



# Centre for Biosystems Science and Engineering

## SEMINAR

at 04:00 PM on March 20, 2017  
MRDG Seminar Hall

Visualizing dynamic cellular machineries with electron microscopy

**Dr. Saikat Chowdhury**

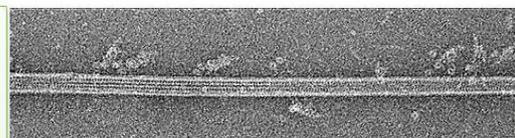
Postdoctoral Associate, The Scripps Research Institute

In spite of recent technological advancements in transmission electron microscopy (EM), determining structures of flexible macromolecules continue to be a major challenge. We show how EM can be used to elucidate structural details of cellular macromolecules involved in diverse biological functions.

The largest of these is the microtubule (MT) associated dynein-dynactin (DD) complex (~2.6 MDa in size), whose conformational flexibility has stymied structure determination for decades. By combining 2D and 3D EM analyses, we obtained structural snapshots of isolated dynein complex and DD complexes attached to MTs. From our analyses we not only discerned the first molecular details of native dynein, but also observed unique orientations of dynactin relative to the dynein tail, with the dynein heads positioned uni-directionally on MTs. This suggests a mechanism of overcoming auto-inhibited conformation of dynein.

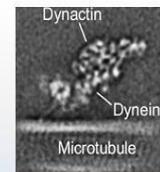
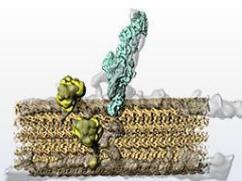
While these studies provide domain-level resolution, we also used EM to determine atomic-level details. To answer how phages evade the CRISPR-based bacterial defense system, we obtained near-atomic resolution reconstruction of phage-encoded anti-CRISPRs (ACRs) bound to the *P. aeruginosa* type I-F CRISPR-crRNA surveillance complex (~450kDa in size). New image processing techniques were used to overcome the intrinsic structural heterogeneity of the complex, revealing for the first time structural details of the type I-F CRISPR surveillance complex, and also explaining the molecular mechanism by which ACRs prevent binding of phage DNA to the CRISPR complex.

Gross molecular organization



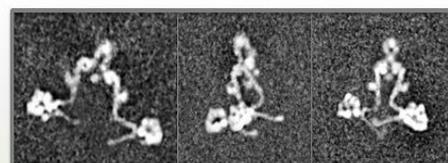
Microtubule with dynein-dynactin complexes attached

Molecular assemblies



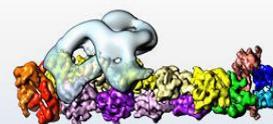
Single microtubule bound dynein-dynactin complex

Domain architectures



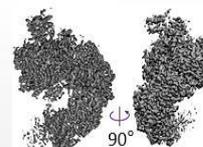
Native dynein molecules

Sub-nanometer resolution molecular details



Dynactin molecule

Near-atomic resolution molecular details



ACR bound type I-F CRISPR surveillance complex



$\beta$ -sheet

$\alpha$ -helix

crRNA

### About the speaker:

Dr. Saikat Chowdhury is currently an Postdoctoral Associate, Dept. of Integrative Structural and Computational Biology at The Scripps Research Institute. Dr. Saikat finished his PhD from Pennsylvania State University with Dr. B. Tracy Nixon. Previously, he did a B.Tech in Bioinformatics, Vellore Institute of Technology.