A Validated 3D Printed Laryngeal Suturing Simulator for Endoscopic Laryngeal Cleft Repair

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Objectives: Endoscopic laryngeal cleft repair (ELCR) with endolaryngeal suturing is an advanced surgical skill. This study objective was to assess the validity of 3-dimensionally (3D) printed laryngeal suturing simulator for ELCR.

Study Design: Development and validation of a simulator for ELCR.

Methods: An ELCR model was developed using 3D printed and readily available materials. Participants were surveyed before and after a simulation session using five-point Likert scale questions. Performance data was assessed using blinded expert video review and rated using a novel objective structured assessment of technical skills (OSATS) for endoscopic laryngeal suturing.

Results: Twenty-one participants ranging from residents to attendings completed the simulation session. Survey respondents reported on a five-point Likert scale that the model was "easy to use" and "quite realistic" (both mean of 4). Confidence improved significantly in 86% of participants (p < 0.01). Overall OSATS scores (out of a total of 55) showed a median improvement in technical skills of 11.7 points (p = 0.004). OSATS demonstrated good intra-rater ($\kappa = 0.689$ and 0.677) and moderate inter-rater ($\kappa = 0.573$) reliability. Completion times improved from the first to the last suture by a median time of 512 to 350 s (decrease of 202 s, p = 0.002). Participants with no prior ELCR experience improved more than those with in vivo experience.

Conclusion: This study demonstrates the validity of a simulator utilizing 3D printed larynges for ELCR. A novel OSATS for endoscopic laryngeal suturing was successfully implemented. Confidence, technical skills, and completion times improved with the use of the model across a variety of participants.

Key Words: 3D printing, education, endoscopic suturing, laryngeal cleft, simulation.

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INTRODUCTION

Laryngeal clefts are congenital upper airway malformations that result in a deficiency of tissue between the laryngotracheal complex and esophagus. They are

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relatively rare, occurring in approximately 1 in 10,000 to 1 in 20,000 live births, but are being diagnosed with increasing frequency.^{1,2} The most common grading system for laryngeal clefts is the Benjamin and Inglis classification, which ranks clefts from type one to type four depending on anatomic depth.³ Type one and type two clefts, which extend either to the level of the vocal cords or partially into the cricoid, respectively, are the most prevalent.⁴

While some laryngeal clefts may be asymptomatic or can be managed conservatively, most undergo surgical intervention for definitive treatment of dysphagia and aspiration.^{2,5} Historically, these were repaired with open surgery, however, the vast majority of laryngeal cleft repairs are now performed endoscopically.^{2,4,5} Similar endoscopic procedures (e.g., interarytenoid suture augmentation) are also becoming more widely used for dysphagia treatment in patients with normal anatomy. Endoscopic endolaryngeal suturing, such as that used in endoscopic laryngeal cleft repair (ELCR), is an advanced and technically challenging surgical skill. It can be difficult to gain competency in ELCR within the course of a residency or fellowship program, placing otolaryngology trainees at risk of graduating with technical deficiencies.⁶

Surgical simulation is widely used in modern surgical education to augment technical training.⁷ It allows for

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the practice of procedural and situational skills in a lowstress environment that does not affect patient morbidity and mortality. Additionally, simulation removes the time constraints of live operating, and allows for repetition, reflection, and real-time feedback. Within the field of otolaryngology, simulators including both physical models and virtual reality have been described in every subspecialty.⁸ Airway surgery, including laryngeal surgery, is no exception to this trend, but no validated simulators have been described for ELCR.

Utilizing a combination of readily available and three-dimensional (3D) printed materials, we developed a realistic training model for ELCR. A novel endoscopic laryngeal suturing objective structured assessment of technical skills (ELS-OSATS) was developed to evaluate the performance of simulated ELCR. We hypothesized that this model would improve both user confidence and competence in technical skills.

METHODS

Model Development

An ELCR simulator was created with two main components: a silicone laryngeal cleft model and a positioning framework for simulated endoscopic surgery.

