

A Commonsense Approach to Pediatric Aspiration:

With a focus on Clefts

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Objectives

- Creating a systematic approach
 - Define the population that you are best suited to manage
- Explore the diagnosis and management of LTEC
 - The good
 - The bad
 - The ugly

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Definitions

- Dysphagia
 - ANY disorder, impairment or variation from norm during the oral, pharyngeal, or esophageal phases of swallow
- Aspiration
 - Secretions, food, or liquid entering the larynx and falling below the level of the vocal cords
 - “Silent” when it does not elicit a cough reflex

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Who do you want to be?

- Consider how you brand yourself
 - Highlight unique aspects of care that can be provided by an otolaryngologist
- Consider the type of pathology that you want to attract
 - As ENT surgeons, we have a specific skill set
 - Slippery slope between being effective and overwhelmed

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Who do you want to be???

- Pediatric Therapeutic Surgical Dysphagia Clinic
“PTSD Clinic”
 - Surgical Treatment of Dysphagia Clinic
“STD Clinic”

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Fine Print: only unexplained aspiration on a swallow study, recurrent bronchialitis, pneumonias, or internal referral from an aerodigestive

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Algorithm for Triage

- When we started
 - Word quickly spread...
 - Many referrals for any dysphagia issue
 - Self referrals from families fed up with their child's feeding problem that was "still not fixed"
 - All of a sudden, every aspirating patient was sent to us to "rule out a cleft"
 - Wait times for new patients became unacceptable (by American standards)

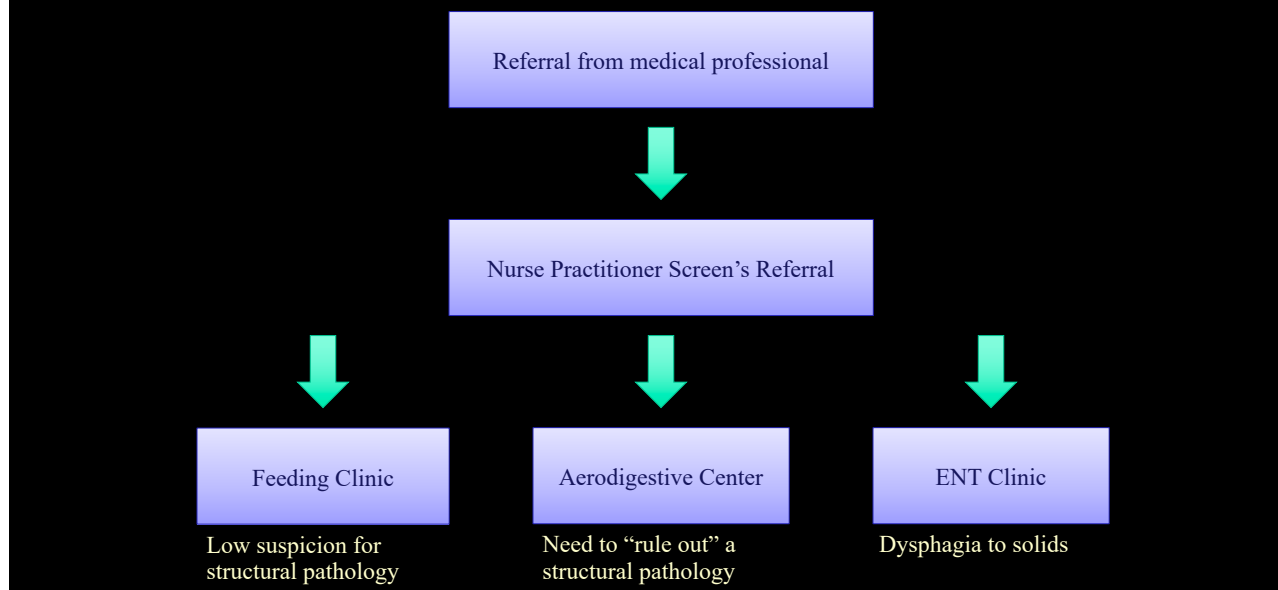
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Algorithm for Triage

- Our Aerodigestive Center
 - Six ENT Surgeons
 - We all see some dysphagia
 - One surgeon (me) who takes on the majority of complex cases
 - One multidisciplinary clinic/month
 - Maximum of 3 patients/hour
 - 20-25 patients per month
 - Each clinic slot is very valuable

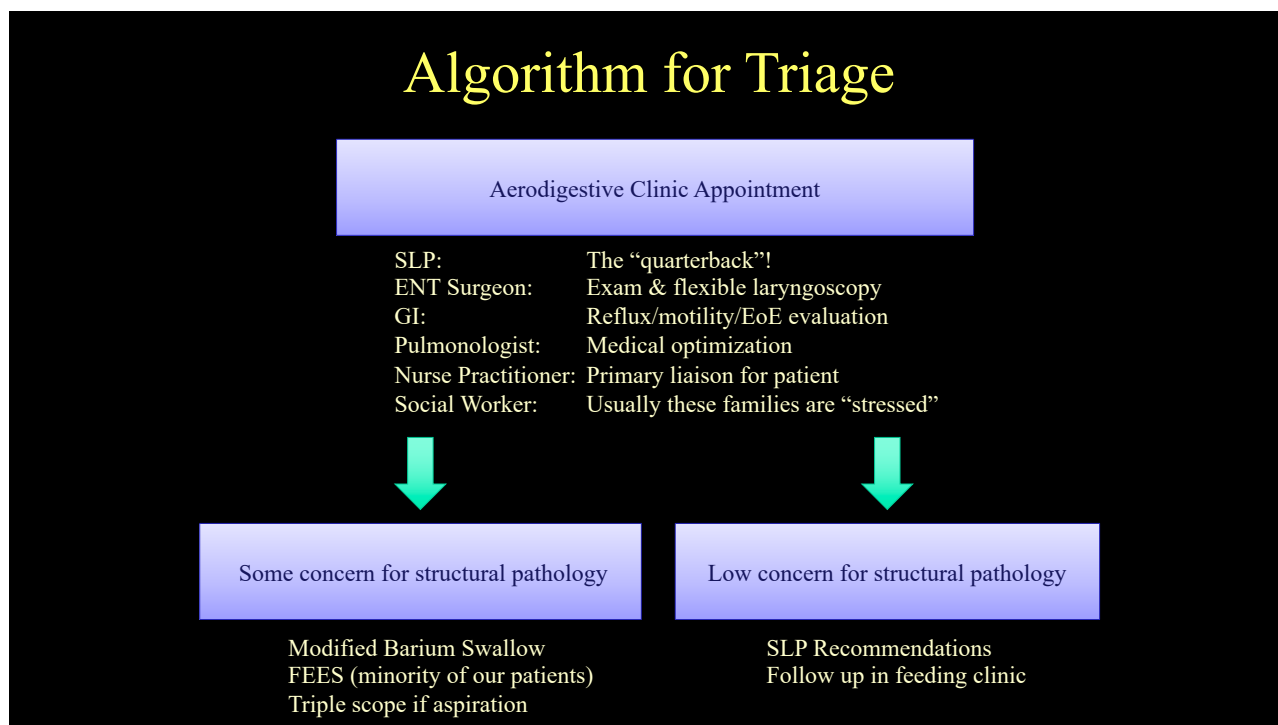
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Algorithm for Triage



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Algorithm for Triage



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Algorithm for Triage

- But this didn’t work....
 - Too many false negatives since screening did not include confirmed (or high suspicion for) aspiration
 - Too many “soft referrals” to “rule out a cleft”
- Current criteria to be seen
 - Confirmed unexplained aspiration
 - Recurrent unexplained respiratory illness
 - Genetic/Syndromic diagnosis with midline defects
 - Direct referral from aerodigestive provider

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Algorithm for Triage

Confirmed unexplained aspiration or recurrent respiratory illness, genetic predisposition for midline defects, referral from aerodigestive provider

SLP:	The “quarterback”!
ENT Surgeon:	Exam & ??? flexible laryngoscopy
GI:	Reflux/motility/EoE evaluation
Pulmonologist:	Medical optimization
Nurse Practitioner:	Primary liaison for patient
Social Worker:	Usually these families are “stressed”



Confirmed concern for structural pathology and optimized

Low concern for structural pathology or not optimized

TRIPLE SCOPE

Medical optimization and F/U
Possible repeat MBSS
Follow up in feeding clinic

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DO

- Collaborate with a medical team
 - Pre-intervention for optimization
 - Post-intervention for continued rehabilitation of the swallow
- Have a defined pathway that will guide surgical candidacy
 - Not every patient needs a “triple scope”

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DON'T

- Make your clinic a “free for all”
 - Your clinic is a limited and precious resource
- Take on pathologies better served in a medical home
 - Many (most) dysphagia patients do not have surgically-amenable pathology
 - Try to limit focus to “aspiration” patients

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DON'T

- ??? Flexible scope every patient
 - We used to do this (pre-COVID)
 - Doesn't add much for the vast majority
 - We look at dynamics anyhow at the time of triple scope
 - We are currently much more selective about doing office scopes

