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Authors

He, Youmin
Jing, Joseph C
Qu, Yueqiao
[et al.](#)

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Spatial mapping of tracheal ciliary beat frequency using real time phase-resolved Doppler spectrally encoded interferometric microscopy

Youmin He^{*}, Joseph C. Jing^{*}, Yueqiao Qu^{*}, Brian J. Wong, Zhongping Chen[†]

Department of Biomedical Engineering, Beckman Laser Institute, University of California, Irvine, 1002 Health Sciences Road East, Irvine, CA 92612, USA

Abstract

Ciliary motion in the upper airway is the primary mechanism by which the body transports foreign particulates out of the respiratory system in order to maintain proper respiratory function. The ciliary beating frequency (CBF) is often disrupted with the onset of disease as well as other conditions, such as changes in temperature or in response to drug administration. Current imaging of ciliary motion relies on microscopy and high-speed cameras, which cannot be easily adapted to in-vivo imaging. M-mode optical coherence tomography (OCT) imaging is capable of visualization of ciliary activity, but the field of view is limited. We report on the development of a spectrally encoded interferometric microscopy (SEIM) system using a phase-resolved Doppler (PRD) algorithm to measure and map the ciliary beating frequency within an *en face* region. This novel high speed, high resolution system allows for visualization of both temporal and spatial ciliary motion patterns as well as propagation of metachronal wave. Rabbit tracheal CBF ranging from 9 to 13 Hz has been observed under different temperature conditions, and the effects of using lidocaine and albuterol have also been measured. This study is the stepping stone to in-vivo studies and the translation of imaging spatial CBF to clinics.

Graphical Abstract

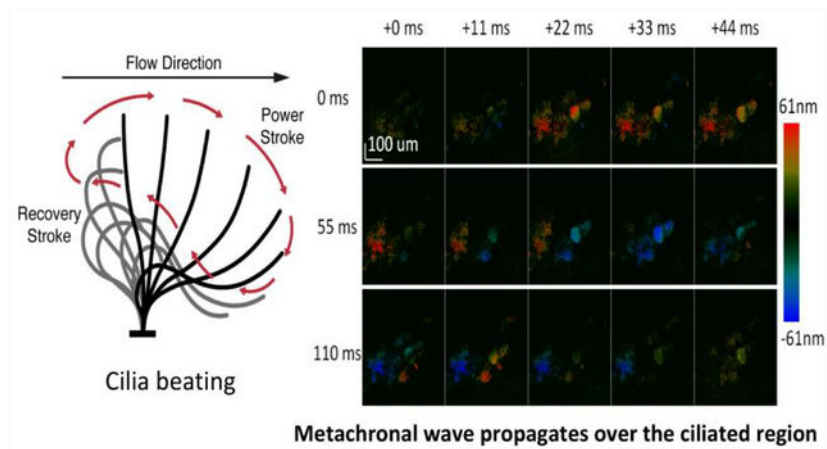
[†]Corresponding author, z2chen@uci.edu.

^{*}First three authors contributed equally to this study.

Disclosures

Dr. Zhongping Chen has a financial interest in OCT Medical Inc., which, however, did not support this work.

Support information: Video of ciliary activity captured with 91 fps for 2.2 s.



Keywords

ciliary beating frequency; ciliary beating pattern; *en face* Doppler imaging; spectrally encoded; microscopy; phase-resolved Doppler

The human airway serves not only as a passage for oxygen intake but also as the first line of defense to protect the lungs from harmful foreign particulates. The main defense mechanism is the mechanical clearance of mucus through ciliary activity and regulation of airway surfaces¹. Specifically, ciliary beat frequency and amplitude are considered the main factors that determine the maximal transportation speed of the mucus from the lower airway to the pharynx and, hence, affect the mucociliary clearance rate (MCR)². In addition, it was reported that the ciliary beating synchronicity would also contribute to the MCR. Dysfunction of cilia, caused by ciliary defects such as primary ciliary dyskinesia (PCD), virus, toxins, or trauma, will usually result in ineffective mucociliary clearance and lead to lung damage³. Although the damage caused by ciliary defects such as PCD cannot be fully treated, previous studies indicate that it is possible to maintain sufficient lung function through appropriate respiratory care⁴. Therefore, it is important to monitor the spatial ciliary beat frequency (CBF) and the ciliary beat pattern (CBP) in human airways for disease diagnosis and management.

The traditional procedure to analyze airway ciliary activity involves three steps: sample preparation, recording, and CBF measurement. A biopsy using a cytology brush is usually performed to harvest the ciliated cells from human respiratory tracts for cell culture^{5,6}. Many techniques, based on the photoelectric method⁷ and video cameras, were reported to be capable of evaluating ciliary activity. Due to the microscopic nature of ciliary structure and movement, a high resolution and high sensitivity imaging modality is essential to visualize the ciliary beating pattern and frequency. Optical microscopy is useful in resolving single cilia movements, but sufficient imaging speed is also required to obtain quantitative results of CBF and record the entire beating cycle. This leads to the utilization of a high speed digital camera microscopy system^{5,8}. However, a conventional digital camera system cannot investigate ciliary functions in-vivo and thereby diminishes the effectiveness of the measurement in physiological settings.

