drinking water available to patient at all times
- If unable to PO:

 Change IVFs to D5 ½NS w/20 mEq KCI/L at maintenance rate

 Once tolerating PO, allow PO intake to thirst and discontinue wean IVFs with goal of matching intake to output
- If unable to maintain PO (ie, input is <50% of output in the last 4 hours), use D5 ½MS to replace 1:1 (in ml) UOP minus PO intake, every 4 hours or sooner if needed (i.e., younger children have larger outputs)

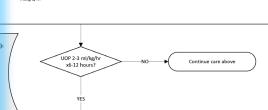
Output: measure q1hr

Strict recording of urine output (UOP) q1hr

**If Na >155 mEq/L:

Place 2nd line for access
Calculate free water deficit and replace with D5W over 12 hours ONE
TIME within 24 hours, to a goal serum Na of 150 mEg/L

Rate of serum sodium decrease should be approximately ≤0.5 mFq/I /hr



MEDICATIONS

If sodium ≥140 mEq/L, start Desmopressin (DDAVP):

≥4 yrs old:

Initial: 0.05 mg PO once to twice daily

Titrate to optimal daily dose range: 0.1 – 0.8 mg/day in 2 divided doses

<4 yrs old:</p>

DDAVP subQ

SubQ initial dosage: 0.05 mcg BID

Dosing range of 0.1 – 1 mcg/day daily-BID

If sodium <135 mEq/L:
O HOLD DDAVP and call endo

Criteria for transfer to Med/Surg:

- Patient is stable with input = output and stable sodium levels on scheduled DDAVP (subQ or PO) for 24 hours after the last titration i the PICU
- See page 3 for Med/Surg care









- Again, meticulous tracking of intake and output is essential.
- The goal would be to ensure that intake = output. Careful monitoring is necessary, especially while on vasopressin.
- While the patient is on vasopressin, an A line and Foley should be maintained.
- If the patient's sodium reaches over 155 mEq/L:
 - A 2nd line should be placed to allow additional fluids
 - Free water deficit should be replaced with D5W once to get to a goal of serum Na 150 mEq/L

MEDICATION

- Order STAT Vasopressin IV infusion at 0.5 mU/kg/hr (max vasopressin dose of 5 mU/kg/ hr)
 - Call pharmacy in order to ensure timely (<30 min) delivery of the medication

Titration of Vasopressin

- Titration for UOP:
 - Increase vasopress in by 0.5 mU/kg/hr every 30-60 min until UOP <3 ml/ kg/hr (max vasopressin dose of 5 mU/kg/hr)
- If UOP <1 ml/kg/hr x2 hrs:
 - Decrease vas opressin by 0.2 mU/kg/hr each hour, to no lower than 0.2 mU/kg/hr
- If UOP increases while decreasing vasopressin:
 - Increase infusion back up to the last rate

*If UOP remains >4 ml/kg/hr after 4 hr on vasopressin: notify endocrine

INTAKE/OUTPUT GOAL: INTAKE = OUTPUT METICULOUS TRACKING OF INTAKE AND OUTPUT IS ESSENTIAL

Access:

Maintain A-line and foley catheter as long as patient is on vasopressin

Baseline intake: record q1hr

- If able to drink PO <u>reliably</u>¹:
 - o Discontinue/wean IVFs with goal of matching intake to output
 - Encourage all PO intake from single volumetric container in order to maintain accurate intake record
 - o Ensure drinking water available to patient at all times
- If unable to PO:
 - Change IVFs to D5 ½NS w/20 mEq KCI/L at maintenance rate
 - Once tolerating PO, allow PO intake to thirst and discontinue/ wean IVFs with goal of matching intake to output
 - If unable to maintain PO (ie, input is <50% of output in the last 4 hours), use D5 ½NS to replace 1:1 (in ml) UOP minus PO intake, every 4 hours or sooner if needed (i.e., younger children have larger outputs)

Output: measure q1hr

• Strict recording of urine output (UOP) q1hr

** If Na >155 mEq/L:

- Place 2nd line for access
- Calculate free water deficit and replace with D5W over 12 hours ONE TIME within 24 hours, to a goal serum Na of 150 mEq/L
 - Rate of serum sodium decrease should be approximately ≤0.5 mEq/L/hr

