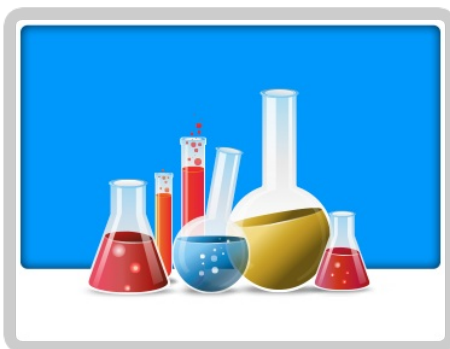


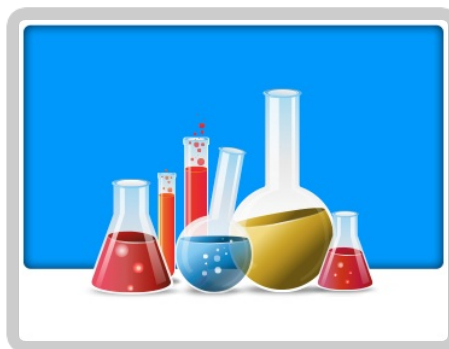
PROCEDURE



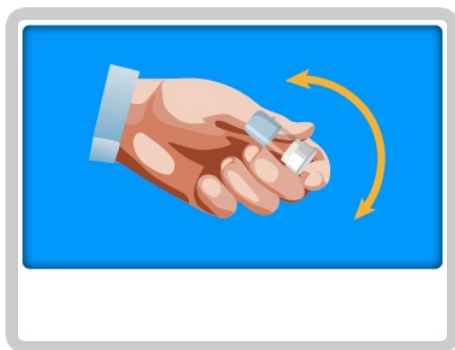
1. Place the cuvettes containing R1 in the incubation cells and perform incubation.



2. Dilute the sample to 1:10. Dilute by using the dedicated dilution kit (*300129).



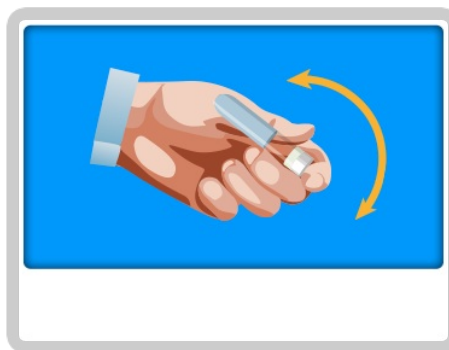
3. Take 100 μ L of oil using the specific pipette and add it to the diluent. Shake the test cuvette by inversion, place it in an incubation cell and leave it to incubate for 2 minutes.



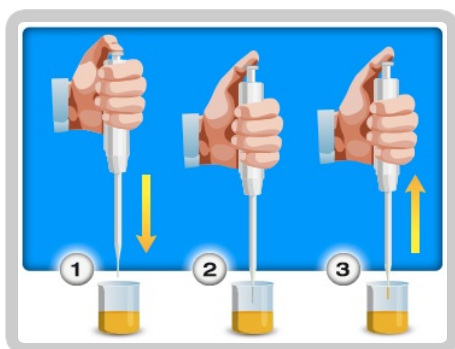
4. Gently shake the cuvette 2-3 times.



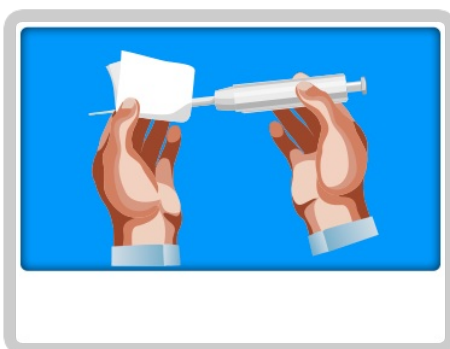
5. Select the cuvette to be read from the display. Place the cuvette in the cell marked with the blue light and press READ.



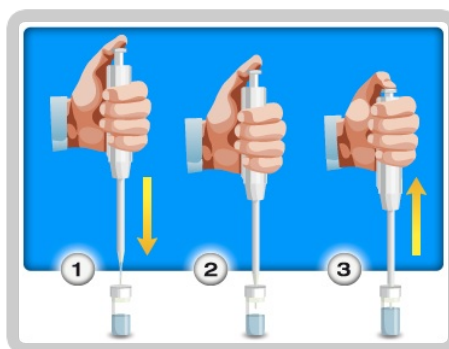
6. Homogenize the sample before collection.



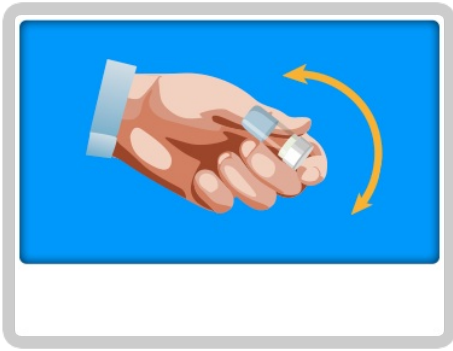
7. Draw the diluted sample with the pipette 2-3 times and release it on blotting paper before collecting it for the test. Then collect 2.5 μ L of diluted sample.



8. Carefully clean the outside of the pipette tip with blotting paper, avoiding contact between the extremity of the tip and the paper.



9. Place the sample in the cuvette. Keeping the tip immersed in reagent, press and release the piston of the pipette several times.



10. Gently shake the cuvette 2-3 times.



11. Place the cuvette in the cell marked with the blue light and press READ to perform the photometric reading.