In order to overcome these limitations and perform in-vivo studies, optical coherence tomography (OCT) was recently applied to ciliary motion imaging, taking advantage of its micron scale resolution, real time imaging capability, and potential for endoscopic imaging⁹. Given these powerful features, the study of ciliary functions moves from the dish plate to its natural environment and demonstrates the feasibility of using OCT to investigate ciliary activity in-vivo¹⁰⁻¹⁶. Specifically, a high-resolution OCT endoscope is inserted into either the respiratory or uterus cavity to perform continuous acquisition at one cross sectional site, so that the intensity fluctuations caused by ciliary activity can be visualized over time¹⁰⁻¹⁶. To further expand the effectiveness of using an OCT system in imaging ciliary activity, our group used a phase resolved Doppler OCT (PRD-OCT) technique to probe the beating direction and speed of cilia. PRD-OCT is well known for its capability of measuring microscopic particle movement with pico-meter sensitivity¹⁷⁻²² and is proven to be a powerful tool in visualizing accurate ciliary dynamics by providing not only its transient position but also the relative speed and direction of the ciliary beat²³. Therefore, the beating phase of different cilia can be simultaneously monitored to study their synchronicity and beating patterns. In addition, the velocity imaging ability of PRD-OCT also benefits ciliary studies by enhancing the contrast of moving cilia versus other stationary tissue components and may possibly aid in locating the ciliated area within the entire cavity during in-vivo experiments. However, to visualize spatial-temporal cilia motion pattern using OCT requires a two-dimensional beam scan to encompass the ciliary motion that occurs in two directions over the ciliated surface²⁴. Because the rate of the spatial scan must be at least two times greater than the CBF for effective measurements according to the Nyquist theory, there is a limitation on the number of scanning points, which limits the lateral field of view (FOV).

A spectrally encoded interferometer (SEI) is capable of lateral 2-D imaging with a single axis beam scan²⁵⁻²⁸ which allows for *en face* imaging at a speed comparable with OCT cross-sectional scans. Recent studies have demonstrated the integration of the PRD method and SEI technique and have achieved phase sensitive *en face* Doppler imaging using a continuous wave laser as the light source^{24,29,30}. Therefore, SEI could be a potential solution to solve the unmet needs in the field of in-vivo ciliary imaging. We report on a swept source spectrally encoded interferometric microscopy (SS-SEIM) system to obtain the *en face* image of cilia beating frequency and pattern at high speed. We have optimized the traditional SEI design by using a swept source laser as the light source and incorporating a microscopy setup. Compared to the continuous laser, the swept source laser has been suggested to offer superior sensitivity, faster imaging speed, and less susceptibility to fringe washout phenomenon³¹⁻³⁵. The microscopy setup allows for high resolution visualization of ciliary structures. With these optimizations, we are able to achieve 91 frames per second (fps) over an *en face* image area up to 1 mm². The novel SS-SEIM system enables, to the best of our knowledge, the first study to map out spatial-temporal ciliary activity and opens up the possibility of investigating spatial ciliary beating frequency and pattern in-vivo.

Method

Spectrally Encoded Microscopy System Design

The SS-SEIM system was developed using a vertical-cavity surface-emitting laser (VCSEL) based on a swept source OCT system that has been explained in a previous publication²³. The center wavelength of the system is 1.3 μm , and the A-line rate is 100 kHz. The lateral resolution and displacement sensitivity are measured to be 1.2 μm and 0.3 nm, respectively. The sample arm is replaced by a spectrally encoded microscopy setup, which allows for simultaneous sample illumination with a single mode fiber input. The light in the sample arm is first collimated and diffracted by the diffraction gratings, yielding a line pattern as the laser sweeps through its full bandwidth. The 1-D galvanometer mirror G1 scans the wavelength encoded line of light in its perpendicular direction to produce *en face* optical imaging. Two optical relays are employed to center the line of light onto the galvanometer mirror and the back focal point of the objective lens, hence ensuring a flat scanning plane on the sample surface. Afterwards, a focused line is illuminated on the sample through the objective where different wavelengths are focused on different locations of the sample based on their diffraction angles. The collected backscattered light interferes with the reference light to form an interferogram which is detected by a balance detector. The system is capable of real time *en face* displacement imaging at up to 100 fps with 1000 A-lines per image and 2048 pixels per A-line, providing a FOV of approximately $480 \times 750 \mu\text{m}$. The system schematic is illustrated in figure 1.

Ex-vivo Tissue Preparation

Rabbit tracheal samples were harvested from freshly euthanized male New Zealand white rabbits under the approval of the Animal Care and Use Committee (IACUC) at UC Irvine. The tissues were immediately submerged in Hanks Balanced Salt Solution (HBSS) and kept at a room temperature of approximately 23 °C. Soft tissue was carefully dissected from the outer surface of the trachea to ensure an even surface for imaging. The tissue was cut along the long axis of the trachea to reveal the mucosal surface for imaging and then secured down with pins onto a silicone-lined petri dish. A layer of HBSS was added to the dish to mimic the natural tissue environment and minimize disruption of ciliary motion. Finally, the sample was moved to the imaging stage where it was placed on a temperature-controlled hot plate for temperature regulation during imaging. For lidocaine treatment, 2% lidocaine was added to the sample while submerged in HBSS, and imaging was performed after 5 minutes. For albuterol experiments, the sample was kept in 2 ml HBSS buffer with 0.3% albuterol, and imaging was performed approximately 1 minute after drug application.

