

The Hong Kong University of Science and Technology
Chemical and Biomolecular Engineering Department

Work Plan: 17038

Title: Gold Recovery from E-waste

Name: Wong Hon Fai & Chiu Yat Ming

Supervisor: Prof. Tom Luo

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1. General Information

Name of Researchers:

Name	ITSC	Contact Number
Chiu Yat Ming	ymchiuaa@connect.ust.hk	51053216
Wong Hon Fai	hfwongad@connect.ust.hk	51675088

Name of Supervisors: Tom Luo

Project Title: Gold Recovery from E-waste

Research Area: Environmental

Location: 7109

Proposed Start Date: 19/6/2017

2. Project Description

In Hong Kong, each year there are over 70,000 tons of computers and electronic wastes deposited. Among those electrical waste, they contain quite a large percentage of precious metals such as Gold, and Silver. Few companies in HK are interested in recycling those precious metals. If 1 million cell phones are collected, 24 kg of gold, 250 kg of silver, 9 kg of palladium, and more than 9,000 kg of copper can be recovered. 24 kg of gold are worth at least 800 thousand HKD. 1 million cell phones weigh around 150 tons, which is only 0.2% of the local electronic waste. This definitely has a tremendous market potential. In this research, recovering gold is our main target.

Gold has a high electricity conductivity and does not tarnish, these properties have made gold very popular in many industries. Gold is used in electronic industry, soldering semi-conductors, glass production or even spacecraft coating. All this usage is related to advanced technologies, and is expected to have paramount importance in those areas in the future. The usage of gold is getting more and more nowadays, it is a must to find an alternative way to get gold resources besides mining, recovery of gold can be a perfect solution to this problem.

3. Equipment List

Equipment
1. pH meter
2. Analytical balance (± 0.001 g accuracy)
3. Volumetric flasks (1000 ml and 2000 ml)
4. Heating plate with magnetic stirring
5. Plastic containers
6. Syringes
7. Borosilicate beakers (250 and 500 ml)
8. Desiccator
9. Spatula
10. Volumetric flask
11. Double Distilled water dispenser
12. Pipette
13. 110mm diameter rapid filter papers, cellulose-based ash-less type
14. Electric Hot plate
15. Burette 50 ml
16. Measuring cylinder
17. filter funnel
18. Glass microfiber filter media of 1 micron meter particle size
19. Suction flask
20. Waring Blender
21. Refrigerator
22. Oil Bath

- 23. Condenser unit
- 24. Vacuum Pump
- 25. Ultrasonic bath
- 26. Carbon Electrodes

4. Experimental Procedures

4.1 Dissolving electronic waste

1. Electronic waste is milled by blender.
2. 50g of electronic waste will be poured into a 100 ml beaker containing 50 ml hydrochloric acid.
3. Take 5 ml of acid solutions for analysis of gold using ICPOES.
4. Repeat step 1 and 3 at different reaction temperature (30°C, 40°C, 50°C), concentration of hydrochloric acid (1M, 2M, 3M, 4M), and reaction time (1hr, 2hrs, 3hrs).
5. Repeat step 1 to 4 using 50 ml acid medium containing hydrochloric acid and nitric acid at a ratio of 1:3 at different reaction temperature (30°C, 40°C, 50°C), ratio of hydrochloric acid and nitric acid (1:1, 1:2, 1:4), and reaction time (1hr, 2hrs, 3hrs).
6. The optimal reaction condition for dissolving gold can be found.

4.2 Solvent Extraction

1. For using hydrochloric acid as acid medium, 50 ml of electronic waste solution is mixed with 50 ml of toluene in a 250 ml beaker for 1 hour.
2. The organic solvent is then extracted and evaporated.
3. Repeat step 1 to 2 using ethanol, butanol, acetone, benzene, and hexane as extraction solvent.
4. For using both hydrochloric acid and nitric acid as acid medium, 50 ml of electronic waste solution is mixed with 50 ml of ethanol for 1 hour.
5. The organic solvent is then extracted and evaporated.
6. Repeat step 4 to 5 using acetone, hexane as extraction solvent.

4.3 Chromatography separation

1. Prepare a column packed with silica gel.
2. Pour 50ml extraction solvent collected from step 4.2.4 into the column.
3. Take 5 ml of separated solution for analysis of gold using ICPOES.
4. Repeat step 1 to 3 using extraction solvent collected from step 4.2.8.

