Oral Atenolol Therapy in the Treatment of Infantile Hemangioma: A Prospective Study
Kh Ahsanul Kabir¹, KMN Ferdous², F Hossain³, MK Islam⁴

Abstract

Background: Infantile haemangiomas (IH) are the most common tumors of infancy with an incidence of >2% of infants in general, and of 10% of Caucasian children, in particular.

Objective: The aim of this study was to assess the efficacy and safety of atenolol in the treatment of proliferating IHs.

Methods: This prospective study was conducted among 120 patients’ up to 30 months of age, clinically diagnosed with IH’s in surgical outpatient department (SOPD) of the Dhaka Shishu Hospital from July 2013 to March 2018. Atenolol was started initially 0.5mg/kg. Then doses were titrated according to clinical response (0.5-3mg/kg/day OD). Parents were instructed to interrupt the administration of the drug if the child had a serious cough with dyspnea or gastroenteritis with vomiting or diarrhea. Patients were advised to come for follow up visit and in each follow up blood pressure, percentage of regression (of size), color change, complications were recorded. The change in the appearance of IH was evaluated on a visual analogue scale (VAS). Regression in the size and color clearance of IHs were evaluated according to 0%-to-100% scale.

Results: Majority of the babies were from 1-10 months age group. Most of the infants were term babies and majority of the infants were girls. Majority of the IHs were located in head and neck regions. Majority of the mothers had multiple gestation pregnancy and few had family history of IH. Excellent, good and fair colour regression was found in 74.17%, 22.50% and 3.33% participants respectively. Excellent, good and fair size regression was found in 57.50%, 38.33% and 4.17% participants respectively. Adverse effects of drug like loose motion (7.50%) and sleep disturbance (3.33%) were found among the participants.

Conclusion: Atenolol is effective in the treatment of IHs with few minor side effects.

Key words: Infantile hemangioma, atenolol, adverse effect.

Introduction

Infantile haemangiomas (IH) are the most common tumors of infancy with an incidence of >2% of infants in general, and of 10% of Caucasian children, in particular. IH typically presents few a weeks after births, and occur more frequently in girls.¹ Many are benign and do not require treatment, but potential complications include airway obstruction, visual

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compromise, disfigurement, ulceration, congestive heart failure, and death.² Previously, corticosteroids were the mainstay of treatment for complicated IHs. However, corticosteroids have undesired side effects, such as temporary growth retardation, an increased risk of infection, and behavioral changes.³ Recently, propranolol, a nonselective β-blocker, became the preferred treatment for complicated or select types of IHs.¹ Still, its use is not risk-free and many adverse events have been documented, including: hypoglycemia, bronchial obstruction, hypotension, seizures, sleep disturbances, and gastrointestinal symptoms, among others.⁴ On the other hand, atenolol, a hydrophilic cardio selective beta-blocker that acts principally on h1 receptors, does not cross the blood-brain barrier and has less h2 effects.⁵ The aim of this study was to assess the efficacy and safety of atenolol in the treatment of proliferating IHs.

Materials and Methods
This prospective study was conducted among 120 patients up to 30 months of age, clinically diagnosed with IHs in surgical outpatient department (SOPD) of the Dhaka Shishu Hospital from July 2013 to March 2018. Patients previously treated for Infantile Hemangioma or having hemangioma presenting life threatening condition, cardiovascular disease contraindicating use of propranolol, known hypersensitivity to atenolol were also excluded from the study. Measurement of lesion was taken along long axis and another one perpendicular to this axis in centimeters with flexible soft rubber tape. Photograph was taken with digital camera. These photographs were taken prior to the treatment (baseline) and after the start of treatment at 2-8 weeks and 11-24 weeks. The change in the appearance of IH was evaluated on a visual analogue scale (VAS). The VAS uses a 100-mm scale on which –100 represented a doubling in the size and extent of the IH, 0 represented no change/baseline and +100 represented complete disappearance.⁶ Atenolol was started initially 0.5mg/kg. Then doses were titrated according to clinical response (0.5- 3mg/kg/day OD). If no relevant adverse effects were detected after administration and if this was well tolerated, treatment was continued. Parents were instructed to interrupt the administration of the drug if the child had a serious cough with dyspnoea or gastroenteritis with vomiting or diarrhoea. They were also advised to call the investigator over phone for any adverse effects. Patients were advised to come for follow up visit at 2nd week, 1st month, 2nd month, 4th month, 6th month and 8th month. In each follow up blood pressure, percentage of regression (of size), color change and complications were recorded.

