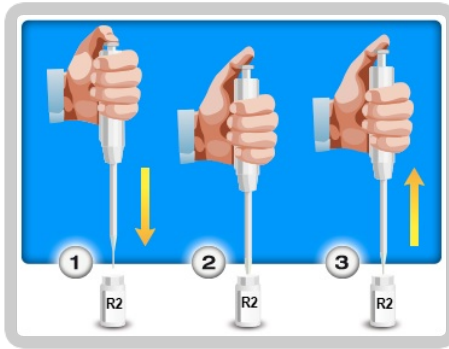


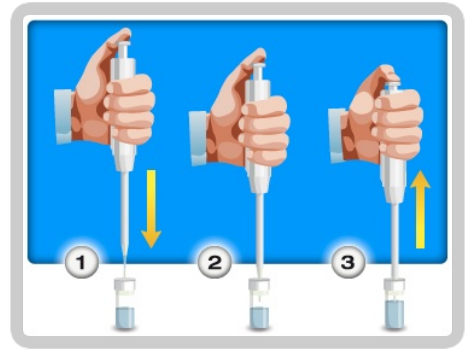
## PROCEDURE



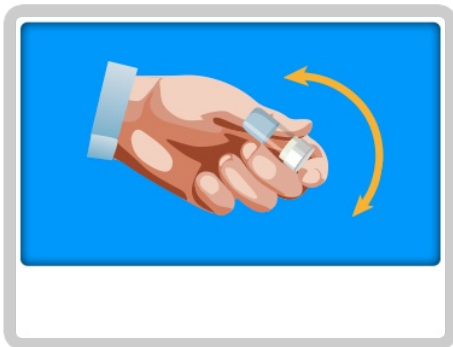
**1.** Place the test tubes containing reagent R1 in one of the incubation cells and let them warm for at least 5 minutes.



**2.** Collect 30  $\mu\text{L}$  of R2 with the pipette.



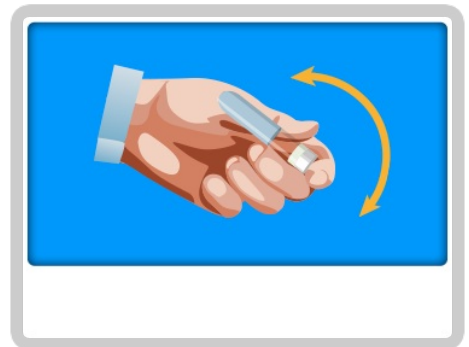
**3.** Add 30  $\mu\text{L}$  of R2 to the cuvette without touching the tip in the liquid, R1 + sample. In case of contamination, replace the tip.



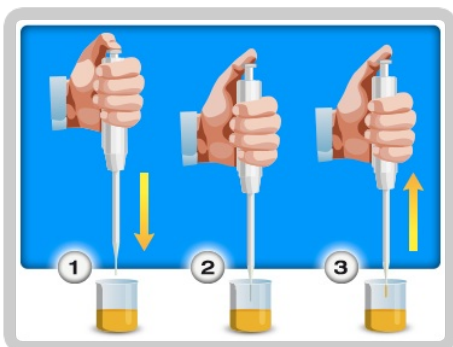
**4.** Gently shake the cuvette 2-3 times.



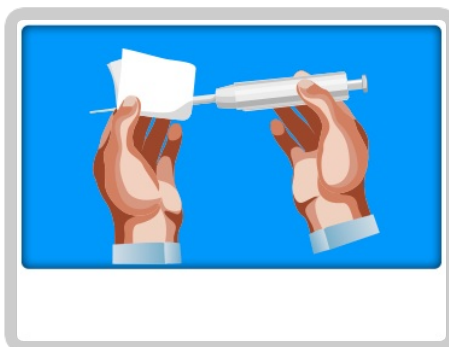
**5.** Select the cuvette to be read from the display. Place the cuvette in the cell marked with the blue light and press READ.



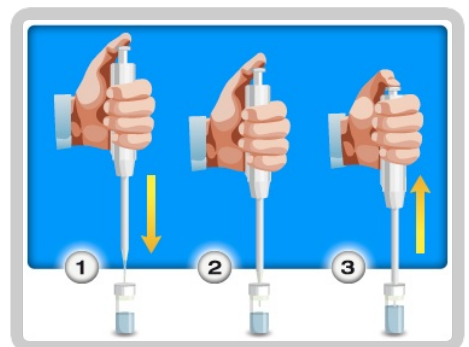
**6.** Homogenize the sample before collection.



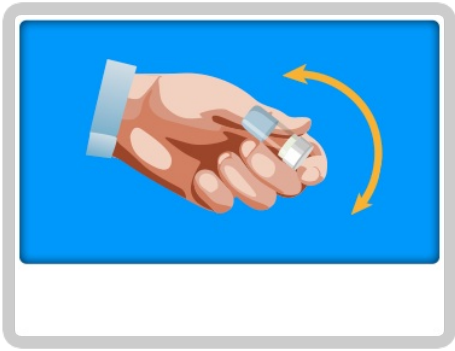
**7.** Draw the sample with the pipette 2-3 times and release it on blotting paper before collecting it for the test. Then collect 5  $\mu\text{L}$  of sample.



**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.