Processing Algorithm

The principle behind ciliary motion induced Doppler OCT was explained in our previous publication²³ where a PRD-OCT algorithm was employed for axial displacement estimation. According to PRD-OCT^{36–38}, the particle displacement Z can be measured by analyzing the temporal phase changes of its corresponding interference component. Under the context of SEIM imaging, the position of the particle is indexed by the wave number due to the laterally dispersed sample illumination, and thereby, the PRD-OCT equation for the SEIM system can be modified as below:

$$\Delta Z(k,t) = \frac{\varphi(k,t + \Delta t) - \varphi(k,t)}{2kn} \quad (1)$$

where k denotes the wave number of the illumination on the particle of interest, n denotes the reflective index of tissue, and Δt represents the sampling interval that was determined by the frame rate. For the purpose of investigating the temporal dynamics of ciliated tissue, the region of interest was oversampled at 91 fps. The instantaneous phase, $\varphi(x,t)$, was derived from the Hilbert transformation on the spectral interference fringe. By performing the algorithm on consecutive *en face* images, spatial-temporal ciliary dynamics can be visualized. The spatial beating frequency of cilia was mapped out by extracting the beating frequency from the spatial-temporal data through a Fourier transformation over time. The flow chart of the processing algorithm is shown in figure 2. The PRD-based processing algorithm and SEM system control program are implemented in a custom C++ software and accelerated by a graphics processing unit (GPU) to provide *en face* displacement images in real-time.

Results

During imaging, the line scan was made to focus on the cilia layer on the top surface of the trachea sample. In addition to synchronized ciliary motion, some random ciliary movements were also observed at certain parts of the sample. For the purpose of measuring the CBF, we focused on the regions with synchronized motion. The ciliary motion was visualized in real-time, and the region of interest could be identified and imaged efficiently. Figure 3 shows the spatial ciliary activity at room temperature from an ex-vivo rabbit trachea sample, where 15 temporally consequent images were arranged from left to right and top to bottom at time intervals of 11 ms. Synchronized ciliary activity can be demonstrated by the metachronal wave propagates over the ciliated region. The synchronicity can be further visualized by the video attached in the appendix which demonstrates the ciliary activity continuously captured within 2.2s at the same region of interest. It has to be noted that the color scale in figure 3 is half of that in the video for better visualization.

Spatially Mapped CBF vs. Temperature

The tracheal sample was acquired as described in the methods section and a region with synchronized ciliary motion was identified. SEIM *en face* imaging was done while increasing the temperature of the sample from room temperature to 33 °C. The raw line-scan Doppler SEIM images are shown in figures 4a–c, where the same area of interest is imaged at 27 °C, 30 °C, and 33°C, respectively. Different colors correspond to the change in the phase amplitude and the direction of the ciliary motion which are directly proportional to the velocity of the motion. For different temperatures, the time at which the cilia complete a single power stroke cycle can be distinguished. The duration of the phase cycle can be estimated to be approximately 0.132 s for 27°C, 0.11 s for 30 °C, and 0.088 s for 33 °C. These results show that the CBF increases with temperature as expected from previous literature²³.

Since the spatially encoded data is continuously collected over 6 seconds, the temporal data can also be extracted. The temporal profile or M-mode at each spatial location can be measured along the depth direction. It is important to note that the line scan system has advantages in visualizing lateral spatial and temporal information but sacrifices depth information and can only image superficial structures. However, for thin samples like cilia, tens of microns in depth are more than sufficient. An example of the temporal profile at one line under the 3 temperature conditions is presented in figures 5a–c. The phase is averaged across the entire depth of view and drawn out in figures 5d–f. Although the phase amplitude varies depending on the angular fluctuations of the sample and noise, the CBF is mostly consistent. In order to quantitatively determine the CBF at that particular location, a FFT is performed on the phase plot to yield the frequency plots in figures 5g–i. The peak frequency occurs at 9 Hz, 12 Hz, and 13 Hz for the 27 °C, 30 °C, and 33 °C cilia samples, respectively.

With the same data processing method, the temporal data is analyzed at each spatial location to yield the spatially coded CBF maps shown in figures 6a–c for the three temperatures, respectively. The same region was imaged for all three conditions where there appeared to be two separate bodies of synchronized cilia, marked by the yellow and red boxes. Small changes in the CBF can be observed between the two spatial regions at each temperature. With the increase in temperature, the CBF of all cilia increased as expected, and the means and standard deviations of both regions at each temperature have been plotted in figure 6d. The mean values ranged between 7.4 Hz and 13 Hz, all reasonable values for mammalian CBF. A general upward trend can be concluded between the CBF and the temperature.

Spatially Mapped CBF vs. Lidocaine Application

For the next experiment, the effects of lidocaine administration were studied. In general, lidocaine is a local anesthetic that is expected to significantly slow down ciliary motion and mucus transport. For this experiment, all samples were kept at 27 °C. As described in the methods section, 2% lidocaine was applied to the sample and imaging took place after 5 minutes. The raw *en face* images for the baseline measurement and after lidocaine application are shown in figures 7a and 7b, respectively. While the baseline cilia completed a full power stroke cycle in approximately 0.14 s, the lidocaine-treated cilia were not even halfway through the cycle in the same amount of time. Again, the temporal or M-mode data is plotted at every spatial location, yielding in the examples shown in figures 7c and 7d for the baseline and lidocaine-treated cilia, respectively. From the temporal data, it is evident that the frequency decreased significantly after lidocaine administration. The displacement plot was generated using the average phase over the entire depth, and FFT was performed to calculate the frequency peak at each location. Finally, the spatial map can be visualized in figures 7e and 7f, which are before and after drug administration, respectively. The mean and standard deviation for the baseline and treatment data were calculated using the CBF from each spatial location where cilia were present and plotted in figure 7g. The CBF before treatment was 7.2 ± 2.2 Hz and decreased to 2.7 ± 0.7 Hz after treatment. As shown in the spatial map, many of the cilia stopped moving altogether after lidocaine was applied, while others merely slowed down. Only the active cilia were taken into account when calculating the mean CBF and standard deviation.

