



Al-Rasheed University College Pharmacy Department

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# Determination of packed cell volume

Physiology lab #4

Done by:

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# Introduction

- **Blood** consist of two portions:

1-Liquid plasma

2-Solid cellular portion.

Solid portion consist of **red blood cells**, **white blood cells** and **platelets**.

**PCV or HCT** is defined as the **volume of packed RBCs** per unit volume of the whole **blood** after [centrifugation](#)

# Introduction

- **PCV** is the percentage of the number of **packed RBCs** in the whole blood.

- Example #1:

If a **100ml** of blood sample contains a **50% PCV** >>

it demonstrates that **100ml of blood have 50ml** of packed RBC!

Question: 100ml of blood sample contains 40% of PCV

What is the volume (ml) of PCV?

# Clinical implications

- **PCV** is affected by the **shape, number of the RBC and plasma volume**.

**High PCV either indicates:**

**(Increase in number of circulating RBCs) OR (Decrease in plasma volume)** (e.g. cholera due to loss of water in the stool ).

**Low PCV either indicates:**

A low PCV indicates either **decrease in RBC or increase in plasma volume**

# Lower number of **PCV** may indicates:

- An insufficient supply of healthy red blood cells (anemia).
- Acute kidney disease (lower Erythropoietin production lead to less RBCs production by the bone marrow).
- Pregnancy may lead to women having additional fluid in blood. This could potentially lead to a small drop in haematocrit level.

# Higher number of **PCV** may indicates:

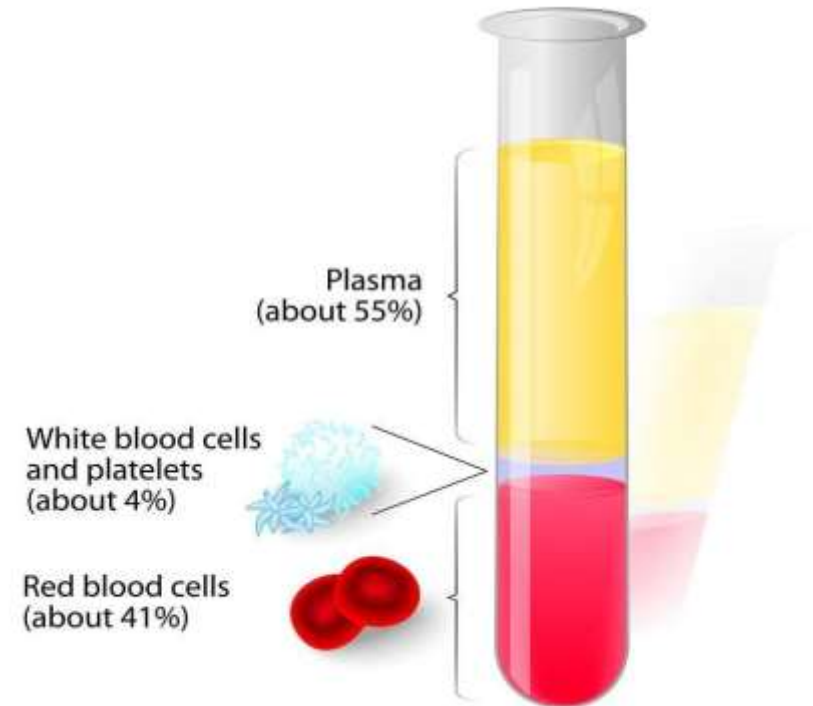
- Abnormal increase in RBC (erythrocytosis)
- A disorder, such as polycythaemia vera causes the body to produce RBCs excessively.
- At higher altitudes → lower oxygen supply in the air → haematocrit levels may increase over time.
- Lung or heart disease, if the body senses low oxygen levels, it will make more RBCs in an effort to increase the amount of oxygen in the blood

# Aim of Experiment

- 1- To assess **PCV values**, methods for determination **PCV value** and clinical importance of **PCV**.
- 2-To determine the volume or the amount of **RBCs** in 100 ml of blood
- 3-To assess whether there is a **sufficient number of circulating RBCs** to transport the required amount of oxygen throughout the body.

