

Cystichyroma with Macroglossia-A Lymphatic Malformation in a Neonate

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Introduction

This 15 days old male infant born at 38 weeks of gestation by spontaneous vaginal delivery was brought to paediatric emergency room with the history of hurried breathing and poor feeding and swelling in the neck (Figure 1). Clinical diagnosis of cystic hygroma with air way compromise was made. Antenatal scans done were normal. In view of deteriorating respiratory distress the child was ventilated. X-ray of neck showed mass on either side of the neck (Figure 2). Ultrasound demonstrated bilateral cystic areas consistent with cystic hygroma

(multicystic) (Figure 3). CT neck was done to rule out any thoracic extension (Figure 4). The mass was infiltrated with sclerosant OK 432 (FDA approved) under ultrasound.

Discussion

Cystic hygroma a lateral neck mass is a shocking scene for any parents. Air way compromise is the major threat to the baby. Early recognition, respiratory care, timely sclerosant therapy, surgery in selected candidates and adequate informative counselling are the main challenges for the treating physician.



Figure 1: A new born with cystic hygroma with macroglossia.



Figure 2: X-ray neck showing bilateral mass in the neck.



Figure 3: Ultrasound neck showing cystic hygroma.



Figure 4: C T neck showing cystic hygroma bilateral.

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