Medication	<5 Years of Age	5–11 Years of Age	≥12 Years of Age and Adults	Potential Adverse Effects	Comments (not all inclusive)
Inhaled Short-Actir	ng Beta ₂ -Agonists				
	Dose applies to Albuterol.	Dose applies to Albuterol/and Levalbuterol.	Dose applies to all four SABAs		Apply to all four (SABAs)
MDI					
Albuterol CFC	1–2 puffs	2 puffs	2 puffs	Tachycardia, skeletal muscle	Drugs of choice for acute bronchospasm.
90 mcg/puff, 200 puffs/canister	5 minutes before exercise	5 minutes before exercise	5 minutes before exercise	tremor, hypokalemia, increased lactic acid, headache, hyperglycemia.	 Differences in potencies exist, but all products are essentially comparable on a puff per puff basis.
Albuterol HFA	2 puffs every	2 puffs every	2 puffs every	Inhaled route, in general,	An increasing use or lack of expected effect indicates diminished control of asthma.
90 mcg/puff, 200 puffs/canister	4–6 hours, as needed for symptoms	4–6 hours, as needed for symptoms	4–6 hours, as needed for symptoms	causes few systemic adverse effects. Patients with preexisting cardiovas- cular disease, especially the	Not recommended for long-term daily treat- ment. Regular use exceeding 2 days/week for symptom control (not prevention of EIB)
Levalbuterol HFA	NA <4 years of			elderly, may have adverse cardiovascular reactions	indicates the need for additional long-term control therapy.
45 mcg/puff, 200 puffs/canister	age			with inhaled therapy.	 May double usual dose for mild exacerbation: For levalbuterol, prime the inhaler by releasin 4 actuations prior to use.
Pirbuterol CFC Autohaler	NA	NA			For HFA: periodically clean HFA actuator, as drug may plug orifice.
200 mcg/puff, 400 puffs/canister					 For autohaler: children <4 years of age may not generate sufficient inspiratory flow to activate an auto-inhaler. Nonselective agents (i.e., epinephrine, isoproterenol, metaproterenol) are not recommended due to their potential for excessive cardiac stimulation, especially in high doses.
Nebulizer solution					May mix with cromolyn solution, budesonide interest and a second control of the least account of the leas
Albuterol					inhalant suspension, or ipratropium solution for nebulization. May double dose for severe
0.63 mg/3 mL 1.25 mg/3 mL 2.5 mg/3 mL 5 mg/mL (0.5%)	0.63–2.5 mg in 3 cc of saline q 4–6 hours, as needed	1.25–5 mg in 3 cc of saline q 4–8 hours, as needed	1.25–5 mg in 3 cc of saline q 4–8 hours, as needed	(Same as with MDI)	exacerbations. Does not have FDA-approved labeling for children <6 years of age.
Levalbuterol (R-albuterol)					Compatible with budesonide inhalant suspension. The product is a sterile-filled preservative-free unit dose vial.
0.31 mg/3 mL 0.63 mg/3 mL 1.25 mg/0.5 mL 1.25 mg/3 mL	0.31–1.25 mg in 3 cc q 4–6 hours, as needed for symp- toms	0.31–0.63 mg, q 8 hours, as needed for symptoms	0.63 mg– 1.25 mg q 8 hours, as needed for symptoms	(Same as with MDI)	

Key: CFC, chlorofluorocarbon; ED, emergency department; EIB, exercise-induced bronchospasm; HFA, hydrofluoroalkane; IM, intramuscular; MDI, metered-dose inhaler; NA, not available (either not approved, no data available, or safety and efficacy not established for this age group); PEF, peak expiratory flor; SABA, short-acting beta₂-agonist

^{*}Dosages are provided for those products that have been approved by the U.S. Food and Drug Administration (FDA) or have sufficient clinical trial safety and efficacy data in the appropriate age ranges to support their use.

Medication	<5 Years of Age	5-11 Years of Age	≥12 Years of Age and Adults	Potential Adverse Effects	Comments (not all inclusive)
Anticholinergics	, ,				
Ipratropium HFA					
MDI					
17 mcg/puff, 200 puffs/canister	NA	NA	2–3 puffs q 6 hours	 Drying of mouth and respiratory secretions, 	Multiple doses in the emergency department (not hospital) setting provide additive benefit
Nebulizer solution				increased wheezing in some individuals, blurred	to SABA. Treatment of choice for bronchospasm due
0.25 mg/mL (0.025%)	NA	NA	0.25 mg q 6 hours	vision if sprayed in eyes. If used in the ED, produces	to beta-blocker medication. Does not block EIB.
Ipratropium with albuterol				less cardiac stimulation than SABAs.	 Reverses only cholinergically mediated bronchospasm; does not modify reaction to antigen.
MDI					May be an alternative for patients who
18 mcg/puff of ipratropium bromide and 90 mcg/puff of albuterol	NA	NA	2–3 puffs q 6 hours		do not tolerate SABA. Has not proven to be efficacious as long-term control therapy for asthma.
200 puffs/canister					
Nebulizer solution					
0.5 mg/3 mL ipratropium bromide and 2.5 mg/3 mL albuterol	NA	NA	3 mL q 4–6 hours		Contains EDTA to prevent discoloration of the solution. This additive does not induce bronchospasm.
Systemic Corticos	teroids				
	Dosages apply t	o first three corti	osteroids.		(Applies to the first three corticosteroids.)
Methylprednisolone 2, 4, 6, 8, 16, 32 mg tablets Prednisolone 5 mg tablets, 5 mg/5 cc, 15 mg/5 cc Prednisone 1, 2.5, 5, 10, 20, 50 mg tablets; 5 mg/cc, 5 mg/5 cc	Short course "burst:" 1–2 mg/kg/ day, maximum 60 mg/day, for 3–10 days	Short course "burst": 40–60 mg/day as single or 2 divided doses for 3–10 days	Short course "burst": 40–60 mg/day as single or 2 divided doses for 3–10 days	 Short-term use: reversible abnormalities in glucose metabolism, increased appetite, fluid retention, weight gain, facial flushing, mood alteration, hypertension, peptic ulcer, and rarely aseptic necrosis. Consideration should be given to coexisting conditions that could be worsened by systemic corticosteroids, such as herpes virus infections, varicella, tuberculosis, hypertension, peptic ulcer, diabetes mellitus, osteoporosis, and <i>Strongyloides</i>. 	 Short courses or "bursts" are effective for establishing control when initiating therapy or during a period of gradual deterioration. Action may begin within an hour. The burst should be continued until patient achieves 80 percent PEF personal best or symptoms resolve. This usually requires 3–10 days but may require longer. There is no evidence that tapering the dose following improvement prevents relapse in asthma exacerbations. Other systemic corticosteroids such as hydrocortisone and dexamethasone given in equipotent daily doses are likely to be as effective as prednisolone.

FIGURE 19. USUAL DOSAGES FOR QUICK-RELIEF MEDICATIONS* (continued)							
Medication	<5 Years of Age	5–11 Years of Age	≥12 Years of Age and Adults	Potential Adverse Effects	Comments (not all inclusive)		
Systemic Corticosteroids (continued)							
Repository injection (Methylprednisolone acetate) 40 mg/mL 80 mg/mL	7.5 mg/kg IM once	240 mg IM once	240 mg IM once		May be used in place of a short burst of oral steroids in patients who are vomiting or if adherence is a problem.		