

PHYSICS LABORATORY MANUAL
for
First semester of B.E. Programme



Department of Physics
BIRLA INSTITUTE OF TECHNOLOGY
Mesra - 835215, Ranchi, India

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EXPERIMENT NO. 0

1.1 Introduction to Uncertainty

In introductory lab work, such as in Physics labs, you usually know in advance what the result is supposed to be. You can compare your actual result with the anticipated result, and calculate an actual error value. In real-world laboratory work, on the other hand, you usually *don't* know in advance what the result is supposed to be. If you *did*, you probably wouldn't be doing the experiment in the first place! When you state your final result, it's important to state also, how much you think you can trust that result, in the form of a numerical uncertainty (or error in measurement). For example, you might state the volume of an object as

$$V = 43.25 \pm 0.13\text{cm}^3 \quad (1.1)$$

When we state the uncertainty in this form, without further elaboration, it generally means that we think that the true value has about a 68% chance of being within that range. A more precise statement would include the confidence level of the uncertainty range, which might be 68% or 95% or even 99%. Usually, in an experiment we measure some number of quantities directly, and combine them mathematically to get a final result. Therefore, estimating the final uncertainty usually involves two steps. First, we must estimate the uncertainties in the individual quantities that we measure directly. Second, we must combine those uncertainties to get the overall uncertainty, in a way that corresponds to the way that we combine individual measurements to get the final result.

1.2 Estimating the Uncertainty in a Single Measurement

1.2.1 Normal analog scale

(e.g. meter stick) Estimate the final digit by interpolating between the smallest scale divisions, and make the uncertainty ± 1 or ± 2 in that last digit (use your judgment in deciding).

1.2.2 Analog scale with vernier

(e.g. vernier caliper or micrometer) Use the vernier scale to get the last digit, and make the uncertainty ± 0.5 of that last digit.

1.2.3 Digital scale

(e.g. digital multimeter) If the reading is steady, make the uncertainty ± 0.5 of the last digit; otherwise take several instantaneous readings, average them, and find the standard deviation of the mean as described below.

1.3 Estimating the Uncertainty in an Averaged Measurement

If you can make several measurements x_1, x_2, \dots, x_N , calculate the mean, \bar{x} , and use that as "the" measurement. Then calculate the standard deviation of

$$\sigma_m = \frac{\sqrt{(x_1 - \bar{x})^2 + (x_2 - \bar{x})^2 + (x_3 - \bar{x})^2 + \dots + (x_N - \bar{x})^2}}{N} \quad (1.2)$$

the mean and use this as the uncertainty, Δx . If your calculator has a standard deviation function, divide its result by \sqrt{N} to get the standard deviation of the mean.

2 Combining Uncertainties in Calculated Results

In the following equations, Δx means the absolute uncertainty in which is the number you get from one of the methods above; it has units just like the measurement itself has. $\Delta x\%$ means the percent (or fractional) uncertainty in x , which is the uncertainty expressed as a percentage or fraction of the measurement; it has no units.

1.4.1 Addition and Subtraction

If $z = x + y$ or $z = x - y$,

$$\Delta z = \sqrt{\Delta x^2 + \Delta y^2} \quad (1.3)$$

If you're adding and subtracting more variables, simply add more terms inside the square root.

1.4.2 Multiplication and Division

If $z = xy$ or $z = x/y$

$$\frac{\Delta z}{z} = \sqrt{\left(\frac{\Delta x}{x}\right)^2 + \left(\frac{\Delta y}{y}\right)^2} \quad (1.4)$$

or (same thing in different notation).

$$\Delta z\% = \sqrt{(\Delta x\%)^2 + (\Delta y\%)^2} \quad (1.5)$$

If you're multiplying and dividing more variables, simply add more terms inside the square root.

1.4.3 Powers, Including Roots

If $z = x^n$

$$\frac{\Delta z}{z} = n \left(\frac{\Delta x}{x}\right) \quad (1.6)$$

1.4.4 More Complicated Calculations

Sometimes you can combine the three rules given above, doing the calculation one step at a time, combining uncertainties as you go along, and switching back and forth between absolute and percent uncertainties as necessary. However, you cannot do this if the same variable appears more than once in the equation or calculation, or if you have situations not covered by the rules given above, such as trig functions. In such cases you must use the general procedure given below.

The following table gives an idea about the relation between error and actual equation:

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Table 1: Some examples

S. no.	Relation between Z and (A, B)	Relation between error Δz and $(\Delta A, \Delta B)$
1.	$Z = A + B$	$(\Delta Z)^2 = (\Delta A)^2 + (\Delta B)^2$
2.	$Z = A - B$	$(\Delta Z)^2 = (\Delta A)^2 + (\Delta B)^2$
3.	$Z = AB$	$\frac{(\Delta Z)^2}{Z^2} = \frac{(\Delta A)^2}{A^2} + \frac{(\Delta B)^2}{B^2}$
4.	$Z = A/B$	$\frac{(\Delta Z)^2}{Z^2} = \frac{(\Delta A)^2}{A^2} + \frac{(\Delta B)^2}{B^2}$
5.	$Z = A^n$	$\frac{(\Delta Z)}{Z} = n \frac{\Delta A}{A}$
6.	$Z = \ln A$	$\Delta Z = \frac{\Delta A}{A}$
7.	$Z = e^A$	$\frac{(\Delta Z)}{Z} = \Delta A$

1.5 General Procedure

If $z = f(x, y)$, first calculate the differences caused by the uncertainty in each variable separately;

$$(\Delta z)_x = f((x + \Delta x), y) - f(x, y) \tag{1.7}$$

$$(\Delta z)_y = f(x, (y + \Delta y)) - f(x, y)$$

Then combine the differences to get the total uncertainty:

$$\Delta z = \sqrt{(\Delta z)_x^2 + (\Delta z)_y^2} \tag{1.8}$$

If there are more variables, extend these equations appropriately by adding more terms. If a variable occurs more than once in the formula for $f(x, y)$, change all occurrences simultaneously when calculating the



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difference for that variable.

To illustrate the procedure for calculation the error in an experiment, we will work out the average (mean) value x and the standard deviation of the mean, σ and the standard deviation of an individual data point, σ , using the position measurements in the accompanying Table 2.

Table 2: Position measurements

x_i (m)	$(x_i - \bar{x}_i)$	$(x_i - \bar{x}_i)^2$
15.68	0.15	0.0225
15.42	0.11	0.0121
15.03	0.50	0.2500
15.66	0.13	0.0169
15.17	0.36	0.1296
15.89	0.36	0.1296
15.35	0.18	0.0324
15.81	0.28	0.0784
15.62	0.09	0.0081
15.39	0.14	0.0196
15.21	0.32	0.1024
15.78	0.25	0.0625
15.46	0.07	0.0049
15.12	0.41	0.1681
15.93	0.40	0.1600
15.23	0.30	0.0900
15.62	0.09	0.0081
15.88	0.35	0.1225
15.95	0.42	0.1764
15.37	0.16	0.0256
15.51	0.02	0.0004

From the above table we can make the following calculations:

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$$N = 21 \sum_{i=1}^N x_i = 326.08\text{m}, \quad \sum_{i=1}^N (x_i - \bar{x})^2 = 1.61998\text{m}$$

And then evaluate the following quantities:

$$\bar{x} = \frac{\sum_{i=1}^N x_i}{N} = \frac{326.08}{21} = 15.53 \text{ m} \quad (1.9)$$

$$\bar{\sigma} = \sqrt{\frac{\sum_{i=1}^N (x_i - \bar{x})^2}{N(N-1)}} = \sqrt{\frac{1.6201}{20.21}} = 0.062 \text{ m} \quad (1.10)$$

(1.10)

$$\sigma = \sqrt{\frac{\sum_{i=1}^N (x_i - \bar{x})^2}{N-1}} = \sqrt{\frac{1.6201}{20}} = 0.063 \text{ m} \quad (1.11)$$

(1.11)

The error or spread in individual measurements is σ 0.28 m. But for the mean $\bar{x} \pm \bar{\sigma}$ 15.53 \pm 0.06 m. This says the average is 15.53 m which has an error of 0.06m. Or putting it another way, there is about a 68% probability that the true value of x falls, in the range 15.47 m to 15.59 m. In some cases, the fractional error σ / \bar{x} , or relative error, is of more interest than the absolute value of σ . It is possible that the size of σ is large while the fractional error is small. Note that increasing the number of individual measurements on the uncertainty of the average reduces the statistical uncertainty (random errors); this improves the "precision". On the other hand, more measurements do not diminish systematic error in the mean because these are always in the same direction; the "accuracy" of the experiment is limited by systematic errors.

1.6 Suggested Experiments:

1. Measure the diameter of a wire using a screw gauge at 10 different places on the wire. Calculate the standard deviation in your measurements.
2. Measure the thickness of a table top at using a scale in cm. Calculate the error in your measurements.

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3. Measure the period of oscillations of a pendulum using your wrist watch and record your data ten times. Estimate the standard deviation and error in your measurements.
4. Ask your partner to drop a solid object at a same height for 10 times. Measure the time of flight with your wrist watch. The same can be repeated by your other partners also. Compare the standard deviation of each of your measurements.

SAMPLE ORAL QUESTIONS

1. What could be the various possible errors in a measurement?
2. What do you mean by significant digits?

EXPERIMENT NO. 1

Aim: To determine the frequency of AC Mains with the help of sonometer.

Apparatus: Sonometer with non-magnetic wire, ammeter, step down transformer (2-10 Volts), key, horse shoe magnet, wooden stand for mounting the magnet, hanger with weight, set of 50 gm masses, two bridges A and B, screw gauge and meter scale (fitted with the sonometer).

Description of the apparatus: As shown in the Figure 1, a uniform non-magnetic wire is stretched on a hollow wooden box (sonometer). A permanent strong horse shoe magnet is kept at the middle of the bridge such that it produces a magnetic field perpendicular to wire. Two moveable sharp edged bridges A and B are provided on the wooden box for stretching wire. One end of the wire is fixed to the point F on the sonometer. The wire passes over the two bridges A and B and pulley P and is fixed to the hook which supports the hanger carrying weights. The position of the magnet may be fixed at the middle of the wire such that the wire lies between edges of the magnet. A step down transformer (2-10V) is connected across the wire to pass alternating current through the wire.

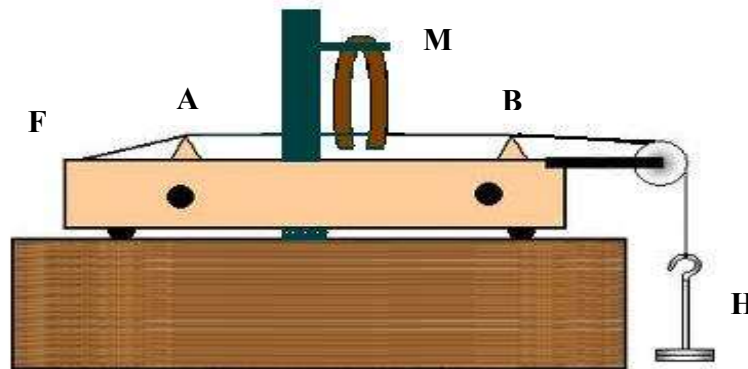


Fig-1: Schematic diagram for sonometer with suggested accessories.

Working Principle: Let the sonometer wire stretched under a constant load be placed in an uniform magnetic field applied at the right angles to the sonometer wire in the horizontal plane and let an alternating current of low voltage (by means of the step down transformer) be passed through the wire. On account of interaction, between the magnetic field and the current in the wire ($F = il \times$

B), the wire will be deflected. The direction of deflection is being given by the Fleming's left hand rule. As the current is alternating, for half the cycle the wire will move upwards and for the next half the wire will move downwards. Therefore, the sonometer wire will receive impulses alternately in opposite directions at the frequency of the alternating current passing through the wire. As a consequence, the wire will execute forced vibrations with a frequency of the AC mains (under the conditions of resonance) in the sonometer wire.

The frequency of AC Mains, which is equal to the frequency of vibration (f) of the sonometer wire in its fundamental mode (only one loop between the two bridges A and B, i.e., having two nodes and one antinode between the two bridges) is given by (under resonance conditions):

$$f = \frac{1}{2l} \sqrt{\frac{T}{m}} \quad (1)$$

where T is the tension applied on the wire and given by $T = Mg$, M being the total mass loaded on the wire (i.e., total mass kept on the hanger and the mass of the hanger), l presents the length of the sonometer wire between the two bridges and g is the acceleration due to gravity. The mass per unit length of the sonometer wire is represented by symbol m and can be calculated in terms of the radius r of the sonometer wire, and the density d of the material wire (nichrome) as

$$m = \pi r^2 d \quad (2)$$

Substitution of value of m , evaluated from the Eq. 2, in Eq. 1, gives the value of frequency of AC mains.

Procedure:

1. Measure the diameter of the wire with a screw gauge at several points along its length. At each point two mutually perpendicular diameters should be measured. Evaluate the radius of the sonometer wire. [See observation Table 1]
2. Connect the step down transformer to AC mains and connect the transformer output (6 V connection) to the two ends of the sonometer wire through a rheostat, ammeter and a key.
3. Place the two movable sharp-edged bridges A and B at the two extremities of the wooden box.
4. Mount the horse shoe magnet vertically at the middle of the sonometer wire such that the wire passes freely in between the poles of the magnet and the face of the magnet is normal

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to the length of the wire. The direction of current flowing through the wire will now be normal to the magnetic field.

5. Apply a suitable tension to the wire, say by putting 100 gm masses on the hanger [tension in the wire = (mass of the hanger + mass kept on the hanger) $\times g$]. Switch on the mains supply and close the key K and then adjust the two bridges A and B till the wire vibrates with the maximum amplitude (in the fundamental mode of resonance) between the two bridges.
6. Measure the distance between the two bridges (l). [See observation Table 2]
7. Increasing the load M in steps of 50 gm, note down the corresponding values of l for maximum amplitude (in the fundamental mode of resonance). Take six or seven such observations.
8. Knowing all the parameters, using the relations given in Eqs. 1 and 2 calculate the frequency of AC mains for each set of observation separately and take average.
9. Also plot a graph between the mass loaded, M along the X-axis and the square of the length (l^2) along Y-axis. This graph should be a straight line. Find the slope of this line and then using the Eqs. 1 and 2, calculate the frequency of AC mains from this graph also. (Frequency (n) = $\sqrt{g/(4 \times \text{slope} \times m)}$).

Observations:

1. **Measurement of radius of sonometer wire (r)**
 - a. Least count of screw gauge = cm
 - b. Zero error of the screw gauge = cm
2. **Measurement of T, l and frequency of the AC Mains**
 - a. Mass of the hanger = 50 gm
 - b. Acceleration due to gravity (g) = 980 cm/sec².
 - c. Density of sonometer wire (nichrome) = 8.18848 gm/cc

Table 1: Measurement of radius of sonometer wire (r)

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S.No	Diameter of wire along one direction (cm)	Diameter of wire in (cm)	Mean observed diameter (cm)	mean corrected diameter (cm)	mean radius r (cm)
1.					
2.					
3.					
4.					
5.					

Table 2: Measurement of T , l and frequency of the AC Mains

S. No	Total Mass Loaded = Mass of hanger + Mass on it M (gm)	Tension in wire $T = Mg$ (gm cm/s ²)	Position of first bridge a (cm)	Position of second bridge b (cm)	Length of wire between two bridges $l=a-b$ (cm)	Frequency (Hz)
1						
2						
3						
4						

1. Mean value of the AC Mains frequency =Hz.
2. Calculations from the graph are also to be given on the left side of paper.
3. The slope of graph plotted between Mass loaded (M) and the square of length $S =$ units
4. AC mains frequency when calculated from the graph = $\sqrt{g/(4 \times \text{slope} \times m)}$ Hz

Results: The frequency of AC Mains as calculated

- a. Experimental calculations: Hz.
- b. Graphical calculations:Hz (Graph to be attached)
- c. Standard Value: 50 Hz (in this country)

d. Percentage Error: %

Sources of errors and precautions

1. The sonometer wire should be uniform and without kinks.
2. The pulley should be frictionless
3. The wire should be horizontal and pass freely in between the poles of magnet.
4. The horse shoe magnet should be placed vertically at the center of the wire with its face normal to the length of wire.
5. The current should not exceed one Ampere to avoid the overheating of the wire.
6. The movement of bridges on the wire should be slow so that the resonance point can be found easily
7. The diameter of the wire must be measured accurately at different points in two mutually perpendicular directions.
8. The sonometer wire and the clamp used to hold the magnet should be non-magnetic.

