

# Ultra High Resolution Optical Coherence Tomography in Laryngology

Aspects of development  
2 parts: background and future

Mette Pedersen MD Prof. Hon. IBC int ENT PhD.

<http://www.mpedersen.org>

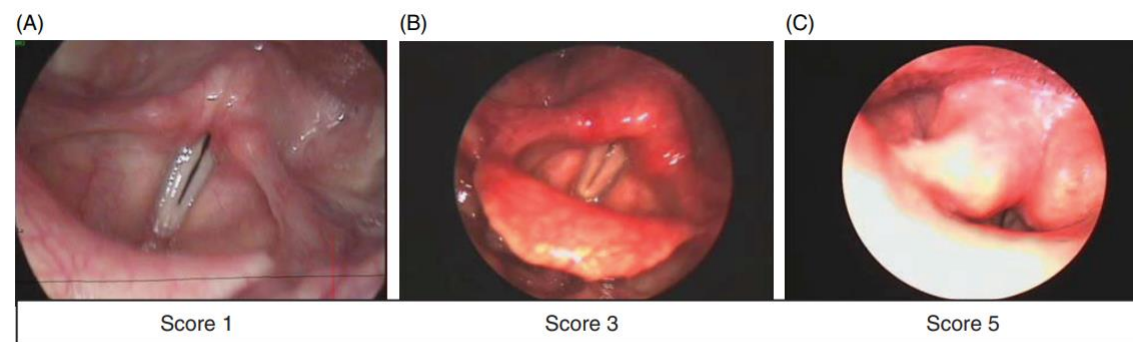
POSITION PAPER

# Optical coherence tomography in the laryngeal arytenoid mucosa for documentation of pharmacological treatments and genetic aspects: a protocol

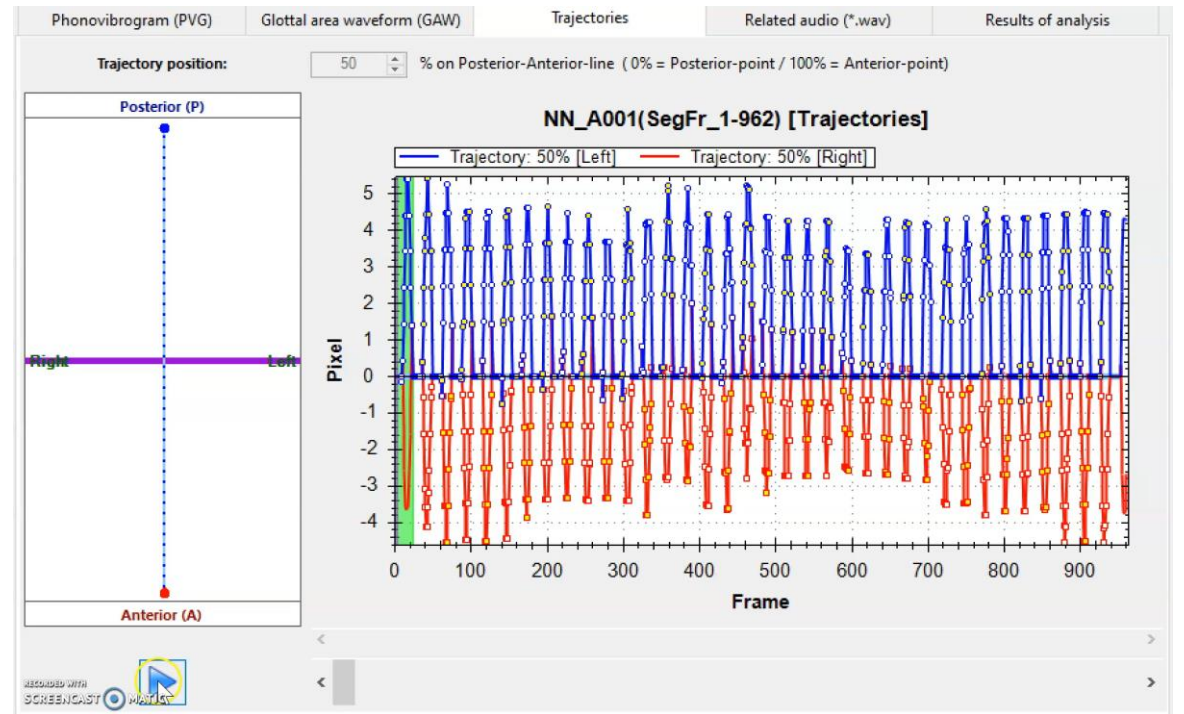
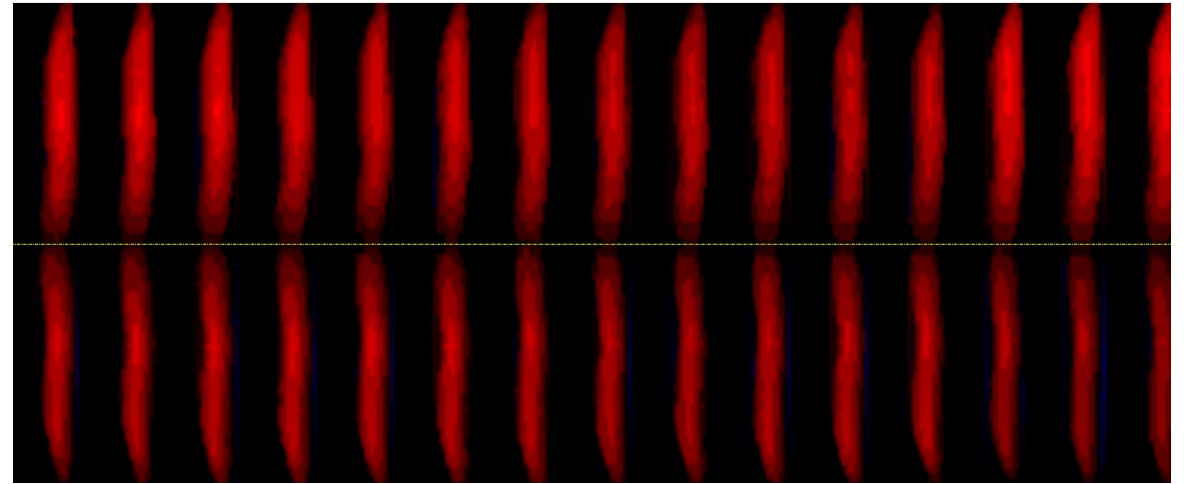
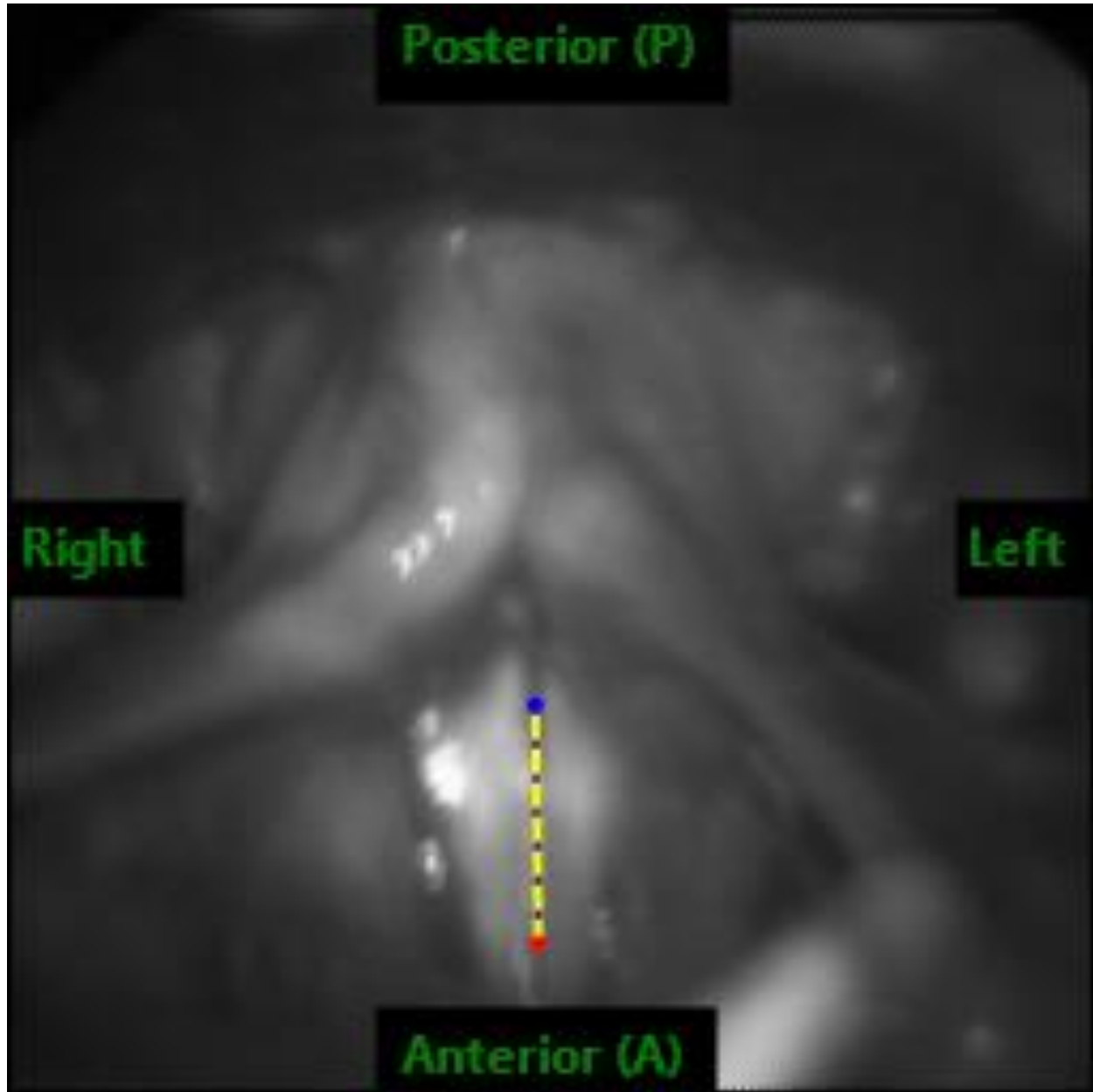
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*Fig. 1.* High-speed film scores with 4,000 pictures/sec of the larynx including the arytenoid regions. Score 1 indicates a normal arytenoid region and normal vocal folds. Score 3 indicates a moderate oedema of the arytenoid region and normal vocal folds. Score 5 indicates almost total closure of the larynx due to arytenoid oedema (4).



