Clinical Pathways

Clinical Pathway: Anaphylaxis

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What is a Clinical Pathway?



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

Objectives of Pathway



- To standardize the approach for patients with anaphylaxis
- To improve the recognition of anaphylaxis and early administration of intramuscular epinephrine
- To improve the safety of patients who develop anaphylaxis
- To improve documentation in patient chart of allergic reactions, including details of the specific reaction and severity of reaction

Why is Pathway Necessary?



- Anaphylaxis is a life threatening condition that requires rapid assessment and treatment
- A clinical pathway can empower early administration of epinephrine when anaphylaxis is suspected
- It can also ensure adherence to the American Academy of Allergy, Asthma and Immunology guideline for management of anaphylaxis





- Anaphylaxis is a multisystem emergency and can progress to a life threatening condition
- Anaphylaxis requires prompt recognition and treatment
- Delayed treatment with epinephrine is associated with increased risk for fatality
- Leading causes of anaphylaxis are medications, foods, Hymenoptera (i.e. bees, wasps, hornets) stings

Background: Signs and Symptoms Anaphylaxis¹

Table II-1

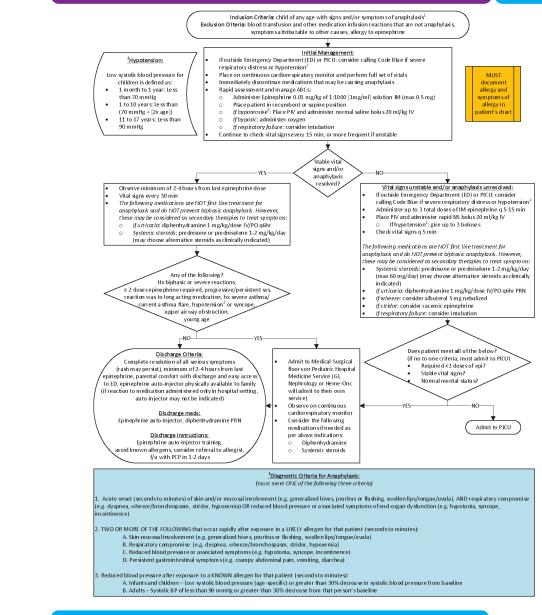
Signs and symptoms of anaphylaxis

System	Symptoms	System	Symptoms	Signs and symptoms of anaphylaxis ^a		
Skip	Flushing local or generalized	Cardiovascular	Chest pain eg substemal tachusardia	Signs and symptoms	Percentage ^b	
Skin Respiratory	Flushing, local or generalized Flushing, local or generalized Localized itching of skin or mucosa (local areas, eg, palms, genitalia, and/or palate) or generalized itching Urticaria Angioedema of skin or mucosa (eg, lips or tongue) morbilliform rash pilar erection Conjunctival itching, redness, tearing, and/or swelling Nasal itching, congestion, rhinorrhea, sneezing Throat itching and tightness Dysphonia, hoarseness, stridor Coughing Increased respiratory rate Shortness of breath Wheezing Chest tightness Cyanosis Respiratory arrest	Cardiovascular Central nervous system Other	Chest pain, eg, substernal, tachycardia, bradycardia, palpitations, arrhythmias, hypotension, feeling faint, urinary or feca incontinence, shock, cardiac arrest Aura of impending doom Uneasiness Sudden behavioral change (eg, irritability) Dizziness Headache (eg, throbbing) Altered mental state Tunnel vision Confusion Seizure Metallic taste in mouth Uterine cramping and/or bleeding	Cutaneous	62-90 45-55 2-5 45-50 50-60 15-20 30-35 25-30 5-8 4-5 1-2 an P, Nicklas R, Oppen- cis practice parameter:	
Gastrointestinal	Abdominal pain (eg, cramping) Nausea Vomiting Diarrhea Dysphagia			Lieberman P, et al. Anaphylaxis in America: results from a national physician surv Ann Allergy Asthma Immunol. 2012;109 (suppl):A20; and Boyle J, Camargo CA, I berman P, et al. Anaphylaxis in America: results from a national telephone surve Allergy Clin Immunol. 2012;129 (suppl):AB132. ^b Percentages are approximations.		