<u>LABS</u>

- Serum sodium q2hr in the first
 24 hrs after diagnosis; then
 can space out to q4hr
 - If Na >155 mEq/L at any time, must obtain 2nd access and replace free water deficit**
- Urine Osm q12hr

Criteria for transfer to Med/Surg:

- Patient is stable with input = output and stable sodium levels on scheduled DDAVP (subQ or PO) for 24 hours after the last titration in the PICU
- See page 3 for Med/Surg care

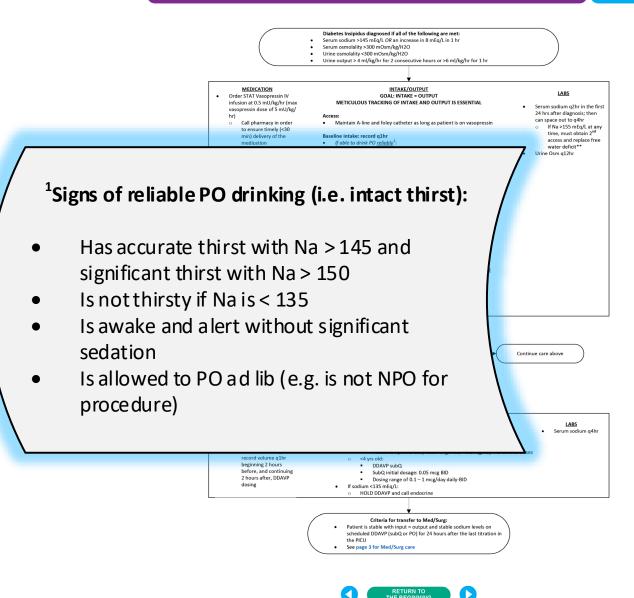








 It is vitally important to understand how to evaluate for intact thirst. Signs of reliable PO drinking are now listed on the pathway.



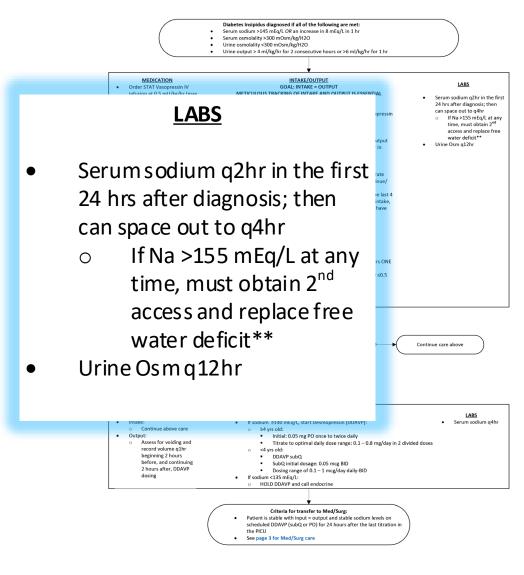
CONTACTS: CEM DEMIRCI, MD | REBECCA RIBA-WOLMAN, MD | DAVID HERSH, MD | JONATHAN MARTIN, MD

ELLIOT MELENDEZ, MD

- Labs are directed at closely monitoring serum sodium and urine osmolality
- Remember, if sodium becomes >155 mEq/L at any time, obtain a 2nd line to replace the free water deficit



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









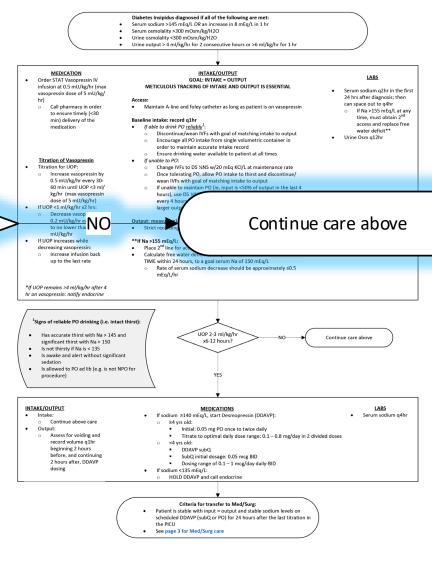




 If UOP has not stabilized to 2-3 ml/kg/hr for 6-12 hours, then the care outlined in previous slides should continue. UOP 2-3 ml/kg/hr x6-12 hours?

