## **Experiment: 08**

**AIM:** To prepare and submit benzotriazole from o-phenylenediamine and calculate its percentage yield.

### **REFERENCES:**

- 1. Vogel's Textbook of Practical Organic Chemistry by Brian S. Furniss, Antony J. Hannaford, Peter W. G. Smith & Austin R. Tatchell; Fifth Edition; Page No.1163.
- Practical in organic chemistry, by Hitesh G. Raval, Sunil L. Baldania and Dimal A. Shah, Nirav Prakashan, Page No. 303.

### **REQUIREMENTS:**

Chemicals: o-phenylenediamine, Glacial acetic acid, and Sodium nitrite.

Apparatus: Beaker, Buchner funnel, Measuring cylinder, Filter paper, etc.

### **Principle:**

Benzotriazole can be prepared by treating o-phenylenediamine with nitrous acid (liberated during the reaction between sodium nitrite and acetic acid) to form monodiazonium salt that follows a spontaneous intramolecular cyclization reaction to produce benzotriazole.

### **Chemical Reaction:**





HNO

Mechanism:



Use:

Used as antifungal, antihypertensive, analgesic etc.

# PROCEDURE

- In a 250-ml beaker, dissolve 10.8 g (0.1 mol) of o-phenylenediamine in a mixture of 12 g (11.5 ml, 0.2 mol) of glacial acetic acid and 30 ml of water. Slight warming may be necessary.
- 2. Cool the clear solution to 15  $^{\circ}$ C and stir magnetically.
- 3. In one portion, add 7.5 g (0.11 mol) of sodium nitrite to 15 ml of water. The mixture will warm up to about 85 °C within 2-3 minutes and then cool down, changing color from deep red to pale brown.
- 4. Continue stirring for 15 minutes, during which the temperature will drop to 35-40 °C. Chill the mixture thoroughly in an ice-water bath for 30 minutes.
- 5. Collect the pale brown solid product by vacuum filtration and wash the product with three 30 ml of ice-cold water.

### **Recrystallization:**

- 1. Dissolve the solid in approximately 130 ml of boiling water and add decolorizing charcoal to the solution.
- 2. Filter the mixture and allow the filtrate to cool to about 50 °C.
- 3. Add a few crystals of the synthesized product (benzotriazole) retained for seeding.
- 4. Let the mixture slowly reach room temperature to avoid separating the material as an oil. Thoroughly chill the mixture in ice.
- 5. Collect the benzotriazole, which separates as pale straw-colored needles with a melting point of 99-100 °C.

# Calculation

Here, the limiting reagent is o-phenylenediamine; yield should be calculated from the amount taken.

 $C_6H_8N_2$  = Molecular formula of o-phenylenediamine

 $C_6H_5N_3$  = Molecular formula of benzotriazole

Molecular weight of o-phenylenediamine = 108 g/mole

Molecular weight of benzotriazole = 119 g/mole

# Theoretical yield:

108 g o-phenylenediamine forms 119 g benzotriazole

Therefore, 10.8 g o-phenylenediamine will form .....? (X) g benzotriazole

 $X = (119 \times 10.8) / 108 = 11.9 \ g$ 

**Theoretical yield** = 11.9 g

**Practical yield** = ——-- g

% Yield = (Practical Yield)/(Theoretical Yield)  $\times$  100

### **Result:**

Benzotriazole was synthesized, and the percentage yield was found to be.....%.

onarmacareerinsider.com