For the model, digital modeling software (Mimics/3-Matics [Materialise, Leuven Belgium]) was used to virtually create atypical airway anatomy from a normal CT scan to develop a laryngeal cleft construct. Cleft depth was placed to a level below true vocal cords, representing a type two cleft. This was then used to produce a three-dimensional (3D) mold and printed on a Flashforge 3D printer (Zhejiang, China) using polylactic acid (PLA) material (Fig. 1A). Dyed silicone and Slacker additive (Macungie, Pennsylvania) were cured in the molds to create the soft tissue framework. True vocal cords were painted white using silicone paint to aid in realism and differentiation from surrounding tissue (Fig. 1B).

A framework was created using a hospital standard plastic supply bin, rolled surgical towels, duct tape, and a large Lindholm laryngoscope (Fig. 2A). Holes were cut in the sides and bottom of the bin to accommodate the laryngeal model. The cleft model was then stabilized using a temporal bone holder (Fig. 2B) and placed into the framework housing the laryngoscope to complete the simulator (Fig. 2C).

When used with a microscope or endoscope, the simulator re-created an ergonomically similar set-up to a patient in suspension laryngoscopy in the operating room (Fig. 2D). Standard micro laryngeal instruments were then used to perform ELCR. The view of the cleft model through the simulator imitated the view of the larynx during an ELCR on a real patient (Fig. 3).

OSATS Development

A novel ELS-OSATS tool was developed for the assessment of endoscopic suturing skills used during ELCR (Figure S1) using a modified Delphi method. The development panel was composed of expert pediatric otolaryngologists from three institutions who perform ELCR regularly in their practices. Similar to other OSATS used in otolaryngology, this included both task-specific and global operative performance sections.^{9,10} The task-specific portion was developed by polling the expert panel and dividing the operative procedure of endoscopic endolaryngeal suturing into compartmentalized steps required to successfully complete the procedure. Finalized steps included microlaryngeal instrument handling, endoscopic suturing (further divided into tissue tension and support and appropriate suture placement with regard to depth and position), endoscopic knot tying, and use of the endoscopic knot pusher. The global-operative performance portion of the scale was developed using the American College of Surgeons OSATS global rating scale.^{11,12} All specific items (five for task-specific performance as listed above, and six for standard global operative performance) were rated on a five-point Likert scale with one representing poor performance and five representing outstanding performance, for a total of 55 possible points.

Data Collection

Internal Review Board approval was obtained from Seattle Children's Hospital and the University of Washington to test the simulator on live participants. The model was implemented at a simulation station for teaching and practice of ELCR at the annual Northwest Airway Course. Participants consisted of a mix of resident, fellow, and attending otolaryngologists. Prior to



Fig. 1. A 3D printed mold of a type 2 laryngeal cleft (A) used to fashion laryngeal cleft models out of dyed silicone (B).



Fig. 2. A framework for housing the model created from readily available materials (A), into which a stabilized laryngeal model (B) is placed to complete the simulator for laryngeal cleft repair (C). When used with a microscope, the simulator re-creates an ergonomically similar set-up to a patient in suspension laryngoscopy in the operating room (D).

simulator use, participants watched a 5-min instructional video on ELCR using the simulator. Study participation was voluntary, and all participants in the course had scheduled time with the simulator.

Each simulation consisted of a 10-min session in which participants performed ELCR on the models. Additional equipment used included an operating microscope (Zeiss, Oberkochen, Germany) and standard microlaryngeal instruments (Karl Storz, Tuttlingen, Germany). Suture repair was completed using 6-0 polydioxanone suture (PDS, Ethicon, Inc., Somerville, New Jersey), with five knots per suture throw. Participants were encouraged to place as many sutures as possible within the time limit. Each participant completed either one or two sessions, which were video recorded. Participants completed surveys before and after sessions rating their confidence in performing laryngeal suturing on a five-point Likert scale, with one being no confidence, to five being complete confidence (Figure S2). Surveys also detailed their training level, prior experience with in vivo ELCR, self-evaluation of each suture throw, and opinion of realism and ease of use of the simulator.