CLINICAL PATHWAY:
Diabetes Insipidus (DI)
PICU Management of DI

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









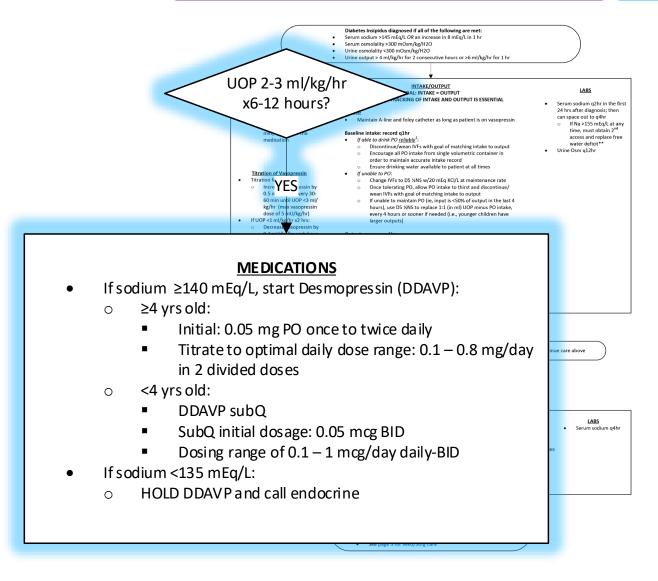
CONTACTS: CEM DEMIRCI, MD | REBECCA RIBA-WOLMAN, MD | DAVID HERSH, MD | JONATHAN MARTIN, MD ELLIOT MELENDEZ, MD



If UOP has reached 2-3 ml/kg/hr for 6-12 hours:

 DDAVP may be started depending on sodium levels. CLINICAL PATHWAY:
Diabetes Insipidus (DI)
PICU Management of DI

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







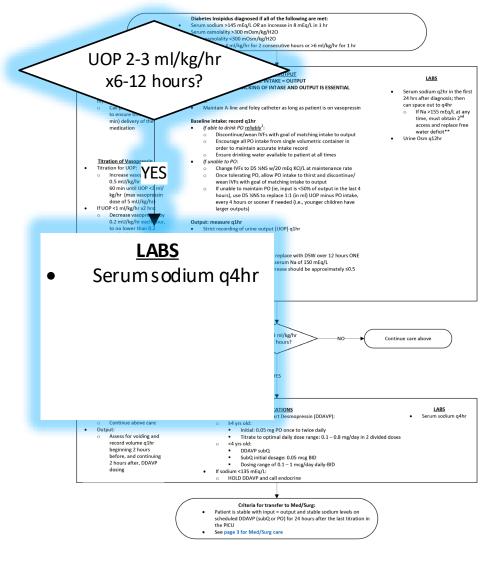




If UOP has reached 2-3 ml/kg/hr for 6-12 hours:

 Serum sodium monitoring can be spaced to every 4 hours if not already done CLINICAL PATHWAY:
Diabetes Insipidus (DI)
PICU Management of DI

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









CONTACTS: CEM DEMIRCI, MD | REBECCA RIBA-WOLMAN, MD | DAVID HERSH, MD | JONATHAN MARTIN, MD ELLIOT MELENDEZ, MD

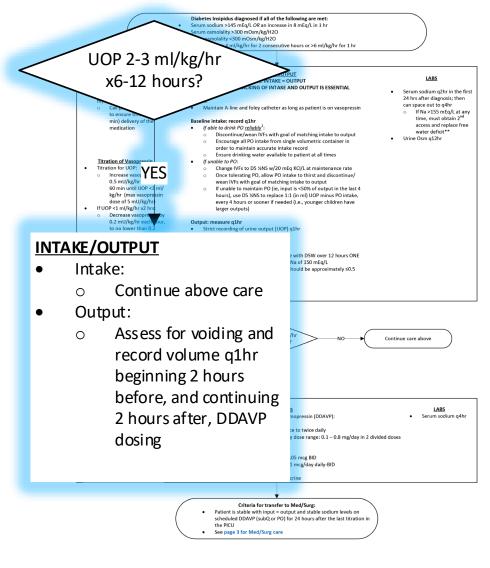


If UOP has reached 2-3 ml/kg/hr for 6-12 hours:

- Close monitoring of Intake and Output should continue
- Output assessment is important around DDAVP dosing

CLINICAL PATHWAY:
Diabetes Insipidus (DI)
PICU Management of DI

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









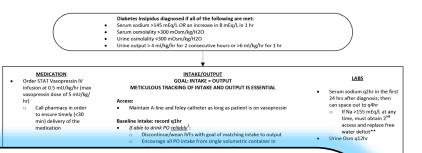
 $\textbf{CONTACTS}: \texttt{CEM DEMIRCI, MD} \mid \texttt{REBECCA RIBA-WOLMAN, MD} \mid \texttt{DAVID HERSH, MD} \mid \texttt{JONATHAN MARTIN, MD} \mid \texttt{ELLIOT MELENDEZ, MD} \mid \texttt{MD} \mid \texttt{MD}$



CLINICAL PATHWAY:
Diabetes Insipidus (DI)
PICU Management of DI

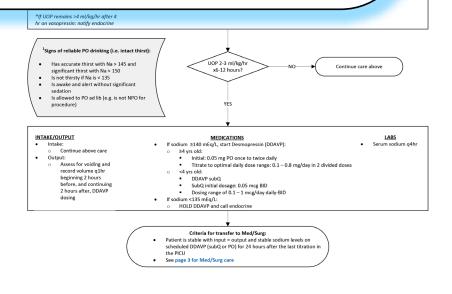
THIS PATHWAY
SERVES AS A GUIDE
AND DOES NOT
REPLACE CLINICAL
JUDGMENT.

 After DDAVP is titrated and the patient remains on scheduled doses for 24 hours, transfer to med/surg can be considered if intake = output and sodium levels are stable.



Criteria for transfer to Med/Surg:

- Patient is stable with input = output and stable sodium levels on scheduled DDAVP (subQ or PO) for 24 hours after the last titration in the PICU
- See page 3 for Med/Surg care











- Once criteria to transfer out of the PICU to the med/surg floors is met, follow page 3 of the pathway: Med/Surg Management of DI
- Endocrinology will direct the care for DI management.

CLINICAL PATHWAY:
Diabetes Insipidus (DI)
Med/Surg Management of DI

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

Med/Surg Management for Diabetes Insipidus:

Patient is transferred from PICU when stable
(input = output and stable sodium levels on scheduled SubQ or PO DAVP for 24 hours after the last titration)

Endocrine to direct care for DI management below

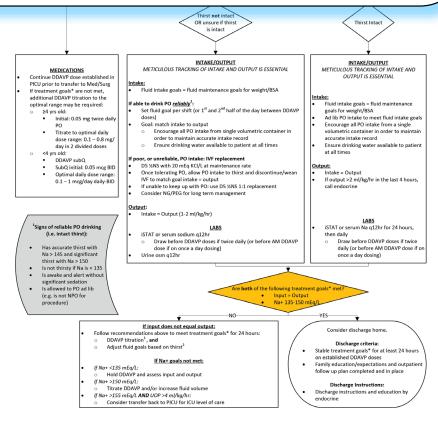
Treatment goals:
 Input = output
 Na+ 135-150 mEq/L

Med/Surg Management for Diabetes Insipidus:

Patient is transferred from PICU when stable

(input = output and stable sodium levels on scheduled SubQ or PO DDAVP for 24 hours after the last titration)

Endocrine to direct care for DI management below





RETURN TO THE BEGINNING



Med/Surg Management for Diabetes Insipidus:

Patient is transferred from PICU when stable

(input = output and stable sodium levels on scheduled Subo or PO DIADV for 24 hours after the last titration