Spatially Mapped CBF vs. Albuterol Treatment

Ciliary activity in response to albuterol, which is commonly used to increase respiratory function, was also investigated to further verify the system effectiveness. According to previous reports about the effects of albuterol on ciliary tissue^{23,39–41}, CBF is expected to increase with the introduction of the drug. Similar to the lidocaine experiment, a rabbit trachea sample was kept at room temperature and the baseline data was first recorded when the sample was placed in HBSS buffer only. Then the buffer was replaced with a 0.3% albuterol solution as mentioned in the methods section, and imaging was performed after approximately 1 minute. As shown in figure 8, the spatial CBF for the experimental group increased across the region of interest from a baseline of 5.6 ± 0.8 Hz to 6.1 ± 0.8 Hz. We then added more albuterol to the buffer to achieve a 0.6% drug concentration. As expected, the overall CBF increased again to 6.7 ± 1.3 Hz. In addition, more regions of ciliated cells were activated and contributed to more synchronized ciliary activity within the same imaging area. The mean and standard deviation values were calculated and plotted in figure 8d.

Discussion & Conclusion:

CBF is one of the most important indicators of respiratory health, and studying the mechanisms of ciliary motion is crucial in understanding diseases of the airway. The currently available methods mostly focus on ex-vivo microscopy and high-speed camera imaging, which do not offer a realistic examination of the CBF in its natural environment. Recent studies have demonstrated in-vivo ciliary studies using a micro-OCT system which visualizes the ciliary beating frequency within a cross-sectional field of view^{10–16}. However, ciliary motion naturally occurs on the surface of ciliated tissue and a cross-sectional field of view may be insufficient to demonstrate the spatial pattern of the coordinated ciliary beats that occurs in two dimensions. Therefore, this study aims to visualize and analyze the CBF both spatially and temporally with potential adaptability for clinical translation and in-vivo analysis.

For this initial feasibility study, variations in the CBF were observed within a spatial region. While some differences in CBF are expected across a sample, it would be worthwhile to more closely analyze the origin of these variations to better understand ciliary motion. It was also obvious that some cilia were beginning to degrade and were no longer moving in a synchronized manner with each other. These randomly moving cilia contributed to noise within the sample and the measured CBF. Future in-vivo studies can help better explain the loss of synchronicity and spatial changes in CBF by eliminating tissue degradation and freshness factors.

The two primary challenges of translating this technology to in-vivo studies are the noise factor and the clinical adaptability of the scanner. Bulk motion from breathing is expected to have a large impact on the data. However, since noise from breathing is largely characterized by a low frequency bulk movement, frequency analysis can be utilized to eliminate these factors. The current system's sample arm and scanner unit are bulky and unsuitable for in-vivo studies. However, a handheld probe based design is currently under fabrication to ensure clinical adaptability in the next study.

In summary, we have demonstrated a novel method of spatially mapping the CBF and ciliary motion with high speed, high resolution Doppler SEIM. With this line-scan system, we were able to acquire real-time *en face* images of the cilia and analyze the synchronicity of the motion. A general trend was observed between increases in the external temperature and increases in the CBF. In addition, the CBF decreased significantly after application of lidocaine, and an increasing trend was observed with the introduction of albuterol. The feasibility of using SS-SEIM technology to map the CBF has been validated across different conditions, and the results serve as a stepping stone to our ongoing translation of the technique for in-vivo and clinical studies.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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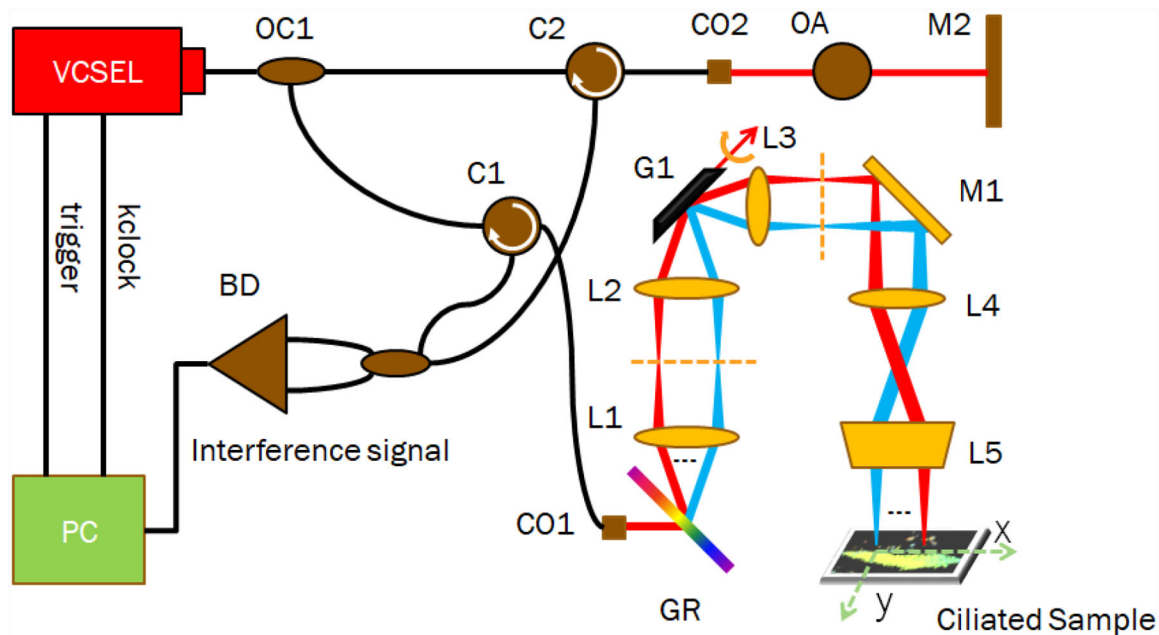


