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Causes and Consequences of Adult Laryngotracheal Stenosis

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Abstract

Objective—Laryngotracheal stenosis is largely considered a structural entity, defined on anatomic terms (i.e. percent stenosis, distance from vocal folds, overall length). This has significant implications for identifying at-risk populations, devising systems-based preventive strategies, and promoting patient-centered treatment. The present study was undertaken to test the hypothesis that LTS is heterogeneous in regard to etiology, natural history, and clinical outcome.

Study Design—Retrospective cohort study of consecutive adult tracheal stenosis patients from 1998–2013.

Methods—Subjects diagnosed with laryngotracheal stenosis (ICD-9: 478.74, 519.19) between January 1, 1998 and January 1, 2013 were identified. Patient characteristics (age, gender, race, follow-up duration), and comorbidities were extracted. Records were reviewed for etiology of stenosis, treatment approach, and surgical dates. Stenosis morphology was derived from intraoperative measurements. The presence of tracheostomy at last follow-up was recorded.

Results—150 patients met inclusion criteria. 54.7% had an iatrogenic etiology followed by idiopathic (18.5%), autoimmune (18.5%), and traumatic (8%). Tracheostomy dependence differed

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based on etiology ($p < 0.001$). Significantly more patients with iatrogenic (66%) and autoimmune (54%) etiologies remained tracheostomy dependent compared to traumatic (33%) or idiopathic (0%) groups. On multivariate regression analysis, each additional point on Charlson Comorbidity Index was associated with a 67% increased odds of tracheostomy dependence (OR 1.67, 95% CI 1.04 – 2.69; $p = 0.04$).

Conclusions—Laryngotracheal stenosis is not a homogeneous clinical entity. It has multiple distinct etiologies that demonstrate disparate rates of long-term tracheostomy dependence. Understanding the mechanism of injury and contribution of comorbid illnesses is critical to systems-based preventive strategies and patient-centered treatment.

Keywords

Tracheal Stenosis; Subglottic Stenosis; Laryngotracheal Stenosis; Intubation; Tracheostomy

INTRODUCTION

Laryngotracheal stenosis (LTS) is a life threatening, fixed, extrathoracic restriction in pulmonary ventilation. LTS is an umbrella term, encompassing luminal compromise at the level of the larynx, subglottis or trachea, which exists in a watershed of specialty care. Diagnosis is frequently delayed as patients rapidly transition from acute inpatient care to outpatient facilities. The majority of patients are not definitively diagnosed until outpatient specialty evaluation¹. Many specialists (e.g. intensivists, otolaryngologists, interventional pulmonologists, thoracic surgeons) initially interact with this population. This makes it difficult to establish the natural history of the disease, define universal predictors of disease outcome and create cogent personalized plans of care. Additionally, long-term sequelae of intensive respiratory support (endotracheal intubation and elective tracheostomy) do not develop on a timescale necessary for recognition by practitioners providing acute care, impeding quality-driven improvement efforts².

Moreover, LTS is generally described in terms of its structural characteristics, defined on anatomic terms (i.e. percent stenosis, distance from vocal folds, overall length). This neglects the unique biology driving luminal compromise in heterogeneous patient populations and has significant implications for identifying at-risk populations, devising systems-based preventive strategies, and promoting patient-centered treatment directed at the diverse pathophysiology driving airway injury. The present study was undertaken to test the hypothesis that LTS is heterogeneous in regard to etiology, natural history, and clinical outcome.

PATIENTS & METHODS

This study was performed in accordance with the Declaration of Helsinki, Good Clinical Practice, and was approved by the Baylor College of Medicine Institutional Review Board (IRB No. H33195).

Patients

Subjects diagnosed with laryngotracheal stenosis (ICD-9: 478.74, 519.19) between January 1, 1998 and January 1, 2013 were identified. Those with a history of tracheal malignancy or isolated laryngeal stenosis were excluded. Laryngeal and tracheal stenosis share an association with both prolonged endotracheal intubation, and many of the same comorbid medical risk factors. However, isolated laryngeal stenosis remains a distinct anatomic and structural injury with a unique treatment algorithm, meriting dedicated independent study, and is not discussed in the present work. Patients meeting inclusion were grouped into 4 categories based on stenosis etiology: idiopathic, iatrogenic, autoimmune, and polytrauma (Table 1).

