

## **Title Page**

### **Predicting Factors of Decannulation in Children with Tracheostomy**

Fusun Unal, MD, Medipol University Faculty of Medicine, Department of Pediatric Pulmonology  
Istanbul, Turkey

Emine Atag, MD, Medipol University Faculty of Medicine, Division of Pediatric Pulmonology  
Istanbul, Turkey

Leyla Telhan, MD, Medipol University Faculty of Medicine, Department of Pediatrics Istanbul,  
Turkey

Burcu Gizem Teber, Medipol University Faculty of Medicine, Department of Pediatrics Istanbul,  
Turkey

Fazilet Karakoc, Professor, Marmara University Faculty of Medicine, Division of Pediatric  
Pulmonology Istanbul, Turkey

Sedat Oktem, Professor, Medipol University Faculty of Medicine, Pediatric Pulmonology Istanbul,  
Turkey

Address correspondence to: Füsün Ünal, Department of Pediatrics, Medipol University Hospital  
Göztepe mah., Metin Sk. No:4, 34214 Bağcılar/İstanbul, Turkey, ([dr\\_fusun@yahoo.com](mailto:dr_fusun@yahoo.com))

Tel: +90 5326569407

Fax: +90212 4607070

Key words: Decannulation, children, colonization

Running title: Predicting Factors of Decannulation in Children with Tracheostomy

Funding Source: No external funding for this manuscript.

Financial Disclosure: The authors have indicated they have no financial relationships relevant to this article to disclose.

Conflict of Interest: The authors have indicated they have no potential conflicts of interest to disclose.

## PREDICTING FACTORS OF DECANNULATION IN CHILDREN WITH TRACHEOSTOMY

### Abstract

**Objectives:** Our aim was to determine the treatable causes to increase the chance of decannulation success. For this purpose we evaluated the differences between the patients who successfully decannulated and the patients who still has tracheostomy.

**Methods:** A retrospective cohort study was conducted based on medical records of all pediatric patients with tracheostomy in a single centre.

**Results:** Decannulation was successfully achieved in 59 patients (34.5%) of total 171 patients with tracheostomy between the years 2012-2019. Median duration of tracheostomy was 41.5 and 12 months in patients who remained with tracheostomy and decannulated respectively. Neurological disorders were higher in patients remained with tracheostomy, congenital heart disease and airway abnormalities were higher in decannulated patients. Presence of bacterial colonization (3.8-fold), history of invasive respiratory support following tracheostomy (2.9-fold), and having any neurological disorder and/or comorbidity (5.2-fold) were significantly associated lower rates of decannulation.

Almost 33 % of patients had bacterial colonization and colonization rates were higher in patients who needed invasive respiratory support following tracheostomy placement ( $p < 0.001$ ), patients with feeding/swallowing problems ( $p = 0.005$ ) and neurological disorders ( $p = 0.002$ ). There was significant correlation between duration of tracheostomy and bacterial colonization rates ( $p = 0.008$ ). But after analysing with logistic regression only having a neurological disorder was associated with bacterial colonization (OR= 2.9; 95% CI: 1.15-7.47  $p = 0.024$ ).

**Conclusion:** While conducting decannulation assessment, the presence of colonization should be considered. Future prospective researchs are necessary in order to determine the role of chronic colonization on decannulation success.

# **PREDICTING FACTORS OF DECANNULATION IN CHILDREN WITH TRACHEOSTOMY**

## **INTRODUCTION**

Tracheostomy is a surgical procedure that provides artificial airway patency. The indications of tracheostomy have been changed over the past few decades from emergency procedure like trauma or infectious diseases to prolonged mechanical ventilation or underlying neurological and cardiological diseases<sup>1-5</sup>.

While the number of patients with tracheostomy remained stable over the years<sup>7</sup>, the age at the time of tracheostomy procedure decreased<sup>5,8</sup>. The accompanying chronic conditions are increased and almost 80% of the patients are accompanied by more than one chronic condition, which is associated with longer hospital length of stay, higher hospital charges and higher in-hospital mortality<sup>4</sup>. Increased number of cumulative comorbidities lower the chances of tracheostomy decannulation<sup>9</sup>. Persistent and recurrent lower respiratory infections due to chronic bacterial colonizations are the most common causes of the hospitalizations in the children with tracheostomy<sup>10-15</sup>. Until now there is no study investigating the effect of bacterial colonization on possibility of decannulation.

