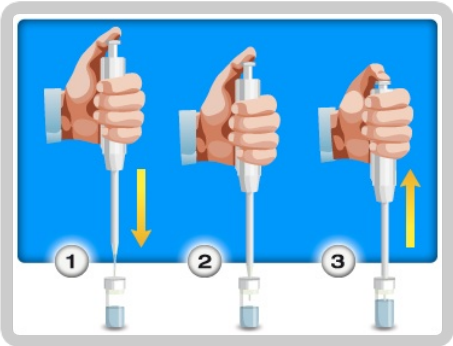
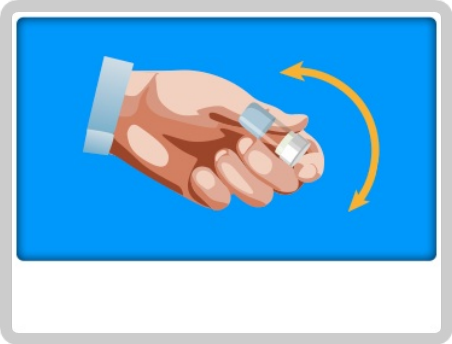
PROCEDURE

1. Centrifuge the must sample before withdrawal.



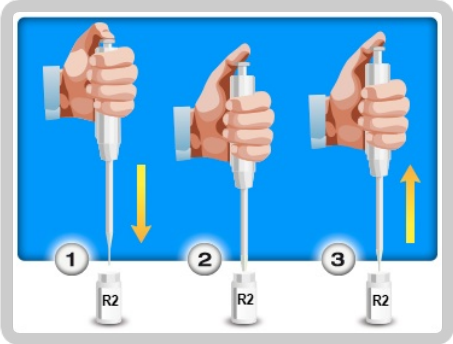
2. Collect 1000 µL of beer with the proper pipette and pour it into an empty cuvette. Close it with the supplied cap.

3. Place the cuvette containing the sample in one of the incubation cells and leave it to incubate for at least 5 minutes.

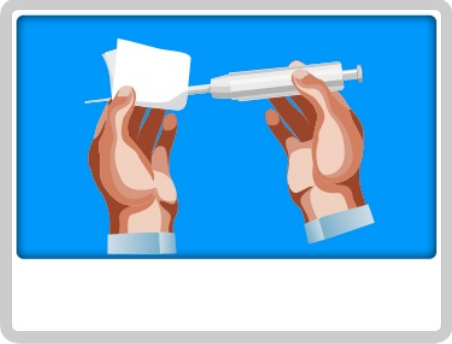


4. Gently shake the cuvette 2-3 times.

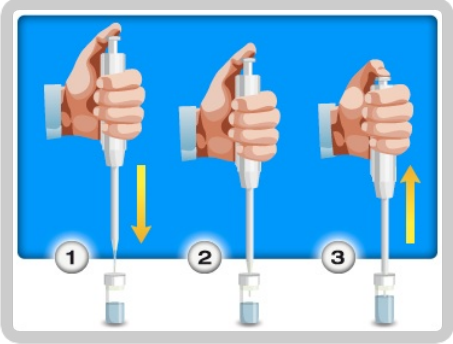
5. Select the cuvette to be read from the display. Place the cuvette in the cell marked with the blue light and press READ.



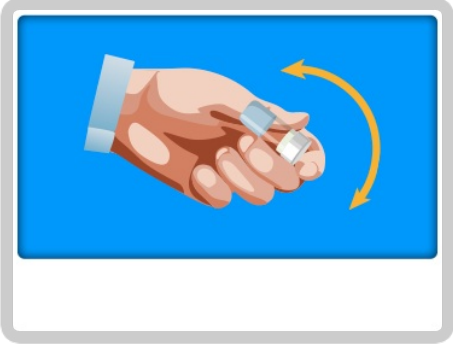
6. Collect 20 µL of R2 with the pipette.



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



8. Place 20 µL of reagent R2 in the cuvette. Keeping the tip immersed in reagent, press and release the piston of the pipette several times.



9. Gently shake the cuvette 2-3 times.

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