

Director of Research

Head of the Microbial Molecular Genetics Laboratory

Dr. Vicky Sophianopoulou

vickybio.demokritos.gr

(+30) 2106503563 -3602

Fax (+30) 2106511767

[Curriculum Vitae](#)

Research Staff

Christos Gournas: Researcher

[Curriculum Vitae](#)

Ada Biratsi, Post-Graduate Fellow (PhD candidate)

Mirsini Charicleous, Undergraduate Student (Diploma Thesis)

Loudres Arias Salazar, Erasmus student

Research Interests

Our group is primarily interested in several aspects concerning: 1) regulation of expression, structure-function relationships, specificity, cell biology and evolution of amino acid transporter proteins and 2) the lateral organization of the fungal plasma membrane. Our model organism of choice is the non-pathogenic ascomycetes *Aspergillus nidulans*, a classic model genetic system since the 1950's. Two

Aspergillus nidulans

amino acid transporters have been cloned and studied in detail in respect to their transcriptional, post-translational/functional and cellular control of expression. In addition, we have introduced

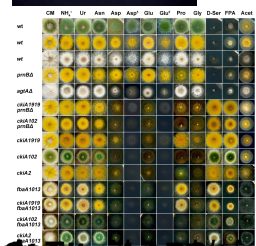
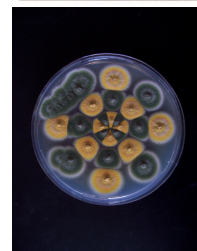
Aspergillus nidulans

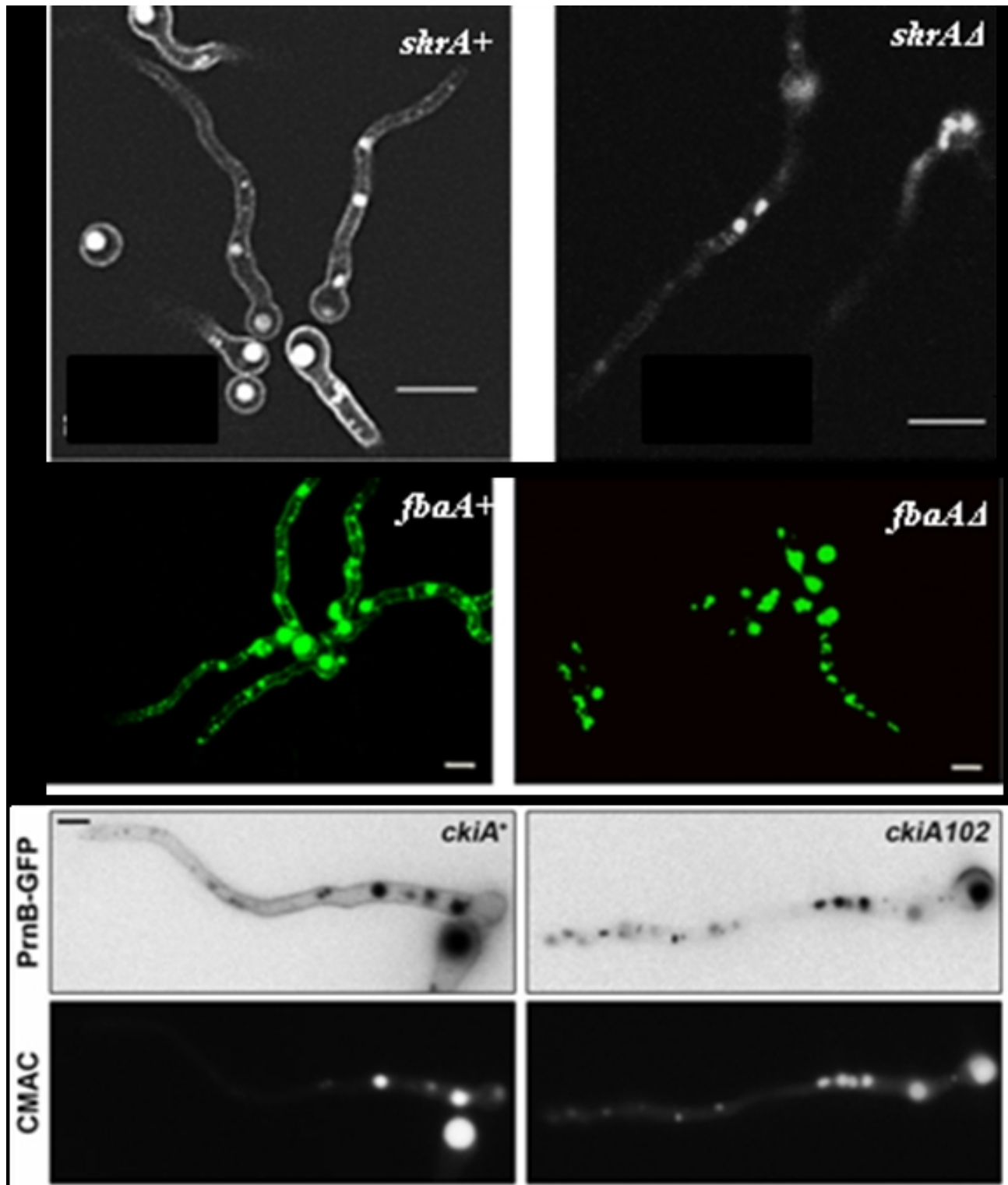
as a novel model system for the expression and/or functional characterization of heterologous transporter genes (Argyrou et al. 2001). In parallel, the last 7 years we have been studying the lateral compartmentation of