Deidentified video recordings were edited to extract the first and last suture throw for each participant. Each participant, therefore, had a pre-simulator use video (first suture throw) and a post-simulator use video (last suture throw of their last session). Videos of each individual suture throw were then reviewed in a randomized, blinded fashion by two pediatric otolaryngology attendings. Each suture throw was ranked using the ELS-OSATS. The videos were re-randomized and re-evaluated by the same reviewers six months after the initial review for intra-rater comparison.

Data Analysis

Statistical analyses were performed on participant selfevaluation of confidence, ELS-OSATS scores, and suture completion time. Independent t-tests were used to compare completion time and ELS-OSATS scores by experience level at the initial attempt. The Wilcoxon signed-rank test was used to compare ELS-OSATS scores as well as changes in survey responses



Fig. 3. Endoscopic view of the laryngeal model (bottom) compared to an in vivo larynx (top) before and after endoscopic suturing.

between attempts (i.e., first to last suture throw). A paired t-test was applied to compare completion times between attempts. Cohen's κ testing was used to assess inter-rater and intra-rater reliability of ELS-OSATS rankings.

RESULTS

Simulation sessions using the ELCR model were performed by a total of 21 otolaryngologists, including 16 residents (76%), two fellows (10%), and three attending physicians (14%). Residents comprised of trainees from clinical post-graduate year (PGY) two (n = 6, 28.5%), three (n = 6, 28.5%), and four (n = 4, 19%). Fellows were both PGY six (n = 2, 10%). Attendings ranged from two to ten years in practice (n = 3, 14%).

All participants completed a simulation session with the model at least one time and all completed at least one suture throw during their session(s). A completed suture throw was defined as an endoscopic passage of the needle through supraglottic tissue and endoscopic tying of one to five knots. Out of the full cohort who completed one throw (n = 21), 76% (n = 16) completed two suture throws, and 14% (n = 3) completed three suture throws. Of the participants who completed only one suture throw (n = 5), three attempted but did not complete a second throw during the allotted time.

All participants completed the survey with a 100% response rate. Overall, the simulator was rated as "easy to use" and "quite realistic", both with a mean response rate of four. All participants either "agreed" or "strongly agreed" (four or five on five-point Likert scale) that otolaryngology trainees would be better prepared for ELCR after using the simulator.

Participants reported confidence increased significantly after using the simulator, improving a mean of one full Likert point from "basic comfort with steps" (2/5) to "intermediate confidence with all steps" (3/5) (p < 0.001, Fig. 4). Self-assessment of individual suture throws attempts also improved by a full point between first and second attempts (p = 0.008, Fig. 4). No statistically significant improvement was seen between the second and third attempts due to insufficient power (n = 3).

ELS-OSATS data (maximum total of 55 points) obtained from the blinded reviews demonstrated a significant overall score improvement in skill of 11.7 points when comparing first to last suture throws (an increase of median score from 26.0 [IQR 12.3–34.3] to 37.7 [IQR 28.9–49.4], p = 0.004) (Fig. 5, Table I). Improvement was also seen when breaking down ELS-OSATS into global operative performance (p = 0.02) and ELCR task-specific performance (p = 0.03). Calculated completion times of each suture throw from the video review also showed improvement from a median time of 512 to 350 s (decrease of 202 s, IQR 15–298, p = 0.02, Fig. 6).

Inter-rater and intra-rater reliability were also calculated from the blinded reviewer data (Table II). Intrarater reliability showed good agreement for rater one ($\kappa = 0.677$) and for rater two ($\kappa = 0.689$). Inter-rater reliability was shown among all OSATS categories, with an overall moderate concordance ($\kappa = 0.573$). The breakdown by OSATS categories is detailed in Table II.

Performance and confidence did differ by participant experience level. When stratifying by PGY year, initial confidence levels varied widely but correlated strongly with prior in vivo ELCR experience (p < 0.0001). Participants with no prior experience, who never observed or participated in ELCR, had both greater subjective and



Fig. 4. Participant survey data ranking self-assessed confidence before and after simulator use on a five-point scale. Median overall improvement was one point.



