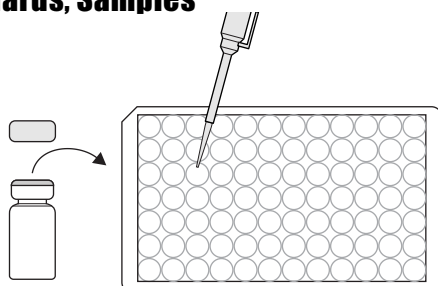


# Sudan Red Plate, Detailed ELISA Procedure

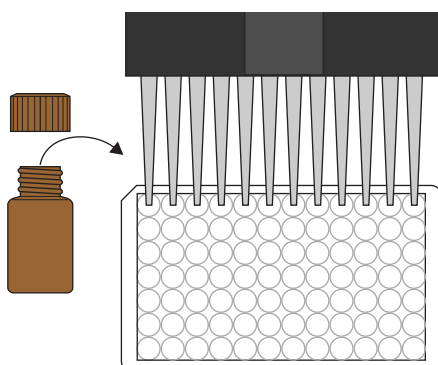
## 1. Addition of Standards, Samples

Add 50  $\mu$ L of the standard solutions or sample extracts into the wells of the test strips according to the working scheme given. We recommend using duplicates or triplicates.



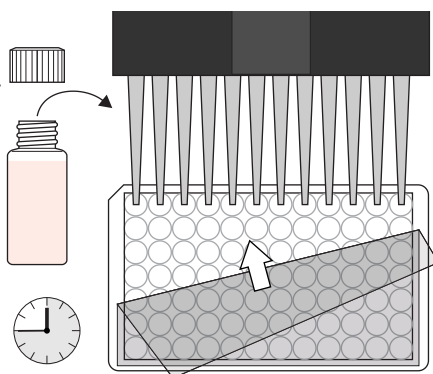
## 2. Addition of Enzyme Conjugate

Add 50  $\mu$ L of the Sudan Red enzyme conjugate to the individual wells successively using a multi-channel pipette or a stepping pipette.



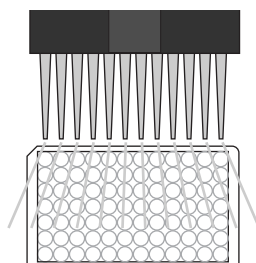
## 3. Addition of Antibody Solution

Add 50  $\mu$ L of the Sudan Red antibody solution to the individual wells successively using a multi-channel 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 45 minutes at room temperature.



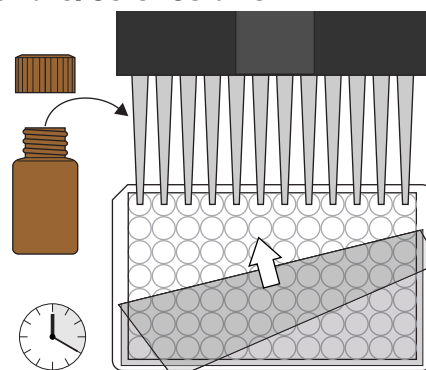
## 4. Washing of Plates

After incubation, remove the covering and vigorously shake the contents of the wells into a sink. Wash the strips four times with a multi-channel pipette or wash bottle using the diluted 1X washing buffer solution. Please use at least 250  $\mu$ L 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.



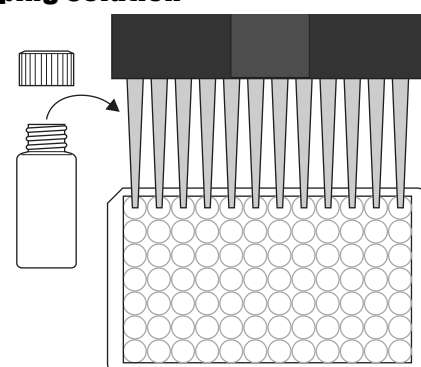
## 5. Addition of Substrate/Color Solution

Add 100  $\mu$ L of substrate/color solution to the individual wells 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. Be careful not to spill contents. Incubate the strips for 20 minutes at room temperature away from direct sunlight.



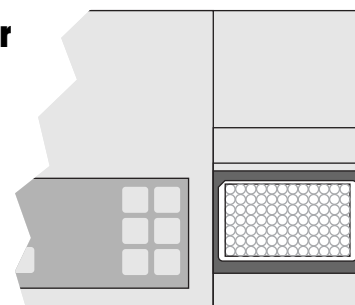
## 6. Addition of Stopping Solution

Add 50  $\mu$ L 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 within 15 minutes. Calculate results.