Idiopathic: No history of significant laryngotracheal injury. No significant history of endotracheal intubation or tracheotomy within 2 years of the presentation. No thyroid or major anterior neck surgery. No neck irradiation. No caustic or thermal injuries to the laryngotracheal complex. No history of vasculitis. Negative titers for angiotensin-converting enzyme (ACE) and antinuclear cytoplasmic antibody (ANCA). The lesion must involve the subglottis.

Iatrogenic: Patients that developed subglottic or tracheal stenosis following tracheostomy. Or subglottic or tracheal stenosis developing within 2 years of intubation.

Autoimmune: Patients with documented clinical along with serologic and/or histologic diagnosis of Wegener's (GPA), Relapsing Polychondritis (RPC), Systemic Lupus Erythematosus (SLE), Rheumatoid Arthritis (RA), Epidermolysis Bullosa (EB), Sarcoidosis, or Amyloidosis

Polytrauma: Patients presenting with Laryngotracheal stenosis following documented traumatic injuries involving multiple organ systems.

Data Collected

Patient characteristics (age, gender, race, follow-up duration), and comorbidities were extracted. Records were reviewed for etiology of stenosis, treatment approach (i.e. endoscopic, open), and surgical dates. Stenosis morphology [% luminal obstruction, distance from glottis (cm), and overall length (cm)] and tracheomalacia were derived from intraoperative findings. Patients were staged with the established Cotton-Myer, Lano, & McCaffrey classification systems as previously described³⁻⁵ (Table 2). The number and frequency between repeat procedures were captured.

Procedures

Treatments for tracheal stenosis included: 1) endoscopic dilations of the stenotic trachea⁶, 2) open surgical resection of the diseased tracheal segment with end-to-end anastomosis⁷, and 3) permanent tracheostomy. The treatment algorithm consisted of initial endoscopic dilation for all patients. In patients who required multiple dilation procedures rigorous selection criteria were applied for consideration of open surgical reconstruction. Patients less than 45 years old, without type II diabetes or connective tissue disease, and with stenosis 2cm or

more below the glottis and less than 2cm in length were offered open surgical reconstruction.

Outcomes

Presence of a tracheostomy at last follow-up was the primary outcome. This represented failure of surgical management to correct airway narrowing.

Statistical Analysis

All data management and analysis were done using STATA/MP version 12.1 software (STATA Corp, College Station, Texas). Univariate analyses were performed using ANOVA, Pearson's chi-squared tests, and Fisher's exact tests as appropriate. Stepwise multivariate logistic regression analysis was used to identify independent risk factors for tracheostomy. A significance level of $p < 0.20$ on univariate analysis was used as the criterion for inclusion in multivariate model. As convention, $p < 0.05$ was required for statistical significance in the model.

RESULTS

A total of 340 patients with a diagnosis of tracheal or laryngeal stenosis were identified. Excluded were those with tracheal malignancy ($N=9$), isolated bilateral vocal fold immobility ($N=181$). In all, 150 patients met inclusion criteria. The most common etiology was iatrogenic (54.7%) followed by idiopathic (18.5%), autoimmune (18.5%), and traumatic (8%; Table 3). Mean follow-up was 39.3 months (95% Confidence Interval (CI) 31.9 – 46.6), but varied significantly by etiology ($p < 0.001$; Table 3).

Univariate Analysis

Patient Characteristics—Age at presentation differed significantly by strata ($p=0.002$) with those in the traumatic group being significantly younger than all others (34.4 years, CI 23.5 – 45.3; Table 3). Gender distribution also differed based on etiology ($p < 0.002$; Table 3). In order, the idiopathic group had a significantly higher percentage of females (93%) than autoimmune (68%) iatrogenic (62%), or traumatic (33%) LTS patients. Charlson Comorbidity Index (CCI) varied between groups ($p < 0.001$). Iatrogenic and autoimmune strata had significantly higher indices than either idiopathic or traumatic strata (Table 3).

