

## **EXPERIMENT**

### **Objective:**

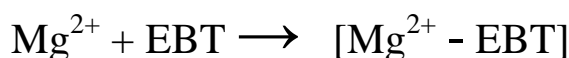
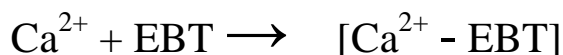
To determine the hardness of given water sample by complexometric titration using EDTA as an intermediate and EBT as an indicator.

### **Apparatus and Chemical required:**

Burette, Pipette, Conical flask, EDTA, water sample, EBT, Buffer solution (pH= 10).

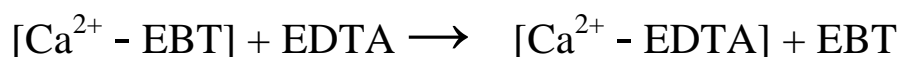
### **Theory:**

Water generally contains  $\text{Ca}^{2+}$  and  $\text{Mg}^{2+}$  salts which are responsible for the hardness of water. When we add EBT to the water sample,  $\text{Ca}^{2+}$  and  $\text{Mg}^{2+}$  salts react with EBT at pH 10 to form an unstable complex which is wine red in color.



Unstable wine- red complexes

Now, when we add EDTA solution to this, the unstable complex converted into stable complex and EBT is liberated which is sky- blue in color.



Stable Complex      Sky blue color

- EDTA (Ethylene diamine tetra acetic acid)
- EBT (Erichrome Black-T)

### **Procedure:**

Pipette out 10 ml of known water sample in a conical flask and 2 ml of buffer solution (pH 10) + 1 drop of EBT → wine red color appears → Titrate with EDTA solution until the color changes to sky- blue color. Perform the above procedure for 5times.

Repeat the same procedure 5 times again by taking unknown water sample (tab water)

### **Observation:**

S.No.	Volume of known water sample (ml)	Volume of EDTA solution used ( $V_1$ ) ml	Volume of unknown water sample (ml)	Volume of EDTA solution used ( $V_2$ ) ml
1.	10		10	
2.	10		10	
3.	10		10	
4.	10		10	
5.	10		10	

### **Calculation:**

Hardness of known water sample is given = 800ppm

So, hardness of unknown water sample is =  $800 \times V_2/V_1 \times 1000$  ppm

### **Precautions:**

- (1) Burette should be vertical throughout the experiment.
- (2) The reaction mixture should continuously be shaken during titration.
- (3) Glass ware should be washed and dried before doing the experiment.

# EXPERIMENT

## Objective:

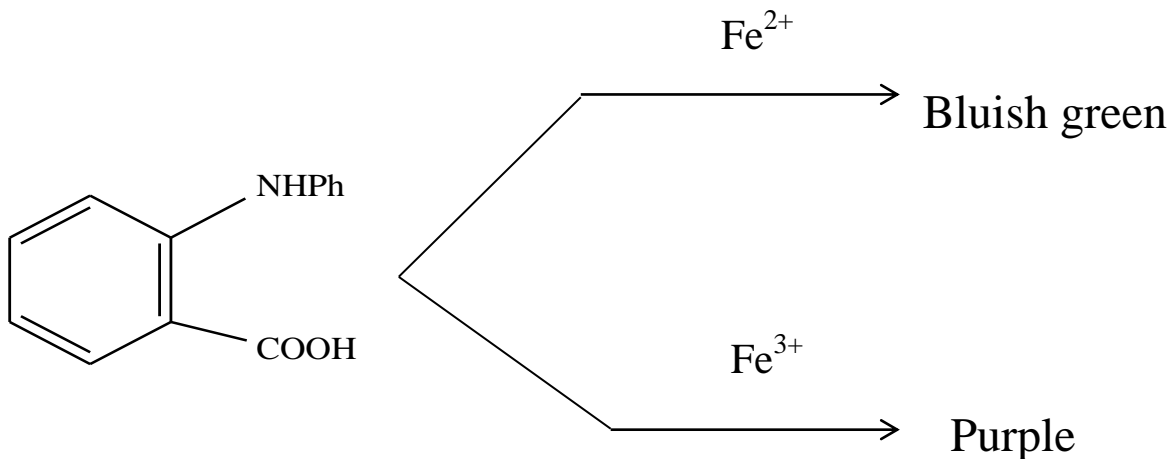
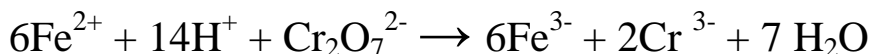
To determine the iron content of a given ferrous ammonium sulphate solution by titrating it against N/50  $K_2Cr_2O_7$  solution using N- phenylanthranilic acid as an internal indicator.

## Apparatus and Chemical required:

Solution of ferrous ammonium sulphate (FAS) or Mohr's salt,  $K_2Cr_2O_7$  solution, N- phenylanthranilic acid, distilled water, burette, Pipette, conical flask, diluted sulphuric acid.

## Theory:

$K_2Cr_2O_7$  acts a strong oxidizing agent in presence of dil.  $H_2SO_4$ .



## Procedure:

Pipette out 10 ml FAS + 2ml of dil.  $H_2SO_4$  + 1 drop of N- phenylanthranilic acid → Bluish green color → titrate it against  $K_2Cr_2O_7$  from burette until the purple color just appears → this will be end point → repeat the same for 5 times.

### **Observation:**

S.No.	Volume of $K_2Cr_2O_7$ used V(ml)
1.	
2.	
3.	
4.	
5.	

### **Calculation:**

Volume of FAS taken = 10 ml.

Normality of potassium dichromate taken = 1/50 N

Volume of potassium dichromate used = V ml.

$$N_{FAS} \times 10 = N/50 \times V$$

Strength of FAS (S) =  $N_{FAS} \times 392.16$  gm/lit.

Iron content =  $S \times 56/392.16$  gm.

### **Result:**

The strength of FAS is = ..... gm/lit.

The iron content is = ..... gm.

### **Precautions:**

- (1) Burette should be vertical throughout the experiment.
- (2) The reaction mixture should continuously be shaken during titration.
- (3) Glass ware should be washed and dried before doing the experiment.