SAMPLE ORAL QUESTIONS

1. What is the scientific and technological importance of the present experiment?
2. What are the practical applications based on the theory and outcome of this experiment?
3. What understanding you build with this experiment?
4. What do you understand by the frequency of AC Mains?
5. Distinguish between AC and DC. What is the use of magnet here?
6. How does the sonometer wire vibrate when AC is passed through it?
7. If you pass a DC through the wire, will it vibrate?
8. What are the chief sources of errors in this experiment?
9. What is the use of magnet here?
10. What is Fleming's left hand rule?
11. What is resonance?
12. What is fundamental mode of vibration?
13. Why do we take the material of wire to be non-magnetic?
14. What is the principle of this experiment?

EXPERIMENT NO. 2

Aim: To determine the wavelength of sodium lines by Newton's rings method.

Apparatus Required: Optical arrangement for Newton's rings, an optically plane glass plate, convex lens, sodium light source, traveling microscope, magnifying lens, reading lamp and spherometer

Description of Apparatus:

The experimental apparatus for obtaining the Newton's rings is shown in the Fig. 1. A Plano-convex lens L of large radius of curvature is placed with its convex surface in contact with a plane glass plate P. At a suitable height over this combination, is mounted a plane glass plate G inclined at an angle of 45 degrees with the vertical. This arrangement is contained in a wooden box. Light from a board monochromatic sodium source rendered parallel with the help of convex lens L_1 is allowed to fall over the plate G, which partially reflects the light in the downward direction. The reflected light falls normally on the air film enclosed between the Plano-convex lens L, and the glass plate P. The light reflected from the upper and lower surfaces of the air film produce interference fringes. At the center the lens is in contact with the glass plate and the thickness of the air film is zero. The center will be dark as a phase change of π radians is introduced due to reflection at the lower surface of the air film (as the refractive index of glass plate P ($\mu = 1.5$)) is higher than that of the air film ($\mu = 1$). So this is a case of reflection by the denser medium. As we proceed outwards from the center the thickness of the air film gradually increases being the same all along the circle with center at the point of contact. Hence the fringes produced are concentric, and are localized in the air film (figure 2). The fringes may be viewed by means of a low power microscope (travelling microscope) shown in the Fig. 1.

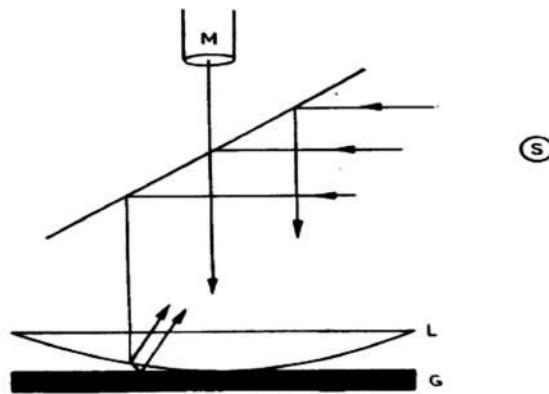


Fig 1

Fig. 1: Schematic representation of the experimental set up

Working Principle:

When a Plano-convex lens of large radius of curvature is placed with its convex surface in contact with a plane glass plate P a thin wedge shaped film of air is enclosed between the two. The thickness of the film at the point of contact is zero and gradually increases as we proceed away from the point of contact towards the periphery of the lens. The air film thus possesses a radial symmetry about the point of contact. The curves of equal thickness of the film will, therefore, be concentric circles with point of contact as the center (Fig.2).

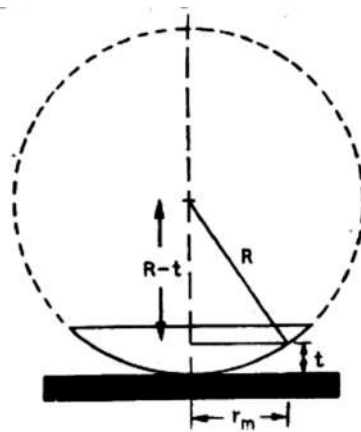


Fig. 2

Fig. 2: Schematic showing the radius of the m th dark ring for an air film thickness, t

In Fig. 3 the rays BC and DE are the two interfering rays corresponding to an incident ray AB. As Newton's rings are observed in the reflected light, the effective path difference x between the two interfering rays is given by:

$$x = 2\mu t \cos(t + \theta) + \lambda/2 \quad (10.1)$$

Where t is the thickness of the air film at B and θ is the angle of film at that point. Since the radius of curvature of the Plano-convex lens is very large, the angle θ is extremely small and can be neglected. The term $\lambda/2$ corresponds to phase change of π radians introduced in the ray DE due to reflection at the denser medium (glass). For air the refractive index (μ) is unity and for normal incidence, angle of refraction is zero. So the path difference x becomes:

$$x = 2t + \lambda/2 \quad (10.2)$$

At the point of contact, the thickness of the film is zero, i.e., $t = 0$, So $x = \lambda/2$. And this is the condition for the minimum intensity. Hence the center of the Newton's rings is dark. Further, the two interfering rays BC and DE interfere constructively when the path difference between the two is given by

$$x = 2t + \frac{\lambda}{2} = 2n \frac{\lambda}{2} \quad (10.3)$$

$$2t = (2n - 1)\lambda/2 \text{ [Maxima]} \quad (10.4)$$

And they interfere destructively when the path difference

$$x = 2t + \frac{\lambda}{2} = (2n + 1)\frac{\lambda}{2} \text{ or } 2t = \frac{2n\lambda}{2} \text{ [Minima]} \quad (10.5)$$

From these equations it is clear that a maxima or minima of particular order n will occur for a given value of t . Since the thickness of the air film is constant for all points lying on a circle concentric with the point of contact, the interference fringes are concentric circles. These are also known as fringes of equal thickness

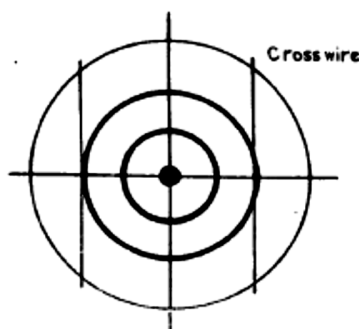


Fig. 3: Schematic representation of Newton's rings and the measurement of its diameter with cross wire.

Experimental Methods:

Calculation of diameters rings:

Let r_n be the radius of Newton's ring corresponding to a point B, where the thickness of the film is t , let R be the radius of curvature of the surface of the lens in contact with the glass plate P, then from the triangle CMB (figure 4), we have:

$$R^2 = r_n^2 + (R - t)^2 \text{ or } r_n^2 = 2Rt - t^2 \quad (10.6)$$

Since t is small as compared to R , we can neglect t^2 and therefore

$$R^2 = 2Rt, \text{ or } 2t = r_n^2/R \quad (10.7)$$

If the point B lies over the n^{th} dark ring then substituting the value of $2t$ from equation (4) we have,

$$\left[\frac{r_n^2}{R} = \frac{2n\lambda}{2} \right], \text{ or } r_n^2 = n\lambda R \quad (10.8)$$

If D_n is the diameter of the ring then,

$$D_n^2 = 4nR\lambda \quad (10.9)$$

Similarly, if the point B lies over a n^{th} order bright ring we have

$$D_n^2 = 2(2n - 1)\lambda R \quad (10.10)$$

Calculation of λ :

From equation (7), if D_{n+1} is the diameter of $(n + p)^{th}$ bright ring, we have

$$D_{n+p}^2 = 2[2(n + 2) - 1]\lambda R \quad (10.11)$$

Subtracting equation (7), from equation (8), we get:

$$D_{n+p}^2 - D_n^2 = 4p\lambda R \quad (10.12)$$

$$\lambda = \frac{D_{n+p}^2 - D_n^2}{4pR} \quad (10.13)$$

By measuring the diameters of the various bright rings and the radius of curvature of the plano convex lens, we can calculate λ from the equation 9.

Formula used

The wavelength λ of the sodium light employed for Newton's rings experiments is given by:

$$\lambda = \frac{D_{n+p}^2 - D_n^2}{4pR}$$

Where D_{n+p} and D_n are the diameter of $(n + p)^{th}$ and n^{th} bright rings respectively, p being an integer number. R is the radius of curvature of the convex surface of the Plano-convex lens.

Methodology

1. Level the travelling microscope table and set the microscope tube in a vertical position. Find the vernier constant (least count) of the horizontal scale of the traveling microscope.
2. Clean the surface of the glass plate p, the lens L and the glass plate G. Place them in position as shown in Figure 1 and as discussed in the description of apparatus. Place the arrangement in front of a sodium lamp so that the height of the center of the glass plate G is the same as that of the center of the sodium lamp. Place the sodium lamp in a wooden box having a hole such that the light coming out from the hole in the wooden box may fall on the Newton's

rings apparatus and adjust the lens L_1 in between of the hole in wooden box and Newton's rings apparatus and adjust the lens L_1 position such that a parallel beam of monochromatic sodium lamp light is made to fall on the glass plate G at an angle of degrees.

- Adjust the position of the travelling microscope so that it lies vertically above the center of lens L . Focus the microscope, so that alternate dark and bright rings are clearly visible.
- Adjust the position of the travelling microscope till the point of intersection of the cross wires (attached in the microscope eyepiece) coincides with the center of the ring system and one of the cross-wires is perpendicular to the horizontal scale of microscope.
- Slide the microscope to the left till the cross-wire lies tangentially at the center of the 20th dark ring (see Figure). Note the reading on the vernier scale of the microscope. Slide the microscope backward with the help of the slow motion screw and note the readings when the cross-wire lies tangentially at the center of the 18th, 16th, 14th, 12th, 10th, 8th, 6th and 4th dark rings respectively [Observations of first few rings from the center are generally not taken because it is difficult to adjust the cross-wire in the middle of these rings owing their large width].
- Keep on sliding the microscope to the right and note the reading when the cross-wire again lies tangentially at the center of the 4th, 6th, 8th, 10th, 12th, 14th, 16th, 18th, 20th dark rings respectively.
- Remove the plano-convex lens L and find the radius of curvature of the surface of the lens in contact with the glass plate P accurately using a spherometer. The formula to be used is :

$$R = \frac{l^2}{6h} + \frac{h}{2} \quad (10.14)$$

Where l is the mean distance between the two legs of the spherometer h is the maximum height of the convex surface of the lens from the plane surface.

- Find the diameter of each ring from the difference of the observations taken on the left and right side of its center. Plot a graph between the number of the rings on X-axis and the square of the corresponding ring diameter on Y-axis. It should be a straight line as given by the equation 9 (see figure). Taken any two points on this line and find the corresponding values of $(D_{n+2}^2 - D_n^2)$ and p for them.

9. Finally calculate the value of wavelength of the sodium light source using the formula.

Observations

Determination of the Least Count:

Determination of the Least Count of the Horizontal Scale of travelling Microscope

1. Value of one division of the horizontal main scale of travelling microscope = cm
2. Total number of divisions on the Vernier scale = which are equal to division of main scale of the Vernier scale =cm
3. Value of one division of the Vernier scale =cm
4. Least count of the horizontal scale of the microscope (given by the value of one division of main scale – the value of one division of Vernier scale) =
5. Pitch of the screw =cm
6. Number of division on circular head =
7. Least count of the spherometer =cm
8. Mean distance between the two legs of the spherometer, l =cm
9. The radius of curvature R of the Plano convex lens is (as given by equation 10): $R = [l^2/(6h + h)/2] = \dots\dots\dots$ cm
10. The wavelength λ of sodium light is (as given by equation 9):

$$\lambda = \frac{D_{n+2}^2 - D_n^2}{4pR}$$

Calculations from the graph:

1. Plot a graph taking squares of the diameters, D_n^2 along the Y-axis and the number of rings along the X-axis.
2. The curve should be a straight line.
3. Take two points P_1 and P_2 on this line and find the corresponding values of $(D_{n+2}^2 - D_n^2)$ and p from it, calculate the value of wavelength of the sodium light from these values.

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Table 1: Determination of $(D_{n+2}^2 - D_n^2)$ and p

Order of The Rings	Reading of the microscope left hand side (a) cm	Right hand side (b) cm	Diameter of the ring ($a \sim b$) cm	Diameter ($a \sim b$) ² cm ²	$(D_{n+2}^2 - D_n^2)$ For p=4 cm ²	Mean value of $(D_{n+2}^2 - D_n^2)$ For p=4 cm ²
20					$D_{20}^2 - D_{16}^2$ =..	
18					$D_{18}^2 - D_{14}^2$ =..	
16					$D_{16}^2 - D_{12}^2$ =..	
14					$D_{14}^2 - D_{10}^2$ =..	
12					$D_{12}^2 - D_8^2$ =..	
10					$D_{10}^2 - D_6^2$ =..	
8					$D_8^2 - D_4^2$ =..	

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Table 2: Determination of R (radius of curvature of the lens L) using a spherometer

Sl. no.	Spherometer reading		h (a~b) cm	Mean h (cm)
	Plane glass plate <i>a</i> (cm)	Convex surface of lens <i>b</i> (cm)		

Results:

The value of the wavelength of the sodium light as calculated

1. Using the observations directly =A°
2. Using the graphical calculations = A°
3. Mean value of the wavelength of sodium light = A°
4. Standard average value of the wavelength of the sodium light = 5893 A°
5. Percentage of error = %

Source of errors and precautions:

- The optical arrangements as shown in Figure 1 should be very clean (use sprit for cleaning these optical elements) and so made that the beam of light falls normally on the plano-convex lens L and glass plate P combinations.
- The plano-convex lens for the production of Newton’s rings should have large value of radius of curvature. This will keep the angle of wedge shape air film very small and therefore the rings will have a larger diameter and consequently the accuracy in the measurements of the diameter of the rings will be increased.
- To avoid any backlash error, the micrometer screw of the travelling microscope should be moved very slowly and moved in one direction while taking observations.
- While measuring diameters, the microscope cross-wire should be adjusted in the middle of the ring.

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- The amount of light from the sodium light should be adjusted for maximum visibility. Too much light increases the general illumination and decrease the contrast between bright and dark rings.

Sample oral questions:

1. What is the scientific and technological importance of the present experiment?
2. What are the practical applications based on the theory and outcome of this experiment?
3. What understanding you build with this experiment?
4. What do you understand by the interference of light?
5. What are essential conditions for obtaining interference of light?
6. What do you understand by coherent source?
7. Is it possible to observe interference pattern by having two independent sources such as two candles?
8. Why should be two sources be monochromatic?
9. Why are the Newton's rings circular?
10. Why is central ring dark?
11. Where are these rings formed?
12. Sometimes these rings are elliptical or distorted, why?
13. What is the difference between the rings observed by reflected light and those observed by transmitted light?
14. What will happen if the glass plate is silvered on the front surface?
15. What will happen when a little water is introduced in between the plano-convex lens and the plate?
16. How does the diameter of rings change on the introduction of liquid?
17. Can you find out the refractive index of a liquid by this experiment?
18. Is it possible to have interference with a lens of small focal length?
19. What will happen if the lens is cylindrical?
20. Why do the rings get closer and finer as we move away from the center?

EXPERIMENT NO. 3

Aim of the Experiment: To determine (a) the resistance per unit length of the wire of a Carey-Foster bridge and then to (b) find the resistivity of the material of a given wire.

Apparatus Required: Carey- Foster bridge, Leclanche cell, Weston galvanometer, standard one – ohm coil, 2 low resistance boxes, plug key, thick copper strip, connecting wires.