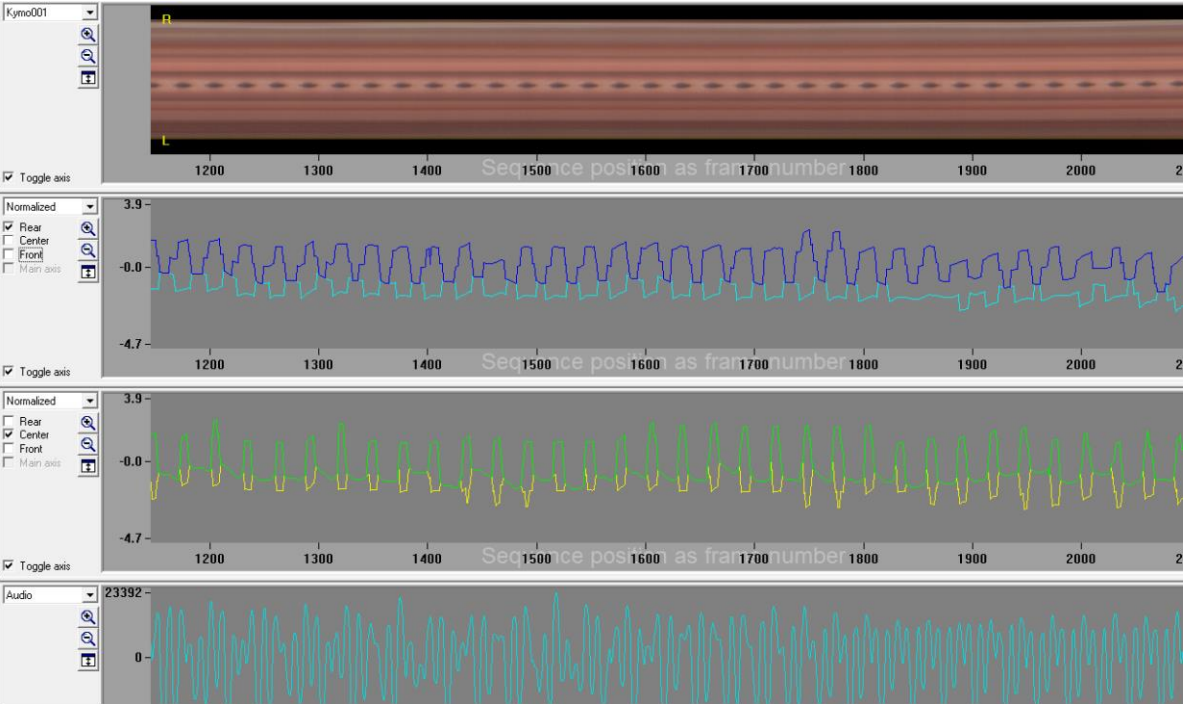
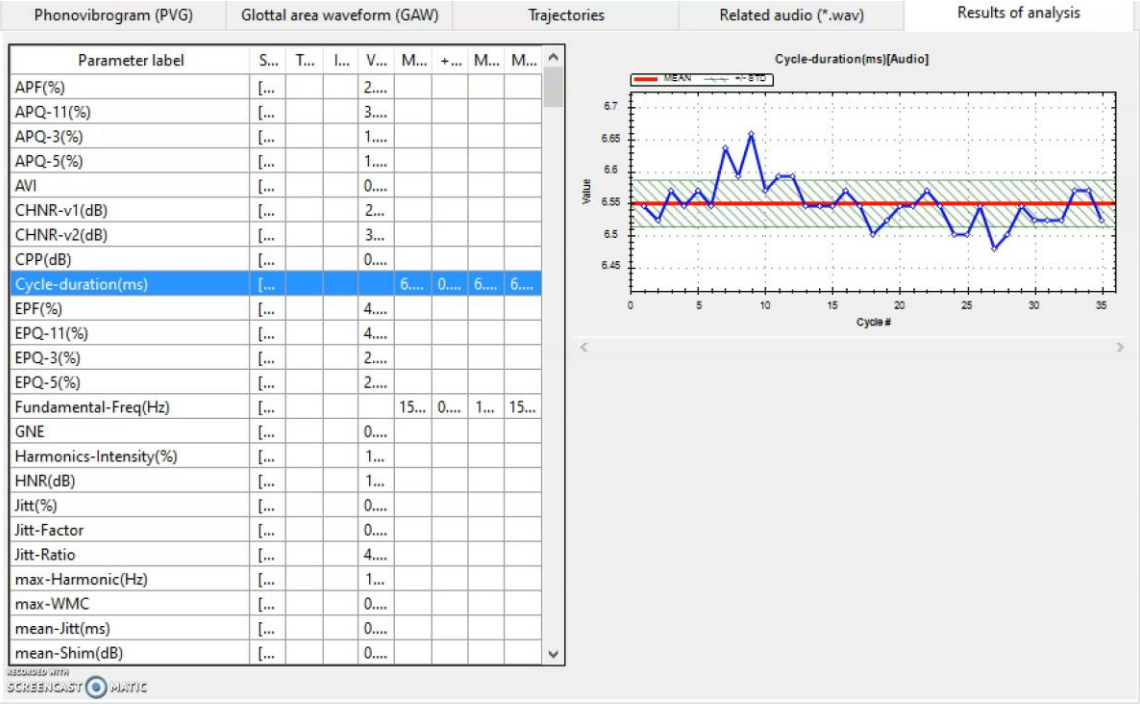


