REACTIONS OF COPPER

Copper will undergo many types of reactions. In this experiment you will observe a sequence of copper reactions. The sequence begins with copper metal and ends with copper metal, so it is called a cycle of copper reactions. Observations will be made for each reaction.

Since no copper is added or removed between the initial and final reaction steps, copper can be quantitatively recovered. In other words to recover the same amount of copper metal that you started with, this will require careful work. The success of the copper metal recovery will be shown by calculating the percent yield. The diagram below shows (in an abbreviated form) the cycle of copper reactions.



The following are the completed but **unbalanced** equations. Each equation is numbered to match each step of the cycle:

I. $HNO_3(aq) + Cu(s) \rightarrow Cu(NO_3)_2(aq) + H_2O(I) + NO_2(g)$ (oxidation-

reduction reaction)

II $Cu(NO_3)_2(aq) + NaOH(aq) \rightarrow Cu(OH)_2(s) + NaNO_3(aq)$

III. $Cu(OH)_2(s) \rightarrow CuO(s) + H_2O(l)$

IV. $CuO(s) + H_2SO_4(aq) \rightarrow CuSO_4(aq) + H_2O(l)$

V.
$$CuSO_4(aq) + Al(s) \rightarrow Al_2(SO_4)_3(aq) + Cu(s)$$

It is useful to classify reactions into different types, because products of reactions can then be predicted.

No one classification scheme can accommodate all known reactions but you will classify the reactions in your experiment based on ideas of double exchange, combination, decomposition, and replacement.

Throughout the experiment careful observations and recordings must be made. Determine the colors of the solids as well as the color of ions in solution. In order to have a good percent yield, it is important to prevent loss by avoiding spattering while boiling, leaving product on sides of beakers, and spilling of product. Purify precipitates by washing efficiently and then drying completely before weighing.



- 1. Safety goggles <u>must</u> be worn at all times
- Concentrated nitric acid, HNO₃, is hazardous. It produces severe burns on the skin and the vapor is a lung irritant. Any acid spilled on the skin or splashed into your eyes must be rinsed with a large volume of water for 15 minutes. While handling nitric acid you must wear gloves. Rinse your hands with tap water after handling HNO₃.
- 3. NaOH solutions are corrosive to the skin.
- 4. Dissolution of the copper wire with concentrated HNO₃ should be carried out in the fume hood. The brown NO₂ gas that is evolved is very irritating to the lungs.
- 5. Dilute hydrochloric acid (HCl) and sulfuric acid (H₂SO₄) can harm eyes, skin, and clothing. Handle with care. Any acid spilled on the skin or splashed into your eyes must be rinsed with a large volume of water.

PROCEDURE

Day 1

Remember: [DO NOT PUT ANY EXCESS REAGENTS BACK INTO THE REAGENT BOTTLES!]

<u>Step I</u>

1. Weigh a clean and dry evaporating dish to the nearest .1 mg. Record the mass on the data sheet (Page 5)

2. Obtain a piece of copper. Weigh the copper plus an evaporating dish to the nearest 0.1 milligram, recording the mass on the data sheet (page 5)

3. Place the copper into a 250 mL beaker; and place it under the fume hood. Your instructor will put on a pair of rubber gloves and will carefully add 4.0 ml of 16M nitric acid, HNO₃ concentrated.

4. Record your observations on the report sheet, page5. After the copper has dissolved swirl the solution to remove the brown NO_2 gas (What is in the solution when the reaction is complete?).

5. After the copper has dissolved, <u>carefully (slowly!</u>) add approximately 100 ml distilled water to your copper solution.

6. Make and record your observations on page 5

Step II (Conducted at your lab bench)

1. While stirring the solution with a glass rod, <u>slowly</u> add 15 mL of Dil (6 M) NaOH to precipitate $Cu(OH)_2$. It is important to add the NaOH slowly, because you are adding a base to an acid. (This type of reaction generates a lot of heat). If your mixture warms up too much, you will skip step II and form the CuO directly-Step III.)

4. Start to preheat hot plates

3. The solution should be alkaline (basic) after the addition. A drop of solution touched to red litmus paper turns it blue. If the solution is not basic, add an additional 5 mLs of NaOH. Wait until the precipitate settles slightly and look at the clear top part of the mixture. There should be no blue color left in the top clear liquid, the solution). (What ion gives the blue color?)