Theory of Carey-Foster bridge: The Carey- Foster bridge is a modified form of the meter bridge (shown in Fig. 1). The Carey- Foster bridge is especially suited for the comparison of two nearly equal resistances whose difference is less than the resistance of the bridge wire.

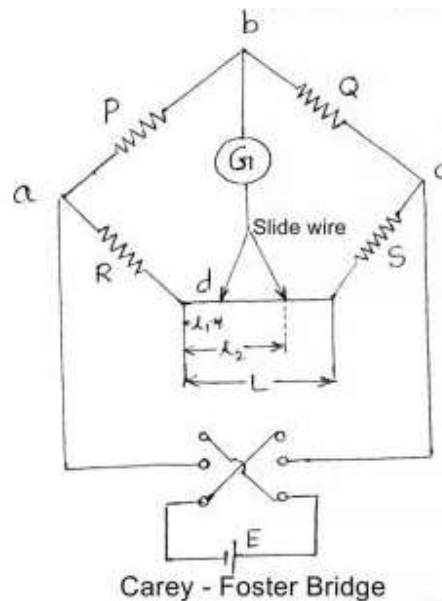


Fig. 2: Circuit diagram of Carey Foster’s bridge

As shown in the Fig. 2, two nearly equal resistances P and Q are connected in the inner gaps of the bridge, and two low resistances X and Y which are to be compared are connected in the outer gaps of the bridge. If l_1 be the position of the null point, we have from the Wheatstone bridge principle:

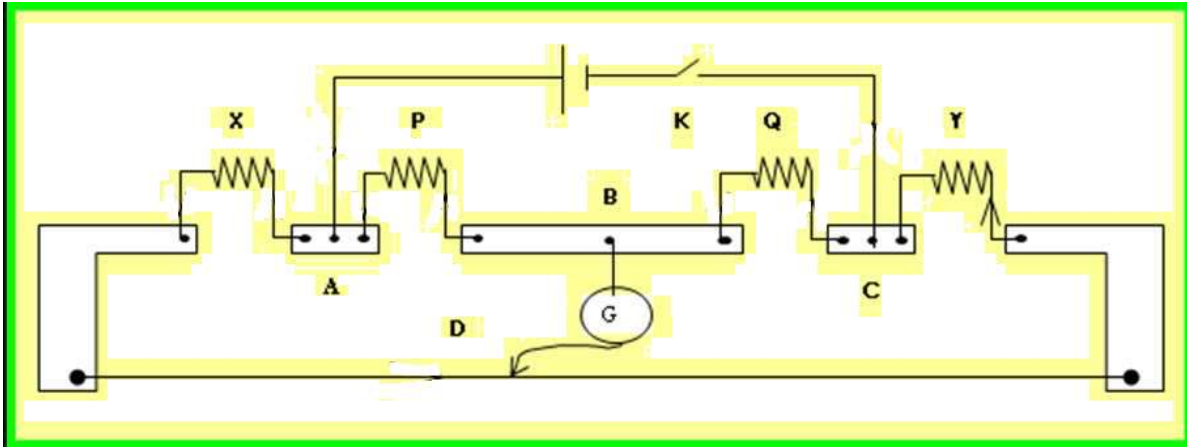


Fig. 1: Schematic representation of Carey-Foster bridge experimental set-up

$$\frac{P}{Q} = \frac{X + \sigma(l_1 + \alpha)}{Y + \sigma(100 - l_1 + \beta)} \quad (1)$$

or

$$\frac{P}{Q} + 1 = \frac{X + Y + \sigma(100 + \alpha + \beta)}{Y + \sigma(100 - l_1 + \beta)} \quad (2)$$

where, σ is the resistance per unit length of the bridge wire, α and β are the end corrections (in cm) at the left and right ends of the bridge wire respectively.

If the resistances X and Y are interchanged and l_2 be the position of the new null point, we have

$$\frac{P}{Q} = \frac{Y + \sigma(l_2 + \alpha)}{X + \sigma(100 - l_2 + \beta)} \quad (3)$$

$$\frac{P}{Q} + 1 = \frac{X + Y + \sigma(100 + \alpha + \beta)}{X + \sigma(100 - l_2 + \beta)} \quad (4)$$

Comparing equations (2) and (4) we see that the fraction on the right hand side are equal and since their numerators are identical their denominators must also be equal. Hence equating the denominators of the right hand sides of equation (2) and (4), we have

$$Y + \sigma(100 - l_1 + \beta) = X + \sigma(100 - l_2 + \beta) \quad (5)$$

$$X - Y = \sigma(l_2 - l_1) \quad (6)$$

[In the above procedure the end-corrections are eliminated from the working equations]

Part A: To find σ , the resistance per unit length of the bridge wire:

Equation (6) can be used to find σ , the resistance per unit length of the bridge wire, if the values of X and Y are known. Let X = a thick copper strip be connected in the outer left gap and Y = the standard 1 Ω be connected in the outer right gap of the bridge, and let l_1 and l_2 respectively be the null points on the bridge wire, before and after interchanging X and Y in the outer gaps. Then we have from Eq. (6) by putting X = 0 and Y = 1 Ω ,

$$\sigma = 1 / (l_1 - l_2) \quad (7)$$

Part B: To find the resistivity ρ of the material of a given wire

Now, replace the copper strip with the wire of unknown resistance = X connected initially in the outer left gap and the standard 1 Ω resistance = Y in the outer right gap of the bridge. Then if l'_1 and l'_2 respectively be the readings on the scale of the positions of the null point before and after interchanging X and Y, we have, from equation (6):

$$X - Y = \sigma(l'_2 - l'_1) \quad (8)$$

$$X = \sigma(l'_2 - l'_1) + Y \quad (9)$$

This equation can be used to calculate X, if σ is determined from equation (7).

Then if L is the length and 'a' the cross-sectional area of the wire, the resistivity ρ is related to X by

$$X = \rho \frac{L}{a} \quad (10)$$

or

$$\rho = \frac{Xa}{L} \quad \text{Ohm-cm.} \quad (11)$$

Procedure for Part A: To find σ , the resistance per unit length of the bridge wire:

Connect a thick copper strip (X = 0) in the outer left gap and a standard 1-ohm resistance (Y = 1 Ω) in the outer right gap of a Carey- Foster bridge as shown in Fig. 1. Connect the two low resistance

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boxes P & Q in the inner gaps as shown. A jockey is connected through the galvanometer as shown. Finally connect the Lechlanche cell and include a plug key K in the circuit. Select nearly equal values of P and Q in the resistance boxes and find the null point l_1 for each of about 10 sets of P and Q. (Note that P and Q need not be exactly equal. Choose values for P, Q such as 3, 3; 3, 4; 4, 3; 4, 4; 4, 5; 5, 4; 5, 5; etc.).

Now connect the thick copper strip (X) in the outer right gap and the standard 1-ohm resistance (Y) in the outer left gap of the bridge and find l_2 for each of these sets of P and Q. Follow the observation table. Calculate the value of σ for each set of observations separately from equation (7) and then find the mean value of σ .

Observations:

Resistance per unit length:

$$\sigma = \frac{1}{(l_1 - l_2)} \Omega/cm$$

Table 1: Determination of σ , resistance per unit length of the bridge wire

S.No.	P (ohm)	Q (ohm)	Position of balance point with X in the		$(l_1 - l_2)$ (cm)	σ (ohm/cm)	Mean σ (ohm/cm)
			Left gap l_1 (cm)	Right gap l_2 (cm)			

Results:

Mean value of $\sigma = \dots \dots \dots$ ohm/cm

Procedure for Part A: To find the resistivity ρ of the material of a given wire:

To find the resistance of the given wire and hence its resistivity, replace the copper strip by the given wire and repeat the above procedure. Find the resistance of the given wire using Eq (6).

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Measure the length of the wire and find its diameter using a micrometer screw gauge. Calculate the resistivity using equation (11).

Length of the wire $L = \dots\dots\dots\text{cm}$

Diameter at three different points on the wire, along perpendicular diameters at each point

Table 2: Determination of resistance and resistivity of the unknown wire (usually Nichrome)

S.No.	P (ohm)	Q (ohm)	Position of balance point with X in the		$(l'_2 - l'_1)$ (cm)	σ (ohm/cm)	Mean σ (ohm/cm)
			Left gap l'_1 (cm)	Right gap l'_2 (cm)			

Results and discussion:

Mean value of the resistance of the given wire is found to be ...

The resistivity ρ of the material of the given wire is

Resistivity of nichrome at room temperature $\sim 100 - 150 \times 10^8 \Omega \cdot \text{m}$

Percentage of error:

Sources of error and precautions:

1. The ends of the connecting wires should be clean and all connections should be tight.
2. The resistances in the four arms of the bridge should be nearly equal, so that the bridge may have high sensitivity.
3. A plug key should be included in the cell circuit and should be closed only when observations are being made.

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4. The galvanometer may be shunted by a low resistance wire to avoid excessive deflection in it when the bridge is out of balance. The exact position of the null point should be determined with full galvanometer sensitivity by removing the shunt wire from it.
5. The jockey should always be pressed gently and the contact between the jockey and the bridge wire should not be made while the jockey is being moved along.

Nichrome is a non-magnetic alloy of nickel, chromium, and often iron, usually used as a resistance wire. Patented in 1905, it is the oldest documented form of resistance heating alloy. A common alloy is 80% nickel and 20% chromium, by mass, but there are many others to accommodate various applications. It is silvery-grey in colour, is corrosion-resistant, and has a high melting point of about 1,400 °C (2,550 °F). Due to its resistance to oxidation and stability at high temperatures, it is widely used in electric heating elements.

Resistivity of nichrome at room temperature $\sim 1.0 - 1.5 \times 10^6 \Omega \cdot m$

(From Wiki)

SAMPLE ORAL QUESTIONS

1. What is the scientific and technological importance of the present experiment?
2. What are the practical applications based on the theory and outcome of this experiment?
3. What understanding you build with this experiment?
4. What is the principle of Carey Foster's bridge?
5. What is a Wheatstone bridge?
6. Why we use Carey-Foster's bridge instead of a meter bridge?
7. When is the Carey-Foster's bridge most sensitive?
8. What is specific resistance?
9. Explain some of the advantages and limitations of a Carey Foster's bridge for measuring resistance.
10. Which other instruments based on the principle of Wheatstone's bridge are used to determine resistances?

EXPERIMENT NO. 4

Aim of the Experiment: Measurement of Mechanical equivalent of heat (Joule's equivalent) by electrical method

1. To understand the phenomena of mechanical equivalent of heat;
2. To measure the mechanical equivalent of heat of water; and
3. To measure the efficiency of a given incandescent lamp.

Equipment required: Regulated DC power supply, digital volt-ammeter, thermometer, stop watch, incandescent lamp, electrical equivalent jar, calorimeter, Indian ink.

Part 1: The Mechanical Equivalent of Heat

Procedure

1. Measure and record the room temperature ()
2. Weigh the EEH jar (with the lid on), and record its mass ().
3. Remove the lid of EEH jar and fill the jar to the indicated water line with cold water. DO NOT OVERFILL. The water should be approximately 10°C below room temperature, but exact temperature is not critical.
4. Add about 10 drops of Indian ink to the water, enough so the filament is just barely visible when the lamp is illuminated.
5. Using lead with banana plug connectors, attach your power supply to the terminals of the EEH Jar. Connect a voltmeter and ammeter as shown in Figure 1.1 so you can measure both the current (I) and voltage (V) going into the lamp.

NOTE-For best results, connect the voltmeter leads directly to the binding posts of the jar.

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6. Turn on the power supply quickly adjust the power supply voltage to about 11.5 volts, then shut the power off. **DO NOT LET THE VOLTAGE EXCEED 13 VOLTS.**
7. Insert the EEH into one of the Styrofoam Calorimeters.
8. Inset your thermometer or thermistor probe through the hole in top of the EEH jar. Stir the water gently with the thermometer or probe while observing the temperature. When the temperature warms to about 6 or 8 degrees below room temperature, turn the power supply on.
9. NOTE: You may want to turn the lamp on to help the cold water reach this starting temperature. If you do, be sure that you turn the lamp off for several minutes before begin your measurements, so you are sure the water temperature is even throughout the jar. Record the starting time () and the temperature ().
10. Record the current, I , and voltage, V . Keep an eye on the ammeter and voltmeter throughout the experiment to be sure these values do not shift significantly. If they do shift, use an average value for V and I in your calculations.
11. When the temperature is as far above room temperatures as it was below room temperature (=Temperature-), shut off the power and record the time (). Continue stirring the water gently. Watch the thermometer or probe until the temperature peaks and start to drop. Record this peak temperature (T_f).
12. Weigh the jar with the water, and record the value ().

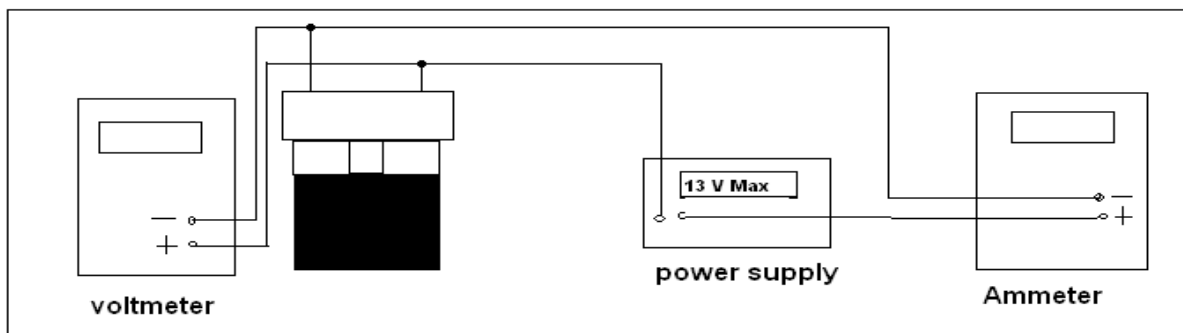


Figure 4.1 Schematic of mechanical connection to bulb, multimeter, power supply.

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Data

Room temperature =

Mass of jar =

Mass of jar with water =

Voltage V =

Current I =

Starting time =

Final time =

Starting temperature =

Final temperature =

Calculation

In order to determine the mechanical equivalent of heat (J), it is necessary to determine both the total electrical energy that flowed into the lamp (E) and the total heat absorbed by the water (H).

E , the electrical energy delivered to the lamp

E = Electrical energy into the Lamp =

= time during which power was applied to the lamp =

H , the heat transferred to the water (and the EEH jar):

.....

= Mass of water heated =

= 23 grams. Some of the heat produced by the lamp is absorbed by EEH jar. For accurate results, therefore, the heat capacity of the jar must be taken into account (The heat capacity of the EEH jar

is equivalent to that of approximately 23 grams of water.)

J_m , the Mechanical Equivalent of Heat

=

Part 2: Efficiency of an Incandescent Lamp

Procedure

Repeat Experiment 1, except do not use the Indian Ink (step 4) or the Styrofoam Calorimeter (step 7). Record the same data as in Experiment 1, and use the same calculations to determine E and H. (Convert H to joule by multiplying by J_m from the first lab.)

In performing the experiment with clear water and no Calorimeter, energy in the form of visible light is allowed to escape the system. However, water is a good absorber of infrared radiation, so most of the energy that is not emitted as visible light will contribute to H, the thermal energy absorbed by the water. The efficiency of the lamp is defined as the energy converted to visible light divided by the total electrical energy that goes into the lamp. By making the assumption that all the energy that doesn't contribute to H is released as visible light, the equation for the efficiency of the lamp becomes:

Efficiency

Data

Room temperature =

Mass of jar =

Mass of jar with water =

Voltage V =

Current I =

Starting time t_i =

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Final time t_f =

Starting temperature =

Final temperature =

Calculation

In order to determine the efficiency of the lamp, it is necessary to determine both the total electrical energy and that flow into the lamp and the total heat absorbed by the water (H).

