Therapeutic effects of Ventolin versus hypertonic saline 3% for acute bronchiolitis in children

Mohammad-Ali Zamani1, Mehran Movahhedi2, Seyyed Mohammad-Kazem Nourbaksh3 Forouzan Ganji4, Mahmoud Rafieian-Kopaei5, Mahmoud Mobasher6, Abolfazl Khoshdel7 Shahram Etemadifar8, Majid Shirani9, Zahra Keivan Hafshejani*10

Received: 16 June 2014 Accepted: 6 January 2015 Published: 6 May 2015

Abstract

Background: Complications of Ventolin as the most common drug used for bronchiolitis are widely known. The present study was conducted to determine the efficacy of hypertonic saline 3%, compared with Ventolin, for treatment of acute bronchiolitis in children.

Methods: This double-blinded clinical trial study was conducted in Hajar Hospital, Shahrekord, Iran, from 2011 to 2012. A total of 70 patients under the age of two years with bronchiolitis were divided into two groups of 35 each. Ventolin nebulizer and hypertonic saline 3% nebulizer three times per day were administered in the first (Ventolin) and second (Hypersaline) group, respectively. The length of recovery was compared between the two groups. The data were analyzed by SPSS software (version 22) using chi-square, t-test, paired t-test, and Mann-Whitney.

Results: The mean±SD length of recovery was 4.14±0.9 and 3.06±0.6 in the Ventolin and hypersaline groups, respectively. The mean duration of recovery was significantly lower in the hypersaline group (p<0.001).

Conclusion: Hypertonic saline 3% nebulizer has more pleasant therapeutic effects on acute bronchiolitis than Ventolin. Therefore, use of hypertonic saline 3% nebulizer is recommended for the treatment of acute bronchiolitis in children under two years old.

Keywords: Hypertonic saline solution, Pediatric, Bronchiolitis, Ventolin-albuterol.


Introduction

Acute bronchiolitis is the most frequent reason for lower respiratory tract infection and hospitalization due to respiratory disease in infancy. The frequent age of the disease incidence is the first two years of life. About 50000-80000 hospitalizations of the infants under the age of one are attributed to this disease in the USA (1). The severe form of this disease is much more prevalent in one- to three-month-old infants and responsible for more than 50% of its incidence, causing respiratory syncytial virus, inflammation and small airway obstruction (1). Clinical symptoms of acute bronchiolitis are similar to those of viral pneumonias, with fever, wheezing, and increased respiratory rate as the most im-
Ventolin versus hypertonic saline 3% in acute bronchiolitis

Research is needed to find the appropriate treatment for this disease. The treatment is often supportive therapy, including fluid therapy, anti-fever drugs, and oxygen (3). Some investigations have been recently conducted on the therapies like bronchodilators including salbutamol and epinephrine as nebulized (2,4). In infants particularly under age of six months, edema and inflammation in bronchioles could lead to respiratory distress because of small diameter of airways (1).

There is no consensus on use of bronchodilators and/or beta-agonists (5,6). Most works have indicated that bronchodilators have no role in treating these patients (7). Today, inhaled salbutamol (Ventolin) is used for treating bronchiolitis patients in Iran’s hospitals while several studies have indicated that this drug does not work (8-12). In many studies, the positive effect of hypertonic saline has been observed on recovering bronchiolitis symptoms (9,10). Also, some studies comparing different concentrations of hypertonic saline have obtained similar results. For example, comparative study of hypertonic saline 3% and 7% indicated a greater therapeutic effect of hypertonic saline 3% than that of hypertonic saline 7% in treating bronchiolitis symptoms and decreasing duration of treatment (13,14). Hypertonic saline 3% was also found as better and more efficient than normal saline and hypersaline 3% nebulizer (15,16).

In addition, a research conducted to compare hypertonic saline 3% with the drugs used for bronchiolitis like Ventolin indicated that hypertonic saline 3% and Ventolin had a greater therapeutic effect than Ventolin and normal saline on treating bronchiolitis symptoms (17). In some studies, Ventolin and hypertonic saline 3% had similar therapeutic effect on symptoms developed by mild to moderate viral bronchiolitis and were free of complications, as well (18). Regarding the widely known complications of Ventolin as the most common drug used for bronchiolitis and that no study has been yet conducted to compare the efficacy of Ventolin and hypertonic saline 3% on bronchiolitis in Iran, the present study was conducted to determine the efficacy of hypertonic saline 3% on treatment of acute bronchiolitis, in comparison with Ventolin.

Methods

This research is a double-blinded clinical trial study conducted in 2011-2012 on 70 referring patients with acute viral bronchiolitis. Registration code of IRCT 2012110510222N2 was issued by Iranian Registry of Clinical Trials for this study. The samples of study were examined for severity of simulated disease prior to enrollment and the patients with acute status and causing a predetermined error in the study were excluded from the investigation. Inclusion criteria were patients with mild to moderate respiratory status, the first wheezing, and obtaining a score of lower than nine in Respiratory Distress Assessment Inventory (RDAI), the most common clinical scoring tool for acute bronchiolitis developed based on wheezing and intercostal retraction (19). All patients were frequently visited and examined by a pediatrician in Hajar Hospital, Shahrekord, Iran till completion of their hospitalization. After the informed consent was obtained from the patients’ guardians, the patients were assigned into two groups of case and control. Group one underwent treatment with Ventolin nebulizer and group two with hypertonic saline 3% nebulizer. The prescribed dose of Ventolin was 0.1 mg/kg and the required concentration was obtained by distilled water. To obtain hypertonic saline 3% solution, 3 ml sodium chloride 5% was mixed with a 5-cc vial of distilled water. The two medicinal combinations were administered to the patients of the two groups every four hours by a nebulizer instrument. The inhaled medicine accompanied with oxygen was nebulized to the patients by a face mask. Other therapies including oxygen and fluid therapy were used for all patients. Clinical symptoms of the patients and the length of recovery since hospitalization were free of complications, as well (18).
zation initiation were calculated for each patient and registered in a special checklist based on the defined RDAI criterion and day. The data were analyzed by SPSS software (version 22) using chi-square, t-test, paired t-test, and Mann-Whitney.

