

Supporting Information

Cord-Based Microfluidic Chips as a Platform for ELISA and Glucose Assays

Laura Y. GALLEGOS, Jenny ELOMAA, and Frank A. GOMEZ*

**Department of Chemistry and Biochemistry, California State University, Los Angeles,
5151 State University Drive, Los Angeles, California 90032, USA*

Detecting Biotinylated Goat anti-Mouse IgG & Rabbit IgG

As seen in Tables S1 and S2, it is described how each reagent was administered onto the platform, how much volume was used, how long the drying time, and overall total time required to run each point of care chip on the μ CAD platform.

Table S1. Reagents and reaction wait times for the detection of goat anti-mouse IgG antibody using the μ CAD platform as a POC device.

Reagents	Administration of fluid	Volume [μL]	Time [min]
NC functionalization of reaction site	Spot	100	15
Biotin labeled IgG	Spot	1.5	10
Wash	Flow	10 x 3	1
Strep-ALP	Flow	15	5
Wash	Flow	10 x 3	1
p-NPP	Spot	1.5	10
p-NPP stop	Spot	1.5	10
Total	--	179.5 μL	52 min

Table S2. Reagents and reaction wait times for the detection of rabbit IgG antibody using the μ CAD platform as a POC device.

Reagents	Administration of fluids	Volume [μL]	Time [min]
Antigen immobilization	spot	1.5	10
Blocking buffer	spot	1.5	10
Antibody complexing	spot	1.5	1
Wash	flow	100 X 20	25
Representative substrate molecule	spot	1.5	30
Total	--	2006 μL	76 min

Detection of Glucose in Urine

As seen in Table S3, it is described how the reagents were administered onto the μ CAD platform, how much volume was used, how long the wait time was for each reagents, and total time it required to run each assay on the μ CAD platform.

Table S3. Reagents and reaction wait times for the detection of glucose using the μ CAD platform.

Reagents	Administration of fluids	Volume [μL]	Time[min]
Glucose	spot	5	10
GO _x , HRP, KI	flow	45	10
--	flow	--	30
Total	--	50	50