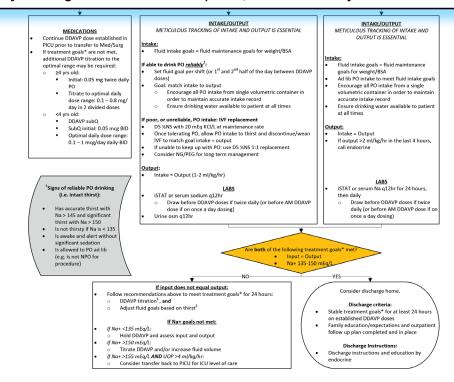
Endocrine to direct care for DI management below

*Treatment goals:
Input = output
Na+ 135-150 mEq/L

Monitoring:

- Meticulous tracking of intake and output q4hr is essential
- Vitals q4hr
- Labs q12hr and are dependent upon thirst² (see below)

If more frequent monitoring of vital signs and I&Os are required, consider transfer back to PICU.





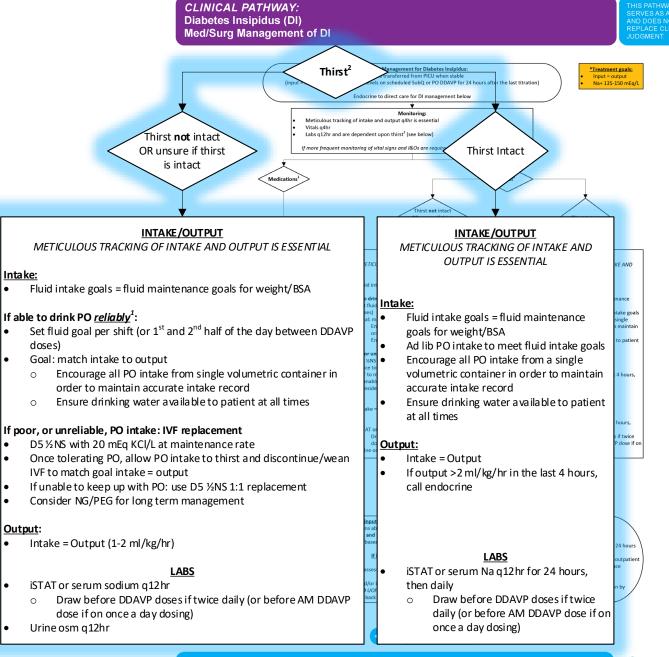
RETURN TO THE BEGINNING

Meticulous I&O tracking is essential.
On the med/surg floors, this is done
every 4 hours in conjunction with
vitals. If more frequent monitoring is
required, then consider transferring
back to the PICU for closer
monitoring.

Labs are dependent upon thirst mechanism.



- Recommendations for I&Os and labs depend on if thirst mechanisms are intact.
- Both require:
 - Meticulous tracking of I&Os
 - · Fluid intake goals to equal fluid maintenance goals for weight/BSA
 - Intake = output



CONTACTS: CEM DEMIRCI, MD I REBECCA RIBA-WOLMAN, MD I DAVID HERSH, MD I JONATHAN MARTIN, MD



Intake:

Output:

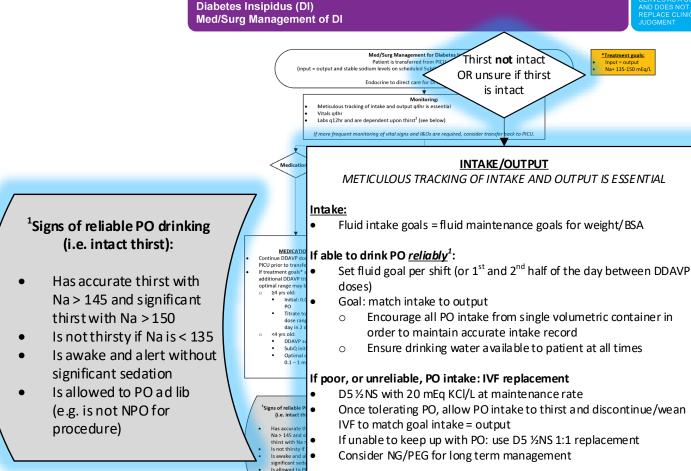
doses)