Fig. 1. System setup for SS-SEIM system. OC: optical coupler; C: circulator; CO: collimator; OA: optical attenuator; M: mirror; BD: balanced detector; GR: diffraction grating; L: lens; G1: galvanometer; FP: focal plane. The beam is dispersed in x-direction based on wavelength and the galvanometer scans the beam along the y-direction.

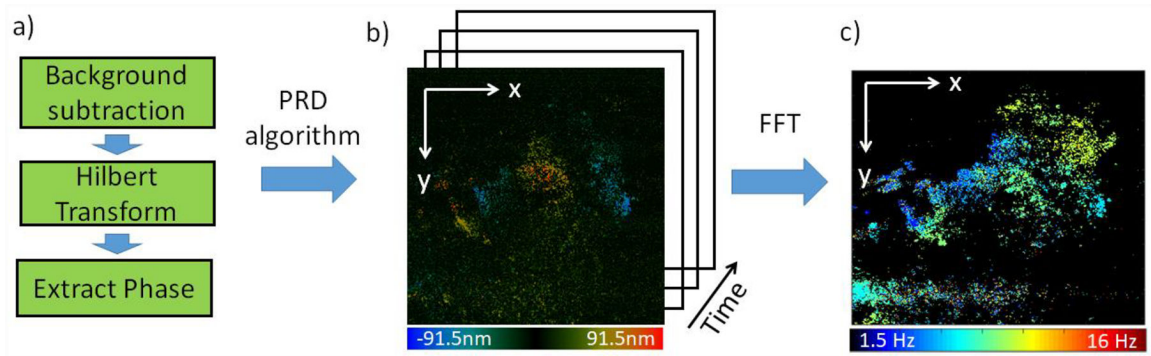


Fig. 2. Processing flow chart for obtaining spatial temporal ciliary activity and spatial CBF. The color bars below images b) and c) encode displacement and beating frequency of ciliary activity, respectively. a) Algorithm to achieve phase value at each imaging point. b) Axial displacement images spaced at same time intervals. c) 2-D CBF map.

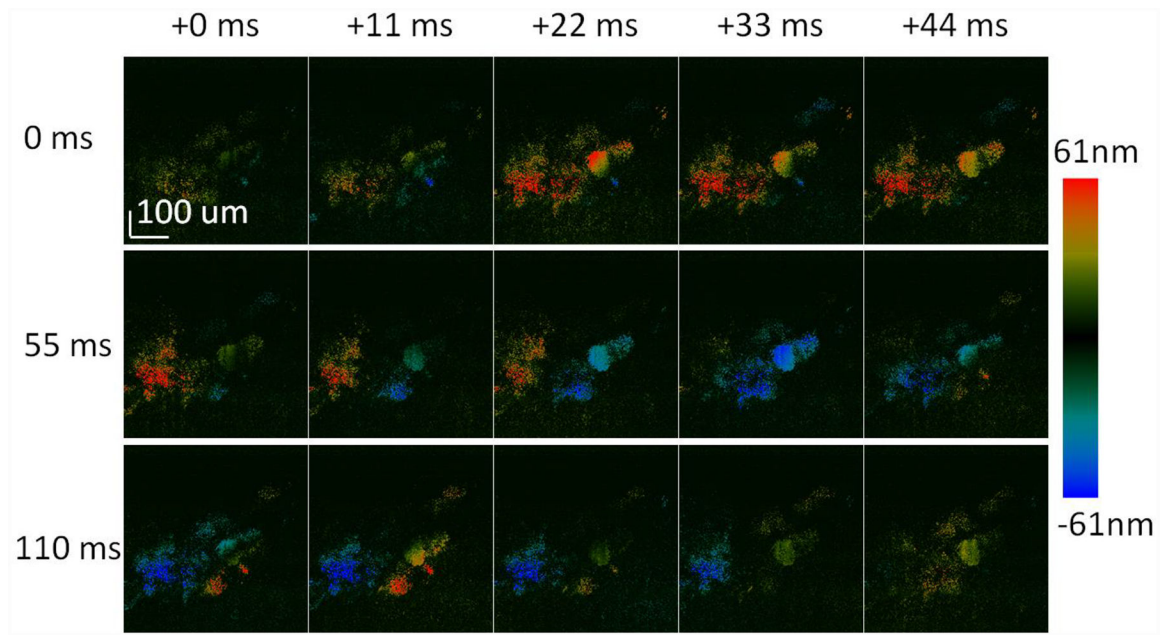


Fig. 3. Consequent images showing metachronal wave propagates over the ciliated region acquired at time intervals of 11ms: the color bar encodes displacement of ciliary activity. The images are arranged from left to right and top to bottom.