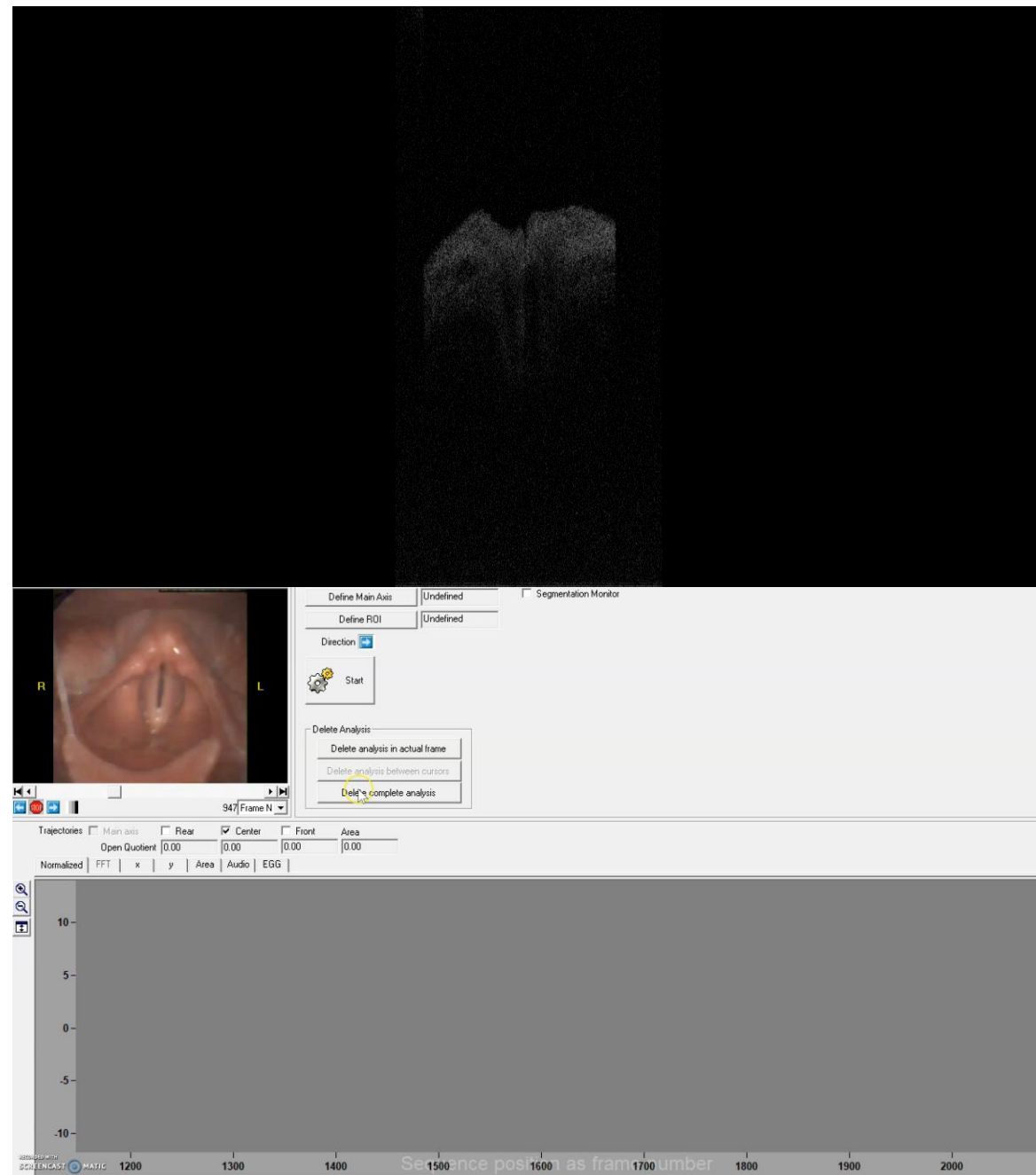
Figure 1 : OCT hand piece.

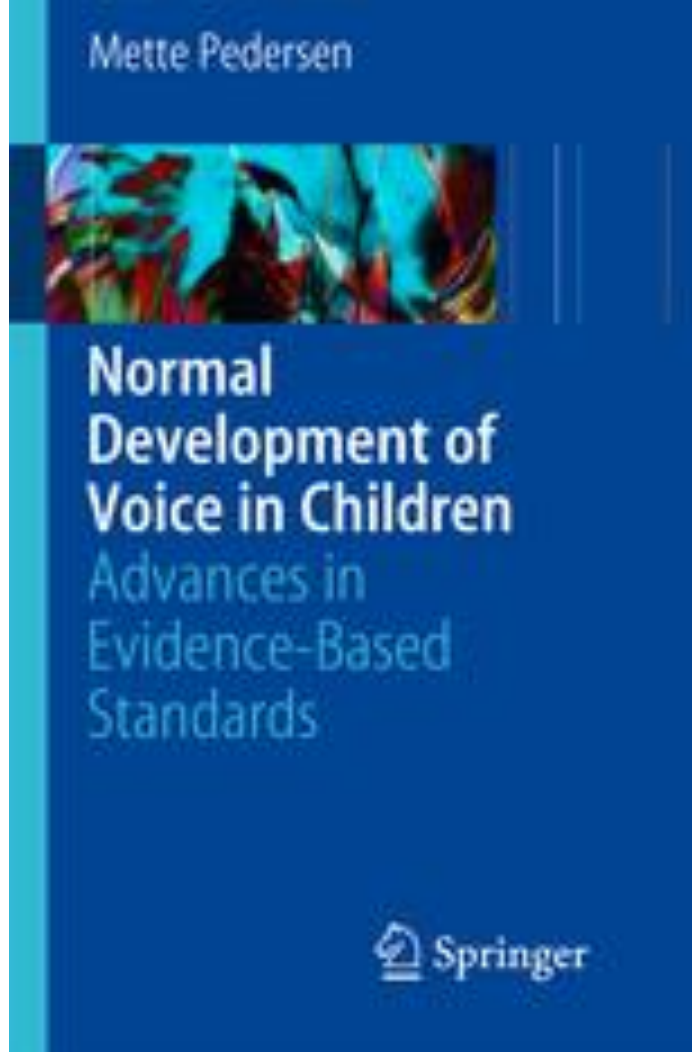
From: In vivo cross-sectional imaging of the phonating larynx using long-range Doppler optical coherence tomography



The OCT device mounted concurrently with a rigid laryngoscope.

Carolyn A. Coughlan, Li-dek Chou, Joseph C. Jing, Jason J. Chen, Swathi Rangarajan, Theodore H. Chang, Giriraj K. Sharma, Kyoungrai Cho, Donghoon Lee, Julie A. Goddard, Zhongping Chen & Brian J. F. Wong





# Die biologische Entwicklung der Stimme in der Pubertät

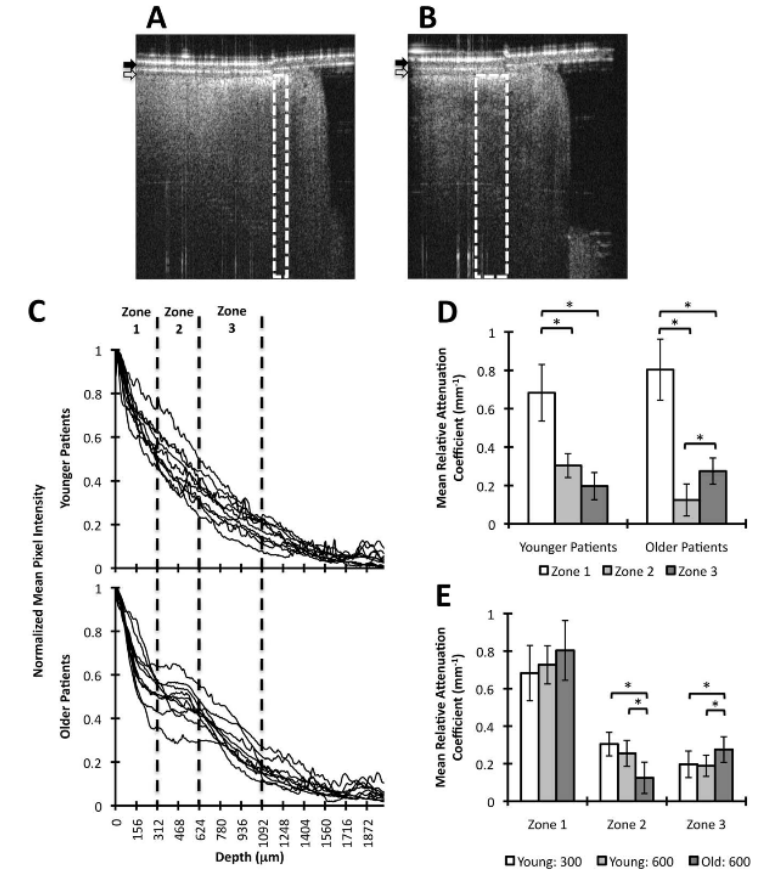
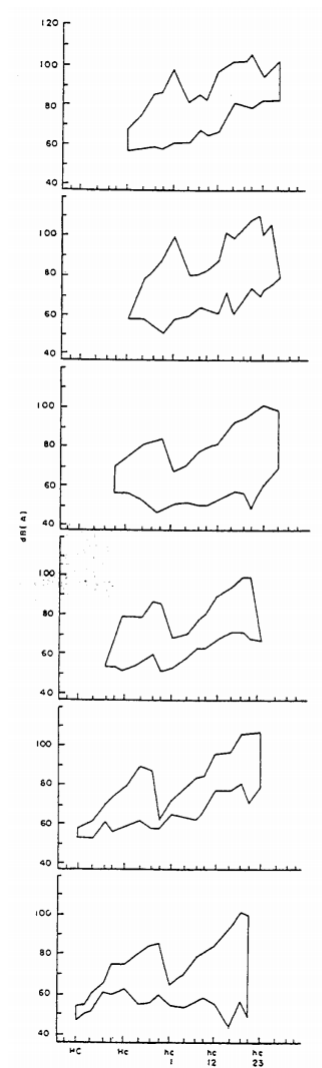