If thirst is not intact (or if there is uncertainty):

Intake:

- Should have fluid goals set and encourage all intake from one container for accurate measurements if patient is able to PO
- If PO intake is poor:
 - Start maintenance IVF
 - May need to replace 1:1
 - Consider using a NG/PEG for long term management



CLINICAL PATHWAY:

Output:

(e.g. is not Ni procedure)

Intake = Output (1-2 ml/kg/hr)

LABS

- iSTAT or serum sodium q12hr
 - Draw before DDAVP doses if twice daily (or before AM DDAVP dose if on once a day dosing)
- Urine osm q12hr







If thirst is not intact (or if there is uncertainty):

Labs:

- Sodium should be measured every 12 hours and drawn before DDAVP doses.
- Urine osm should also be monitored

CLINICAL PATHWAY: Diabetes Insipidus (DI) Med/Surg Management of DI

¹Signs of reliable PO drinking

(i.e. intact thirst):

Has accurate thirst with

Na > 145 and significant

Is not thirsty if Na is < 135

Is awake and a lert without

thirst with Na > 150

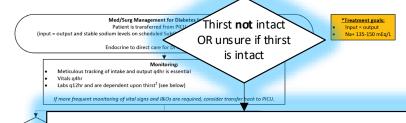
significant sedation

(e.g. is not NPO for

procedure)

Is allowed to PO ad lib

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



INTAKE/OUTPUT

METICULOUS TRACKING OF INTAKE AND OUTPUT IS ESSENTIAL

<u>Intake:</u>

Continue DDAVP de

PICU prior to trans

If treatment goals*

optimal range ma

≥4 yrs old:

Initial: 0

Titrate

DDAVP

SubQ in

Signs of reliable (i.e. intact t

Has accurate Na > 145 and

Is not thirsty

Is awake and significant se Is allowed to (e.g. is not Ni procedure) Fluid intake goals = fluid maintenance goals for weight/BSA

If able to drink PO <u>reliably</u>1:

- Set fluid goal per shift (or 1st and 2nd half of the day between DDAVP doses)
- Goal: match intake to output
 - Encourage all PO intake from single volumetric container in order to maintain accurate intake record
 - o Ensure drinking water available to patient at all times

If poor, or unreliable, PO intake: IVF replacement

- D5½NS with 20 mEq KCl/L at maintenance rate
- Once tolerating PO, allow PO intake to thirst and discontinue/wean
 IVF to match goal intake = output
- If unable to keep up with PO: use D5 ½NS 1:1 replacement
- Consider NG/PEG for long term management

Output:

Intake = Output (1-2 ml/kg/hr)

LABS

- iSTAT or serum sodium q12hr
 - Draw before DDAVP doses if twice daily (or before AM DDAVP dose if on once a day dosing)
- Urine osm q12hr



RETURN TO THE BEGINNING

CONTACTS: CEM DEMIRCI, MD | REBECCA RIBA-WOLMAN, MD | DAVID HERSH, MD | JONATHAN MARTIN, MD ELLIOT MELENDEZ, MD



©2019 Connecticut Children's Medical Center. All rights reserved.

If thirst is intact

Intake:

 PO should be ad lib and monitored from one container

Output:

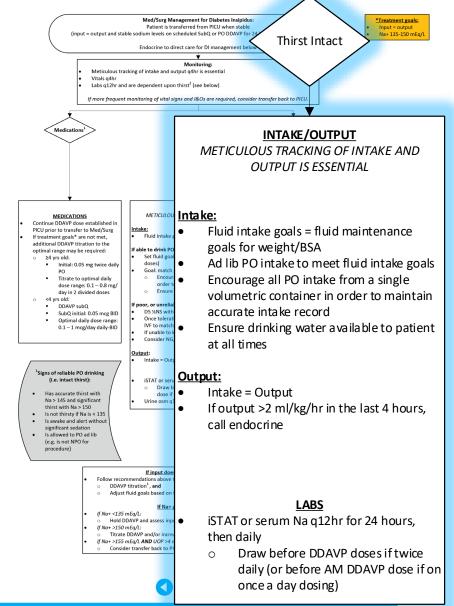
 Closely monitor output and call endocrine if output exceeds 2 ml/kg/hr in the last 4 hours