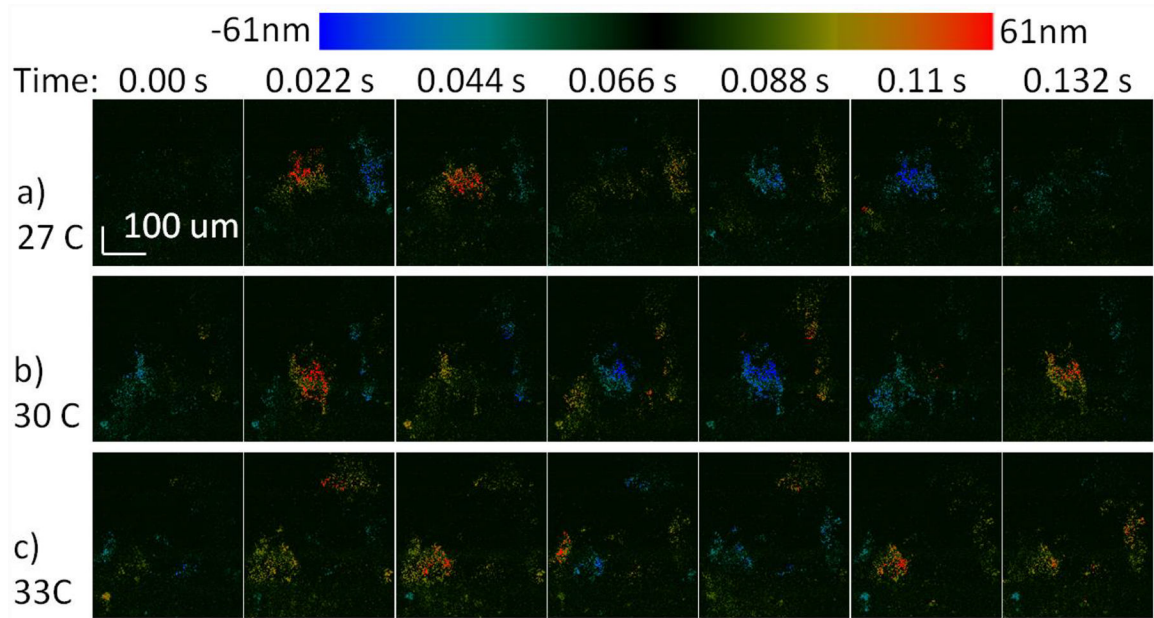


Fig. 4. Cilia motion at different temperatures: the color bar encodes displacement of ciliary activity.
a) Synchronized cilia cycle from 0 to 0.2 s at 27 °C, b) at 30 °C, c) at 33 °C.

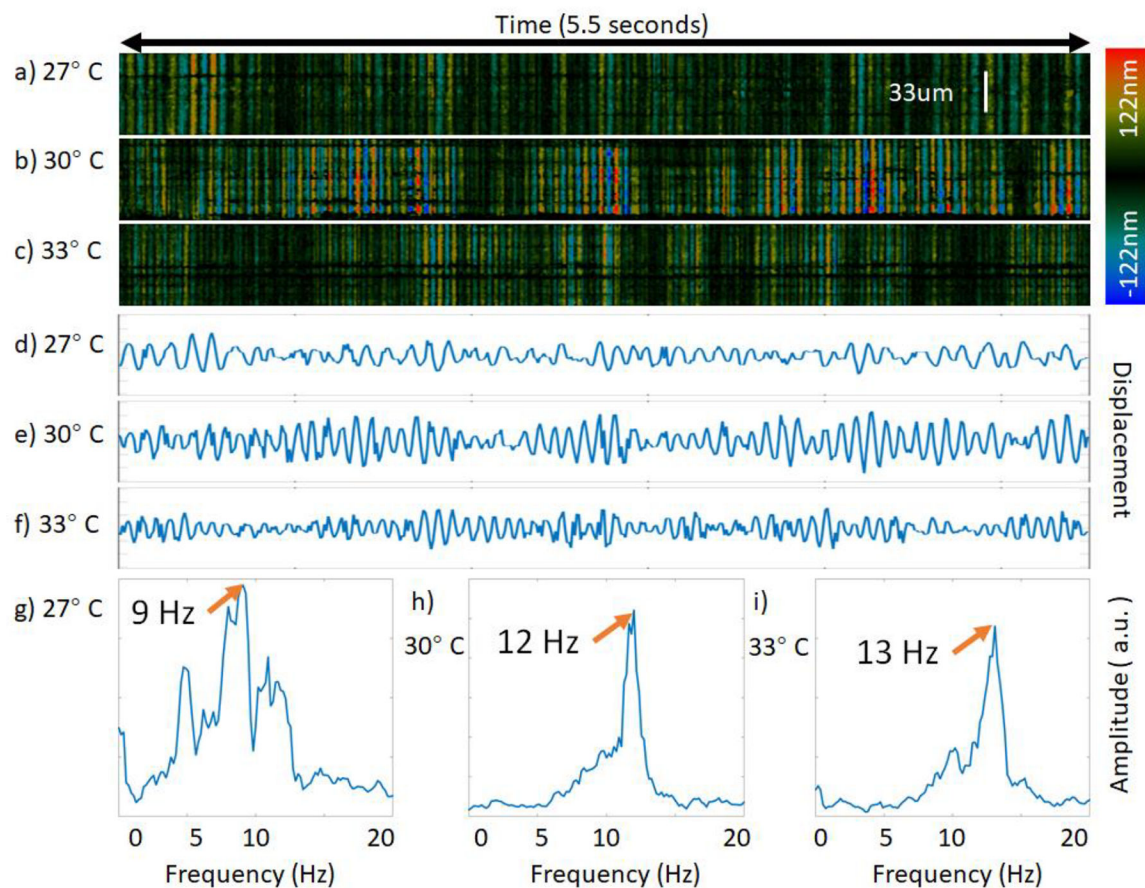


Fig. 5. Ciliary beat frequency measurements at a single location on synchronized cilia; the color bar encodes displacement of ciliary activity in images a-c). a-c) Raw phase data over depth, d-f) mean phase plot of raw data, g-i) frequency distribution after FFT.

