PHASE I (ED/Urgent Care): ORT

Inclusion Criteria
- Vomiting and/or diarrhea of recent onset not due to chronic disease, with or without fever, nausea, or abdominal pain

Exclusion Criteria
- Patient < 3 months of age
- Toxic appearance (consider sepsis)
- Diarrhea > 7 days (consider chronic disease, bacterial enteritis)
- Bloody diarrhea (consider HUS)
- Comorbid conditions (Medically Complex Children (MCC), renal failure, cardiac disease)
- Bilious emesis (consider bowel obstruction)
- On diuretic therapy
- Hyponatremia (<130 mEq/L) or Hypernatremia (>155 mEq/L)
- Acute surgical abdomen

Give Ondansetron if Moderately Dehydrated
- Consider holding if diarrhea is chief complaint

Initial ORT Challenge
- 5 mL q 5 mins if <10 kg
- 10 mL q 5 mins if ≥10 kg

Emesis after initial ORT?
- No

Increase ORT
- 10 mL q 5 mins if <10 kg
- 20 mL q 5 mins if ≥10 kg
- Assess in 30-60 mins

Overt Shock (Dehydration >10%)

Moderate Dehydration (5-10%)
- Prolonged capillary refill (>2 seconds)
- Abnormal skin turgor ('tenting' or inelastic skin)
- Tachypnea

Hold ORT for 20 minutes
- Then restart ORT at initial rate

No

Yes

Antimicrobials not recommended

Routine testing for stool pathogens not recommended

Consider oral challenge (if attempted)

Passed oral challenge (if attempted); Educate and prepare for discharge

Discharge Criteria
- Clinical status improved
- Tolerating ORT or regular diet
- Adequate family teaching
- Follow-up established

Discharge Instructions
- Continue ORT at home for 4-6 additional hours; then resume regular diet, or lactobacillus if predominantly formula fed
- ED Acute Gastroenteritis discharge instructions and ORT worksheet
- Lactobacillus for 5 days

Emesis after initial ORT?

Yes

Continue to Phase II

No

Discharge Criteria
- Clinical status improved
- Tolerating ORT or regular diet
- Adequate family teaching
- Follow-up established

Discharge Instructions
- Continue ORT at home for 4-6 additional hours; then resume regular diet, or lactobacillus if predominantly formula fed
- ED Acute Gastroenteritis discharge instructions and ORT worksheet
- Lactobacillus for 5 days

No

Increased ORT

Failed oral challenge (if attempted)

Passed oral challenge (if attempted); Educate and prepare for discharge
PHASE II (ED/Urgent Care): IV or NG Rehydration

Check vital signs
- Check BP, HR, RR
- Evaluate heart and lung sounds

Involve caregiver and patient in decision to hydrate with IV or NG

1st IV Fluid Bolus
- NS 20 mL/kg over 30-60 mins (Max dose: 1,000 mL)
- Upon initiation of IV, check electrolytes
- Consider ondansetron if not already given or tolerated

Recheck Vital Signs; Re-examine
- If improved, consider returning to ORT

2nd IV Fluid Bolus
- NS 20 mL/kg over 30-60 mins

1st NG Fluid Bolus
- Pedialyte 20 mL/kg over 60 mins (Max dose: 600 mL)
- Consider ondansetron if not already given or tolerated

Recheck Vital Signs; Re-examine
- If improved, consider returning to ORT

2nd NG Fluid Bolus
- Pedialyte 20 mL/kg over 60 mins

Evaluate for Discharge
- Clinical status improved
- IV or NG fluids not required
- Tolerating ORT or regular diet
- Adequate family teaching
- Follow-up established

Re-examine Exclusion Criteria
- Patient < 3 months age
- Toxic appearance (consider sepsis)
- Diarrhea > 7 days (consider chronic disease, bacterial enteritis)
- Bloody diarrhea (consider HUS)
- Comorbid conditions (MCC, renal failure, cardiac disease)
- Bilious emesis (consider bowel obstruction)
- On diuretic therapy
- Hyponatremia (<130 mEq/L) or Hypernatremia (>155 mEq/L) (if electrolytes checked)
- Acute surgical abdomen

Yes, meets discharge criteria
- Continue to Phase III
- Admit on AGE pathway

No, does not meet discharge criteria
- Urgent Care Transfer Criteria:
  - Not tolerating ORT following 2nd IV/NG bolus (send by BLS)
  - Need for a 3rd IV/NG bolus (send by BLS)
  - Overt shock (IV access, bolus started, ALS transport)
  - Worsening clinical status
  - Continue treatment as outlined by pathway while awaiting transport

Discharge Instructions
- Continue ORT at home for 4-6 additional hours; then resume regular diet, or lactose free formula if predominantly formula fed
- ED Acute Gastroenteritis discharge instructions and ORT worksheet
- Lactobacillus for 5 days

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Last Updated: June 2015
Next Expected Review: June 2020
**Acute Gastroenteritis v. 2.0: Inpatient Management**

**PHASE III: INPATIENT MANAGEMENT**

**Admit on AGE powerplan**
- Start IV D5 NS + 20 mEq KCl/liter OR NG Pedialyte at maintenance rate
- Assess need for additional NS or Pedialyte bolus

**Assess at 0800 (or after 4 hours of IV/NG fluids if admitted between 0800-1600)**
- Vomiting: ≤1 episodes in the past 4 hours?
- Intake > Output?