4.4 Electrolysis

1. Prepare a tank for electrolysis
2. Add 0.6 g agarose to 40 ml DI water to prepare 1.5% gel.
3. Melt the agarose in a microwave oven for 1 minute.
4. Let the agarose cool to 55°C and pour the melted agarose in a casting tray.
5. Place the gel in the middle of the tank to separate the anode and cathode.
6. Pour 50 ml of electronic waste solution collected from 4.1.2 into anode, and pour the equal amount of DI water into cathode.
7. Run the electrolysis for 10 minutes.
8. Take 5 ml of solution collected from cathode for ICPOES analysis.

9. Repeat the step 6 at different reaction time (15 min, 30 min, and 45 min)
10. Repeat step 1 to 9 using the different % gel
11. Repeat step 1 to 10 using acrylamide gel
12. Add 0.2 ml of acrylamide and 0.4 ml of bisacrylamide to 40 ml DI water.
13. Pour the solution into a casting tray, and add 5ml of butanol and 5 ml of ammonium persulfate.
14. Repeat step 1 to 13 using electronic waste solution collected from 4.1.5.

4.5 Neutralization

1. Add 50 ml of solution from cathode in step 4.4 and 50 ml of Sodium hydroxide in to a 250 ml beaker for different reaction time (10 min, 20 min, and 30 min).
2. Separate the precipitate by vacuum filtration.
3. Take 5 ml of acid solutions for analysis of gold using ICPOES.

4.6 Replacement reaction

1. Add 10 g of copper bar into 50 ml of solution from cathode in step 4.4 into a 100 ml beaker and wait for 1 hour.
2. Separate the precipitate by vacuum filtration.
3. Take 5 ml of acid solutions for analysis of gold using ICPOES.
4. Repeat step 1 and 2 using different metal such as Iron, Magnesium, and lead.

4.7 Redox reaction

1. Add 10 ml of hydrogen peroxide into 20 ml of solution from cathode in step 4.4 into a 100 ml beaker and wait for 1 hour.
2. Separate the precipitate by vacuum filtration.
3. Take 5 ml of acid solutions for analysis of gold using ICPOES.

5. Procedure Template

4.1 Dissolving electronic waste					
Experimental Procedure No.	Experimental Procedure Description	Scale (Mass/Volume)	Location (Fumehood, benchtop, etc)	Method (New or Existing)	
4.1.1	Electronic waste is milled by blender.	Not available	benchtop	Existing	
4.1.2	50g of electronic waste will be poured into a 100 ml beaker containing 50 ml hydrochloric acid.	50ml HCl	Fumehood	Existing	
4.1.3	Take 5 ml of acid solutions for analysis of gold using ICPOES.	Not available	7101	Existing	
4.1.4	Repeat step 1 and 3 at different reaction temperature (30°C, 40°C, 50°C), concentration of hydrochloric acid (1M, 2M, 3M, 4M), and reaction time (1hr, 2hrs, 3hrs).	150 ml HCl	Fumehood	Existing	
4.1.5	Repeat step 1 to 4 using 50 ml acid medium containing hydrochloric acid and nitric acid at a ratio of 1:3 at different reaction temperature (30°C, 40°C, 50°C), ratio of hydrochloric acid and nitric acid (1:1, 1:2, 1:4), and reaction time (1hr, 2hrs, 3hrs).	50ml HNO ₃	Fumehood	Existing	
4.1.6	The optimal reaction condition for dissolving gold can be found.	Not available	benchtop	Existing	

4.2 Solvent Extraction					
Experimental Procedure No.	Experimental Procedure Description	Scale (Mass/Volume)	Location (Fumehood, benchtop, etc)	Method (New or Existing)	
4.2.1	For using hydrochloric acid as acid medium, 50 ml of electronic waste solution is mixed with 50 ml of toluene in a 250 ml beaker for 1 hour.	50 ml toluene	Fumehood	Existing	
4.2.2	The organic solvent is then extracted and evaporated.	Not available	Fumehood	Existing	
4.2.3	Repeat step 1 to 2 using ethanol, butanol, acetone, benzene, and hexane as extraction solvent.	50 ml ethanol, butanol, acetone, benzene, and hexane	Fumehood	Existing	
4.2.4	For using both hydrochloric acid and nitric acid as	50 ml ethanol	Fumehood	Existing	

	acid medium, 50 ml of electronic waste solution is mixed with 50 ml of ethanol for 1 hour.			
4.2.5	The organic solvent is then extracted and evaporated.		Fumehood	Existing
4.2.6	Repeat step 4 to 5 using acetone, hexane as extraction solvent.	Not available 50 ml acetone, and hexane	Fumehood	Existing

