INTRODUCTION

Clinicians are constantly faced with management of complex surgical cases requiring advanced and extensive tissue management. Successful management of these clinical scenarios for periodontal, oral surgery, or implant indications relies on a profound knowledge of the anatomical structures (Greenstein, Cavallaro, Romanos, & Tarnow, 2008; Tavelli, Barootchi, Namazi, et al., 2019; Tavelli, Barootchi, Ravida, Oh, & Wang, 2019). In particular, due to an increasing demand for augmenting atrophic posterior mandible and
subsequent implant rehabilitation (Urban et al., 2017), a firm understanding of the biological structures in this region cannot be overemphasized. Surgical complications in the lingual posterior mandible may include mucosal tissue laceration, intrusion in the sublingual space, trauma to the branches of the lingual artery, and injury to the lingual nerve (Annibali, Ripari, La Monaca, Tonoli, & Cristalli, 2009; Camargo & Van Sickels, 2015; Greenstein et al., 2008; Isaacson, 2004; Longoni et al., 2007; Urban et al., 2017). These unfortunate adverse events may result in unfavorable bone augmentation outcomes, massive hemorrhage (Askar et al., 2019; Camargo & Van Sickels, 2015), neurosensory disturbances, and impaired mastication function. Therefore, successful flap release and meticulous tissue management for achieving primary wound closure depend on a thorough anatomical understanding (Chan et al., 2010; Ritter et al., 2012; Urban et al., 2017, 2018).

In addition, it is now known that the quality and quantity of soft tissues greatly influence the healing of periodontal and implant procedures (Chao, Chang, Fu, Wang, & Chan, 2015; De Bruyckere, Eghbali, Younes, De Bruyn, & Cosyn, 2015; Fu et al., 2010; Lin, Chan, & Wang, 2013). This determines the tissue phenotype, which is currently evaluated through visual examination and probing (De Rouck, Eghbali, Collys, De Bruyn, & Cosyn, 2009; Eghbali, De Rouck, De Bruyn, & Cosyn, 2009). For instance, in the management of an extraction socket, tissue phenotype has been correlated with the amount of horizontal and vertical bone resorption that occurs following immediate implant placement (Ferrus et al., 2010). Other studies have shown its phenotypic feature in correlation with peri-implant marginal bone remodeling as well (Linkevicius, Apse, Grybauskas, & Puisys, 2009a, 2009b; Suarez-Lopez Del Amo, Lin, Monje, Galindo-Moreno, & Wang, 2016). Soft tissue features are a determinant of success of ridge augmentation and should be carefully evaluated before the surgery (Chen et al., 2017).

In medicine, the use of non-ionizing ultrasound has been established and advocated for many years (Bhaskar, Chan, MacEachern, & Kripfgans, 2018; Hoskins & Kenwright, 2015; Moskalik et al., 1995; Oelze & Mamou, 2016). In dentistry, its advantage for providing low-cost real-time cross-sectional images can be quite useful as it relates to providing optimal soft tissue contrast of pertinent anatomical structures and the peri-implant tissues (Bhaskar et al., 2018; Chan, Sinjab, et al., 2017; Chan et al., 2018). Additionally, ultrasound has been validated for measuring tissue thickness in different locations of the oral cavity (Chan, Sinjab, et al., 2017; Chan, Wang, Fowlkes, Giannobile, & Kripfgans, 2017). A recent study from our group applied and validated the use of ultrasound for accurate assessment of peri-implant tissues on human cadavers in comparison with direct visual and cone-beam computed tomography (CBCT) (Chan et al., 2018). In light of an increasing importance of the lingual anatomy, for the first time, we applied our ultrasound probe prototype to characterize the lingual structures, that is, the dimensions of the mucosa, mylohyoid muscle and lingual nerve, in comparison with histology. Feasibility of ultrasound to image the lingual nerve on live human patients was also investigated.