**ORT or Regular Diet Challenge**
- Discontinue IV or NG fluids
- ORT: 5 mL q 5 mins if <10kg; 10 mL q 5 mins if ≥10kg, or resume regular diet (lactose-free formula if predominantly formula fed)
- Reassess after 20 minutes
- Send prescription for Lactobacillus GG

**Discharge Instructions**
- Continue ORT, or
- Resume regular diet, or lactose-free formula if predominantly formula fed
- Acute Gastroenteritis And Oral Rehydration Therapy handout (PE636)
- Lactobacillus for 5 days

**Discharge Criteria**
- Sufficient rehydration as indicated by weight gain OR normal respiratory rate, capillary refill, and skin turgor
- IV or NG fluids not required
- Tolerating ORT or regular diet
- Adequate family teaching
- Follow-up established

**Consider secretory diarrhea if there is copious output in the absence of oral intake**

**Anti-emetics are not recommended outside of the ED / Urgent Care**

**Executive Summary**

**Explanation of Evidence Ratings**

**Test your knowledge**

**Summary of Version Changes**

**Citation Information**

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Last Updated: June 2015
Next Expected Review: June 2020
Phase II (ED/UC): IV or NG Rehydration (Cont'd)

- Do not routinely obtain stool cultures in cases of uncomplicated AGE. Consider performing stool studies in specific cases, such as: infants <3 months of age, patients with bloody or mucoid stools, diarrhea for >7 days, immunocompromised patients, if there is a known outbreak of a bacterial pathogen, if the child has recently been on antibiotics, if the diagnosis of simple AGE is in doubt, if there is concern for sepsis, and in cases of diarrhea after foreign travel.1, 5, 8 [Guideline]

- If the patient meets any of the above criteria, they would be off pathway.
Phase I (ED/UC): ORT

- It is recommended that the history and physical examination be the primary basis for the diagnosis of AGE [Expert Opinion].
- **Acute body weight change is considered the gold standard measure of dehydration in a child but is often impractical for the initial assessment due to lack of an accurate pre-illness weight measurement. When a reliable pre-illness weight is available, it is recommended to compare the pre-illness to the current weight as the best assessment for degree of dehydration.**\(^5\), \(^8\) [Guideline]
- **Prolonged capillary refill time, abnormal skin turgor, and respiratory rate are the best individual examination measures.**\(^1\), \(^5\), \(^6\) [Guideline]

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Phase I (ED/UC): ORT (Cont'd)

- After using the Inclusion and Exclusion criteria to determine if a patient is appropriate for the Acute Gastroenteritis: Clinical Standard Pathway, the next step is to determine the level of dehydration of the patient
- Determining the level of dehydration of the patient will guide the amount of intervention required of the patient
- The level of dehydration can broadly be categorized into 3 levels:
  1. minimal to no dehydration (loss of < 5% body weight)
  2. mild to moderate dehydration (loss of 5 – 10% body weight)
  3. overt shock (loss >10% body weight)
Phase I (ED/UC): ORT (Cont'd)

- If a patient is in overt shock (>10% dehydration), the patient should be off pathway and treated for shock.
- In a patient with moderate dehydration (5-10% dehydration), you may see corresponding clinical symptoms, and these patients should receive oral ondansetron.

Phase I (ED/UC): Oral Rehydration Therapy (ORT)

- If a patient has minimal to no dehydration (<5% dehydration), there is no evidence for routine use of oral ondansetron in this population, and the caregiver and child should be instructed on proper use of ORT.
- Some providers may consider giving a child with minimal to no dehydration an oral challenge in the ED/UC, and if the child fails this challenge if given, the child may receive oral ondansetron at this point.
Discharge Instructions: Lactobacillus

- Use probiotics in the form of Lactobacillus GG, Saccharomyces boulardii, and L acidophilus LB as an adjunctive treatment in the management of children with diarrhea from acute gastroenteritis. Start immediately and treat for 5 days.4, 5, 7, 8, 11-13 [Guideline/_ISO_O]

- A combination of systematic reviews and guidelines were reviewed to address the question of the efficacy of probiotics in the adjunctive treatment of diarrhea in children with AGE.