4.3 Chromatography separation

Experimental Procedure No.	Experimental Procedure Description	Scale (Mass/Volume)	Location (Fumehood, benchtop, etc)	Method (New or Existing)
4.3.1	Prepare a column packed with silica gel.	100g silica gel	benchtop	Existing
4.3.2	Pour 50ml extraction solvent collected from step 4.2.4 into the column.	Not available	benchtop	Existing
4.3.3	Take 5 ml of separated solution for analysis of gold using ICPOES.	Not available	7101	Existing
4.3.4	Repeat step 1 to 3 using extraction solvent collected from step 4.2.6.	100g silica gel	benchtop	Existing

4.4 Electrolysis

Experimental Procedure No.	Experimental Procedure Description	Scale (Mass/Volume)	Location (Fumehood, benchtop, etc)	Method (New or Existing)
4.4.1	Prepare a tank for electrolysis	Not available	Benchtop	Existing
4.4.2	Add 0.6 g agarose to 40 ml DI water to prepare 1.5% gel.	0.6 g agarose	Benchtop	Existing
4.4.3	Melt the agarose in a microwave oven for 1 minute.	Not available	Benchtop	Existing
4.4.4	Let the agarose cool to 55°C and pour the melted agarose in a casting tray.	Not available	Benchtop	Existing
4.4.5	Place the gel in the middle of the tank to separate the anode and cathode.	Not available	benchtop	Existing
4.4.6	Pour 50 ml of electronic waste solution collected from 4.1.2 into anode, and pour the equal amount of DI water into cathode.	Not available	benchtop	Existing
4.4.7	Run the electrolysis for 10 minutes.	Not available	Fumehood	Existing
4.4.8	Take 5 ml of solution collected from cathode for	Not available	7101	Existing

	ICPOES analysis.			
4.4.9	Repeat the step 6 at different reaction time (15 min, 30 min, and 45 min)	3 g agarose	Fumehood	Existing
4.4.10	Repeat step 1 to 9 using the different % gel	10 g agarose	Benchtop, fumehood	Existing
4.4.11	Repeat step 1 to 10 using acrylamide gel	Not available	Benchtop, fumehood	Existing
4.4.12	Add 0.2 ml of acrylamide and 0.4 ml of bisacrylamide to 40 ml DI water.	0.2 ml acrylamide 0.4 ml bisacrylamide	Fumehood	Existing
4.4.13	Pour the solution into a casting tray, and add 5ml of butanol and 5 ml of ammonium persulfate.	5ml butanol 5 ml ammonium persulfate		Existing
4.4.14	Repeat step 1 to 13 using electronic waste solution collected from 4.1.5.	13 g agarose 0.2 ml acrylamide 0.4 ml bisacrylamide 5ml butanol 5 ml ammonium persulfate	Fumehood	Existing

4.5 Neutralization

Experimental Procedure No.	Experimental Procedure Description	Scale (Mass/Volume)	Location (Fumehood, benchtop, etc)	Method (New or Existing)
4.5.1	Add 50 ml of solution from cathode in step 4.4 and 50 ml of Sodium hydroxide in to a 250 ml beaker for different reaction time (10 min, 20 min, and 30 min).	150 ml Sodium hydroxide	Fumehood	Existing
4.5.2	Separate the precipitate by vacuum filtration.	Not available	Fumehood	Existing
4.5.3	Take 5 ml of acid solutions for analysis of gold using ICPOES.	Not available	Fumehood	Existing

4.6 Replacement reaction

Experimental Procedure No.	Experimental Procedure Description	Scale (Mass/Volume)	Location (Fumehood, benchtop, etc)	Method (New or Existing)
4.6.1	Add 10 g of copper bar into 50 ml of solution from cathode in step 4.4 into a 100 ml beaker and wait for 1 hour.	10 g copper	benchtop	Existing
4.6.2	Separate the precipitate by vacuum filtration.	Not available	benchtop	Existing
4.6.3	Take 5 ml of acid solutions for analysis of gold	Not available	7101	Existing

	using ICPOES.			
4.6.4	Repeat step 1 and 2 using different metal such as Iron, Magnesium, and lead.	10 g Iron, Magnesium, and lead	benchtop	Existing