Table I-1

Connecticut

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This is the Anaphylaxis Clinical Pathway.

We will be reviewing each component in the following slides.

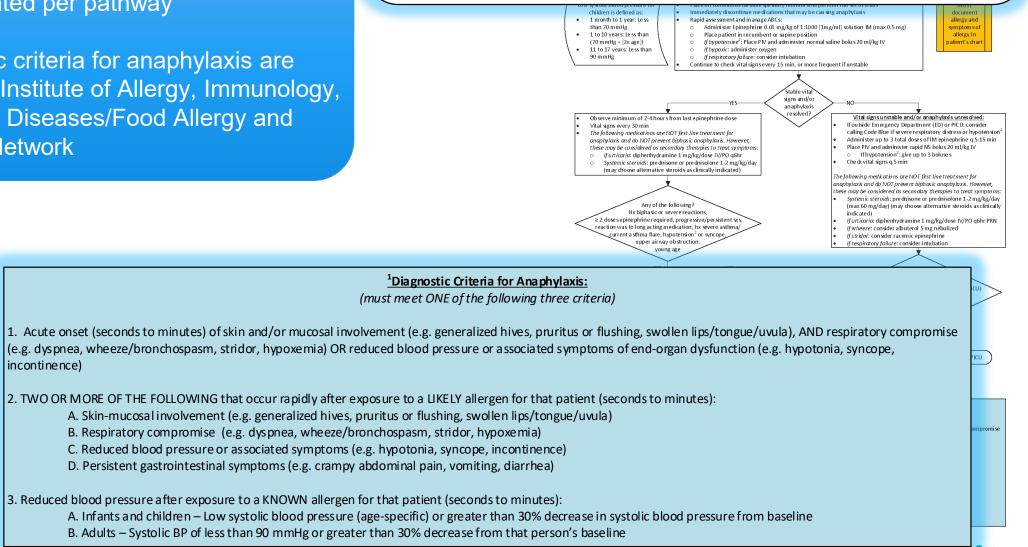
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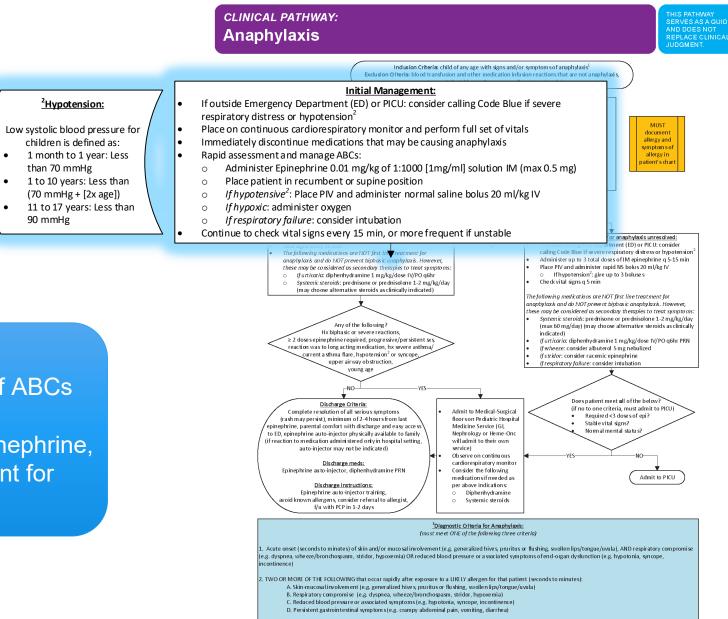
CLINICAL PATHWAY: Anaphylaxis

- Inclusion criteria: Anyone with anaphylaxis should be treated per pathway
- The diagnostic criteria for anaphylaxis are from National Institute of Allergy, Immunology, and Infectious Diseases/Food Allergy and Anaphylaxis Network

Inclusion Criteria: child of any age with signs and/or symptoms of anaphylaxis¹ **Exclusion Criteria:** blood transfusion and other medication infusion reactions that are not anaphylaxis, symptoms attributable to other causes, allergy to epinephrine