- Three strains in particular seemed to be useful in reducing the duration of diarrhea, reducing the risk of protracted diarrhea (>7 days), and reducing the risk of hospitalization

- Lactobacillus GG, in particular, was reviewed most extensively, and has proven efficacy against rotavirus, one of the main causes of protracted diarrhea and the associated morbidities.
Medication Use: Anti-diarrheal Agents

- Anti-diarrheal agents are NOT recommended in the routine management of children with AGE and use of them may be associated with increased serious adverse effects.¹ ⁸ [Guideline]
Use of Antimicrobials for Acute Gastroenteritis

• The etiology of acute gastroenteritis is typically viral, and therefore the routine use of antimicrobial therapies is not indicated.

• It is recommended that antimicrobial therapies be used ONLY for selected children with acute gastroenteritis [Expert Opinion]:
  o Children with evidence of a serious bacterial infection (SBI)
  o Children with persistent diarrhea and evidence of infection with Giardia lamblia or Cryptosporidium; if found, treatment is available with metronidazole or nitazoxanide
Medication Use: Ondansetron

- Oral or IV ondansetron in moderately dehydrated children with AGE may decrease the number of episodes of vomiting, decrease the time to cessation of vomiting, facilitate earlier oral rehydration, decrease the need for IV rehydration, decrease hospital admission rates, decrease the risk of receiving IV fluids up to 72 hours post discharge from the ED, and improve parental satisfaction as compared to placebo.\(^2\)\(^5\)\(^8\) [Guideline]

- The efficacy of ondansetron for hospitalized children has not adequately been studied.

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Antiemetic Use Outside of the ED / Urgent Care

- Several high quality guidelines and systemic reviews demonstrate some benefit with the single use of ondansetron in the ED or Urgent Care settings. However, no such evidence exists for routine use in the inpatient setting.

- *Routine, ongoing use of ondansetron in the inpatient setting is not supported by current literature and therefore is not currently recommended.*\(^5\)\(^6\) [Guideline]
**Phase I (ED/UC): ORT**

- **Give Ondansetron if Moderately Dehydrated**
  - Consider holding if diarrhea is chief complaint

**Begin ORT**

**Initial ORT Challenge**
- 5mL q 5 mins if <10 kg
- 10mL q 5 mins if ≥10 kg

- The amount of ORT to be given during the challenge is based on the patient’s admitted weight:
  - 5 mL every 5 minutes if < 10 kg
  - 10 mL every 5 minutes if ≥ 10 kg

- The amount of ORT started is limited at first, and gradually increased as tolerated.\(^8\) [Guideline]

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**Oral Rehydration Therapy**

- It is recommended that dehydration be treated with oral rehydration solutions (ORS), if tolerated and if intake exceeds losses, for a period of 4 to 6 hours or until an adequate degree of rehydration is achieved.\(^1, 5, 8\) \(^9\) [Guideline]
Phase I (ED/UC): ORT Moderate Dehydration

• If, after 20 minutes, the patient tolerates the initial ORT challenge (there is no emesis), then the amount of ORT is increased:
  o 10 mL every 5 minutes if < 10 kg
  o 20 mL every 5 minutes if ≥ 10 kg

• If, during the initial 20 minute challenge, the patient does have emesis, then:
  o the ORT challenge is held for 20 minutes
  o the ORT challenge is restarted after 20 minutes at the initial, lower rate
    ▪ 5 mL every 5 minutes if < 10 kg
    ▪ 10 mL every 5 minutes if ≥ 10 kg

Phase I (ED/UC): ORT Moderate Dehydration (Cont'd)

• If the patient tolerates the increase in ORT without emesis, then the patient can be prepared for discharge

• Patients should be encouraged to continue on their usual or regular diet and should be educated on the prevention of dehydration

• If the patient does not tolerate the increase in ORT, and continues to have emesis even after holding ORT for 20 minutes and restarting at the lower rate, then the patient will change phases on the Acute Gastroenteritis: Clinical Standard Pathway and move to:
  o Phase II (ED/UC): IV or NG Rehydration
Obtaining Lab or Stool Studies

- Supplementary laboratory studies, including serum electrolytes, are usually unnecessary in children with mild to moderate dehydration not requiring IV or NG fluids. They have not been shown to be a reliable surrogate for determining the degree of dehydration or rehydration. [Guideline]

- Perform electrolytes in moderately dehydrated children who are requiring the initiation of IV fluids. [Guideline]

- A normal bicarbonate concentration may be useful in ruling out dehydration. [Guideline]

- An abnormal bicarbonate concentration has not been studied as a predictor for hospitalization after correction of dehydration [Expert Opinion].

