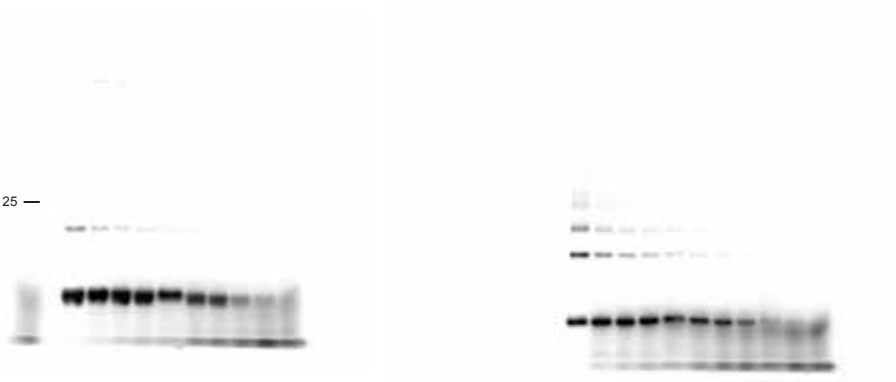


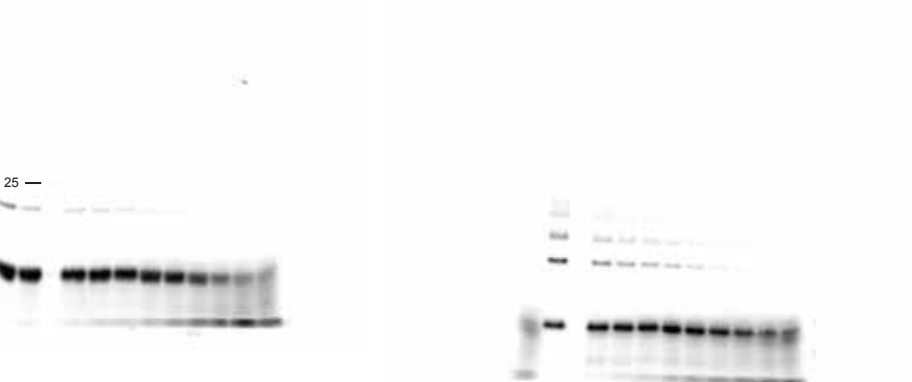
Molecular weight marker lane position indicates concentration of thermolysin used in the experiment.
Many gels were imaged at different exposures. For other exposures of gels contact corresponding author.
Time points are as indicated in Figure 2 and Methods.
Quantification is as describe in Methods.
All gels were imaged for fluorescein fluorescence.

0.2 mg/mL Thermolysin

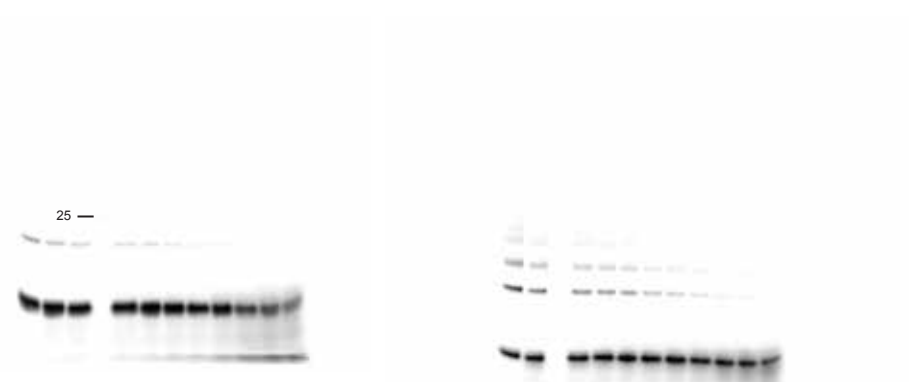


Final Replicate is presented in Figure 2 and Supplementary Figure 5

0.1 mg/mL Thermolysin



0.06 mg/mL Thermolysin



0.04 mg/mL Thermolysin



Molecular weight marker lane position indicates concentration of thermolysin used in the experiment.
Many gels were imaged at different exposures. For other exposures of gels contact corresponding author.
Time points are as indicated in Figure 2 and Methods.
Quantification is as describe in Methods.
All gels were imaged for fluorescein fluorescence.