Labs:

Only sodium will be monitored

CLINICAL PATHWAY:
Diabetes Insipidus (DI)
Med/Surg Management of DI

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



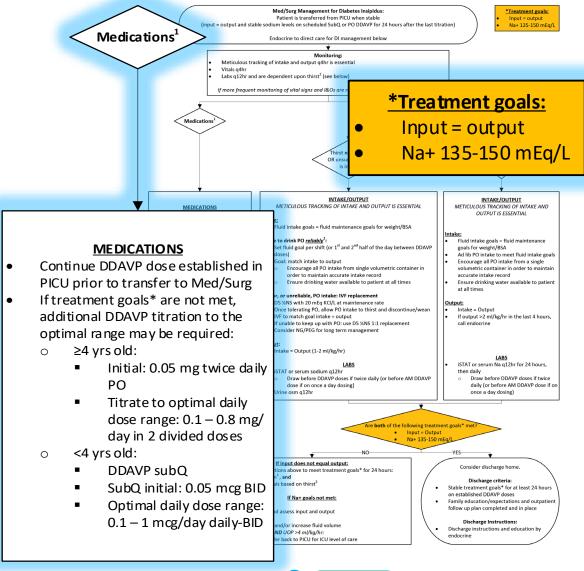
CONTACTS: CEM DEMIRCI, MD | REBECCA RIBA-WOLMAN, MD | DAVID HERSH, MD | JONATHAN MARTIN, MD ELLIOT MELENDEZ, MD



- All patients will continue their scheduled DDAVP dose that was established in the PICU.
- Treatment goals are outlined in the yellow box. DDAVP may need to be further titrated to reach these optimal ranges.

CLINICAL PATHWAY:
Diabetes Insipidus (DI)
Med/Surg Management of DI

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

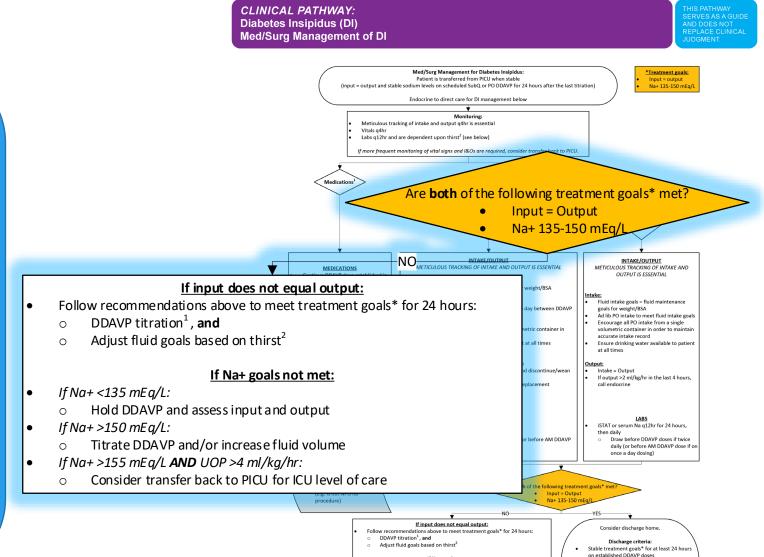




RETURN TO THE BEGINNING



- Remember that treatment goals are input=output and sodium levels within 135-150 mEq/L
- If input does not equal output, consider DDAVP titration and adjusting fluid goals.
- If sodium goals are not met, guidelines for DDAVP and/or input adjustments are listed.
- If there is concerning Na of >155
 mEq/L and UOP >4 ml/kg/hr,
 consider transfer back to the PICU.





Hold DDAVP and assess input and output

If Na+ >155 mEq/L AND UOP >4 ml/kg/hr:

Consider transfer back to PICU for ICU level of car.

