

DATE: July 2021

TEXAS CHILDREN'S HOSPITAL EVIDENCE-BASED OUTCOMES CENTER Acute Gastroenteritis (AGE) (Acute Vomiting and/or Diarrhea) Evidence-Based Guideline

<u>**Definition:**</u> Acute gastroenteritis (AGE) is a decrease in the consistency of stools (loose or liquid) and/or an increase in the frequency of stools (typically \geq 3 in 24 hours), with or without fever or vomiting. However, vomiting alone is typical of early presentation. Duration of illness is typically less than 14 days. ⁽¹⁻³⁾

Epidemiology: In the United States, an estimated 9.4 million episodes of foodborne illness, 55,961 hospitalizations, and 1,351 deaths occur each year due to foods consumed that were contaminated with 31 known agents of foodborne disease. ⁽⁴⁾ In February 2006, routine use of a pentavalent human-bovine rotavirus vaccine was recommended. ⁽⁵⁻⁷⁾ Since these recommendations have been implemented, there has been a delayed and shorter season. ^(8,9) In children seen in the Emergency Center at Texas Children's Hospital, pentavalent rotavirus vaccine (RV5) was noted to be highly effective in preventing rotavirus disease. ⁽⁸⁾

Etiology: The most common causes of AGE are infectious agents. In the developed world, viruses are responsible for 70 to 80% of infectious diarrhea cases. Rotavirus and norovirus are the leading viral pathogens with nearly every child in the U.S. being infected with rotavirus by 5 years of age. ^(5,10) Since the introduction of the Rotavirus vaccine, norovirus has become the leading cause of medically attended acute gastroenteritis. ⁽⁹⁾ Various bacterial pathogens account for another 10 to 20% of cases and as many as 10% may be attributable to diarrheagenic *Escherichia coli*. ⁽¹¹⁾ Parasitic organisms such as *Giardia* species cause fewer than 10% of cases. Incidence is affected by climate and season. Other factors that increase the risk of AGE in children include day care attendance and impoverished living conditions with poor sanitation. Community-acquired *Clostridium difficile* infections are on the rise.

Inclusion Criteria

Age ≥60 days to 17 years Healthy children without underlying conditions Clinical findings of AGE

Exclusion Criteria

Toxic appearance Episodes of diarrhea lasting >14 days

Differential Diagnosis

Ingestion Food-borne illness Intussusception Appendicitis Urinary tract infection (UTI) Bowel obstruction Extra-intestinal infection Allergic reaction

Diagnostic Evaluation (12)

- History: Assess for
- Age of child
- Developmentally appropriate behavior
- ≥3 loose or watery stools/day
- Onset, frequency, quantity, and character (e.g., black, bloody) of vomiting/diarrhea

- Travel and/or day care exposure
- Dietary changes
- Vaccination status (especially Rotavirus vaccine)
- Last episode of vomiting
- Volume and frequency of urine output
- Use of antibiotics

Rotavirus disease typically begins abruptly. Vomiting often precedes the onset of diarrhea. ⁽⁵⁾ Norovirus is characterized by acute onset of nausea, vomiting, abdominal cramps, and diarrhea. Vomiting can appear alone in norovirus illness. ⁽¹³⁾

Physical Examination

Severity of dehydration (none/mild, moderate, or severe) is the key factor in determining the severity of AGE which is primarily based on the child's dehydration status. Management requires a rapid risk assessment of dehydration. (12,14,15)

A complete physical exam should be performed assessing for:

- Weight loss (pre-illness weight minus acute body weight)
- Prolonged capillary refill time (>2 seconds)*
- Dry mucous membranes*
- Absent tears*
- Poor overall appearance*
- Abnormal skin turgor
- Sunken eyes
- Abnormal radial pulse
- Tachycardia (HR >150; scale validated in children 1 month to 5 years) ⁽¹⁶⁾
- Abnormal respirations
- Decreased urine output

Accurate body weight is considered the gold standard in determining fluid deficit (pre-illness weight minus acute body weight). ^(12,16)

*The presence of \geq 3 of 4 predicts a fluid deficit \geq 10%. ⁽¹⁶⁾ Combination of clinical findings improves diagnosis. ^(14,16,17)

Laboratory Tests

Routine laboratory tests are **NOT** recommended for children with mild/moderate dehydration. ^(14,18,19,21-24)

Consider stool studies (stool culture and stool ova and parasites) with bloody stools, prolonged symptoms, suspicion of epidemic, travel exposure, and/or age <3 months. ^(23,25-28)

Consider *C. difficile* if age >2 years, previous use of antibiotics, previous *C. difficile* infection, and/or hospitalization within the last 30 days.

Serum sodium bicarbonate is an unreliable predictor for determining the severity of dehydration. ⁽¹⁴⁾

Consider urinalysis (UA) with micro and culture when concerned for UTI.

Urine Specimen for Urinalysis and Culture§ (UTI Guideline)

Non-toilet trained children: transuretheral catherization

Toilet trained children: midstream clean catch

§Obtained by non-invasive method. If positive, invasive method may be necessary.