2 | MATERIAL AND METHODS

This project was prepared in accordance with the EQUATOR guidelines Standards for Reporting Qualitative Research (SRQR) (O’Brien, Harris, Beckman, Reed, & Cook, 2014).

2.1 | Study design of the cadaver research

Nine fresh un-embalmed fully/partially edentulous human cadaver heads were provided by the Department of Anatomy to the Department of Periodontics and Oral Medicine of the University of Michigan. To reduce the occurrence of any structural tissue damages, all specimens were kept frozen at a controlled temperature of -20°C (without formalin fixation) after harvesting from the human donors. Immediately prior to utilization for the experiments, the specimens were thawed to room temperature. For inclusion in the present research, it was required that specimens were either completely edentulous or partially edentulous particularly in the mandibular arch (past the mandibular canine). No other eligibility criteria were imposed in regard to the cadavers. This study was exempted by the University of Michigan Institutional Review Board (IRB) under the application number HUM00168533.

2.2 | Ultrasound imaging and measures

The ultrasound equipment setups and the scanning procedures were performed by two experienced investigators (HC and OK) (Chan, Sinjab, et al., 2017; Chan, Wang, et al., 2017). Three distinct sites, the premolar, molar, and retromolar sites, were selected for imaging (Figure 1). The initial scan was performed at the premolar site, identified in relation to the mental foramen, followed by the molar site, measured 20 mm from the premolar site, and lastly the retromolar site, 15 mm posterior to the molar site. The distances were gauged with a periodontal probe (University of North Carolina [UNC] Probe, Hu-Friedy) accurate to the nearest 1 mm. The ultrasound probe (L8-25; Zonare/Mindray) was placed at each selected site to obtain a cross-sectional image in DICOM format. The built-in function in the ultrasound device for spatial compounding was selected to obtain well-resolved images (ZS3 Zonare/Mindray). Acoustic coupling was achieved with the application of ultrasound gel (Aquasonic, Parker Inc.) and the use of a gel-based stand-off pad (Parker Inc.).

The captured ultrasound images were read with a commercially available software package (Osirix) to obtain thickness measurements of the mucosal, mylohyoid, and the lingual nerve. At each site, the thickness of the mucosa and the mylohyoid muscle were measured at two distinct locations, 5 and 10 mm distances lingual to the muscle attachment to the mandible. In addition to the stated measurements, at the retromolar site, the lingual nerve diameter was also obtained. All measurements were carried out by a single calibrated examiner (SB) with a built-in device caliper.
accurate to 0.01 mm. The examiner calibration was performed, by measuring 10 random samples by the senior investigator (HC) and the assessment of accuracy and reproducibility in measurements by the chosen examiner (SB) to reach an agreement value of at least 0.86.

2.3 Biopsy sample collection and measurements

Immediately after the ultrasound images were captured from each cadaver head, a biopsy sample from the same imaged sites was collected by an operator with expertise in cadaveric tissue handling and biopsy collection (SN) from the University of Michigan Anatomical Department of the Medical School. From each site, two samples, the mucosa and muscle tissues, were carefully collected. From the retromolar area, a cross-sectional slice of the lingual nerve was also obtained (as imaged with the ultrasound). All collected samples were promptly placed in 10% formalin and sent to the Histology Core at the University of Michigan Health System, Department of Pathology, Immunohistochemistry Laboratory, where they were embedded in paraffin, and sectioned to three 5 micron-thick slices at every 5 mm interval from the attachment, as specifically marked at the harvesting procedure using a tissue coloring marker. Subsequently, all samples were stained with hematoxylin–eosin (H&E).

