45th National Chemistry Olympiad

Maastricht University

PRACTICAL TEST

Assignment booklet

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Directions/resources

- This practical test consists of two parts:
 - The determination of the amount of crystal water (x) in a mixture of $Na_2CO_3.xH_2O$ and $NaHCO_3$.
 - Kinetic study of the decomposition of tris-(oxalato)-manganate(III) ions.
- The practical test ends after 4 hours. During this time:
 the attached answer sheets need to be completed;
 - $\circ~$ all questions must be answered.
- After the practical test, when you have handed in everything, the glassware still needs to be cleaned and tidied up.
- The maximum score for the practical test is 80 points.
- The score is determined by:
 - practical skills, working clean, safety

maximum 20 points

- \circ results of the determinations and answers to the questions maximum 60 points
- Required tools: (graphic) calculator, ruler/protractor and Binas or ScienceData.
- First read the introduction and all assignments before you start working.
- Write the answers to the questions in the boxes on the answer sheets provided. If you don't have enough space, you can ask for extra paper.

Additional:

- This is a test; it is not permitted to consult with other participants.
- If you have a question, you can ask the supervisor.
- If something is wrong with your glassware or equipment, report it to the supervisor as soon as you discover it. Don't borrow someone else's things!

Experiment 1 The determination of the amount of crystal water (x) in a mixture of $Na_2CO_3.xH_2O$ and $NaHCO_3$

40 points

Introduction

Sodium carbonate is mined mostly from a mineral named Trona. It is a mixture of hydrated sodium carbonate ($Na_2CO_3.xH_2O$) and sodium hydrogen carbonate ($NaHCO_3$).



The goal of this task is to determine the amount of crystal water bound to sodium carbonate in a solid mixture containing hydrated sodium carbonate ($Na_2CO_3.xH_2O$) and sodium hydrogen carbonate ($NaHCO_3$).

In this determination two separate titrations with a solution of hydrochloric acid are performed:

- a titration with methyl orange as the indicator
- a titration with phenolphthalein as the indicator.

Each titration is executed in duplicate. In advance, you have to execute a test titration.

Chemicals

- a weighed mixture of Na₂CO₃.xH₂O and NaHCO₃
- 0.1000 M hydrochloric acid
- a solution of methyl orange
- a solution of phenolphftalein
- demineralized water

Safety

- wear safety glasses
- if your skin comes into contact with any of the chemicals, rinse it of immediately

Equipment and glassware

- 250 mL volumetric flask
- funnel for the volumetric flask
- 50 mL burette
- funnel for the burette
- four beakers
- four small plastic pipettes
- pipette-balloon
- 25 mL pipette
- 50 mL Erlenmeyer flask for the titrations
- marker
- magnetic stirrer
- magnetic stir bar

Procedure

- Prepare a solution of your solid sample in the 250 mL volumetric flask.
- **D**2 Fill the burette with the 0.1000 M hydrochloric acid.
 - Note: Probably you have to refill the burette a few times during the determination.
- B With each of the titrations below, first perform a test titration. Afterwards perform each titration in duplicate.

Titration in presence of phenolphtalein

- ^{D4} Bring 25.00 mL of the solution that is in the volumetric flask, into the 50 mL Erlenmeyer flask.
- **D**5 Add three drops of the phenolphftalein solution.
- Put the magnetic stir bar in the solution and start stirring (don't heat).
- ^{D7} Titrate until the pink color disappears totally.
- **B** Repeat the points 4 to 7.

Titration in presence of methyl orange

- Bring 25.00 mL of the solution that is in the volumetric flask, into the 50 mL Erlenmeyer flask.
- ID Add three drops of the methyl orange solution.
- I1 Titrate until the color becomes red.
- □12 Repeat the points 9 to 11.

Questions - write the answers on the answer sheets

- 1 Write down:
 - the mass of the sample
 - the molarity of the hydrochloric acid
 - all burette readings
- 2 Give the equations of the reactions that occur during the titrations.
- Calculate the amount of mmoles of Na₂CO₃.xH₂O and the amount of mmoles of NaHCO₃ in your sample.
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- 4 Calculate the value of x in $Na_2CO_3.xH_2O$.
- It is also possible to perform the determination by using a precipitation reaction (and weighing the precipitate) instead of one of the titrations.
 Explain which solution could be used and which titration does not have to be executed in that case.

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Experiment 2 Kinetic study of the decomposition of tris-(oxalato)manganate(III) ion 40

40 punten

Introduction

When solutions of manganese(II) sulfate, potassium permanganate and oxalic acid are poured together, so called tris-(oxalato)-manganate(III) ions, $\{Mn(C_2O_4)_3\}^{3-}$ are formed according the reaction equation below:

 $MnO_{4}^{-} \ + \ 4 \ Mn^{2+} \ + \ 15 \ H_2C_2O_4 \ \rightarrow \ 5 \ \{Mn(C_2O_4)_3\}^{3-} \ + \ 22 \ H^+ \ + \ 4 \ H_2O \ \ \ reaction \ 1$

As soon as these ions are formed, they begin to decompose and Mn^{2+} , $C_2O_4^{2-}$ and CO_2 are formed:

$$2 \{Mn(C_2O_4)_3\}^{3-} \rightarrow 2 Mn^{2+} + 5 C_2O_4^{2-} + 2 CO_2$$
 reaction 2

The rate equation of reaction 2 can be given as:

$$s = -\frac{d[\{Mn(C_2O_4)_3\}^{3^-}]}{dt} = k[\{Mn(C_2O_4)_3\}^{3^-}]^n$$

The goal of this experiment is:

- to show that reaction 2 is a first order reaction;
- to determine the value of the rate constant *k*.

