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## Die Physikalisch-Medizinische Sozietät Erlangen und das Max-Planck-Institut für die Physik des Lichts

lädt Sie zu folgendem Vortrag ein:

## "Nuclear biomechanics and cellular phenotype"

## Professor Kris Noel Dahl, Ph.D.

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The sequencing of the human genome has provided a wealth of scientific information, but this information is limited by the poor understanding of the mechanisms which control gene expression. In addition to containing the code for the cell, the genome within the nucleus is a complex, self-assembled polymeric structure with unique rheological properties. The genome of metazoan cells is surrounded by an intermediate filament network known as the nucleoskeleton. Using spectroscopy, imaging, micromanipulation and computational techniques, we measure the mechanics of the nucleoskeleton and the nuclear interior at various length scales. Specifically, I will focus on recent techniques we have developed comparing particle tracking microrheology with fluorescence lifetime imaging microscopy (FLIM) of the chromatin in the nuclear interior. Both measurements provide information regarding the structural and mechanical state of the chromatin. We are particularly interested in the role that force and cytokine treatment play in altering nuclear mechanics and gene expression in primary human cells. Motor activity from the cytoskeleton transduced through the nucleoskeleton impacts the driving force for nuclear and subnuclear movement, and altered chromatin condensation shifts the resistance and propagation of forces. We also quantify nuclear stiffness in a broad spectrum of cell types: cells with less regulated gene expression patterns, including stem cells and cancer cells, have much softer nuclei whereas aged cells have stiffer nuclei. While the mechanisms directing stiffness are still being elucidated, we have quantified dramatic downstream impacts of nuclear stiffness on cellular migration. Generally, nuclear architecture and mechanics impacts cell fate directly by altering cell stiffness and indirectly by modulating gene expression. These results have broad implications in cell biology, inhibition of cancer metastasis, and for applications in cellular therapies.

Kris Noel Dahl is an Associate Professor in the Departments of Biomedical Engineering and Chemical Engineering at Carnegie Mellon University. Her group is interested in structure and mechanics of materials inside cells including the nucleus and cytoskeleton. By studying these structures, it is possible to provide insight into cell function and adaptation including stem cell differentiation, cancer metastesis and interactions of cells with nanomaterials. Dahl received her BS degree from Carnegie Mellon in 1998, and Ph. D. degree in Chemical Engineering at University of Pennsylvania in 2004. She performed her postdoctoral fellowship in Cell Biology at Johns Hopkins University before joining Carnegie Mellon in 2007. She is a recipient of a Whitaker Fellowship and NIH Postdoctoral Fellowship for her training. She received an NSF CAREER award as well as a Young Investigator Award from the World Congress of Biomechanics. She has published 40 peer reviewed papers and 10 review papers and book chapters.

Mittwoch, 22. Juni 2016, 10:00 Uhr

(45 Minuten Vortrag plus Diskussion)

<u>Veranstaltungsort:</u> <u>Seminarraum TRC (</u>Erdgeschoss), TRC Erlangen Schwabachanlage 12, Erlangen

Für Rückfragen wenden Sie sich bitte an: Prof. Dr. med. Christian Bogdan Mikrobiologisches Institut - Klinische Mikrobiologie, Immunologie und Hygiene Universitätsklinikum Erlangen, Wasserturmstraße 3-5, D-91054 Erlangen Telefon: 09131 / 852-2551/-2281 · Fax: 09131 / 852-2573 · E-mail: <u>christian.bogdan@uk-erlangen.de</u>