The specimens were viewed using an E800 Microscope (Nikon Instruments Inc.) with a 2× objective to perform the measurements. Images were captured using a CoolSNAP EZ camera (Photometrics) and saved using a software (NIS-Elements Advanced, Nikon Corporation). To obtain the thickness measurements, each sample was measured at every third of the total sample length and then averaged to obtain the measurement representative of that slide. This was performed for the collected samples of the mucosa and muscle. While for the nerve measurements, the diameter of each sample was measured twice in a way that the two measurements would be perpendicular to one another and then averaged to obtain the cross-sectional (diameter) thickness of that nerve. All measurements were performed by a single calibrated examiner (SB). The software was able to conduct measurements with an accuracy 0.001 mm. The examiner calibration was performed prior to initiation of the measurement by randomly selecting 10 samples for measurement by the senior investigator (HC) and assessment of precision of the chosen examiner (SB) for reaching an agreement value of at least 0.86.

2.4 Clinical feasibility of imaging the lingual nerve

To assess the feasibility of imaging the lingual nerve with ultrasound, the second part of this study consisted of recruitment of healthy adult patients. The live human investigation part of the current study was approved by the Institutional Review Board for Human Studies (HUM00139630). The study was conducted at the Graduate Periodontal Clinic, Department of Periodontology and Oral Medicine, University of Michigan. It was conducted according to the principles embodied in the Helsinki Declaration of 1975, as revised in 2000 for biomedical research involving human subjects. The device setups and the scanning protocol followed the above-mentioned methods. One experienced examiner (HC) performed the scanning of the lingual nerve; while the other examiner (OK) operated the scanning machine. Acquired ultrasound images were saved
in DICOM files and interpreted with the same commercially available software (Osirix). The lingual nerve dimension was measured with the built-in caliper accurate to 0.01 mm.

### 2.5 Data management and statistical analysis

All recorded measurements were entered into a spreadsheet and checked for entry errors. For cadaverous data, descriptive statistics were used for qualitative presentation of the ultrasound and histology measurements by computation of means and standard deviations. To test the presence of statistically significant differences among the two modes of measurements, independent *t* tests were utilized and a *p* value threshold of .05 was set for significance. For live human data, the ultrasound lingual nerve dimension was measured, presented as the mean and standard deviation. All analyses were conducted in Rstudio (Rstudio version 1.1.383; RStudio, Inc.) for Macintosh. Inter-examiner reliability calibration tests were performed with the DescTools package (Signorell, 2019).

### 3 RESULTS

The lingual anatomical structures of nine human cadaver heads were imaged using ultrasound. Biopsy samples corresponding to the imaged sites were also successfully collected from every site. The means and standard deviations of obtained measurements from the ultrasound and histology were summarized in Table 1.

### 3.1 Imaging interpretation

The mylohyoid is a hypoechoic (dark) band with relatively uniform thickness along its length (Figure 2—left). Within it is hyperechoic (white) strips. It attaches to the mandible at one end and extends apically and lingually toward the tongue. Above it is the mucosal layer and the sublingual space, containing the sublingual gland; below it is the submandibular space. The lingual nerve has its characteristic hyperechoic continuous bundles of neuronal fascicles separated from surrounding hypoechoic connective tissue (Figure 2—right). Anatomically, it lies above the mylohyoid muscle at the retromolar area. In this case, it is located superficially, just below the mucosal layer. Figure 3 depicts representative histologic images of the lingual mucosa, mylohyoid muscle, and lingual nerve.

### 3.2 Dimension comparisons

The overall mean ultrasound mucosal thickness was 1.45 ± 0.49 mm at 5 mm distance to the muscle attachment (1.44 mm in the premolar, 1.31 in the molar, and 1.58 in the retromolar region), and 1.54 ± 0.48 mm at the 10 mm distance to the attachment (1.46 mm in the premolar, 1.35 in the molar, and 1.76 mm in the retromolar region). The corresponding histologic mucosal thickness at the respective sites averaged to 1.39 ± 0.51 mm at the 5 mm distance to the attachment (1.46 mm in the premolar, 1.30 in the molar, and 1.44 mm in the retromolar region), and 1.37 ± 0.46 mm at 10 mm distance to the attachment (1.28, 1.16, and 1.61 mm in the premolar, molar, and retromolar region, respectively). The differences among the obtained values from the ultrasound compared to the biopsied samples did not reach statistical significance (*p* > .05) when tested as a whole, and among each respective region.