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Reduced blood pressure a fter exposure to a KNOWN allergen for that patient (second sto minutes): A. Infants and children - Low systolic blood pressure (age-specific) or greater than 30% decrease in systolic blood pressure from baseline B. Adults - Systolic BP of less than 90 mmHg or greater than 30% decrease from that person's baseline

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Initial Management:

 Rapid assessment and management of ABCs are key

than 70 mmHg

90 mmHg

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Do NOT delay administration of IM epinephrine, as this is the definitive first line treatment for anaphylaxis

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

It is VERY important to document any allergic reaction in the patient chart.

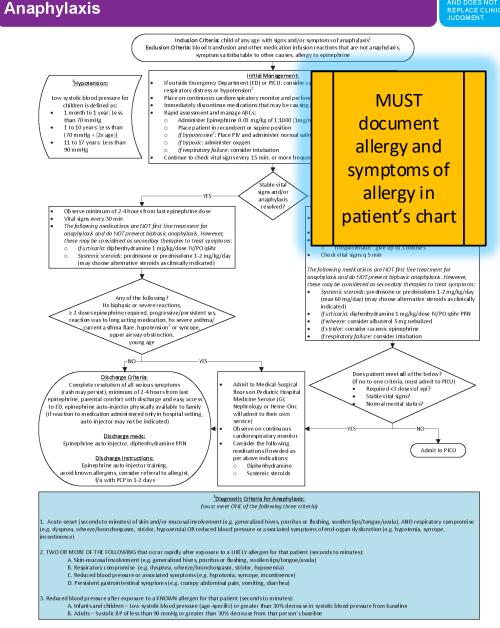
Document allergen AND allergic reaction associated with the exposure

Table I-2

Essential features of history in the evaluation of a patient who has experienced an episode of anaphylaxis

- A Detailed history of ingestants (foods/drugs) taken within 6 h before the event
- B Activity in which the patient was engaged at the time of the event
- C Location of the event (home, school, work, indoors/outdoors)
- D Exposure to heat or cold
- E Any related sting or bite
- F Time of day or night
- G Duration of event
- H Recurrence of symptoms after initial resolution
- Exact nature of symptoms (eg, if cutaneous, determine whether flush, pruritus, urticaria, or angioedema)
- In a woman, the relation between the event and her menstrual cycle
- K Was medical care given and what treatments were administered
- L How long before recovery occurred and was there a recurrence of symptoms after a symptom-free period

From Liberman P, Nicklas RA, Randolph C, et al. Anaphylaxis - a practice parameter update 2015. *Ann Allergy Asthma Immunol.* 2015 Nov;115(5):341-84.

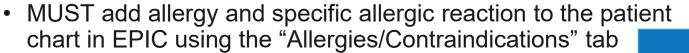


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CLINICAL PATHWAY:

Document detailed history of allergic reaction



- For an allergic reaction that • occurs to a medication given while at Connecticut Children's:
 - Document under \cap allergies as above
 - Consider documenting Ο by using SmartPhrase ".MEDREACTION" in event note to document details of reaction