E, the electrical energy delivered to the lamp

E = electrical energy into the lamp =

= the time during which power was applied to the lamp =

H, the heat transferred to the water (and calorimeter)

= Mass of water heated =

=

= 23 grams. Some of the heat produced by the lamp is absorbed by the EEH jar. For accurate results, therefore, the heat capacity of the jar must be taken into account (The heat capacity of the EEH jar is equivalent to that of approximately 23 grams of water.)

Efficiency

=

Sample Questions

1. What is the scientific and technological importance of the present experiment?
2. What are the practical applications based on the theory and outcome of this experiment?
3. What understanding you build with this experiment?
4. Define mechanical equivalent of heat.
5. Define heat transfer.
6. What are the various modes of heat transfer?
7. What is conduction?
8. State Fourier's law of conduction.
9. Define convection.
10. State Newton's law of convection.
11. Define radiation.
12. Define emissive power
13. What effect are the following factors likely to have on the accuracy of your determination of the mechanical equivalent of heat? Can you estimate the magnitude of the effect?
 - (a) The inked water is not completely opaque to visible light.
 - (b) There is some transfer of thermal energy between the EEH jar and the room atmosphere.
(What is the advantage of beginning the experiment below room temperature and ending it an equal amount above room temperature?).
14. Is an incandescent lamp more efficient as a light bulb or as a heater?

EXPERIMENT NO. 5

Aim of the experiment: Determination of refractive index of the material of a prism using spectrometer and sodium light.

Apparatus required: source of light (sodium vapor lamp or mercury lamp), spectrometer, prism, spirit level.

Basic description of a Spectrometer: In some of the optics experiments, spectrometers are used. The spectrometer is an instrument for studying the optical spectra. Light coming from a source is usually dispersed into its various constituent wavelengths by a dispersive element (prism or grating) and then the resulting spectrum is studied. A schematic diagram of prism spectrometer is shown in Fig.1. It consists of a collimator, a telescope, a circular prism table and a graduated circular scale along with two verniers. The collimator holds an aperture at one end that limits the light coming from the source to a narrow rectangular slit. A lens at the other end focuses the image of the slit onto the face of the prism. The telescope magnifies the light dispersed by the prism (the dispersive element for your experiments) and focuses it onto the eyepiece. The angle between the collimator and telescope are read off by the circular scale. The detail description of each part of the spectrometer is given below.

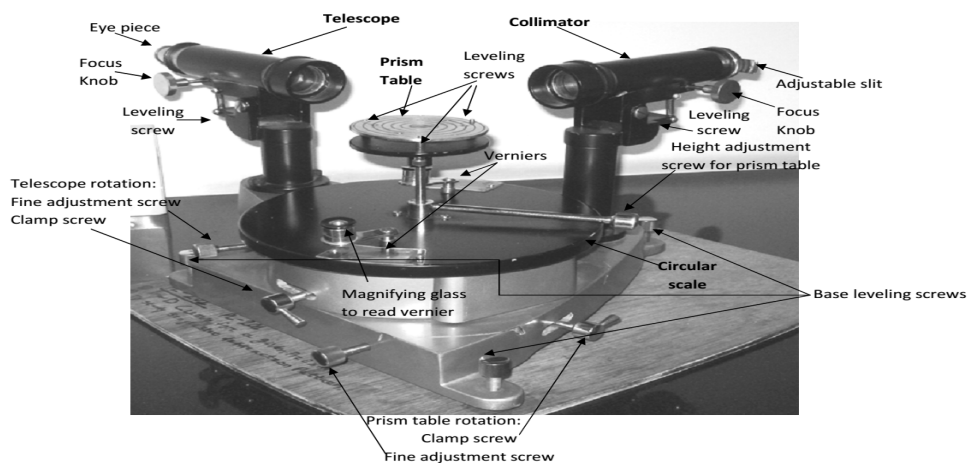


Fig. 1: Different parts of spectrometer

- a) Collimator (C):** It consists of a horizontal tube with a converging achromatic lens at one end of the tube and a vertical slit of adjustable width at the other end. The slit can be moved in or out of the tube by a rack and pinion arrangement using the focus knob and its width can be adjusted by turning the screw attached to it. The collimator is rigidly fixed to the main part of the instrument and can be made exactly horizontal by adjusting the leveling screw provided below it. When properly focused, the slit lies in the focal plane of the lens. Thus the collimator provides a parallel beam of light.
- b) Prism table (P):** It is a small circular table and capable of rotation about a vertical axis. It is provided with three leveling screws. On the surface of the prism table, a set of parallel equidistant lines parallel to the line joining two of the leveling screws, is ruled. Also, a series of concentric circles with the centre of the table as their common centre is ruled on the surface. A screw attached to the axis of the prism table fixes it with the two verniers and also keep it at a desired height. These two verniers rotate with the table over a circular scale graduated in fraction of a degree. The angle of rotation of the prism table can be recorded by these two verniers. A clamp and a fine adjustment screw are provided for the rotation of the prism table. It should be noted that a fine adjustment screw functions only after the corresponding fixing screw is tightened.
- c) Telescope (T):** It is a small astronomical telescope with an achromatic doublet as the objective and the Ramsden type eye-piece. The eye-piece is fitted with cross-wires and slides in a tube which carries the cross-wires. The tube carrying the cross wires in turn, slides in another tube which carries the objective. The distance between the objective and the cross-wires can be adjusted by a rack and pinion arrangement using the focus knob. The Telescope can be made exactly horizontal by the leveling screws. It can be rotated about the vertical axis of the instrument and may be fixed at a given position by means of the clamp screw and slow motion can be imparted to the telescope by the fine adjustment screw.
- d) Circular Scale (C.S.):** It is graduated in degrees and coaxial with the axis of rotation of the prism table and the telescope. The circular scale is rigidly attached to the telescope and turned

with it. A separated circular plate mounted coaxially with the circular scale carries two verniers, V1 and V2, 180° apart. When the prism table is clamped to the spindle of this circular plate, the prism table and the verniers turn together. The whole instrument is supported on a base provided with three leveling screws. One of these is situated below the collimator.

1. Measurement of angles with the help of spectrometer:

The spectrometer scales are angle measuring utilities for the positions of the telescope which can be rotated about the central axis of the instrument. The main circular scale is attached with the telescope so that when the telescope is rotated, the main circular scale also rotates with it. The angle, through which the telescope is rotated, can be measured by reading the positions of the verniers attached to the prism table and sliding over the main scale. In a spectrometer there are two sets of main circular scales (fitted with the telescope) and vernier scale (attached with the prism table). Both sets are diagonally (left hand and right hand sides) fixed in the instrument and measures angle for a particular telescope position with a difference of 180 degrees. These scales can be used in a similar manner as a simple Vernier Caliper or traveling microscope is used. The vernier Caliper or traveling microscope is used to measure small distances (in centimeters and fractions whereas spectrometer scales are used to measure small angular displacements (in degrees, minutes, and seconds) (1 degree is equal to 60 minutes, and 1 minute is equal to 60 seconds; ($1^\circ = 60'$ and $1' = 60''$))

2. Least Count of the Spectrometer Scale:

BIT, Physics Laboratory has two types of spectrometers in which

- (i) 60 divisions of vernier Scale are equal to 59 divisions of the Main Scale, and
- (ii) 30 divisions of vernier Scale are equal to 29 divisions of the Main scale.

Now, we will find out the least count in first case which 60 divisions of Vernier scale are equal to 59 divisions of the main scale. The method is as follows:

- a) Value of one division of circular main scale = $0.5^\circ = 30'$ (as $1^\circ = 60'$)
- b) Value of one division of sliding vernier scale = $(59/60) \times 0.5^\circ$
- c) Least count of spectrometer scale = Value of 1 div. of main scale – value of 1 div. of vernier = $0.5^\circ - [(59/60) \times 0.5]^\circ$

$$= [0.5 \times 1/60]^\circ = 0.5' = 30'' \text{ (THIRTY SECONDS)}$$

- d) Similarly, the least count of the spectrometer scale in second case in which 30 divisions of Vernier scale are equal to 29 divisions of the circular main scale can also be calculated. In this case the value of least count will be 1' or 60''.

3. Taking Readings on the two Spectrometer scales:

Following is an illustration for taking observation reading using the left hand side set of the circular main scale (attached with the telescope) and the corresponding vernier scale (sliding over the circular main scale and attached with the prism table). Assuming that we are using the spectrometer in which 60 divisions of vernier scale are equal to 59 divisions of the main scale.

The 0th division of the vernier scale precedes the circular main scale division whose value is 234° and 30'. Therefore, the main scale division reading is 234° 30'. Let 13th division of vernier scale coincides completely with a main scale division. Therefore, the vernier scale reading would be = 13x Least count of vernier scale

$$= 13 \times 30'' = 390'' = 6' 30''$$

$$\begin{aligned} \text{Total Spectrometer Scale Reading} &= \text{Circular Main scale Reading} + \text{Vernier Reading} \\ &= 234^\circ 36' 30'' \end{aligned}$$

Reading of the right hand side scale can be similarly observed. The readings taken from left hand side and right hand side should ideally differ by 180°

Theory: A prism is a portion of refracting material bounded by three planes. A cross-section of prism by a horizontal plane is triangular in form. The prisms we use in the lab are equilateral prisms, but other configurations are also possible. The faces of a prism are called refracting faces. In the lab spectrometer experiment one face of the prism is etched or frosted so that light does not pass through it. The line at which two refracting faces meet is called a refracting edge. Since there are only two refracting faces, there is only one refracting edge.

A ray of light EF incident on one of the refracting faces gets refracted along the path FG through the prism and emerges along the path GH as shown in the Fig.2. The angle between the original direction of the incident ray and the emergent ray is called the angle of deviation *D*. For refraction through a prism,

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$$i + e = A + D$$

where i & e are the angle of incidence and angle of emergence respectively, A being the angle of the prism and D is the angle of deviation.

Angle of deviation (D) depends upon the angle of incidence i . For a certain angle of incidence, the deviation will be minimum. It is denoted by D_m . The refractive index μ of the material of the prism is related to the angle of prism A and the angle of minimum deviation D_m through the relation

$$\mu = \frac{\sin\left(\frac{A + D_m}{2}\right)}{\sin\frac{A}{2}}$$

Where μ = refractive index of the material of the prism, A = angle of the prism, D_m = angle of minimum deviation. When the prism is at the minimum deviation position, the angle $i = e$ and the angles of refraction at both surfaces $r_1 = r_2$

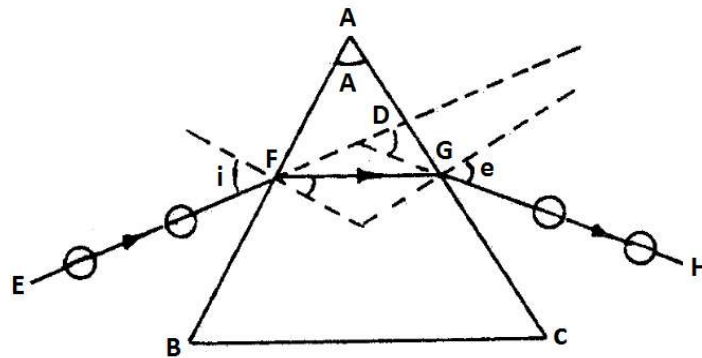


Fig. 2: Ray diagram of a ray of light through a prism

$$\text{From geometry } D = i + e - A$$

$$r_1 + r_2 = A$$

$$D_m = 2i - A$$

or,

$$2i = A + D_m$$

$$i = A + D_m/2$$

$$\& r = A/2.$$

Procedure: Measurement of refractive index of material of the prism consists of two parts *i. e.*

- A. Determination of the angle of prism A (For an equilateral prism we may take $A = 60^\circ$)
- B. Determination of angle of minimum deviation D_m

Measurement of the angle of minimum deviation:

There are two procedures we may use to find D_m – a direct but approximate procedure, and a rigorous graphical procedure. In both cases we may use either monochromatic light from a sodium lamp, or polychromatic light from a mercury lamp. Note that the refractive index μ will differ for the different colours from the mercury lamp.

A) Direct procedure to find D_m for light from a sodium lamp, or from a mercury lamp

Initial procedures to practice and perform before main experiment:

Level the spectrometer base, prism table, telescope and collimator, using procedures explained elsewhere. Adjust the telescope and collimator for parallel rays as described elsewhere. Place the prism so that its center coincides with the center of the prism table (use the circles on the prism table to centre the prism). Adjust the prism so that light from the given source passes through the collimator and falls on one of the refracting faces of the prism and emerges out of the other refracting face after refraction. An image of the slit or the spectrum of light from the source is obtained, and may be viewed either directly by your eye or through the telescope.

Now remove the prism and bring the telescope in the line of the collimator. Observe the slit directly through the telescope and rotate the telescope so that the vertical crosswire coincides with the image of slit. Note the reading of any one of the two verniers – V_1 say – and continue to use only this one vernier scale throughout the experiment. Let the direct ray be D_o .

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Place and centre the prism on the prism table as shown in the Fig. 3. If the telescope is placed as shown, keep the frosted surface to the left and the refracting edge to the right. We start with the procedure for monochromatic light from a sodium vapour lamp. Switch on the source and view the emergent light – image of the slit - with one unaided eye. Now adjust the prism for minimum deviation in the following way. Starting with the refracting face nearly normal to the light, rotate the prism table clockwise so that the refracting edge moves towards you, all the time keeping the image of the slit in view. You will easily find a position where the image or spectral line begins to recede in the opposite direction although you continue to rotate the table in the same direction. Once you find this position, you may rotate the prism both clockwise and anti- clockwise to find the same thing happening. This position where the spectral line begins to recede in opposite direction is the position of minimum deviation. Now while looking at the slit with the unaided eye, rotate the telescope so that it comes between the prism and the eye. Rotate the prism and if needed the telescope to find the position of minimum deviation. Note the reading of the vernier V_1 . Let it be R . The difference between the D_o and R is the minimum deviation D_m . Calculate refractive index from the equation (1) above.

If we use light from a mercury lamp please note that the light is polychromatic, and several coloured lines red, yellow, green, blue, will be seen. Focus on one colour at a time and repeat the above procedure for all colours (wavelengths).

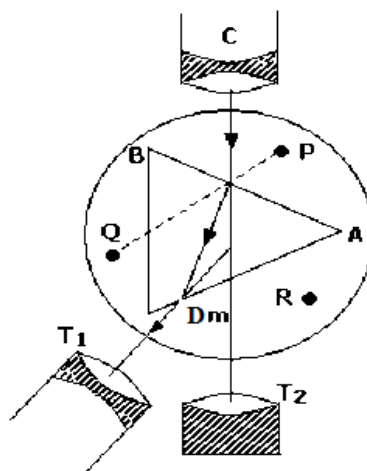


Fig. 3: Ray diagram for determination of angle of minimum deviation

B) Rigorous graphical procedure to find D_m for light from a sodium lamp

Initial procedures to practice and perform before main experiment:

Level the spectrometer base, prism table, telescope and collimator, using procedures explained elsewhere. Adjust the telescope and collimator for parallel rays as described elsewhere. Place the prism so that its center coincides with the center of the prism table (use the circles on the prism table to centre the prism). Adjust the prism so that light from the given source passes through the collimator and falls on one of the refracting faces of the prism and emerges out of the other refracting face after refraction. An image of the slit or the spectrum of light from the source is obtained, and may be viewed either directly by your eye or through the telescope.

Now remove the prism and bring the telescope in the line of the collimator. Observe the slit directly through the telescope and rotate the telescope so that the vertical crosswire coincides with the image of slit. Note the reading of any one of the two verniers – V_1 say – and continue to use only this one vernier scale throughout the experiment. Let the direct ray be D_o .

Place and centre the prism on the prism table. Facing the slit, keep the frosted surface of the prism to the left and the refracting edge to the right. Switch on the monochromatic light source - sodium vapour lamp. Now take the readings for angle of incidence I on the prism and corresponding angle of deviation in the following way. Starting from the direct reading position D of telescope, rotate the telescope anti-clockwise by about 60° and then while looking through the telescope rotate the prism clockwise (so that the refracting edge moves towards you), until the image of the slit comes into view. Make accurate adjustments and take the reading P . Obviously $2I + (D \sim P) = 180$ and I can be calculated. Now without disturbing the prism, rotate the telescope clockwise and find the deviated ray. Take the reading R . Obviously $D_o \sim R = D$. Additional values of I and corresponding D may be taken by rotating the telescope in steps through $60^\circ + 2x$ from direct position (so that I increases by x , where $x = 10^\circ$ say), and finding the corresponding R and D .

