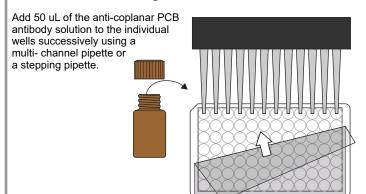


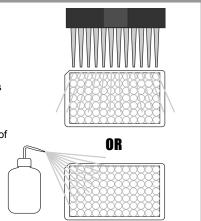
ABRAXIS® Coplanar PCB ELISA 530011

1. Addition of Antibody Solution

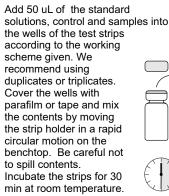


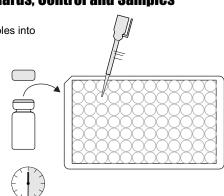
4. Washing of Plates

After incubation, remove the covering and vigorously shake the contents of the wells into a sink. Wash the strips three times with a multi-channel pipette or wash bottle using the diluted 1X washing buffer solution. Please use at least a volume of 250 uL of washing buffer for each well and each washing step. Remaining buffer in the wells should be removed by patting the plate dry on a stack of paper towels.



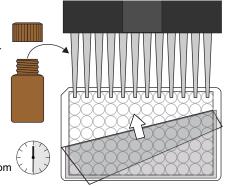
2. Addition of Standards, Control and Samples





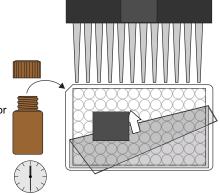
5. Addition of Substrate/Color Solution

Add 150 uL of substrate/color solution to the individual wells successively using a multichannel pipette or a stepping pipette. Cover the wells with parafilm or tape and mix the contents by moving the strip holder in a rapid circular motion on the benchtop. Be careful not to spill contents. Incubate the strips for 20 - 30 min at room temperature.



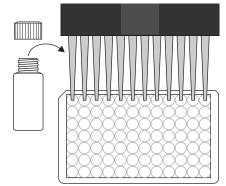
3. Addition of Reconstituted Enyzme Conjugate

Add 50 uL of the reconstituted enzyme conjugate to the individual wells successively using a multi- channel pipette or a stepping pipette. Cover and mix for 30 seconds by rotating on benchtop. Incubate for 90 minutes at room temperature.



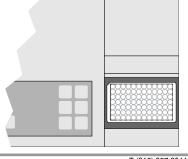
6. Addition of Stopping Solution

Add 50 uL of stop solution to the wells in the same sequence as for the substrate solution using a multi- channel pipette or a stepping pipette.



7. Measurement of Color

Read the absorbance at 450 nm using a microplate ELISA reader. Calculate results.



Gold Standard Diagnostics 795 Horsham road Horsham, PA 19044 WEB: www.abraxiskits.com T (215) 357 3911

F (215) 357 5232

Ordering: info.abraxis@us.goldstandarddiagnostics.com Technical Support: support.abraxis@us.goldstandarddiagnostics.com

Date this Flow Chart is effective: 05/17/2024

Version: 0