Quantitative optical coherence tomography imaging of intermediate flow defect phenotypes in ciliary physiology and pathophysiology

Brendan K. Huang
Ute A. Gamm
Stephan Jonas
Mustafa K. Khokha
Michael A. Choma
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Abstract. Cilia-driven fluid flow is a critical yet poorly understood aspect of pulmonary physiology. Here, we demonstrate that optical coherence tomography-based particle tracking velocimetry can be used to quantify subtle variations in cilia-driven flow performance in *Xenopus*, an important animal model of ciliary biology. Changes in flow performance were quantified in the setting of normal development, as well as in response to three types of perturbations: mechanical (increased fluid viscosity), pharmacological (disrupted serotonin signaling), and genetic (diminished ciliary motor protein expression). Of note, we demonstrate decreased flow secondary to gene knockdown of *kif3a*, a protein involved in ciliogenesis, as well as a dose-response decrease in flow secondary to knockdown of *dnah9*, an important ciliary motor protein. © The Authors. Published by SPIE under a Creative Commons Attribution 3.0 Unported License. Distribution or reproduction in any form, in whole or in part, is allowed only under terms of the license.

Keywords: mucus; *Xenopus*; optics; particle tracking; *dnah9*, *kif3a.*

Cilia-driven fluid flow clears mucus, bacteria, and particulates from the lungs. Despite its importance, it is a poorly understood aspect of respiratory physiology. Severe dysfunction of cilia in conditions such as primary ciliary dyskinesia (PCD) is associated with impaired respiratory mucus clearance and recurrent pulmonary infections. It is unknown, however, if subtle variations in cilia-driven mucus clearance underlie clinically significant changes in respiratory diseases severity, such as in asthma.

Prior measurements of ciliary flow have shown that flow speed can vary considerably between different specimens within an experimental population. Additionally, within a single specimen, flow has been previously described to vary as a function of distance from the cilia. Thus, when quantifying subtle variations in ciliary performance, it can be helpful to make cross-sectional measurements localized near a ciliated surface. Given these considerations, there has been increasing interest in applying optical coherence tomography (OCT), a modality that offers both spatial and phase information, to study ciliary physiology.

We previously demonstrated that OCT-based particle tracking velocimetry (OCT-PTV) could be used to estimate the velocity flow field in the *Xenopus* animal model system. In this letter, we build on these results and show that OCT-PTV can be used to quantify subtle changes induced by physical, chemical, and genetic perturbations. We quantify changes in ciliary flow due to changes in the viscosity of the fluidic environment, disruption of the serotonin signaling pathway, and diminished molecular expression of two important ciliary proteins, *dnah9* and *kif3a*. Additionally, we use OCT-PTV to characterize the developmental process of a ciliated surface.

*Xenopus* embryos (tadpoles), including *X. laevis* and *X. tropicalis* species, express cilia on their epithelial surface during development. The cilia themselves provide a time-varying backscattered signal that can be detected by OCT. Using speckle variance processing, we identified ciliated patches of the epithelium [Fig. 1(a) and Video 1], consistent with the methods described in Ref. 6. The ciliated patches in turn drive a microfluidic flow that can also be imaged using OCT. After immobilizing the embryo with benzocaine and seeding 10 μm microspheres in the fluid, we used OCT-PTV to estimate the microfluidic vector flow field, a spatial map showing the direction and magnitude of flow velocity at each location relative to a ciliated surface [Fig. 1(b)]. Of note, although chemical anesthetics have the potential to alter flow, benzocaine was previously described to have no discernable effect on ciliary performance, and we observed no visible effects.

Flow field estimation using OCT-PTV provides a two-dimensional (x, y), two-component (v<sub>x</sub>, v<sub>y</sub>) description of steady-state flow. Flow near the surface of the embryos is consistently directed head to tail, but flow more than several hundreds of microns from the surface varied in magnitude and directionality depending on the exact positioning of the embryo in the well. As such, we extracted only the component of flow spatially near the surface and directed along it a metric we denote as the average tangential flow speed. The tangential flow speed was calculated by manually drawing a line tangent to the surface of the embryo, extracting the flow field measurements <100 μm above the line and projecting each velocity vector along the tangent vector. These tangential flow measurements over the length of the embryos were then averaged to give the average tangential flow speed.