- If a patient is found to be hypoglycemic, hyponatremic, or hypernatremic, correct with appropriate fluids accordingly.
Phase II (ED/UC): IV or NG Rehydration (Cont'd)

1st IV Fluid Bolus
- NS 20 mL/kg over 30-60 mins (Max dose: 1,000 mL)
- Upon initiation of IV, check electrolytes
- Consider ondansetron if not already given or tolerated

Recheck Vital Signs; Re-examine
- If improved, consider returning to ORT

2nd IV Fluid Bolus
- NS 20 mL/kg over 30-60 mins

1st NG Fluid Bolus
- Pedalyte 20 mL/kg over 60 mins (Max dose: 600 mL)
- Consider ondansetron if not already given or tolerated

Recheck Vital Signs; Re-examine
- If improved, consider returning to ORT

2nd NG Fluid Bolus
- Pedalyte 20 mL/kg over 60 mins

- Patients who advance to Phase II (ED/UC): IV or NG Rehydration on the AGE Clinical Standard Pathway will, after shared decision-making between the caregivers and providers, be given either a first bolus of:
  - IV normal saline (NS) at a dose of 20 mL/kg body weight over 30-60 minutes (max dose 1000mL) [Expert Opinion]
  - or, Pedalyte through an NG over 60 minutes (max dose 600mL) [Expert Opinion].

Phase II (ED/UC): IV or NG Rehydration (Cont'd)

- Consider giving a dose of oral, IV (if initiating IV fluids), or NG (if initiating NG fluids) ondansetron at this time if not previously given or if the patient vomited within 15 minutes of the initial dose.
- Providers may consider a bedside glucose check at this time, usually during the placement of the peripheral IV [Expert Opinion].
- Perform a full assessment before and after the fluid bolus, that includes vital signs (HR, RR, BP) and evaluation of heart tones and lung sounds, for consideration of the potential for heart failure and/or myocarditis.
Phase II (ED/UC): IV or NG Rehydration (Cont'd)

1st IV Fluid Bolus
- NS 20 mL/kg over 30-60 mins (Max dose: 1,000 mL)
- Upon initiation of IV, check electrolytes
- Consider ondansetron if not already given or tolerated

Recheck Vital Signs; Re-examine
- If improved, consider returning to ORT

2nd IV Fluid Bolus
- NS 20 mL/kg over 30-60 mins

1st NG Fluid Bolus
- Pedialyte 20 mL/kg over 60 mins (Max dose: 600 mL)
- Consider ondansetron if not already given or tolerated

Recheck Vital Signs; Re-examine
- If improved, consider returning to ORT

2nd NG Fluid Bolus
- Pedialyte 20 mL/kg over 60 mins

- Recheck vital signs and re-examine the patient after first IV or NG fluid bolus, and, if improved, consider returning to an ORT trial at this time.
  - 5 mL every 5 minutes if < 10 kg
  - 10 mL every 5 minutes if ≥ 10 kg

- Advance ORT after 20 minutes if tolerated.

- If the patient has not improved, or if emesis continues, give a second bolus of fluid, consisting of, either:
  - IV normal saline (NS) at a dose of 20 mL/kg body weight over 30-60 minutes (max dose 1000mL) [Expert Opinion]
  - or, Pedialyte through an NG over 60 minutes (max dose 600mL) [Expert Opinion].
Phase II (ED/UC): IV or NG Rehydration

- A comparison between NG versus IV fluids for rehydration found that NG rehydration was as efficacious, no more labor intensive, and was associated with less complications than the use of IV fluids. Because of these reasons, it is advisable to involve the parents early on in the decision of how to appropriately rehydrate their child. [Guideline]
Myocarditis

Patients with AGE who deteriorate after an IV fluid bolus should be assessed for evidence of myocarditis or heart failure [Expert Opinion]. Signs and symptoms might include:

- Worsening tachypnea or increase in heart rate over baseline
- Hypotension or a decrease in blood pressure over baseline
- Muffled or hard-to-hear heart sounds
- Coarser (wetter) lung sounds
- An enlargement of liver span

Patients with signs and/or symptoms of myocarditis or heart failure should be removed from the AGE pathway.

Return to Phase II
Phase II (ED/UC): Urgent Care Transfer Criteria

Urgent Care Transfer Criteria:

- Not tolerating ORT following 2nd IV/NG bolus (send by BLS)
- Need for a 3rd IV/NG bolus (send by BLS)
- Overt shock (IV access, bolus started, ALS transport)
- Worsening clinical status

**Continue treatment as outlined by pathway while awaiting transport**

- Additionally, a patient who is being treated for AGE according to this pathway may be transferred from the Urgent Care to the ED at any time if:
  - A decision is made with the caregivers to initiate NG rehydration and the ability to place an NG tube is not possible at the UC.
Phase II (ED/UC): IV or NG Rehydration (Cont'd)

- If the patient has tolerated ORT or their regular diet for 20 minutes, is no longer requiring IV or NG fluids, and appears clinically improved, the provider may instruct the family on continued ORT use at home, and discuss establishing follow up care with their primary care provider.