Because the ions $\{Mn(C_2O_4)_3\}^{3-}$ give the solution a light brown color ($\lambda_{max} = 440$ nm) and the reaction products are colorless, the course of the reaction can be followed spectrophotometrically by measuring regularly the absorbance (extinction) A of the solution.

The average reaction rate in a time interval Δt can be calculated by using the expression below:

$$s = -\frac{\Delta A}{\Delta t \times \varepsilon \times l} = k \left[\left\{ \mathsf{Mn}(\mathsf{C}_2\mathsf{O}_4)_3 \right\}^{3-} \right]^n$$

In this expression ΔA is the change in the absorbance (extinction) in the time interval Δt and is $[{Mn(C_2O_4)_3}^{3-}]$ the concentration of the complex at the beginning of that time interval.

Further applies that $\varepsilon = 70 \text{ Lmol}^{-1} \text{ cm}^{-1}$ and l = 1.00 cm.

The experiment is executed in two variants, with two sets of solutions that differ in concentration.

To execute the measurements with the spectrophotometer you can get help from the labassistent.

Chemicals

- solutions of $MnSO_4$: 0.20 M and 0.075 M
- solutions of $H_2C_2O_4$: 0.20 M and 0.075 M
- solutions of KMnO₄: 0.020 M and 0.0075 M

Safety

- wear safety glasses
- if your skin comes into contact with any of the chemicals, rinse it of immediately.

Materials

- a 2 mL pipette
- a 10 mL pipette
- a 25 mL pipette
- three plastic pipettes
- three UV cuvettes
- pipette-balloon
- two 50 mL Erlenmeyer flasks
- four beakers
- magnetic stirrer
- magnetic stir bar
- stopwatch

Procedure

Variant I

- ^a1 Introduce successively into the 50 mL Erlenmeyer flask: 2.0 mL of 0.20 M solution of MnSO₄ and 14.0 mL of 0.20 M solution of $H_2C_2O_4$.
- ² Put the magnetic stir bar into the solution and start stirring (don't heat).
- $^{\tt D}3$ Add 4.0 mL 0.020 M solution of KMnO4 and start time measurement.
- Immediately after starting time measurement, take a sample from the reaction vessel, by means of a plastic pipette, bring it into a cuvette and measure the absorbance (extinction).
- ^{D5} Write down, in the designated table of your answer sheet, the measured absorbance and the time of the measurement.
- Pour back the solution from the cuvette into the reaction vessel.
- After about 60 s, take another sample from the solution and measure the absorbance.
 Write down, in the table of your answer sheet, the absorbance and the time of the measurement.
- Pour back the solution from the cuvette into the reaction vessel.
- $\square 9$ Repeat the points 7 and 8 three times every ca. 60 s.

Variant II

^a10 Repeat the points 1 to 7 of the procedure of variant I, but now use 0.075 M solution of MnSO₄, 0.075 M solution of $H_2C_2O_4$ and 0.0075 M solution of KMnO₄.

Questions - write the answers on the answer sheets

- 6 Write down, in the designated tables of your answer sheet, all measured absorbances and the times of the measurements of the absorbances.
- 7 Calculate the reaction rate in the first time interval for both variants. Write down your calculation in the designated tables of your answer sheet.

The order of the reaction can be calculated by means of the formula below:

 $n = \frac{\log s_{||} - \log s_{|}}{\log[\{Mn(C_2O_4)_3\}^{3-}]_{||} - \log[\{Mn(C_2O_4)_3\}^{3-}]_{||}}$

Herein s_1 and s_{II} are the reaction rates in the first time interval of variant I and variant II, respectively. [{ $Mn(C_2O_4)_3$ }^{3-}]_1 and [{ $Mn(C_2O_4)_3$ }^{3-}]_II are the initial concentrations of the complex in the first time interval of both variants. These approximately equal the concentrations of the complex on t = 0: [{ $Mn(C_2O_4)_3$ }^{3-}]_1 = 0.020 mol L⁻¹ and [{ $Mn(C_2O_4)_3$ }^{3-}]_II = 0.0075 mol L⁻¹.

- 8 Calculate *n* using your measurements.
- 9 Does it follow from your measurements that reaction 2 is a first order reaction? Motivate your answer.
- 10 Calculate the reaction rates in the second, third and fourth time interval of variant I. Write down your calculations in the designated table of your answer sheet.
- 11 Calculate the average rate constant k that follows from this, assuming that reaction 2 is a first order reaction.

Write down your calculations in the designated table of your answer sheet.

12 Show, by means of a calculation, that the initial concentration of $\{Mn(C_2O_4)_3\}^{3-}$ in variant I equals 0.020 mol L⁻¹.

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