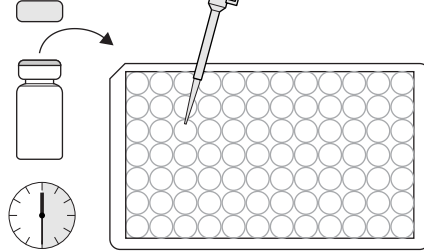


Ochratoxins ES ELISA Kit, Detailed Procedure

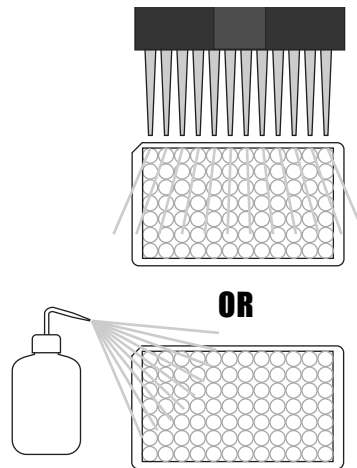
1. Addition of Standards, Samples

Add 100 μ L of the standard solutions or samples into the wells of the test strips according to the working scheme given. We recommend using duplicates or triplicates. Incubate the strips for 30 min. at room temperature.



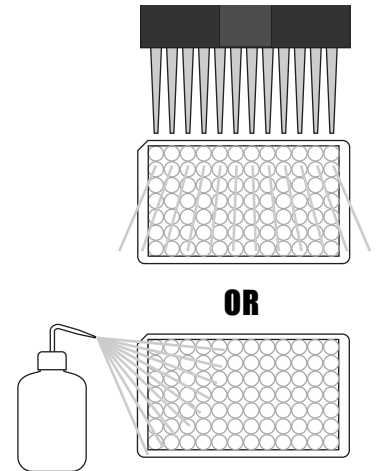
2. Washing of Plates

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 μ 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.



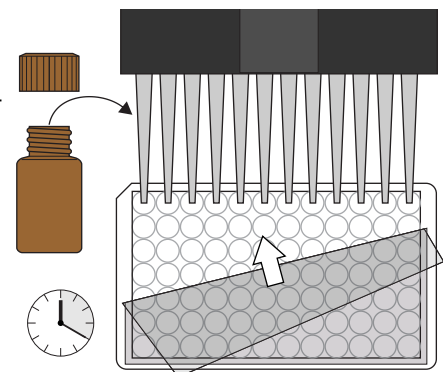
5. Washing of Plates

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 μ 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.



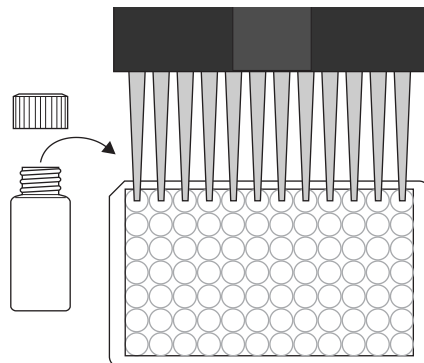
6. Addition of Substrate/Color Solution

Add 150 μ 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 min. at room temperature.



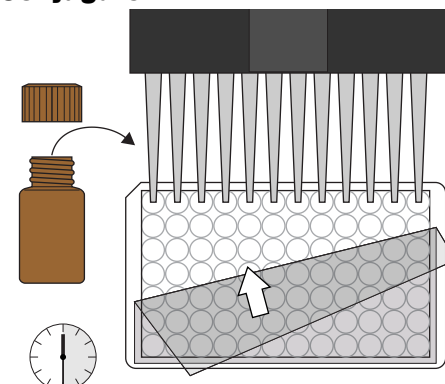
3. Addition of HRP Assay Buffer

Add 50 μ L of the HRP assay buffer to the individual wells successively using a multi-channel pipette or a stepping pipette.



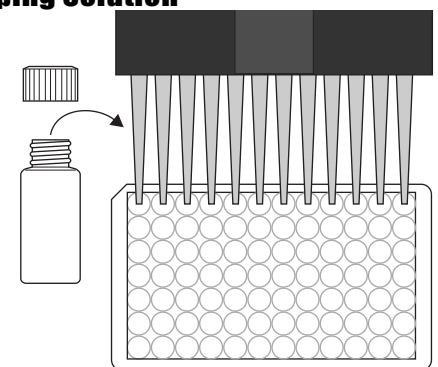
4. Addition of HRP Conjugate

Add 50 μ L of the Ochratoxin HRP conjugate 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 30 min. at room temperature.



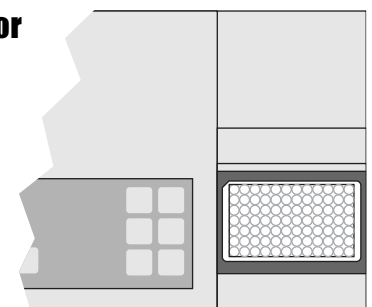
7. Addition of Stopping Solution

Add 100 μ 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.