In order to verify that average tangential flow speed could be used to quantify changes in ciliary flow, we first tested the effects of a simple physical perturbation, an increase in viscosity. We increased the viscosity of the physiologic solution [1/9× modified Ringer’s (MR) solution] surrounding *X. tropicalis* embryos by adding high molecular weight dextrans (Sigma 95771, MW 2,000,000) to final concentrations of 1.3% and...
Due to the existence of previously characterized gene expression on the flow phenotype, we used morpholino oligonucleotides, a type of antisense technology that can decrease protein expression by preventing mRNA splicing or translation. Due to the existence of previously characterized morpholinos in X. laevis, we chose to investigate knockdown in X. laevis, a comparable system to X. tropicalis with a higher baseline flow speed. Using the dnah9 splice-blocking morpholino as previously described in Ref. 8, we injected varying doses into single-cell zygotes, ranging from 1 to 4 picomoles (pmol) per embryo. We coincubated an Alexa488 (Invitrogen) tracer into the embryos to verify proper delivery after 24 h. As shown in Fig. 2(c), increased morpholino dosing diminished average tangential speed in a dose-dependent manner when compared with both the uninjected controls, as well as a negative control injected with 4 pmol of a scramble morpholino sequence. Thus, we observed intermediate decreases in ciliary flow due to intermediate decreases in gene expression. This result highlights how subtle variations in ciliary function can be modulated by plausible molecular mechanisms, and how quantitative imaging can enable the detection of these intermediate phenotypes.

We also investigated the effects of disrupting the kinesin motor protein kif3a [Fig. 2(c)]. Under-expression of kif3a has been associated with a more severe asthma phenotype. Using sequence information available on Xenbase, we designed a morpholino to bind to the first splice site of kif3a in X. laevis (sequence 5′-AGAGCCTCTCTTACCAGGCATTTGTT-3′). Noting that an 8 pmol dosage leads to nonspecific embryo toxicity, we injected single-cell zygotes with 1 to 4 pmol. We found that, in contrast to dnah9, there was not a significant difference between the 1 and 2 pmol groups of kif3a (the power of comparison was 0.81). Only the highest dose of 4 pmol leads to a statistically significant decrease in flow (p = 0.03 for 2 to 4 pmol comparison, as calculated by the Mann–Whitney U test). Indeed, not all genes would necessarily be expected to generate intermediate phenotypes, but some may lead to binary phenotypes after knockdown past a given threshold. More studies increasing the number of dosing levels as well as quantifying protein expression would further elucidate this hypothesis.

Lastly, we used OCT to investigate how a ciliated surface transitions from the absence of flow to the presence of flow. Like other organs, ciliated surfaces undergo a developmental process. The development of coordinated ciliary flow in...
Xenopus embryos is known to occur over approximately 24 h and to involve the interplay between tissue patterning and hydrodynamic signaling. In spite of having been extensively studied from a molecular perspective, however, to our knowledge, the flow speed during this period has not yet been described in a quantitative manner. Thus, we quantified the average tangential speed during the onset of flow during 24 to 50 h, corresponding approximately to Nieuwkoop–Faber (NF) stages 21 to 34.

Fig. 3 (a) Longitudinal measurements of developmental flow on nine different embryos (Video 2, MOV, 312 KB) [URL: http://dx.doi.org/10.11171.JBO.20.3.030502.2] and (b) aggregate data showing mean ± standard deviation during 24 to 50 h postfertilization, or approximately Nieuwkoop–Faber (NF) stages 21 to 34.

In conclusion, the physiology of cilia-driven fluid flow is important for respiratory disease, but it is currently unknown whether small perturbations can cause intermediate defects in ciliary flow. Here, we have used OCT to show that genetic perturbations can cause intermediate flow phenotypes. We were also able to characterize flow in a developmental context. OCT-PTV is well-suited toward quantifying subtle changes because it can be used to interrogate flow near a ciliated surface with high spatial resolution. Based on our results, we believe that OCT-PTV will continue to find an important role in quantitatively investigating microfluidic ciliary physiology.

Acknowledgments

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