Fig. 3. Quantitative analysis of patient A-lines reveals distinct patterns in relative attenuation coefficients in young and old patients. (A) Example image of a 2-year-old true vocal fold edge. (B) Example image of a 16-year-old true vocal fold edge. Representative regions of interest with a height of 2000  $\mu\text{m}$  (white boxes) are shown. The endosheath (black arrow) and epithelium (white arrow) are indicated. (C) Young and old patient A-lines. Demarcation of zones 1, 2, and 3 are shown in both populations. (D) Mean relative attenuation coefficients for zones 1, 2, and 3 for each patient group. Zone 1 was distinct relative to zones 2 and 3 for each group. Zone 2 was distinct from zone 3 in older patients but not in younger patients. (E) Placing the region of interest series 600  $\mu\text{m}$  inferior to the vocal fold edge (young: 600) did not significantly change relative attenuation coefficients when compared to analysis by placing the region of interest series 300  $\mu\text{m}$  inferior to the vocal fold edge (young: 300). The relative attenuation coefficients between groups young: 300 and young: 600 were indistinct and had the exact same pattern of statistical significance when compared to older patients (old: 600). \* $P < .05$ .

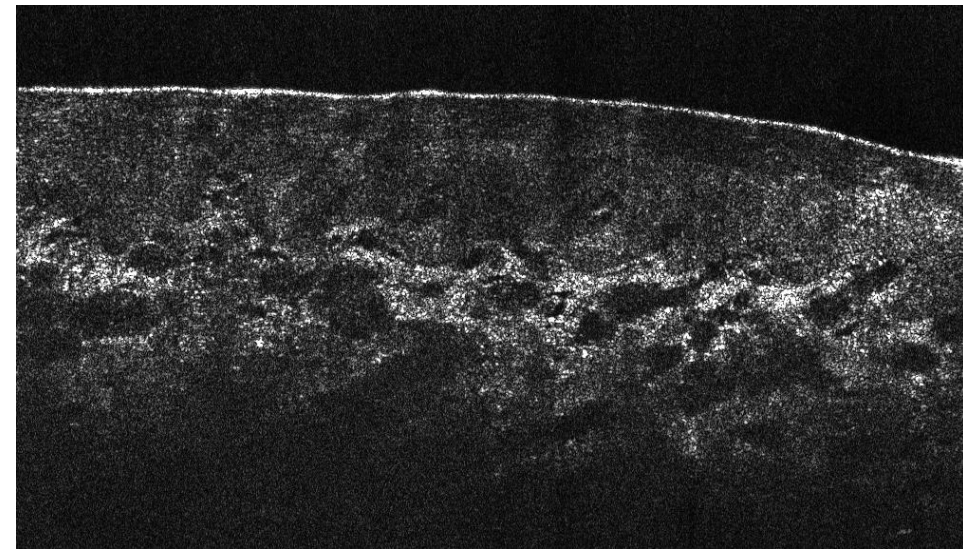
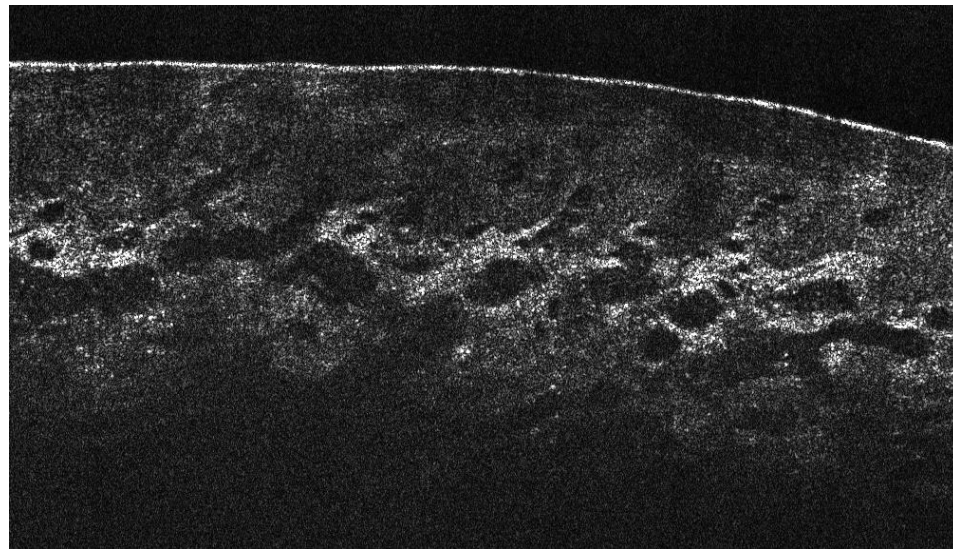
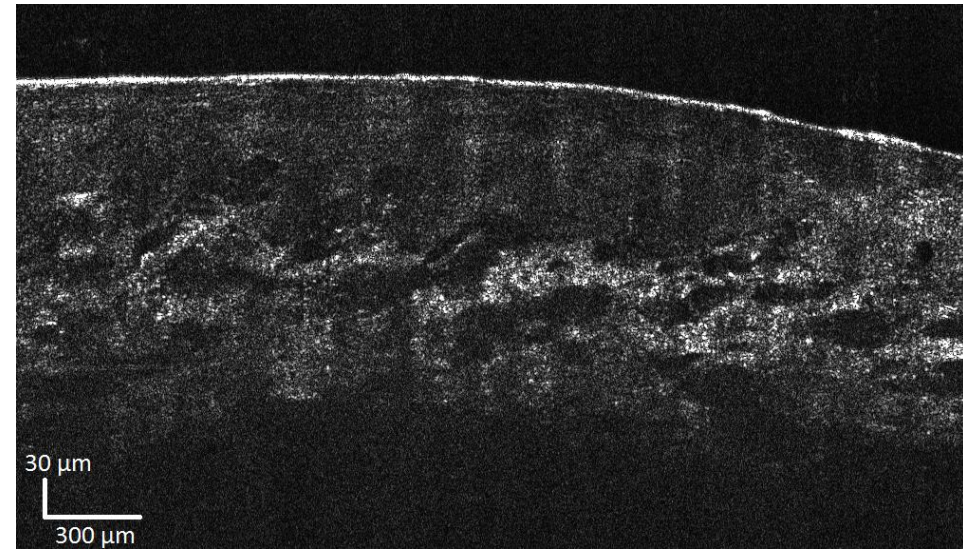


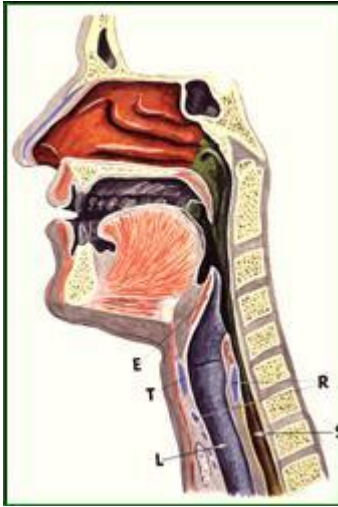
Future

ULTRAHIGH RESOLUTION OPTICAL  
COHERENCE TOMOGRAPHY FOR DETECTING  
TISSUE ABNORMALITIES OF THE ORAL AND  
LARYNGEAL MUCOSA: A PRELIMINARY STUDY.