Critical Points of Evidence

Evidence Supports

Use of oral rehydration solutions (ORS) for treatment of mild/moderate gastroenteritis if tolerated ^(29,30) Using low-osmolarity ORS (270 mmol/L or less) for oral rehydration therapy ⁽³¹⁾ Resuming the child's regular age-appropriate diet when tolerated, continuing breastfeeding during rehydration therapy, and restricting fruit juices and carbonated beverages ⁽³²⁻³⁴⁾ Use of antiemetics in the management of children with AGE who are vomiting ⁽³⁵⁻⁴¹⁾ Probiotics as adjunctive therapy ⁽⁴²⁻⁴⁹⁾ Use of the Gorelick score to evaluate dehydration ^(17,20,50-55)

Evidence Lacking/Inconclusive

Administration of IV maintenance fluids at 1.5 or 2 times maintenance ^(56,57) Preferred routes of rehydration between nasogastric tube versus IV ⁽³⁰⁾ Use of specific barrier creams to prevent diaper dermatitis in non-toilet trained children

Evidence Against

Use of ondansetron in patients with prolonged QT or medications that may prolong QT interval ⁽⁵⁸⁾ Use of routine lab tests including tests for specific pathogens ^(14,18-24) Use of stool studies to predict patient outcomes ^(23-27,59-62) Use of anti-motility agents for routine management of acute diarrhea ⁽⁶³⁾ Using high osmolarity (>300 mmol/L) liquids for oral rehydration therapy ⁽³⁰⁾

Condition-Specific Elements of Clinical Management

<u>General</u>: The clinical picture of children with acute gastroenteritis is highly variable and determination of etiologies is difficult. The severity of dehydration is an important factor to consider in managing this disease.

Treatment Recommendations

Mild Dehydration, Rehydration

Infants/children should continue their regular diet \pm ORS if tolerating oral fluids. ⁽³¹⁻³³⁾ If a child is vomiting, begin small frequent feedings of ORS 1ml/kg every 5 minutes and consider an antiemetic. ^(31-40,67)

Moderate Dehydration, Rehydration

Infants/children should be given ORS. ^(28,29) If a child is vomiting, begin small frequent feedings of ORS 1ml/kg every 5 minutes and consider an antiemetic. ^(31-40,67) Reassess at 1 hour or until sufficient rehydration is achieved.

Oral Rehydration Therapy Failed or Severe Dehydration If an infant/child is unable to tolerate oral feeding and/or is at risk for being unable to maintain hydration status, then a bolus of isotonic intravenous fluids (IVF) should be considered. May require up to 3 boluses. Each bolus should be followed by a repeat physical exam. In addition to boluses, maintenance fluid therapy should be considered (See Table I).⁽⁶⁴⁾ Consider admission to Observation for continued rehydration if output is greater than intake.

Table I. Maintenance Rates

Holliday-Segar Method ⁽⁶⁴⁾
4 mL/kg/h for 1 st 10 kg of body weight
2 mL/kg/h for 2 nd 10 kg of body weight
1 mL/kg/h for each kg over 20 kg

Oral Feeding During Rehydration

- Continued administration of ORS for infants/children who are mildly/moderately dehydrated is recommended. An ageappropriate diet includes complex carbohydrates and foods that are low in sugar, fat, and caffeine can be given when tolerated (See Table II).
- For infants, the use of breastmilk or formula should continue on demand.

<u>Admission Criteria</u> Unable to maintain hydration status via oral route

Consults and Referrals

Consultation with a Gastroenterology specialist and/or Infectious Disease is appropriate for chronic diarrhea or *C. difficile* infection.

Infection Control

Contact precautions are required for all children with diarrhea.

Other Therapy Alternatives

- In children, antibiotic therapy is typically not required for most cases of dysentery and should be driven by stool culture results.
- If UTI is suspected, refer to UTI Guideline.
- If the patient is not responding to fluid therapy, auscultate heart and lung sounds and evaluate for heart failure. Refer to <u>Acute</u> <u>Decompensated Heart Failure Guideline</u>.
- Depending on stool culture results Antibiotic treatment is indicated for infants <6 months.
- Consider the use of antiemetic for treatment of nausea and vomiting ⁽³⁴⁻⁴⁰⁾
- Use probiotics for treatment of diarrhea (See Table III.) Effective organism and dosing for treatment of acute gastroenteritis:
- Saccharomyces boulardii of at least 250 mg per day (equivalent to 5 billion colony forming units per day), ^(42,43,46)
- Lactobacillus GG at 10 billion colony forming units per day (41,42,45,47)
- Lactobacillus reuteri of at least 100 million colony forming units per day. ^(44,48)
- In non-toilet trained children, maintain skin integrity with episodes of frequent diarrhea with frequent diaper changes and skin care.

Caregiver Education

Encourage frequent hand-washing for the patient, family, and caregivers ^(65, 66)
Encourage continued breastfeeding with infants ⁽³¹⁾
Limit exposure to other children (e.g., day care)
Encourage fluid intake until symptoms resolve.
Advance to regular diet once tolerated.
In incontinent children, maintain skin integrity with frequent diaper changes.

