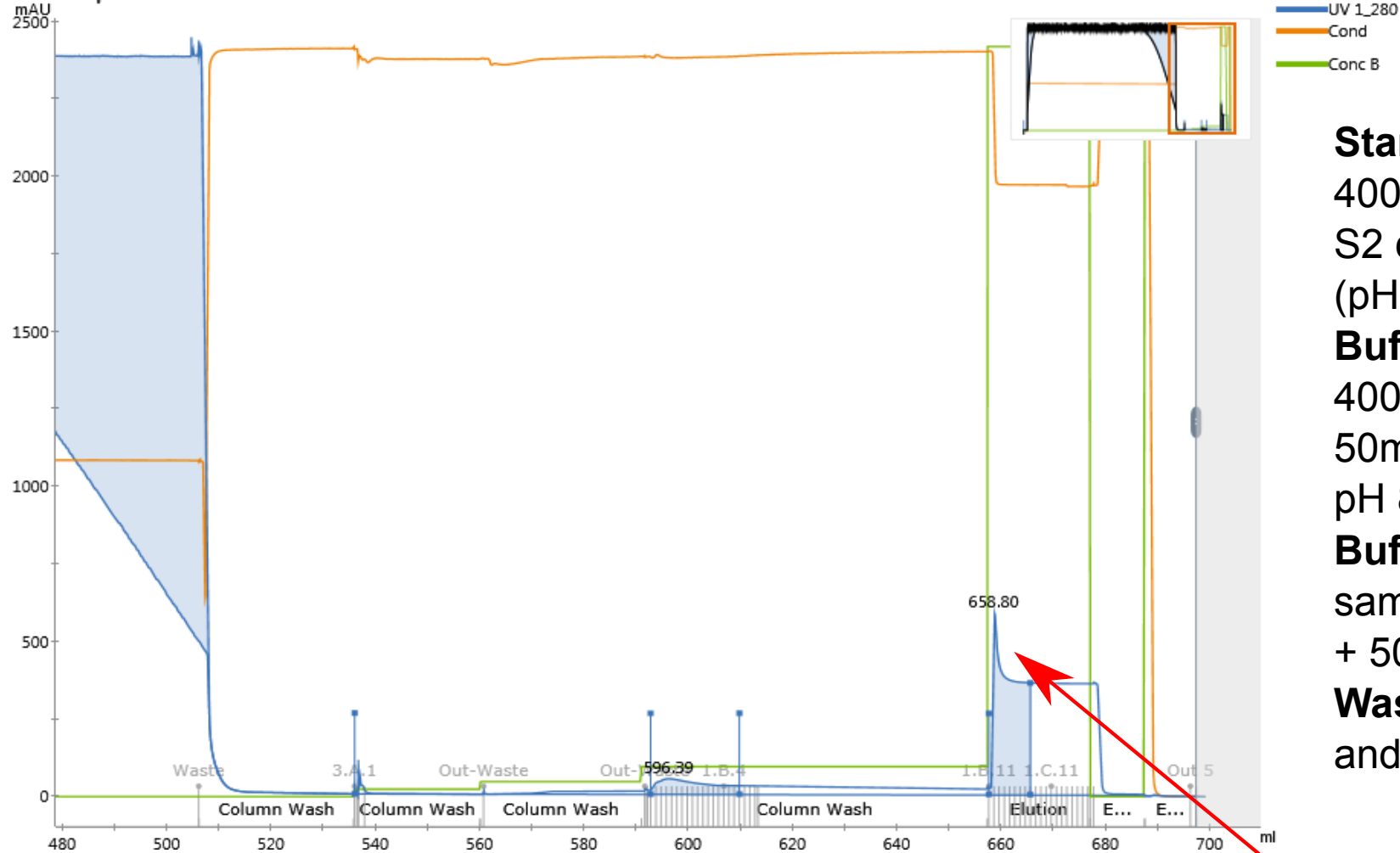


Group 1 & 8, Step 1: Ni²⁺ Sepharose (Excel) Affinity Chromatography of H₁₀-hHepsin

H10-Hepsin 2015-12-16 Ni2+



Starting material:
400ml of conditioned S2 cell supernatant (pH adjusted to 7.4)

Buffer A (binding buffer):
400mM NaCl,
50mM phosphate,
pH 8.0

Buffer B (elution buffer):
same as binding buffer
+ 500mM imidazole

Washing steps: 0, 1, 2
and 4% of Buffer B

Very tiny elution peak