# Principle of the experiment

- Hematocrit is derived from Greek words (Haima = blood) and (krites =separate).
- “Hematocrit” means separation of blood where blood cells and plasma are separated by centrifugation.
- When a known volume of blood is centrifuged → Cells being **heavier** will **settle down** leaving a **clear column of plasma above**.





# PCV determination methods

**Microhematocrit  
Method**



**1-Requires less  
blood  
2-less time to get  
values of PCV  
(commonly used).**

**Macrohematocrit  
Method OR  
Wintrobe method**



**1-Time consuming  
2-Requires large  
amount of blood  
3-Has higher degree of  
plasma trapping**

**Automated  
Method**



**Automated  
hematology  
analyzer**

# Materials and instruments

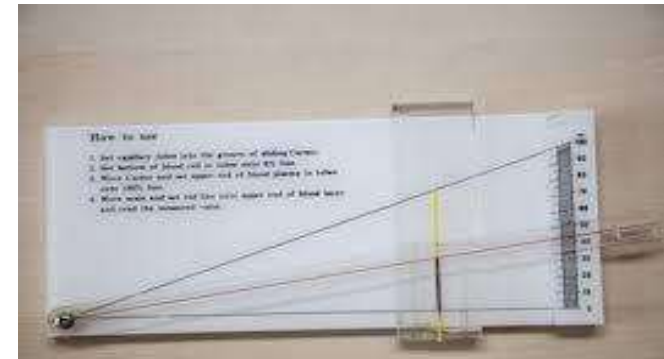
1-Microhematocrit tube (Red capillary tube).

2-Microhematocrit centrifuge device.

3-Plastic seal to seal one end of the capillary tube.

4-Microhematocrit reader.

5-Lancet, Alcohol 70%, and Cotton.



# Procedure

- 1- Clean your finger with 70% alcohol and let it dry for 20-30 sec
- 2-Blood is drawn into the tube by capillary phenomenon.
- 3-Seal the dry end of the tube by plastic seal
- 4-The sealed tube then is placed in the radial grooves of the Microhematocrit centrifuge for 5 min at 10,000 RPM.
- 5-Balance the tubes in the centrifuge with the clay ends facing the outside away from the centre (place the tubes opposite each other in the centrifuge)

# Observation

***A tall upper layer of clear plasma*** - slightly yellow coloured.

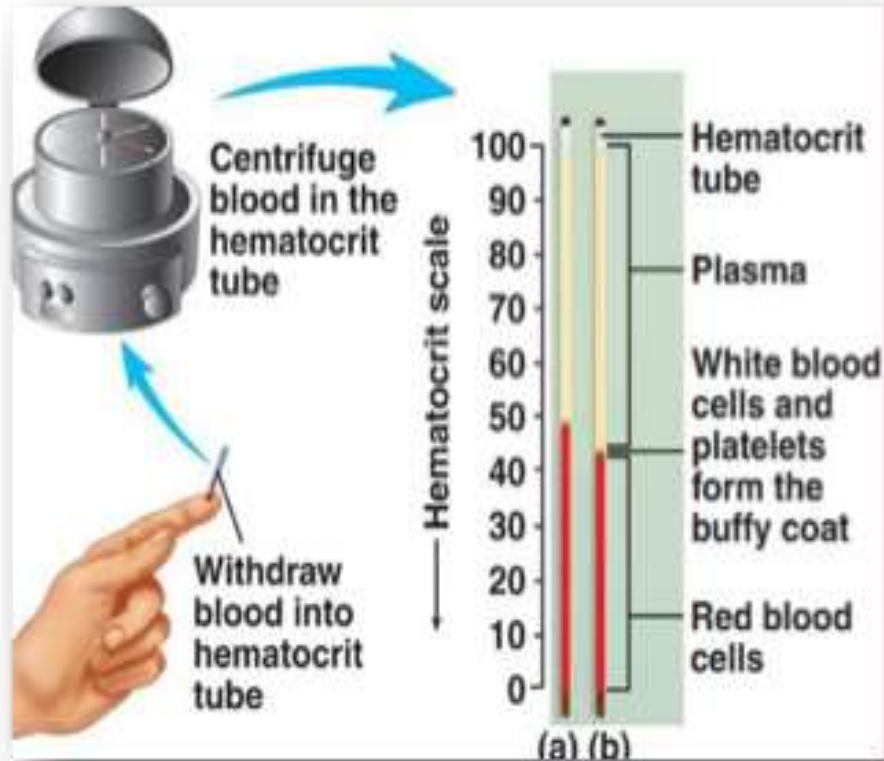
\*Disclaimer\* It shouldn't be pink or red which would indicate hemolysis