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While rotating the prism table note down the different angles from the vernier and read as angle of deviation and calculate the angle of incidence using the equation $i = A + D_m/2$.

1. Plot a graph (i vs D) taking angle of incidence i along the X-axis and angle of deviation D along the Y-axis. The nature of the graph is shown in Fig.4.
2. Draw a horizontal line as a tangent to the lowest point of the curve. Intersection of this horizontal line on Y-axis gives the angle of minimum deviation D_m (Fig.4).

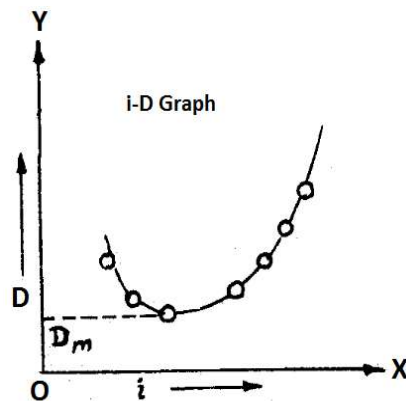


Fig. 4: Graph of angle of deviation vs angle of incidence

Observations:

The angle of given prism is 60°

Table 1: Calculation of the angle of minimum deviation (D_m):

S.No.	Colour	Vernier	Telescope Reading for minimum deviation			Telescope Reading for direct image			Difference $D_m = a-b$	Mean Value of D_m
			MSR	VSR	TR(a)	MSR	VSR	TR(b)		
1.	Violet	V_1								
		V_2								

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2.	Red	V ₁				
		V ₂				
3.	Yellow	V ₁				
		V ₂				

MSR = Main Scale Reading, VSR = Vernier Scale Reading, TR = MSR+VSR = Total Reading

Calculations: Putting the mean value of A from Table 1 and the angle of minimum deviation D_m from the graph, the refractive index of the material of the prism can be found out.

Result: Refractive Index for the material of the prism:

S.No.	Color	Calculated Refractive Index	Standard Refractive Index	%Error

Precautions and Sources of Error:

1. The telescope and collimator should be individually set for parallel rays.
2. Slit should be as narrow as possible.
3. While taking observations, the telescope and prism table should be clamped with the help of clamping screws.
4. Both verniers should be read.
5. The prism should be properly placed on the prism table for the measurement of angle of the prism as well as for the angle of minimum deviation.

Sample questions

1. What is the scientific and technological importance of the present experiment?
2. What are the practical applications based on the theory and outcome of this experiment?
3. What understanding you build with this experiment?
4. What is the least count of spectrometer used?
5. Name the various parts of a spectrometer. What are their use?
6. Define deviation and angle of deviation.
7. How the angle of minimum deviation is ascertained experimentally?
8. How does the angle of deviation vary with the wave length?
9. What is the relation between the angle of incidence and the angle of deviation?
10. How angle of prism is related with the angle of deviation?
11. When light enters into the prism is there any change in the frequency of wave length?
12. What is unit of refractive index?
13. Which of the colours have maximum and minimum angle of deviations?
14. How does the refractive index changes with wavelength?
15. What do you mean by angular dispersion of a prism?

EXPERIMENT NO . 6

Aim of the experiment: To determine the frequency of electrically maintained tuning fork by Melde's experiment.

Apparatus required: Electrically maintained tuning fork, light weight pan, weight box, analytical balance, power supply, light weight string, stand with clamp and pulley.

Theory:

1. Standing waves in strings and normal modes of vibration:

When a string under tension is set into vibrations, transverse harmonic waves propagate along its length. The speed of the wave in the stretched string depends on the tension in the string and mass per unit length of the string and is given by:

$$v = \sqrt{\frac{T}{\mu}} \quad (1)$$

where T is the tension in the string which is equal to Mg . M is the mass suspended on the string and g is the acceleration due to gravity and μ is the mass per unit length of the string, given by $\mu = m_s/L_o$. m_s is the mass of the string and L_o is the total length of the string.

A string can be set into vibrations by means of an electrically maintained tuning fork. When the other end of the string is clamped to a rigid support (pulley in present case), reflected waves will also exist. The incident and reflected waves will superimpose to produce transverse stationary waves in the string. The string will vibrate in such a way that the clamped points of the string are nodes and the anti-node exists at the middle.

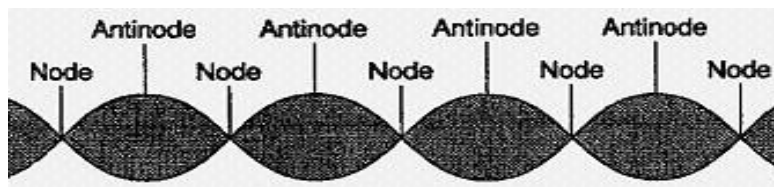


Fig. 1: Schematic representation of standing waves showing nodes and antinodes.

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The loops are formed from the end of the rigid support where it touches the pulley to the position where it is fixed to the prong of tuning fork. If l is the length of the string between two successive nodes, then

$$l = \frac{\lambda}{2} \quad (2)$$

where, λ is the wavelength of the traversing wave. The frequency (f) of the vibration is given by

$$f = \frac{v}{\lambda} = \frac{v}{2l} \quad (3)$$

substituting the value of v from Eq. (1), we get

$$f = \frac{1}{2l} \sqrt{\frac{T}{\mu}} \quad (4)$$

2. Transverse mode arrangement:

In this arrangement, the vibrations of the prongs of the tuning fork are in the direction perpendicular to the length of the string. The experimental setup with transverse mode arrangement is shown in Figure 2.

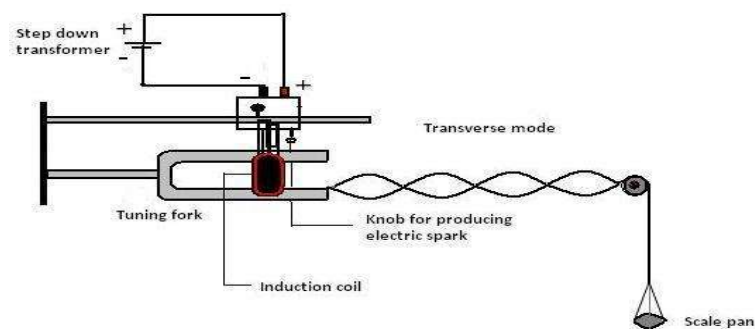


Fig. 2: Experimental setup with tuning fork in transverse mode arrangement.

In transverse mode, the string also completes one vibration when the tuning fork completes one. Hence in this mode, frequency of the tuning fork is equal to the frequency of the string which is same as Eq. 4.

$$f = \frac{1}{2l} \sqrt{\frac{T}{\mu}} = \frac{1}{2l} \sqrt{\frac{Mg}{\mu}} \quad (5)$$

where, M is the mass suspended on the string. $M = m + M'$. m is the mass of the weights placed on the scale pan and M' is the mass of the scale pan attached to the string. If ' P ' loops are formed in length ' L ' (between two fixed ends) of the thread, then $l = L/P$. Thus, Eq. 5 can also be expressed as,

$$f = \frac{P}{2L} \sqrt{\frac{Mg}{\mu}} \quad (6)$$

3. *Longitudinal mode arrangement:*

In this arrangement, the tuning fork is set in such a manner that the vibrations of the prongs are parallel to the length of the string. The experimental setup with longitudinal mode arrangement is shown in Figure 3.

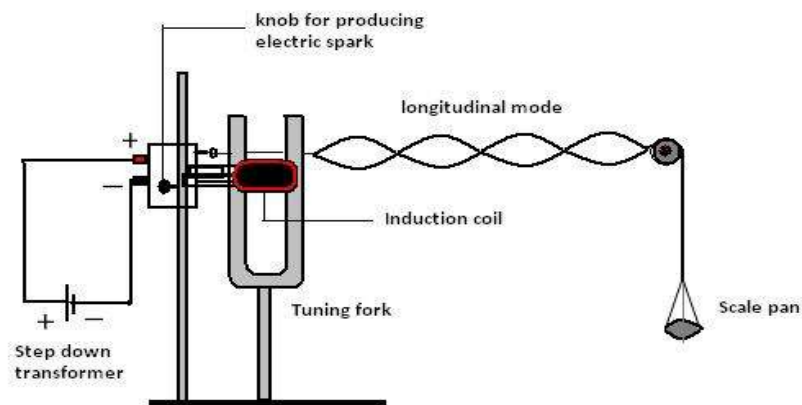


Fig. 3: Experimental setup with tuning fork in longitudinal mode arrangement.

In longitudinal mode, the string completes half of its vibration when the tuning fork completes one. Hence in this mode, frequency of the tuning fork is double the frequency of the string and is given as

$$f = \frac{P}{L} \sqrt{\frac{Mg}{\mu}} \quad (7)$$

Procedure:

1. Find the mass of the scale pan M' and arrange the apparatus as shown in figure.
2. Excite the tuning fork by switching on the power supply (advisable to use voltage more than 6V)
3. Adjust the position of the pulley in line with the tuning fork.
4. Change the load in the pan attached to the end of the string.
5. Adjust the applied voltage so that vibrations and well defined loops are obtained.
6. The tension in the string increases by adding weights in the pan slowly and gradually. For finer adjustment, add milligram weight so that nodes are reduced to points at the edges.
7. Count the number of loop and the length of each loop. For example, if 4 loops formed in the middle part of the string. If ' L ' is the distance in which 4 loops are formed, then distance between two consecutive nodes is $L/4$.
8. Note down the weight placed in the pan and calculate the tension T .
9. Tension, $T = (\text{wts. on the pan} + \text{wt. of pan}) g$.
10. Repeat the experiment for longitudinal and transverse mode of vibrations.
11. Measure one-meter length of the thread and find its mass to find the value of mass produced per unit length (m_s).

Observations and calculations:

Mass of the pan, $M' = \dots\dots\dots$ gm

Mass per unit length of the string,

$\mu = \dots\dots\dots$ gm/cm

For transverse mode arrangement:

Frequency

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$$f = \frac{P}{2L} \sqrt{\frac{Mg}{\mu}} \quad (8)$$

Table-1: Frequency of transverse mode arrangement

S.No.	Weight (<i>M</i>) gm	No. of loops (<i>P</i>)	Length of thread (<i>L</i>) cm	Length of each loop (<i>L/P</i>) cm	Tension (<i>T</i>) (<i>m + M'</i>) gm	Frequency (<i>f</i>) Hz
1						
2						
3						

Mean frequency= ----- Hz

For longitudinal mode arrangement

Frequency

$$f = \frac{P}{L} \sqrt{\frac{Mg}{\mu}} \quad (9)$$

Table-2: Frequency of longitudinal mode arrangement

S.No.	Weight (<i>W</i>) gms	No. of loops (<i>P</i>)	Length of thread (<i>L</i>) cms	Length of each loop (<i>L/P</i>) cms	Tension (<i>T</i>) (<i>W+w</i>) gms	Frequency (<i>f</i>) Hzs
1						
2						
3						

Mean frequency= ----- Hz

Precautions:

1. The thread should be uniform and inextensible.
2. Well defined loops should be obtained by adjusting the tension with milligram weights.
3. Frictions in the pulley should be least possible.

Sample Oral Questions

1. What is the scientific and technological importance of the present experiment?
2. What are the practical applications based on the theory and outcome of this experiment?
3. What understanding you build with this experiment?
4. What are standing waves?
5. Give few other examples of standing waves.
6. What is the condition of formation of standing waves?
7. What is Resonance?
8. Explain the difference between longitudinal and transverse waves.
9. In our experiment which type of wave passing along the thread?
10. Why the frequency is different in longitudinal mode when compared with the transverse mode? Which mode is greater and why?

EXPERIMENT NO. 7

Aim of the experiment: Measurement of voltage and frequency of a given signal using a cathode ray oscilloscope

Apparatus Required: Cathode Ray Oscilloscope, Function Generator(s), a pair of BNC Connectors

Introduction: A cathode ray oscilloscope (CRO) can be used to measure the voltage and frequency of given unknown signal. (A brief description of CRO is given at the end of this manual). A RC oscillator can be used to generate an electrical signal of desired frequency and amplitude. In the given experiment the RC oscillator has to be used to generate the signal and the CRO will be used to measure its voltage and frequency.

Procedure: Switch on the oscillator. Place the time base knob in horizontal input position and wait for a couple of minutes. Notice a bright spot of light on the screen of the CRO. The spot can be moved in vertical or horizontal direction by using the horizontal position knob and vertical position knob respectively. Place the time base in appropriate position (i.e. 1ms/cm or 0.1 ms/cm or any other value). A bright line can be noticed on the CRO screen. Your CRO is now ready to measure voltage and frequency of the unknown signal.

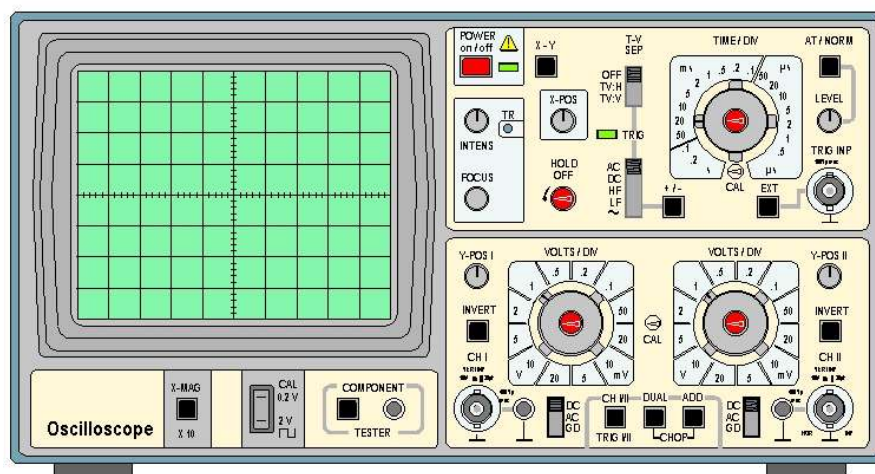


Fig. 1: The Cathode ray oscilloscope used in Physics Laboratory.



Fig. 2: The function generator used in Physics Laboratory.

Selection of frequency:

The RC oscillator is having several knobs, which can be used to select frequency of the signal to be generated. In the top left hand corner, you would see three knobs. These knobs can be used to select frequency value, which can be represented by three digits. For example, suppose you are setting the left knob to 6, the middle knob to 5 and the extreme right knob to 4. Then the selected frequency will be 654Hz. Below these three knobs you will get a multiplier. The multiplier will multiply the above selected frequency. Thus if you select 654Hz and multiplier position is 10 then the overall frequency will be 6540Hz.

Selecting amplitude of the signal:

The RC oscillator provides you option to vary the voltage of the signal to be generated. This can be done using two voltage selecting knobs (Fine and Coarse). Therefore, using different knobs a signal of given amplitude and given frequency can be generated by the RC oscillator. This signal can now be used as an input to the CRO and its frequency and voltage can be measured.

Voltage Measurements:

Use the signal generated by RC oscillator as an input to CRO. Place the Y amplifier in proper value. From vertical scale measure the peak to peak value. This will give the value of peak to peak voltage of the signal.

Lissajous Figures:

Theory: When two simple harmonic motions are plotted against each other at right angles, the resulting configuration is called a Lissajous figure. Simple harmonic motions plotted against time gives sinusoidal configurations. Two sinusoidal electrical inputs given to an oscilloscope will give a Lissajous pattern on the screen. The particular pattern depends upon the frequency, amplitude and phase of the applied inputs. The frequency ratio of the inputs may be determined from an analysis of the Lissajous figure produced. If a Lissajous figure is enclosed in a rectangle whose sizes are parallel to the formation axes of the figure, the frequency ratio of the two inputs may be determined by counting the points of tangency to the sides of the rectangle enclosing the pattern. Once the frequency ratio is known, the input frequency can also be determined from the same.