4. Record your observations

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Step III

- 1. Heat the Cu(OH)₂ solution just barely to the boiling point using a **hot-plates**, while stirring continuously gently with a glass rod to prevent "bumping" (a phenomenon caused by the formation of a large steam bubble in a locally overheated area of the beaker)
- 2. Continue to heat gently while stirring to better coagulate the CuO.

- 3. When the transformation is complete, remove the hot solution from the hotplate to cool, continue stirring for a minute or two.
- 4. Record your observations.
- 5. Cover your solution with a watch glass and store it in the designated cabinet under the Fume Hood on top of a paper towel with your name until the next lab period.

Day 2 Step IV

- 1. Carefully decant as much supernatant as possible without losing ppt.
- 2. Now filter the solution using a clean vacuum filtration system. Then rinse the black solid thoroughly three times using 7 mL of DI water.
- Place a clean 125 mL Erlenmeyer flask under the stem of the funnel. Slowly pour 5 mL of 3M H₂SO₄. Repeat this step if black solid remains. If the entire solid is not dissolved, ask instructor for help. (What is in the solution now?).
- 4. Transfer the solution into a clean 250 mL beaker and place it into the hood.
- 5. Record your observations.

Step V

This part of the experiment is to be conducted in the fume hood.

1. While in the hood, place two 1 in. squares of aluminum foil into the solution. Ask instructor to add 4-5 drops concentrated HCl to your solution. Immediately start to stir for 1 min. (What happens? - What gas is produced?).

- 2. Check to see that the solution is colorless.
- 3. Record your observations

4. Wash the solid with two separate 50 mL portions of distilled water, decanting carefully after each washing.

5. Transfer all the copper to the evaporating dish using a little water to wash all the copper from the beaker. Decant the water used in transferring.

6. Place the evaporating dish with the copper on a steam bath to **dry to constant mass-** within 10 mg. (What color is the metal?)

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7. After the evaporating dish has cooled to room temperature, weigh the dish plus the copper metal.

8. Place copper into waste.

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Repo	ort Sheet	Name		F '
		Instructor's initi	Las [.] al	t First
	Experiment: Reactions	of Copper		
DATA	A AND CALCULATIONS 1. Mass of the empty evaporating dish			
	2. Initial mass of conner wire plus the out			
	2. Initial mass of copper wire plus the ev			
		Da	ata approva	al
	3. Calculate initial mass of Copper wire			
	Setup:			
	4. Final mass of recovered copper			
	Mass of evaporating dish plus dr	y copper #1		
		#2		
		#3		
		Da	ata approva	al
	5. Mass of recovered copper Setup:			
	6 Percentage recovery (show calculation Setup:	ns)		
<u>EQU</u>	ATIONS AND OBSERVATIONS			
and w each s <u>STE</u> I	For each step of the cycle, write the province physical states. Classify each reaction step and answer the questions posed ear <u>PI</u>	oducts of the reactior on. Also, record your lier in the Experimen	n, balance f observatio tal Procedu	he chemical equations, hos of what happens at ure.
	$Cu + HNO_3 \rightarrow$			
	What ions are in the solution after the rea	action is complete?		
	Observations:			
1.				
2.				
3.				

Data approval_____

<u>STEP II</u>

	Cu(NO ₃) ₂ + NaOH →		
	Reaction type		_
	HNO_3 + NaOH \rightarrow		
	Reaction type		_
	The gelatinous precipitate is	s Cu(OH) ₂ .	
	What ions are in solution?		_
	Observations:		
1.			
2.			
		Da	ta Approval
<u>Ste</u>	<u>EP III</u>		
	heat Cu(OH) ₂ →		
	Reaction type		_
	Observations:		
		Da	ta Approval
<u>STE</u>	<u>PIV</u>		
	CuO + H ₂ SO ₄ →		
	Reaction type		_
	Observations:		
1.			
2.			
			Data approval

