

Multi-scale phenotyping of developing cotton fibers

Alexander H Howell¹, Youngwoo Lee¹, Corrinne E Grover², Sivakumar Swaminathan³, Heena Rani^{1,4}, Pengcheng Yang², Elena Yu¹, Anika Sood¹, Jun Xie², Eileen Mallery¹, Jonathan F Wendel², Olga A Zabokna³, and Daniel B Szymanski^{1,5}

¹Department of Botany and Plant Pathology

²Affiliation not available

³Roy J Carver Department of Biochemistry, Biophysics and Molecular Biology, Iowa State University

⁴Department of Biochemistry, Punjab Agricultural University

⁵Department of Biological Sciences, Purdue University

October 30, 2023

Abstract

Cotton fibers in the *Gossypium* genus are the foundation of a multi-billion-dollar textile industry. Fiber development begins as unicellular trichoblasts emerge from the seed coat epidermis. This hemispherical trichoblast subsequently tapers and executes a complex cell elongation program. The trichoblast transitions to a cellulose-generating machine as it puts down layers of secondary cell wall before cell death and desiccation. Elucidating the multi-scale interactions and feedback controls among cytoskeletal systems, cell wall properties, and changing cell geometries will provide an abundance of opportunities to engineer more favorable traits during fiber development. To meet this long-term goal, we are conducting a “multi-omic” systems level analysis to better understand the fiber elongation process from 5 to 24 days post anthesis. An evolutionarily conserved microtubule-cellulose synthase control module is central to the processes of fiber tapering and anisotropic cell elongation. As such, molecular signatures of transcripts and proteins using quantitative proteomics were profiled and integrated across fiber development. Concurrently, a multi-scale image analysis pipeline was developed. Whole organs and fiber growth was measured under a stereomicroscope, fiber geometry and cellulose microfibril anatomy was characterized with confocal microscopy, and wall thickness was measured via TEM. Changes in the microfibril system are being analyzed in the context of the microtubule-CESA-module of gene expression dynamics. Though correlation of the phenotypic and molecular data is ongoing, these analyses are generating models to predict mechanisms of cellular pathway integration and phenotypic control. Finally, the structural information provides a robust dataset to refine finite element models of fiber growth.

Multi-scale phenotyping of developing cotton fiber

Alexander H. Howell^{1,2}, Youngwoo Lee^{1,2,◦}, Corrinne E Grover^{5,◦}, Sivakumar Swaminathan^{6,◦}, Heena Rani^{1,2,7}, Pengcheng Yang³, Elena Yu^{1,2}, Anika Sood^{1,2}, Jun Xie³, Eileen Mallery^{1,2}, Jonathan F Wendel^{5,*}, Olga A Zabolina^{6,*}, Daniel B Szymanski^{1,2,4,*},

¹Center for Plant Biology, ²Department of Botany and Plant Pathology, ³Department of Statistics, ⁴Department of Biological Sciences, Purdue University, West Lafayette, IN

⁵Department of Ecology, Evolution and Organismal Biology, and ⁶Roy J Carver Department of Biochemistry, Biophysics and Molecular Biology, Iowa State University, Ames, IA, USA and ⁷Department of Biochemistry, Punjab Agricultural University, Ludhiana, India

◦ These authors contributed equally to this work. * Corresponding authors

Cotton fibers in the *Gossypium* genus are the foundation of a multi-billion-dollar textile industry. Fiber development begins as unicellular trichoblasts emerge from the seed coat epidermis. This hemispherical trichoblast subsequently tapers and executes a complex cell elongation program. The trichoblast transitions to a cellulose-generating machine as it puts down layers of secondary cell wall before cell death and desiccation. Elucidating the multi-scale interactions and feedback controls among cytoskeletal systems, cell wall properties, and changing cell geometries will provide an abundance of opportunities to engineer more favorable traits during fiber development. To meet this long-term goal, we are conducting a “multi-omic” systems level analysis to better understand the fiber elongation process from 5 to 24 days post anthesis. An evolutionarily conserved microtubule-cellulose synthase control module is central to the processes of fiber tapering and anisotropic cell elongation. As such, molecular signatures of transcripts and proteins using quantitative proteomics were profiled and integrated across fiber development. Concurrently, a multi-scale image analysis pipeline was developed. Whole organs and fiber growth was measured under a stereomicroscope, fiber geometry and cellulose microfibril anatomy was characterized with confocal microscopy, and wall thickness was measured via TEM. Changes in the microfibril system are being analyzed in the context of the microtubule-CESA-module of gene expression dynamics. Though correlation of the phenotypic and molecular data is ongoing, these analyses are generating models to predict mechanisms of cellular pathway integration and phenotypic control. Finally, the structural information provides a robust dataset to refine finite element models of fiber growth.