The Decomposition of Hydrogen Peroxide

The decomposition of hydrogen peroxide in aqueous solution proceeds very slowly. A bottle of 3% hydrogen peroxide sitting on a grocery store shelf is stable for a long period of time. The decomposition takes place according to the reaction below.

\[ 2 \text{H}_2\text{O}_2(\text{aq}) \rightarrow 2 \text{H}_2\text{O} + \text{O}_2(\text{g}) \]

A number of catalysts can be used to speed up this reaction, including potassium iodide, manganese (IV) oxide, and the enzyme catalase. If you conduct the catalyzed decomposition of hydrogen peroxide in a closed vessel, you will be able to determine the reaction rate as a function of the pressure increase in the vessel that is caused by the production of oxygen gas. If you vary the initial molar concentration of the H\text{2}O\text{2} solution, the rate law for the reaction can also be determined. Finally, by conducting the reaction at different temperatures, the activation energy, \( E_a \), can be calculated.

**OBJECTIVES**

In this experiment, you will

- Conduct the catalyzed decomposition of hydrogen peroxide under various conditions.
- Calculate the rate constant for the reaction.
- Determine the rate law expression for the reaction.
- Calculate the activation energy for the reaction.

![Figure 1](image)

**MATERIALS**

- Vernier computer interface
- computer
- Vernier Gas Pressure Sensor
- thermometer or Temperature Probe
- one-hole rubber stopper with stem tubing with two Luer-lock connectors
- solid rubber stopper (#1)
- distilled water
- 3% hydrogen peroxide, H\text{2}O\text{2}, solution
- 0.5 M potassium iodide, KI, solution
- 18 × 150 mm test tube
- two 10 mL graduated cylinders
- graduated plastic Beral pipet
- 1 liter beaker
- ~800 mL room temperature water
PRE-LAB EXERCISE

The hydrogen peroxide solution that you are using in this experiment is labeled as a 3% solution, mass/volume. However, in order to complete the calculations, the concentration must be in molarity. Calculate the molarity of a 3% mass/volume H₂O₂ solution (Part I, II, and IV) and a 1.5% mass/volume H₂O₂ solution (Part III) and record these values in the table below.

<table>
<thead>
<tr>
<th>Part</th>
<th>Volume H₂O₂ (mL)</th>
<th>[H₂O₂] before mixing</th>
<th>Volume KI (mL)</th>
<th>[KI] before mixing</th>
</tr>
</thead>
<tbody>
<tr>
<td>I</td>
<td>4</td>
<td></td>
<td>1</td>
<td>0.50 M</td>
</tr>
<tr>
<td>II</td>
<td>4</td>
<td></td>
<td>1</td>
<td>0.25 M</td>
</tr>
<tr>
<td>III</td>
<td>4</td>
<td></td>
<td>1</td>
<td>0.50 M</td>
</tr>
<tr>
<td>IV</td>
<td>4</td>
<td></td>
<td>1</td>
<td>0.50 M</td>
</tr>
</tbody>
</table>

PROCEDURE

Part I  Decompose 3% H₂O₂ solution with 0.5 M KI solution at ~20°C.

1. Obtain and wear goggles.

2. Prepare the reagents for temperature equilibration.
   a. Obtain room temperature water to set up a water bath to completely immerse the test tube. Use a thermometer or a Temperature Probe to measure the temperature of the bath. Record this temperature in your data table for Parts I-III; presume that the water bath temperature remains constant throughout.
   b. Measure out 4 mL of 3% H₂O₂ solution into the test tube. Seal the test tube with the solid rubber stopper and place the test tube in the water bath.
   c. Measure out 2 mL of 0.5 M KI solution in a graduated cylinder. Draw about 1 mL of the KI solution into a graduated Beral pipet. Invert the pipet and immerse the reservoir end of the pipet in the water bath.

3. Connect a Gas Pressure Sensor to Channel 1 of the Vernier computer interface. Connect the interface to the computer using the proper cable.

4. Use the plastic tubing to connect the one-hole rubber stopper to the Gas Pressure Sensor, as shown in Figure 1. About one-half turn of the fittings will secure the tubing tightly.

5. Start the Logger Pro program on your computer. Open the file “12 Peroxide” from the Advanced Chemistry with Vernier folder.

6. Prepare to run the reaction and collect pressure data.
   a. Remove the test tube from the water bath and remove the solid stopper.
   b. Remove the plastic Beral pipet from the water bath and quickly transfer 1 mL of KI solution into the test tube. Tap or lightly shake the test tube to mix the reagents.
   c. Seal the test tube with the one-hole stopper connected to the Gas Pressure Sensor.
   d. Place the test tube back in the water bath.