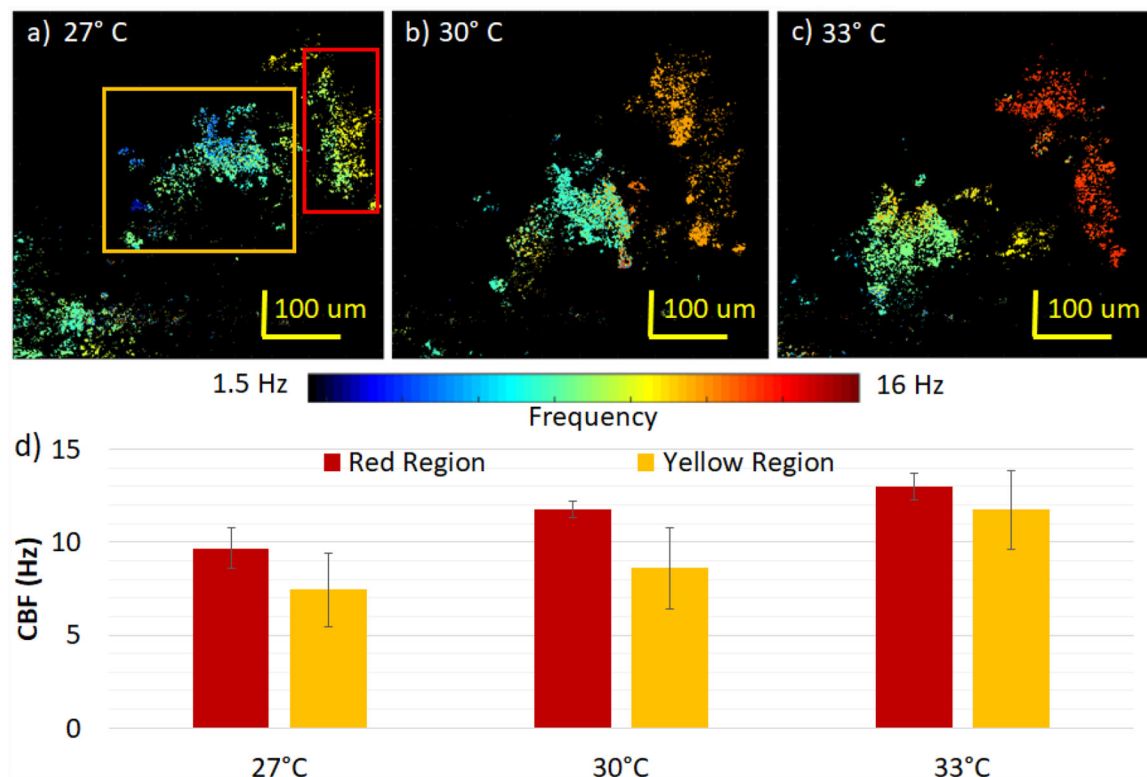


Fig. 6. Spatial distribution of ciliary beat frequency with changes in temperature. The color bar encodes the CBF for images a-c): a) at 27 °C, b) at 30 °C, c) at 33 °C with the same corresponding region. d) CBF analysis of yellow and red regions. The CBF at 27 °C, b) at 30 °C, c) at 33 °C for the red region is 9.7 ± 1.1 Hz, 11.8 ± 0.5 Hz, 13.0 ± 0.7 Hz and for the yellow region, 7.4 ± 1.9 Hz, 8.6 ± 2.2 Hz, 11.7 ± 2.1 Hz, respectively.

