# **ENE 310 ENGINEERING LABORATORY – I HYDROSTATIC BENCH**

The *Hydrostatic Bench* enables the study of the main properties and the behavior of such liquids under hydrostatic conditions, with the aid of some accessories to make the different experiments.

### **Equipment Description**

The equipment consists of a metallic structure assembled on wheels with a panel at the top. In the lower part of the bench there is a tank where water is stored. Water is then sent to a methacrylate tank placed at the upper part of the bench and to other plastic deposit. Two hand-operated pumps are used for such distribution. The methacrylate tank is connected to two communicating tubes on the front panel, enabling to perform some practices; the other deposit placed on the horizontal surface of the bench is necessary for performing the rest of the practices. All water in excess is sent back to the storage tank by the drain. The rest of the equipment consists of the following different elements and independent accessories:

- $\bullet$  Barometer (10)
- Thermometer  $(3)$
- Ubbelohde capillary viscosimeter,  $0.6-3$  cp  $(0c)$
- Ubbelohde capillary viscosimeter, 2-10 cp (I)
- Ubbelohde capillary viscosimeter, 10-50 cp (Ia)
- Ubbelohde capillary viscosimeter, 60-300 cp (IIc)
- 3 graduated cylinders
- Accessory for demonstration of free surface in static conditions (7)
- Bourdon manometers calibration (13)
- $\bullet$  Mercury manometers (9)
- Accessory to determine the metacentric height (FME11)
- Accessory for studying Archimedes´ principle
- Accessory for studying the hydrostatic pressure (FME08) (14)
- Fluid level gauge calibrator  $(16)$
- Set of weights (5, 10, 20, 50, 100, 400, 1000, 2000, 5000 gr.)
- Air pump
- 2 water pumps  $(11 \text{ and } 12)$
- Universal hydrometer (1)
- Chronometer
- Set of measurement cylinders (2 of 600 ml) (4)
- Spare parts for the viscosimeter elements





Figure 2.0.1. Main parts of the Hydraulic Bench

### **Experiment – 1**

### **Density and Specific Gravity Measurements**

#### **Theory**

The density of a substance is defined as the mass of such substance in relation with the volume that it occupies, and it is called " $\rho$ ".

$$
\rho = \frac{mass \ of \ liquid}{volume \ occupied} = \frac{M}{L^3}
$$

We must have into account that the density of a liquid is practically constant, since the volume occupied by a given mass of a liquid is almost invariable. But in the case of gases, the density varies in accordance with the volume occupied (for a mass of such a gas). As a result of that, a liquid can be considered virtually incompressible (except when it is working in critical conditions), while gasses are compressible.

The specific gravity or relative density of a fluid is defined as the ratio between its mass and the mass of the same volume of water at 4ºC, and "S" represents it.

$$
S = \frac{given \, mass \, of \, the \, liquid}{mass \, of \, water \, (same \, volume)}
$$

If V is the volume of a liquid and  $V_w$  the volume of water,  $\rho_1$  is the density of the liquid and  $\rho_a$  is the density of water, then:

$$
S = \frac{\rho_1.V}{\rho_w.V} = \frac{\rho_1}{\rho_w}
$$

#### **Hydrometer**

The two previous properties can be studied using the hydrometer placed in the left hand end of the front panel.

The operation of the hydrometer is based on Archimedes´ principle, which consists in that, when a body is submerged into a liquid, it becomes subject to a vertical force equal to the weight of the liquid that the body displaces.

Thus, a simple hydrometer can consist of a glass tube, closed by an end having a scale within. A small amount of lead, sand or mercury is placed at the bottom for preventing flotation.



**Figure 1.** Scheme of a simple hydrometer

First, the tube must be submerged in water, and the scale marked in the submerged part. Then, the operation is repeated by submerging the tube in another liquid, and the submerged length is marked again

If  $L_w$  = submerged length in water of density  $\rho_w$ 

If  $L_1$  = submerged length in a liquid of density  $\rho_1 \square$ 

then, the weigh of water displaced is equal to  $\rho_w \Box \Box \Box g \cdot A \cdot L_w$  and the weigh of the liquid displaced is equal to  $\rho_l$ ,  $g$ ,  $A$ ,  $L_l$ 

By applying Archimedes´ principle, we shall get:

$$
\rho_w g A L_w = \rho_1 g A L_1
$$

And so,

$$
S = \frac{\rho_1}{\rho_w} = \frac{L_w}{L_1} = \frac{length \, submerged \, in \, water}{length \, submerged \, in \, the \, liquid}
$$

The immersion depth in water was marked in the paper scale as 1.00 and the liquid passed  $L_w/L_1$ .