Examination of the individual components of the CCI showed cardiovascular comorbidities (i.e. myocardial infarction, congestive heart failure, peripheral vascular disease, and cerebrovascular disease) and type II diabetes were significantly more prevalent in the iatrogenic strata than in other etiologies (Figure 1A and Table 3). There were not significant differences in the rate of gastroesophageal reflux disease (GERD) between strata (Table 3).

Disease Morphology—Degree of stenosis differed between etiologic strata ($p=0.01$). Idiopathic LTS involved less of the tracheal lumen (mean 57%; CI 52 – 63%) than those in the autoimmune or iatrogenic groups (Table 3). There were no differences in the mean distance from the glottis ($p=0.11$) or the length of stenoses between strata ($p=0.44$). In the iatrogenic group, LTS occurred in the subglottis (1.5cm from the glottis) in 59% of patients

(49/82) (Figure 1B). Even in those patients presenting with iatrogenic LTS following tracheostomy, 41% (16/39) had subglottic injuries on intra-operative examination.

Treatment—There was no difference in number of surgeries per year of follow-up ($p=0.49$) or the types of surgeries performed by etiologic strata ($p=0.14$; Table 3). Most patients were treated with tracheal dilation (84%), followed by T-tube placement (8%), resection (6%), and no treatment (2%).

Tracheostomy Dependence—Tracheostomy dependence differed based on etiologic strata ($p<0.001$) (Figure 1C). Significantly more patients in the iatrogenic (66%) and autoimmune (54%) groups were tracheostomy dependent at last follow-up compared to those in either the traumatic (33%) or idiopathic (0%) groups. Tracheostomy dependence also differed based on established staging systems (Figure 2A). When stratified via Cotton-Myer staging (based on the degree of luminal obstruction), significantly more patients with grade III (90%) and grade IV (90%) lesions were tracheostomy dependent at last follow-up compared to those in either the grade II (38%) or grade I (32%) groups. ($p<0.001$; Figure 2A). When stratified by the Lano classification (based on the stenosis location), increasing subsite involvement was significantly associated with a higher rate of tracheostomy ($p<0.001$; Figure 2A). When staged according to the McCaffrey classification system (based on both stenosis location and length), increased stage was associated with progressively increased risk of tracheostomy ($p<0.01$) (Figure 2A).

All 3 established adult LTS staging systems accurately stratified patients' outcomes based on the severity of their structural injury. Overall (consistent with prior reports), patients in our series with more severe luminal compromise, those with longer stenosis, and those with lesions spanning multiple subsites (glottis, subglottis &/or trachea) had a much higher incidence of tracheostomy. However, this observation did not hold when patients were stratified by etiology of injury (Table 4.) No patients in the idiopathic group required tracheostomy (even those with lengthy, severe stenosis, involving multiple subsites). Conversely, patients with iatrogenic injuries had a significantly higher rate of tracheostomy even when matched at lower stenosis grades when compared with the other etiologic strata. The non-uniform rate of tracheostomy observed in different etiologic groups was seen in all three established LTS classification systems (Table 4).

Tracheal Structural Instability—Patients with iatrogenic injuries had a significantly higher rate of tracheomalacia seen on bronchoscopic evaluation (37% vs. 8%, $p<0.001$; Figure 2B). Given the retrospective nature of this work, it is not possible to establish a causative relationship between the initial injury and the loss of structural integrity associated with tracheomalacia. However, it is interesting that among the iatrogenic group, 45% of patients without malacia required tracheostomy, while 97% of those with malacia necessitated long-term tracheostomy ($p<0.001$; Figure 2C).

Multivariate Analysis

Multivariate regression analysis was performed to determine independent predictors of ultimate tracheostomy dependence. Each additional point on CCI was associated with a 67%

increased odds of tracheostomy dependence (OR 1.67, 95% CI 1.04 – 2.69; $p=0.04$). Moreover, there was a 3% increased odds of tracheostomy dependence for each additional percentage of airway compromise (OR 1.03, 95% CI 1.01 – 1.06; $p=0.001$). LTS patient characteristics (etiology, age, sex, race) were not significantly associated with odds of tracheostomy dependency.