Results
In this study, 70 patients were randomly assigned to two groups of 35 each. Group one underwent treatment with Ventolin nebulizer (Ventolin group) and group two with hypertonic saline 3% nebulizer (Hypersaline group). The mean±SD age of the patients was 14.1±5.6 months in the Ventolin group and 12.6±5.6 months in the hypersaline group.

The mean±SD length of recovery was 4.14±0.9 days in the Ventolin group and 3.06±0.6 in the hypersaline group. The mean length of recovery was significantly lower in the hypersaline group (p<0.001). The mean±SD RDAI criterion on the days two, three, four, and five was respectively 5.68±1.3, 4.85±1.6, 3.62±1.6, and 1.42±0.8 in the Ventolin group and 4.25±1.5, 3.2±1.5, 2.54±1.6, and 0.9±0.54 in the hypersaline group; the mean RDAI criterion was significantly lower in the hypersaline 3% group (p<0.001).

Discussion
The general purpose of conducting this study was to compare the efficacy of hypertonic saline 3% with that of Ventolin on treating acute bronchiolitis in the children under two years. In this study, two groups (35 patients in each group) of infants hospitalized for bronchiolitis in Hajar Hospital, Shahrekord, Iran were investigated. According to the results of this study, the mean length of recovery was 4.14±0.4 days in the Ventolin group and 3.0±0.6 in the hypersaline group. The patients under hypersaline treatment recovered more rapidly, which is clinically and economically important as 50000-80000 hospitalizations of under one-year-old infants has led to expending about 300 million dollars per year in the USA (1). Other studies reported the therapeutic effect of hypertonic saline 3% on acute bronchiolitis in children.

In Kuzik et al study, use of hypertonic saline 3% nebulizer had more pleasant outcomes than the normal saline nebulizer (15). Also, Anil et al who used some medicinal combinations (Ventolin nebulizer with normal saline, epinephrine with normal saline and hypertonic saline 3%, and normal saline alone) for treating acute bronchiolitis in the children found no significant difference among the groups; the effect of hypertonic saline 3% was similar to that of other medicinal combinations (9). In Zhang et al study, hypertonic saline 3% was more effective than normal saline (16). Mandelberg et al compared hypertonic saline 3% nebulizer, epinephrine, and normal saline and found hypertonic saline 3% as therapeutically more effective on bronchiolitis symptoms (13). In Ater et al study to compare Ventolin nebulizer and normal saline with Ventolin and hypertonic saline 5%, hypertonic saline 5% had a better therapeutic effect in the children with bronchiolitis than normal saline (20).

In addition, the decreased length of hospitalization after hypertonic saline 3% was observed in some studies (10,13,17,20).

The results of this study indicated that the mean RDAI criterion on the day of hospitalization and prior to treatment initiation had no significant difference between the two groups. The RDAI criterion indicating the disease severity had no significant difference between the two groups of study. In addition, the mean RDAI criterion was significantly lower in the hypertonic saline 3% group, indicating that use of hypertonic saline 3% nebulizer was obviously more efficient than Ventolin after hospitalization (the day one), i.e. from the day two to the last day of hospitalization. The greatest efficacy was noted on the day two after hypertonic saline 3%, irrespective of the last day of hospitalization. In Sarrell et al study, there was no significant difference prior to study between the two groups, but an obviously better effect was noted by the use of hypertonic saline 3% nebulizer between the
day two and the last day of hospitalization. However, the highest effect was observed in the day two. The effect of hypertonic saline 3% was obviously better than that of normal saline on the day one in Sadbhavna et al study, but no difference was observed between the two groups on the day two (17, 21).

The length of hospitalization in our study was 2-6 days, similar to other studies (13, 15, 16). Although the mechanism of hypertonic saline 3% was not examined in the present work, hypertonic saline 3% was demonstrated to increase mucus clearing in Dasgupta et al study (22). Also, Tomooka et al examined the effects of hypertonic saline on recovery of respiratory symptoms. These effects included improvement of mucus cilia’s function, decrease in mucous edema and inflammatory mediators, mechanical clearing of secretions, and improvement of mucus function (23). On the other hand, no complication was observed by the use of hypertonic saline 3% nebulizer throughout the study in the studied patients, which is consistent with other studies (9, 13, 18).

Conclusion

Regarding the obtained results in this study and comparing them with those of other studies, we found hypertonic saline 3% nebulizer as having more pleasant therapeutic effects than Ventolin on treating acute bronchiolitis in the children under two years. Moreover, use of hypertonic saline 3% leads to no remarkable drug-related complications in the children; the reason is that it is free of medicinal compounds. In addition, the treatment costs would decline considerably and hence use of hypertonic saline 3% nebulizer is recommended for treatment of acute bronchiolitis in children under the age of two.

Acknowledgements

We gratefully thank Research and Technology Deputy of Shahrekord University of Medical Sciences.

Conflict of interest

The authors report no conflict of interest.

References

17. Sarrell EM, Tal G, Witzling M, Someck E,


