### **CLIL - Argentometric Titrations**

Argentometric Titrations are titration of precipitate compounds. Precipitation titrations are based on reactions that yield ionic compounds of limited solubility. The most widely used and important precipitating reagent is silver nitrate (**AgNO**<sub>3</sub>), which is used for the determination of the halogens anions (chlorides **Cl**<sup>-</sup>, bromides **Br**<sup>-</sup>, iodides **I**<sup>-</sup>), thiocyanates (**SCN**<sup>-</sup>), cyanides (**CN**<sup>-</sup>)etc.

#### 1. Mohr Method

In the Mohr method, potassium chromate ( $\mathbf{K}_{2}\mathbf{CrO}_{4}$ ) serves as the indicator for the argentometric titration of chloride, bromide, and cyanide ions. Silver ions react with chromate to form the brick-red silver chromate ( $\mathbf{Ag}_{2}\mathbf{CrO}_{4}$ ) precipitate in the equivalence-point region. The Mohr method should be carried out at pH 7÷10. At low pH the concentration of the chromate ion in the acidic solutions is so low that it can not form a precipitate with  $\mathbf{Ag}^{+}$  at the equivalence point. Because in acidic solutions  $\mathrm{CrO}_{4}^{2-}$  is converted to dichromates  $\mathrm{Cr}_{2}\mathrm{O}_{7}^{2-}$ . Chromate ion is conjugated base of chromic acid. At high pH the silver ions can be precipitated as AgO. Titration Reaction:

 $Ag^+ + Cl^- \rightarrow AgCl_{(s)}$ 

Silver chloride (AgCl) is a white caseous precipitate (similar to ricotta cheese). at the equivalent point the indicator turns brick-red, with the following reaction:

$$2Ag^{+} + CrO_{4^{2-}} \rightarrow Ag_2CrO_{4(s)}$$

#### 2.Volhard Method

In this method, Fe (III) ion is used as an indicator. The most important application of the Volhard method is the indirect determination of halide ions. First, a measured excess of standard silver nitrate solution is added to the sample:

$$Ag^+ + Cl^- \rightarrow AgCl_{(s)}$$

The excess silver is determined by back-titration with a standard thiocyanate solution:

$$Ag^+ + SCN^- \rightarrow AgSCN_{(s)}$$

Back-Titration Reaction (excess  $AgNO_3$ ) white Iron(III) serves as the indicator. The solution turns red with the first slight excess of thiocyanate ion due to the formation of the complex  $[Fe(SCN)]^{2+}$ :

$$Fe^{3+} + SCN^{-} \rightarrow [Fe(SCN)]^{2+}$$

The strongly acidic environment of the Volhard titration is a distinct advantage over other titrations of halide ions because such ions as carbonate, oxalate, and arsenate do not interfere. The silver salts of these ions are soluble in acidic media but only slightly soluble in neutral media. Silver chloride is more soluble than silver thiocyanate. As a result, in chloride determinations using the Volhard method, the reaction

$$AgCl(s) + SCN^{-} \rightarrow AgSCN(s) + Cl^{-}$$

occurs to a significant extent near the end of the back-titration. This reaction causes the endpoint to fade and results in overconsumption of thiocyanate ion. The resulting low results for chloride can be overcome by filtering the silver chloride before undertaking the back-titration. Filtration is not required for other halides because they form silver salts that are less soluble than silver thiocyanate.



### 3. Fajans Method

The Fajans method uses an adsorption indicator, an organic compound that adsorbs onto or desorbs from the surface of the solid in a precipitation titration. Ideally, the adsorption or desorption occurs near the equivalence point and results not only in a color change but also in the transfer of color from the solution to the solid or vice versa. Fluorescein is a typical adsorption indicator useful for the titration of chloride ion with silver nitrate. In the aqueous solution, fluorescein is partially ionized to hydronium ions and yellowish green colored negatively charged fluoresceinate ions. The fluoresceinate ion forms a dark red silver salt. Before the equivalence point, **Cl**<sup>-</sup> is in excess, and the primary adsorbed layer is **Cl**<sup>-</sup>. This repels the indicator anion, and the more loosely held secondary (counter) layer of adsorbed ions is cations, such as **Na**<sup>+</sup>. Beyond the equivalence point, **Ag**<sup>+</sup> is in excess, and the surface of the precipitate becomes positively charged, with the primary layer being **Ag**<sup>+</sup>. This will now attract the indicator anion and adsorb it in the counterlayer. Titrations involving adsorption indicators are rapid, accurate, and reliable, but their application is limited to the few precipitation titrations that form colloidal precipitates rapidly.

Adapted from: acikders.ankara.edu.tr

# TEACHING AIMS:

- Understanding the meaning of argentometric titration;
- Understanding the meaning of titrant;
- Understanding the meaning of adsorption indicator;
- Understanding the differences between the titrations and back titrations;
- Understanding the different method of argentometric titrations;

## EXERCISE

#### 1 Read the text and complete the scheme below