DISCUSSION

Although most airway stenosis appears similar on anatomic imaging and clinical examination, we present data supporting the hypothesis that different mechanisms of injury are associated with differing rates of long-term tracheostomy dependence. The relationships between the anatomic stenosis characteristics (% stenosis, location, & length) and endoscopic or open surgical “success” have been established through pioneering work in both children^{8,9} and adults⁷. In advanced centers, procedural intervention for LTS offers a high rate of long-term tracheostomy free survival^{4,10,11}. However, success in these large published series remains critically dependent on patient selection. With our consecutive series of both inpatient and outpatient consultations, we believe this study captured a more representative cross-section of symptomatic LTS patients than many prior adult surgical case series. In the “real world”, those patients deemed poor operative candidates (e.g. sicker patients) are often left with limited therapeutic options regardless of the structural morphology of their stenosis.

Endotracheal intubation and tracheostomy can be lifesaving, but should not be considered benign procedures. They harbor significant long-term risks to communication¹², swallowing¹³, and breathing¹⁴, particularly in the subset of patients with comorbid illness¹⁵. Ironically, this is also the population that more frequently requires intensive respiratory support. In our series each additional point on CCI was associated with a 67% increased odds of tracheostomy dependence. Although this association doesn’t appear surprising, we believe it is powerful. It demonstrates the suitability of the CCI to serve as a systems-based protocol to identify patients who mandate a heightened awareness of complication from these procedures.

Consistent with previously published series^{4,16,17}, despite many risk factors for iatrogenic injury being clarified over the past 40 years^{15,18–20}, more than half the LTS burden in our cohort was potentially preventable. Overall, 59% of iatrogenic injuries occurred within the subglottis and, therefore, are attributable to intubation. In a *post hoc* analysis, 83% (15/18) “healthy” patients (those without DMII or cardiovascular disease) with iatrogenic LTS were women. This previously reported observation²¹ suggests that endotracheal tube size may contribute to tracheal injury and should be carefully considered in the smaller female trachea²².

As has been consistently shown across other large series¹⁵, patients with DMII are particularly vulnerable to airway injury and have a higher likelihood of long-term tracheostomy dependence when injury occurs. Interestingly, the rate of gastroesophageal reflux disease (GERD) was not significantly different between the etiologic subgroups. Although other investigators have suggested a tight relationship between GERD and adult

idiopathic LTS, this was not seen in our patient population. The limits of retrospective review prevent us from direct comparison of the objective data on the frequency and severity of reflux episodes between individuals and subgroups. Increased body mass index (BMI) also has a suggested association with increased risk of tracheal injury with intubation and worse response to procedural intervention. Our series lacked the biometric data to address this concept. Additionally, the limits of our tertiary care referral center (with limited out-of-network medical records) prevented us from exploring the relationship between the length of intubation, or type of tracheostomy procedure (open vs. percutaneous) and the ultimate injury severity or treatment outcome.

A strong association between the degree of stenosis and ultimate decannulation has previously been reported in children²³. Our series supports these prior observations in the pediatric population and now extends them to adults. As previously reported in adults, the location of injury and the length of stenosis are also essential components to predict long-term tracheostomy dependence. Critically, we now also offer data supporting an additional relationship between the cause of upper airway injury and its ultimate response to therapy. This relationship had been assumed; we offer the first formal demonstration.

Anatomic staging systems are numerous^{3-5,24-28} yet, the ideal system in adult LTS remains unresolved. The most established allow some degree of prognosis, promote individualized treatment planning, and facilitate multi-institutional comparison. In this work we utilized three separate, established LTS classification systems. As expected, they all effectively stratify patient's risk of long-term tracheostomy. However, interestingly, in adult LTS it appears the McCaffrey and Lano systems offer more precision than does the Cotton-Myer scale.

While in general, patients in our series with more severe luminal compromise, longer stenoses, and lesions spanning multiple subsites had a much higher incidence of tracheostomy, this observation did not hold in the idiopathic group (whom never required tracheostomy), suggesting a unique injury. Conversely, while lower LTS stages (in all 3 systems) overall had a lower rate of tracheostomy, patients with iatrogenic injuries had a significantly higher rate even when matched at lower stenosis grades (identically in all 3 systems). Grouping LTS patients solely by an anatomic classification of their injury neglects a critical component of the heterogeneous biology responsible for tracheal scar. Patients with iatrogenic stenosis appear to possess unique medical comorbidities. Their disease ultimately behaves differently as evidenced by their disparate rate of long-term tracheostomy dependence even when matched for similar degree of luminal compromise. These separate subgroups likely merit tailored treatment strategies.