Procedure:

1. Connect one signal generator to the vertical input and the other to the horizontal input of the oscilloscope. Switch controls so that the oscilloscope accepts the output of the signal generator instead of the horizontal sweep. Set both the generators for 1000 cycles (say) and make gain adjustments until an ellipse of satisfactory size is observed on the screen. Adjust controls as necessary to stop the ellipse. By switching one of the generators off and on, cause the ellipse to change phase, noting the various shapes it assumes. By phase changes and amplitude adjustments, one may try to get a circular configuration.
2. Leaving the vertical input at 1000 cycles and assuming it to be the standard, adjust the horizontal input generator (the variable) approximately 500 c.p.s. to obtain the 1-2 Lissajous figure, a figure 8 on its side.
3. Next obtain the 2:1 pattern by varying the horizontal input frequency. This is an upright figure 8.
4. In like manner, obtain Lissajous figures down to 1:5 and upto 5:1. Sketch all the figures obtained and compare the frequency. Obtained from the Lissajous ratios with the dial reading of the horizontal input signal generator.

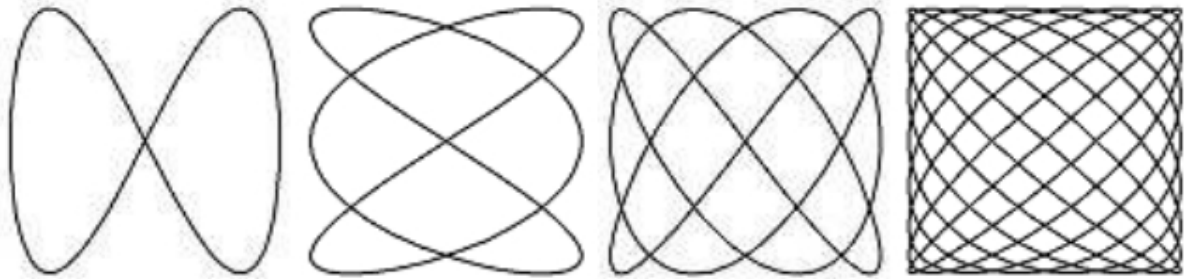


Figure 3: Sample Lissajous Figures

5. Now, remove one signal generator from the oscilloscope and connect the given unknown source. Changing the frequency of the signal generator, various Lissajous figures may be obtained (e.g. circle, 8 shape, etc.). Hence, from the known ratio of the respective Lissajous figures, the frequency of the AC source can be measured.

Observation Table:

Table 1: Measurement of voltage by CRO

Sl. No.	Voltage from Source (V_{p-p})	Peak-to-peak voltage measured by CRO			Ratio of V_{p-p}/V_{CRO}
		Y-amplifi setting (V/div)	Vertical scale No of div.	V_{CRO} (V)	

Table 2: Measurement of frequency

Sl. No.	Freq. of function generator (f in Hz)	Freq. measurement using CRO (f_o in Hz)			Ratio f/f_o
		Value of time base	No. of div.	f_o (in Hz)	

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Table 2: Measurement of unknown frequency with Lissajous figures

Horizon input Frequency on dial	Shape of figure	No. of tangency Points on X-axis on Y-axis	Vertical/	Horizontal

Precautions: The precautions to be taken while performing the experiment are:

1. Correct output/input terminals of function generators and CRO should be chosen.
2. Make sure that correct types of wave output have been chosen from function generator (Sine or Triangular or Square or DC)
3. Don't crank up the voltage level from function generator at its maximum limit.
4. The image produced over the CRO screen should be sharp and as thin as possible to take the correct readings.

Sample Oral Questions

1. What is the scientific and technological importance of the present experiment?
2. What are the practical applications based on the theory and outcome of this experiment?
3. What understanding you build with this experiment?
4. What is a cathode ray? How is it generated?
5. On what principle a cathode ray oscilloscope works?
6. How the screen gets discharged from excessive negative charges due to incoming electrons?
7. Is the cathode ray striking the CRO screen a line or a spot?
8. Why the screen of a CRO glows green when cathode ray strikes on it?
9. What is a function generator?
10. What type of functions are generated using the function generator of our lab?
11. What are Lissajous figures?

EXPERIMENT NO. 8

Aim of the Experiment: To determine the wavelength of prominent spectral lines of mercury light by a plane transmission grating using normal incidence.

Apparatus Required: A spectrometer, Mercury lamp, Transmission grating, Reading lamp and Reading lens.

Theory: Basic description of spectrometer is given in page no. 35.

The wavelength λ of any spectral line using plane transmission grating can be calculated from the formula $(e+d) \sin\theta = n\lambda$, where $(e+d)$ is the grating element, θ is the angle of diffraction, and n is the order of the spectrum. If there are N lines per inch ruled on the grating surface, then the grating element is given by $(e+d) = 2.54/N$ cm. Hence $(2.54/N) \sin\theta = n\lambda$, or $\lambda = 2.54 \sin\theta / nN$ cm.

Procedure

1) Adjustment of the grating for normal incidence:

The initial adjustment of the spectrometer is made as usual. The plane transmission grating is mounted on the prism table. The telescope is released and placed in front of the collimator. The direct reading is taken after making the vertical cross-wire to coincide with the fixed edge of the image of the slit, which is illuminated, by a monochromatic source of light. The telescope is then rotated by an angle 90° (either left or right side) and fixed. The grating table is rotated until on seeing through the telescope the reflected image of the slit coincides with the vertical cross-wire. This is possible only when the reflected image of the slit coincides with the vertical cross-wire. This is possible only when a light emerging out from the collimator is incident at an angle 45° to the normal to the grating. The vernier table is now released and rotated by an angle 45° towards the collimator. Now light coming out from the collimator will be incident normally on the grating. (Fig 1).

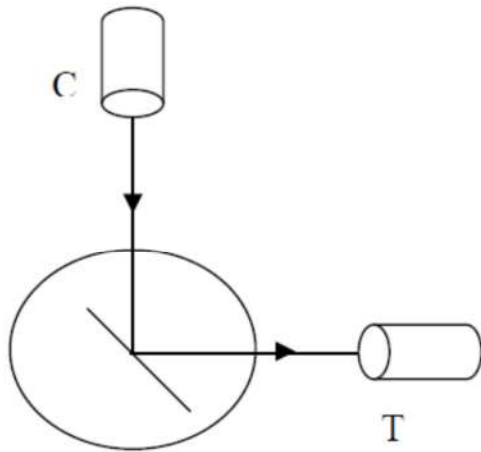


FIG.1

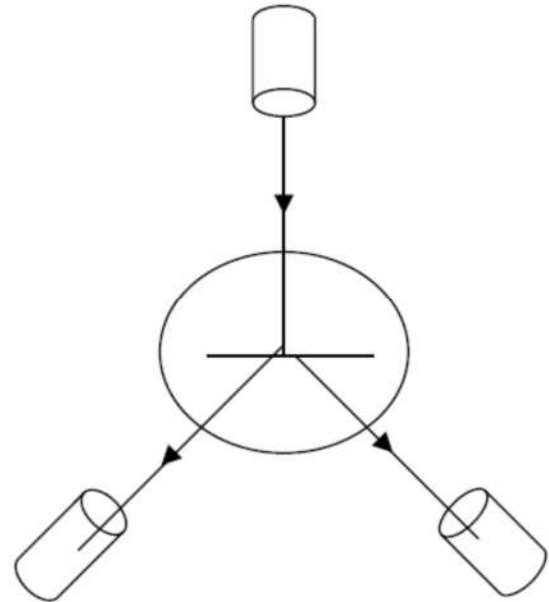


FIG.2

2) Standardization of the grating:

The slit is illuminated by sodium light. The telescope is released to catch the different image of the first order on the left side of the central direct image. The readings in the two vernier are noted. It is then rotated to the right side to catch the different image of the first order, the readings are noted. (Fig.2). The difference between the positions of the right and left sides given twice the angle of diffraction 2θ . The number of lines per meter of the grating (N) is calculated by using the given formula assuming the wavelength of the sodium light as 589.3 nm.

3) Wavelengths of the spectral lines of the mercury spectrum:

The sodium light is removed and the slit is now illuminated by white light from mercury vapour lamp. The telescope is moved to either side of the central direct image, the diffraction pattern of the spectrum of the first order and second order are seen. The readings are taken by coinciding the prominent lines namely violet, blue, bluish – green, yellow1, yellow2 and red with the vertical wire. The readings are tabulated and from this, the angles of diffraction for different colours are

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determined. The values are tabulated in Table 1. The wavelengths for different lines are calculated by using the given formula.

Observations:

A. For the adjustment of grating for normal incidence:

1. Least count of the Spectrometer scale:

Value of 1 division of main scale =

Divisions of main scale are equal to divisions of vernier scale.

Value of 1 division of vernier scale.

Least count of Spectrometer scale:

= value of 1 division of main scale – 1 division of vernier scale.

2. Reading of the telescope for direct image of the slit:

V1 = V2 =

3. Reading of the telescope after rotating it through 90° :

V1 = V2 =

4. Reading of prism table when reflected image of the slit coincides with the vertical cross wire V1 = V2 =

1. Reading of prism table when rotated through 45° or 135° :

V1 = V2 =

1. Number of lines N ruled per inch on the grating

2. Grating element $(e+d) = 2.54/N$

B. Calculations:

For first order, $n=1$, $\lambda = (e+d) \sin\theta/2$ cm.

λ for Violet I colour =cm.

Calculate λ for all visible spectral lines also.

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Table 1: Table for the measurement of the angle of diffraction θ

Order	Color of the spectral line	Spectrum to the left of the direct images			Spectrum to the right of the direct images			$2\theta = X - Y$	Angle $= \theta$
		M.S.R	V.S.R	T.R=(X)	M.S.R	V.S.R	T.R=(Y)		
1 st order	Violet Window 1 Window 2								
	Blue Window 1 Window 2								
	Green Window 1 Window 2								
	Yellow Window 1 Window 2								
	Red Window 1 Window 2								
2 nd order	Violet Window 1 Window 2								
	Blue Window 1 Window 2								
	Green Window 1 Window 2								
	Yellow Window 1 Window 2								
	Red Window 1 Window 2								

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Results: The experimental values of the wavelength of spectrum colours along with the percentage errors are finally tabulated in Table 2.

Table 2: Experimental values and verification of wavelengths of spectrum colours.

Colour of the Spectral line	Wavelength as obtained by experiment	Standard value of wavelength	Percentage error %
Violet I		4047 Angstroms	
Blue		4358 Angstroms	
Green		5461 Angstroms	
Yellow I		5770 Angstroms	
Red		6234 Angstroms	

Source of Error and Precautions

1. The axes of the telescope and the collimator must intersect at and be perpendicular to the main axis of the spectrometer.
2. The collimator must be so adjusted as to give out parallel rays.
3. The telescope must be so adjusted as to receive parallel rays and form a well-defined image of the slit on the crosswire.
4. The prism table must be optically leveled.
5. The grating should be so mounted on the prism table that its ruled lines are parallel to the main axis of the spectrometer.
6. The plane of the grating should be normal to the incident light and its ruled surface must face the telescope so that the error due to non-parallelism of the incident rays is minimum.
7. The slit should be as narrow as possible and parallel to the ruled surface of the grating.
8. While handling the grating one should not touch its faces but hold it between the thumb and the fingers by edges only.
9. While taking observations of the spectral lines, the prism table must remain clamped.
10. The reading of both the verniers should be recorded. This eliminates the error due to non-coincidence of the center of the graduated scale with the main axis of the spectrometer.

Sample Oral Questions:

1. What is the scientific and technological importance of the present experiment?
2. What are the practical applications based on the theory and outcome of this experiment?
3. What understanding you build with this experiment?
4. What do you understand by diffraction of light?
5. How does it differ from interference of light?
6. What is a diffraction grating?
7. How do you measure the wavelength of light using grating?
8. What is grating element?
9. How do you adjust telescope and collimator for parallel rays?
10. How do you set the grating for normal incidence?
11. Why the ruled surface of grating face should forward the telescope?
12. How many orders of spectra are you getting with the grating?
13. What is the difference between a prism spectrum and a grating spectrum?
14. What are the various series of lines observed in hydrogen spectrum?
15. What is Rydberg constant?
16. A transmission grating with 2000 lines/cm is illuminated by a beam of 694.3 nm light from a laser. Spots of light, on both sides of the undeflected beam, appear on a screen 2.0 m away.
 - ❖ How far from the central axis is either of the two nearest spots?
 - ❖ Find how much difference it makes whether you use the approximation $\sin \theta \approx \theta$

EXPERIMENT NO: - 9

Aim of the experiment: To determine the electromotive force (emf) of an unknown cell using a stretched wire potentiometer.

Apparatus required: Stretched wire potentiometer, jockey, galvanometer, Power source, Cell (Leclanché), Standard cell, rheostat, resistance box, connecting wires, plug and key.

Theory: If a current i flows through the potentiometer wire of L cms and Resistance R_p ohms. If R is the series resistance series with the potentiometer, then

$$i = \frac{E}{(R + R_p)} \quad (1)$$

Where E is the e.m.f of the cell C. The potential drop across the end A and B of the potentiometer wire is

$$V = i \times R_p = \frac{ER_p}{(R+R_p)} \quad (2)$$

Hence the potential drop per centimeter of the wire v is

$$v = \frac{V}{L} = \frac{E}{(R+R_p)} \times \frac{R_p}{L} = \frac{E}{(R+R_p)} \times \frac{R_p}{1000} \text{ volts} \quad (3)$$

If the cells C_1 and C_2 of e.m.f.s E_1 and E_2 respectively are connected to the circuit and the required lengths of the potentiometer wire for balance are l_1 and l_2 then the e.m.f. E_1 and E_2 are given by

$$E_1 = vl_1 = \frac{E}{(R+R_p)} \times \frac{R_p}{1000} \times l_1 \text{ volts} \quad (4)$$

$$E_2 = vl_2 = \frac{E}{(R+R_p)} \times \frac{R_p}{1000} \times l_2 \text{ volts} \quad (5)$$

The ratio of emf of two cell is

$$\frac{E_1}{E_2} = \frac{l_1}{l_2} \quad (6)$$

The e.m.f. of single cell is given by

$$E_1 = \frac{l_1}{l_2} \times E_2 \quad (7)$$

By simply determining the balancing length emf of unknown cell can be calculated from the above equation.

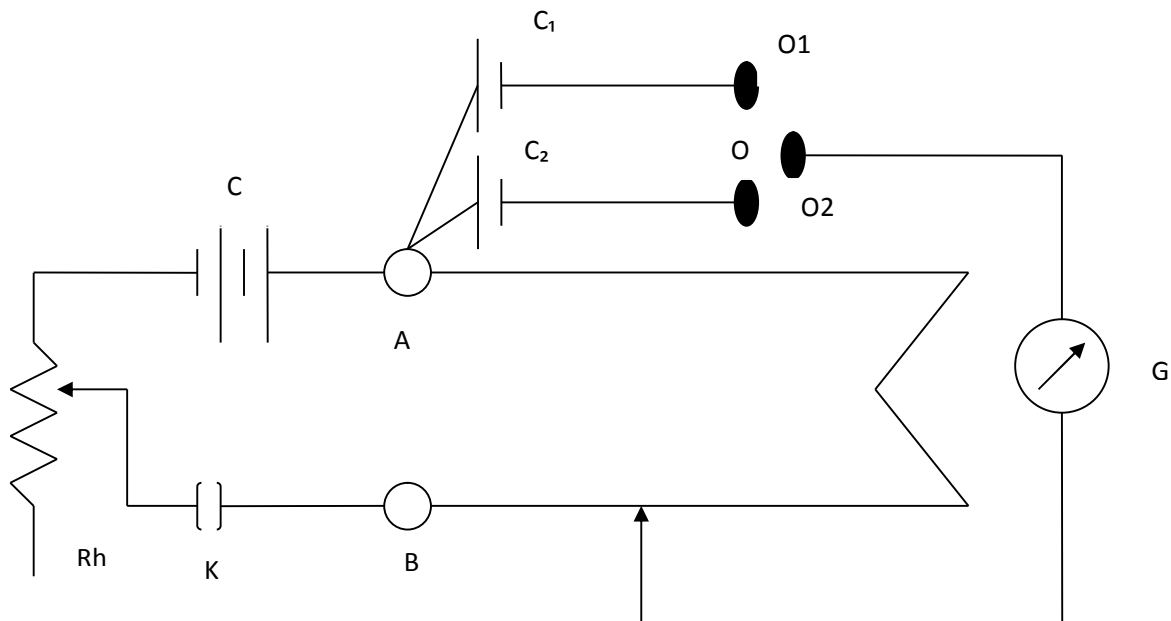


Fig. 1: Circuit diagram to determine the emf of an unknown cell

Procedure:

1. Connect the apparatus as is shown in Fig 1. Connect the positive terminal of the power supply to the end A of the potentiometer wire and the negative through the rheostat.
2. Connect the positive terminal of two cells C_1 and C_2 to A and negative terminals to the binding screws O_1 and O_2 of the two-way key K1.
3. Join the third binding screw to one terminal of the galvanometer to the jockey J.
4. Make the resistance R_h zero and resistance R maximum. Put the jockey J in contact with the first and last. If the galvanometer deflection is opposite, then the connection for C_1 is correct.