### **Aim of this Experiment**

To determine density and specific gravity.

### **Necessary devices**

Universal hydrometer.

Open precipitate tubes or cylinder



**Figure 2.** Schematic representation of density and dpecific gravity measurement set-up

# **Procedure**

1. Fill the precipitate tube or cylinder with water in such a way that the hydrometer floats. Check that the submerged length corresponds to 1.00 in the graduated scale.

2. Fill the other three cylinders with the liquids to work with, and note down the scale mark for each one. This value in the scale indicates the specific gravity.

3. Note down the results obtained in the following graph, taking into account the values of the atmospheric pressure and temperature in the moment of performing the practice.



# **Sample Test Results**



### **Experiment -2.1**

### **Viscosity measurement with Ubbelohde viscometer**

### **Theory**

The viscosity of a liquid is defined as the grade or measure in which a fluid opposes the changes of shape when an external force is applied on it. Viscosity depends on molecular cohesion and the activity of the fluid. The gas viscosity, where cohesion of the atoms is low, increases when temperature rises. In liquids, due to the fact that molecular cohesion is higher than their activity (mainly at low temperature), viscosity decreases when temperature rises.

In order to obtain a measure of viscosity, it is necessary to consider the viscous flow of a liquid; for this purpose, two considerations have to be taken into account:

- 1. Maybe there is not a sliding or relative movement.
- 2. Maybe the applied effort is directly proportional to the movement.

In the first case, we have that, in a fluid that is at rest, by definition, there are not shearing stresses between the fluid and the solid in contact with it, neither between the adjacent layers of the fluid itself. Nevertheless, when the fluid is in movement, different speeds appear between the outer faces and the inner faces of the fluid, causing forces that exert a friction. If one of the body faces moves with a speed *u* and the other does it with a speed *u+du*, then the ratio of the force applied to the movement or the speed gradient equals to *du/dy.*

In the second case, the effort applied is proportional to  $du/dy$ , that is,  $\tau = \ln 2 \text{.}du/dy$ , where  $\eta$ is a proportionality coefficient called viscosity coefficient.

### **Viscosity measurement**

The measurement of flux velocity through a capillary tube of known radius allow us obtain the viscosity  $\Box$  for a liquid or solid, using the equations:

$$
\frac{V}{t} = \frac{\pi r^4}{8\eta} \frac{P_1 - P_2}{y_1 - y_2}
$$
 liquid laminar flow (1)  

$$
\frac{dn}{dt} = \frac{\pi r^4}{16\eta RT} \frac{P_1^2 - P_2^2}{y_1 - y_2}
$$
 ideal gas laminar flow (2)

where V is liquid volume that cross a transversal section of the tube in t time and  $\Box$  P<sub>2</sub>  $\Box$  P<sub>1</sub>)/ (y<sub>2</sub>  $\Box$  y<sub>1</sub>) is the pressure gradient along the tube (P<sub>1</sub> pressure at point y<sub>1</sub> and P<sub>2</sub> pressure at point y<sub>2</sub>, therefore y<sub>2</sub>-y<sub>1</sub> is tube longitude), n viscosity, R ideal gas constant, r tube radius, T temperature in K, dn/dt flux velocity in moles per time unit.



**Figure 3.** Fluid moving in a cylindrical tube. The shadowed section is used to probe the Poiseuile´s law.

A convenient method for viscosity determination of a liquid is using **Ostwald viscosimeter**, a capillary tube joined to a lower bulb L and an upper bulb A, as you can see in next figure forms it.



Figure 4. Ostwald viscometer. Is measured the time that use the liquid going from level M2 to M1.

First, lower bulb L is filled with sample solution, getting it into the viscometer through A. Liquid is sucked through branch B until liquid level goes up upper bulb, taking care of avoiding the formation of air bubbles.

It is measured the time t that a liquid delay on passing from mark M1 to mark M2 in liquid level, while the liquid flows through the capillary tube.