***A greyish-white ( buffy layer)*** thin layer (about 1 mm) in thickness consisting of platelets and WBCs.

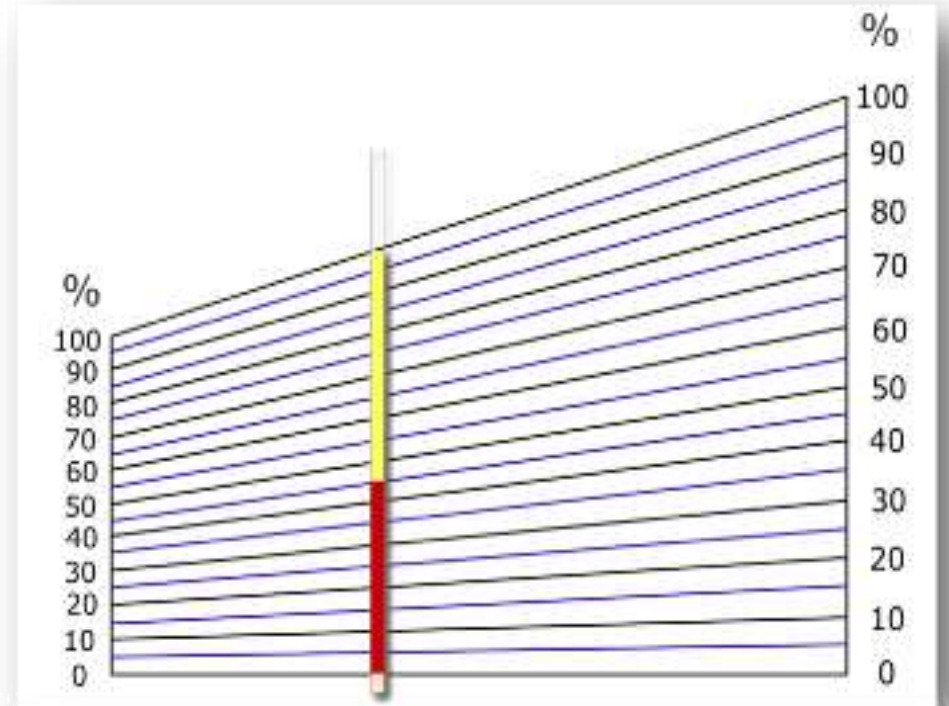
***A tall bottom layer of RBCs*** which have been closely packed together.

# Reading PCV Values

- The capillary tube should be **parallel** to graduation and you should do the following:
- **lower level of the scale (packed RBCs) on zero line of the scale.**
- **Upper level of the scale (plasma) on 100% line of the scale**
- **Do not** include the buffy coat (WBCs and platelets) when reading PCV value.



***Procedure of PCV experiment***



***Reading of PCV value***

# Sources of errors

- 1-Improper sealing of the capillary tube
- 2-Time and speed of centrifugation.
- 3-The buffy coat of the specimen should not be included in the PCV reading, **because its inclusion would falsely elevate the result.**
- 4-Decrease or increase in the readings may be seen if the micro- hematocrit reader is not used properly.
- 5-The microhematocrit centrifuge should never be forced to stop by applying pressure to the metal cover plate!

This will cause → RBCs layer to “sling” forward → falsely elevated value.



**IT'S OVER**

**IT'S FINALLY OVER**