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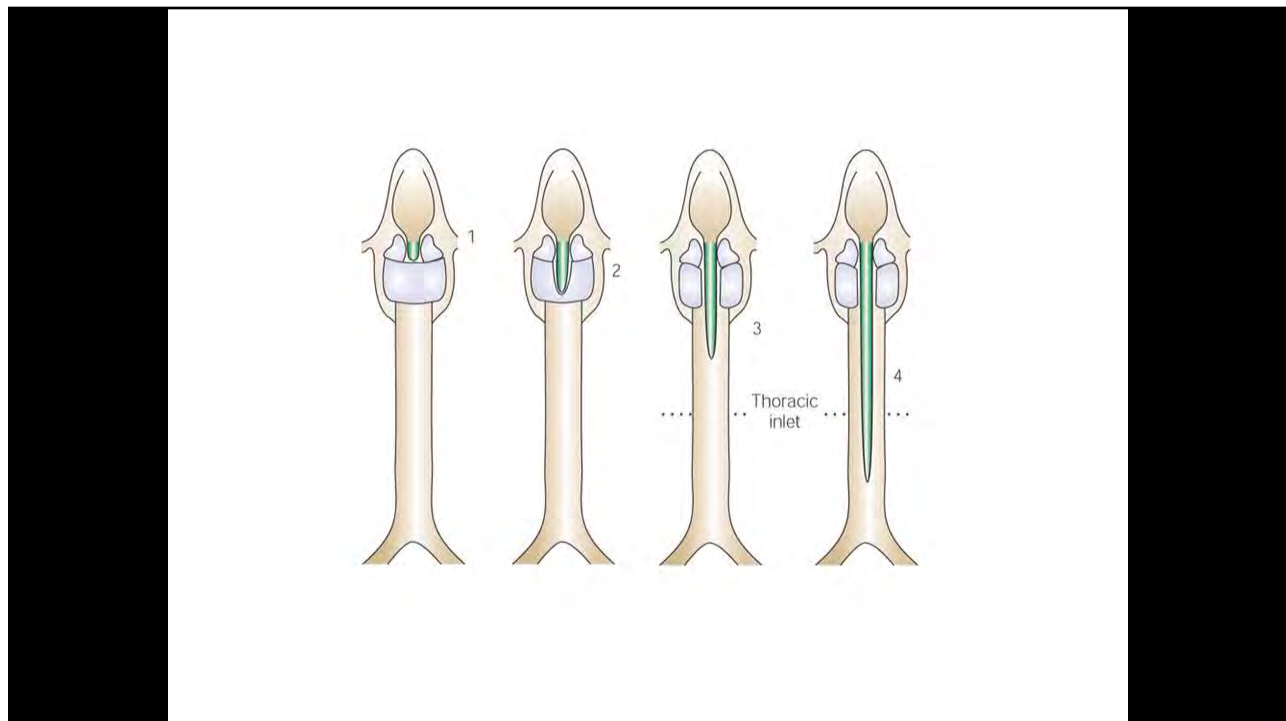
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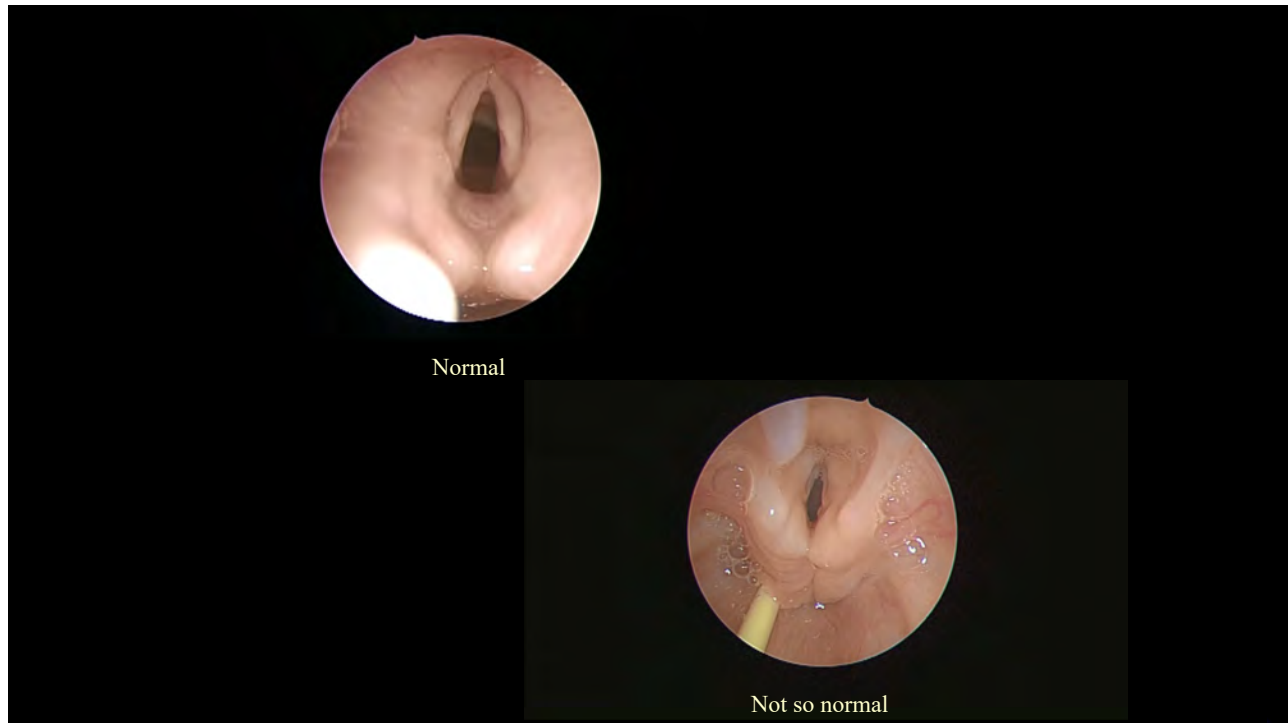
Laryngotracheoesophagealclefts

- Presentation
 - Type I/II: Cough with feeds, Feeding issues, URI symptoms, recurrent pneumonias
 - Type III: Above +/- Stridor, raspy cry, frank distress, completely unable to PO feed
 - Type IV: Distress++, Difficulty maintaining intubation
 - Other midline anomalies or genetic syndrome
 - Associated TEF in about 1/3 of patients