Decannulation should be planned as early as possible to minimise tracheostomy related complications, to improve the quality of life of the patients and caregivers, as well as to reduce the health care costs<sup>16</sup>.

In children, the overall successful decannulation rates vary between 24 to 60%<sup>5,8</sup>. Decannulation decision has been made by clinical judgement according to the preliminary assessment in line with the consensus reports and checklist<sup>17-19</sup>. There is no consensus protocol for decannulation practices that changes based on the patient characteristics and conditions of center.

Our aim was to investigate the differences between the patients who successfully decannulated and the patients who still has tracheostomy in terms of underlying diseases, comorbidities, bacterial colonization and previous need for respiratory support. We also aimed to evaluate the factors associated with decannulation success.

## **MATERIAL-METHOD**

### **Patients:**

We retrospectively reviewed the medical records of patients with tracheostomy at our institution between July 2012 to December 2019. Data including age, gender, underlying disease, presence of comorbidities, bacterial colonization and invasive respiratory support following tracheostomy were obtained from patient's medical records.

While the main causes of tracheostomy are classified as primary disorders, other systemic static problems are defined as comorbidities.

Primary disorders that cause tracheostomy were classified as, neurologic disorders (ND), congenital heart diseases (CHD), chronic pulmonary diseases (CPD), airway abnormalities, metabolic/genetic disorders and other causes. ND group was involved mostly cerebral palsy, spinal muscular atrophy (SMA Type 1 and 2), neuromuscular disorders, refractory epilepsy, central nervous system trauma and cancer.

Comorbidities were classified as, neurological, cardiological and feeding and swallowing problems. Neurological comorbidities were concomitant problems including minor static disabilities and controlled epilepsy except than the primary neurologic disorder. Cardiological comorbidities were included hemodynamic stable structural defects as VSD (ventricular septal defect), ASD (atrial septal defect), MVP (mitral valve prolapses).

Feeding status of the patients (Tube feeding or oral feeding) and presence of feeding/swallowing problems, (which was defined as abnormal swallowing study or respiratory symptom such associated with feeding) were also obtained from patients' medical records.

Tracheal aspirate cultures were obtained every three months during routine clinical visits regardless of any clinical signs of infection or any time with presence of respiratory infections at our clinic. Bacterial colonization was defined as the presence of the same microorganism ( $< 100.000$

colonies per ml) in more than 3 consecutive cultures of tracheal aspirate without any infection (Lep-  
aiteur, Morar). We included the results of trach aspirates in the last year before decannulation in  
decannulated patients. In patients with remained tracheostomy dependency we evaluated the  
cultures of last one year before December 2019 by the end of the study period.

Required surgical procedures before decannulation, post decannulation complications, need for res-  
piratory support following tracheostomy and decannulation failure were also recorded. Decannula-  
tion failure was defined as an invasive airway need within 48-96 hours following planned decannu-  
lation attempt<sup>20</sup>.

### **Tracheostomy decannulation protocol:**

Patients who do not need respiratory support more than 16 hours a day, have no recent history of  
aspiration, can control their airway secretions and have an effective cough reflex were considered  
candidate for decannulation. Patients with a neurologic disease who can use cough assist successfully  
were also included to decannulation protocols.

Before decannulation decision, first step was to evaluate airway anatomy and patency by flexible  
bronchoscopy. Airway pathologies requiring surgical correction were treated before decannulation.

Tracheostomy tube was downsized to a smaller diameter and capped for a 24-48 hours according to  
patients' clinical condition. Decannulation was performed in children who was stable during tube  
capping period who has adequate airway patency, stable night/day time oxygen saturation, normal  
blood gas results and absence of any sign of infection. Patient was monitored in the intensive care  
unit or pediatric ward for at least 48 hours according to patient's clinical status. Oxygen  
supplementation and chest physiotherapy were applied if necessary.