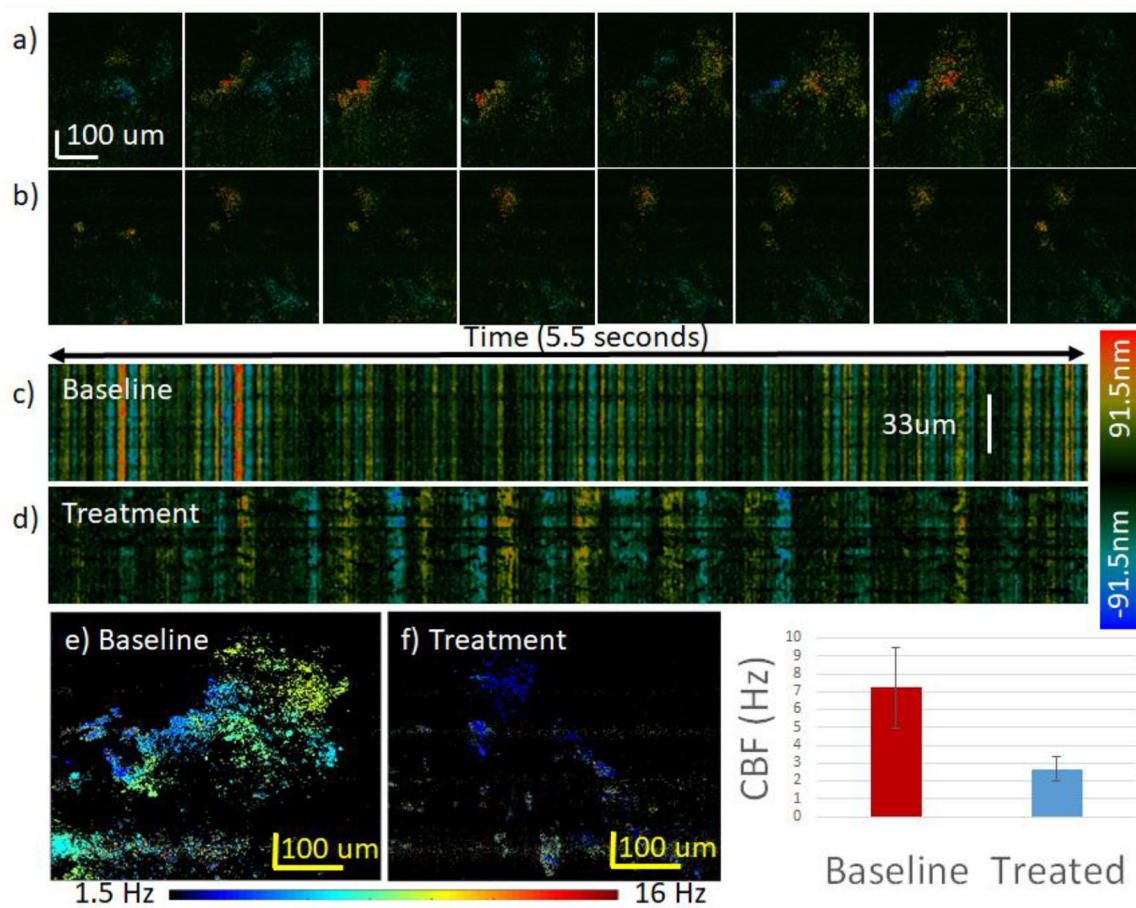


Fig. 7. Effect of 2% lidocaine treatment on CBF. The color bar on the right of images c-d) encodes ciliary displacement for images a-d). The color bar on the bottom encodes the CBF for images e-f). a) Synchronized cilia cycle from 0 to 0.2 s for baseline sample at 23 °C. b) Synchronized cilia cycle from 0 to 0.2 s for treatment sample at 23 °C. c) Raw phase data of baseline data. b) Raw phase data after treatment. Note that most of the cilia were no longer active after treatment, and a small region of active cilia was used for analysis. e) Spatial distribution of CBF for baseline sample. f) Spatial distribution of CBF for treatment sample. g) CBF analysis of entire region. The CBF before treatment was 7.2 ± 2.2 Hz and decreased to 2.7 ± 0.7 Hz after treatment.

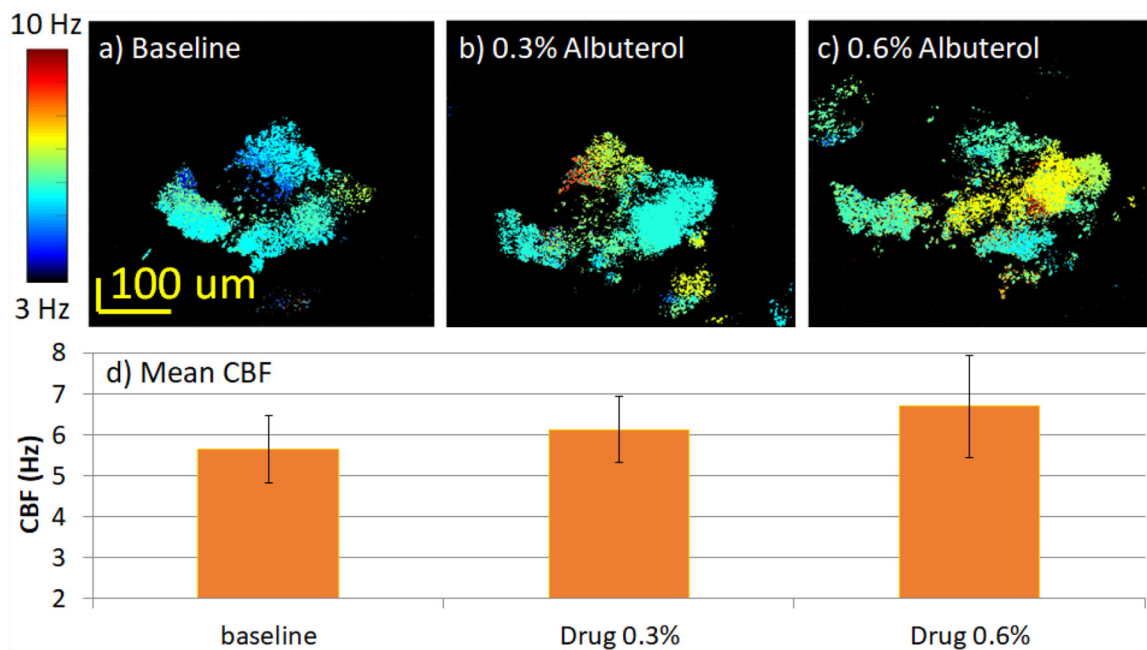


Fig. 8. Effect of albuterol treatment on CBF. The color scale for images a-c) encodes CBF. a) Spatial distribution of CBF for baseline. b) Spatial distribution of CBF for sample treated with 0.3% albuterol. c) Spatial distribution of CBF for sample treated with 0.6% albuterol. d) CBF analysis of entire region. The CBF of images a-c) are 5.6 ± 0.8 Hz, 6.1 ± 0.8 Hz, and 6.7 ± 1.3 Hz, respectively.