In regard to the mylohyoid muscle thickness measurements, the overall ultrasound mean value was 2.31 ± 0.56 at 5 mm from the attachment (2.03 mm for the premolar area, 2.59 mm at the molar, and 2.22 mm at the retromolar region), and 2.46 ± 0.56 mm at 10 mm distance to the attachment (2.28 mm at the premolar, 2.75 mm at the molar, and 2.33 mm at the retromolar area). The corresponding histologic values have an overall mean of 2.25 ± 0.47 mm at the 5 mm distance (2.04 mm for premolar, 2.46 for the molar, and 2.22 for the retromolar regions), and 2.36 ± 0.50 mm at the 10 mm distance to the attachment (2.23, 2.56, and 2.22 mm at the premolar, molar, and retromolar regions, respectively). Again, there was no

<table>
<thead>
<tr>
<th>Structure</th>
<th>Site</th>
<th>5 mm</th>
<th>10 mm</th>
<th><strong>p Value</strong></th>
<th>5 mm</th>
<th>10 mm</th>
<th><strong>p Value</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Ultrasound</td>
<td>Histology</td>
<td></td>
<td>Ultrasound</td>
<td>Histology</td>
<td></td>
</tr>
<tr>
<td>Mucosa</td>
<td>Premolar</td>
<td>1.448 ± 0.402</td>
<td>1.460 ± 0.441</td>
<td>.957</td>
<td>1.467 ± 0.505</td>
<td>1.280 ± 0.501</td>
<td>.469</td>
</tr>
<tr>
<td></td>
<td>Molar</td>
<td>1.317 ± 0.536</td>
<td>1.301 ± 0.58</td>
<td>.959</td>
<td>1.358 ± 0.391</td>
<td>1.168 ± 0.391</td>
<td>.391</td>
</tr>
<tr>
<td></td>
<td>Retromolar</td>
<td>1.581 ± 0.531</td>
<td>1.440 ± 0.512</td>
<td>.531</td>
<td>1.761 ± 0.512</td>
<td>1.619 ± 0.512</td>
<td>.521</td>
</tr>
<tr>
<td></td>
<td>Overall</td>
<td>1.453 ± 0.496</td>
<td>1.398 ± 0.507</td>
<td></td>
<td>1.541 ± 0.489</td>
<td>1.370 ± 0.496</td>
<td></td>
</tr>
<tr>
<td>Muscle</td>
<td>Premolar</td>
<td>2.037 ± 0.449</td>
<td>2.044 ± 0.441</td>
<td>.976</td>
<td>2.282 ± 0.631</td>
<td>2.231 ± 0.579</td>
<td>.862</td>
</tr>
<tr>
<td></td>
<td>Molar</td>
<td>2.590 ± 0.646</td>
<td>2.461 ± 0.466</td>
<td>.598</td>
<td>2.751 ± 0.538</td>
<td>2.567 ± 0.472</td>
<td>.414</td>
</tr>
<tr>
<td></td>
<td>Retromolar</td>
<td>2.227 ± 0.471</td>
<td>2.225 ± 0.475</td>
<td>.827</td>
<td>2.334 ± 0.471</td>
<td>2.227 ± 0.455</td>
<td>.781</td>
</tr>
<tr>
<td></td>
<td>Overall</td>
<td>2.316 ± 0.564</td>
<td>2.256 ± 0.478</td>
<td></td>
<td>2.467 ± 0.568</td>
<td>2.367 ± 0.505</td>
<td></td>
</tr>
<tr>
<td>Nerve</td>
<td>Retromolar</td>
<td>2.386 ± 0.441</td>
<td>2.432 ± 0.423</td>
<td>.785</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Note: All reported values are in mm ± standard deviation. *p* values are from independent *t* tests.
significant differences between ultrasound and histology in any of the obtained mylohyoid muscle thickness measurements ($p > .05$ for all comparisons).

