Comparative Degradation Kinetic Studies of Tetrabutylammonium Tribromide (TBATB) and Cetyltrimethylammonium Tribromide (CTMATB) - A Route to Reactivity Assessment for Solvent-Free Reactions

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Abstract: Tetrabutylammonium tribromide (TBATB) and cetyltrimethylammonium tribromide (CTMATB) are important alternatives to the hazardous reagent bromine (Br2) because of their efficiency in various chemical transformations, especially organic bromination reactions. In the context of green chemistry, development of solvent-free reaction protocols is much needed because of the hazards posed by organic solvents. In order to assess the comparative behavior of these tribromides in solvent-free reactions at elevated temperatures, thermogravimetric studies came in as a useful tool. Thus, the kinetics of the thermal degradation of TBATB and CMTATB were investigated through thermogravimetric analysis and the thermal stability order of the two bromides based on TG followed the order: CTMATB > TBATB. Non isothermal multiple heating rate model free methods Ozawa-Flynn-Wall and Kissinger methods are used to calculate degradation activation energy of the tribromides. The reaction mechanism based on activation energy calculation at different conversions has been proposed.

Keywords: Tetrabutylammonium tribromide, Cetyltrimethylammonium tribromide, Thermal analysis, Brominations, Solvent-free, Environmentally benign

Introduction

Acknowledging the importance of bromo-organic compounds1-2 and the need for environmentally benign brominating reagents, substantial developments have been made through the synthesis and use of quaternary ammonium tribromides (QATBs) as alternatives of bromine3-10. Among the numerous tribromides that have been synthesized so far, tetrabutylammonium tribromide (TBATB) and cetyltrimethylammonium tribromide (CTMATB) have been found to be useful for a number of different types of reactions, including solvent-free brominations conducted at elevated temperature and under microwave
conditions. Ability for any reagent to perform at high temperature greatly broadens the scope of the reagent for other solvent-free reactions and thereby it becomes important to understand their degradation pattern at elevated temperature. It has been observed that in the case of QATBs, other than a brief study on CTMATB there seems to be no report on their thermal analysis, especially in terms of a comparative assessment. Acknowledging the importance of such kind of a study, cetyltrimethylammonium tribromide (CTMATB) and tetrabutylammonium tribromide (TBATB) have been taken as candidates for estimating the kinetic parameters of degradation processes such as activation energies (Ea) and the Arrhenius pre-exponential factor (A). Accordingly, this paper deals with the comparative thermal and kinetic study of TBATB and CTMATB using simultaneous thermal analysis by the Kissinger and Flynn-Wall-Ozawa methods. The Kissinger method derives from the Arrhenius Equation, which correlates the rate constant with the activation energy, through an equation:

\[
K = A \exp\left[-\frac{E_a}{RT}\right]
\]  

Where, \(k\) is the rate constant; \(A\) is the pre-exponential factor also known as frequency factor; \(E_a\) is the activation energy; \(R\) is gas constant and \(T\) is the temperature. The Kissinger method can be used in calculating activation energy at different heating rate (\(\beta\)) which is given by the equation (2) as:

\[
\ln\left(\frac{\beta}{T_{\text{max}}^2}\right) = \ln A - \frac{E_a}{R T_{\text{max}}} + \ln \left(n \left[1 - \alpha_{\text{max}}\right]^n \right) - \frac{E_a}{R T_{\text{max}}}
\]  

Where \(\beta\) is heating rate, \(T_{\text{max}}\) is temperature related to maximum reaction rate, \(A\) is the pre-exponential factor, \(\alpha_{\text{max}}\) is the maximum degradation fraction, \(n\) is reaction order. Plotting \(\ln(\beta/T_{\text{max}}^2)\) versus \(1/T_{\text{max}}\) gives activation energy from slope.

The Flynn-Wall-Ozawa method is an integral method that can be used to determine the activation energy without the knowledge of reaction mechanism, pre-exponential factor (A) and activation energy (Ea) do not depend on degradation fraction, but they depend on the temperature. This method uses equation (3) as:

\[
\log g(\alpha) = \log \left(\frac{AE_a}{R}\right) - \log \beta + \log P \left(\frac{E_a}{RT}\right)
\]  

Doyle approximation is used and equation (3) can be obtained.