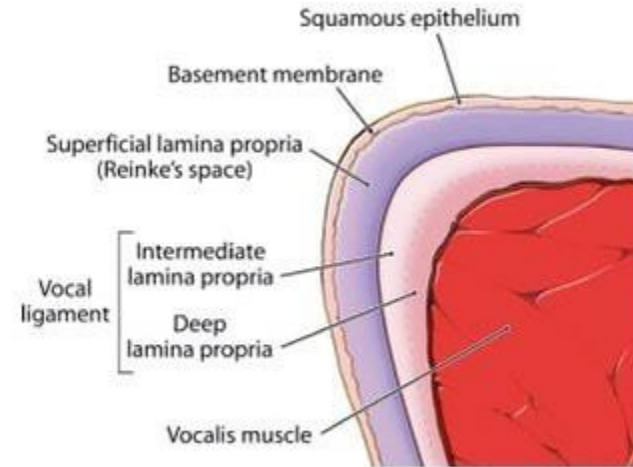
Niels Møller Israelsen, Mikkel Jensen, Anders  
Overgård Jønsson, Mette Pedersen.

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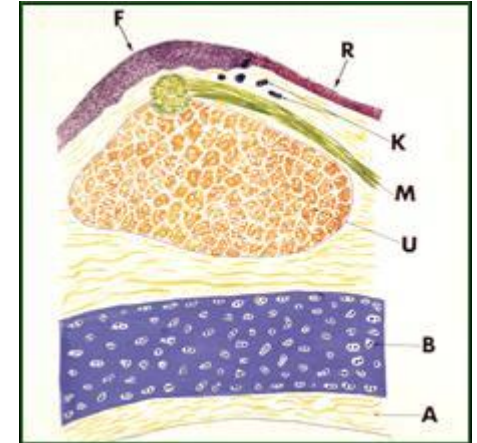




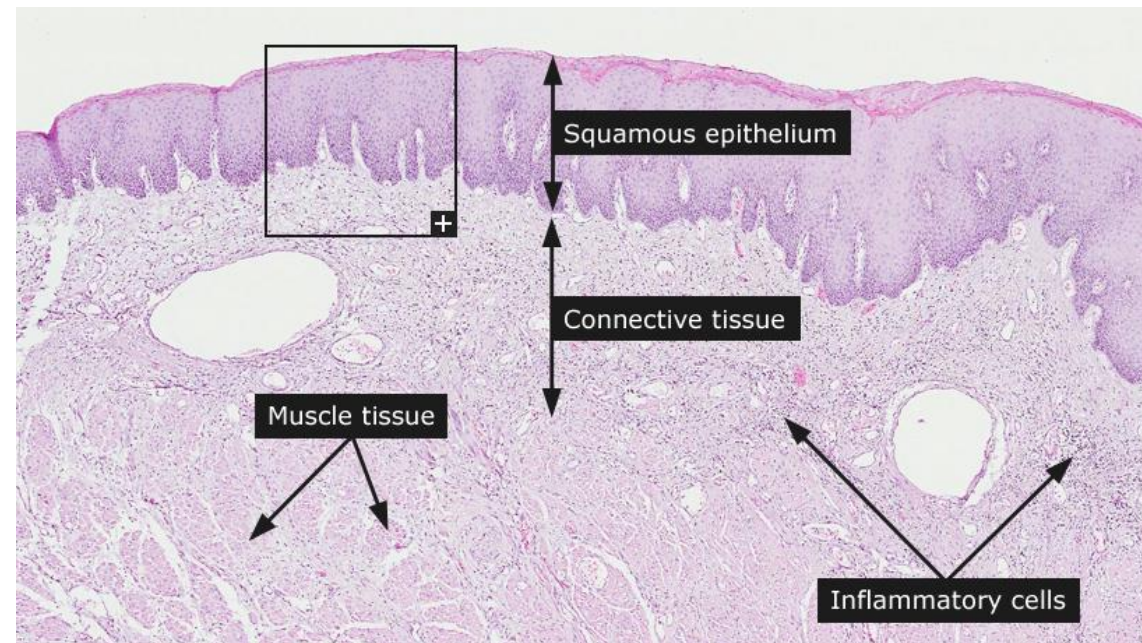
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anatomionline.dk



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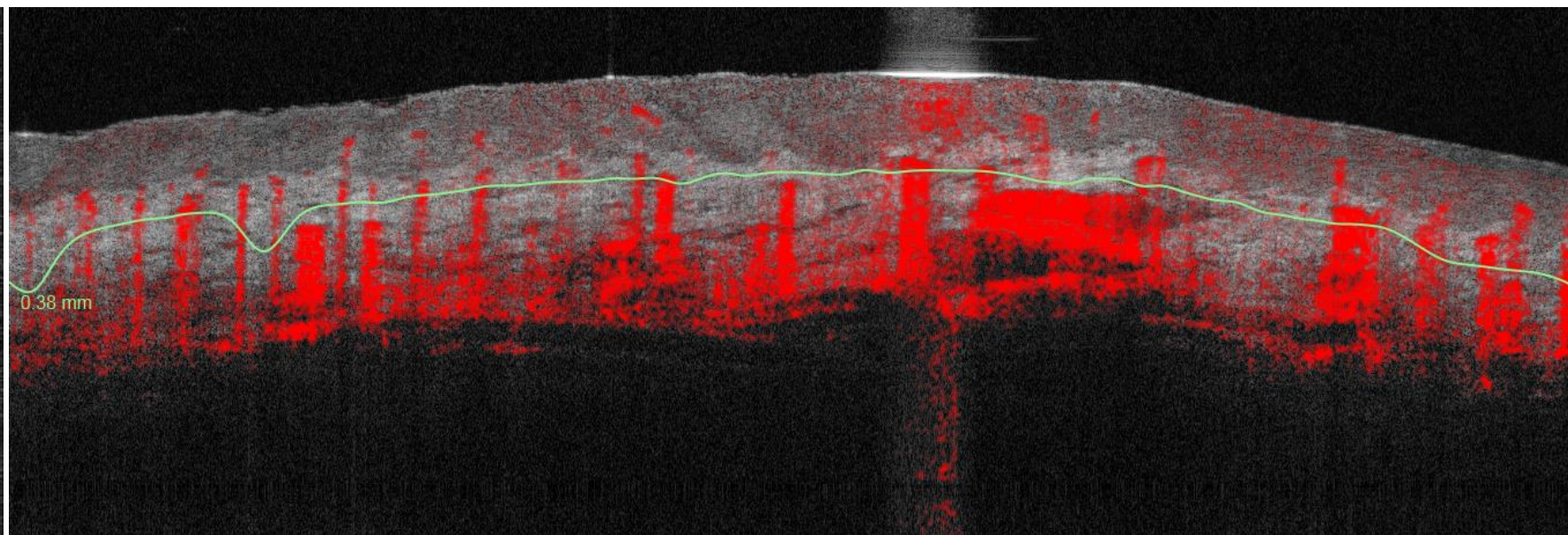
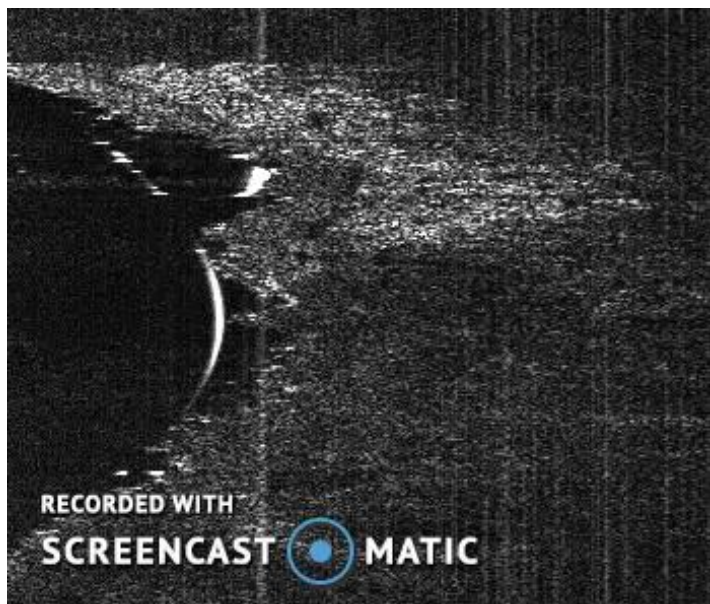
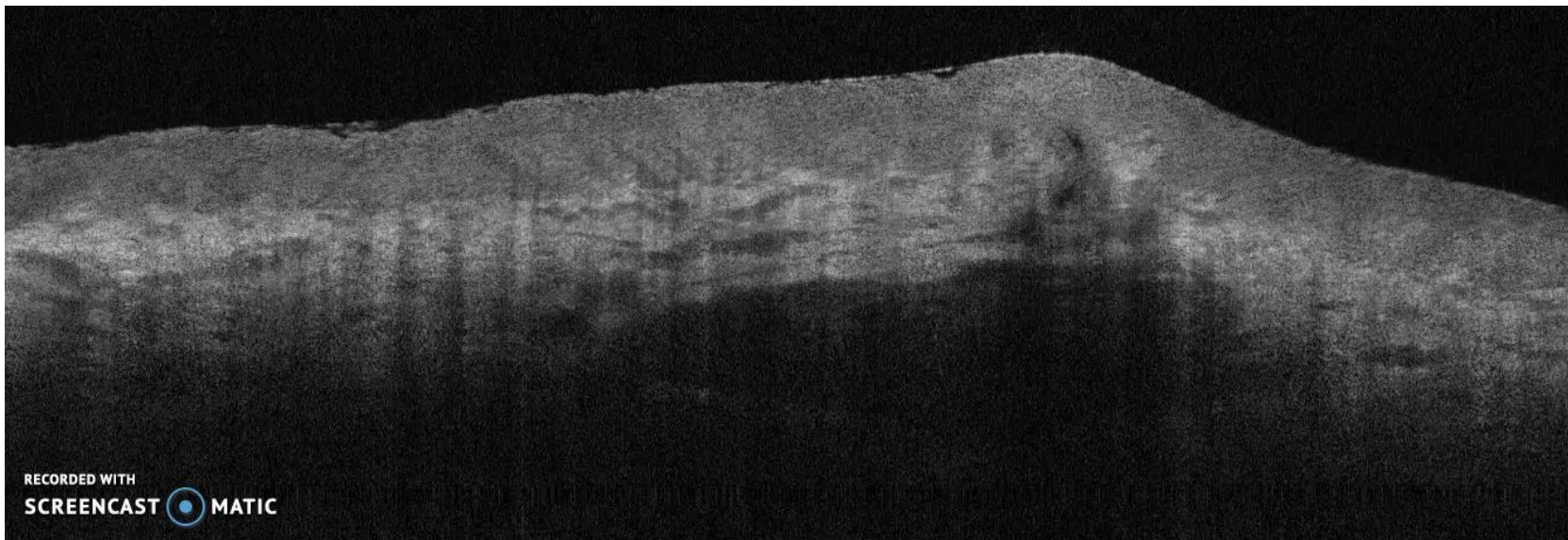