4.7 Redox reaction				
Experimental Procedure No.	Experimental Procedure Description	Scale (Mass/Volume)	Location (Fumehood, benchtop, etc)	Method (New or Existing)
4.7.1	Add 10 ml of hydrogen peroxide into 20 ml of solution from cathode in step 4.4 into a 100 ml beaker and wait for 1 hour.	10 ml hydrogen peroxide	Fumehood	Existing
4.7.2	Separate the precipitate by vacuum filtration.	Not available	Fumehood	Existing
4.7.3	Take 5 ml of acid solutions for analysis of gold using ICPOES.	Not available	7101	Existing

6. HAZOP Analysis

HAZOP Template

Hazard and Operability Analysis (4.1 Dissolving electronic waste)

NO	HAZARD	HAZARD EFFECT	SEVERITY	PROBABILITY	RISK	MINIMISE RISK BY	RESIDUAL RISK
4.1.1	Blade from blender	Physical injuries of fingers	M	L	L	Get training for proper operation of equipment	L
4.1.2	Contact with hydrochloric acid	Skin, eye and respiratory irritation	M	H	H	Prepare stock solution and carry out reaction in fume hood. Always wear lab coat, gloves and safety goggles	M
4.1.3	Not available						
4.1.4	Hot heat plate	Burned skin	M	M	M	Wear heat protective gloves, lab coat and safety goggles. Put up warning sign Collect samples only when they are cooled to room temperature	L
4.1.5	Contact with hydrochloric acid and nitric acid	Skin, eye and respiratory irritation	M	H	H	Prepare stock solution and carry out reaction in fume hood. Always wear lab coat, gloves and safety goggles	M
4.1.6	Not available						
4.1.7	Contact with sulfuric acid	Skin, eye and respiratory irritation	M	H	H	Prepare stock solution and carry out reaction in fume hood. Always wear lab coat, gloves and safety goggles	M
4.1.8	Contact with hydrochloric acid	Skin, eye and respiratory irritation	M	H	H	Prepare stock solution and carry out reaction in fume hood. Always wear lab coat, gloves and safety goggles	M
FINAL ASSESSMENT: User can protect themselves with proper training and take precaution					OVERALL RISK:		
					M		

Remark: Severity-Low:1 (Minor injuries, first aid); Medium:2 (Hospitalization, medical leave); High:3 (Serious injuries, fatality) Probability-
Unlikely:1; Possible:2; Very Likely: 3

HAZOP T-14-

Hazard and Operability Analysis (4.2 Solvent Extraction)

NO	HAZARD	HAZARD EFFECT	SEVERITY	PROBABILITY	RISK	MINIMISE RISK BY	RESIDUAL RISK
4.2.1	Contact with hydrochloric acid and toluene	Skin, eye and respiratory irritation. Toxic to health if swallowed or breathed in.	H	H	H	Prepare stock solution and carry out reaction in fume hood. Always wear lab coat, gloves and safety goggles	L
4.2.2	Not available						
4.2.3	Contact with ethanol, butanol, acetone, benzene, or hexane	Skin, eye and respiratory irritation. Toxic to health if swallowed or breathed in.	H	H	H	Prepare stock solution and carry out reaction in fume hood. Always wear lab coat, gloves and safety goggles	L
4.2.4	Contact with hydrochloric acid and nitric acid	Skin, eye and respiratory irritation. Toxic to health if swallowed or breathed in.	H	H	H	Prepare stock solution and carry out reaction in fume hood. Always wear lab coat, gloves and safety goggles	L
4.2.5	Not available						
4.2.6	Contact with hydrochloric acid, nitric acid, acetone and hexane	Skin, eye and respiratory irritation. Toxic to health if swallowed or breathed in.	H	H	H	Prepare stock solution and carry out reaction in fume hood. Always wear lab coat, gloves and safety goggles	L
FINAL ASSESSMENT: User can protect themselves with proper training and take precaution						OVERALL RISK:	L

FINAL ASSESSMENT: User can protect themselves with proper training and take precaution	OVERALL RISK: 1
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Comments from proper planning and take precautions	OVERALL RISK:	L
Remark: Severity-Low:1 (Minor injuries, first aid); Medium:2 (Hospitalization, medical leave); High:3 (Serious injuries, fatality) Probability--Unlikely:1; Possible:2; Very Likely: 3.		