0.4 mg/mL Thermolysin



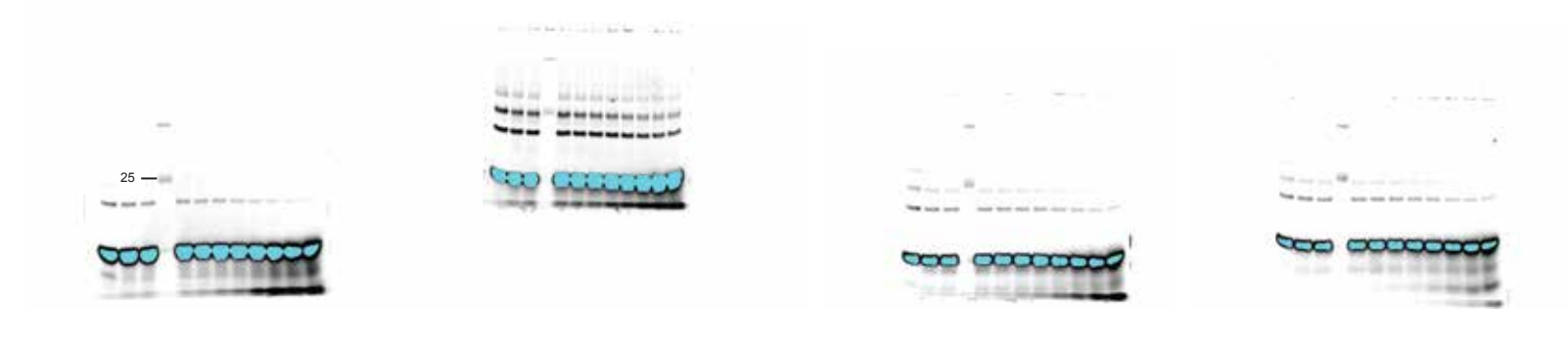
0.2 mg/mL Thermolysin



0.1 mg/mL Thermolysin

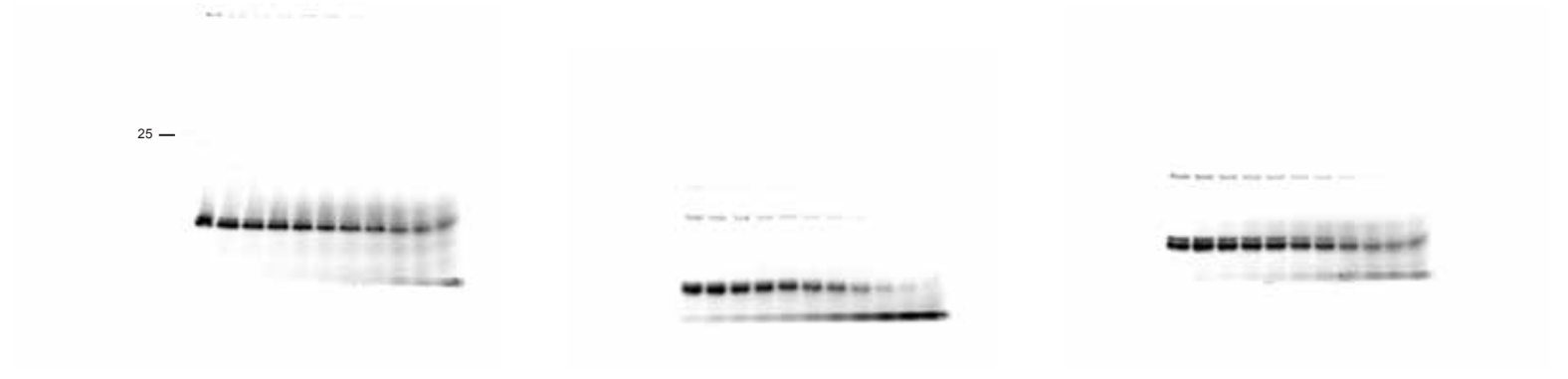


0.04 mg/mL Thermolysin

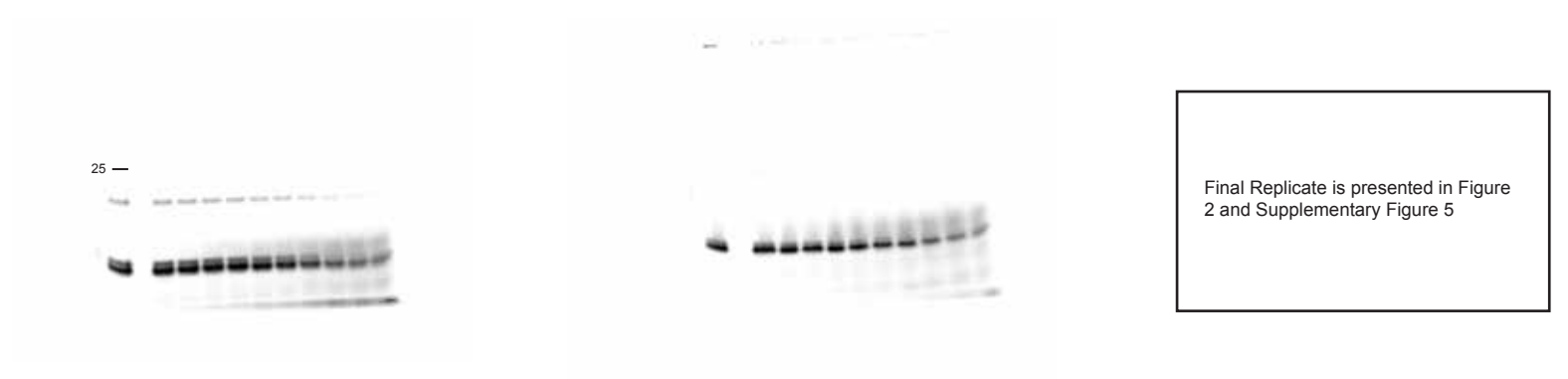


Molecular weight marker lane position indicates concentration of thermolysin used in the experiment.
Many gels were imaged at different exposures. For other exposures of gels contact corresponding author.
Time points are as indicated in Figure 2 and Methods.
Quantification is as describe in Methods.
All gels were imaged for fluorescein fluorescence.