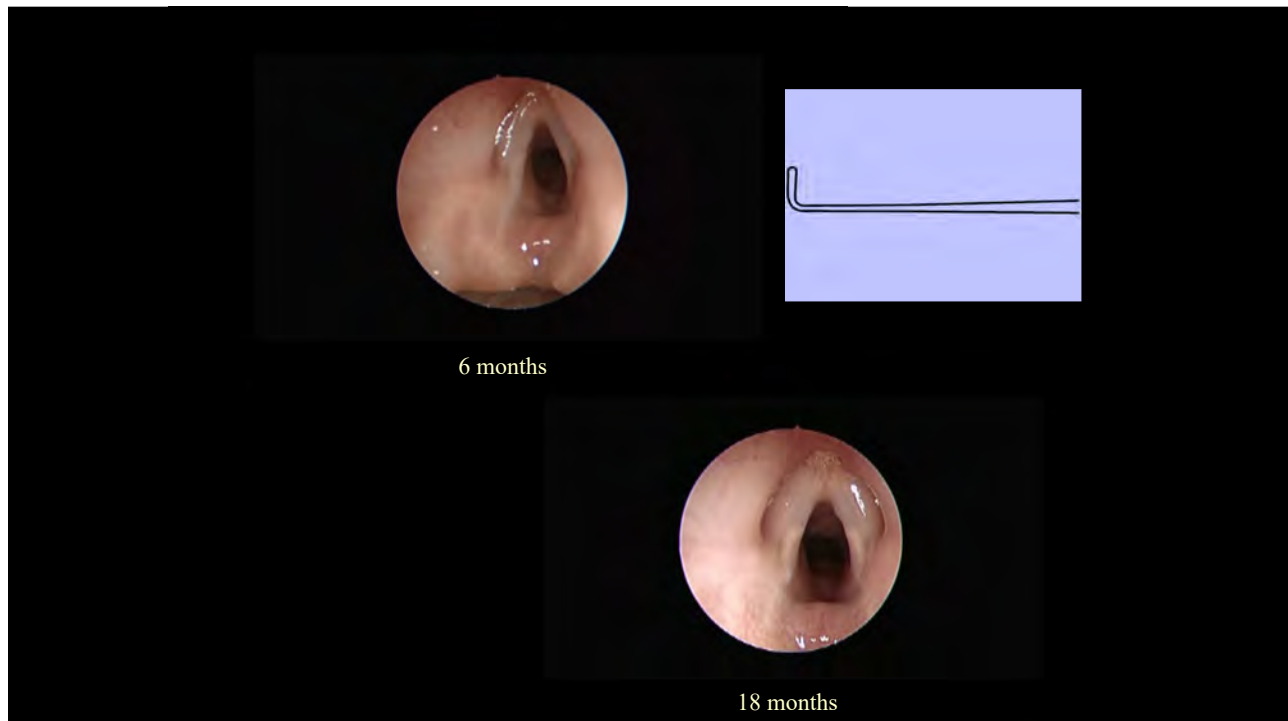
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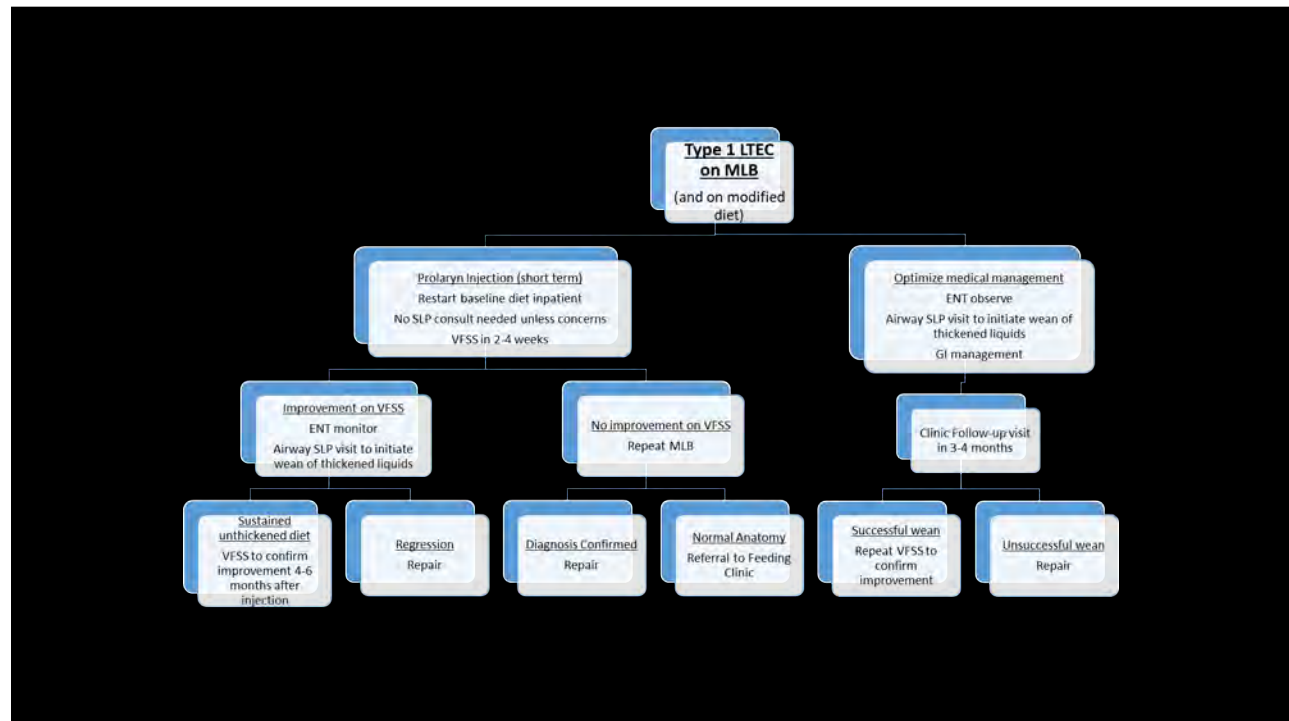
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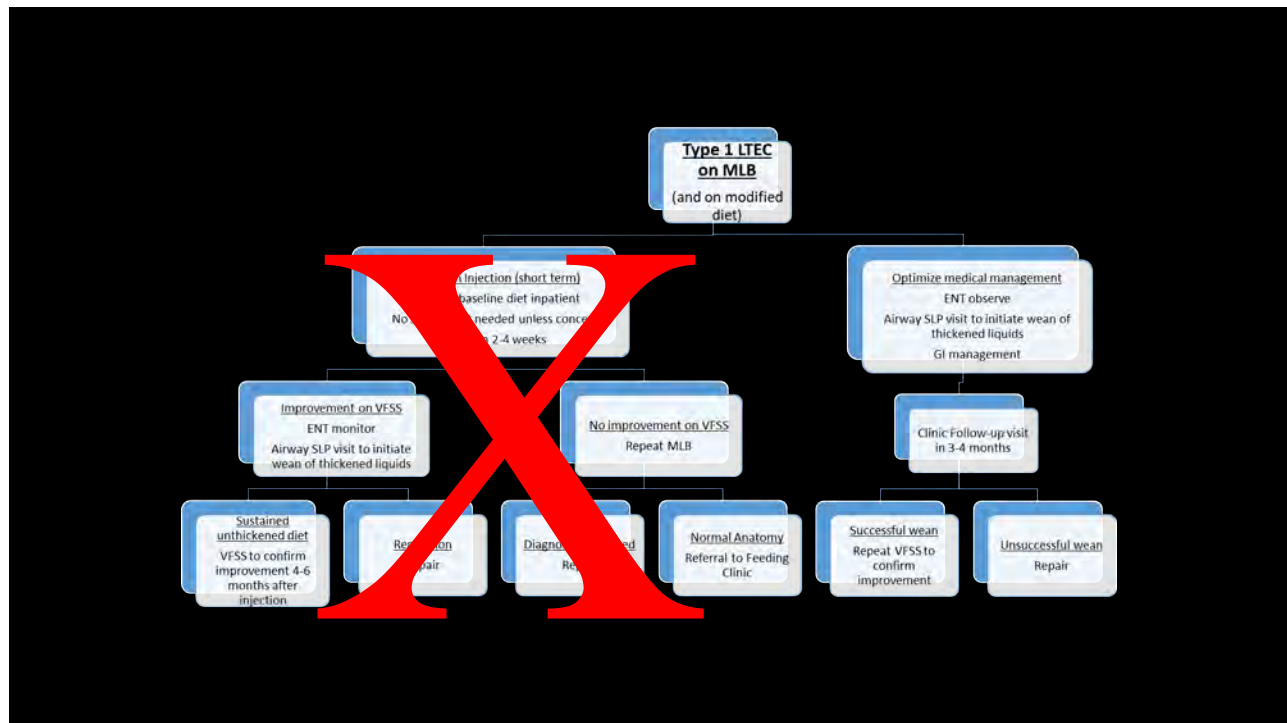
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- Is the "Type I Cleft" to the airway surgeon what the "Posterior Tongue Tie" is to the dentist?