Discharge Criteria

Sufficient rehydration achieved Tolerating oral fluids Appropriate support system (e.g., primary care physician [PCP], caregivers)

Follow-Up Care

Children diagnosed with AGE with persistent or worsening symptoms (e.g., decreased urine output, diarrhea, and/or vomiting returns) should follow up with their PCP

Prevention

Importance of strict hand-washing ^(65,66) Rotavirus vaccine per recommended immunization schedule ⁽⁵⁾

Measures

- # of children who received rotavirus vaccine prior to this illness
- Predictive accuracy of dehydration assessment scoring
- % of children receiving antibiotics inappropriately
- % of positive stool cultures when stool cultures are obtained
- Unscheduled visits to the Emergency Center/PCP during the course of this illness
- # of unnecessary diagnostic tests performed (e.g., labs, stool cultures) when criteria not met
- Results and treatment indications of stool cultures



Clinical standards are developed for 80% of the patient population with a particular disease. Each practitioner must use his/her clinical judgment in the management of any specific patient.

© Evidence-Based Outcomes Center Texas Children's Hospital

Acute Gastroenteritis: Diet Suggestions

For the next 2 weeks at home, children should return to eating a regular healthy diet. Sometimes, greasy and high sugar foods are not tolerated well and should be avoided.

- For infants: Resume breastfeeding or drinking formula
- For toddlers and older: Resume formula or milk, and age appropriate foods (i.e. pureed foods, solids)

Stay hydrated by drinking enough fluids, especially water, throughout the day.

- Avoid caffeine (tea, energy drinks)
- Avoid all carbonated (bubbly drinks such as sodas, carbonated waters)
- Avoid any sweet drinks (fruit juice, lemonade, fruit punch, even diluted beverages)
- Avoid sports drinks

It is OK to have sugar-free liquids (Crystal Light, sugar free popsicles). It is also OK to have "hydration solutions" such as Infalyte, Pedialyte, and Drip Drops.

Food Group	Тір	Suggestions
Fruit	Select fresh, frozen, canned, and dried fruit more often than juice	Banana, apple slices, cantaloupe, watermelon, pears, grapes, kiwi, mango, strawberries
Vegetables	Aim for variety every day; pick vegetables from several subgroups: dark green, red & orange, beans and peas, starchy, and other veggies	Steamed broccoli, sweet potato, spinach, squash, green beans, peas, mushrooms, bell peppers, cucumber
Dairy	Include fat-free and low-fat dairy foods every day	2% milk (whole milk if under 2 years), cheese, yogurt, cottage cheese, smoothies
Protein Foods	Aim for variety—choose seafood, lean meat and poultry, beans, peas, nuts, and seeds each week	Turkey, baked fish, peanut butter, nuts, tuna, black beans, lentils, eggs, tofu
Grains	Make at least half your grains whole grains	Whole grain cereal, whole wheat bread, whole grain crackers, rice, pasta, museli, oatmeal

Regular Healthy Age-Appropriate Diet

https://www.myplate.gov/

Table III. Probiotics Recommendations

Recommended Probiotic Products Meeting Efficacy Criterion for Acute Gastroenteritis					
Brand Name	Active Ingredient	Amount	Preparation	Dose	Administration
Culturelle [®] Capsules	Lactobacillus rhamnosus GG	10 billion CFUs per capsule	Capsule	1 capsule daily	Swallow whole
Culturelle [®] Kids Chewable Tablets	Lactobacillus rhamnosus GG	5 Billion CFUs per chewable tablet	Chewable Tablet	2 tablets daily	Chew tablet
Culturelle [®] Kids Packets	Lactobacillus rhamnosus GG	5 Billion CFUs per packet	Granules	2 packets daily	Empty packet into cool food or drink. Mix until dissolved. Do not add to warm or hot food or beverage.
Florastor [®] Capsules	Saccharomyces boulardii Iyo	250 mg per capsule	Capsule	1-3 capsules daily	 For immediate administration, capsules may be: Swallowed whole Emptied directly onto tongue and followed by 4 oz of water/juice Sprinkled over semi-solid food (i.e. applesauce, yogurt, etc.) Dissolved in liquid
Florastor [®] Kids Packets with tutti- frutti flavoring	Saccharomyces boulardii Iyo	250 mg per packet	Powder	1-3 packets daily	 For immediate administration, packets may be: Emptied directly onto tongue and followed by 4 oz of water/juice Sprinkled over semi-solid food (i.e. applesauce, yogurt, etc.) Dissolved in liquid
Fleet [®] Pedia-Lax Probiotic Yums Chewable Tablet	Lactobacillus reuteri DSM 17938	100 million CFUs per tablet	Chewable Tablet	1 tablet daily	Chew tablet

Note: Use dietary supplements (probiotics) containing live bacteria or yeast with caution in immunocompromised patients. The following products do not meet the efficacy criterion to reduce the duration of symptoms of acute gastroenteritis: yogurt, kefir, Nature's Bounty[®] Advanced Probiotic 10, Align[®], Bacid[®], Floranex[®], Garden of Life[®] Raw Probiotics Powder for Kids, Lactinex[®], Lacto-Pectin[®], Megadophilus[®], MoreDophilus[®], Pearls[®], RisaQuad[®], Superdophilus[®], Ultimate Flora[®] Kids Probiotic Supplement, VSL[®] #3 (or Junior), or Zarbee's Naturals[®] Children's Probiotic Supplement. If the product in question is not listed, please review the active ingredients to see if it contains the recommended probiotic strains and amount of culture forming units.