Fig. 5. ELS-OSATS results (maximum of 55 points) compared between first and last suture throws as calculated via blinded review. Median overall improvement was 11.7 (21%).

objective improvement than those with some experience (≥ 1 in vivo ELCR). The group with no experience (n = 7) had an even greater improvement in confidence (1.5 points), compared to the group with experience (n = 9, p < 0.01). This group also increased in median overall ELS-OSATS scores by 12.1 points (IQR 1.3–18.5) as opposed to 10.5 points (IQR 0–18) for experienced participants (p = 0.04) and improved in completion time by 269 s (IQR 254–421) as opposed to 55 s (IQR 0–192) (p < 0.001, Table II).

DISCUSSION

Endoscopic suturing for ELCR requires advanced surgical skills which can be difficult both to learn and to teach. Limitations in training opportunities exist due to the single-operator nature, operative time constraints, and low case volumes. Additionally, the vast majority of patients undergoing these operations are infants or young children, making other factors such as anesthetic time even more important. Even over the course of an entire residency or fellowship, otolaryngology trainees may lack sufficient exposure to in vivo ELCR to gain competency. Surgical simulation using ELCR models offers a realistic way to develop these skills, which may ultimately translate to improved surgical proficiency and more effective training when applied to live patient opportunities. Our described surgical simulator represents the first report of a validated tool to improve user confidence and surgical skills for ELCR.

The simulator was successfully implemented among a group of otolaryngologists ranging from junior residents to attendings, indicating its feasibility. Participants overall rated the simulator as "quite realistic" (Likert scale

TABLE I.

Median ELS-OSATS and Completion Time Data for All Participants Overall and Compared between Participants with No Prior Experience and Participants with Some Experience (≥1 in vivo ELCR).

| | First suture throw | | | Last suture throw | | | Change from first to last | | |
|-------------------------------------------|--------------------|-------------------------|-------------------------------|-------------------|-----------------------------|-------------------------------|---------------------------|-----------------------------|-------------------------------|
| | Total (n = 16) | No experience $(n = 7)$ | Some experience (n = 9) | Total (n = 16) | No experience (n = 7) | Some experience (n = 9) | Total (n = 16) | No experience (n = 7) | Some experience (n = 9) |
| Total ELS-OSATS (maximum 55) | 26.0 | 21.1 | 28.6 | 37.7 | 33.2 | 39.1 | 11.7* | 12.1* | 10.5* |
| Task-specific performance (maximum 25) | 11.0 | 9.9 | 11.8 | 16.4 | 14.6 | 17.8 | 5.4* | 4.7* | 5.9* |
| Global operative performance (maximum 30) | 14.3 | 11.3 | 16.7 | 20.1 | 18.6 | 21.3 | 5.8* | 7.3* | 4.5* |
| Completion time (seconds) | 512 | 630 | 428 | 350 | 361 | 373 | 202* | 269* | 55* |

Asterisk (*) indicates p < 0.05.



Fig. 6. Completion time between first and last suture throws. Median overall improvement was 202 s (3 min 22 s).

median of 4) and the majority (n = 15, 72%) strongly agreed (Likert scale 5) that otolaryngology trainees would be better prepared for ELCR after using the simulator. These patterns were corroborated by expert attending physicians who perform ELCR regularly in their practice (n = 3), showing face validity.

Both an increase in subjective confidence and a statistically significant improvement in objective measures using a novel ELS-OSATS were observed after the use of the simulator. ELCR-specific aims were identified and successfully performed on models, including laryngeal tissue handling, endoscopic suture placement, endoscopic knot tying and use of the endoscopic knot pusher, and endoscopic suture cutting. The ELS-OSATS showed sufficient inter-rater and intra-rater reliability to indicate its utility in assessing these ELCR task-specific performances as well as global operative performance for endoscopic suturing. Objective improvement was seen in overall scores (p = 0.004), and both for global operative performance (p = 0.02) and ELCR task-specific