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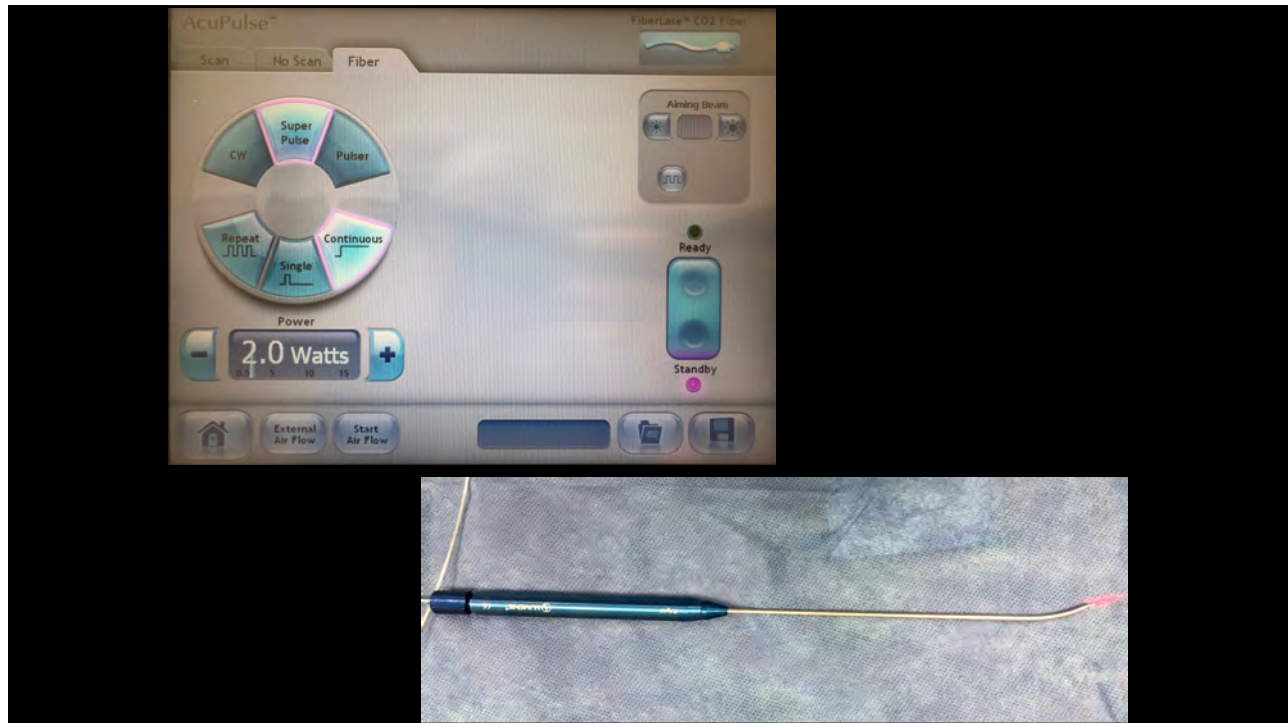
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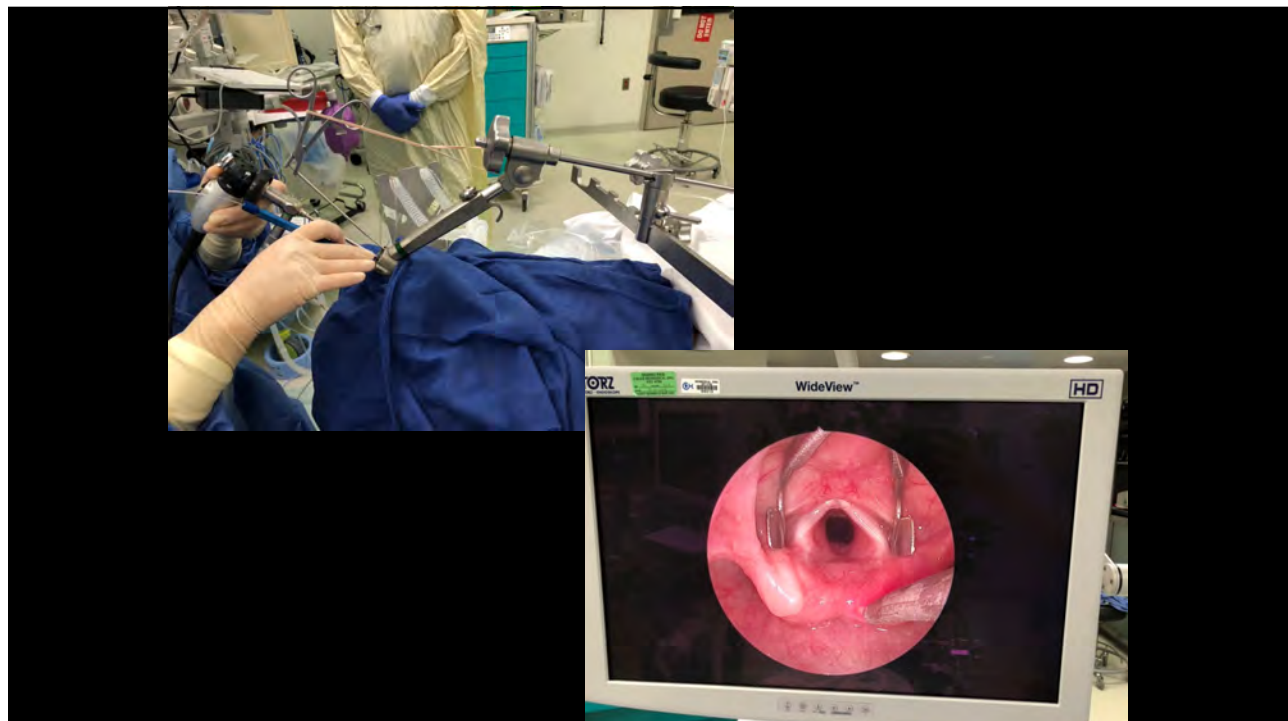
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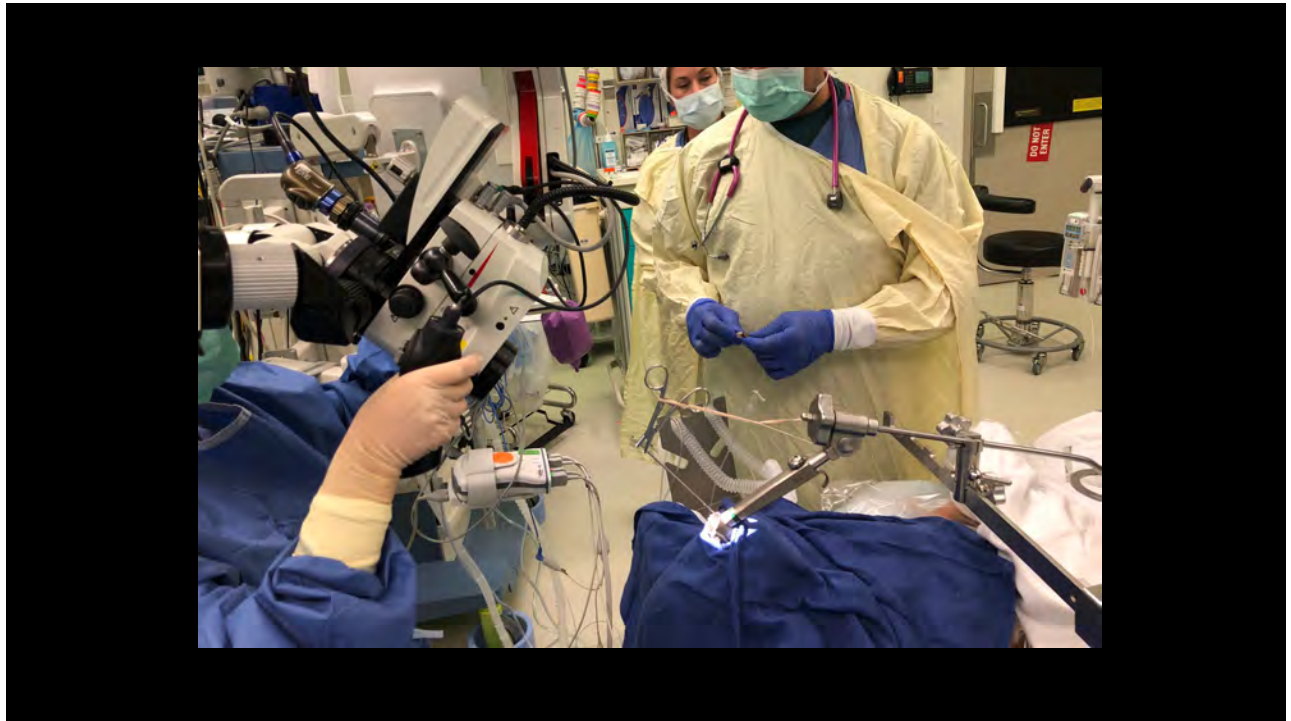
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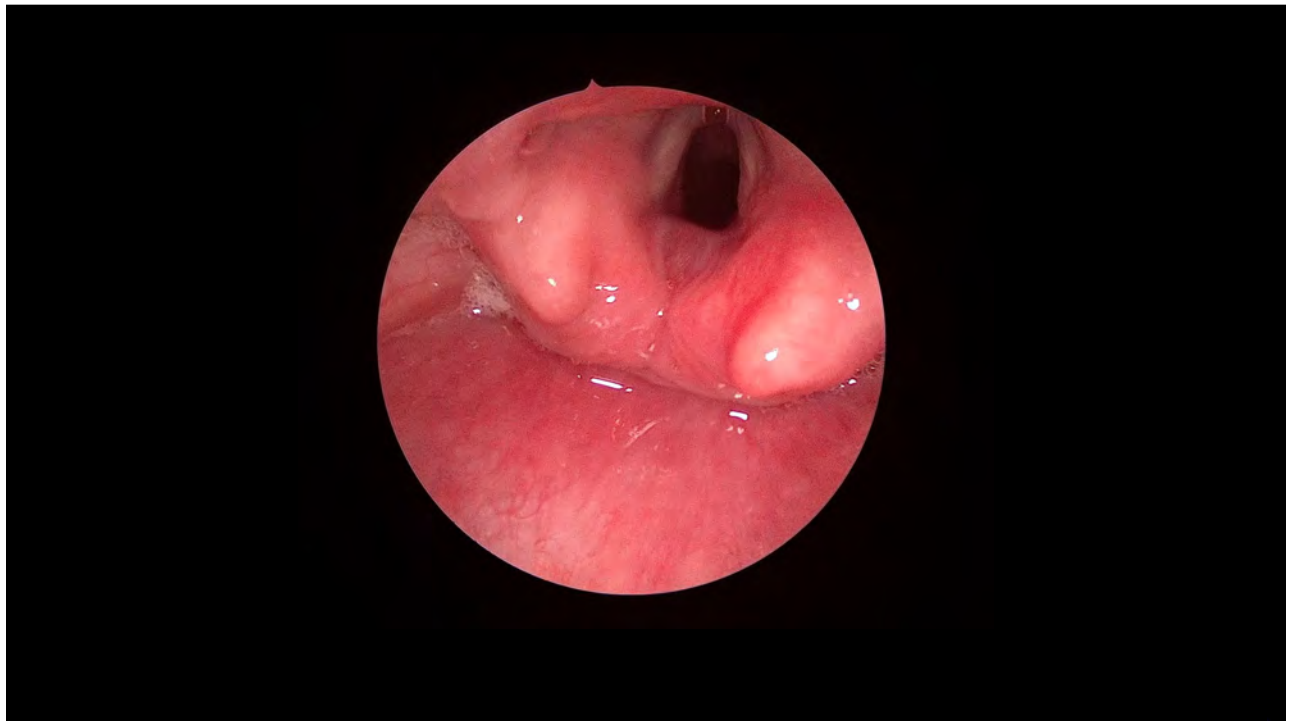
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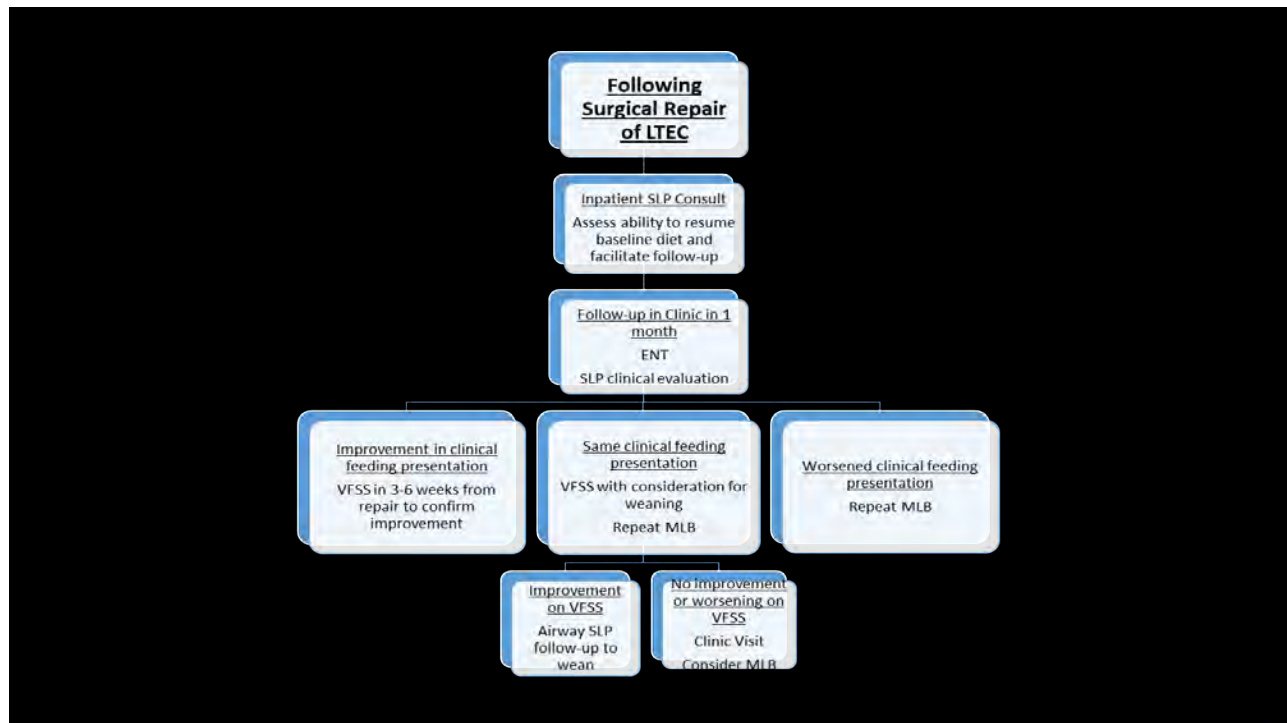
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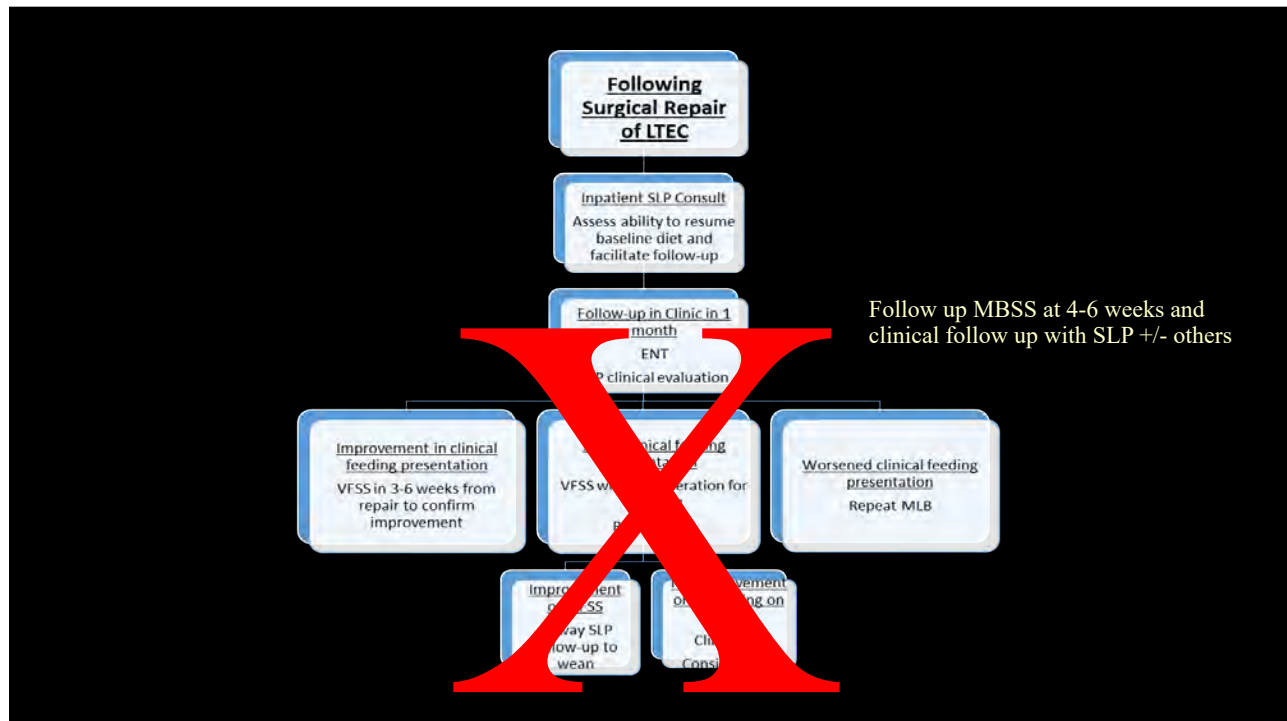
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DO