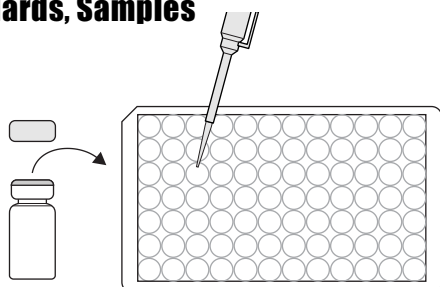


For Ordering or Technical Assistance Contact:  
ABRAXIS, LLC  
54 Steamwhistle Drive, Warminster, PA 18974  
Phone: 215-357-3911 Fax: 215-357-5232  
[www.abraxiskits.com](http://www.abraxiskits.com)

# Sudan Red Plate, Concise ELISA Procedure

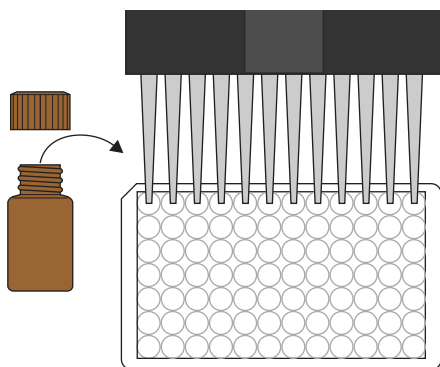
## 1. Addition of Standards, Samples

Add 50  $\mu$ L of standard solutions or sample extracts.



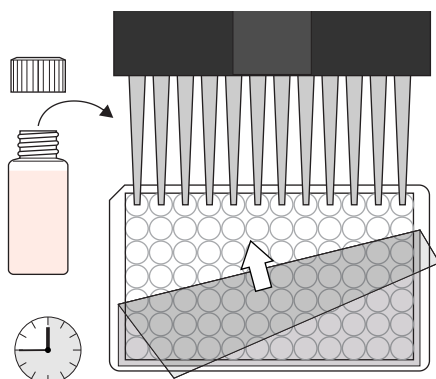
## 2. Addition of Enzyme Conjugate

Add 50  $\mu$ L of enzyme conjugate.



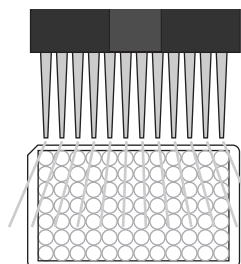
## 3. Addition of Antibody Solution

Add 50  $\mu$ L of the antibody solution. Cover and mix for 30 seconds by moving strip holder in a circular motion on benchtop. Incubate for 45 minutes at room temperature.



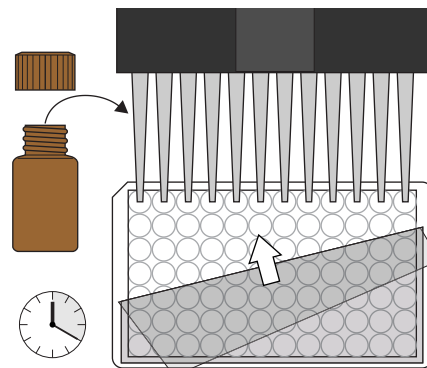
## 4. Washing of Plates

Wash the wells four times with 250  $\mu$ L of diluted 1X washing buffer.



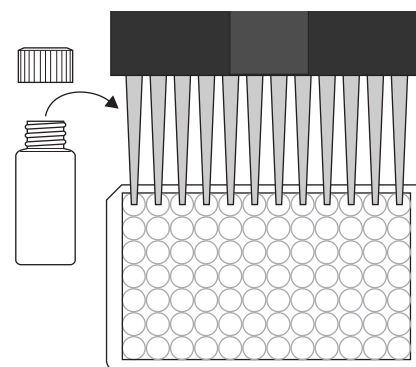
## 5. Addition of Substrate/Color Solution

Add 100  $\mu$ L of substrate/color solution. Cover and mix for 30 seconds by moving strip holder in a circular motion on benchtop. Incubate 20 minutes at room temperature away from direct sunlight.



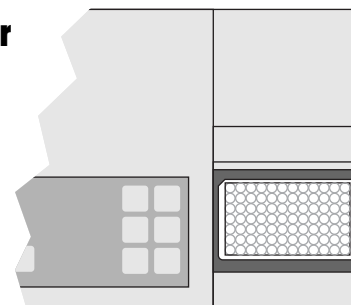
## 6. Addition of Stopping Solution

Add 50  $\mu$ L of stop solution.



## 7. Measurement of Color

Measure color at 450 nm within 15 minutes. Calculate results.



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](http://www.abraxiskits.com)