Then the viscometer is filled with a liquid of knowing viscosity, using the same volume, and again is measured t. Pressure exercised by liquid through the tube is  $\rho gh$  (where  $\rho$  is liquid density, g gravity acceleration and h the difference between the arms of the viscometer).  $\rho gh$ substitute  $P_1-P_2$  Poiseuille's law (Equation 1). Because h change in time, flux velocity change and it should be written like:

$$
\frac{dV}{dt} = \frac{\pi r^4}{8\eta} \rho g h (y_2 - y_1)
$$

Consider  $h_0$  as the level difference when t=0 and the liquid level in left arm is in mark A. Since in all experiment is used the same liquid volume,  $h_0$  is constant. h variation from its initial valour  $h_0$  is function of the volume V that has fluxed through the viscosimeter: hh0=f(V), where f function depends on the viscosimeter geometry. We have:

$$
[h_o + f(V)]^{-1}dV = \left[\frac{\pi r^4 \rho g}{8\eta(y_2 - y_1)}\right]dt
$$

$$
\int_0^V \frac{1}{h_o + f(V)} \, dV = \frac{\pi r^4 g}{8(y_2 - y_1)} \frac{\rho}{\eta} \, t
$$

where V' is the volume that flux in t time when the liquid level goes from A to B. Since V' and f(V) are the same in all the experiments made on the same viscosimeter, the previous volume integral is constant, therefore  $p t / \eta$  is constant in all experiments. If we consider two different liquids a and b, we have,

$$
\frac{\rho_a}{\eta_a} t_a = \frac{\rho_b}{\eta_b} t_b
$$

regrouping

$$
\frac{\eta_b}{\eta_a} = \frac{\rho_b}{\rho_a} \frac{t_b}{t_a}
$$

If we know  $\eta_a$ ,  $\rho_a$  and  $\rho_b$ , we can find  $\eta_b$ .

# **Aim of this Experiment**

Determine the viscosity of different liquids at atmospheric pressure and environmental temperature.

## **Necessary devices**

- Ubbelohde capillary viscosimeter, 0.6-3 cp
- Ubbelohde capillary viscosimeter, 2-10 cp
- Ubbelohde capillary viscosimeter, 10-50 cp
- Ubbelohde capillary viscosimeter, 60-300 cp
- Chronometer
- Hydrometer
- Thermometer



**Figure 2.** Viscosimeter provided with the equipment

# **Procedure**

The liquids to be studied are:

- Car Motor oil

- Glycerol

- Castor oil

1. Find in tables four liquids of known viscosity, each one inside the measurement range of each viscosimeter.

2. Fill each Ubbelhode capillary viscosimeter, with the same volume of liquid of known viscosity and density, and write down the time used by the liquid of going down the viscosimeter.

3. Make four problem samples aliquots, with the same volume as used with known viscosity solutions. Measure their falling time in each viscosimeter. In some cases the liquid will fall too fast to take any measurement, and in others it will probably spend too much time. Avoid these liquids.

4. Write down the existing atmospheric pressure and temperature in that moment in the laboratory. With the aid of the data and expressions given hereafter, complete the following table:

Barometric Pressure .. mm Hg

Temperature .. º C

Car Motor oil density (depending on the manufacturer) ...............................g/cm³

Glycol density  $1.25$  g/cm<sup>3</sup>

Castor oil density  $0.95$  g / cm<sup>3</sup>

5. Write down in next table the values obtained with solutions of known viscosity and density.



6. Now, repeat again the experience, with the problem samples, and fill next table with the obtained values, using previous obtained data and equations.



### **Experiment -2.2**

### **Viscosity measurement with falling body method**

#### Falling Sphere Method

The falling sphere viscometer is one of the earliest and least involved methods to determine the absolute shear viscosity of a Newtonian fluid. In this method, a sphere is allowed to fall freely a measured distance through a viscous liquid medium and its velocity is determined. The viscous drag of the falling sphere results in the creation of a restraining force, F, described by Stokes' law:

$$
F=6\pi\eta r_{s}U_{t}
$$

where rs is the radius of the sphere and Ut is the terminal velocity of the falling body. If a sphere of density  $\rho_2$  is falling through a fluid of density  $\rho_1$  in a container of infinite extent, then by balancing the below equation with the net force of gravity and buoyancy exerted on a solid sphere, the resulting equation of absolute viscosity is:

$$
\eta = 2gr_s^2 \frac{(\rho_2 - \rho_1)}{9U_t}
$$

The above equation shows the relation between the viscosity of a fluid and the terminal velocity of a sphere falling within it. Having a finite container volume necessitates the modification of this equation to correct for effects on the velocity of the sphere due to its interaction with container walls (W) and ends (E). Considering a cylindrical container of radius r and height H, the corrected form of given equation can be written as:

$$
\eta = 2gr_s^2 \frac{(\rho_2 - \rho_1)W}{(9U_t E)}
$$

$$
W = 1 - 2.104 \left(\frac{r_s}{r}\right) + 2.09 \left(\frac{r_s}{r}\right)^3 - 0.95 \left(\frac{r_s}{r}\right)^5
$$

$$
E = 1 + 3.3 \left(\frac{r_s}{H}\right)
$$

The wall correction was empirically derived and is valid for  $0.16 \leq \text{rs/r} \leq 0.32$ . Beyond this range, the effects of container walls significantly impair the terminal velocity of the sphere, thus giving rise to a false high viscosity value. The following figüre contains a schematic diagram of the falling sphere method and demonstrates the attraction of this method — its simplicity of design. The simplest and most cost-effective approach in applying this method to transparent liquids would be to use a sufficiently large graduated cylinder filled with the liquid. With a distance marked on the cylinder near the axial and radial center (the region least influenced by the container walls and ends), a sphere (such as a ball bearing or a material that is nonreactive with the liquid) with a known density and sized to within the bounds of the container correction, free falls the length of the cylinder. As the sphere passes through the marked region of length d at its terminal velocity, a measure of the time taken to traverse this distance allows the velocity of the sphere to be calculated. Having measured all the parameters of the corrected equation, the shear viscosity of the liquid can be determined. This method is useful for liquids with viscosities between 10–3 Pa·s and 105 Pa·s. Due to the simplicity of design, the falling sphere method is particularly well suited to high pressure– high temperature viscosity studies.



Figure: A schematic demonstration of falling sphere viscometer.

# **Experiment -4**

## **Pressure center in a smooth surface**

### **Aim of this Experiment**

To determine the position of the pressure center on the rectangular face of the float

### **Necessary devices**

Hydrostatic Pressure device or hydrostatic device.







**Figure 5.** Hydrostatic Pressure device or hydrostatic device

### **Procedure**

1. Measure and note down the dimensions designed as a, L, d, and b; the last corresponding to the flat surface placed at the end of the quadrant.

2. With the receiver placed on the bench, place the balance arm on the support (sharp profile). Hang the pan at the end of the arm.

3. Connect a length of flexible hose to the receiver draining cock and connect the other end to drain.

4. Level the receiver by properly acting on the support feet, which is adjustable, while the "bubble level" is observed.

5. Displace the counterweight of the arm until getting the arm to be horizontal.

6. Close the drain cock in the bottom of the receiver.

7. Introduce water in the receiver until its free surface is tangent to the lower edge of the quadrant. The fine adjustment of that level can be achieved by slightly overreaching the established filling and then slowly draining through the cock.

8. Place a calibrated weight on the balance pan and slowly add water until the balance arm recovers the horizontal position. Record the water level, indicated in the quadrant, and the value of the weight placed on the pan.

9. Repeat the operation above several times, increasing progressively the weight in the pan until, the balance arm is at level, the level of the free water surface becomes flush with the upper edge of the flat rectangular surface that the end of the quadrant presents.

10. From this point on, and in the order inverse to the operation above of placing the weights on the pan, the weight increments given in each step are removed, the arm is leveled (after every removal) by using the drain cock and the weight in the pan and the water level values are recorded.



For y < d (partial immersion), calculate the practical ant the theoretical value of  $m/v^2$  using the equation:

 $m/y^2 = p.b/2L (a+d-y/3).$ 

The slope of this graph must be  $-p.b/2L$ , and its intersection with the coordinate axis  $p.b$  $(a+d)/2L$ .

See the discrepancies in a reasoned way, if any, between the average values measured and the values obtained with the equations above



# **Appendix – I Useful Data**

**Table 1.** Table of the atmospheric pressure in function of the height

