



Fig. S3. Commassie stain of HiTrap Benzamidine trapping of EspP used for espB degradation. Grown  $\triangle espB$  or  $\triangle espB\triangle espP$  in 200 ml low-glucose DMEM, 37C, 5% CO $_2$  for 4.5 hrs; filtered and concentrated with 10 kDa cut-off to 1 ml; buffer-exchanged 500 ul with PBS; buffer-exchanged other 500 ul with Binding Buffer and ran over HiTrap Benzamidine column; eluted with 7.5 ml glycine buffer, pH 3.0 into 2 ml Tris buffer, pH 9.0  $\Rightarrow$  buffer-exchanged with PBS to volume of 200 ul. Incubated samples +/- 50 ug of EspB overnight, 37°C; ran on 12% SDS-PAGE and Coomasie stained