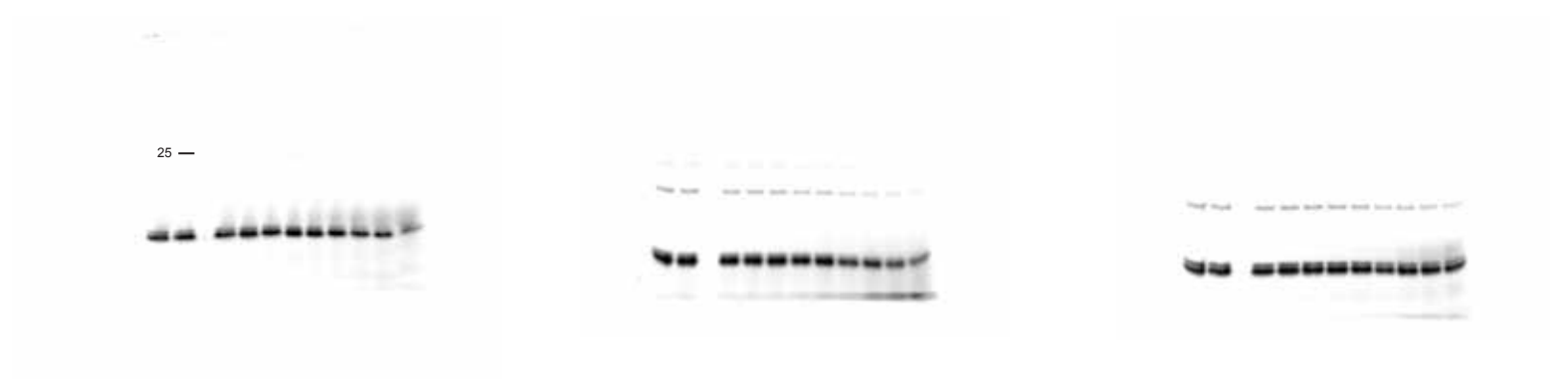
0.4 mg/mL Thermolysin



0.2 mg/mL Thermolysin



0.1 mg/mL Thermolysin



Molecular weight marker lane position indicates concentration of thermolysin used in the experiment.
Many gels were imaged at different exposures. For other exposures of gels contact corresponding author.
Time points are as indicated in Figure 2 and Methods.
Quantification is as describe in Methods.
All gels were imaged for fluorescein fluorescence.

0.4 mg/mL Thermolysin



0.2 mg/mL Thermolysin



0.1 mg/mL Thermolysin



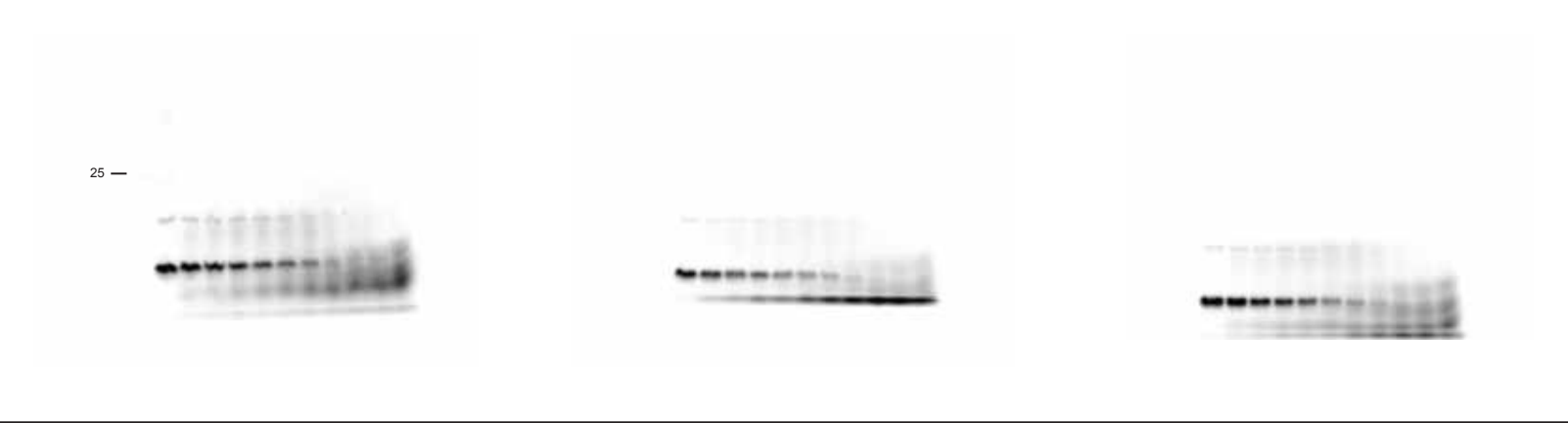
0.04 mg/mL Thermolysin



Molecular weight marker lane position indicates concentration of thermolysin used in the experiment.
Many gels were imaged at different exposures. For other exposures of gels contact corresponding author.
Time points are as indicated in Figure 2 and Methods.
Quantification is as describe in Methods.
All gels were imaged for fluorescein fluorescence.

Source Data for *M. smegmatis* DHFR gels are presented in Figure 2, Supplementary Figure 5, and Supplementary Figure 6.

0.4 mg/mL Thermolysin



0.2 mg/mL Thermolysin



0.1 mg/mL Thermolysin



0.06 mg/mL Thermolysin