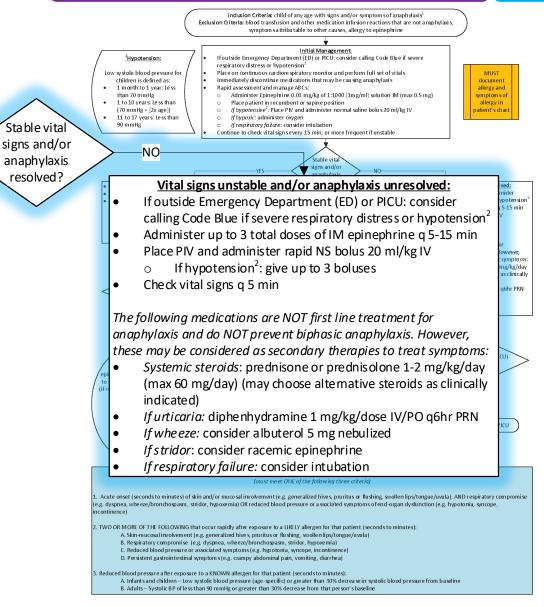
		Add a new agent		Full Search	R view Drug-Allerg	y Interactions
ontraindications"	tab		umns		Show: Deleted	Expired ;
			Reaction	Severity	Updated	
		Allergies				
		Peanut				
My Note	Se <u>n</u> sitive	Agent:	Peanut			
Type: Event Note 🔎 Service:	Pediatric Hospit 🔎	Reactions:	Anaphylaxis	Severity:	High 🔎	
	02 (]			O Noted:	1/12/2022	
Cosign Required						
Summary:		Reaction Type:	Food Allergy	0		
🛠 🖪 🗩 🦥 ᅿ 🎣 🕇 Insert Sr	martText 📑 😓 🔿 🛼 📿 🔳	Comments:				
Medication Reaction Event N	ote	€ 🕫 €	😋 🕄 ∔ 🛛 Insert Sm	artText 🖷 🗲 🕇	- 🍫 🛼 🛛 100% 👻 🖍	
Medication: ***		Stridor, hypo	tension, vomiting, hives			
Dose: *** Date of Reaction: ***						
Time of Reaction: ***						
Signs and symptoms: ***						
CT-CAE Grade: ***					✓ Accept	t X Cancel
Anaphylaxis (Y/N): ***						
Interventions: ***wa Infusion resumed (Y/N): ***		Amoxicillin	Nausea And V	omiting Low	Past Up	odates
Infusion start time: ***		Noted: 6/4/2	2016			
First dose (Y/N): ***						

Allergies/Contraindications



Management Continued (UNSTABLE PATIENT OR ANAPHYLAXIS UNRESOLVED):

- Must determine if symptoms/signs of anaphylaxis resolve and patient is stable or if requires repeated doses of IM
- If unstable at any point AND outside the Emergency Department, consider calling Code Blue if severe respiratory distress or hypotension while addressing these issues
- See next slide to learn about secondary therapies



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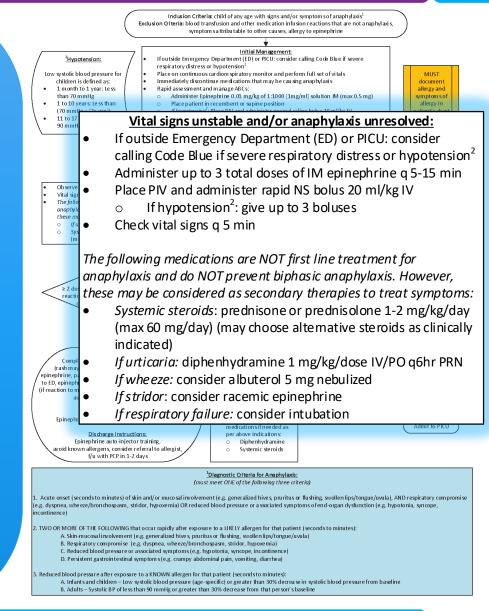
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CLINICAL PATHWAY: Anaphylaxis

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Adjunctive or Second Line Therapies ¹⁻⁶:

- They do NOT prevent or treat upper airway obstruction or hypotension! IM epinephrine is the first line treatment
- Antihistamines, H1 and H2 blockers, are second line, because there is no evidence to support their use in first line management of anaphylaxis – may be used symptomatically to treat pruritus/hives. These do NOT prevent biphasic or prolonged reactions
- Literature review demonstrates that systemic corticosteroids do NOT prevent biphasic or prolonged reactions. Again, may be used for symptom control/comfort in dose of 1-2 mg/kg/day
- Patients with complete resolution of symptoms after treatment with epinephrine do NOT require prescription for antihistamines or corticosteroids