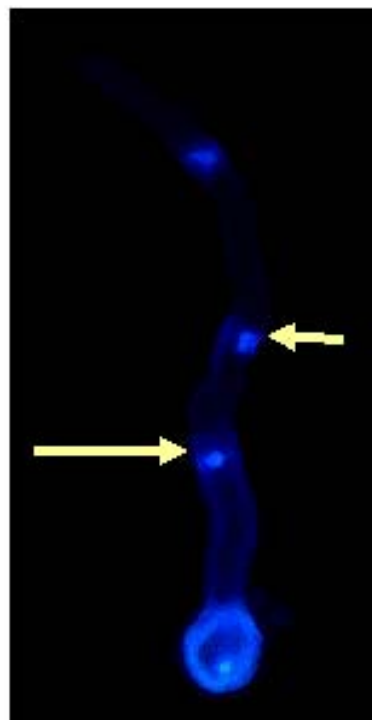
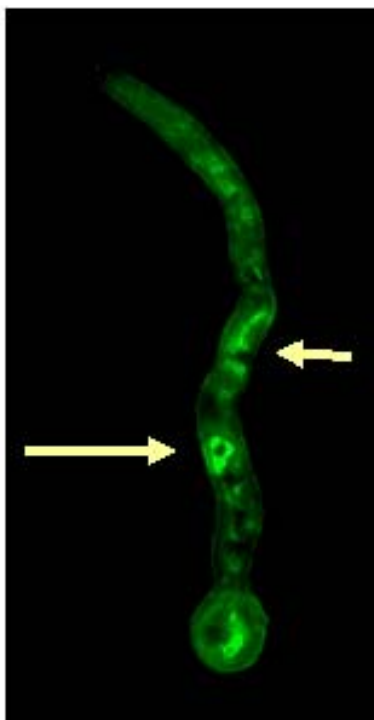
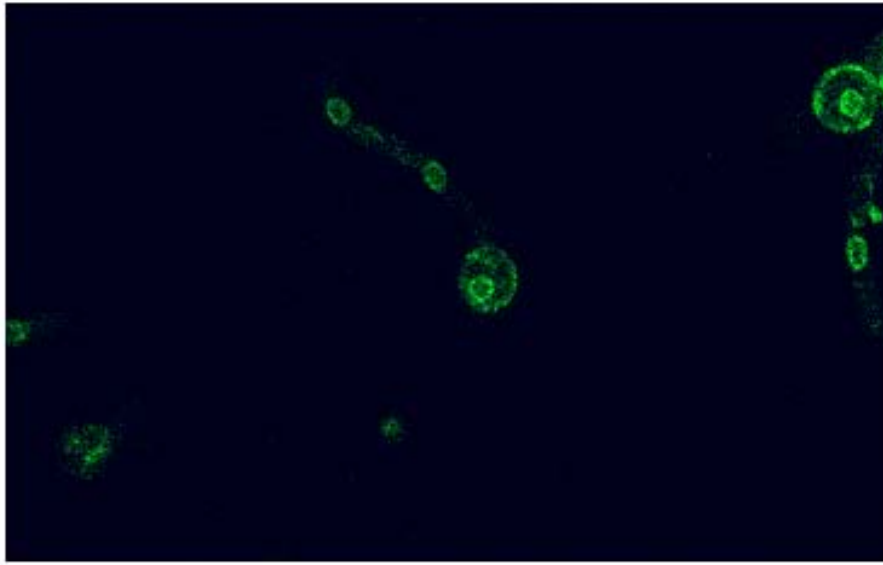
Aspergillus nidulans