Recheck Vital Signs; Re-examine

Evaluate for Discharge
- Clinical status improved
- IV or NG fluids not required
- Tolerating ORT or regular diet
- Adequate family teaching
- Follow-up established
Phase II (ED/UC): IV or NG Rehydration (Cont'd)

- If a patient does not meet discharge criteria from the ED or Urgent Care, re-examine if the patient meets any of the exclusion criteria and does not belong on this pathway.
- If no exclusion criteria are present, admit to Phase III: Inpatient Management.

Re-examine Exclusion Criteria:
- Patient < 3 months age
- Toxic appearance (consider sepsis)
- Diarrhea > 7 days (consider chronic disease, bacillary enteritis)
- Bloody diarrhea (consider HUS)
- Comorbid conditions (MCC, renal failure, cardiac disease)
- Bilious emesis (consider bowel obstruction)
- On diuretic therapy
- Hyponatremia (<130 mEq/L) or Hypermagnesemia (>155 mEq/L) if electrolytes checked
- Acute surgical abdomen

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Return to Phase II
Secretary Diarrhea

- No clear guidelines exist for when to consider treatment for secretory diarrhea.
- *Special consideration should be made for children with high stool output exceeding 30 mL/kg/day, especially if coupled with low or no oral intake* [Expert Opinion]
- *These patients may have secretory diarrhea, and might require a period of hyperalimentation, as well as nothing by mouth for a period of time.* [Expert Opinion]
Patients will be assessed for transition from IV or NG rehydration to oral rehydration therapy at 08:00am.

- This assessment will be performed by nursing at 08:00am for any patient admitted after 16:00 the previous day.
- This nursing assessment will determine if a patient is ready for an ORT challenge.
- Patients admitted between 08:00am and 16:00pm will be assessed after 4 hours of IV or NG fluid rehydration.

Inpatient Management: Assessment at 08:00am (Cont’d)

- During the nursing assessment, the patient will be assessed for:
  o total input and output, and
  o number of episodes of emesis or vomiting.
- If the total input > output and there is ≤ 1 episodes of vomiting, then IV or NG fluid rehydration will be stopped and the patient will be transitioned to an ORT challenge.
Inpatient Management: Not ready for oral challenge

- If, during the nursing assessment, the patient is deemed not ready for an ORT challenge, then the patient will continue with the previously ordered IV or NG rehydration.
- The patient will then be reassessed every 4 hours for potential ORT challenge.
- On-going losses from stool or emesis can be replaced at a 1:1 ratio with normal saline (NS) if the patient is receiving IV rehydration or Pedialyte if the patient is receiving NG rehydration.

Inpatient Management: Ready for oral challenge

- If, during the nursing assessment, the patient is determined to be ready for an ORT challenge, then:
  - the IV or NG fluids will be stopped, and
  - the ORT challenge will begin at the same doses as previously noted in Phase I (ED/Urgent Care): ORT
    - 5 mL every 5 minutes if < 10 kg
    - 10 mL every 5 minutes if ≥ 10 kg
  - Patients make choose to skip ORT and resume a regular diet, or lactose-free formula if predominantly formula fed

NOTE: The starting amount of ORT to be given during the ORT challenge is based on the patient’s admitted weight

Inpatient Management: ORT or Regular Diet Challenge

- Success or failure of the ORT/regular diet challenge is determined by the presence or absence of emesis
- If there is no emesis after 20 minutes, then the ORT challenge volumes can be doubled:
  - 10 mL every 5 minutes if < 10 kg
  - 20 mL every 5 minutes if ≥ 10 kg
- If emesis occurs at any point during the ORT challenge, then hold ORT for 20 minutes and restart at the initial rate.
Inpatient Management: Discharge Diet

- The goal in feeding during AGE is to return the patient to their regular and usual diet as soon as possible.

- *Continue to offer feedings (child’s preferred diet) to children with AGE.*
  
  5 [Guideline] If a child has previously been breast feeding, continue breast feeding.  
  
  5 [Guideline] Do not use the bread, rice, apple, toast (BRAT) diet; similarly, do not use other restrictive or progressive diets.  
  
  5, 8 [Guideline] Avoid use of fruit juices and carbonated sodas.  
  
  1, 5, 9 [Guideline] If diarrhea is the chief complaint in a child who is predominantly formula-fed, use a lactose-free formula until symptoms improve.  