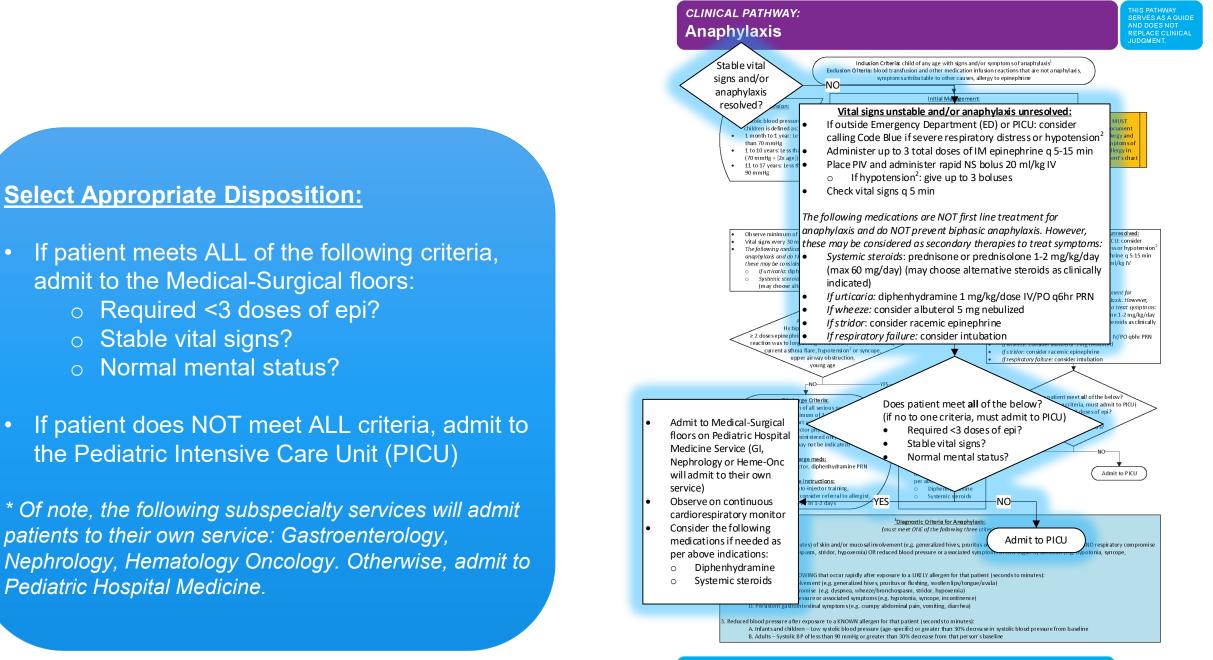


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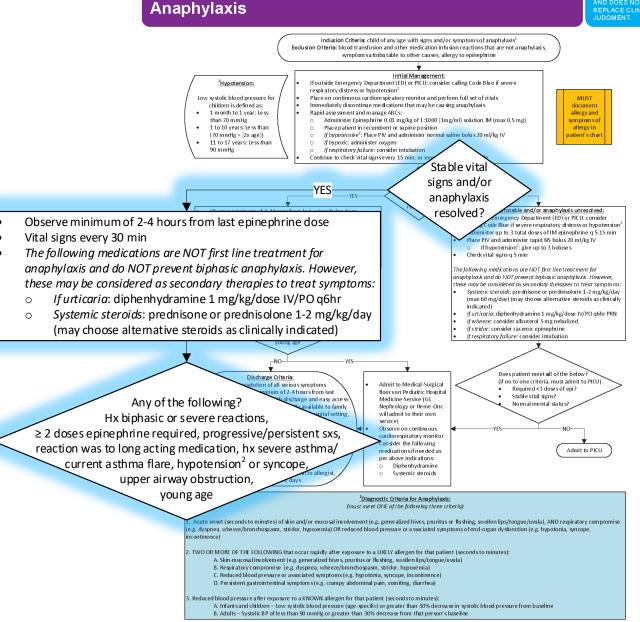
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Management Continued (STABLE PATIENT):

- If patient has stable vital signs and/or anaphylaxis resolved, observe for 2-4 hours from last epinephrine dose
- If NO risk factors, may discharge home

• Risk factors:

- History biphasic or severe reactions
- $\circ \geq 2$ doses epinephrine required
- Progressive/persistent symptoms
- Reaction was to long acting medication
- History of severe asthma/current asthma flare
- Hypotension or syncope
- Upper airway obstruction
- Young age



