Use of Codeine- and Dextromethorphan-Containing Cough Remedies in Children

ABSTRACT. Numerous prescription and nonprescription medications are currently available for suppression of cough, a common symptom in children. Because adverse effects and overdosage associated with the administration of cough and cold preparations in children have been reported, education of patients and parents about the lack of proven antitussive effects and the potential risks of these products is needed.

INDICATIONS AND CONTRAINDICATIONS
Cough is a reflex response to mechanical, chemical, or inflammatory irritation of the tracheobronchial tree mediated by sensory neurons in the airways reflexly through neurons in the brainstem. Cough serves as a physiologic function to clear airways of obstructive or irritating material or to warn of noxious substances in inspired air.

In some pathologic states (eg, asthma, bronchopulmonary dysplasia, cystic fibrosis, and a variety of inflammatory conditions), excessive and/or abnormal airway secretions may be produced. The cough reflex serves to maintain airway patency by clearing these secretions. Clearing of pathologic tracheobronchial secretions is essential to patient management and may be enhanced by chest physiotherapy. Cough suppression may adversely affect patients with these conditions by promoting pooling of secretions, airway obstruction, secondary infection, and hypoxemia.

Many common respiratory conditions in which cough is prominent (eg, respiratory viral infections) are self-limited (lasting a few days). Cough may be an expression of airway reactivity or asthma. The cough that is associated with these conditions may be satisfactorily managed with fluids and increased ambient humidity (especially of value with croup). When cough is persistent, it is usually secondary to infection, allergy (including asthma), environmental irritants (eg, cigarette smoke, dust particles) or, occasionally, a foreign body. Therapy should be directed at the underlying condition for lasting benefit.

ANTITUSSIVE AGENTS
Most cough suppressant preparations are marketed as mixtures of dextromethorphan or codeine with antihistamines, decongestants, expectorants, and/or antipyretics. Some nonprescription preparations substitute diphenhydramine or eucalyptus oil in place of codeine or dextromethorphan. Prescription medications may substitute other narcotic agents (hydrocodone or hydromorphone) for codeine and may be more addictive than codeine. In addition, many of these cough products are elixirs, which may contain up to 25% alcohol by volume.

The over-the-counter availability of numerous cough and cold preparations promotes the perception that such medications are safe and efficacious. Although codeine and dextromethorphan are efficacious for cough suppression in adults, similar efficacy has not been demonstrated in children. Taylor et al conducted a randomized, controlled trial of codeine, dextromethorphan, and placebo in children with acute nocturnal cough without evidence of chronic underlying lung disease (asthma, cystic fibrosis, or bronchopulmonary dysplasia). Neither dextromethorphan nor codeine in the dosages used was significantly more effective than placebo in reduction of acute cough. Studies using larger dosages have not been performed. Other studies focusing exclusively on children with cough have not been placebo-controlled trials. To our knowledge, studies of the use of other purportedly antitussive agents in children, such as diphenhydramine, have not been reported in the literature.

Demonstration of the efficacy of antitussive preparations in children is lacking, and these medications may be potentially harmful. Decongestant (sympathomimetic) components of these mixtures administered to children have been associated with irritability, restlessness, hallucination, hypertension, and dystonic reactions. The clearance and metabolism of the components of cough mixtures may vary with age and disease state. Great variability in the enterohepatic circulation of these drugs is noted in adults, which affects drug response, especially with repeated dosing. The relative immaturity of hepatic enzyme systems that metabolize drugs in young children may enhance the risk of adverse effects of such medications, especially in infants younger than 6 months. Metabolism and/or toxicity also may be altered by concurrent use of medications such as acetaminophen. Unfortunately, the dosing guidelines for these agents are based on extrapolation from adult data without consideration of their potentially unique metabolism and disposition in children.

Codeine
In adults, codeine and dextromethorphan have been shown to suppress both artificially induced and
disease-related cough, mainly through central nervous system mechanisms. A linear relationship has been shown to exist between a codeine dosage in the range of 7.5 to 60 mg/d and a decrease in the frequency of chronic cough. Complete suppression of cough was not achieved in these trials, even at the highest daily dose of codeine.

Dosage
Pharmacokinetic studies of codeine therapy in children are lacking. The published dosage recommendation for codeine in children is 1 mg/kg/d in four divided doses, not to exceed 60 mg/d. To our knowledge, no well-controlled studies have documented the safety and efficacy of this dosage.

Adverse Reactions and Overdosage
The principal clinical manifestations of codeine toxicity are respiratory depression and obtundation. In children, antitussive dosages of 3 to 5 mg/kg/d have produced somnolence, ataxia, miosis, vomiting, rash, facial swelling, and pruritus. Respiratory depression requiring mechanical ventilation occurred in 3% of children receiving dosages greater than 5 mg/kg/d; two of these patients died. Doses of codeine less than 2 mg/kg are unlikely to be associated with significant adverse reactions. Reports of adverse reactions to codeine are based on single dose ingestions; the repetitive administration of codeine for therapeutic purposes may be associated with adverse symptoms at doses lower than a single dose of 5 mg/kg. In adults, glucuronide conjugation in the liver apparently inactivates codeine, but 10% of an oral dose is demethylated to form morphine, which is believed by some to be the active form of the drug. The hepatic glucuronidation pathway is incompletely developed in infants, which places them at particular risk for adverse dose-related effects. Furthermore, alteration of hepatic enzyme pathways by illness or concurrent drug therapy (such as acetaminophen) may further alter metabolism of this drug and increase the risk of drug toxicity.

Other narcotic antitussives that are available in cough preparations, such as hydrocodone and hydromorphone, have no demonstrated advantage as antitussive agents compared with codeine, have similar adverse effects, and have a greater risk of dependency. Hydrocodone and hydromorphone are classified as Schedule III drugs under the Controlled Substances Act.

Dextromethorphan
The addictive potential of codeine encouraged the marketing of dextromethorphan in a variety of cough and cold preparations. Although dextromethorphan is chemically derived from the opiates, it has no analgesic or addictive properties. The cough suppression potency of dextromethorphan in adults is nearly equal to that of codeine. The drug, like codeine, acts on the central nervous system to elevate the threshold for coughing.
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