Lastly, as for the measurements of the lingual nerve diameter, similar values were obtained via ultrasound (2.38 ± 0.44 mm) and histologic assessments (2.43 ± 0.42 mm) without statistically significant difference ($p > .05$).

### 3.3 Outcomes of live human scans

A total of 19 individuals, corresponding to 30 sites (18 on the right, and 12 on the left) were available for ultrasound imaging of the lingual nerve in the retromolar area. The mean diameter was 2.11 ± 0.35 mm (ranging from 1.49 to 3.14 mm). The mean values between the right and left sides were not significantly different.
(2.02 ± 0.25 mm vs. 1.99 ± 0.47 mm, *p* = .79). Figure 4 displays a cross-sectional image of a lingual nerve in a live human participant, and Figure 5 shows the distribution of the lingual nerve diameter.

**4 | DISCUSSION**

For the first time in the literature, ultrasound was found accurate in imaging the human mucosal, mylohyoid muscle, and lingual nerve because of (a) dimensional consistency with histology and the literature and (b) imaging characters in accordance with other nerves and muscles. Our study concluded ultrasound and histologic dimensions of the abovementioned structures are not statistically different. A recent study (Kikuta, Iwanaga, Kusukawa, & Tubbs, 2019) showed the mean diameter of the lingual nerve is 2.2 mm (range 1.61–2.95 mm), which is in agreement with our measurements on human cadavers as well as live humans. The sheath-like hyperechoic appearance of the lingual nerve on ultrasound images is consistent with nerve fascicles. The hypoechoic band representing mylohyoid muscle is also characteristic of muscles in the rest of the body (Engel, Harn, & Cohen, 1987; Koolstra & van Eijden, 1999).

Anatomy of the mandibular lingual region has become more important than ever because of the popularity of performing ridge augmentation for implant placement in this region. Lingual flap releasing for achieving primary wound closure requires detachment of the lingual mucosa from the underlying mylohyoid muscle. Knowledge about the lingual mucosa thickness, sublingual salivary glands, and mylohyoid muscle attachment is key to successful lingual flap management. This study showed that the mean mucosal thickness is approximately 1.5 mm. Histology also showed that the submucosa is mainly composed of adipose tissue. The thin dimension and loose tissue consistency reaffirmed difficulties in managing the lingual flap clinically. The mean mylohyoid muscle dimension is approximately 2.5 mm. What may be more important is the location of the muscle attachment because it determines the degree of difficulty in releasing the lingual flap. When the attachment is high, that is, closer to the alveolar crest, flap releasing is more challenging and vice versa. Knowledge about the mylohyoid muscle location and lingual mucosa features pre-surgery could be beneficial to assess the risk of wound opening after ridge augmentation.