plasma membrane and how this organization is implicated in different cellular processes. In particular we are focusing on “eisosomes” static plasma membrane compartments that constitute nanoscale furrow-like invaginations of the plasma membrane, where proteins and lipids segregate and their implication in fundamental cellular processes (sphingolipid biosynthesis, endocytosis of membrane proteins, function of efflux pumps and fungal pathogenicity).









Shree Anand, PhD student in Microbiology (SRMIST), is a member of carbon metabolism (FbaA) and a casein kinase



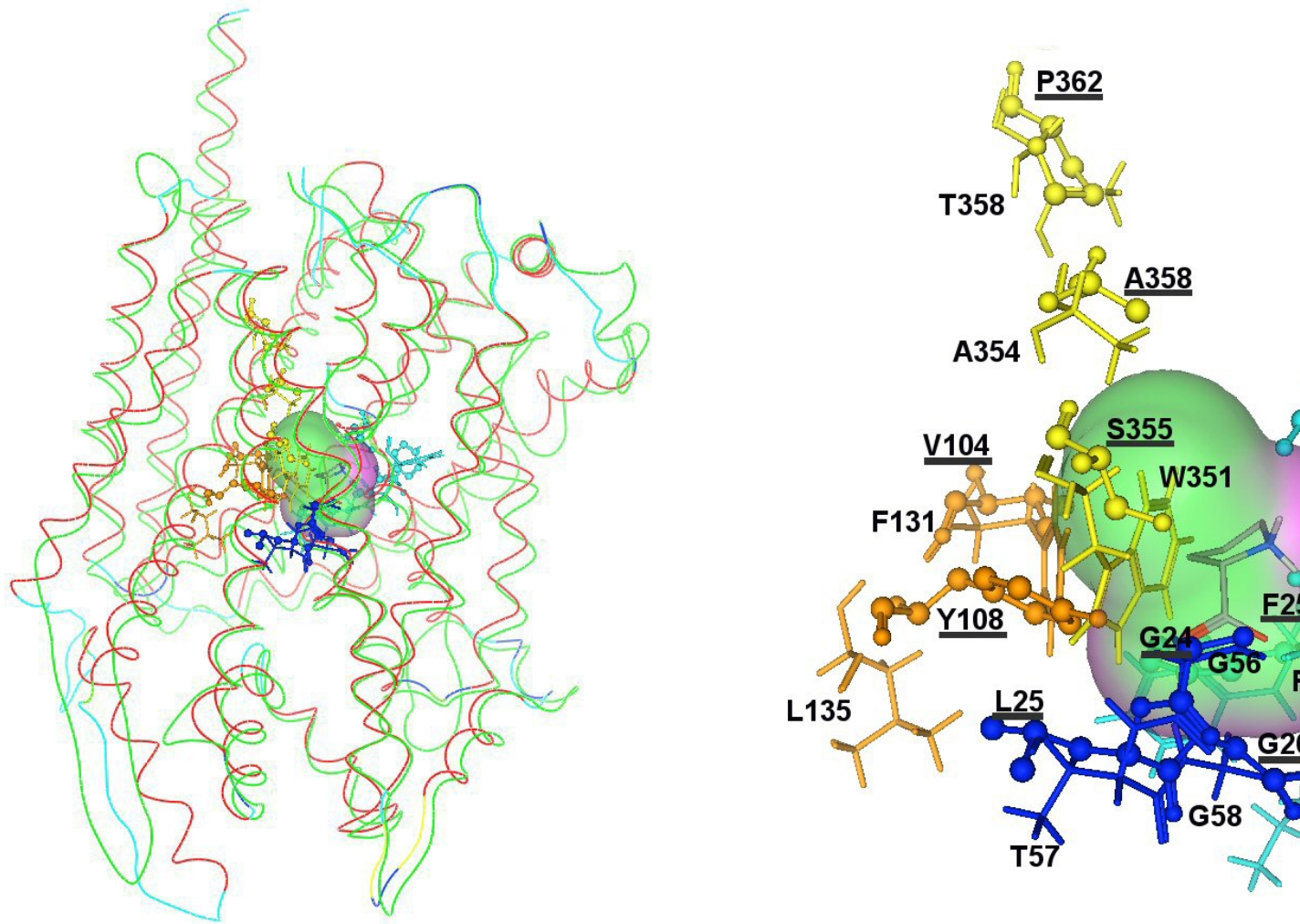
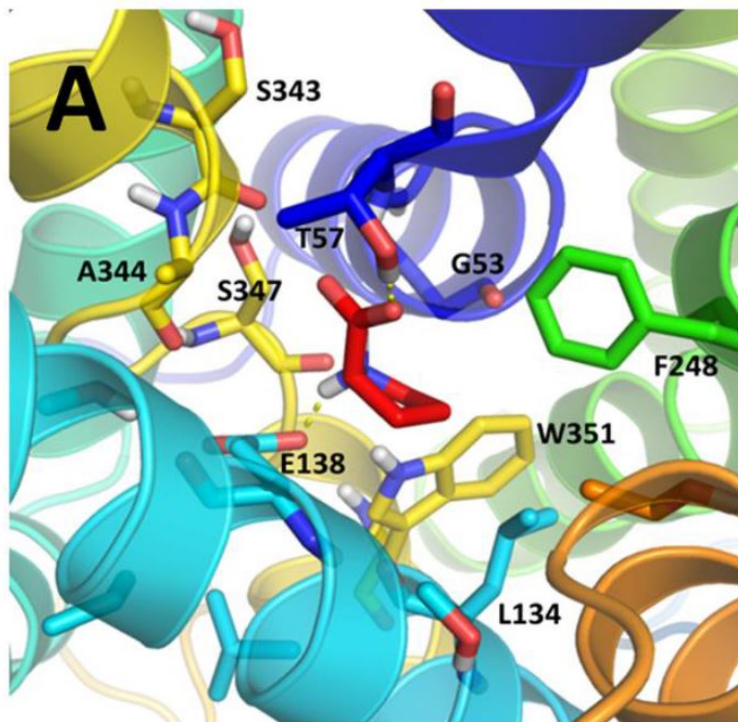
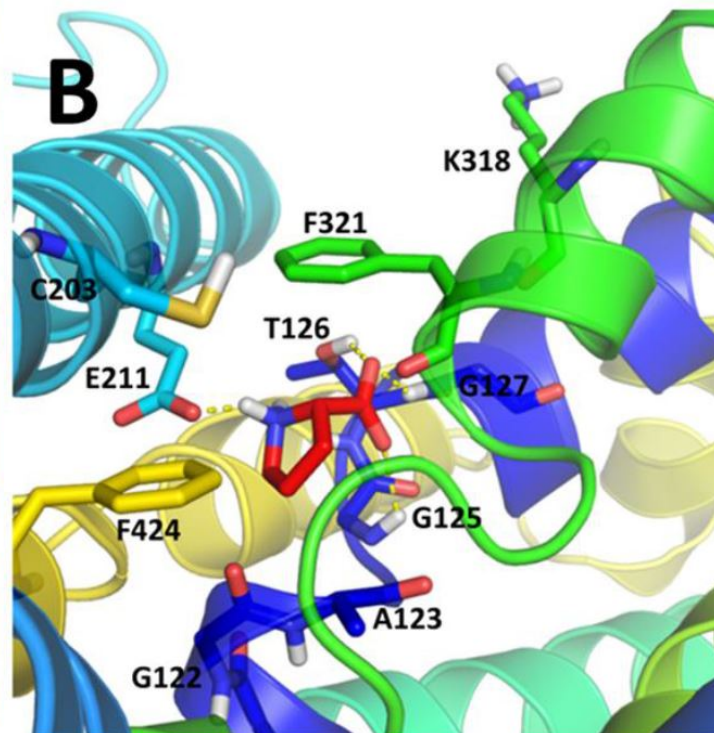