Regression in the size and color clearance of IHs were evaluated according to 0%-to-100% scale. An excellent response denotes 80% to 100% regression or color clearance. A good response denotes 50% to 80%. A fair response denotes 25% to 50% and finally a poor response denotes 25% or less.⁷ Frequency distributions for categorical variables were used to describe the characteristics of the total sample. The findings of the study were presented by frequency, percentage in tables and graphs.

Results
Majority of the babies were from 1-10 months age group. Most of the infants were termed babies and majority of the infants were girls. Majority of the IHs were located in head and neck regions. Majority of the mothers had multiple gestation pregnancy and few had family history of IH (table I). Excellent, good and fair colour regression was found in 74.17%, 22.50% and 3.33% participants respectively (figure 1). Excellent, good and fair size regression was found in 57.50%, 38.33% and 4.17% participants respectively (figure 2). Adverse effects of drug like loose motion (7.50%) and sleep disturbance (3.33%) were found among the participants (table II).

<table>
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<th>Table I</th>
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<td><strong>Base line characteristics of the participants (n=120)</strong></td>
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<td>Characteristics</td>
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<td><strong>Age (in months)</strong></td>
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<td>11-20 months</td>
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<td>&gt;20 months</td>
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<td><strong>Term baby</strong></td>
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<td><strong>IH in head and neck region</strong></td>
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<td><strong>Multiple gestation pregnancy</strong></td>
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<th>Table II</th>
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<tr>
<td><strong>Adverse effect of drugs (n=120)</strong></td>
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<td>Adverse effect</td>
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Infantile hemangiomas (IHs), also known as “hemangiomas of infancy,” are common benign tumors of endothelial cells characterized by a unique pattern of rapid proliferation that occurs in the first months of life. Infertility (IHs) are more common among female infants, however, although older data suggest female to male ratios ranging from 3:1, more recent studies suggest a range of 1.4:1 to 3:1. Similar result found in the current study where it was found that majority of the participants were female. The present study also that most of the infants were termed baby and product of multiple gestations. Other study also reported that infants with hemangiomas are more likely to be female, premature, and product of multiple gestations. Hemangioma can be found in all regions of the baby, but they occur most commonly in the head and region (60%), followed by the trunk (25%) and then the extremities (155). Majority of the infants (67.5%) of the current study also had IH located in the head and neck regions.

The first noticeable effects on treatment were the changes in color and softening of IH, followed by regression of their sizes. Color regression of IH was determined by blind investigator using digital photograph. The photographs were taken prior to be treatment, after the start of treatment at 4 months and at 8 months. Excellent colour regression was found in 74.17% infants, Excellent size regression was found in 57.50% infants. This result was consistent with other studies. Abraz, ua-Araya et al. had conducted a prospective comparative study to evaluate the effectiveness of atenolol against propranolol for the treatment of IH where they reported that patients treated with atenolol had a complete response of 53.8%. The study of Jiet al. evaluated the efficacy and safety of atenolol in the treatment of proliferating IH. They observed an “excellent” treatment response (complete or nearly complete resolution of the IH) in 56.5% of patients at week 24.

Clinical studies that addressed the adverse effects of atenolol in children with IH have generated conflicting results. In a study by Abarzua-Araya et al, the authors found no significant adverse event in atenolol.
treatment during the 6-month follow up. In the present study, any adverse events were recorded by parents between study visits and were documented by investigations at each visit. Among the 120 infants, few developed minor adverse effects like loose motion (7.50%) and sleep disturbance (3.33%). No patients developed any major adverse effects. In the study of de Graaf M, et al, mild side effects occurred in 40% (12/30) of patients and severe side effects occurred in 3% (1/30). Other study most common side effects of atenolol in the treatment of IHs were diarrhea, followed by central nervous system effects. These minor side effects occurred in the present study were treatment and ceased after cessation of the drug.

Conclusion
Atenolol is effective in the treatment of IHs with few minor side effects.

References