Tracheoesophageal fistula with complex injury to the existing anomaly - Youngest neonate to survive: A case report

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KEYWORDS

Esophageal atresia, Tracheoesophageal fistula, Neonate, Preterm, Tracheoesophageal trauma

ABSTRACT

Background: Complex injury to the esophagus, trachea, and tracheoesophageal fistula in a case of esophageal atresia is rarely reported that requires prompt identification and management.

Case Presentation: A very-low-birth-weight preterm with a tracheoesophageal fistula on ventilatory support for respiratory distress syndrome was transferred to our hospital. The baby developed air-leak through the tracheoesophageal fistula into the stomach causing gross distension of the stomach and reduction in oxygen saturation. Repositioning and reintubation failed to reduce the abdominal distension and to improve the saturation. Subsequent intubation with large volume, low pressure cuffed tube, and replacement of Replogle tube resulted in pneumothorax and further deterioration requiring emergency thoracotomy. Complex tracheoesophageal injury to the esophagus, trachea, and fistulous tract was repaired.

Conclusion: The baby survived the early repair of the tracheoesophageal fistula and the iatrogenic complex tracheoesophageal injury. Pre-operative tracheo-esophagoscopy with stenting of the fistula with Fogarty catheter helped to carry out a successful repair technically.

INTRODUCTION

Esophageal atresia (EA) is one of the most common life-threatening congenital anomalies of the newborn and occurs in 2.44 in 10,000 births. The current survival rate of EA with Tracheoesophageal fistula (TEF) in term neonates with birth weight more than >2500 gm - normal birth weight (NBW) and >1500 g to <2500 g - low birth weight (LBW), without major congenital cardiac anomaly, is 95% and can be achieved with early primary repair.[1] The very low birth weight (VLBW) babies (>1000 g to <1500 g) had poor survival with a mortality ranging from 100% in the early fifties to 20% in the late nineties which was attributed to intrauterine growth retardation, respiratory distress syndrome (RDS), associated anomalies, cardiopulmonary vulnerability, necrotizing enterocolitis, intracranial hemorrhage, and immature immune system.[2] Since the late nineties, the survival in this group has increased to 87%. [3] VLBW babies with EA are more commonly associated with isolated EA (10%) than EA with TEF. VLBW babies with isolated EA have better survival. VLBW babies with EA and TEF are a special group because of the intrauterine growth retardation, related RDS, the risk of gross air-leak through the fistula into the stomach, and its complication- the gastric perforation, which carries a mortality rate as high as 70%.[2] The iatrogenic injuries caused during the preoperative period of management in preterm babies with TEF are uncommon, life-threatening, under-recognized, and had poor survival. This report is about a baby belonging to such a special group resulting in a gross air leak and cardiovascular collapse requiring repositioning of the endotracheal tube (ETT). Several attempts to replace ETT, suctioning, and replacement of Replogle tube resulted in complex tracheoesophageal trauma and pneumothorax

CASE REPORT

A preterm female baby born by emergency LSCS for decreased fetal movements at 32 weeks weighing 1.1
kg with EA/TEF was transported on ventilatory support to our institution at three days of age. The baby developed poor chest rise, desaturation, and bradycardia due to a large air leak into the stomach through the fistula. Initial attempts by adjusting the position of the ETT failed. Further attempts by changing the ETT and subsequently with a large volume low pressure cuffed tube as well as replacing the Replogle tube ended up in pneumothorax (Fig.1).

Intercostal drainage (ICD) was inserted, and the baby was shifted to OR for emergency surgery. Preoperative laryngo-tracheoscopy (PLT) showed a fistulous opening at the usual site proximal to the carina. The opening was wide and the edges were irregular suggestive of trauma (Fig.2a,2b & 2c). A Fogarty catheter was introduced into the fistulous opening and the balloon was placed in the stomach, which was verified by fluoroscopy (Fig.2d).

PLT with a cystoscope with an offset lens and placement of Fogarty catheter has been routinely carried out by us for the last five years.[4] Right posterolateral thoracotomy by trans-pleural approach was carried out. The feeding tube passed into the upper pouch at the time of induction replacing the Replogle tube had come out through a tear in the upper pouch into the pleural cavity. The fistula was torn along its entire length except for a sliver of tissue medially holding the trachea and the lower pouch exposing the Fogarty catheter. The endotracheal tube could be seen through a tear extending into the trachea from the proximal end of the fistula (Fig.3a). After a careful dissection with the help of the exposed stent which was seen bridging across the tracheal end of the fistula and lower pouch through the torn fistulous tract, the remaining fistulous tract was disconnected from the trachea, and the lower pouch and was excised. The fistulous opening and the adjacent tear on the trachea were closed. The torn tissues of the lower end of the upper pouch were trimmed. The anastomosis of the upper and lower pouches was carried out with slight tension (Fig.3b).