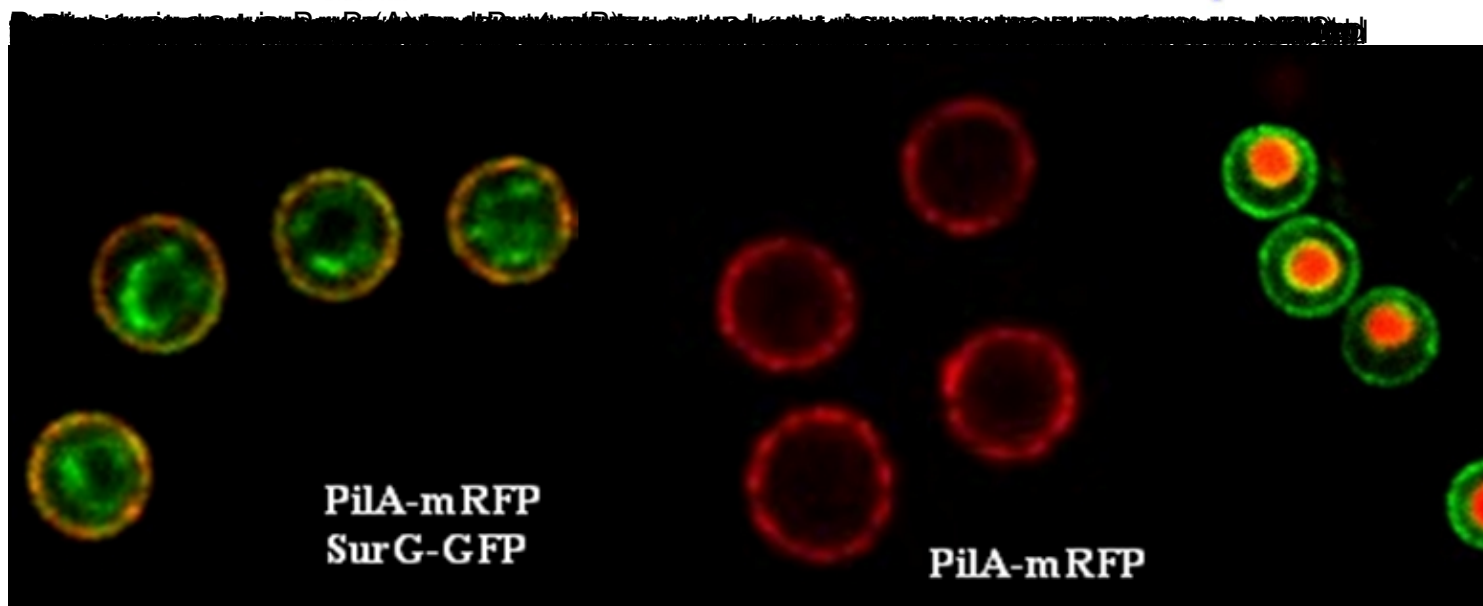
Figure 1. (a) Ribbon diagram of the protein structure with the bound ligand. (b) Surface representation of the protein structure with the bound ligand. (c) Detailed view of the binding site showing the interaction between the protein and the ligand.