7. Click [Collect] to begin data collection. Data will be gathered for three minutes. If necessary, gently hold the test tube so that it stays completely immersed in the water bath.
8. When the data collection is complete, carefully remove the stopper from the test tube to relieve the pressure. Dispose of the contents of the test tube as directed.

9. Examine the graph of Part I. Select a linear region just beyond the initial flat portion of the graph that covers one minute of the reaction. Click the Linear Regression button, $\text{Linear Regression}$, to calculate the best-fit line equation. Record the slope as the initial rate of the reaction in your data table. Store the results from the first trial by choosing Store Latest Run from the Experiment menu.

10. Rinse and clean the test tube for the second trial.

**Part II Decompose 3% $\text{H}_2\text{O}_2$ solution with 0.25 M KI solution at ~20°C**

11. Measure out 4 mL of 3% $\text{H}_2\text{O}_2$ solution into the test tube. Seal the test tube with the solid rubber stopper and place the test tube in the water bath.

12. Add 1 mL of distilled water to the remaining 1 mL of KI solution in the graduated cylinder. Swirl the mixture gently to mix the solution.

13. Draw 1 mL of the KI solution into a plastic Beral pipet. Invert the pipet and immerse the reservoir end of the pipet in the water bath. Allow both the test tube and the Beral pipet to remain in the water bath for at least two minutes before proceeding.

14. Repeat Steps 6–10 to complete Part II. Remember to store the data.

**Part III Decompose 1.5% $\text{H}_2\text{O}_2$ solution with 0.5 M KI solution at ~20°C**

15. Prepare a 1.5% $\text{H}_2\text{O}_2$ solution by mixing 2 mL of distilled water with 2 mL of 3% $\text{H}_2\text{O}_2$ solution. Transfer the resulting 4 mL of the 1.5% $\text{H}_2\text{O}_2$ solution to the test tube, seal the test tube with the solid stopper, and place the test tube in the water bath.

16. Rinse and clean the graduated cylinder that you have used for the KI solution. Add a fresh 2 mL of 0.5 M KI solution to the graduated cylinder.

17. Draw 1 mL of the KI solution into a plastic Beral pipet. Invert the pipet and immerse the reservoir end of the pipet in the water bath. Allow both the test tube and the Beral pipet to remain in the water bath for at least two minutes before proceeding.

18. Repeat Steps 6–10 to complete Part III, and store the data.

**Part IV Decompose 3% $\text{H}_2\text{O}_2$ solution with 0.5 M KI solution at ~30°C**

19. Conduct Part IV identically to the procedure in Part I, with one exception: set the water bath at 30°C.
### DATA TABLE

<table>
<thead>
<tr>
<th>Part</th>
<th>Reactants</th>
<th>Temperature (°C)</th>
<th>Initial rate (kPa/s)</th>
</tr>
</thead>
<tbody>
<tr>
<td>I</td>
<td>4 mL 3.0% H₂O₂ + 1 mL 0.5 M KI</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>II</td>
<td>4 mL 3.0% H₂O₂ + 1 mL 0.25 M KI</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>III</td>
<td>4 mL 1.5% H₂O₂ + 1 mL 0.5 M KI</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>IV</td>
<td>4 mL 3.0% H₂O₂ + 1 mL 0.5 M KI</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

### DATA ANALYSIS

<table>
<thead>
<tr>
<th>Part</th>
<th>Initial rate (mol/L-s)</th>
<th>[H₂O₂] after mixing</th>
<th>[I⁻] after mixing</th>
<th>Rate constant k</th>
</tr>
</thead>
<tbody>
<tr>
<td>I</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>II</td>
<td>-</td>
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<tr>
<td>III</td>
<td>-</td>
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<td>-</td>
</tr>
<tr>
<td>IV</td>
<td>-</td>
<td>-</td>
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</tr>
</tbody>
</table>

1. Calculate the rate constant, \( k \), and write the rate law expression for the catalyzed decomposition of hydrogen peroxide. Explain how you determined the order of the reaction in H₂O₂ and KI.

2. The following mechanism has been proposed for this reaction:

\[
\begin{align*}
H₂O₂ + I⁻ & \rightarrow IO⁻ H₂O \\
H₂O₂ + IO⁻ & \rightarrow I⁻ + H₂O + O₂
\end{align*}
\]

If this mechanism is correct, which step must be the rate-determining step? Explain.

3. Use the Arrhenius equation (shown below) to determine the activation energy, \( E_a \), for this reaction.

\[
\ln \frac{k_1}{k_2} = \frac{E_a}{R} \left( \frac{1}{T_2} - \frac{1}{T_1} \right)
\]
Vernier Lab Safety Instructions Disclaimer

THIS IS AN EVALUATION COPY OF THE VERNIER STUDENT LAB.

This copy does not include:

- Safety information
- Essential instructor background information
- Directions for preparing solutions
- Important tips for successfully doing these labs