The lingual nerve is a branch of the mandibular nerve, providing sensory innervation to the mucous membranes of the anterior two-thirds of the tongue and the lingual tissues. After entering the oral cavity, it is located at a mean distance of 3 mm apical to the osseous crest and 2 mm horizontally from the lingual cortical plate in the third molar area. (Behnia, Kheradvar, & Shahrokhi, 2000). Nevertheless, the nerve may be situated at or above the crest of bone in 15–20% cases (Pogrel & Goldman, 2004). Furthermore, 22% of the time the nerve may contact the lingual cortical plate (Behnia et al., 2000). Once passing the 3rd molar, it travels mesially, apically, and medially toward the tongue. The vertical distance between the nerve and the cementoenamel junction (CEJ) of the second molar, first molar, and the second premolar was 9.6, 13, and 14.8 mm, respectively (Chan et al., 2011). Because it has superficial location in the 3rd molar region, precaution has to be exercised when performing a flap surgery in this area. A 0.6%–2% incidence of lingual nerve injury has been reported following third molar extraction (Bataineh, 2001; Gomes, Vasconcelos, de Oliveira e Silva, & da Silva, 2005; Gulicher & Gerlach, 2001; Hillerup & Stoltze, 2007; Valmaseda-Castellon, Berini-Aytes, & Gay-Escoda, 2000). Ultrasound is an optimal imaging modality for this nerve because it cannot be seen on radiographs. Our group published a proof-of-principle study showing ultrasound can image the intact lingual nerve. (19) The present study with a larger sample size and application in live humans, further confirmed the accuracy of ultrasound in imaging this nerve. Earlier reports of investigating the lingual nerve, while with a different methodology, can also been noted in the literature. Olsen et al. using a slightly larger ultrasound device (25 mm transducer, 10–5 MHz) analyzed the lingual nerve in nine pig cadaveric specimens, in an attempt to correctly identify an intact, partially transected or fully transected injury to the nerve (Olsen et al., 2007). Later on, Al-Amery and colleagues, using ultrasound on previously dissected and harvested lingual nerves from six human cadavers were able to visualize the bur-induced lacerations at the damaged sites (Al-Amery, Ngeow, Nambiar, & Naidu, 2018).

Another clinical indication for locating the lingual nerve is for its block anesthesia. The most common target for local anesthesia of the lingual nerve is the pterygomandibular space. However, inadequate anesthesia of the lingual nerve is common because of unreliable landmarks (Balasubramanian et al., 2017). Exclusive lingual nerve block at the 3rd molar region could be an effective alternative because of the following advantages: (a) easier and closer access, (b) aspiration is not required because of no major vessels in this area, and (c) less chance of post-injection trismus. Visualization of the nerve with ultrasound may improve clinician confidence, increase anesthesia success rate and working time, and reduce injection quantity. Moreover, ultrasound could be a learning tool for dental students to practice lingual nerve anesthesia.
5 | CONCLUSION

This study successfully characterizes important anatomical structures in the lingual mandible, including the mucosa, the mylohyoid muscle, and the lingual nerve with non-invasive, non-radiation, and chairside ultrasound. This novel imaging modality may become a useful tool to evaluate lingual anatomy and assess the risk of developing complications, particularly prior to a surgery.

ACKNOWLEDGEMENTS

The authors would like to thank the body donors and their families, Mr. Dean Mueller, Coordinator of the Anatomical Donations Program for preparing the specimens, Mrs. Alicia Baker, Clinical Coordinator, and Ms. Cynthia Miller, Dental Assistant, for kindly coordinating the experiment. We also give thanks to Kenneth Rieger, Multimedia Designer, the University of Michigan for his effort in the drawing of the illustration.

CONFLICT OF INTEREST

The authors do not have any financial interests, either directly or indirectly, in the products or information listed in the paper.

AUTHOR CONTRIBUTIONS

SB contributed to conception and design, drafted and critically revised the manuscript, performed the ultrasound and histology measurements, and analyzed the data; HLC contributed to conception and design, mentored the project, performed the ultrasound scans, and drafted and critically revised the manuscript; SN performed the biopsy specimens and drafted and critically revised the manuscript; HLW contributed to conception, design, and critically revised the manuscript; OK contributed to conception and design, performed the ultrasound scans, and drafted and critically revised the manuscript. All authors gave final approval and agreed to be accountable for all aspects of the work.

ORCID

Shayan Barootchi https://orcid.org/0000-0002-5347-6577
Hsun-Liang Chan https://orcid.org/0000-0001-5952-0447
Hom-Lay Wang https://orcid.org/0000-0003-4238-1799

REFERENCES


SUPPORTING INFORMATION

Additional supporting information may be found online in the Supporting Information section.