Source:  
anatomionline.dk



Biopsy of mucosal tissue in the mouth

Vivosight billeder 6x6 mm



To finish we present 2 abstracts to show you that we are not the only ones focusing on improving tissue diagnostics in the larynx.

14 March 2018

# WIDE-FIELD AND LONG-RANGING-DEPTH OPTICAL COHERENCE TOMOGRAPHY MICROANGIOGRAPHY OF HUMAN ORAL MUCOSA

(Conference Presentation)

[Wei Wei](#); [Woo June Choi](#); [Shaojie Men](#); [Shaozhen Song](#); [Ruikang K. Wang](#)

Univ. of Washington (United States)

Novel use of long ranging swept source optical coherence tomography (SS-OCT) system with unprecedented **imaging field of view of 1600 mm<sup>2</sup>** for anatomic and microvascular imaging of human oral mucosa tissue **in vivo**. This system is further applied to identify and analyze oral mucosa lesions in situ without a need for invasive biopsy. Qualitative assessment of the structure characteristics (i.e. collagen fibrosis, volume of salivary glands, and tissue scattering) during wound healing delineates the anatomical lesion development accompanied with tissue inflammation. **Quantitative assessment of the vasculature network** (i.e. capillary loop density and vessel morphological orientations) reveals pathological and nutritional underpinnings of microcirculation for oral lesion recovery. Specifically, the **progression of oral capillary angiogenesis**, indicated by elevations in capillary loop density, occurs **within 12 hours of disease onset and peaks at day 7 thereafter**, which provide invaluable information for the time course of therapeutic treatment. Our demonstration shows potential movement of oral cavity OCT microangiography toward **clinical translations**.

4 March 2019

# OPTICAL BIOPSY OF VOCAL FOLDS DURING PHONATION USING PARALLEL OCT (Conference Presentation)

[Gopi N. Maguluri](#); [Daryush D. Mehta](#); [James B. Kobler](#); [Jesung Park](#); [Nicusor V. Iftimia](#)

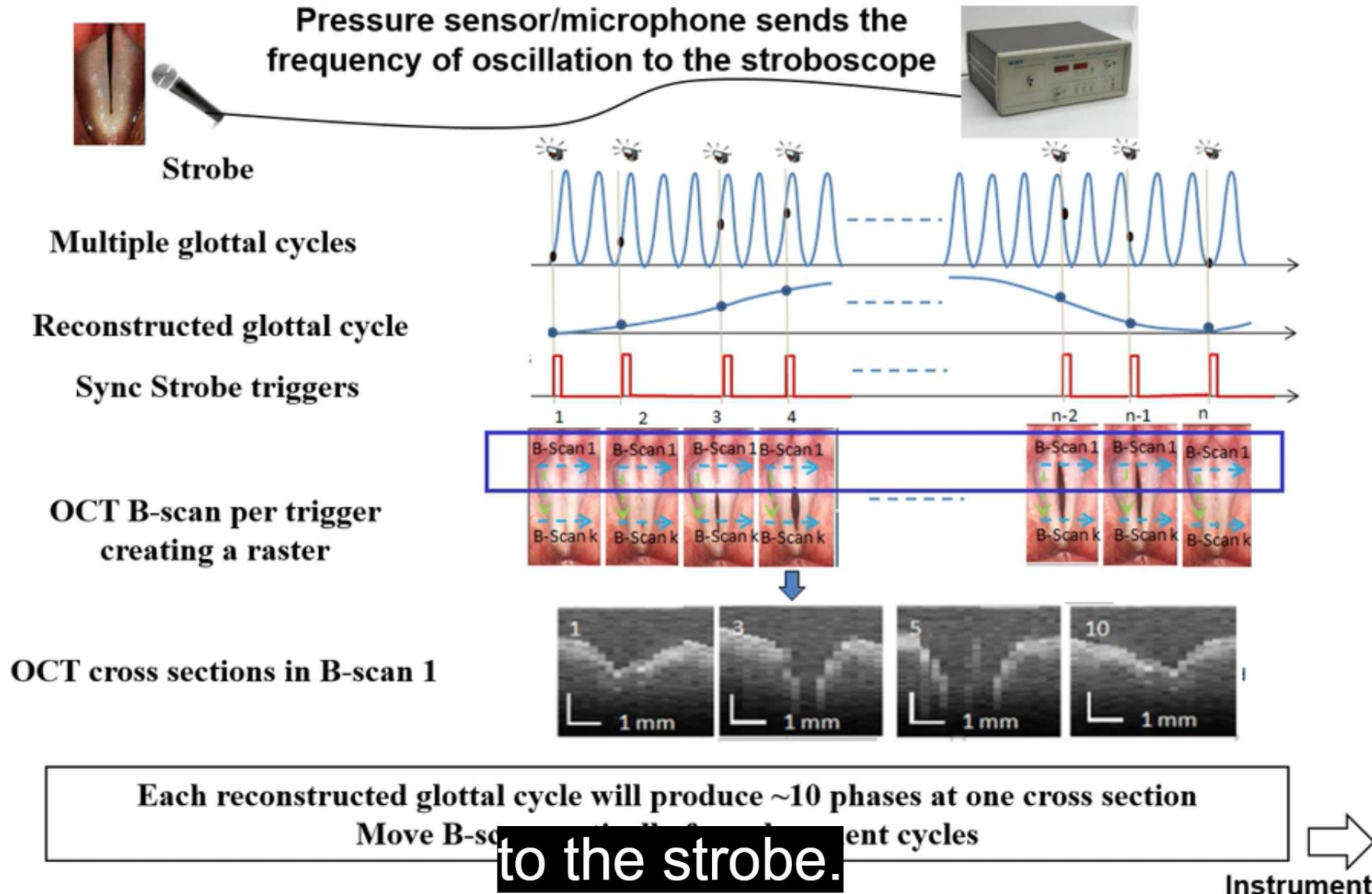
[Author Affiliations](#) -Gopi N. Maguluri,<sup>1</sup> Daryush D. Mehta,<sup>2</sup> James B. Kobler,<sup>2</sup> Jesung Park,<sup>1</sup> Nicusor V. Iftimia<sup>1</sup>