© Evidence-Based Outcomes Center Texas Children's Hospital

References

- Acute Gastroenteritis Guideline Team, Cincinnati Children's Hospital Medical Center: Evidence-based care guideline for prevention and management of acute gastroenteritis in children age 2 months to 18 years, http://www.cincinnatichildrens.org/service/j/anderson-center/evidence-basedcare/gastroenteritis/ Guideline 5, pages 1-20, Dec 21, 2011.
- Guarino, A., Ashkenazi, S., Gendrel, D., Lo Vecchio, A., Shamir, R., & Szajewska, H. (2014). European Society for Pediatric Gastroenterology, Hepatology, and Nutrition/European Society for Pediatric Infectious Diseases evidence-based guidelines for the management of acute gastroenteritis in children in Europe: update 2014. *Journal of Pediatric Gastroenterology and Nutrition, 59*(1), 132-152. doi: 10.1097/mpg.00000000000375
- 3. NICE, National Institute for Health and Clinical Excellence (2009). *Diarrhoea and Vomiting Caused by Gastroenteritis: Diagnosis, Assessment and Management in Children Younger than 5 Years* (Vol. 2009). London: National Collaborating Centre for Women's and Children's Health.
- Scallan, E., Hoekstra, R. M., Angulo, F. J., Tauxe, R. V., Widdowson, M. A., Roy, S. L., Griffin, P. M. (2011). Foodborne illness acquired in the United States--major pathogens. *Emerging Infectious Diseases*, 17(1), 7-15. doi: 10.3201/eid1701.091101p1
- 5. Centers for Disease Control and Prevention. Prevention of rotavirus gastroenteritis among infants and children. MMWR 2006;56(No. RR-12):1-13.
- Committee on Infectious Diseases. (2007). Prevention of rotavirus disease: Guidelines for use of rotavirus vaccine. *Pediatrics*, *119*(1), 171-182.
 Centers for Disease Control and Prevention. Delayed onset and diminished magnitude of rotavirus activity- United States, November 2007- May 2008. MMWR Early Release 2008;57:1-3.
- Boom, J., Tate, J., Sahni, L., Rench, M., Hull, J. J., Gentsch, J. R., et al. (2008). Effectiveness of pentavalent rotavirus vaccine (RV5) in a large urban population in the United States. Pediatrics, 125(2), 199-207.
- 9. Payne, D. C., Vinje, J., Szilagyi, P. G., Edwards, K. M., Staat, M. A., Weinberg, G. A., Parashar, U. D. (2013). Norovirus and medically attended gastroenteritis in U.S. children. New England Journal of Medicine, 368(12), 1121-1130. doi: 10.1056/NEJMsa1206589
- Patel, M., Widdowson, M.-A., Glass, R. I., Akazawa, K., Vinje, J., & Parashar, U. D. (2008). Systematic literature review of role of noroviruses in sporadic gastroenteritis. *Emerging Infectious Diseases*, 14(8), 1224-1231.
- 11. Cohen, M. B., Nataro, J. P., Bernstein, D. I., Hawkins, J., Roberts, N., & Staat, M. A. (2005). Prevalence of diarrheagenic Escherichia coli in acute childhood enteritis: A prospective controlled study. *Journal of Pediatrics*, 146(1), 54-61.
- Centers for Disease Control and Prevention. Managing acute gastroenteritis among children: Oral rehydration, maintenance, and nutritional therapy. MMWR 2003;52(No. RR-16):1-16.
- Centers for Disease Control and Prevention. Norwalk-like viruses: Public health consequences and outbreak management. MMWR 2001;50(No. RR-09):1-18.
- 14. Steiner, M. J., DeWalt, D. A., & Byerley, J. S. (2004). Is this child dehydrated? Journal of the American Medical Association, 291(22), 2746-2754.
- 15. Goldman, R. D., Friedman, J. N., & Parkin, P. C. (2008). Validation of the clinical dehydration scale for children with acute gastroenteritis. *Pediatrics*, 122(3), 545-549.
- 16. Gorelick, M. H., Shaw, K. N., & Murphy, K. O. (1997). Validity and reliability of clinical signs in the diagnosis of dehydration in children. *Pediatrics*, *99*(5), E6.
- 17. Jauregui, J., Nelson, D., Choo, E., Stearns, B., Levine, A. C., Liebmann, O., & Shah, S. P. (2014). External validation and comparison of three pediatric clinical dehydration scales. *PLoS One*, *9*(5), e95739. doi: 10.1371/journal.pone.0095739
- Hayajneh, W. A., Jdaitawi, H., Al Shurman, A., & Hayajneh, Y. A. (2010). Comparison of clinical associations and laboratory abnormalities in children with moderate and severe dehydration. *Journal of Pediatric Gastroenterology and Nutrition*, 50(3), 290-294. doi: 10.1097/MPG.0b013e31819de85d
- 19. Hoxha, T. F., Azemi, M., Avdiu, M., Ismaili-Jaha, V., Grajqevci, V., & Petrela, E. (2014). The usefulness of clinical and laboratory parameters for predicting severity of dehydration in children with acute gastroenteritis. *Medical Archives*, *68*(5), 304-307. doi: 10.5455/medarh.2014.68.304-307