<u>STEP V</u>

	Rea	ctior	i type						_				
	AI	+	H ₂ SO	4	\rightarrow								
	Rea	ctior	ı type						_				
	AI	+	HCI	\rightarrow									
	Rea	ctior	ı type						_				
	Obs	erva	tions:										
1.													
2.													
3.													
4													
	Wha	at ga	s is pro	oduc	ed in th	e reaction?							
	Wha Wha	at ga at ior	s is pro ns are r	oduc emc	ed in th oved by	e reaction? the washin	g and de	cantatio	— n nea	r the e	end of	STER	⊳V?
	Wha Wha Wha	at ga at ior at is t	s is pro ns are r the colo	oduc emc or of	ed in th oved by recove	the vashin	g and de	cantatio	 n nea	r the e	end of	STEF	⊳V?
	Wha Wha Wha	at ga at ior at is t	s is pro	oduc emc or of	ed in th oved by recove	the vashin	g and de ?	cantatio	 n nea 	r the o	end of	STEF	⊳ Vš
QU	Wha Wha Wha <u>ESTI</u>	at ga at ior at is t <u>ON</u> :	s is pro hs are r the colo <u>S:</u>	emc	ed in th oved by recove	the washin	g and de ?	cantatio	 n nea 	r the e	end of	⁻ STEF	o ∨?
<u>QU</u> 1. G	Wha Wha Wha <u>ESTI</u> Give po	at ga at ior at is t <u>ON</u> ossib	s is pro ns are r the colo <u>S:</u> lle reas	oduc emo or of ons	ed in th oved by recove why th	the washin the washin red copper	g and de ? ge recove	cantatio	n nea	r the e	end of pprov	[:] STEF	₽ \?
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<u>QU</u> 1. G	Wha Wha Wha <u>ESTI</u> Give po a. b.	at ga at ior at is t <u>ON</u>	s is pro ns are r the colo <u>S:</u> lle reas	oduc emo or of ons	ed in th wed by recove why th	e reaction? the washin ered copper	g and de ? ge recove	cantatio	n nea	T the o	end of	[:] STEF	₽ V?
QU 1. G	Wha Wha Wha ESTI Give po a. b. c.	at ga at ior at is t <u>ON</u>	s is pro hs are r the colo <u>S:</u> lle reas	oduc emc or of ons	ed in th oved by recove why th	e reaction? the washin ered copper	g and de ? ge recove	cantatio	n nea	r the e	end of	⁻ STEF	₽ V?
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QU 1. G ; ; 2. G a.	Wha Wha Wha ESTI Give po a. b. c. Give po	at ga at ior at is t ON: ossib	s is pro ns are r the colo <u>S:</u> le reas	oduc emc or of ons	ed in th oved by recove why th	e reaction? the washin red copper e percentag	g and de ? ge recove	ry might	t be o	ow.	pprov	[:] STEF	₽ \?
QU 1. G ; ; 2. G a. b.	Wha Wha Wha <u>ESTI</u> Give po a. b. c. Give po	at ga at ior at is t ON	s is pro hs are r the colo <u>S:</u> lle reas	oduc emc or of ons	ed in th oved by recove why th	e reaction? the washin ered copper e percentag	g and de ? ge recove	ery might	n nea	ow.	pprov	STEF	₽ V?
QU 1. G ; ; ; 2. G a. 5.	Wha Wha Wha <u>ESTI</u> Give po a. b. c. Give po	at ga at ior at is t ON	s is pro is are r the colo <u>S:</u> ile reas	oduc emc or of ons	ed in th oved by recove why th	e reaction? the washin ered copper e percentag	g and de ? ge recove	cantatio	n nea	r the o	pprov	STEF	₽ V?

3. How many milliliters of 3.0 M NaOH are required to react with 4.0 mL of 16.0 M HNO3?

Equation:

Answer____

4. How many milliliters of 3.0 M NaOH are required to react with 0.50 g of Cu²⁺ to form Cu(OH)₂?

Equation:

Answer____

5. What is the theoretical percentage copper in Cu(OH)₂?

Setup:

Answer_____6. What theoretical weight of CuO can be obtained from 2.00 g of Cu?

Setup:

Answer_

7. Iron is obtained from iron ore according to the following reaction:

 Fe_2O_3 + 3 CO \rightarrow 2 Fe + 3 CO₂ (unbalanced)

a. Assuming the blast furnace is 90.0 % efficient in recovering the iron, what is the actual mass of iron obtainable from a ton of ore?

1 ton = 2000.0 lb 1 lb = 453.6 g

Answer_____

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Problem #7 continued

b. What is theoretical yield of iron if 200.0 g of Fe_2O_3 reacts with 100.0 g of CO?

Answer_____

c.How many grams of excess reactant are left at the end of the reaction?

Answer_____

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