HAZOP Template							
Hazard and Operability Analysis (4.3 Chromatography separation)							
NO	HAZARD	HAZARD EFFECT	SEVERITY	PROBABILITY	RISK	MINIMISE RISK BY	RESIDUAL RISK
4.3.1	Break of glass apparatus	scratch on skin	L	L	L	be careful with placing and handling those instruments	L
4.3.2	Not available						
4.3.3	Not available						
4.3.4	Not available						
FINAL ASSESSMENT: User can protect themselves with proper training and take precaution					OVERALL RISK:		
Remark: Severity-Low:1 (Minor injuries, first aid); Medium:2 (Hospitalization, medical leave); High:3 (Serious injuries, fatality)					Probability-		
					Unlikely:1; Possible:2; Very Likely: 3.		

HAZOP Template							
Hazard and Operability Analysis (4.4 Electrolysis)							
NO	HAZARD	HAZARD EFFECT	SEVERITY	PROBABILITY	RISK	MINIMISE RISK BY	RESIDUAL RISK
4.4.1	Not available						
4.4.2	Contact with agarose	Skin, eye and respiratory irritation	M	H	H	Prepare stock solution and carry out reaction in fume hood. Always wear lab coat, gloves and safety goggles	M
4.4.3	Hot solution from microwave	Burned skin	M	M	M	Wear heat protective gloves, lab coat and safety goggles. Put up warning sign Collect samples only when they are cooled to room temperature	L
4.4.4	Not available						
4.4.5	Not available						
4.4.6	Not available						
4.4.7	Electronic device	Electric shock	M	L	L	Keep hands dry, wear insulated gloves	L
4.4.8	Not available						
4.4.9	Not available						
4.4.10	Contact with agarose	Skin, eye and respiratory irritation	M	H	H	Prepare stock solution and carry out reaction in fume hood. Always wear lab coat, gloves and safety goggles	M
4.4.11	Not available						
4.4.12	Contact with acrylamide or bisacrylamide	Skin, eye and respiratory irritation	M	H	H	Prepare stock solution and carry out reaction in fume hood. Always wear lab coat, gloves and safety goggles	M
4.4.13	Contact with butanol or ammonium persulfate.	Skin, eye and respiratory irritation	M	H	H	Prepare stock solution and carry out reaction in fume hood. Always wear lab coat, gloves and safety goggles	M
4.4.14	Not available						
FINAL ASSESSMENT: User can protect themselves with proper training and take precaution						OVERALL RISK:	M

Remark: Severity-Low:1 (Minor injuries, first aid); Medium:2 (Hospitalization, medical leave); High:3 (Serious injuries, fatality) Probability-Unlikely:1; Possible:2; Very Likely: 3.

HAZOP Template Hazard and Operability Analysis (4.5 Neutralization)							
NO	HAZARD	HAZARD EFFECT	SEVERITY	PROBABILITY	RISK	MINIMISE RISK BY	RESIDUAL RISK
4.5.1	Contact with Sodium hydroxide	Skin, eye and respiratory irritation	M	H	H	Prepare stock solution and carry out reaction in fume hood. Always wear lab coat, gloves and safety goggles	M
4.5.2	Not available						
4.5.3	Not available						
4.5.4	Contact with Sodium hydroxide	Skin, eye and respiratory irritation	M	H	H	Prepare stock solution and carry out reaction in fume hood. Always wear lab coat, gloves and safety goggles	M
4.5.5	Contact with agarose	Skin, eye and respiratory irritation	M	H	H	Prepare stock solution and carry out reaction in fume hood. Always wear lab coat, gloves and safety goggles	M
FINAL ASSESSMENT: User can protect themselves with proper training and take precaution						OVERALL RISK:	M

Remark: Severity-Low:1 (Minor injuries, first aid); Medium:2 (Hospitalization, medical leave); High:3 (Serious injuries, fatality) Probability-Unlikely:1; Possible:2; Very Likely: 3.

HAZOP Template							
Hazard and Operability Analysis (4.6 Replacement reaction)							
NO	HAZARD	HAZARD EFFECT	SEVERITY	PROBABILITY	RISK	MINIMISE RISK BY	RESIDUAL RISK
4.6.1	Not available						
4.6.2	Not available						
4.6.3	Not available						
4.6.4	Not available						
FINAL ASSESSMENT: User can protect themselves with proper training and take precaution					OVERALL RISK:		
Remark: Severity–Low:1 (Minor injuries, first aid); Medium:2 (Hospitalization, medical leave); High:3 (Serious injuries, fatality) Probability–					L		

Unlikely:1; Possible:2; Very Likely: 3.

