

## ATW0015

Aptamer to Human Serum Albumin

### Selection Information

**Target for Selection:** Human Serum Albumin (HSA), Sigma Cat# A3782

**Number of DNA Nucleotides:** 39

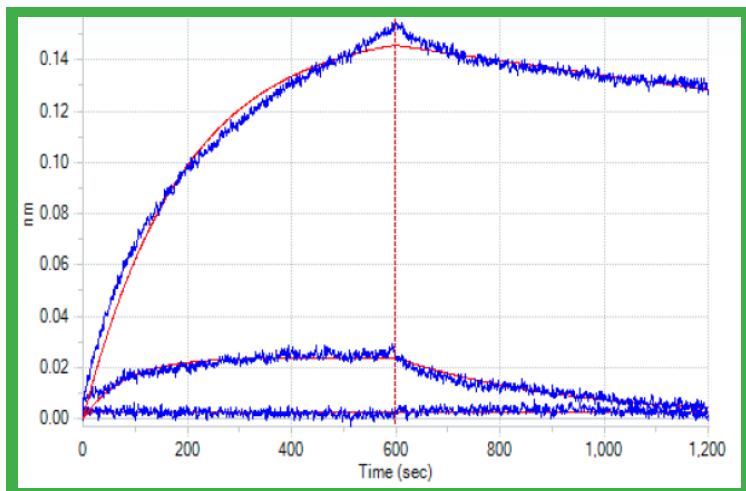
Aptamers were selected from a randomized Base Pair 39-mer DNA library against the target protein. Proprietary methods were used to select this specific aptamer sequence.

### Affinity Determination

**Affinity Determination Method:** Bio-Layer Interferometry (BLI)

**Buffer Used for Affinity Determination:** 20mM Tris, 100mM NaCl, 0.005% Tween 20, in nuclease-free water, pH 7.4

**Average  $K_D$ :** 29.1 nM



**Figure 1. Aptamer-HSA Binding**

Association and dissociation graph of 1:1 fitting model of HSA aptamer (biotinylated) to HSA protein concentrations 500, 125 and 31.25 nM, by single reference method.

**Table 1.  $K_D$ ,  $R^2$ , and  $\chi^2$  values by local fitting for single reference method. Avg  $K_D$  = 29.1 nM**

Immobilized Aptamer	Analyte	Concentration	Response	Full $\chi^2$	Full $R^2$
ATW0015 + biotin	Hu HSA Protein	500	0.512	0.016463	0.985181
ATW0015 + biotin	Hu HSA Protein	125	0.0268	0.004003	0.933033

### Aptamer Folding

For optimal binding, aptamers must be folded into their tertiary structure prior to use. Dilute to 10x working concentration in Folding Buffer, heat to 90-95°C for 5 minutes, then cool to room temperature (~15 minutes). Final application buffers used for dilution of aptamer to working concentration and washing should contain 1 mM MgCl<sub>2</sub>.