- Always double check your diagnosis before proceeding
- Question the value of “injections”
- Simplify your postoperative approach
 - This is where your team approach can really shine

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DON'T

- Label everything a “Type 1 Cleft”!
 - What really is a “deep notch” anyhow?
- Operate unless medically optimized
- Promise families immediate success
 - Swallow improvement requires rehabilitation

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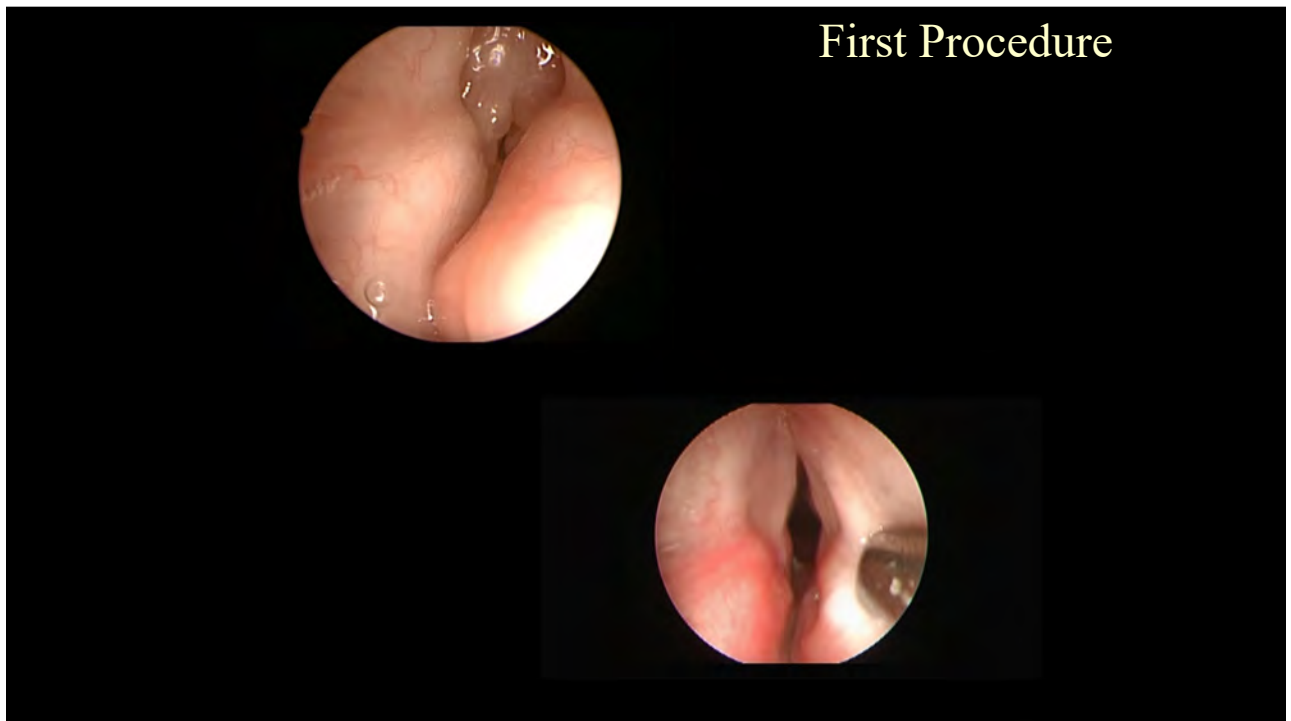
Let's CHARGE forward!

- 32 mo male
 - CHARGE
 - TEF with esophageal atresia repaired
 - Multiple cardiothoracic vessel anomalies
 - PDA repaired, transient unilateral VC paralysis
 - Diverticulum of Kommerell with esophageal impingement
 - G-tube, GERD, Nissan
 - Adenotonsillar hypertrophy with OSA
 - CSOM, SNHL

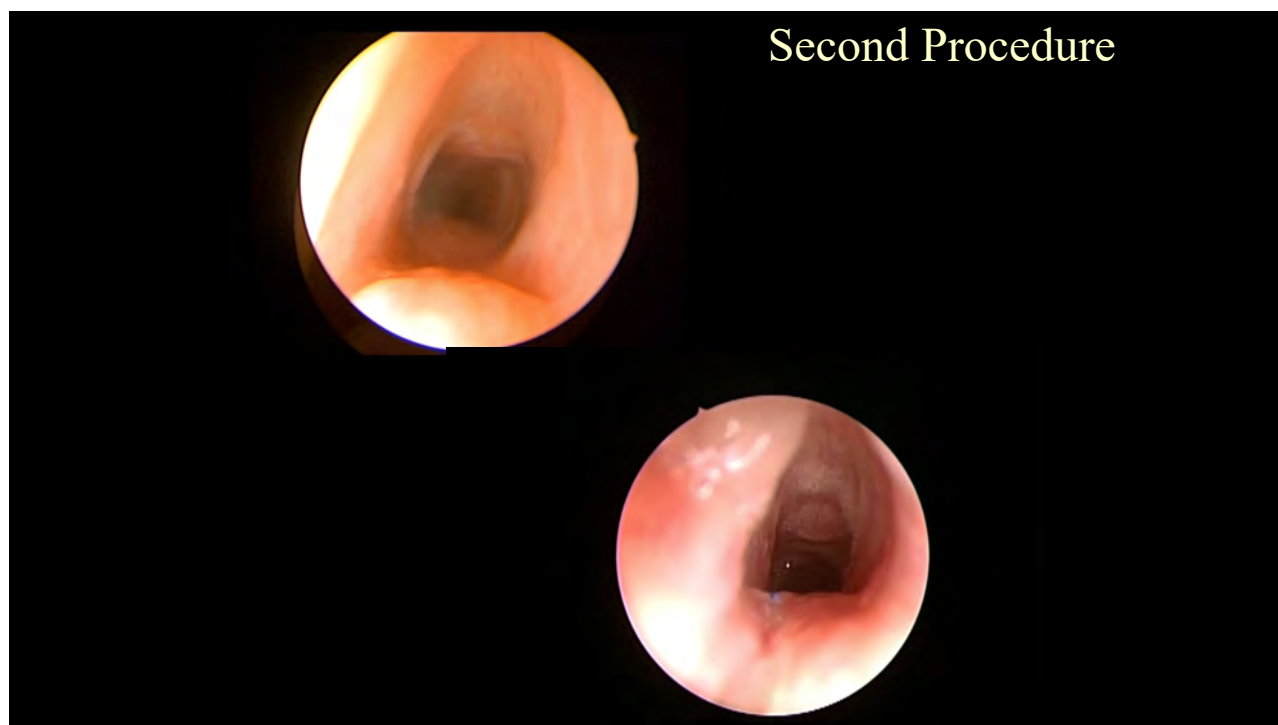
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DO