- Marcus, N., Mor, M., Amir, L., Mimouni, M., & Waisman, Y. (2007). The quick-read C-reactive protein test for the prediction of bacterial gastroenteritis in the pediatric emergency department. *Pediatric Emergency Care, 23*(9), 634-637. doi: 10.1097/PEC.0b013e31814a6a52
- Parkin, P. C., Macarthur, C., Khambalia, A., Goldman, R. D., & Friedman, J. N. (2010). Clinical and laboratory assessment of dehydration severity in children with acute gastroenteritis. *Clinical Pediatrics (Philadelphia), 49*(3), 235-239. doi: 10.1177/0009922809336670
- Steiner, M. J., Nager, A. L., & Wang, V. J. (2007). Urine specific gravity and other urinary indices: Inaccurate tests for dehydration. *Pediatric Emergency Care*, 23(5), 298-303. doi: 10.1097/01.pec.0000270162.76453.fa
- Sykora, J., Siala, K., Huml, M., Varvarovska, J., Schwarz, J., & Pomahacova, R. (2010). Evaluation of faecal calprotectin as a valuable non-invasive marker in distinguishing gut pathogens in young children with acute gastroenteritis. *Acta Paediatrica, 99*(9), 1389-1395. doi: 10.1111/j.1651-2227.2010.01843.x
- 24. Kaiser, P., Borte, M., Zimmer, K. P., & Huppertz, H. I. (2012). Complications in hospitalized children with acute gastroenteritis caused by rotavirus: A retrospective analysis. *European Journal of Pediatrics*, 171(2), 337-345. doi: 10.1007/s00431-011-1536-0
- Chen, C. C., Chang, C. J., Lin, T. Y., Lai, M. W., Chao, H. C., & Kong, M. S. (2011). Usefulness of fecal lactoferrin in predicting and monitoring the clinical severity of infectious diarrhea. World Journal of Gastroenterology, 17(37), 4218-4224. doi: 10.3748/wjg.v17.i37.4218
- Chen, C. C., Huang, J. L., Chang, C. J., & Kong, M. S. (2012). Fecal calprotectin as a correlative marker in clinical severity of infectious diarrhea and usefulness in evaluating bacterial or viral pathogens in children. *Journal of Pediatric Gastroenterology and Nutrition*, 55(5), 541-547. doi: 10.1097/MPG.0b013e318262a718
- 27. Duman, M., Gencpinar, P., Bicmen, M., Arslan, N., Ozden, O., Uzum, O., et al. (2015). Fecal calprotectin: Can be used to distinguish between bacterial and viral gastroenteritis in children? *American Journal of Emergency Medicine*, 33(10), 1436-1439. doi: 10.1016/j.ajem.2015.07.007
- 28. Liu, J., Kabir, F., Manneh, J., Lertsethtakarn, P., Begum, S., Gratz, J., et al. (2014). Development and assessment of molecular diagnostic tests for 15 enteropathogens causing childhood diarrhoea: A multicentre study. *Lancet Infectious Diseases*, 14(8), 716-724. doi: 10.1016/s1473-3099(14)70808-4
- 29. Craven, J. A., Campbell, L., & Martin, C. T. (2009). Waiting room oral rehydration in the paediatric emergency department. Irish Medical Journal, 102(3), 85-87.
- Hartling, L., Bellemare, S., Wiebe, N., Russell, K., Klassen, T. P., & Craig, W. (2006). Oral versus intravenous rehydration for treating dehydration due to gastroenteritis in children. *Cochrane Database of Systematic Reviews*(3), CD004390. doi: 10.1002/14651858.CD004390.pub2
- 31. Hahn, S., Kim, S., & Garner, P. (2002). Reduced osmolarity oral rehydration solution for treating dehydration caused by acute diarrhoea in children. *Cochrane Database of Systematic Reviews* (1), CD002847. doi: 10.1002/14651858.cd002847
- 32. Gregorio, G. V., Dans, L. F., & Silvestre, M. A. (2011). Early versus delayed refeeding for children with acute diarrhoea. *Cochrane Database of Systematic Reviews* (7), CD007296. doi: 10.1002/14651858.CD007296.pub2
- MacGillivray, S., Fahey, T., & McGuire, W. (2013). Lactose avoidance for young children with acute diarrhoea. Cochrane Database Syst Rev, 10, CD005433. doi: 10.1002/14651858.CD005433.pub2
- 34. Valois, S., Costa-Ribeiro, H., Jr., Mattos, A., Ribeiro, T. C., Mendes, C. M., & Lifshitz, F. (2005). Controlled, double-blind, randomized clinical trial to evaluate the impact of fruit juice consumption on the evolution of infants with acute diarrhea. *Nutrition Journal*, *4*, 23. doi: 10.1186/1475-2891-4-23
- 35. Al-Ansari, K., Alomary, S., Abdulateef, H., Alshawagfa, M., & Kamal, K. (2011). Metoclopramide versus ondansetron for the treatment of vomiting in children with acute gastroenteritis. *Journal of Pediatric Gastroenterology and Nutrition*, *53*(2), 156-160. doi: 10.1097/MPG.0b013e3182132d8d