Where \(\beta\) is the heating rate and \(g(\alpha)\) is the conversion temperature. The plot of \(\log \beta\) versus 1000/T should be linear with the slope \(E_a/R\) from which \(E_a\) can be calculated.

\[
\log \beta = \log \left(\frac{AE_a}{R}\right) - \log \left(\alpha\right) - 2.314 - 0.4567 \left(\frac{E_a}{RT}\right)
\]  

**Experimental**

All reagents of highest purity were purchased from commercial sources and used without further purification. Melting points were determined in open capillaries and are uncorrected.

**Synthesis of tetrabutylammonium tribromide (TBATB)**

0.057 g (0.53 mmol) sodium carbonate Na2CO3 was added to 50% H2O2, (10 mL; 47 mmol) and the mixture was stirred at room temperature for about 5 minutes until Na2CO3 completely dissolved and the solution attained a clear colourless solution. To this a solution of 5 g (15.53 mmol) tetrabutyl ammonium bromide (TBAB) and 3.7 g (31.90 mmol) potassium bromide were added and the mixture was heated at 110°C until the color of the solution changed to yellow-greenish. The reaction mixture was then cooled to room temperature and poured into 50 mL of ice water, and the precipitate was filtered off and washed with cold water. The product was dried under vacuum to yield 0.25 g of the desired product (95% yield).
(KBr), both dissolved in 50 mL of water was added. To the resultant reaction mixture, 50 mL of 2 M H₂SO₄ was added in small portions, upon which the precipitate appeared immediately. The mixture was allowed to be stirred for further 1 h whereby a bright yellow coloured compound precipitated out completely. The compound was filtered under suction using Whatman-40 filter paper and dried in vacuum desiccators using self-indicating coarse silica gel. It was further recrystallized in acetonitrile, isolated by suction filtration and dried in vacuo yielding 7.38 g (98%) of the pure compound.

**Synthesis of cetyltrimethylammonium tribromide (CTMATB)**

0.057 g (0.53 mmol) sodium carbonate Na₂CO₃ was added to 50% H₂O₂ (10 mL; 47 mmol) and the mixture was stirred at room temperature for about 5 minutes until Na₂CO₃ completely dissolved and the solution attained a clear colourless solution. To this a solution of 5 g (15.53 mmol) cetyltrimethylammonium bromide (CTMAB) and 3.7 g (31.90 mmol) potassium bromide (KBr), both dissolved in 50 ml of water was added. To the resultant reaction mixture, 50 mL of 2 M H₂SO₄ was added in small portions, upon which the precipitate appeared immediately. The mixture was allowed to be stirred for further 1 h whereby a bright yellow coloured compound precipitated out completely. The compound was filtered under suction using Whatman-40 filter paper and dried in vacuum desiccators using self-indicating coarse silica gel. It was further recrystallized in acetonitrile, isolated by suction filtration and dried in vacuo. The yield of the isolated compound was 7.12 g (99%).

**Thermogravimetric analysis**

Thermogravimetric (TG) experiments were conducted on a Perkin Elmer STA-6000 (Simultaneous Thermal Analyser) instrument. Samples weighing about 12 mg were taken in the crucible and heated from 30 to 300 °C at a heating rate of 5, 10, 15 and 20 °C/min in nitrogen atmosphere.

**Data processing and activation energy calculation**

TG curves were analyzed by using Pyris software from TG Analyzer and data was used in MS Excel software to calculate activation energy (Ea) by the method of least square. Activation energy of samples was calculated by “model free” iso-conversional methods. Kissinger method was used in calculating activation energy at different heating rate (β) which is given by the equation (2):

\[
\ln \left( \frac{\beta}{T_{\text{max}}^2} \right) = \ln \frac{A R}{E_a} + \ln \left( n(1-\alpha_{\text{max}})^n-1 \right) - \left( \frac{E_a}{R T_{\text{max}}} \right)
\]

Where \( \beta \) is heating rate, \( T_{\text{max}} \) is temperature related to maximum reaction rate, \( A \) is pre-exponential factor, \( \alpha_{\text{max}} \) is maximum degradation fraction, \( n \) is reaction order. Plotting \( \ln(\beta/T_{\text{max}}^2) \) versus \( (1000/T_{\text{max}}) \) gives activation energy from slope. Flynn-Wall-Ozawa method was also used to determined activation energy at a fixed conversion with the slope of such a line being \(-0.4567 E_a/RT\)