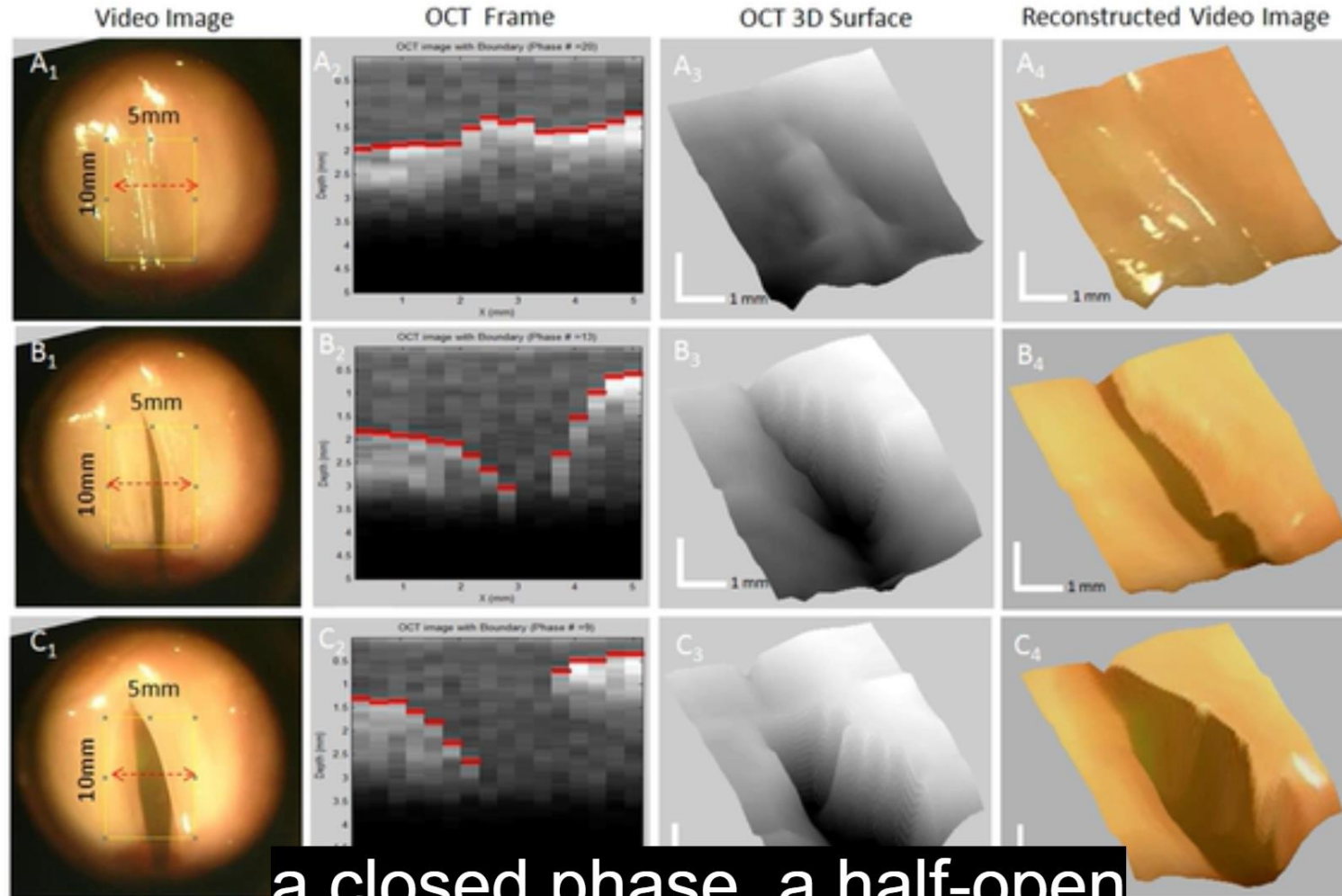
<sup>1</sup>Physical Sciences Inc. (United States)

<sup>2</sup>Ctr. for Laryngeal and Voice Rehabilitation, Massachusetts General Hospital (United States)

There is a constant need of an endoscopic imaging tool in voice clinics that can directly capture the three-dimensional (3D) surface motion of the vocal folds in real time as patients phonate. To address the need, we will present a Parallel OCT/Video Stroboscopy imager that will combine parallel swept source OCT technique with Video Stroboscopy to provide a real time display of the vocal folds in all three axial dimensions during phonation. The results will yield cross sections (B-Scans) with ~16 co-linear sampling locations spread over ~5mm on a **phonating ex-vivo calf larynx** showing fluid periodic cyclic motion of the vocal folds (~A-scan rate) in real time enabled by Parallel OCT approach. We will also capture **Video Stroboscopy images in sync with Parallel OCT B-scans** validating the real time cross sectional probing of Parallel OCT/Video Stroboscopy imager.

# OCT-VS Synchronization





a closed phase, a half-open  
phase, and a fully-open phase.