| TABLE II. Inter-rater and Intra-rater Reliability for ELS-OSATS Compared between Blinded Video Reviewers. | | | | | | | | | |
|-----------------------------------------------------------------------------------------------------------------|------------------------------|---------------------------------|---------------------|--|--|--|--|--|--|
| | Cohen's κ | | | | | | | | |
| | Task specific performance | Global operative performance | Overall performance | | | | | | |
| Inter-rater reliability | | | | | | | | | |
| Reviewer 1 versus Reviewer 2 | 0.560 | 0.581 | 0.573 | | | | | | |
| Intra-rater reliability | | | | | | | | | |
| Reviewer 1 | 0.682 | 0.665 | 0.677 | | | | | | |
| Reviewer 2 | 0.688 | 0.680 | 0.689 | | | | | | |

performance (p = 0.03). These results collectively demonstrate the content validity of the ELCR simulator.

Construct validity, defined as the degree to which the test items identify the quality, ability, or trait it was designed to measure,¹³ was also shown by comparing users of varying experience levels. Participants with no experience with in vivo ELCR started off with lower skill levels on ELS-OSATS and confidence and improved in both categories by a larger margin than those with more experience. Time to completion also showed greater improvement in novice participants. These results indicate that incorporating surgical simulation for ELCR early on in training may be valuable.

Model validity was also shown when assessed by the Messick framework, another proposed method of demonstrating simulator validity. This system assesses five specific sources: content, response process (i.e., quality control), internal structure (e.g., reliability), relations with other variables, and consequences of the assessment.¹⁴ Evaluation of the content of the simulator and OSATS was conducted by expert otolaryngologists who regularly perform ELCR in their practice. The response process was ensured by providing a standardized instructional video to participants as well as by having the videos randomized and blinded prior to review. The internal structure was assessed using Cohen's kappa metrics for inter-rater and intra-rater reliability. Relation to other variables was determined statistically by comparing scores and completion times between groups with different training levels and in vivo ELCR experience.

One other laryngeal cleft model has been described in the literature by Kavanaugh et al.¹⁵ This model included a laryngeal cleft among a library of pediatric laryngeal pathologies constructed with various 3D printing materials. They showed that silicone-based material was the most mechanically similar to live tissue but did not evaluate the use of the simulator for training purposes.

Our model provides not only a validated way to practice ELCR but also offers several other benefits over other described airway simulation models. Printing the mold as opposed to the larynges themselves allowed for faster, more cost-effective production of multiple models. Material cost for the 3D printed mold and silicone larynges was less than \$10, not including production labor costs, dependent on who is producing the model. Each model can be reused for multiple simulations. In our experience, the tissue quality was adequate for 30–40 suture throws. The mold also makes model development portable and easily replicable, and they can be shared with institutions that may not have advanced 3D-printing capabilities.

Limitations in this study exist mainly in its relatively small sample size and the subjective nature of survey-based results. In particular, including more experienced attending surgeons in future work may help strengthen the validity of the model. Additionally, data was only collected from one institution which may contribute to additional implicit bias. We also acknowledge that the creation of models like this requires high-cost resources including a 3D printer. While these technologies are becoming more widely available and more cost-effective, they still involve a moderate upfront investment, space dedication, and specialized technical expertise for development. Options including inter-institutional collaborations and design sharing may help mitigate these constraints. Further testing is needed to demonstrate the predictive validity of this model showing how it affects operative performance for in vivo ELCR.

CONCLUSION

This study demonstrates face, construct, and content validity of a simulator for endoscopic ELCR. A novel OSATS for endoscopic laryngeal suturing was concurrently developed and implemented. Both subjectively perceived confidence and objective technical skills improved with use of the model across a variety of participants. Completion times per endoscopic suture throw decreased by over 3 min on average between the first and last attempt with the simulator. This data shows that the use of the simulator is a useful adjunct in surgical training and practice for ELCR.

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CONFLICT OF INTEREST

David A. Zopf is a co-founder of MakeMedical LLC, a University of Michigan-affiliated start-up.

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