Return to Phase III
**Discharge Criteria**

- Sufficient rehydration as indicated by weight gain OR normal respiratory rate, capillary refill, and skin turgor
- IV or NG fluids not required
- Tolerating ORT or regular diet
- Adequate family teaching
- Follow-up established

- Although weight gain can be a sign of readiness for discharge, it is not a required criteria
- Readiness for discharge can also be indicated by resolution of clinical symptoms such as tachypnea, prolonged capillary refill, and abnormal skin turgor

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**Discharge Instructions**

- Patients will be discharged:
  - To continue either ORT, or their regular, preferred diet, or lactose-free formula if predominantly formula fed
  - With an “Acute Gastroenteritis and Oral Rehydration Therapy” handout (PE636)
  - With a prescription for lactobacillus, to be taken for a total of 5 days

- Continue ORT, or
- Resume regular diet, or lactose free formula if predominantly formula fed
- Acute Gastroenteritis And Oral Rehydration Therapy handout (PE636)
- Lactobacillus for 5 days
Inpatient Management: Discharge Diet—Lactose-free formula

- There is a theoretical concern that intestinal damage during an episode of AGE will result in a loss of the intestinal brush border and result in a temporary inability to tolerate lactose.

- Although previous guidelines did not recommend the use of lactose-free formulas, a recent Cochrane review suggests that for children who are primarily bottle fed, a change to a lactose-free formula/diet will result in earlier resolution of diarrhea. 6

NOTE: The recommendation to use a lactose-free formula in predominantly formula fed infant is new for 2015.
Objective
To standardize the care of patients presenting to the emergency department or admitted to the hospital with acute gastroenteritis.

Additional Objectives
- Decrease use of inappropriate medications and laboratory studies
- Standardize the process for ORT challenge (timing, amounts, escalation schedule)
- Facilitate rapid transition from IV or NG fluids back to regular diet

Recommendations

**Laboratory testing**
1. Do not routinely perform stool cultures.
2. Check electrolytes only in moderately dehydrated children who are requiring the initiation of IV fluids.

**Assessment of dehydration**
3. Use percentage loss of body weight to best measure the degree of dehydration.
4. Use the following 3 individual examination signs to best assess for dehydration of 5% or greater: prolonged capillary refill time, abnormal skin turgor, and abnormal respiratory pattern.
5. Classify patients into subgroups (no or minimal dehydration, moderate dehydration, or overt shock) to guide appropriate treatment.

**Use of IV or NG fluid resuscitation**
6. Involve the caregivers in the decision to use IV fluids versus NG rehydration to treat dehydration in patients who are unsuccessful at oral rehydration.

**Return to regular diet**
7. Continue to offer feedings (child’s preferred diet) to children with AGE. If a child has previously been breast feeding, continue breast feeding.
8. Do not use the bread, rice, apple, toast (BRAT) diet or other restrictive or progressive diets.
9. Avoid use of fruit juices, sports drinks, and carbonated sodas.
10. If diarrhea is the chief complaint in a child who is predominantly formula-fed, use a lactose-free formula until symptoms improve.

**Medications**
11. Use ondansetron (oral, NG, or IV) only in moderately dehydrated patients presenting to the ED or Urgent Care.
12. Prescribe probiotics in the form of Lactobacillus GG as an adjunctive treatment in the management of children with diarrhea from acute gastroenteritis. Start immediately and treat for 5-7 days.
13. Do not prescribe anti-diarrheal medications for the treatment of AGE.

Rationale
- **Safety** will be enhanced by reducing the need for unnecessary medication use, laboratory testing, or hospitalization.
- **Cost** per episode of care is predicted to be reduced by decreasing use of medications and laboratory studies as clinically appropriate, as well as by standardizing the use of ORT in moderately dehydrated patients before initiating IV or NG fluids.
Executive Summary

• **Delivery** of care will be improved by standardizing the process for ORT challenge and by expediting patient flow on the inpatient unit through frequent nursing reassessment of the patient’s hydration status.

• **Quality** of care will be improved by promoting timely management of AGE in the ED, UC, and inpatient units, while reducing the need for unnecessary lab work and medication use, and by facilitating close follow up between the primary care provider and the patient upon discharge.

• **Engagement** is founded on the premise that this pathway was developed by providers from multiple disciplines and practice settings and incorporates feedback from family members.

**Evidence**
Studies were identified by searching electronic databases using search strategies developed and executed by a medical librarian. A search was performed for topics including gastroenteritis, dysentery, enteritis, adenoviridae infections, rotavirus infections or rotavirus, looking back from 2007 to 2014. The results were systematically reviewed by the clinical pathway team, and 12 articles total were included to inform the recommendations.