- Take these patients very seriously!
 - Often have comorbidities
 - May look great awake but misbehave after anesthesia
 - Counsel families that revisions may be necessary

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DON'T

- Give up if the first crack at it doesn't work
 - Optimize
 - Optimize again
 - Take care of all other surgical pathology before tackling the cleft

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LTEC III

- Virtually every one of these cases has thrown me a curve ball
- The two intraoperative “codes” that I have had have been in these patients
- Both endoscopic and open repair techniques have been described for these cases

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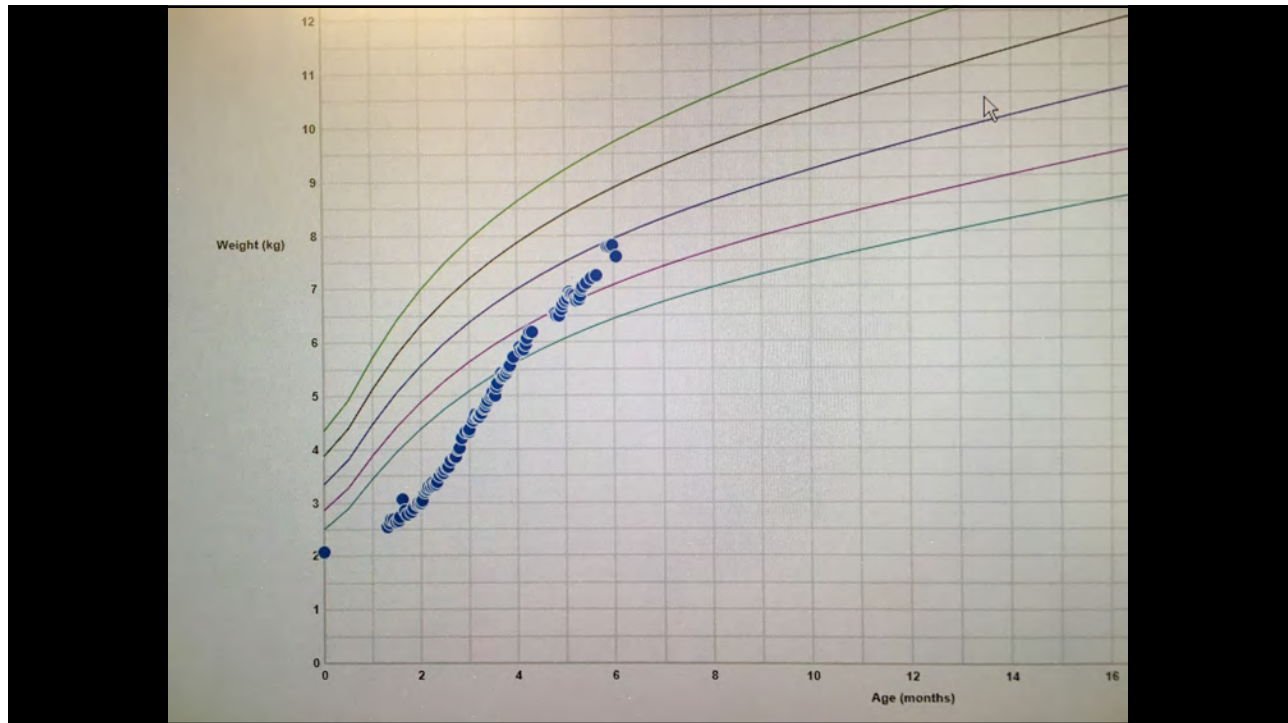
Such Disrespect!!!

- 6 week old male
 - 33 week preme
 - Airway distress at birth, difficulty ventilating
 - MLB DOL#3 diagnosed with LTEC Type III
 - Extubated and eventually weaned off oxygen
 - Transferred to CHOP for second opinion on management
 - Exam: Room air, NO STRIDOR, Weight: 2.5 kg

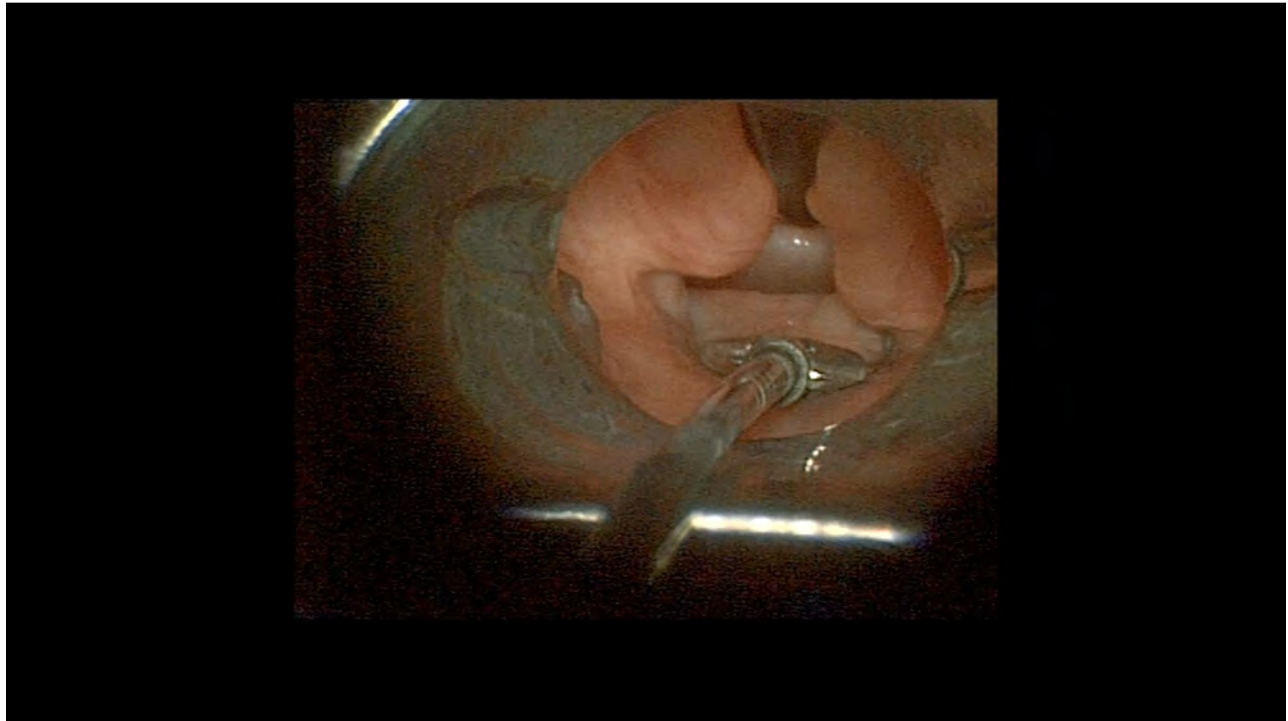
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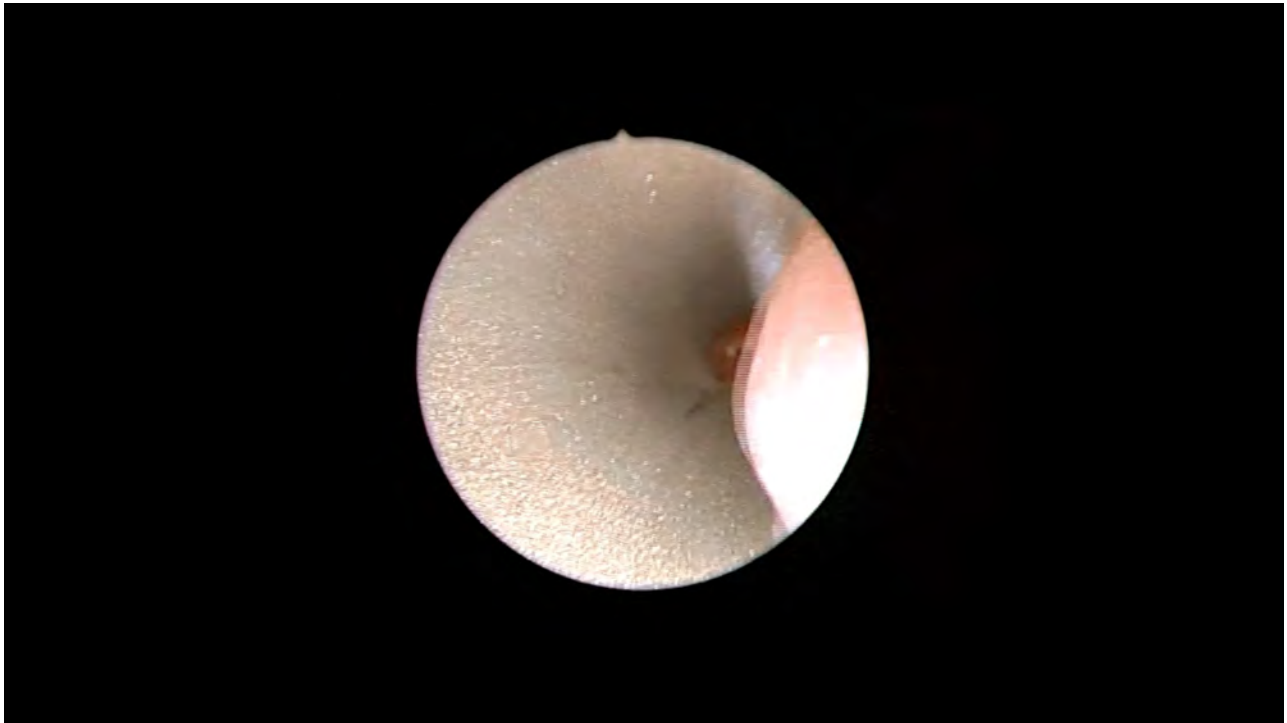
Such Bravado!

