

PhD Personal Statement Example

Field: Molecular Biology and Biochemistry

***What this essay does:** Opens with a specific dateable lab moment. Names the PI, institution, technique, and result in the first paragraph. States the research question precisely in paragraph two. Explains the choice of doctoral program over alternatives. Closes with a named trajectory, not a generic aspiration.*

The protein band should not have been there. I had spent six weeks designing an assay to detect San1 ubiquitin-ligase activity in a reconstructed *E. coli* system, and the western blot I was running that Tuesday afternoon in Dr. Rachel Kim's lab at Northwestern was supposed to show a clean negative control. Instead, there was a faint but unmistakable band at 34 kDa, exactly where I would expect to see it if the ubiquitin pathway had reconstituted spontaneously. Dr. Kim looked at the image for a long moment and then said: "Run it again with three biological replicates. If it holds, that is your thesis." It held.

I am applying to the PhD program in Molecular Biology and Biochemistry at Stanford University because I want to build on that finding under the mentorship of Professor David Chen, whose work on substrate specificity in the ubiquitin-proteasome system directly intersects with the questions my data raised. My central research question is why the ubiquitin modification pathway, which evolved exclusively in eukaryotes, can achieve partial reconstitution in a prokaryotic host, and what that reconstitution reveals about the minimal molecular requirements for ubiquitin-substrate recognition.

My undergraduate research in the Kim lab gave me the technical foundation to pursue this question. Over three years, I became proficient in bacterial and yeast transformation, PCR-based cloning, site-directed mutagenesis, co-immunoprecipitation, and quantitative western blotting. I also wrote the grant proposal that secured a Mary Gates Research Scholarship to fund my independent *E. coli* reconstitution project, which gave me direct experience in experimental design, budget management, and scientific writing under review. The work resulting from that scholarship was accepted for a poster presentation at the 2026 American Society for Biochemistry and Molecular Biology annual meeting.

What drew me specifically to doctoral research rather than a master's program or industry position is the timeline. The question of ubiquitin reconstitution cannot be answered in two years, and it cannot be answered in a setting where project goals are determined by commercial relevance. It requires the freedom to follow unexpected results like the one I found that Tuesday afternoon, the institutional support to run three hundred more replicates if necessary, and advisors who understand that the most

important findings are often the ones that contradict the experimental design. I am looking for five years of exactly that.

My long-term goal is to establish an independent research program investigating the evolution of protein quality control mechanisms, with a focus on how organisms repurpose conserved molecular machinery under novel cellular pressures. I intend to use Stanford's structural biology core facilities to combine my biochemical approach with cryo-EM data on ubiquitin-ligase conformational states, a methodological expansion that is not possible at my current institution. I am also interested in Professor Chen's lab's collaboration with the Stanford Neurodegenerative Disease Research Center, because aberrant ubiquitin pathway activity is implicated in both Parkinson's and Huntington's disease, and I want my basic science work to remain in contact with clinical questions.

I am ready to commit the next five years to a single question. The protein band that appeared in the wrong lane has not left my mind in eight months. I want to find out what it means.