### **Statistical Analysis**

The IBM SPSS Statistics (version 22.0 IBM Corp., Armonk, NY) software program was used to  
analyze the data. Categorical variables are presented as numbers (n) or percentages (%). Continuous  
variables are shown as medians with 25 and 75 percentages. For comparisons; chi-square test, Mann

Whitney u test and logistic regression analysis were used as appropriate. Odds Ratio (OR) were estimated using logistic regression. A p-value < 0.05 was considered significant.

### **Ethical Approval**

The study was approved by the Ethical Committee of Medipol University Medical Faculty (Number: 10840098-604.01.01).

## **RESULTS**

### ***Clinical Demographics and Indications***

Between July 2012 to December 2019, we evaluated 171 tracheostomized patients (54% male). Median age of all the patients was 66 (42-93) months. Neurological disorders (49%), congenital heart diseases (17,5%) and airway abnormalities (11,7%) were the most common primary disorders. Decannulation was successfully achieved in 59 patients (34.5%). Median age of the patients who remained with tracheostomy and decannulated were 66 and 67 months, respectively.

Eightytwo (48%) of the patients needed invasive respiratory support following tracheostomy. Fiftynine of the 112 tracheostomy patients and 23 of the 59 decannulated patients needed invasive respiratory support following tracheostomy (Table 1). Seven patients (3 patients with congenital central hypoventilation, 3 patients with SMA, and 1 patient with severe airway malacia) required non-invasive ventilation following decannulation. Sixteen of the 23 patients detached from ventilator before decannulation during follow-up period.

Median duration of tracheostomy was 41.5 (25-66.5) and 12 (5-20) months in patients who remained with tracheostomy and decannulated respectively. At decannulation 24 % of patients were under 1 year old while 55% of them were under two years old.

While neurological disorders, invasive respiratory support following tracheostomy, bacterial colonization were higher in patients remained with tracheostomy, congenital heart disease and airway abnormalities were higher in decannulated patients (Table 1).

### ***Bronchoscopic Findings***

Bronchoscopic evaluation were performed before decannulation in 59 patients and 15 patients (25%) had normal bronchoscopic findings. Also fifteen (25%) of the patients have more than one abnormal bronchoscopic findings. Abnormal bronchoscopic findings at 44 patients were listed at Table 2. Patients with subglottic hemangiomas were treated medically before decannulation. Granulation tissue was required surgical excision in 6 patients. Six patients with grade 3 subglottic stenosis according to Cotton Myer scoring system, underwent laser surgery before decannulation. Three



patients with supraglottic collaps (2 patients with SMA type 1, one patient with SMA 2) and one patient with severe tracheomalasia required NIV after decannulation. Also another 3 patients with congenital central hypoventilation syndrome were decannulated with NIV and one of these patients had mild granulation tissue and two of them were normal in the bronchoscopic evaluation.

### ***Risk Factors for Affecting Decannulation***

Bacterial colonization was present in 56 (33%) patients. *Pseudomonas aeruginosa* (60 %) and *Staphylococcus aureus* (13%) were the most commonly identified pathogens. Bacterial colonization was significantly higher in patients remained with tracheostomy ( $p < 0.001$ ), underlying neurological disorder ( $p=0.002$ ), who required invasive respiratory support after tracheostomy placement ( $p<0.001$ ) and patients with feeding and swallowing problems ( $p=0.005$ ). There was significant correlation between duration of tracheostomy and bacterial colonization rates ( $p=0.008$ ). Tracheostomy duration was 70 months and 52 months respectively, in colonized and noncolonized patients. Having a neurological disorder was an independent risk factor of colonization after the analysis with logistic regression (Table 3).

Logistic regression analysis revealed affecting factors related with decannulation; presence of bacterial colonization (3.8-fold), history of invasive respiratory support following tracheostomy (2.9-fold) and having any neurological disorder and/or comorbidity (5.2-fold) were significantly associated lower rates of decannulation. Having a neurological disorder was also an independent risk factor of bacterial colonization (Table 4).