**Implementation Items**
• Revised algorithm for care, including Urgent Care setting
• Revised web-based training module for physicians and ARNPs
• Revised patient handouts and discharge teaching/planning
• Revised nursing teaching protocol

**Metrics Plan**
• CSW Core Metrics
  o Count of inpatient and observation discharges
  o Median Length of Stay
  o Percent of patients with any of the specified powerplans
  o Average charges per case
  o Readmission
• Electrolytes utilization
• Lactobacillus prescription rate
• Use of NG hydration
• ED LOS or Admit LOS when using NG hydration
• Appropriate formula ordered

**PDCA Plan**
The clinical pathway team will meet quarterly to review metrics, medical literature, and any use of pathway-related tools.

**Revision History**
Date Approved: June 2015
Next Review Date: June 2020
Executive Summary

Approved by the CSW Acute Gastroenteritis Team on June 17th, 2015

CSW Acute Gastroenteritis Team:

<table>
<thead>
<tr>
<th>Position</th>
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<tbody>
<tr>
<td>Hospital Medicine, Owner</td>
<td>James O’Callaghan, MD, FAAP, SFHM</td>
</tr>
<tr>
<td>Emergency Medicine, Owner</td>
<td>Sabreen Akhter, DO, DTM</td>
</tr>
<tr>
<td>Urgent Care, Consultant</td>
<td>Bryan Dryer, MD</td>
</tr>
<tr>
<td>Emergency Medicine CNS</td>
<td>Sara Fenstermacher, RN</td>
</tr>
<tr>
<td>Medical CNS</td>
<td>Kristi Klee, MSN, RN-BC</td>
</tr>
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Clinical Effectiveness Team:

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<tr>
<td>Consultant</td>
<td>Jeffrey Foti, MD</td>
</tr>
<tr>
<td>Project Leader</td>
<td>Elizabeth Austin, MPH</td>
</tr>
<tr>
<td>KM Analyst</td>
<td>James Johnson</td>
</tr>
<tr>
<td>CIS Informatician</td>
<td>Michael Leu, MD, MS, MHA</td>
</tr>
<tr>
<td>CIS Analyst</td>
<td>Heather Marshall</td>
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<tr>
<td>Librarian</td>
<td>Susan Klawansky, MLS</td>
</tr>
<tr>
<td>Program Coordinator</td>
<td>Ashlea Tade</td>
</tr>
</tbody>
</table>
Title: Acute Gastroenteritis (AGE) Pathway

Authors:
- Seattle Children’s Hospital
- James O’Callaghan
- Sabreen Akhter
- Elizabeth Austin
- Bryan Dryer
- Sara Fenstermacher
- Jeffrey Foti
- Kristi Klee
- Michael Leu

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Example:
Completion qualifies you for 1 hour of Category II CME credit. If you are taking this self-assessment as a part of required departmental training at Seattle Children’s Hospital, you MUST logon to Learning Center.

(1) Which one of the following is NOT an exclusion criteria?
   a) toxic appearance
   b) bloody diarrhea
   c) rotavirus infection
   d) diarrhea for more than 7 days
   e) bilious emesis

(2) Which of the following statements concerning probiotics is/are TRUE?
   a) probiotics are considered adjunctive therapy.
   b) probiotics may be more effective for rotavirus diarrhea, compared to all-cause diarrhea
   c) yogurt is considered a good source of probiotics
   d) a and b
   e) a and c
   f) b and c
   g) all of the above

(3) The historical BRAT diet (consisting of bananas, rice, applesauce, and toast) is unnecessarily restrictive.
   a) true
   b) false

(4) Which of the following statements is/are TRUE concerning the clinical assessment of dehydration in AGE?
   a) Prolonged capillary refill time, abnormal skin turgor, and respiratory rate are the best individual examination measures
   b) A normal bicarbonate concentration may be useful in ruling out dehydration
   c) Acute body weight change is considered the gold standard measure of dehydration
   d) It is recommended that the history and physical examination be the primary basis for the diagnosis of AGE
   e) all of the above

(5) Which of the following discharge criteria are TRUE?
   a) tolerating ORT and/or regular diet
   b) tolerating BRAT diet
   c) medical follow up is available via telephone or office visit
   d) a and b
   e) a and c
   f) b and c
   g) all of the above

(6) Ondansetron has been shown to be a safe and effective therapy in hospitalized children.
   a) true
   b) false

(7) Which of the following is(are) the recommended diet(s) to prevent or limit dehydration?
   a) BRAT diet
   b) clear liquid diet
   c) Paleo diet
   d) regular diet
   e) a and b
   f) a and d
   g) b and d
   h) a, b, and d

(8) Which of the following liquids are appropriate for use in oral rehydration therapy?
   a) Pedialyte®
   b) Gatorade©
   c) Gatorade© with added salt, in the ratio of 1L Gatorade© + 1 tbsp salt
   d) a and b
   e) a and c
   f) all of the above
(1) The correct answer is (c); all other choices except rotavirus infection are exclusion criteria.

(2) The correct answer is (d); probiotics are considered adjunctive therapy and may be more effective for rotavirus diarrhea, compared to all-cause diarrhea.