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5. Repeat the process for Cell C₂
6. After verifying the deflection find the null point for both C₁ and C₂
7. If the length for C₁ is greater than that of C₂ then the emf for E₁ is greater than emf for E₂ for C₂.
8. Adjust the rheostat for different values and take the reading

Observation:

No. of obs.	Cell	Null points			Total length in (cm)	$E_1/E_2 = l_1/l_2$	Mean E_1/E_2
		Wire number (cm)	Scale reading (cm)	Mean scale reading (cm)			
1	First (E ₁)	10th	 = (x ₁)	900+x ₁		
	Second (E ₂)						
2	First (E ₁)	9th	 = (x ₃)	800+x ₃		
	Second (E ₂)						
3	First (E ₁)	10th	 = (x ₅)	900+x ₁		
	Second (E ₂)						

The e.m.f. of an unknown cell is calculated by

$$E_1 = \frac{l_1}{l_2} \times E_2 \quad (8)$$

Result & Discussion:

Precautions:

1. All the connection terminals should be clean and tight.
2. Jockey should be held vertical.

Sample Oral Questions:

1. What is the scientific and technological importance of the present experiment?
2. What are the practical applications based on the theory and outcome of this experiment?
3. What understanding you build with this experiment?
4. What is electromotive force? How is it different from potential difference?
5. Is the voltage marked in a AAA battery potential difference or emf? Why?
6. What is the emf of a completely discharged 12 V battery?
7. Can we determine the emf of a cell using a multimeter or voltmeter? Support your answer with logical reasoning.

EXPERIMENT NO. – 10

Aim of the experiment: To study the frequency response and quality factor of series LCR circuit.

Apparatus required: Cathode ray oscilloscope (CRO) with probe, Function generator, Inductor, Capacitor, decade resistance box, a $1\text{k}\Omega$ resistor, connecting wires with BNC and crocodile clip terminations.

Theory: If we apply a sinusoidal voltage to a series LCR circuit the net impedance offered by the circuit to the flow of current will be the vector sum of that offered by the resistive (frequency independent) part as well as the reactive (frequency dependent) part, i.e.

$$Z = \sqrt{R^2 + X^2}$$
$$\text{or, } Z = \sqrt{R^2 + (X_L - X_C)^2}$$
$$\text{or, } Z = \sqrt{R^2 + \left(\omega L - \frac{1}{\omega C}\right)^2}$$

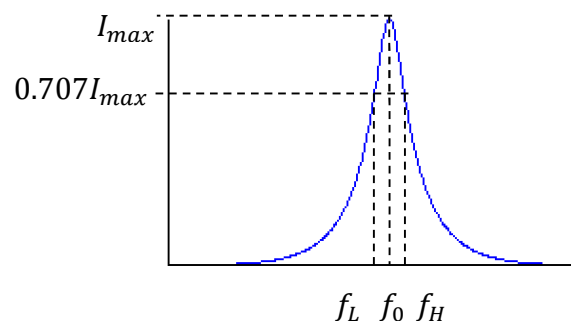
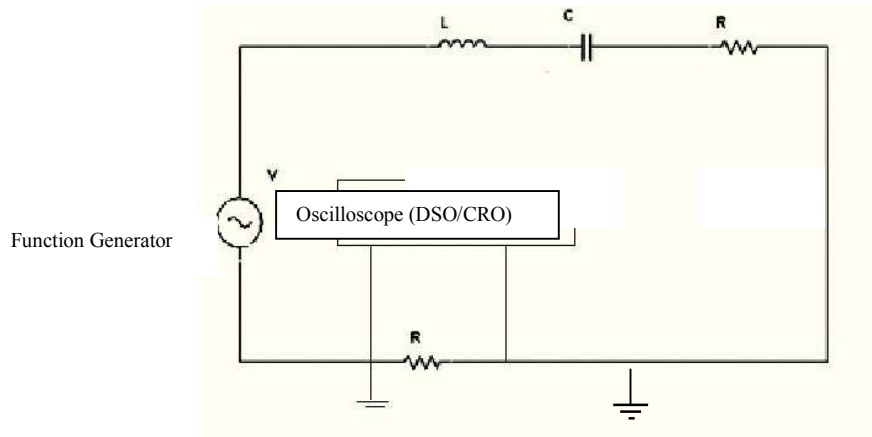
The current in the circuit is given by

$$I = \frac{E}{Z}$$
$$\text{or, } I = \frac{E}{\sqrt{R^2 + \left(\omega L - \frac{1}{\omega C}\right)^2}}$$

and the phase angle between the applied voltage and the current through the circuit is given by,

$$\phi = \tan^{-1} \left(\frac{\omega L - \frac{1}{\omega C}}{R} \right)$$

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If the frequency (ω) of applied voltage, matches the natural frequency (ω_0), of the circuit then the inductive reactance and the capacitive reactance equals each other i.e.,

$$\omega_0 L = 1/\omega_0 C$$

and the current in the circuit is solely decided by the value of R, i.e.,

$$I = \frac{E}{R}$$

The frequency at which the inductive reactance equals the capacitive reactance is the natural frequency or the resonant frequency of the circuit. The resonant frequency of an LCR circuit depends upon the values of L and C by the relation

$$\omega_0^2 = \frac{1}{LC}$$
$$f_0 = \frac{1}{2\pi\sqrt{LC}}$$

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The sharpness of resonance for a particular value of L and C depends upon the value of R and is computed from the plot of I versus f by the relation

$$Q = \frac{f_0}{f_H - f_L}$$

The 'Quality factor' or the 'Q factor' is a dimensionless parameter that describes how under-damped an oscillator or resonator is. Higher values of Q indicate lower rate of energy loss relative to the energy stored in the oscillator.

There are two separate definitions of the quality factor that are equivalent for high Q resonators but are different for strongly damped oscillators.

Generally, Q is defined in terms of the ratio of the energy stored in the resonator to the energy being lost in one cycle:

$$Q = 2\pi \times \frac{\text{Energy stored}}{\text{Energy dissipated per cycle}}$$

The factor of 2π is used to keep this definition of Q consistent (for high values of Q) with the second definition:

$$Q = \frac{f_0}{\Delta f} = \frac{\omega_0}{\Delta\omega}$$

where, f_0 is the resonant frequency, Δf is the bandwidth, ω_0 is the angular resonant frequency, and $\Delta\omega$ is the angular bandwidth.

The definition of Q in terms of the ratio of the energy stored to the energy dissipated per cycle can be rewritten as:

$$Q = \omega \times \frac{\text{Energy Stored}}{\text{Power loss}}$$

where, ω is defined to be the angular frequency of the circuit (system), and the energy stored and power loss are properties of a system under consideration.

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Procedure:

1. Before making any connections (except mains power cords) switch on the oscilloscope and ensure that the sensitivity of the oscilloscope is set at a low value, for example 10V/div.
2. Switch on the function generator and adjust its voltage amplitude to a level $\leq 5V$ and let it remain constant throughout the experiment.
3. Also, make sure you are using a compatible probe with the CRO / DSO and function generator. Please note that the probes may appear similar but are not interchangeable.
4. Set the function generator in sinusoidal signal mode.
5. Connect the L, C, R and a $1K\Omega$ resistor in series and the two extreme ends to the function generator output through a BNC (as shown in figure).
6. Ensure that the $1K\Omega$ resistor is connected to the function generator output ground.
7. Connect the ground terminal of the oscilloscope probe to the ground of function generator, and the other terminal of the probe in such a way that the oscilloscope gets connected just across the $1k\Omega$ resistor as shown in the preceding circuit diagram.
8. Set $R=0$ from the resistance box.
9. Adjust the frequency output of the function generator to 10Hz.
10. Record the voltage amplitude of the signal shown by the oscilloscope.
11. Go on incrementing the frequency logarithmically and record your observation, i.e., repeat step-9 till you reach 1MHz.
12. In next set of experiment increase R to $1k\Omega$ and repeat steps 8, 9 &10.
13. Once again increase R to $4k\Omega$ and repeat steps 8, 9 &10.
14. Current (in ampere) can be obtained by dividing the voltage amplitude by the resistor value (1000Ω in our case). Value of current in mA is numerically equal to the voltage amplitude itself as long as the value of resistor across the oscilloscope is maintained.

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Observations:

$$L = 100\text{mH}$$

Frequency ν (Hz)	R = 0 Ω		R = 1k Ω		R = 4k Ω	
	V (volt)	I (mA)	V (volt)	I (mA)	V (volt)	I (mA)
100						
200						
..						
..						
900						
1k						
2k						
..						
..						
9k						
10k						
20k						
..						
..						
90k						
100k						

Plot: Plot the current versus frequency for the three resistor values (i.e., 0 Ω , 1 k Ω and 4 k Ω) on the same graph sheet neatly and mark the resonant frequency (f_0), the peak amplitude I_{max} and $\frac{I_{max}}{\sqrt{2}}$ for each plot. Sketch a dotted horizontal line corresponding to $\frac{I_{max}}{\sqrt{2}}$ and mark that it cuts the plot at two points. Drop perpendiculars from these points on the axis and mark the lower cutoff f_L and upper cutoff frequency f_H points. Note that the gap between the cutoff frequencies widens as the value of R increases.

Calculations:

Calculating the value of capacitor

$$C = \frac{1}{4\pi^2 L f_0^2}$$

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Calculating quality factor (for different values of R)

$$Q = \frac{f_0}{f_H - f_L}$$

Result & discussion:

L = 100 mH					
R	f_0 (Hz)	C (μ F)	f_L (Hz)	f_H (Hz)	Q
0 Ω					
1 K Ω					
4 K Ω					

Compare the resonant frequency and quality factors for the three resistor values. Comment on the results obtained by you.

Sample Oral Questions

1. What is the scientific and technological importance of the present experiment?
2. What are the practical applications based on the theory and outcome of this experiment?
3. What understanding you build with this experiment?
4. Define capacitance, inductance and resistance.
5. What is the purpose of capacitors?
6. What is the purpose of inductors?
7. What is the purpose of resistor?
8. What is an impedance?
9. What is time constant?
10. What will happen if both capacitor and inductor are connected in a circuit?
11. What is quality factor?
12. What is resonant frequency?
13. LCR series circuit is called Acceptor. Why?
14. What is the relation between impedance and current in this series circuit?
15. What is the status of a current in series and parallel connections?

EXPERIMENT NO. 11

Aim of the Experiment: To find the specific rotation of sugar solution by using a polarimeter.

Apparatus Required: Polarimeter, Volume flask, Sugar, Distilled water, weighting balance.

Theory: Optical activity is an intriguing property of certain molecules. It was found that solution of sugar and certain other naturally occurring chemicals would rotate a beam of polarized light passing through solution. They called it optically active substance. The instrument used to measure the rotation of polarized light is called Polarimeter and shown in Fig. 1.



Fig. 1: Polarimeter apparatus

Fig. 2 shows a schematic diagram of a polarimeter. The polarimeter is made up of two Nicol prisms (the polarizer and analyzer). The polarizer is fixed and the analyzer can be rotated. The light waves may be considered to correspond to waves in the string. The polarizer allows only those light waves which move in a single plane. This causes the light to become plane polarized. When the analyzer is also placed in a similar position it allows the light waves coming from the polarizer to pass through it. When it is rotated through the right angle no waves can pass through the right angle and the field appears to be dark. If now a glass tube containing an optically active solution is placed between the polarizer and analyzer the light now rotates through the plane of polarization through a certain angle, the analyzer will have to be rotated in same angle.

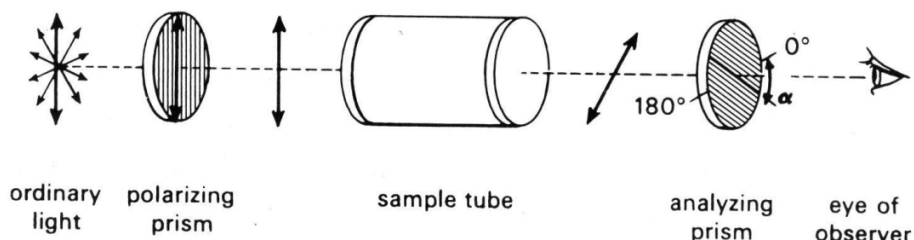


Fig. 2 Schematic diagram of Polarimeter with sample tube containing an optically active substance

If a column of solution of length l cm and at temperature $t^\circ\text{C}$, contains m gms of active substance per c.c. of the solution then it will produce a rotation of the plane of polarization of a plane-polarised light of wavelength λ transmitted through it, by an amount θ given by

$$\theta = \frac{slm}{10} \quad (1)$$

Here s is the specific rotation of the substance and is defined as the rotation produced by a column of solution of one decimeter (=10 cm) in length containing 1 gm of active substance per c.c. of solution. If c be the percentage strength of the solution (i.e. c gms. of active substance are present in 100 c.c of the solution) then

$$m = \frac{c}{100} \quad (2)$$

Thus, Eq. (1) becomes

$$s = \frac{1000 \theta}{cl} \quad (3)$$

Procedure:

(A) To Prepare the solution of $c\%$ strength

The mass (m_1) of a dry and empty flask is determined by a balance. Then by introducing a certain quantity of the substance (with which the solution of a given strength is to be prepared) in a flask, the total mass (m_1+w) is determined. Thus the mass of the substance taken is w gms. Now distilled

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water of volume $100 w / c_1$ c.c. is added to the flask to get a solution of $c_1\%$ strength by volume (c_1 should be the highest strength say 10 % to be used). By shaking the flask, the substance should be completely dissolved in water and if necessary the solution should be filtered.

Table 1: Preparation of sugar solution of $c_1\%$ strength by volume

No. of Obs.	Vol. in c.c. of sol. to be prepared (v)	Mass of solid required to prepare v c.c. of $c_1\%$ strength $w = \frac{c_1 v}{100}$	Mass of		Vol. of water added to the flask to make $c\%$ strength $= \frac{100 w}{c_1}$
			Dry and empty flask (ml)	Flask + subs. ($m_1 + w$)	

(B) To find the angle of rotation (θ) for the solution $c_1\%$ strength

1. Take the polarimeter tube and clean well both the sides such that it is free from dust. Now fill the tube with pure water and see that no air bubble is enclosed in it. Place the tube in its position inside the polarimeter. Switch on the source of light and look through the eyepiece.
2. In case of half shade polarimeter, two halves of unequal intensity is observed. Left half may be bright and the right half may be dark, or *vice versa*. By rotating the analyzer eyepiece system, the bright-dark pair gets interchanged to dark-bright pair, or *vice versa*. Rotate the analyzer (first in clockwise direction and then in anticlockwise direction) until the intensity of two halves is about to interchanged and circular field of view appears equally bright.
3. Take the first reading at equal intensity position (either bright or gray) and also record the second reading at 180° apart from this position, in both the directions (clockwise and anticlockwise). Find the mean of two directions reading separately for both the position.
4. Prepare a sugar solution of known strength by dissolving the known amount of sugar. Take the polarimeter tube and remove the pure water. Fill it with the prepared sugar solution and again place it in the polarimeter.
5. Rotate the analyzer eyepiece system to obtain the equal intensity position, first in clockwise direction and then in anticlockwise direction. Note down the first position of the analyzer scale in the two directions. Find the mean reading. Repeat similarly, for second position at 180° apart.