The finding of the high rate of tracheomalacia in the subgroup with iatrogenic injuries, and the significant association between tracheomalacia and long-term tracheostomy dependence in this subgroup, raises questions regarding the relative contributions of mucosal injury and cartilaginous injury in LTS. Ultimately, we believe the degree of tracheal wall injury (what we term "superstructure instability") may have significant prognostic power for overall response to therapy. However, this is difficult to quantify at present with our current diagnostic modalities and remains an area of active research.

Our study represents one of the largest published adult LTS series in the scientific literature. The data supports the hypothesis that laryngotracheal stenosis is a common endpoint to multiple pathophysiologic processes. While different mechanisms of airway injury physiologically affect the patient in similar ways, we show they occur in unique populations and have divergent responses to therapy. Management and prevention strategies should carefully consider this heterogeneous pathophysiology. This difference is not reflected in staging systems limiting themselves to an anatomic description of the tracheal scar.

CONCLUSION

Relief through endoscopic dilation, or open tracheal resection is attainable in some cases of LTS; however, treatment is not universally successful. It is incumbent on the scientific community to move beyond viewing LTS as a purely anatomic problem, remedied only through surgical reconstruction. Rather, the management of airway stenosis should transition from reactive surgical intervention to prevention through early recognition¹ of at-risk populations and increasingly personalized care through careful consideration of the pathophysiology driving airway injury in the unique subgroups affected with LTS.

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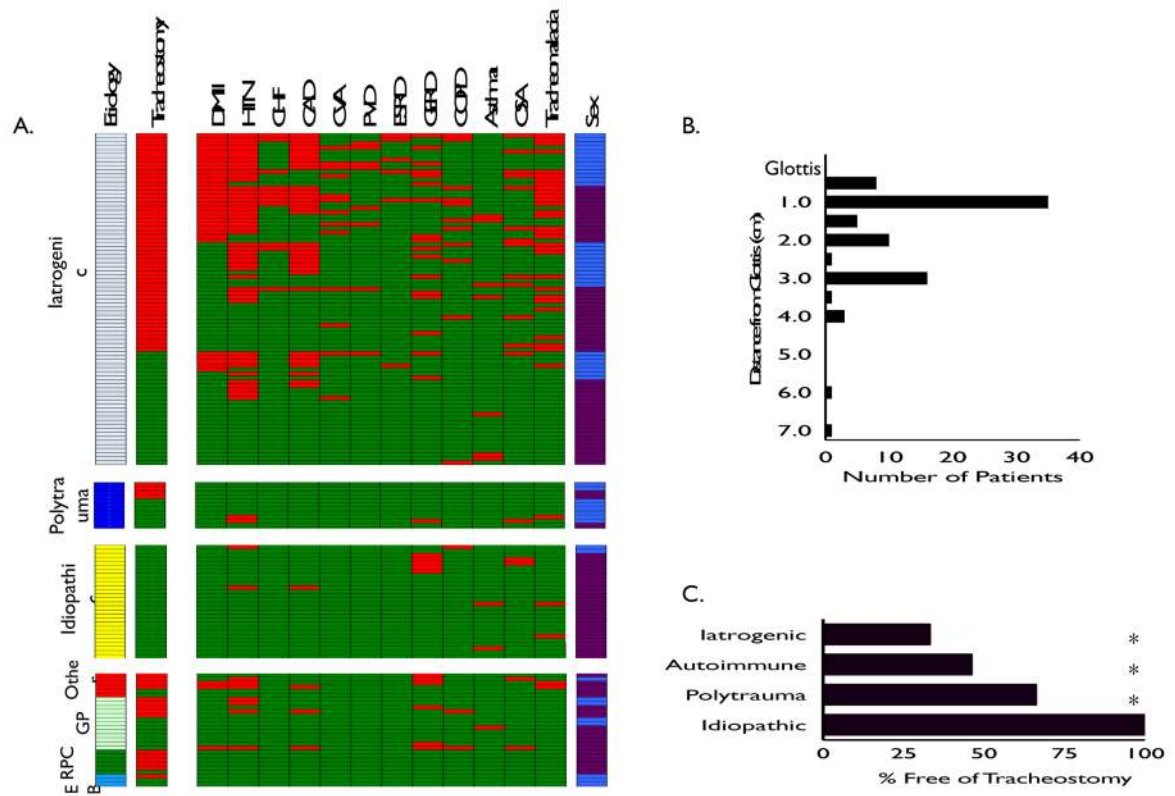