(3) The correct answer is (a).

(4) The correct answer is (e); all of the above statements are true about the clinical assessment of dehydration in AGE.

(5) The correct answer is (e); tolerating ORT and/or a regular diet and ensuring that adequate medical follow up is available by telephone or office visit are discharge criteria for patients with a diagnosis of AGE.

(6) The correct answer is (b); while studies suggest a benefit of ondansetron in the outpatient or emergency room setting, ondansetron has not been shown to be an effective therapy in hospitalized children.

(7) The correct answer is (d); only a patient’s usual or regular diet is the recommended diet to prevent or limit dehydration.

(8) The correct answer is (a); only Pedialyte® is appropriate for use in oral rehydration therapy.
This pathway was developed through local consensus based on published evidence and expert opinion as part of Clinical Standard Work at Seattle Children’s. Pathway teams include representatives from Medical, Subspecialty, and/or Surgical Services, Nursing, Pharmacy, Clinical Effectiveness, and other services as appropriate.

When possible, we used the GRADE method of rating evidence quality. Evidence is first assessed as to whether it is from randomized trial or cohort studies. The rating is then adjusted in the following manner (from: Guyatt G et al. J Clin Epidemiol. 2011;4:383-94.):

- Quality ratings are **downgraded** if studies:
  - Have serious limitations
  - Have inconsistent results
  - If evidence does not directly address clinical questions
  - If estimates are imprecise OR
  - If it is felt that there is substantial publication bias

- Quality ratings are **upgraded** if it is felt that:
  - The effect size is large
  - If studies are designed in a way that confounding would likely underreport the magnitude of the effect OR
  - If a dose-response gradient is evident

Guideline – Recommendation is from a published guideline that used methodology deemed acceptable by the team.

Expert Opinion – Our expert opinion is based on available evidence that does not meet GRADE criteria (for example, case-control studies).

**Quality of Evidence:**
- 🌟🌟🌟🌟 High quality
- 🌟🌟🌟 Moderate quality
- 🌟🌟 Low quality
- 🌟🌟🌟 Very low quality

Guideline
Expert Opinion
Summary of Version Changes

- **Version 1.1 (11/08/2011)**: Go Live
- **Version 1.2 (07/28/2014)**: Administrative update
- **Version 2.0 (06/17/2015)**: Periodic review; updated literature search, recommendations, and pathway tools
Medical Disclaimer

Medicine is an ever-changing science. As new research and clinical experience broaden our knowledge, changes in treatment and drug therapy are required.

The authors have checked with sources believed to be reliable in their efforts to provide information that is complete and generally in accord with the standards accepted at the time of publication.

However, in view of the possibility of human error or changes in medical sciences, neither the authors nor Seattle Children’s Healthcare System nor any other party who has been involved in the preparation or publication of this work warrants that the information contained herein is in every respect accurate or complete, and they are not responsible for any errors or omissions or for the results obtained from the use of such information.

Readers should confirm the information contained herein with other sources and are encouraged to consult with their health care provider before making any health care decision.
Literature Search Strategy

Search Methods, Acute Gastroenteritis, Clinical Standard Work

Studies were identified by searching electronic databases using search strategies developed and executed by a medical librarian, Susan Klawansky. Searches were performed in November 2014 in the following databases – on the Ovid platform: Medline and Cochrane Database of Systematic Reviews; elsewhere: Embase, Clinical Evidence, National Guideline Clearinghouse, TRIP and Cincinnati Children’s Evidence-Based Care Recommendations. Retrieval was limited to 2007 to current, ages 0-18, and English language. In Medline and Embase, appropriate Medical Subject Headings (MeSH) and Emtree headings were used respectively, along with text words, and the search strategy was adapted for other databases as appropriate. Concepts searched were gastroenteritis, dysentery, enteritis, adenoviridae infections, rotavirus infections or rotavirus. All retrieval was further limited to certain evidence categories, such as relevant publication types, index terms for study types and other similar limits. Additional articles were identified by team members and added to results.

Identification

129 records identified through database searching

1 additional records identified through other sources

Screening

130 records after duplicates removed

130 records screened

84 records excluded

Eligibility

46 records assessed for eligibility

34 full-text articles excluded, 5 did not answer clinical question, 6 did not meet quality threshold, 1 outdated relative to other included study, 19 duplicates, 3 other

Included

12 studies included in pathway

Flow diagram adapted from Moher D et al. BMJ 2009;339:bmj.b2535


8. National GC. Evidence-based care guideline for prevention and management of acute gastroenteritis (AGE) in children aged 2 months to 18 years. http://www.guideline.gov/content.aspx?id=35123&search=%22acute+gastroenteritis%22+and+(child*+or+pediatr*+or+paediatr*):.


