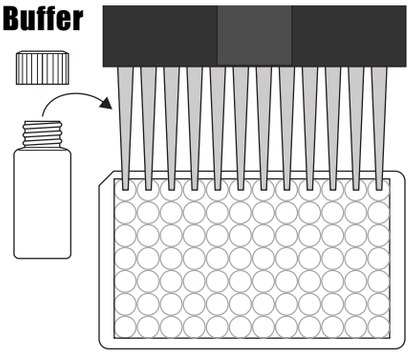
Tetracyclines Plate, Detailed ELISA Procedure

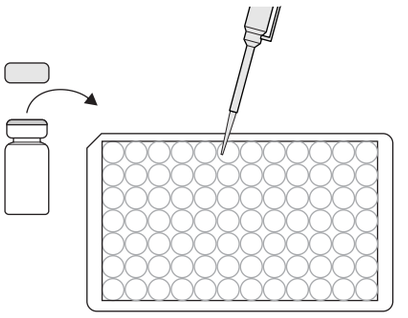
1. Addition of Assay Buffer

Add 50 ul of Assay Buffer to the wells of the test strips successively using a multi-channel pipette or a stepping pipette according to the working scheme given. We recommend using duplicates or triplicates.



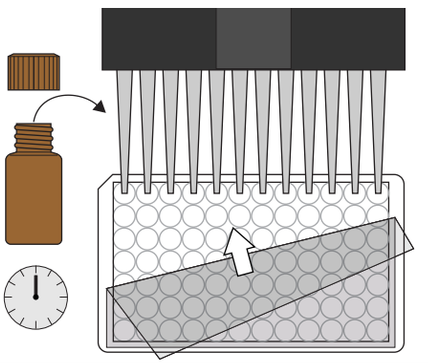
2. Addition of Standards, Samples

Add 100 ul of the standard solutions, control, or samples to the wells of the test strips according to the working scheme given.



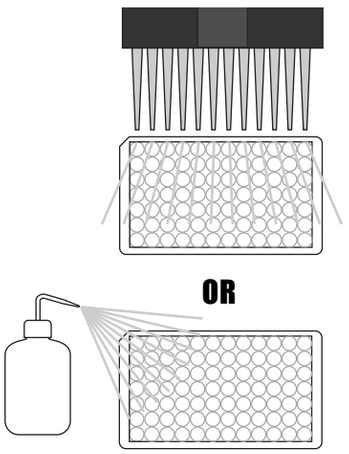
3. Addition of Conjugate Solution

Add 50 uL of the enzyme conjugate 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 60 min at room temperature.



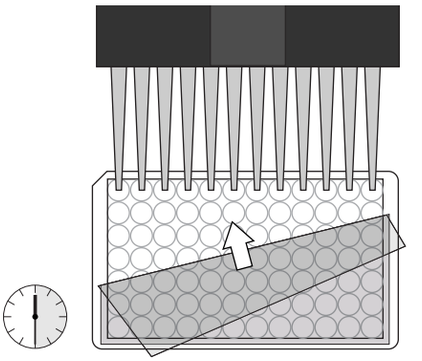
4. Washing of Plates

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 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.



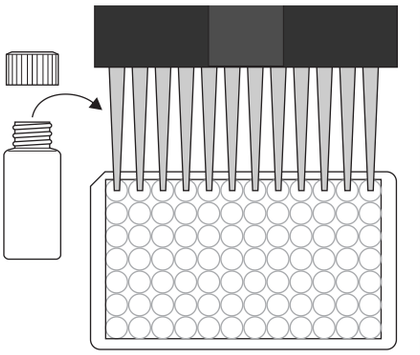
5. Addition of Substrate/Color Solution

Add 150 uL 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 20-30 min at room temperature.



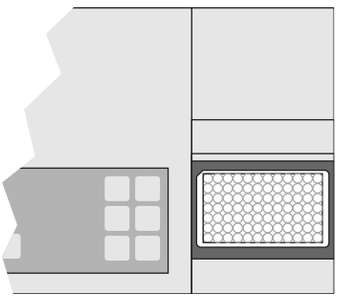
6. Addition of Stopping Solution

Add 100 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.