### ***Decannulation Failure, Complications and Mortality***

Three patients required tracheostomy placement in 24-72 hours following decannulation. Two of these three patients couldn't tolerated decannulation due to congenital heart disease and hemodynamic instability. Second decannulation attempt after 6 months was successful in the other patient who had traumatic brain injury and improvement after rehabilitation period. Consequently, 2 patients (3,3 %) were accepted as decannulation failure in our study.

No complication was found in 42 (71 %) of the 59 patients following decannulation. Atelectasis,

pneumonia and stridor were the most common complications (Table 5).

Eighteen patients (11%) were died during study period because of underlying medical conditions before decannulation. No decannulated patient died during study period.

## **DISCUSSION**

Decannulation under safe circumstances, should be considered in all patients with tracheostomy as soon as possible when the initial indication is no longer exists. Determination of the criterias that predict successful decannulation is important in order protect patients and their families from unnecessary risks and false expectations. Only few studies evaluated clinical factors affecting decannulation success rate in patients with tracheostomy. In these studies, there are big variations in terms of sample size and underlying conditions<sup>21-24</sup>.

For the first time in the literature, our study showed that colonization effects decannulation rates.

### ***Indications***

Indications of tracheostomy may vary according to the specialization of centre and patient population. Most frequent tracheostomy indications were ND (49%) and CHD (18%) in our study. The indications of pediatric tracheostomy are shifting to dependency of prolonged ventilation in last 30 years<sup>8,25</sup> and the most frequent indications are neurological (% 39-51) and cardiological diseases (% 21-34) in large pediatric series<sup>5,6</sup>.

### ***Bronchoscopic Findings***

Almost in all studies bronchoscopic evaluation is recommended before decannulation for avoiding complications after decannulation and reducing decannulation failure<sup>17,18,26</sup>. In our study, airway granulation tissue was the most common bronchoscopic finding which was detected 39% of bronchoscopic evaluation. The incidence of airway granulation tissue in patients with tracheostomy ranges from 50 to 80% and also granulation tissue is the most common pre-decannulation bronchoscopic finding up to 51% in literature<sup>27,28</sup>. There was no correlation between duration of tracheostomy and existence of grannulation.

### ***Risk Factors Affecting Decannulation***

Studies suggested some predictive factors for decannulation, including age at time of tracheostomy, neurologic disease, and anatomic airway disease<sup>29-31</sup>. In our study, 59 patients (34.5 %) were decannulated and the median duration of tracheostomy at decannulation was 22 months. Our results

are consistent with the literature in which decannulation rates are ranging between 24 -60 %<sup>5,6,8</sup>. Seventy percent of our patients were less than 1 year of age at the time of tracheostomy placement and were 22 months of age at the time of decannulation, which are also consistent with literature rates<sup>5,8</sup>. Presence of genetic abnormalities, GERD, dysphagia, irregular respiratory pattern (breath holding spells), false expectations of the parents, and the absence of a multidisciplinary team follow-up were the main factors for decannulation failure<sup>21,22,24</sup>.

Our study suggest that bacterial colonization affect the decannulation success in addition to neurological disorders and invasive ventilation requirement following tracheostomy placement.

### *Bacterial Colonization*

Bacterial colonization is a common problem in patients with tracheostomy. In our study almost 33 % of patients had bacterial colonization and the most frequent identified pathogen was *Pseudomonas aeruginosa* (60 %) in accordance with previous studies<sup>23,32-34</sup>. Natural protective effect of the upper airways is eliminated in patients with tracheostomy. Additionally, altered anatomy and recurrent aspiration of airway secretion may lead to chronic colonization and subsequent infections<sup>13,35</sup>. In patients with tracheostomy persistent detection of *Pseudomonas aeruginosa* in tracheal aspirate cultures is known to increase the risk of pulmonary exacerbations<sup>11</sup> and associated with worse clinical outcomes, increase the number of hospitalizations and length of stay in intensive care unit<sup>12</sup>.