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6. The difference between water and sugar solution reading gives the specific rotation.
7. The experiment can be repeated with sugar solutions of different concentrations.
8. Measure the length of the tube in centimeters and change it in decimeters.

(C) Length of the polarimeter tube $l = \frac{\text{----} + \text{----} + \text{----}}{3}$ cm =decimeter

(D) Determination of the vernier constant of the vernier scale attached to the polarimeter:

Suppose n divisions of vernier scales are equal to n_1 division of the main scale. The method is as follows:

1. Value of one division of circular main scale = x° . Value of one division of sliding vernier scale = $\frac{n_1}{n} \times x^\circ$
2. Least count of the polarimeter scale = value of 1 div. of main scale – value of 1 div. of vernier scale

$$= x^\circ - \left[\left(\frac{n_1}{n} \right) \times x^\circ \right]$$

(E) Table for the specific rotation:

Value of one division of the main scale =

No. of division on vernier scale =

Least count of vernier =

Table 2: Reading with water

No. of Obs	Analyzer Reading for position I			Mean	Analyzer Reading for position II			Mean
	M.S.	V.S	T		M.S.	V.S.	T	
Mean A=					Mean B=			

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Table 3: Reading with sugar solution

No. of Obs.	% strength of solution (c)	Vernier	Analyzer Reading			Rotation for the vernier	Mean Rotation $\theta = \frac{\theta_1 + \theta_2}{2}$	Specific Rotation $s = \frac{1000 \theta}{cl}$	Mean (s)
			M.S.	V.S.	T Mean				
	c_1	1 st position				$\theta_1 = C - A$			
			Mean C						
		2 nd position				$\theta_2 = D - B$			
			Mean D						
	c_2	1 st position				$\theta_1 = C - A$			
			Mean C						
		2 nd position				$\theta_2 = D - B$			
			Mean D						

M.S.: Main Scale Reading **V.S.:** Vernier Scale Reading, **T**=Total Reading=**M.S.**+**V.S.**

Calculation: According to the analyzer's reading

- For I position rotation produced by sugar solution, $\theta_1 = C - A$
- For II position rotation produced by sugar solution, $\theta_2 = D - B$

Thus, the mean rotation produced by cane sugar solution,

$$\theta = \frac{\theta_1 + \theta_2}{2} \quad (4)$$

Thus, the specific rotation of cane sugar solution is calculated from eq. (3).

(F) To draw (C- θ) graph

The different values of θ obtained for different known values of c are plotted on a graph paper with the percentage strength c along x-axis and its corresponding rotation θ along y-axis. The graph would be a straight line. The value of s is also be found from the coordinate of a point for this graph.

Results and discussion:

At a temperature° C and wavelength Å;

The specific rotation for cane sugar solution =

Standard value of specific rotation for cane sugar solution =

Percentage Error =%

Precautions:

1. The polarimeter tube should be well cleaned.
2. Water used should be dust free.
3. Whenever a solution is changed, rinse the tube with the new solution under examination.
4. There should be no air bubble inside the tube.
5. The position of analyzer should be set accurately.
6. The temperature and wavelength of light used should be stated.
7. Reading should be taken when halves of the field of view becomes equally illuminated.

Sample Oral Questions:

1. What is the scientific and technological importance of the present experiment?
2. What are the practical applications based on the theory and outcome of this experiment?
3. What understanding you build with this experiment?
4. What do you mean by unpolarized light, plane polarized light, circularly polarized light and elliptically polarized light?
5. How are plane polarized light, circularly polarized light and elliptically polarized light produced experimentally?
6. How do you analyze and differentiate between a plane polarized light, circularly polarized light and elliptically polarized light?
7. What is a half-wave plate and a quarter-wave plate?
8. Define optically active substance.
9. What do you mean by specific rotation?
10. What is a polarimeter?
11. What is the principle of Laurent's Half-shade polarimeter?
12. How does a biquartz polarimeter works?
13. What are the merits and demerits of Half-shade and biquartz polarimeter?

EXPERIMENT NO. 12

Aim of the Experiment: To determine the Hall voltage and calculate the Hall coefficient and carrier concentration of a semiconductor sample.

Apparatus Required: Electromagnet, Constant current power supply, Hall probe with Digital Gauss meter, *n*-type or *p*-type Germanium crystal, Hall Effect setup consisting of a constant current source and a milli-voltmeter.

Theory of Hall Effect: When a current carrying semiconductor is placed in a perpendicular magnetic field, a voltage is developed across the specimen in a direction perpendicular to both the current and the magnetic field. The phenomenon is called the “Hall Effect”.

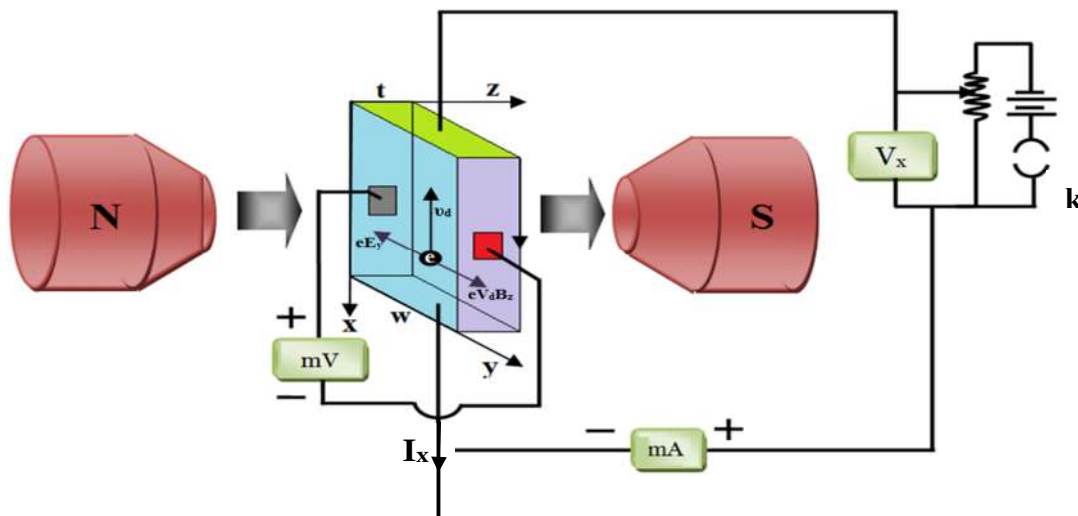


Fig. 1: Schematic diagram of Hall effect measurement set up.

Consider a rectangular semiconductor sample of width (w) and thickness (t) placed in xy plane. An electric field (E_x) is setup in the x -direction by allowing current (I_x) from a constant current source to flow through the sample. If the current carrying sample is placed in a magnetic field (B_z) applied perpendicular (along the z -direction) to the electric field E_x , Lorentz force acts on the charge

carriers (holes and/or electrons) moving with drift velocity v_d causing the charge carriers to curve along the y -direction. This results in accumulation of charge carriers at the side edges of the sample. This charge separation induces a transverse electric field (E_y) which develops a potential difference along the y -axis, known as Hall voltage (V_H). If a charge (e) moves along the x -axis (opposite of the direction of conventional current I_x for electrons while is as the direction of I_x for holes) and the magnetic field is along the z -axis then the force (F) is along the y -axis, as per the following Lorentz force equation,

$$F = ev_d \times B_z \quad (1)$$

If the charges are negative, then the force acts in the direction of the positive y -axis. Negative charges will therefore move left (see Figure 1) and hence, the Hall voltage, develops in the negative y -direction (E_y). The Hall voltage increases until the force due to the transverse electric field exactly balances the force due to the magnetic field and equilibrium is reached. i.e.

$$eE_y = ev_d B_z$$

Thus,
$$E_y = v_d B_z \quad (2)$$

The current I_x is the current density J times of the cross-sectional area of the conductor wt and the current density is ne times of the drift velocity; where n is the charge carrier number density (i.e., number of carriers per unit volume). So,

$$I_x = Jwt = neV_d wt \quad (3)$$

The Hall voltage related to the Hall field by,

$$V_H = \int_0^w E_y dy = -E_y w \quad (4)$$

Thus from equation (2), (3) and (4) we obtain,

$$V_H = -\left(\frac{1}{ne}\right) \frac{IB_z}{t} \quad (5)$$

Again, the *Hall-coefficient* is defined by,

$$R_H = \frac{E_y}{JB_z} \quad (6)$$

R_H measures the resulting Hall-field, along y, per unit transverse applied current and magnetic field. The larger is R_H the greater is E_y and B_z . In other words, R_H gauges the magnitude of the Hall effect. From equation (2), (3) and (6) we obtain,

$$R_H = -\frac{1}{ne} \quad (8)$$

Thus, from equation (5) and (7) we have the *Hall-coefficient* is,

$$R_H = \frac{V_H t}{I_x B_z} \quad \text{Ohm.m/G} \quad (9)$$

Working formula: $R_H = (\text{slope}) \times \frac{t}{B_z} \text{ Ohm.m/G}$

Procedure: The experiment can be divided into three sections:

- Calibration of magnetic field with current
- Measurement of Hall Voltage and determination of Hall coefficient and carrier concentration,
- Tracing the variation of Hall voltage with magnetic field.

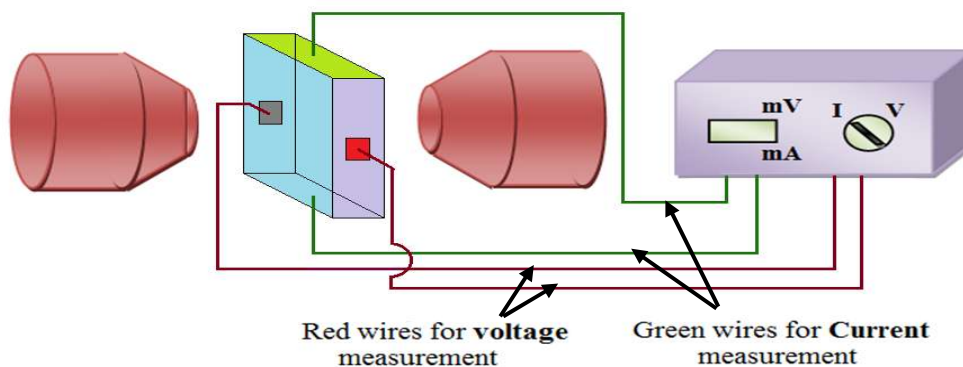


Fig. 2: Schematic diagram of the connections in Hall Effect measurement set up.

A. Calibration magnetic field with current

1. Connect the electromagnet leads to the constant current power supply.
2. Switch on the digital Gauss meter. Its reading is adjusted to zero using the 'ZERO' control key. This adjustment is to be done keeping the digital panel meter on X 1 range.
3. Switch on the electromagnet power supply. Start with the minimum current value to the electromagnet coil using the current knob.
4. Select the appropriate range of the Gauss meter. Insert the digital Hall probe (red in colour) into the magnetic field between the two poles of the electromagnet. The flat face of the probe is kept perpendicular to the direction of the magnetic field. The reading of the digital gauss panel meter multiplied by the range value gives the flux density of the magnetic field in Gauss.

(Note: If the display of the field strength is not shown, start with the $\times 10$ range).

5. Vary the current flowing through the electromagnets by adjusting the control knob of the constant current source.
6. Put the digital Gauss meter probe in the electromagnet poles and read the magnetic field B_z
7. Take at least 10 readings and note them in a format shown in Table 1. After taking reading, switch off the digital Gauss meter and insert the probe within its cap.

B. Measurement of Hall Voltage:

1. Check the semiconductor sample, i.e. the lightly doped Germanium crystal. The germanium crystal is mounted on a sunmica-decorated Bakelite strip. The crystal is connected with four leads. The two green wires are attached length-wise and are meant for supplying current. Whereas the remaining two red leads are attached width-wise and are for measuring the Hall voltage developed.
2. Connect the width-wise red contacts of the Ge crystal in Hall effect setup to the 'Voltage' terminal and length-wise green contacts to the 'Current' terminals.
3. Switch on the Hall effect set up and adjust the Hall current keeping the display switch on the current side.

4. Switch over the display to voltage side. There may be some voltage reading even outside the magnetic field. This is due to imperfect alignment of the four contacts of the Hall probe and is generally known as 'Zero Field Potential'.
5. Now place the sample vertically between the poles of the electromagnet so the magnetic field direction is perpendicular to the plane of the sample. Adjust the magnetic field to a certain value by adjusting the current of the constant current power supply.
6. Measure the Hall Voltage (V_H) as the function of current I_x keeping the magnetic field constant. For this, vary the Hall current and measure the respective Hall voltages (V_H) by switching over the range to 'voltage' side in the Hall Effect setup. Never exceed the current value more than 8 mA.
7. Plot the I_x versus V_H graph for each values of magnetic field B_z .
8. Find the slope V_H/I_x value.

C. Tracing the variation of Hall voltage with magnetic field.

1. First, adjust the current in Constant Current Power Supply to 0 Amp by adjusting the current control Nob.
2. Check the magnetic field by using the digital Gauss-meter. If it shows any value other than zero, make it zero by adjusting the 'zero adjustment Nob'.
3. Fix the magnetic field (by increasing the current) at some value (say 100 G). Remove the Gauss-meter and place the semiconducting sample.
4. Measure the Hall Voltage (V_H) by switching over the display to voltage side.
5. Repeat the steps (3) and (4) with an increment of 100 G in the values of the magnetic field.
6. Plot the V_H vs B_z graph.

Observations:

The thickness of the given Germanium crystal (*n*-type or *p*-type), $t = 0.5$ mm

The conductivity of the given crystal, $\sigma = 0.1$ ohm/cm

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Table-1 Calibration of Magnetic field with Current

Sl. No.	Current through the electromagnet (A)	Magnetic field generated between the poles (G)
1		
2		
:		
10		

Table-2 Observation of I_x vs V_H for $B_z = 1000$ G

Sl. No.	Current I_x (mA)	Hall Voltage V_H (mV)
1		
2		
:		
10		

Table-3 Observation of V_H vs B_z

Sl. No.	Magnetic field B_z (G)	Hall Voltage V_H (mV)
1		
2		
:		
10		

Calculations and Results:

$$\text{Slope, } \frac{V_H}{I_x} = \dots\dots\dots \text{ Ohm}$$

$$\text{So, Hall Coefficient, } R_H = \left(\frac{V_H}{I_x}\right) \frac{t}{B_z} = \dots\dots\dots \text{ Ohm. m/G}$$

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and carrier concentration, $n = -\frac{1}{R_H e} = \dots\dots\dots \text{m}^{-3}$

Precautions:

1. Electromagnet power supply should be connected to a 3-pin 15Amp AC main's socket having good earth connection.
2. Switch 'ON' or 'OFF' the current supply at zero current position.
3. The gauss meter probe is very delicate and should be used at temperature well below 50⁰ C.
4. The crystal contacts in Hall probe should neither be loose nor too tight. The crystal is thin and very brittle.
5. The current through the crystal should not be large enough to cause heating. It should not be exceeding 10mA.

Sample Oral Questions:

1. What is the scientific and technological importance of the present experiment?
2. What are the practical applications based on the theory and outcome of this experiment?
3. What understanding you build with this experiment?
4. What is Hall effect?
5. What are p- and n-type semiconductor?
6. Why is Hall potential developed?
7. If Hall coefficient is negative what does it indicate?
8. What is the significance of Hall coefficient?
9. What is Fleming's left hand rule?
10. Can Hall effect be observed in metals? Support your answer with proper reasoning.