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CLINICAL PATHWAY:

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Discharge Criteria:

- Complete resolution of all serious symptoms
- (rash may persist)
- Minimum of 2-4 hours from last epinephrine
- Parental comfort with discharge and easy access to ED
- Epinephrine auto-injector physically available to family (if reaction to medication administered only in hospital setting, auto-injector may not be indicated)

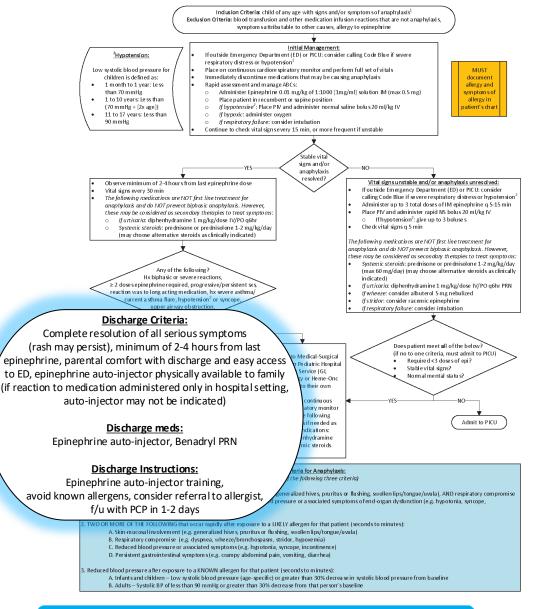
Discharge meds:

- Epinephrine auto-injector (consider dispensing at least 2)
- Diphenhydramine PRN pruritic rash

Discharge Instructions:

- Epinephrine auto-injector training
- Avoid known allergens
- Consider referral to allergist,
- F/u with PCP in 1-2 days





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Review of Key Points



- This clinical pathway serves to standardize management of anaphylaxis across the institution (ED, inpatient, clinics, etc.) and across different triggers (medications, food, environmental, etc.)
- Anaphylaxis requires rapid assessment and treatment
- The primary treatment for anaphylaxis is rapid administration of IM epinephrine
- Secondary therapies may be used for symptomatic management, but they do NOT prevent or treat upper airway obstruction or hypotension and they do NOT prevent biphasic or prolonged reactions
- Documentation of allergic reactions, including details of the specific reaction and severity should occur in a standardized manner

Quality Metrics



- Percentage of patients with utilization of pathway order set
- Time from reaction (if in a clinic or hospitalized) or time from arrival to ED to administration of epinephrine (if not already given prior to arrival)
- Percentage of patients on pathway with documentation of allergy AND reaction in chart
- Number of patients discharged from the ED who return within 72 hours and 7 days

Pathway Contacts







- 1. Campbell RL, Li J, Nicklas RA, Sadosty AT. Emergency department diagnosis and treatment of anaphylaxis: a practice parameter. *Ann Allergy Asthma Immunol*. 2014 Dec;113(6):599-608.
- 2. Choo KJ, Simons E, Sheikh A. Glucocorticoids for the treatment of anaphylaxis: Cochrane systematic review. *Allergy*. 2010 Oct;65(10):1205-11.
- 3. Lieberman P. Biphasic anaphylactic reactions. *Ann Allergy Asthma Immunol*. 2005 Sep;95(3):217-26.
- 4. Liberman P, Nicklas RA, Randolph C, et al. Anaphylaxis a practice parameter update 2015. *Ann Allergy Asthma Immunol.* 2015 Nov;115(5):341-84.
- 5. Shaker MS, et al. Anaphylaxis a 2020 practice parameter update, systematic review, and Grading of Recommendations, Assessment, Development and Evaluation (GRADE) analysis. *J Allergy Clin Immunol*. 2020 April;145(4):1082-1123.
- 6. Sheikh A, Ten Broeck V, Brown SG, Simons FE. H1-antihistamines for the treatment of anaphylaxis: Cochrane systematic review. *Allergy*. 2007 Aug;62(2):830-7.

Thank You!



About Connecticut Children's Clinical Pathways Program

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