Our study shows that presence of a feeding and swallowing disorders, respiratory support requirement following tracheostomy placement, concomitant neurological disorders and duration of tracheostomy were the main factors for bacterial colonization. Aspiration of the gastric contents, upper airway secretions and food into airways may cause airway inflammation, facilitate the colonization of the airways with pathogenic bacterias which may lead to lower respiratory tract infections and bronchiectasis<sup>33-36</sup>. Also feeding dysfunction decreased the odds of a successful first decannulation attempt (5.5-fold) and cause decannulation failure<sup>21,37</sup>.

Colonization rates were also higher in patients who needed invasive respiratory support following tracheostomy placement and patients with feeding/swallowing problems ( $p<0.001$  and  $p=0.005$ ).

In a recent unpublished study conducted at our center, revealed that the nebulized antibiotic treatment reduced the number of hospitalizations, length of stay in the intensive care unit and the bacterial load in the tracheal aspirate cultures in patients with tracheostomy who has persistent bacterial colonization <sup>38</sup>. In line with previous studies, the most common underlying disease in children with tracheostomy were also neurological disorders in our study and colonization rates with pathologic microorganisms were more common this group of patients (65%) <sup>5,6,23,32-34</sup>. Neurological disorders were independent risk factor of colonization regardless of tracheostomy duration (2.93 fold). Bacterial colonization rate was associated with the longer duration of tracheostomy, median duration of tracheostomy was 70 and 52 months in colonized and non-colonized patients respectively. Pozzi et al evaluated the respiratory colonization rates among 65 pediatric tracheostomized patients after acute brain injury in a long-term rehabilitation center. Their study revealed that these patients had high colonisation rates in which *Pseudomonas aeruginosa* and *Staphylococcus aureus* were the most commonly identified microorganisms. Of the 23% and 26 % of the patients colonized with one and more additional microorganisms respectively during rehabilitation period <sup>23</sup>.

This is the first study in literature which shows that the presence of colonization reduces the odds of decannulation rate by 3.8-fold.

#### *Invasive Ventilation Requirement Following Tracheostomy*

Almost half of our patients (48%) required invasive ventilation following tracheostomy placement which was associated with lower decannulation rates (2.9 folds). In literature the need for respiratory support in patients with tracheostomy were 33-44 % and there was no correlation between the duration of respiratory support via tracheostomy and decannulation rate<sup>5,39,40</sup>. Our high

rate of ventilator dependent patient at discharge was related to the high rate of patients with neurological disorders ( $p < 0.001$ ). Even in patients with underlying neurological disorders and / or ventilation dependency, decannulation could be done with NIV.

### *Neurological Problems*

As reported in previous studies, the most common underlying disorder in children with tracheostomy was neurological disorders as in our study<sup>5,6</sup>. Studies have found that neurologic diseases were associated with the inability to decannulate successfully<sup>39,41</sup>. In our study, patients with underlying neurological disorders or presence of a concomitant neurological comorbidities in addition to primary disease was associated with lower decannulation rates which was 25 %.

### *Noninvasive Ventilation need After Decannulation*

In previous studies NIV was used immediately following decannulation as a weaning tool (elective NIV) or in children who develops obstructive symptoms following decannulation (rescue NIV)<sup>29,42</sup>. Decannulation with NIV rates vary between 4-30 %<sup>29,42</sup>. Majority of patients requiring NIV following decannulation are patients with airway problems, congenital central hypoventilation syndrome and neurologic disorders<sup>43</sup>. In our study 7 of 59 (12%) decannulated patients need NIV after decannulation. Six of these patients had ND and one patient had severe tracheomalacia.

NIV was used for the increase of success rate in patients with upper airway obstruction during decannulation<sup>42</sup>, but there is no study reporting the need for NIV during decannulation of patients with ND. An explanation of the need for NIV in our study may be the tendency to upper airway obstruction due to hypotonia.

None of our patients decannulated with NIV required invasive ventilation during follow-up and one patient with tracheomalacia could be independent from NIV.