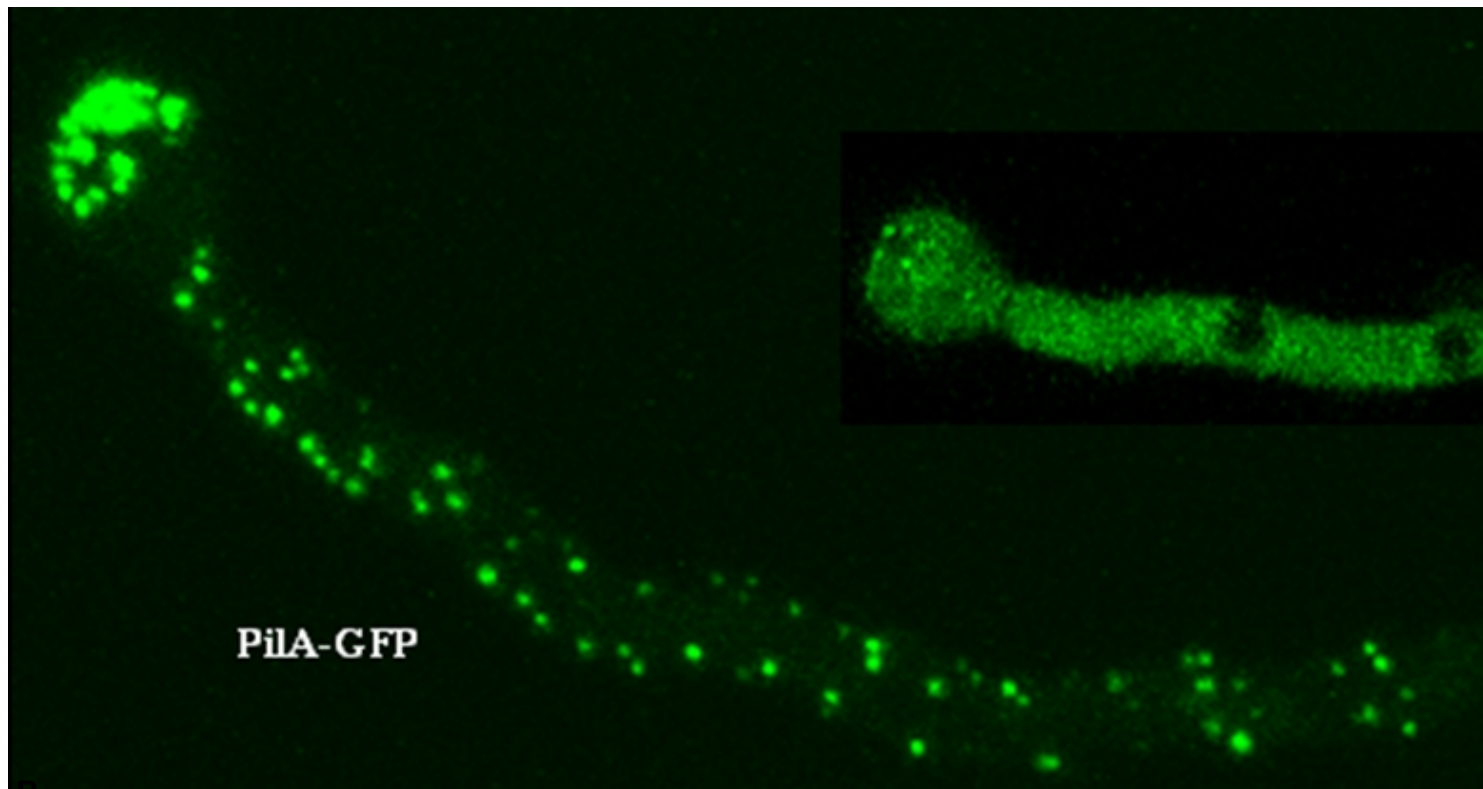


PrnB

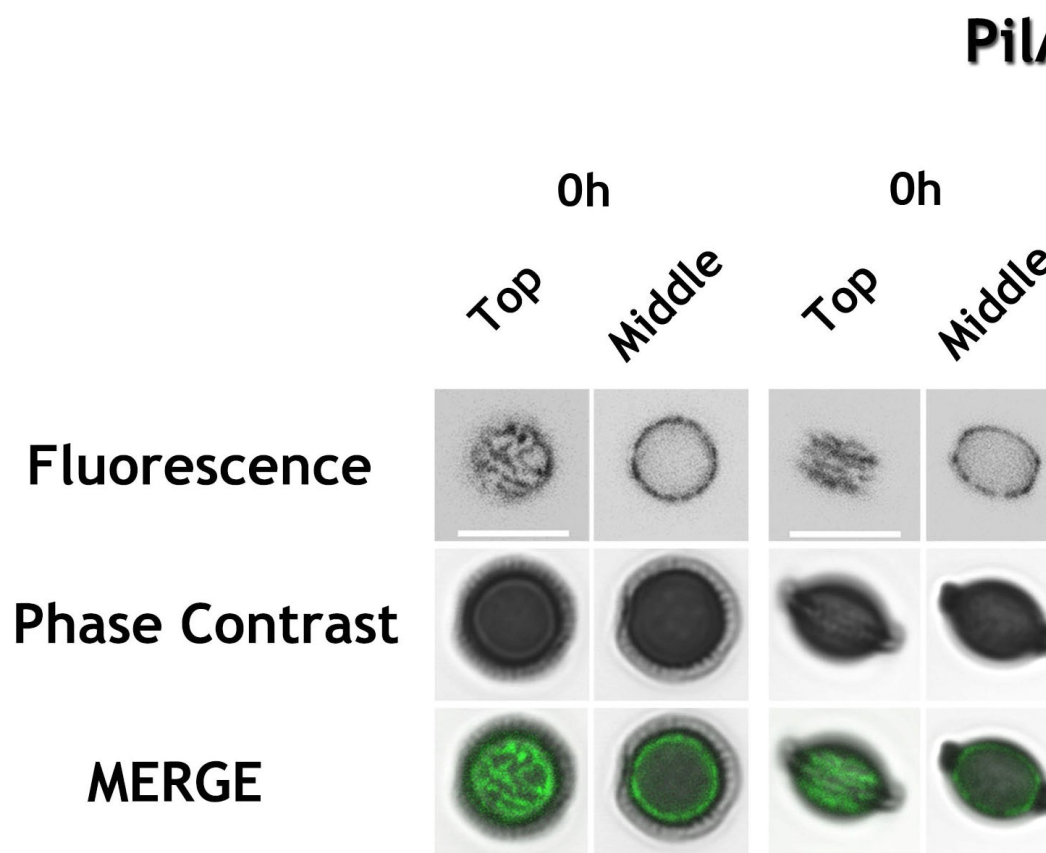


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B.



Fluorescence microscopy images of a bacterial cell expressing PiLA-GFP. The main image shows a long, curved chain of green fluorescent spots. The inset shows a higher magnification view of the cell's tip, which is densely packed with green fluorescent spots. The text 'PiLA-GFP' is overlaid on the main image.