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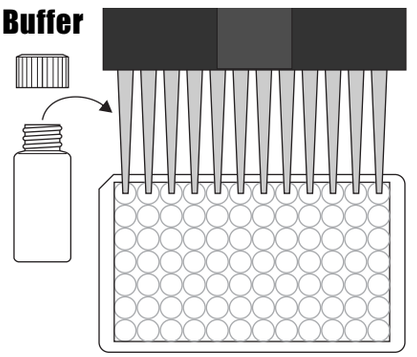
Tetracyclines Plate, Concise ELISA Procedure

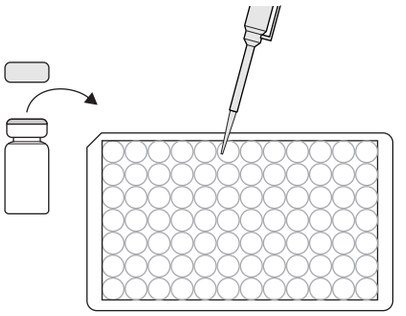
1. Addition of Assay Buffer

Add 50 ul of Assay Buffer.



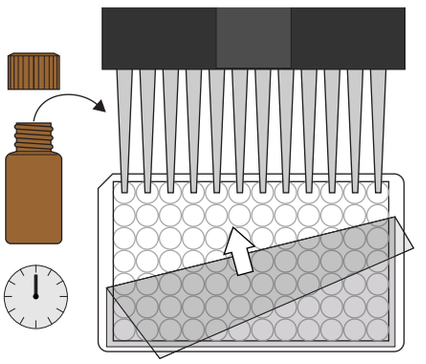
2. Addition of Standards, Samples

Add 100 ul of the standard solutions, control, or samples.



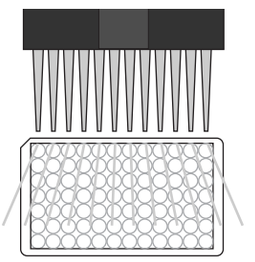
3. Addition of Conjugate Solution

Add 50 uL of the enzyme conjugate. Cover and mix for 30 seconds by rotating on benchtop. Incubate for 60 min at room temperature.

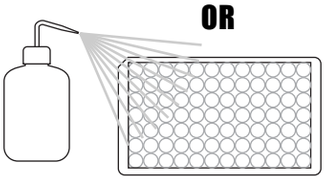


4. Washing of Plates

Wash the plates four times with 250 uL of diluted 1X washing buffer.

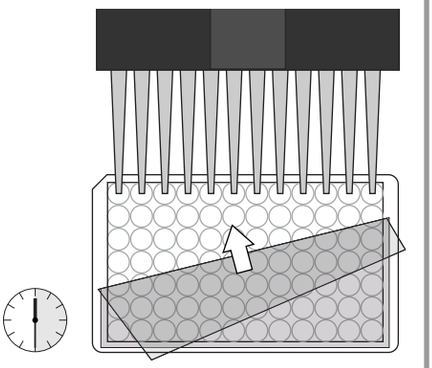


OR



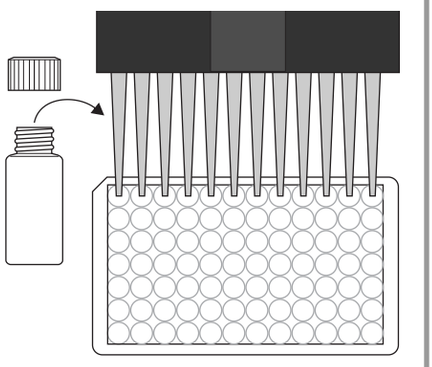
5. Addition of Substrate/Color Solution

Add 150 uL of substrate/color solution. Incubate 20-30 minutes at room temperature and away from direct sunlight.



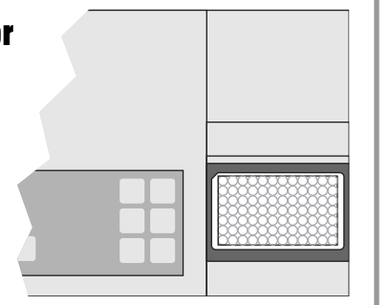
6. Addition of Stopping Solution

Add 100 uL of stop solution.



7. Measurement of Color

Measure color at 450 nm. Calculate results.



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