### *Decannulation Failure, Complications and Mortality*

In our study only 2 (3.3%) patients required tracheostomy replacement in 24-72 hours following decannulation due to decompensation of underlying cardiologic disease.

Decannulation failure rates vary according to the decannulation protocols <sup>26,44</sup>. At Wirtz et al's decannulation and observation protocol without any capping or downsizing which they evaluate patient's airway under sedation with spontaneous breathing, decannulation failure rate was % 6. Decannulation failure rates were between 0-13% in studies with capped PSG protocol <sup>44</sup>.

While signs of upper airway obstruction were prominent in the first 4 hours following decannulation, problems related to retention of secretions were more common after 24 hours <sup>20</sup>. To our knowledge, there is no study reporting the early complications after decannulation that not cause decannulation failure. Our complication rate was 29% and the most frequent complication was stridor that resolved within few hours.

Children with tracheostomy have higher mortality rates which may be related with underlying disease (12.5-20 %) and preventable complications of tracheostomy (1-3.2%) itself <sup>5,8,39</sup>. In our study 18 patients (11%) died, mortality was related to the underlying condition for all of these patients and none of our decannulated patients died during study period.

### ***Limitations***

The retrospective design based on medical records was the limitation of our study. Second limitation, we did not evaluate the patients with PSG before decannulation. But inpatient sleep monitorization that involve sleep and awake oxygen monitorization, morning blood gas analysis for detecting nocturnal hypoventilation were performed to all patients.

## **CONCLUSION**

In order to prevent tracheostomy related morbidity and mortality, decannulation, under safe circumstances, should be considered in all patients with tracheostomy as soon as possible. The lack of evidence-based decannulation guidelines make difficult to predict decannulation success. In our study, we showed for the first time in the literature that the presence of chronic colonization affects decannulation rates, as well as neurological disorders and the need for respiratory support after tracheostomy. While conducting decannulation assessment, the presence of colonization should be investigated and necessary measures should be taken. Future prospective researchs are necessary in order to develop evidence based decannulation guidelines and determine the role of chronic colonization on decannulation success.



## REFERENCES

1. Resen MS, Grønhøj C, Hjuler T. National changes in pediatric tracheotomy epidemiology during 36 years. *Eur Arch Otorhinolaryngol* 2018;275(3):803-808.
2. Grønhøj C, Charabi B, Buchwald CV, Hjuler T. Indications, risk of lower airway infection, and complications to pediatric tracheotomy: report from a tertiary referral center. *Acta Otolaryngol* 2017;137(8):868-871.
3. Berry JG, Graham DA, Graham RJ, Zhou J, Putney HL, O'Brien JE, et al. Predictors of clinical outcomes and hospital resource use of children after tracheotomy. *Pediatrics* 2009;124(2):563-572.
4. Zhu H, Das P, Roberson DW, Jang J, Skinner ML, Paine M, et al. Hospitalizations in children with preexisting tracheostomy: a national perspective. *Laryngoscope* 2015;125(2):462-468.
5. Pérez-Ruiz E, Caro P, Pérez-Frías J. et al. Paediatric patients with a tracheostomy: a multicentre epidemiological study. *Eur Respir J* 2012; 40 (6) 1502-1507.
6. Funamura JL, Durbin-Johnson B, Tollefson TT, Harrison J, Senders CW. Pediatric tracheotomy: indications and decannulation outcomes. *Laryngoscope* 2014;124(8):1952-8.
7. Pavone M, Verrillo E, Onofri A et al. Characteristics and outcomes in children on long-term mechanical ventilation: the experience of a pediatric tertiary center in Rome. *Ital J Pediatr* 2020;46(1):12.
8. de Trey L, Niedermann E, Ghelfi D, Gerber A, Gysin C. Pediatric tracheotomy: a 30-year experience. *J Pediatr Surg* 2013;48:1470–1475.
9. Schweiger C, Manica D, Lubianca Neto JF, Sekine L, Krumenauer R, Caixeta JA, Maunsell R, Gomes Avelino M. Determinants of successful tracheostomy decannulation in children: a multicentric cohort study. *J Laryngol Otol* 2020 Jan;134(1):63-67.