**Results and Discussion**

**Comparative TG and DTG studies**

TG curves for TBATB and CTMATB at different heating rates in nitrogen atmosphere are shown in Figures 1-2. The shapes of TG curves are quite similar, shifting towards higher temperatures at higher heating rates. This is probably due to the heat transfer lag with increased heating rate. The initial decomposition temperature of TBATB and CTMATB occurs around 174 °C and 176 °C respectively. Although the values differ slightly the thermal stability of the samples follows the order: CTMATB>TBATB as indicated by Figure 3.
The decomposition of TBATB and CTMATB occur in single step as confirmed by DTG curves (Figure 4). The main decomposition of TBATB occurs around 230 °C with a percentage mass of 3% remaining as unburned whereas in the case of CTMATB it occurs around 231 °C. The higher decomposition value again reiterates the higher stability of cetyltrimethylammonium tribromide.

**Figure 1.** TG curve of TBATB at 5, 10, 15 and 20 °C

**Figure 2.** TG curve of CTMATB at 5, 10, 15 and 20 °C

**Figure 3.** TG curve of TBATB and CIMATB at 10 °C/min

**Figure 4.** DTG curves of TBATB and CIMATB at 10 °C/min

**Comparison of degradation activation energies of the two bromides**

Activation energies for TBATB and CTMATB were calculated using Kissing method at varying heating rates (β) and are given in Table 1 and 2. The linear plots of Kissinger method for TBATB and CTMATB are given in Figure 5. Activation energies for TBATB and CTMATB were calculated using O-F-W procedure at varying degree of conversions and are given in Table 3. The iso-conversational plot of O-F-W show a general trend of activation energy. For illustration iso-conversational plot of O-F-W for TBATB and CTMATB are shown in Figure 6 and 7. From the figure we observed that the fitted lines for the two bromides are almost parallel which indicates the activation energy at different conversions and consequently, implying the possibility of single reaction mechanism.
Table 1. Calculation of activation energies for decomposition of TBATB at varying degree of conversions

<table>
<thead>
<tr>
<th>β</th>
<th>ln β</th>
<th>T_{max}</th>
<th>T_{max} K</th>
<th>T_{max} sq</th>
<th>β/T_{max} sq</th>
<th>lnβ/T_{max} sq</th>
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Table 2. Calculation of activation energies for decomposition of CTMATB at varying degree of conversions

<table>
<thead>
<tr>
<th>β</th>
<th>ln β</th>
<th>T_{max}</th>
<th>T_{max} K</th>
<th>T_{max} sq</th>
<th>β/T_{max} sq</th>
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Figure 5. Plot of Kissinger method for TBATB and CTMATB at varying degree of conversions

Figure 6. Iso-conversional plot of O-F-W method for TBATB at varying degree of conversions

Figure 7. Iso-conversional plot of O-F-W method for CTMATB at varying degree of conversions
It can be seen from the Table 3 the activation energies calculated by iso-conversional method shows that the degree of conversion i.e. at 0.33 the Ea values for CTMATB was greater than that of TBATB. This means that the initial thermal stability of CTMATB was comparatively higher than TBATB which also matches well with the initial thermal degradation order observed for TG curves. The values calculated by Kissinger method for TBATB and CTMATB are 0.0936 kJmol\(^{-1}\) and 0.0947 kJmol\(^{-1}\) respectively. The order of E\(_a\) values calculated by Kissinger method was same as evaluated by initial thermal degradation order i.e. CTMATB>TBATB.

Table 3. Activation energies for TBATB and CTMATB calculated by O-F-W method at varying degree of conversions

<table>
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<tr>
<th>(\alpha)</th>
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<th>CTMATB</th>
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<td>0.33</td>
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Acknowledgement

The author (NK) is grateful to Bendang STA, Nagaland University, Lumami, for recording the TGA data.

Conclusion

TBATB and CTMATB have been synthesized by environmental friendly method. The thermal stability of the synthesized reagent has then been studied in order to understand the basic theory and kinetics of thermal degradation of TBATB and CTMATB.

- On the basis of initial thermal decomposition temperature calculated from TG curves, the thermal stability of samples follows the order CTMATB>TBATB.
- The activation energy values calculated for the tribromides also shows that CTMATB is more stable than TBATB.
- The thermal and kinetic data presented in this study would help to find out new applications of these tribromides either as binary or ternary blends.

References