Titrate DDAVP and/or increase fluid volume

If Na+ <135 mFa/I

If Na+ >150 mEq/L:





follow up plan completed and in plac

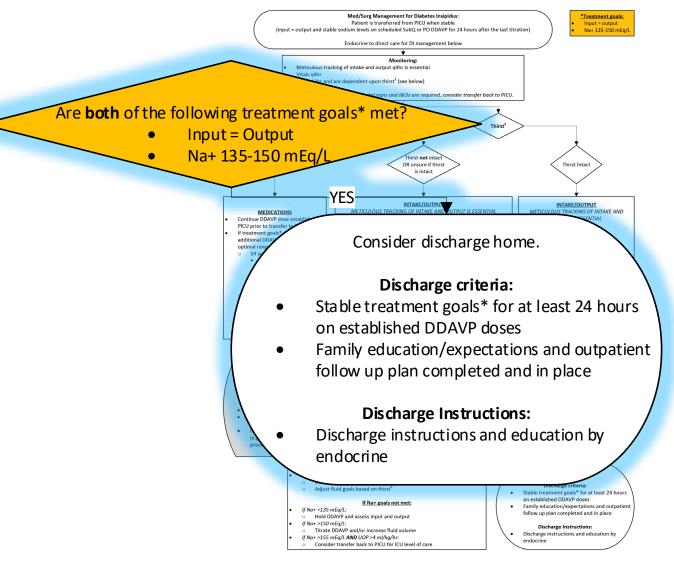
Discharge instructions and education b

Discharge Instructions

- If treatment goals are met for both input=output and Na 135-150 mEq/L while on established DDAVP for at least 24 hours, then patient can be considered for discharge.
- Discharge education and instructions will be provided by endocrinology.



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





CONTACTS: CEM DEMIRCI, MD | REBECCA RIBA-WOLMAN, MD | DAVID HERSH, MD | JONATHAN MARTIN, MD





ELLIOT MELENDEZ. MD

Review of Key Points



- Central DI can develop post-operatively after a neurosurgical procedure following sellar and parasellar regions
- DI diagnosis depends on serum sodium, serum osmolality, urine osmolality and urine output measures
- The main goal is for the patient to maintain intake = output, thus careful monitoring of I&Os are essential
- A second goal is to achieve desirable sodium levels. Frequency of monitoring depends on the clinical situation.

Use of Order Set



 An associated order set in Care Navigator is undergoing completion. We will make an announcement when it is available.

Quality Metrics



- Percentage of eligible patients with pathway order set usage (PICU, Med-Surg Unit)
- Average time (minutes) from arrival to PICU to administration of Vasopressin if DI suspected/confirmed
- Number of transfers from Med-Surg unit back to PICU (all cause)
- Number of patients with DI post-operatively
- Average length of stay (PICU, days)
- Average length of stay (hospital, days)

Pathway Contacts



- Cem Demirci, MD
 - Division of Endocrinology
- Rebecca Riba-Wolman, MD
 - Division of Endocrinology
- David Hersh, MD
 - Division of Neurosurgery
- Jonathan Martin, MD
 - Division of Neurosurgery
- Elliot Melendez, MD
 - Division of Critical Care

References



- Christ-Crain M, Biche DG, Fenske WK, et al. Diabetes insipidus. *Nat Rev Dis Primers*. 2019 Aug;5(1): 54.
- Di lorgi N, Napoli F, Allegri AE, et al. Diabetes Insipidus Diagnosis and Management. Horm Res Paediatr. 2012;77(2):69-84.
- Melmed S, Koenig R, Rosen C, Auchus R, Goldfine A. Williams Textbook of Endocrinology. 14th ed. Elsevier; c2019. 1792 p.
- Pratheesh R, Swallow DM, Rajaratnam S, et al. Incidence, predictors and early post-operative course of diabetes insipidus in paediatric craniopharygioma: a comparison with adults. *Childs Nerv Syst*. 2013 Jun;29(6):941–9.
- Sperling M, et al. Sperling Pediatric Endocrinology. 5th ed. Elsevier; c2020.
 1072 p.

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.