- Look at us, closing up a Type III LTEC endoscopically. Yay!! Woohoo!! Winning!!! We ROCK...
- Let's just drop him off in the PICU and go grab a drink!
- Plan will be to extubate in the morning after we round on him...

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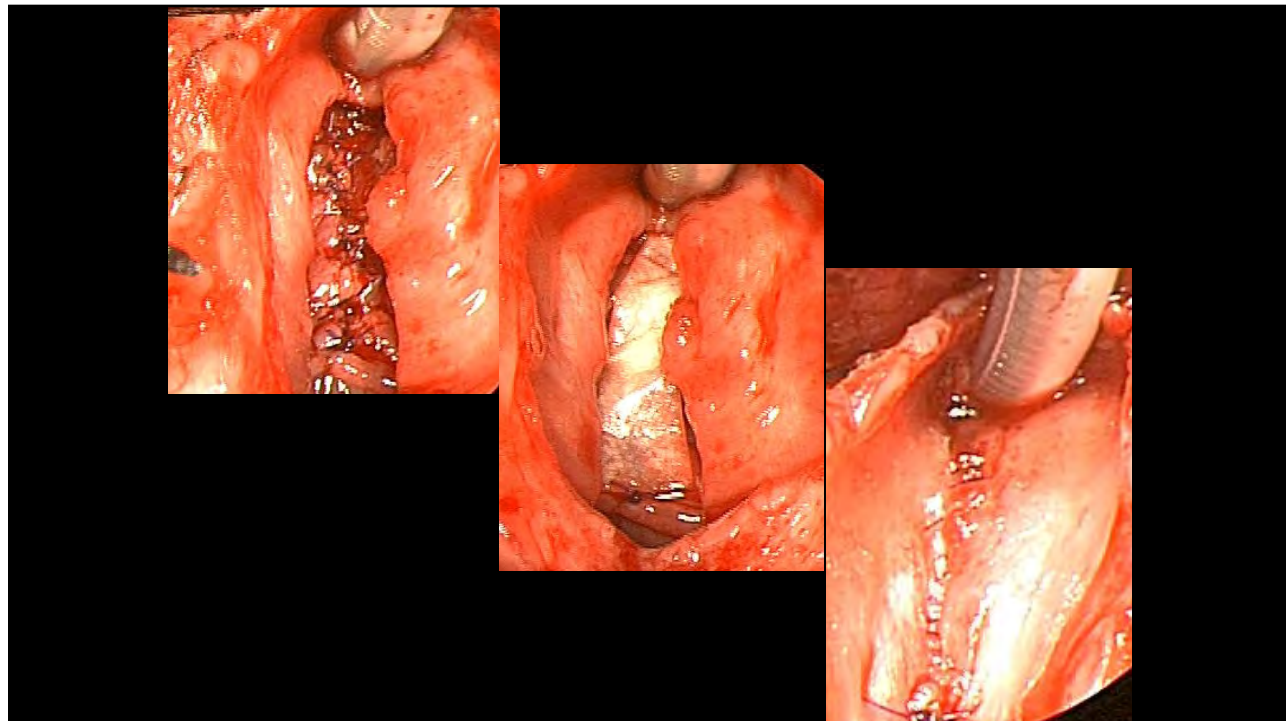
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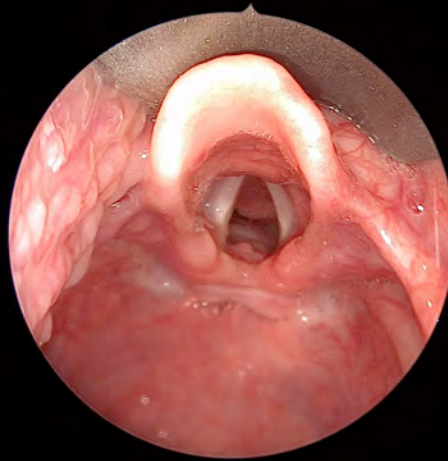


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3 years post repair



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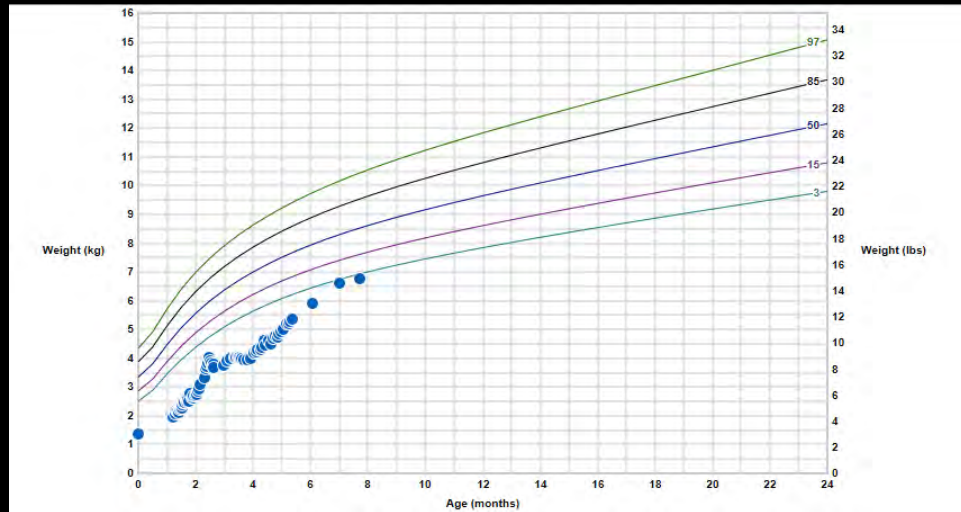
This one is really the (O)pitz!!!

- 5 week old male
 - 29 week gestation
 - Respiratory distress at birth → Intubated
 - Thought to have TEF → Transferred to CHOP
 - Cleft lip and plate
 - Hypospadias
 - Diagnosed with Optiz GBBB

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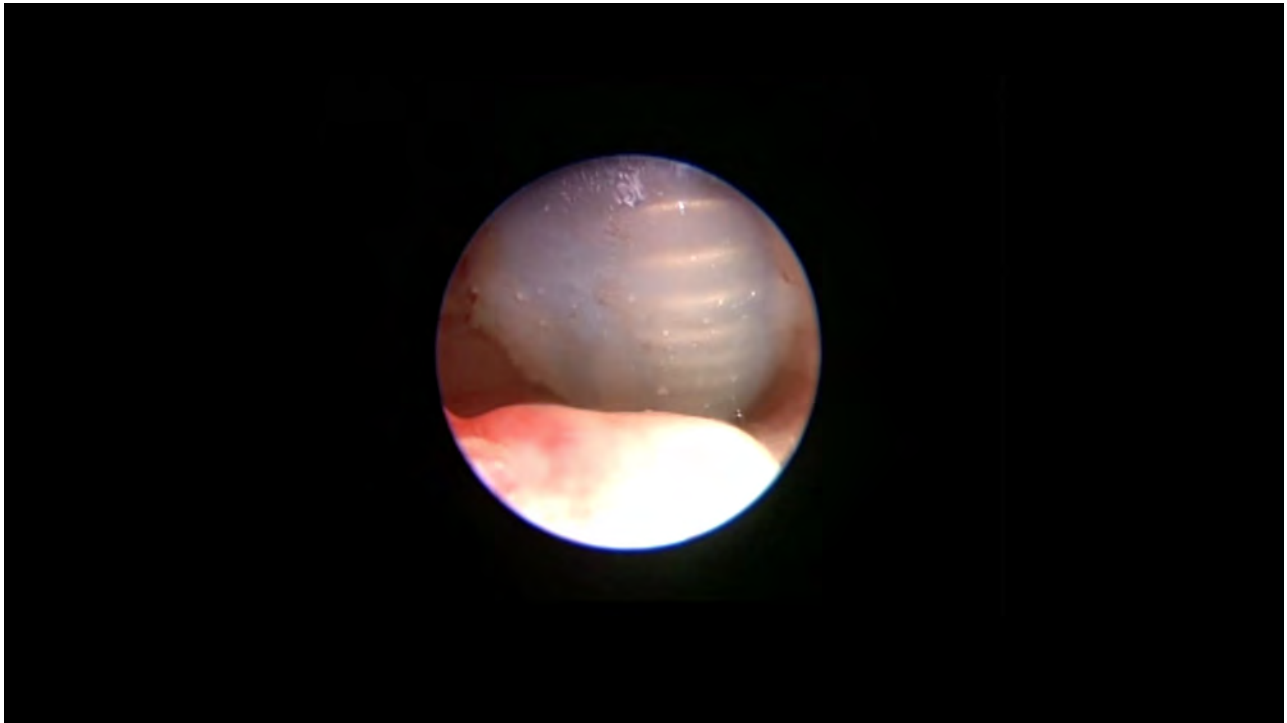
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This one is really the (O)pitz!!!

- So, we are doing great here and ultimately decannulate our patient. Yay!! Woohoo!! Winning!!! We ROCK...
- ...And a year later try and close the TCF...
- Leading to immediate distress and replacement of the trach...