10. Al-Samri M, Mitchell I, Drummond DS, Bjornson C. Tracheostomy in children: a population-based experience over 17 years. *Pediatr Pulmonol* 2010;45:487–493.
11. Niederman MS, Ferranti RD, Zeigler A, Merrill WW, Reynolds HY. Respiratory infection complicating long-term tracheostomy. The implication of persistent gram-negative tracheobronchial colonization. *Chest* 1984; 85: 39–44.
12. Sanders CD, Guimbellot J, Muhlebach MS, Lin FC, Gilligan P, Esther CR Jr. Tracheostomy in children: Epidemiology and clinical outcomes. *Pediatr Pulmonol* 2018;53(9):1269–1275.
13. Morar P, Singh V, Jones AS, Hughes J, van Saene R. Impact of tracheotomy on colonization and infection of lower airways in children requiring long-term ventilation: a prospective observational cohort study. *Chest*. 1998;113:77–85.
14. Rao AR, Splaingard MS, Gershan WM, Havens PL, Thill A, Barbieri JT. Detection of *Pseudomonas aeruginosa* type III antibodies in children with tracheostomies. *Pediatr Pulmonol* 2005;39:402–407.
15. McCaleb R, Warren RH, Willis D, Maples HD, Bai S, O'Brien CE. Description of respiratory microbiology of children with long-Term tracheostomies. *Respir Care* 2016;61:447–452.
16. Seligman KL, Liming BJ, Smith RJH. Pediatric Tracheostomy Decannulation: 11-Year Experience. *Otolaryngol Head Neck Surg* 2019 Sep;161(3):499-506
17. Sherman JM, Davis S, Albamonte-Petrack S, Chatburn RL et al. Care of the child with a chronic tracheostomy. This official statement of the American Thoracic Society was adopted by the ATS Board of Directors. July 1999. *Am J Respir Crit Care Med* 2000;161:297– 308.
18. Mitchell RB, Hussey HM, Setzen G. et al. Clinical consensus statement: tracheostomy care. *Otolaryngol Head Neck Surg* 2013;148:6–20.
19. Heffner JE. The technique of weaning from tracheostomy. Criteria for weaning; practical measures to prevent failure. *J Crit Illn* 1995;10:729–33.
20. Stelfox HT, Crimi C, Berra L, Noto A, Schmidt U, Bigatello LM, Hess D. Determinants of tracheostomy decannulation: an international survey. *Crit Care* 2008;12:R26.

21. Takahashi N, Takano K, Mitsuzawa M et al. Factors associated with successful decannulation in pediatric tracheostomy patients. *Acta Oto-Laryngologica* 137:10. 1104-1109.
22. Garrubba M, Turner T, Grieveson C. Multidisciplinary care for tracheostomy patients: a systematic review. *Crit Care* 2009;13:R177.
23. Pozzi M, Pellegrino P, Galbiati S. et al. Prevalence of respiratory colonisations and related antibiotic resistances among paediatric tracheostomised patients of a long-term rehabilitation centre in Italy. *Eur J Clin Microbiol Infect Dis* 2015;34:169–175.
24. Pozzi M, Galbiati S, Locatelli F, Clementi E, Strazzer S. Performance of a tracheostomy removal protocol for pediatric patients in rehabilitation after acquired brain injury: Factors associated with timing and possibility of decannulation. *Pediatr Pulmonol* 2017;52(11):1509-1517.
25. Gergin O, Adil EA, Kawai K, Watters K, Moritz E, Rahbar R. Indications of pediatric tracheostomy over the last 30 years: Has anything changed? *International Journal of Pediatric Otorhinolaryngology* 2016;87:144-147.
26. Wirtz N, Tibesar RJ, Lander T et al. A Pediatric Decannulation Protocol: Outcomes of a 10-Year Experience. *Otolaryngology–Head and Neck Surgery* 2016;154:731–734.
27. Sachdev A, Ghimiri A, Gupta N, Gupta D. Pre-decannulation flexible bronchoscopy in tracheostomized children. *Pediatr Surg Int.* 2017;33(11):1195-1200.
28. Prescott CAJ. Peristomal complications of paediatric tracheostomy. *Int J Pediatr Otorhinolaryngol* 1992;23:141–149
29. Lee JH, Smith PB, Quek MBH, Laughon MM, Clark RH, Hornik CP. Risk factors and in-hospital outcomes following tracheostomy in infants. *J Pediatr* 2016;173:39-44.e31.
30. Namachivayam P, Shann F, Shekerdemian L, et al. Three decades of pediatric intensive care: who was admitted, what happened in intensive care, and what happened afterward. *Pediatr Crit Care Medicine* 2010;11:549-555.
31. Corbett HJ, Mann KS, Mitra I, Jesudason EC, Losty PD, Clarke RW. Tracheostomy—a 10-year experience from a UK pediatric surgical center. *J Pediatr Surg* 2007;42:1251-1254.