Cochrane Database of Systematic Reviews(9), CD005506. doi: 10.1002/14651858.CD005506.pub5

- Freedman, S. B., Powell, E. C., Nava-Ocampo, A. A., & Finkelstein, Y. (2010). Ondansetron dosing in pediatric gastroenteritis: A prospective cohort, dose-response study. *Paediatric Drugs*, 12(6), 405-410. doi: 10.2165/11537770-00000000-00000
- Freedman, S. B., Tung, C., Cho, D., Rumantir, M., & Chan, K. J. (2012). Time-series analysis of ondansetron use in pediatric gastroenteritis. *Journal of Pediatric Gastroenterology and Nutrition*, 54(3), 381-386. doi: 10.1097/MPG.0b013e31822ecaac
- Gouin, S., Vo, T. T., Roy, M., Lebel, D., & Gravel, J. (2012). Oral dimenhydrinate versus placebo in children with gastroenteritis: A randomized controlled trial. *Pediatrics*, 129(6), 1050-1055. doi: 10.1542/peds.2011-2945
- Hervas, D., Armero, C., Carrion, T., Utrera, J. F., & Hervas, J. A. (2012). Clinical and economic impact of oral ondansetron for vomiting in a pediatric emergency department. *Pediatric Emergency Care*, 28(11), 1166-1168. doi: 10.1097/PEC.0b013e3182717358
- Qazi, K., BinSalleeh, H. M., Shah, U. H., AlGhamedi, N., Tamim, H., Mubasher, M., et al. (2014). Effectiveness of granisetron in controlling pediatric gastroenteritis-related vomiting after discharge from the ED. American Journal of Emergency Medicine, 32(9), 1046-1050. doi: 10.1016/i.aiem.2014.06.018
- 42. Aggarwal, S., Upadhyay, A., Shah, D., Teotia, N., Agarwal, A., & Jaiswal, V. (2014). Lactobacillus GG for treatment of acute childhood diarrhoea: An open labelled, randomized controlled trial. *Indian Journal of Medical Research, 139*(3), 379-385.
- 43. Allen, S. J., Martinez, E. G., Gregorio, G. V., & Dans, L. F. (2010). Probiotics for treating acute infectious diarrhoea. *Cochrane Database of Systematic Reviews*(11), CD003048. doi: 10.1002/14651858.CD003048.pub3
- 44. Feizizadeh, S., Salehi-Abargouei, A., & Akbari, V. (2014). Efficacy and safety of Saccharomyces boulardii for acute diarrhea. *Pediatrics*, 134(1), e176-e191. doi: 10.1542/peds.2013-3950
- Dinleyici, E. C., & Vandenplas, Y. (2014). Lactobacillus reuteri DSM 17938 effectively reduces the duration of acute diarrhoea in hospitalised children. Acta Paediatrica, 103(7), e300-e305. doi: 10.1111/apa.12617
- Nixon, A. F., Cunningham, S. J., Cohen, H. W., & Crain, E. F. (2012). The effect of Lactobacillus GG on acute diarrheal illness in the pediatric emergency department. *Pediatric Emergency Care*, 28(10), 1048-1051. doi: 10.1097/PEC.0b013e31826cad9f
- 47. Savas-Erdeve, S., Gokay, S., & Dallar, Y. (2009). Efficacy and safety of Saccharomyces boulardii in amebiasis-associated diarrhea in children. *Turkish Journal of Pediatrics*, 51(3), 220-224.
- 48. Szajewska, H., Skorka, A., Ruszczynski, M., & Gieruszczak-Bialek, D. (2013). Meta-analysis: Lactobacillus GG for treating acute gastroenteritis in children--updated analysis of randomised controlled trials. *Alimentary Pharmacology & Therapeutics, 38*(5), 467-476. doi: 10.1111/apt.12403
- Szajewska, H., Urbanska, M., Chmielewska, A., Weizman, Z., & Shamir, R. (2014). Meta-analysis: Lactobacillus reuteri strain DSM 17938 (and the original strain ATCC 55730) for treating acute gastroenteritis in children. *Beneficial Microbes*, 5(3), 285-293. doi: 10.3920/bm2013.0056
- Bailey, B., Gravel, J., Goldman, R. D., Friedman, J. N., & Parkin, P. C. (2010). External validation of the clinical dehydration scale for children with acute gastroenteritis. Academic Emergency Medicine, 17(6), 583-588. doi: 10.1111/j.1553-2712.2010.00767.x
- 51. Goldman, R. D., Friedman, J. N., & Parkin, P. C. (2008). Validation of the clinical dehydration scale for children with acute gastroenteritis. *Pediatrics*, 122(3), 545-549. doi: 10.1542/peds.2007-3141