Figure 1. Heatmap grouped by different etiologies of stenosis. Each line represents an individual patient. Tracheostomy status (red indicating trach), medical comorbidities (presence highlighted in red), and sex (blue indicating male, purple indicating female). In Autoimmune subgroup: GPA (granulomatosis with polyangiitis, i.e. Wegener’s granulomatosis), RPC (relapsing polychondritis), EB (epidermolysis bullosa) (A.). Location of tracheal stenosis in iatrogenic injuries. Histogram showing location of stenotic lesion in iatrogenic subgroup in relation to distance from glottis (B.). Tracheostomy status of different etiologies at last f/u. Asterisk denotes statistical significance from idiopathic group (C.)

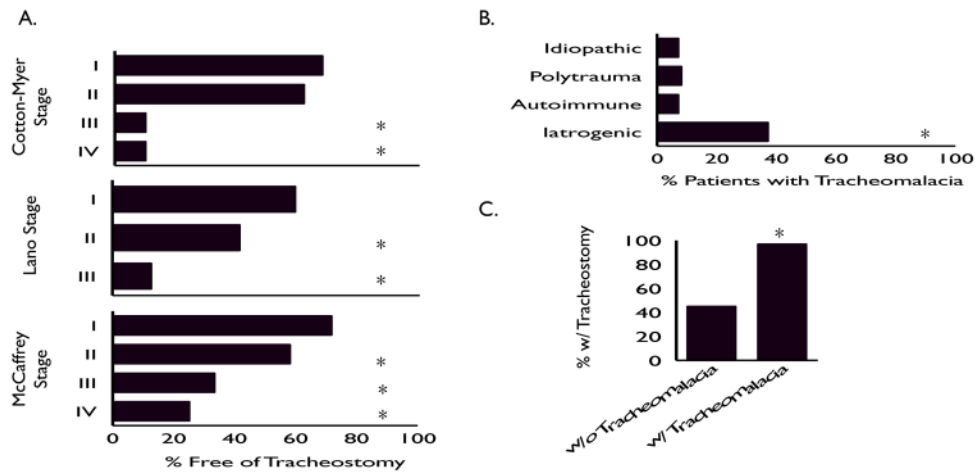


Figure 2. Tracheostomy status of different Cotton-Myer, Lano, & McCaffrey grades at last f/u. For Cotton-Myer staging, asterisk denotes statistical significance between Grade I&II vs. III&IV (A.). Diagnosis of tracheomalacia stratified by etiology. Asterisk denotes statistical significance between iatrogenic etiology and all other groups (B.). Rate of tracheostomy in iatrogenic etiology patients with and without a diagnosis of tracheomalacia. Asterisk denotes statistical significance (C.).

Table 1

Definitions of laryngotracheal stenosis etiology of injury utilized in this study.

Idiopathic	No history of significant laryngotracheal injury. No significant history of endotracheal intubation or tracheotomy within 2 years of the presentation. No thyroid or major anterior neck surgery. No neck irradiation. No caustic or thermal injuries to the laryngotracheal complex. No history of vasculitis. Negative titres for angiotensin-converting enzyme (ACE) and antinuclear cytoplasmic antibody (ANCA). The lesion must involve the subglottis.
Autoimmune	Patients with documented clinical along with serologic and/or histologic diagnosis of Wegeners (GPA), Relapsing Polychondritis (RPC), Systemic Lupus Erythematosus (SLE), Rheumatoid Arthritis (RA), Epidermolysis Bullosa (EB), Sarcoidosis, or Amyloidosis
Polytrauma	Patients presenting with Laryngotracheal stenosis following documented traumatic injuries involving multiple organ systems.
Iatrogenic	Patients that developed subglottic or tracheal stenosis following tracheostomy. Or subglottic or tracheal stenosis developing within 2 years of intubation.