32. Lepointeur M, Ogna A, Clair B, Dinh A, Tarragon C, Prigent H, Davido B, Barbot F, Vaugier I, Afif M, et al. Risk factors for respiratory tract bacterial colonization in adults with neuromuscular or neurological disorders and chronic tracheostomy. *Respir Med* 2019;152:32-36.
33. McCrea N. et al. *Arch Dis Child Educ Pract Ed* 2013;98:84–91.
34. Gerdung CA, Tsang A, Yasseen AS, et al. Association Between Chronic Aspiration and Chronic Airway Infection with *Pseudomonas aeruginosa* and Other Gram-Negative Bacteria in Children with Cerebral Palsy. *Lung* 2016;194:307–314.
35. Tan CY, Chiu NC, Lee KS, et al. Respiratory tract infections in children with tracheostomy. *Journal of Microbiology. Immunology and Infection* 2020;53:315-320.
36. Chang AB, Bush A, Grimwood K. Bronchiectasis in children: diagnosis and treatment. *Lancet* 2018;392:866-879.
37. Bandyopadhyay A, Cristea AI, Davis S, et al. Retrospective Analysis of Factors Leading to Pediatric Tracheostomy Decannulation Failure. A Single-Institution Experience. *Ann Am Thorac Soc* 2017;14: 70–75.
38. Arslan H, Unal F, Teber BG, et al. The effects of nebulized antibiotics in children with tracheostomy. Paper presented at 23. Turkish Thoracic Society International Congress. 2020;EPS-049:224.[https://kongre2020.toraks.org.tr/wp-content/uploads/2020/11/23.-TORAKS-2020\\_L-v6.pdf](https://kongre2020.toraks.org.tr/wp-content/uploads/2020/11/23.-TORAKS-2020_L-v6.pdf).
39. Dursun O, Ozel D. Early and long-term outcome after tracheostomy in children. *Pediatrics International* 2011;53: 202-206.
40. Chia AZH, Ng ZM, Pang YX, Ang AHC, Chow CCT, Teoh OH, Lee JH. Epidemiology of Pediatric Tracheostomy and Risk Factors for Poor Outcomes: An 11-Year Single-Center Experience. *Otolaryngol Head Neck Surg* 2020;162(1):121-128.
41. McPherson ML, Shekerdemian L, Goldsworthy M, et al. A decade of pediatric tracheostomies: indications, outcomes, and long-term prognosis. *Pediatr Pulmonol*

2017;52:946-953.

42. Fauroux B, Leboulanger N, Roger G, et al. Noninvasive positive-pressure ventilation avoids recannulation and facilitates early weaning from tracheotomy in children. *Pediatr Crit Care Med* 2010;11:31–7.
43. Kam K, Bjornson C, Mitchell I. Congenital central hypoventilation syndrome; safety of early transition to non-invasive ventilation. *Pediatr Pulmonol* 2014;49(4):410-413.
44. Robison JG, Thottam PJ, Greenberg LL, Maguire RC, Simons JP, Mehta DK. Role of polysomnography in the development of an algorithm for planning tracheostomy decannulation. *Otolaryngol Head Neck Surg* 2015;152:180-184.