- 52. Kinlin, L. M., & Freedman, S. B. (2012). Evaluation of a clinical dehydration scale in children requiring intravenous rehydration. *Pediatrics, 129*(5), e1211-1219. doi: 10.1542/peds.2011-2985
- 53. Madati, P. J., & Bachur, R. (2008). Development of an emergency department triage tool to predict acidosis among children with gastroenteritis. *Pediatric Emergency Care*, 24(12), 822-830. doi: 10.1097/PEC.0b013e31818ea004
- Schnadower, D., Tarr, P. I., Gorelick, M. H., O'Connell, K., Roskind, C. G., Powell, E. C., et al. (2013). Validation of the modified Vesikari score in children with gastroenteritis in 5 US emergency departments. *Journal of Pediatric Gastroenterology and Nutrition*, 57(4), 514-519. doi: 10.1097/MPG.0b013e31829ae5a3
- 55. Tam, R. K., Wong, H., Plint, A., Lepage, N., & Filler, G. (2014). Comparison of clinical and biochemical markers of dehydration with the clinical dehydration scale in children: A case comparison trial. *BMC Pediatrics, 14*, 149. doi: 10.1186/1471-2431-14-149
- 56. Levy, J. A., & Bachur, R. G. (2007). Intravenous dextrose during outpatient rehydration in pediatric gastroenteritis. *Academic Emergency Medicine*, 14(4), 324-330. doi: 10.1197/j.aem.2006.10.098
- 57. Freedman, S. B., Thull-Freedman, J. D., Rumantir, M., Atenafu, E. G., & Stephens, D. (2013). Emergency department revisits in children with gastroenteritis. *Journal of Pediatric Gastroenterology and Nutrition*, 57(5), 612-618. doi: 10.1097/MPG.0b013e3182a1dd93
- 58. FDA Drug Safety Communication: Abnormal heart rhythms may be associated with use of Zofran (ondansetron) 09-15-2011 www.FDA.gov
- 59. Albano, F., Bruzzese, E., Bella, A., Cascio, A., Titone, L., Arista, S., et al. (2007). Rotavirus and not age determines gastroenteritis severity in children: a hospital-based study. European Journal of Pediatrics, 166(3), 241-247. doi: 10.1007/s00431-006-0237-6
- Barletta, F., Ochoa, T. J., Mercado, E., Ruiz, J., Ecker, L., Lopez, G., et al. (2011). Quantitative real-time polymerase chain reaction for enteropathogenic Escherichia coli: A tool for investigation of asymptomatic versus symptomatic infections. *Clinical Infectious Diseases*, 53(12), 1223-1229. doi: 10.1093/cid/cir730
- 61. Mast, T. C., DeMuro-Mercon, C., Kelly, C. M., Floyd, L. E., & Walter, E. B. (2009). The impact of rotavirus gastroenteritis on the family. *BMC Pediatrics*, 9, 11. doi: 10.1186/1471-2431-9-11
- 62. Valentini, D., Vittucci, A. C., Grandin, A., Tozzi, A. E., Russo, C., Onori, M., et al. (2013). Coinfection in acute gastroenteritis predicts a more severe clinical course in children. *European Journal of Clinical Microbiology & Infectious Diseases, 32*(7), 909-915. doi: 10.1007/s10096-013-1825-9
- 63. Li, S. T., Grossman, D. C., & Cummings, P. (2007). Loperamide therapy for acute diarrhea in children: Systematic review and meta-analysis. *PLoS Medicine*, *4*(3), e98. doi: 10.1371/journal.pmed.0040098
- 64. Robertson, J., & Shilkofski, N. (Eds.). (2005). The Harriet Lane handbook: A manual for pediatric house officers (17th ed.). Philadelphia, PA: Mosby.
- 65. Aiello, A. E., Coulborn, R. M., Perez, V., & Larson, E. L. (2008). Effect of hand hygiene on infectious disease risk in the community setting: A metaanalysis. *American Journal of Public Health*, 98(8), 1372-1381.
- 66. Ejemot-Nwadiaro, R. I., Ehiri, J. E., Arikpo, D., Meremikwu, M. M., & Critchley, J. A. (2015). Hand washing promotion for preventing diarrhoea. *Cochrane Database of Systematic Reviews*, 9, CD004265. doi: 10.1002/14651858.CD004265.pub3
- 67. American Academy of Pediatrics. Committee on Infectious Diseases. Red Book : Report of the Committee on Infectious Diseases. Elk Grove Village, IL :American Academy of Pediatrics, 2021.