8. Measurement of Color

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

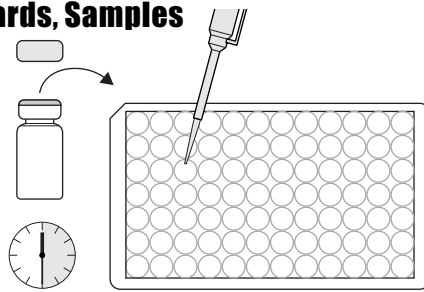


For Ordering or Technical Assistance Contact:
ABRAXIS, LLC
54 Steamwhistle Drive, Warminster, PA 18974
Phone: 215-357-3911 Fax: 215-357-5232
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Ochratoxins ES ELISA Kit, Concise Procedure

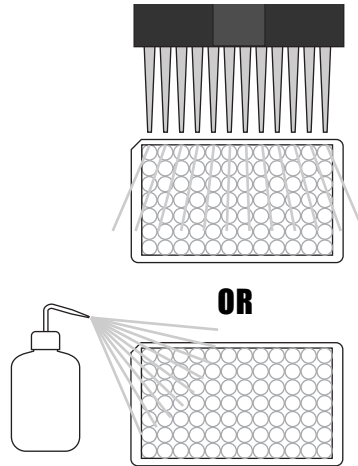
1. Addition of Standards, Samples

Add 100 μ L of the standard solutions or samples. Incubate the strips for 30 min. at room temperature.



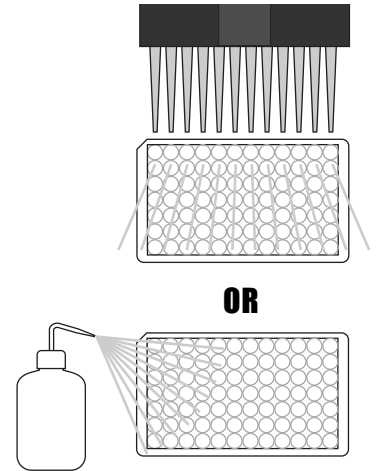
2. Washing of Plates

Wash the plates four times with 250 μ L of diluted 1x washing buffer.



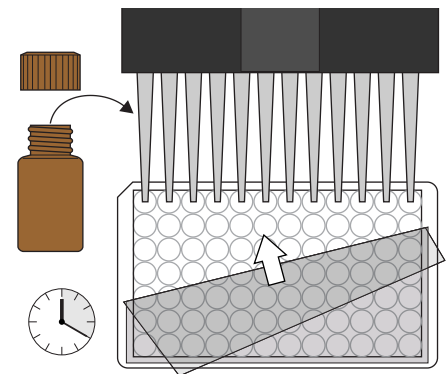
5. Washing of Plates

Wash the plates three times with 250 μ L of diluted 1x washing buffer.



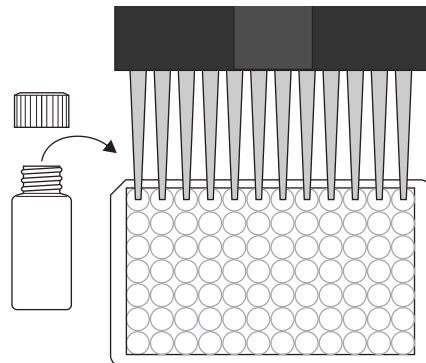
6. Addition of Substrate/Color Solution

Add 150 μ L of substrate/color solutions. Incubate the strips for 20 min. at room temperature.



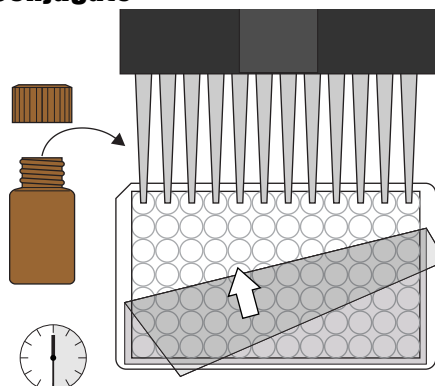
3. Addition of HRP Assay Buffer

Add 50 μ L of the HRP assay buffer.



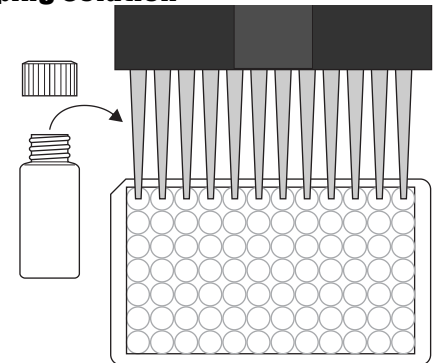
4. Addition of HRP Conjugate

Add 50 μ L of the Ochratoxin HRP conjugate solution. Cover and mix the contents. Incubate the strips for 30 min. at room temperature.



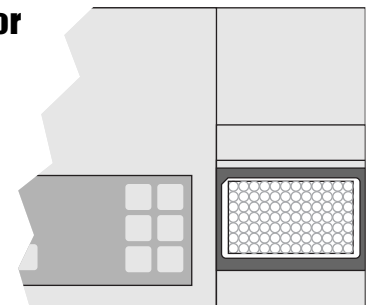
7. Addition of Stopping Solution

Add 100 μ L of stop solution.



8. Measurement of Color

Read the absorbance at 450 nm. Calculate results.



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