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DO

- Have a very healthy respect for these patients
 - Many are syndromic
 - Tracheomalacia may present intra or postoperative issues
- Question the ability of endoscopic repair to fix these

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DON'T

- Operate until you sort out and optimize comorbidities
- Underestimate the changes in the airway dynamics post-repair
- Assume that your repair is “successful” without longitudinal endoscopic follow up

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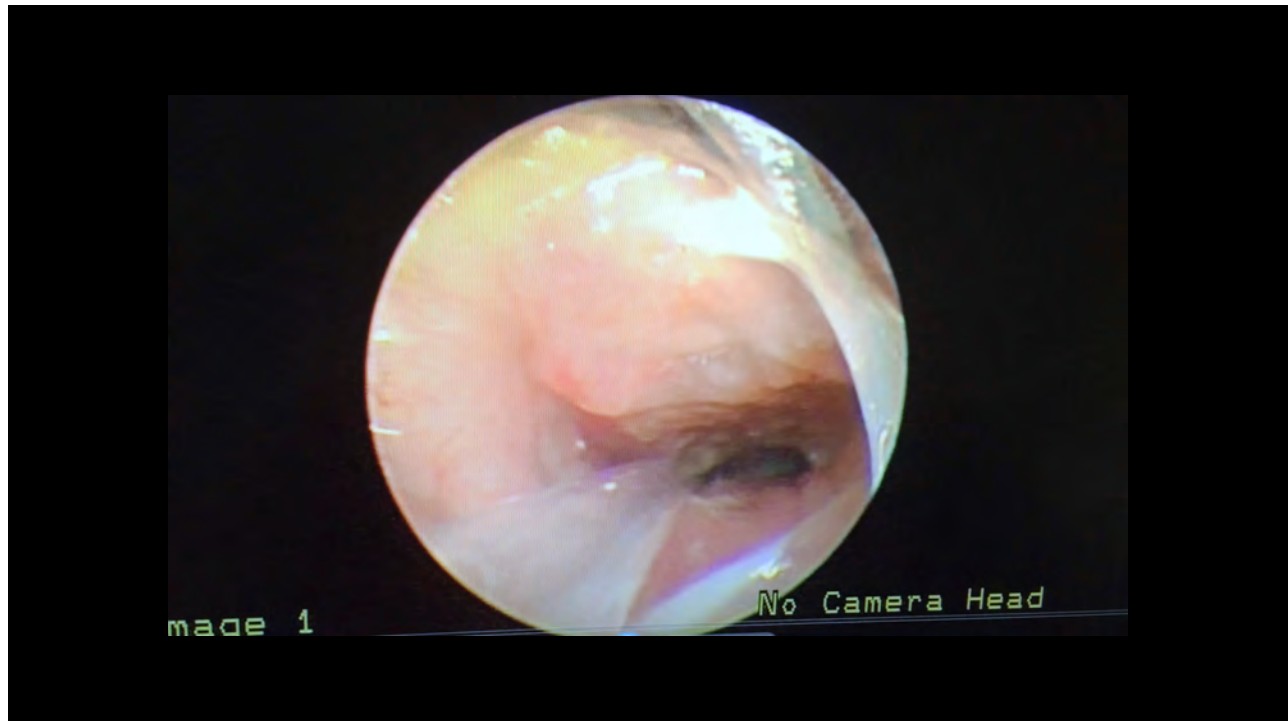
LTEC IV

- Newborn female
 - 27 week, 830 gram, twin
 - Respiratory distress at birth, positional ETT
 - Presumptive diagnosis of TEF
 - Distress at DOL #3
 - Bedside MLB concerning for LTEC
 - Pulmonary reserve poor
 - Requires oscillator

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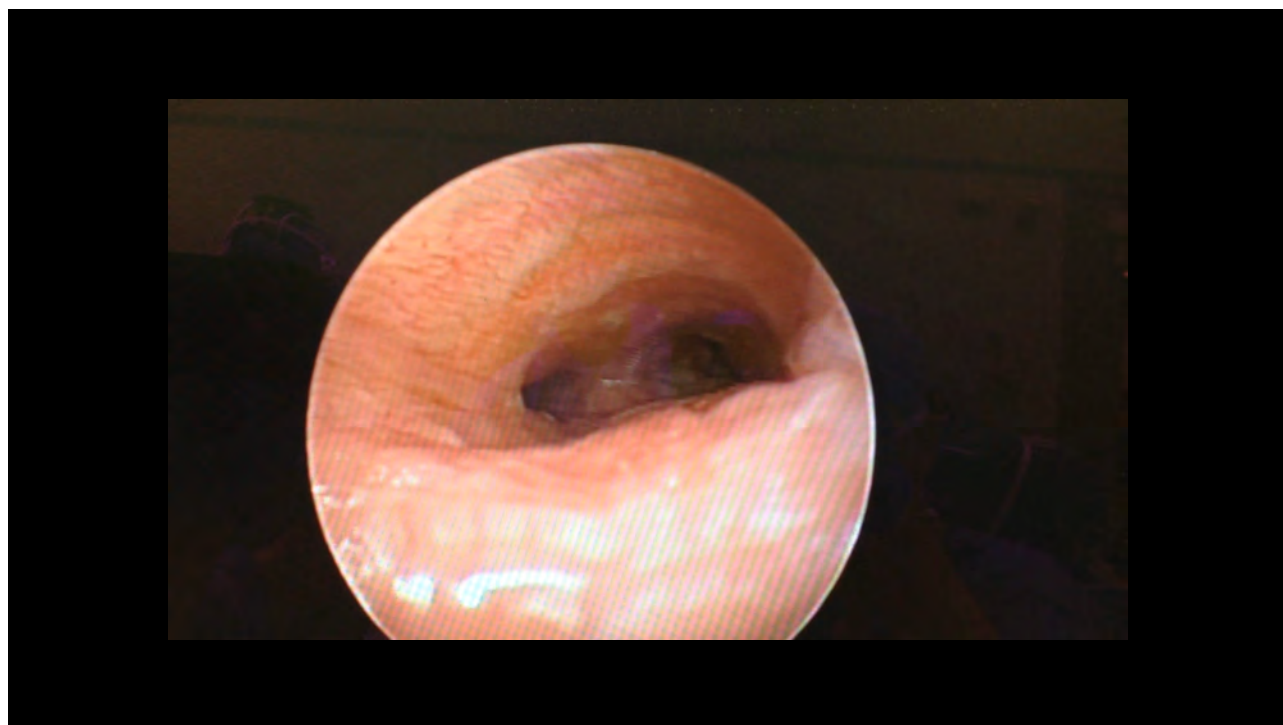


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LTEC IV

- By 5 months of age...
 - “Stable”
 - Fellow at bedside 2-3x/week to reposition ETT and scope
 - Parents want everything done
 - Mom: Lawyer
 - Dad: Son of an orthopedic surgeon

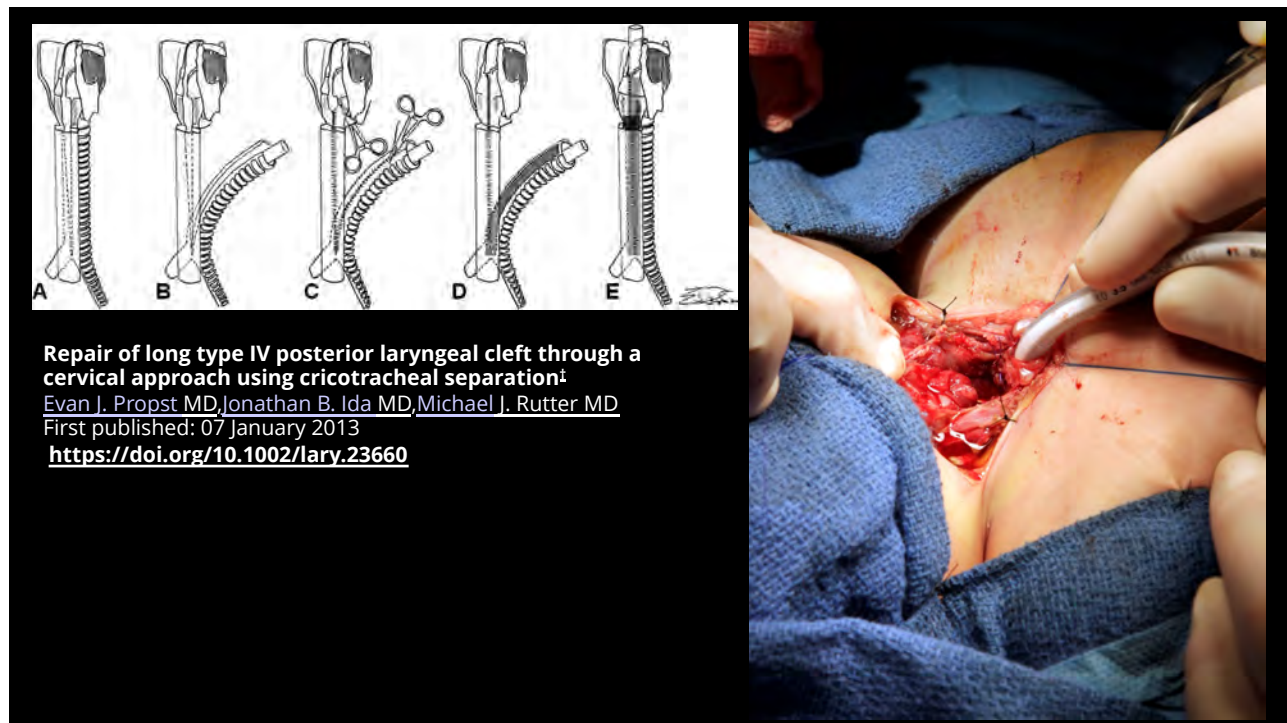
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Would you offer an operation???

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Conclusions

- Define who you want to be
 - We are surgeons and like to fix things
- Create a systematic triage process
 - And stick to it!
- Define and question what is and when to repair a Type I cleft
- Have a very healthy respect for advanced LTECs

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THANK YOU!



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