Guideline Preparation

This guideline was prepared by the Evidence-Based Outcomes Center (EBOC) team in collaboration with content experts at Texas Children's Hospital. Development of this guideline supports the TCH Quality and Patient Safety Program initiative to promote clinical guidelines and outcomes that build a culture of quality and safety within the organization.

Content Expert Team Members

Dana Cerminara, PharmD Sheebu Chacko, MD, Emergency Medicine Melissa Chladek, MD, Pediatric Hospital Medicine Kenneth Cohen, MD, Texas Children's Pediatric Associates Lindsay Day, MD, Emergency Medicine James Dunn, PhD, Medical Director, Pathology Ankhi Dutta, MD, Pediatric Hospital Medicine Ashley Joshi-Patel, MD, Pediatric Hospital Medicine Mindy Fein, MD, Emergency Medicine Michele Kuslich, LVN, TCPA Quality Buffy Orndorff, RN, Inpatient Nursing Julia Shelburne, MD, Pediatric Hospital Medicine Seema Walsh, MD, Gastroenterology Matthew Wilber, MD, Texas Children's Pediatric Associates Elizabeth Wuestner, RN, Clinical Nurse Specialist Darleen Yepes, RN, Infection Control

EBOC Team Members

Sheesha Porter, MSN, RN, Evidence-Based Practice Specialist Binita Patel, MD, Chief Medical Quality Officer

Additional EBOC Support

Anne Dykes, MSN, RN, ACNS-BC, Assistant Director Warren Boudreau, MSN, RN, Director

Development Process

This guideline was developed using the process outlined in the EBOC Manual (2021). The literature appraisal documents the following steps:

- 1. Review Preparation
 - PICO questions established
 - Evidence search confirmed with content experts
- 2. Review of Existing Internal and External Guidelines
- TCH Acute Gastroenteritis Clinical Guideline, Cincinnati Evidence-Based Care Guideline Prevention and Management of Acute Gastroenteritis, NICE Guideline on Diarrhoea and vomiting caused by gastroenteritis, and the European Society for Pediatric Gastroenterology, Hepatology, and Nutrition/European Society for Pediatric Infectious Diseases Evidence-Based Guideline for the Management of Acute Gastroenteritis in Children in Europe.
- 3. Literature Review of Relevant Evidence
 - Searched: PubMed, Cochrane Library, CINAHL, and Up to Date.
- 4. Critically Analyze the Evidence
 - 11 systematic reviews and meta-analyses, 8 randomized controlled trials, 29 non-randomized studies.
- 5. Summarize the Evidence
 - Materials used in the development of the guideline, evidence summary, and order sets are maintained in an Acute Gastroenteritis evidence-based review manual within EBOC.

Evaluating the Quality of the Evidence

Published clinical guidelines were evaluated for this review using the **AGREE II** criteria. The summary of these guidelines are included in the evidence summary. AGREE II criteria evaluate Guideline Scope and Purpose, Stakeholder Involvement, Rigor of Development, Clarity and Presentation, Applicability, and Editorial Independence using a 4-point Likert scale. The higher the score, the more comprehensive the guideline.

This guideline specifically summarizes the evidence *in support* of or *against* specific interventions and identifies where evidence is

lacking/inconclusive. The following categories describe how research findings provide support for treatment interventions.

"Evidence Supports" the guideline provides clear evidence from welldesigned randomized controlled trial(s) (RCT[s]) that the benefits of the intervention exceed harm.

"Evidence Against" provides clear evidence from more than one welldone RCT that the intervention is likely to be ineffective or that it is harmful.

"Evidence Lacking/Inconclusive" indicates there is currently insufficient data or inadequate data to support or refute a specific intervention.

The **GRADE** criteria were utilized to evaluate the body of evidence used to make clinical recommendations. The table below defines how the quality of the evidence is rated and how a strong versus weak recommendation is established. The evidence summary reflects the critical points of evidence.

Recommendation			
STRONG	Desirable effects clearly outweigh undesirable effects or vice versa		
WEAK	Desirable effects closely balanced with undesirable effects		
Quality	Type of Evidence		
High	Consistent evidence from well-performed RCTs or exceptionally strong evidence from unbiased observational studies		
Moderate	Evidence from RCTs with important limitations (e.g., inconsistent results, methodological flaws, indirect evidence, or imprecise results) or unusually strong evidence from unbiased observational studies		
Low	Evidence for at least 1 critical outcome from observational studies, RCTs with serious flaws or indirect evidence		
Very Low	Evidence for at least 1 critical outcome from unsystematic clinical observations or very indirect evidence		

Recommendations

Practice recommendations were directed by the existing evidence and consensus amongst the content experts. Patient and family preferences were included when possible. The Content Expert Team and EBOC team remain aware of the controversies in the management of acute gastroenteritis in children. When evidence is lacking, options in care are provided in the clinical standard and the accompanying order sets (if applicable).

Approval Process

Clinical standards are reviewed and approved by hospital committees as deemed appropriate for its intended use. Clinical standards are reviewed as necessary within EBOC at Texas Children's Hospital. Content Expert Teams are involved with every review and update.

Disclaimer

Practice recommendations are based upon the evidence available at the time the clinical standard was developed. Clinical standards (guidelines, summaries, or pathways) <u>do not</u> set out the standard of care and are not intended to be used to dictate a course of care. Each physician/practitioner must use his or her independent judgment in the management of any specific patient and is responsible, in consultation with the patient and/or the patient's family, to make the ultimate judgment regarding care.

Version History				
Date	Comments			
Feb 2009	Original guideline completed			
Mar 2016	Updated			
Jun 2017	Added links to Heart Failure Guideline			
Aug 2019	Algorithm updated with link to EC stool testing guideline			
July 2021	ORT revised to 1ml/kg q 5 mins for guideline & algorithm; note added for cautious use of probiotics in immunocompromised patients.			