Table 2

Definitions of Clinical Laryngotracheal Stenosis Classification Systems.

Cotton-Myer	I	< 50% obstruction
	II	51–70% obstruction
	III	71–99% obstruction
	IV	Complete obstruction
Lano	I	One subsite involvement
	II	Two subsite involvement
	III	Three subsite involvement
		* subsites defined as glottis, subglottis, & trachea
McCaffrey	I	Subglottis or trachea < 1cm
	II	Subglottis > 1cm
	III	Subglottis & trachea > 1cm
	IV	Any lesion involving glottis

Table 3
Demographics, Stenosis characteristics, Comorbidities, and Treatment of LTS grouped by etiology of injury.

	Idiopathic (n=28)	Polytrauma (n=12)	Autoimmune (n=28)	Introgenic (n=82)	Significance (p)
<i>Demographics</i>					
Follow up (Mean months, 95% CI)	56.07 (41.5 – 70.6)	12.3 (7.2 – 17.5)	69.1 (39.7 – 98.6)	27.05 (20.9 – 33.1)	<0.001
Age (Mean years, 95% CI)	50.35 (45.9 – 54.8)	35.7 (24.1 – 47.4)	45.1 (39.7 – 50.4)	51 (48.0 – 54.7)	0.002
Sex (% Female)	93	33	68	62	0.002
<i>Race (%)</i>					
Caucasian	89	50	71	63	
African American	7	17	14	16	0.330
Asian	0	8	0	2	
Hispanic	4	17	14	17	
<i>Disease Morphology</i>					
% Stenosis (Mean %, 95% CI)	57.86 (52.3 – 63.4)	69.6 (55.1 – 84.1)	68.5 (60.6 – 76.4)	72.8 (68.1 – 77.6)	0.010
Distance below glottis (Mean cm, 95% CI)	1.289 (1.0 – 1.6)	2.17 (1.29 – 3.05)	1.94 (1.38 – 2.51)	1.77 (1.5 – 2.02)	0.110
Stenosis Length (Mean cm, 95% CI)	1.657 (1.3 – 1.99)	1.95 (0.99 – 2.91)	2.12 (1.62 – 2.62)	2.167 (1.91 – 2.42)	0.440
<i>Comorbidities</i>					
Charlson Index (Mean, 95% CI)	0.07 (0 – 0.16)	0.00 (0)	1.28 (0.99 – 1.58)	1.32 (0.94 – 1.7)	<0.001
DMII (%)	0	0	11	39	<0.001
MI (%)	0	0	3.6	28	<0.001
CHF (%)	0	0	0	13	0.027
CVA (%)	0	0	0	7	0.008
COPD (%)	4	0	7	13	0.390
Connective tissue (%)	0	0	100	0	<0.001
* GERD (%)	18	8	21	20	0.859
<i>Treatment</i>					
# of Procedures / year (Mean, 95% CI)	1.75 (0.8 – 2.6)	3.41 (1.6 – 5.2)	1.8 (0.9 – 2.7)	2.65 (1.7 – 3.6)	0.490

Percentage of LTS patients with tracheostomy by Cotton-Myer, Lano, & McCaffrey stage, grouped by etiology of injury (n/a refers to an absence of patients within a given stage).

Table 4

	Cotton-Myer			
	I	II	III	IV
Idiopathic	0	0	0	n/a
Polytrauma	0	25	100	100
Autoimmune	36	50	100	100
Iatrogenic	57	44	92	88

	Lano		
	I	II	III
Idiopathic	0	0	n/a
Polytrauma	27	100	n/a
Autoimmune	50	42	83
Iatrogenic	60	75	100

	McCaffrey			
	I	II	III	IV
Idiopathic	0	0	0	0
Polytrauma	0	60	50	n/a
Autoimmune	63	50	50	n/a
Iatrogenic	36	65	82	80