



September 27, 2023

Dr. XXX XXXX  
 Company  
 Address

Dear Dr. XXXX,

Enclosed is a report detailing the results of one HCP antibody analysis obtained from 2D electrophoresis and western blotting of your protein sample and commercial antibody. Also enclosed are the silver-stained gels, ECL films, Western blotted PVDF membranes, and a CD containing copies of the files. A summary of the results is listed in Table 1. Gel loading and sample preparation are given in Table 2.

Film ID	Sample	Antibody Used	Percent Coverage	Figure Page #
LF1762 #8	K12 MG1655 <i>E. coli</i>	Cygnus Goat Ani- <i>E. coli</i>	73% (1550/2120)	p. 2 - 5

Table 1. Summary of Results of One HCP Antibody Analysis.

Gel ID	Sample	µl loaded	µg loaded	Treatment/Antibody
LF1762 #8	K12 MG1655 <i>E. coli</i>	150	250	Cygnus Goat Ani- <i>E. coli</i>
LF1762 #12	K12 MG1655 <i>E. coli</i>	150	25	Silver
LF1762 #13	K12 MG1655 <i>E. coli</i>	150	25	Silver

Table 2. Key to 2D Gel Loading and Sample Preparation. The sample was lysed in 5 ml of SDS boiling buffer without reducing agents and 5 ml osmotic lysis buffer containing nucleases, protease inhibitors, and phosphatase inhibitors. The sample was sonicated for 5 minutes, heated in a boiling water bath for 5 minutes, and treated with omnicleave. The protein concentration of the sample (16.2 mg/ml) was then determined using the BCA Assay (Smith et. al. *Anal. Biochem.* 150: 76-85, 1985, and Pierce Chemical Co., Rockford, IL). Sample was diluted to 2.5 and 0.25 mg/ml in SDS boiling buffer diluted 1:1 in Urea sample buffer before loading.

Please feel free to contact me if you have any questions.

Sincerely,

Isaac O'Malley-Laursen  
 Senior 2D Analyst