To tide over the tension at the anastomosis, a trans anastomotic nasogastric 8F Silastic Foley balloon catheter was placed as a stent. The balloon inside the stomach was inflated and gentle traction was given at the nasal end of the Foley catheter till a slackness was felt at the anastomosis and the catheter was fixed to the nose. This technique to reduce the tension at the anastomosis has been used by us for the last 10 years.[5] There was no leak in the postoperative contrast esophagogram (Fig.3c). The baby required prolonged CMV and TPN. The baby was discharged at the age of 4 months with adequate weight (2.94 kg) for the age and normal oral feeding and is doing well 14 months after the surgery.
The survival of VLBW babies has improved in recent years. This improved survival is related to the consideration of early primary repair in stable cases and delayed esophageal anastomosis or staged surgery in unstable cases, availability of endotracheal intubation, and advances in neonatal, respiratory, surgical, and anesthetic care. There are controversies in the method of management of the VLBW babies with EA and TEF between primary repair and delayed primary anastomosis and staged surgery among the institutions.[2,3,6]

VLBW babies with EA with TEF are a special group and may require methods of surgical management other than the primary anastomosis which was first performed by Haight in 1943. This group is considered special because of the intrauterine growth retardation, related RDS, and the risk of progressive atelectasis, loss of functional residual capacity, increased pulmonary vascular resistance, and frequent requirement of continuous mechanical ventilation (CMV). High pressures in the pulmonary parenchyma and the relatively lower pressure of the fistula creates a ventilatory leak through the TEF. This worsens the respiratory distress secondary to the loss of ventilation and resultant gastric distention leading to cardiovascular collapse and gastric perforation which carries a mortality rate as high as 70%.[2,7] The aggressive non-surgical methods adopted to tackle the air leak through the fistula into the stomach before surgical correction are positioning of ETT tip distal to the fistulous opening, positioning with affected side down, placement of Fogarty catheter either by bronchoscopy or by laparotomy to occlude the fistula with the balloon, banding of the esophagus, gastric division, nec-

### Table 1: Cases of iatrogenic tracheoesophageal injury in EA with or without TEF

<table>
<thead>
<tr>
<th>Year / Author</th>
<th>Age / EA type / Age of onset of symptoms</th>
<th>Site of perforation</th>
<th>Cause</th>
<th>Aggressive non-surgical methods</th>
<th>Type of repair</th>
<th>Surgical procedures</th>
<th>Survival / Death</th>
</tr>
</thead>
<tbody>
<tr>
<td>1978 Wright VM[11]</td>
<td>LWB 1800 g 34 wks. TEF Type C. 18 Hours of life</td>
<td>Upper pouch</td>
<td>Replogle Plastic sump catheter</td>
<td>Nil</td>
<td>Staged surgery</td>
<td>Fistula ligation, Cervical esophageostomy</td>
<td>Survived</td>
</tr>
<tr>
<td>1986 McLeod BJ[9]</td>
<td>Preterm, LBW,1600 g. TEF Type C Day 1</td>
<td>Trachea at the level of fistula</td>
<td>Cole’s tracheal tube 3 mm (Thick) without stylet</td>
<td>Nil</td>
<td>Staged Surgery</td>
<td>Fistula ligation and gastrostomy</td>
<td>Died.</td>
</tr>
<tr>
<td>2013 Acker SN[12]</td>
<td>ELBW, 755 g. TEF Type C. Day 2 of life</td>
<td>Upper pouch and isolated ileal perforation</td>
<td>Orogastic tube</td>
<td>Nil</td>
<td>Primary anastomosis</td>
<td>Thoracotomy – Primary esophageal anastomosis and gastrostomy</td>
<td>Alive and thriving</td>
</tr>
<tr>
<td>2013 Parekar S[13]</td>
<td>VLBW 1400 g. Pure EA. Day 3 of life</td>
<td>Upper pouch</td>
<td>Stiff 10F rubber catheter used for contrast study of upper pouch</td>
<td>Nil</td>
<td>Cervical esophageostomy and feeding gastrostomy without thoracotomy</td>
<td>Results not mentioned</td>
<td></td>
</tr>
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</table>

**DISCUSSION**

The survival of VLBW babies has improved in recent years. This improved survival is related to the consideration of early primary repair in stable cases and delayed esophageal anastomosis or staged surgery in unstable cases, availability of endotracheal intubation, and advances in neonatal, respiratory, surgical, and anesthetic care. There are controversies in the method of management of the VLBW babies with EA and TEF between primary repair and delayed primary anastomosis and staged surgery among the institutions.[2,3,6]

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REFERENCES


