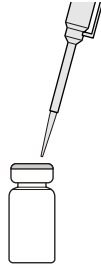


LAS Plate Kit, Detailed ELISA Procedure

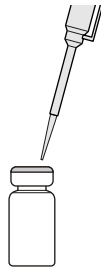
1. Sample Pretreatment

Filter the samples if necessary and add methanol to obtain a final methanol concentration of 10% (v/v)



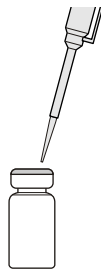
2. Preparation of Standard Solution

Dilute the LAS standard concentrate with 10% methanol/water solution to prepare designated standard concentrations from 0.02mg/L to 1mg/L.



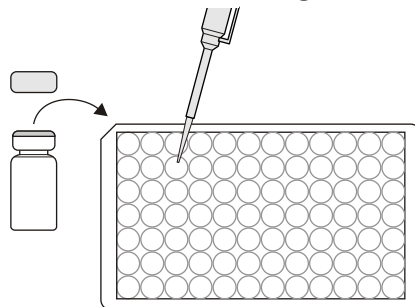
3. Reconstitution of Antigen-enzyme Conjugate

Reconstitute antigen-enzyme conjugate powder with 7mL of buffer solution. Mix by filling tip and expelling the contents with a pipette.



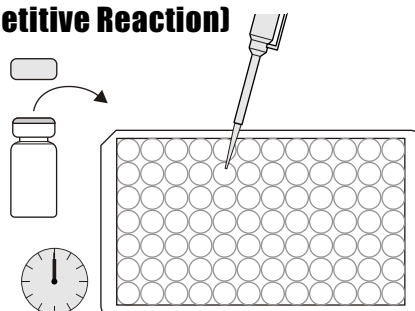
4. Addition of Standard/Sample and Conjugate

Dispense aliquots of 100uL of the above mixture into each **antibody coated** well of a microplate. Cover the wells with parafilm or tape and mix the contents by moving the strip holder in a rapid circular motion on the benchtop for 30 seconds. Be careful not to spill contents. Incubate for 60 minutes at room temperature.



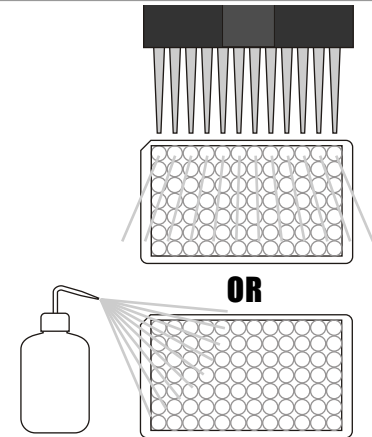
5. Incubation (Competitive Reaction)

Dispense aliquots of 100uL of the above mixture into each coated well of a microplate. Incubate for 60 minutes at room temperature.



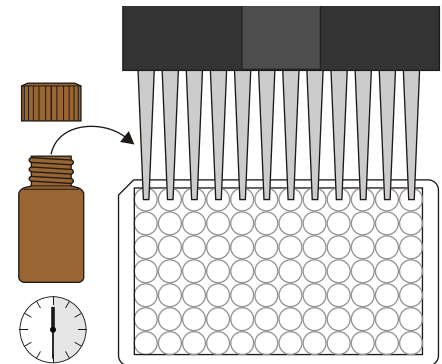
5. Washing Solution

Dilute 6-fold wash solution with 5 volumes of distilled water. Rise each microplate well with 300uL of the 1x wash solution and repeat 3 times.



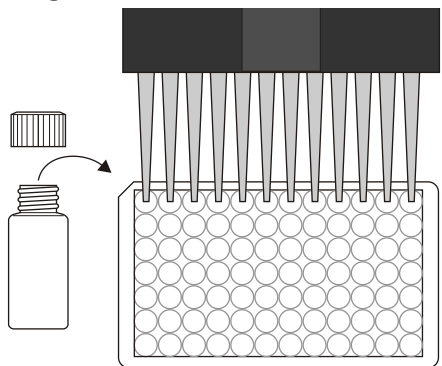
6. Color Development

Dispense 100uL of the Color Solution into each well successively using a multi-channel 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 for 30 seconds. Be careful not to spill contents. Incubate for 30 minutes at room temperature.



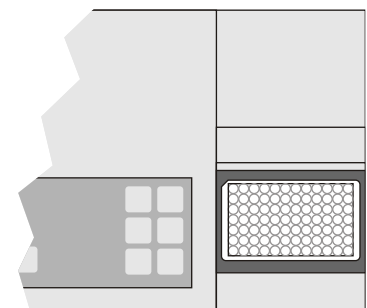
7. Addition of Stopping Solution

Add 100uL of stop solution to the wells in the same sequence as for the substrate solution using a multi-channel pipette or a stepping pipette.