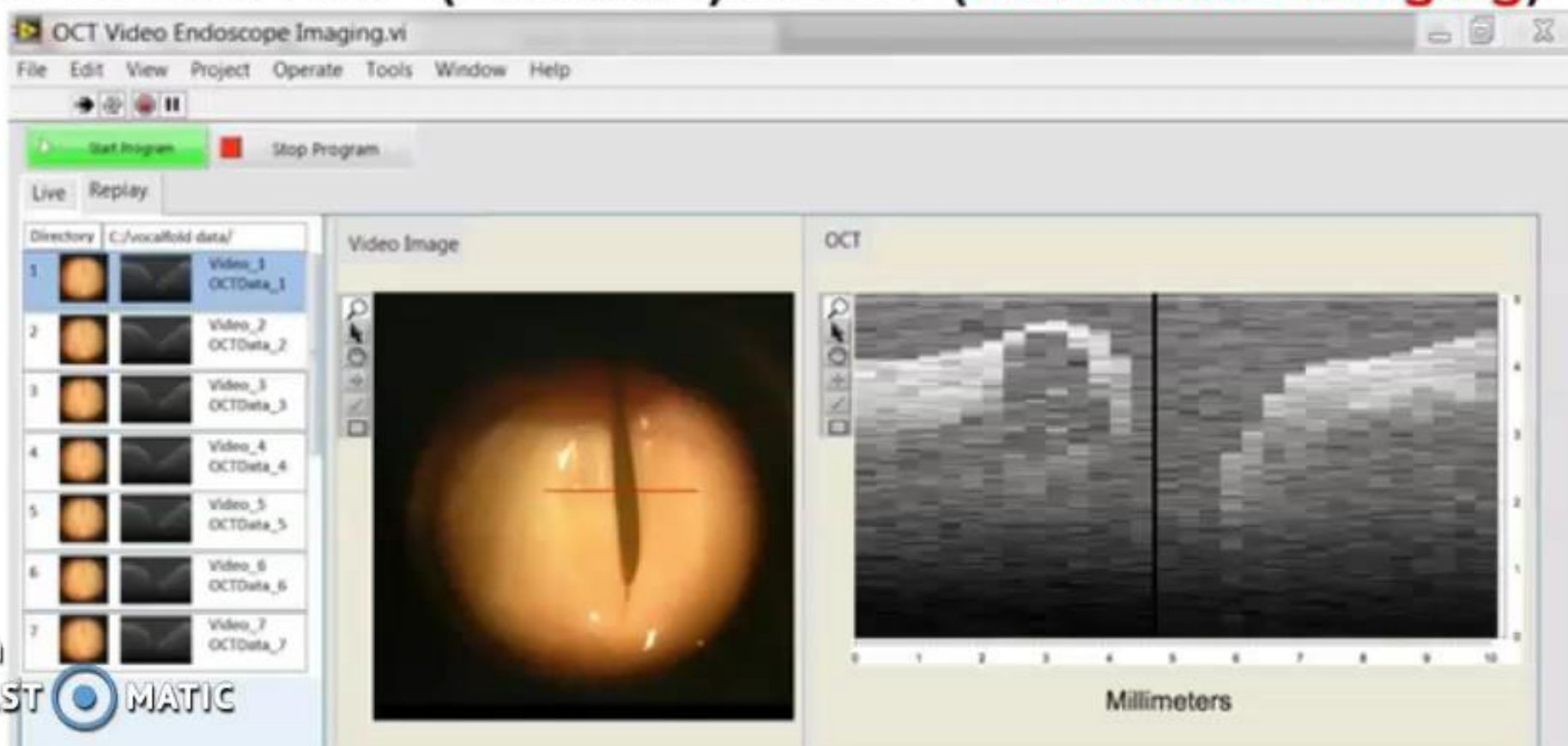
Video

Superimposed  
T 3D

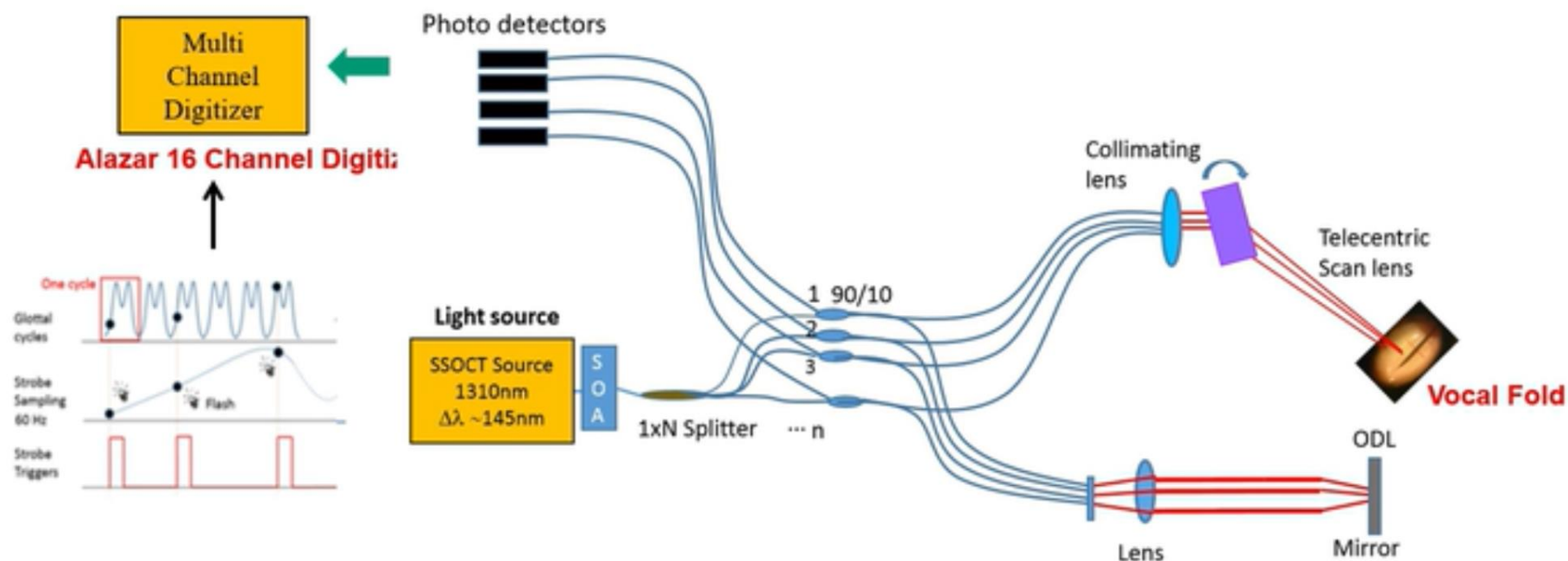
Pilot Data

# Pilot Data OCT+VS (Dynamic)

Restylane injected under the surface can be clearly seen as circular zone (~1.25mm) in OCT (**Subsurface imaging**)



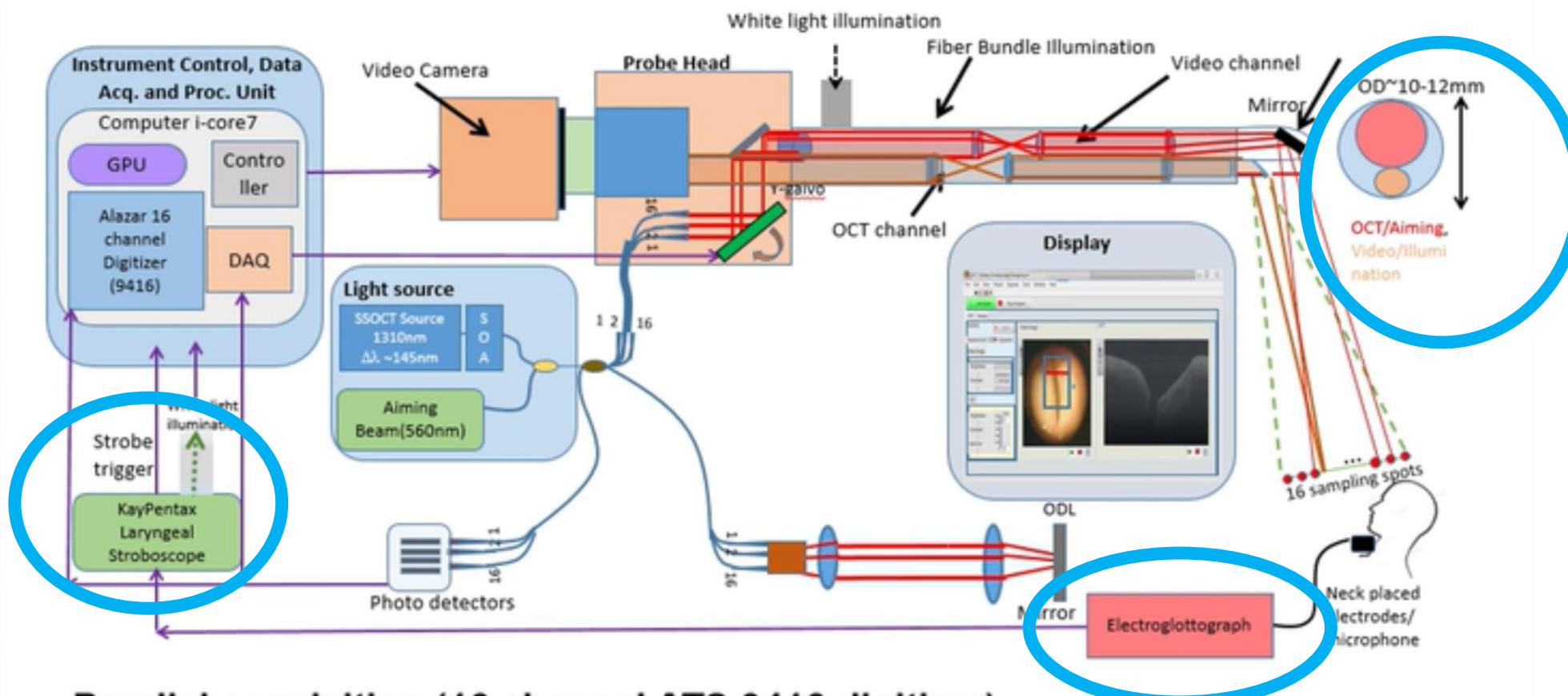
- Address with parallel OCT approach
- No galvanometer for B-scan
- Instead, 16 beamlets spanning ~5-10mm; each an individual OCT system



- Source: Santec 20 kHz; (lets us use low cost low speed OCT systems)
- SOA- Reduces the bandwidth but will provide the power needed ~50mW
- 16 channel Acquisition (Alazar at 100MS/sec)

# Proposed pOCT Probe Head

OD ~10mm



- **Parallel acquisition (16 channel ATS 9416 digitizer)**
  - Clinically adaptable (OD 10-12mm); produces fluid mucosal wave motion
  - All beam paths, (OCT, Video, white light illumination and aiming beam within OD
  - Synchronous with VS (video);
- Disadvantage: No morphological mode (No dense sampling but it is not vital)
- pOCT Hardware

- **Designed and developed Gen-1 OCT-VS Imaging instrument**
- **Demonstrated synchronous OCT/VS imaging on ex-vivo vocal folds.**
  - 3D imaging with spatial co-registration and temporal synchronization is demonstrated
  - Dynamic cross-sectional imaging is demonstrated during phonation.
- **Designed next generation probe head using parallel OCT technique to achieve clinical viability**
  - OD ~10-12mm
  - Capability to achieve cross sectional (B-scan) imaging at up to A-scan rate
- **By next year, hope to demonstrate in-vivo imaging**