8. Quantification

Measure the absorbance at 450nm for each standard solution and generate a standard curve. Calculate the quantity of AE in a sample from the absorbance reading.



For Ordering or Technical Assistance Contact:

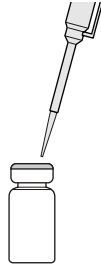
ABRAXIS, LLC
54 Steamwhistle Drive, Warminster, PA 18974
Phone: 215-357-3911 Fax: 215-357-5232
www.abraxiskits.com

LAS Plate Kit Part # 520031

LAS Plate Kit, Concise ELISA Procedure

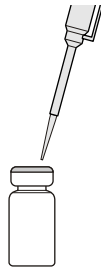
1. Sample Pretreatment

Filter the samples if necessary and add methanol.



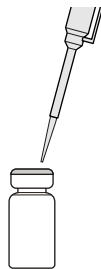
2. Preparation of Standard Solution

Dilute standards with 10% methanol/ water solution.



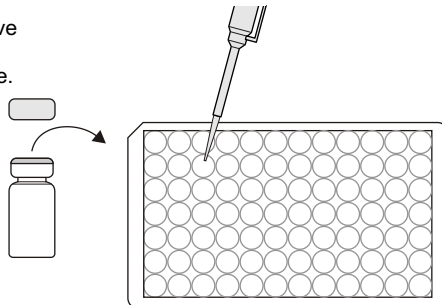
3. Reconstitution of Antigen-enzyme Conjugate

Reconstitute conjugate bottle with 7mL of buffer solution.



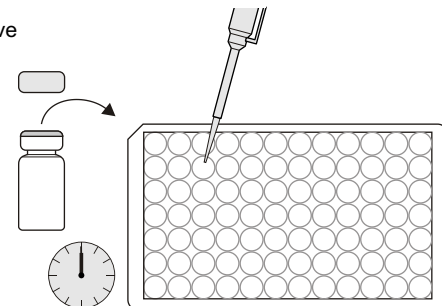
4. Addition of Standard/Sample and Conjugate

Dispense 100uL of the above mixture into each well of an antibody coated microplate. Incubate for 60 minutes.



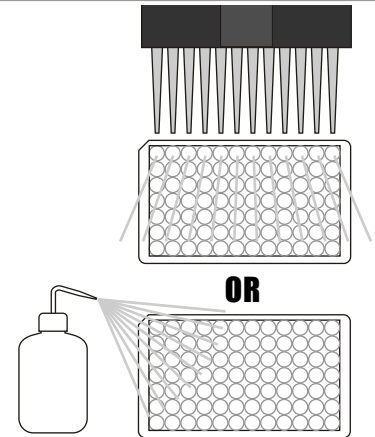
5. Incubation (Competitive Reaction)

Dispense 100uL of the above mixture into each well. Incubate for 60 minutes.



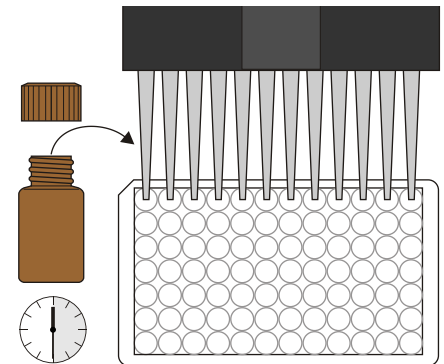
5. Washing Solution

Rise with 300uL of the 1x wash solution and repeat 3 times.



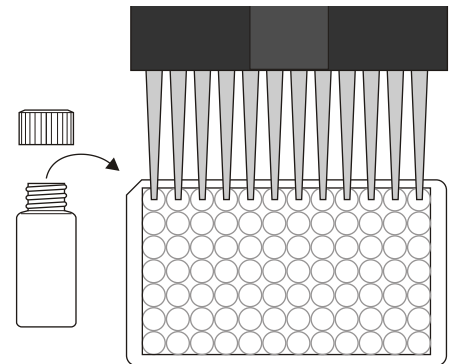
6. Color Reagent and Development

Add 100uL of the Color Solution into each well. Incubate for 30 minutes away from sunlight.



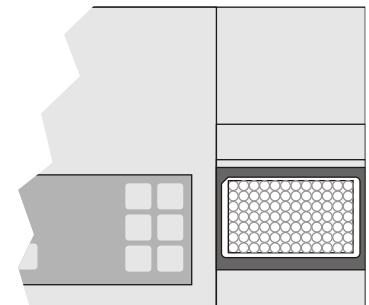
7. Addition of Stopping Solution

Add 100uL of stop solution into each well.



8. Quantification

Measure the absorbance at 450nm.



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