HAZOP Template Hazard and Operability Analysis (4.7 Solvent Extraction)							
NO	HAZARD	HAZARD EFFECT	SEVERITY	PROBABILITY	RISK	MINIMISE RISK BY	RESIDUAL RISK
4.7.1	Contact with hydrogen peroxide	Skin, eye and respiratory irritation	M	H	H	Prepare stock solution and carry out reaction in fume hood. Always wear lab coat, gloves and safety goggles	M
4.7.2	Not available						
4.7.3	Not available						
FINAL ASSESSMENT: User can protect themselves with proper training and take precaution						OVERALL RISK:	M

Remark: Severity-Low:1 (Minor injuries, first aid); Medium:2 (Hospitalization, medical leave); High:3 (Serious injuries, fatality) Probability-Unlikely:1; Possible:2; Very Likely: 3.

7. Operating condition

Temperature: 20°C- 70°C

Pressure: atmospheric

Flow rates: Batch operation, N/A

8. Services List

Electric power: 220V, 15A

Water: distilled water, tap water, DDI water

Gas: Compressed air

Fume hood in 7109

9. Chemical List

No.	Chemical	Purity	Quantity per Experiment
(1)	Hydrochloric acid	37%	250mL
(2)	Nitric acid	65%	250mL
(3)	Hexane	95%	250mL
(4)	Ethanol	95%	250mL
(5)	Toluene	95%	250mL
(6)	Agarose	100%	50g
(7)	Silica gel	100%	50g
(8)	Hydrogen peroxide	30%	250mL
(9)	Sodium hydroxide	30%	250mL
(10)	Copper	90%	10g
(11)	• • • Acrylamide	90%	100mL
(12)	Bisacrylamide	90%	100mL
(13)	Ammonium persulfate	90%	100mL
(14)	Iron	90%	10g
(15)	Magnesium	90%	10g

(16)	Lead	90%	10g
(17)	Acetone	95%	100ml
(18)	Benzene	95%	100mL

10. Biological Agent List

Nil

11. Summary of Relevant Hazard and Incompatibilities

Material	Summary of Hazards	Incompatibilities
Hydrochloric acid	Corrosive to metals. Severe skin and eye burns. Respiratory Irritation.	Bases, Amines, Alkali metals, Metals, permanganates, e.g. potassium permanganate, Fluorine, metal acetylides, hexalithium disilicide
Nitric acid	May intensify fire; oxidizer. May be corrosive to metals. Causes severe skin burns and eye damage.	Alkali metals, Acetic anhydride, Organic materials, Alcohols, Acetonitrile, Acrylonitrile
Hexane	Highly flammable, Harmful, Irritant, Dangerous to environment	Strong oxidizing agent
Ethanol	Highly flammable liquid and vapor. May form explosive mixtures with air	Alkalis and oxidizing agents
Toluene	Flammable, Severe respiratory Irritation.	Oxidizing agent
Agarose	Slightly hazardous in case of skin contact or eye contact, will lead to irritation.	Nil
Silica gel	Slightly hazardous in case of skin contact or eye contact, will lead to irritation.	Nil
Hydrogen peroxide	Severe skin irritation and eye inflammation. Respiratory Irritation.	Oxidizing agents, reducing agents, combustible materials, organic materials, metals, acids, alkalis.

Sodium hydroxide	Severe skin and eye burn. Severe respiratory Irritation.	Oxidizing agents, reducing agents, metals, acids, alkalis, moisture.
Copper	Slightly skin and eye irritation.	Nil
Acrylamide	Severe skin and eye burns. Respiratory Irritation.	Oxidizing agent, acids, alkalis and moisture.
Bis-acrylamide	Skin and eye burns. Respiratory Irritation. Prolonged or repeated exposure affects the nervous system.	Strong oxidizing agent, strong reducing agent, strong acid and strong alkali
Ammonium persulfate	Severe skin burn, and inhalation irritation.	Reducing agents, combustible materials, organic materials, metals.
Iron	Skin and eye irritation.	Oxidizing agent and acid
Magnesium	Skin and eye Irritation.	Oxidizing agent, acid and moisture
Lead	Skin and eye Irritation.	Oxidizing agent.
Acetone	Severe skin burn, inhalation and skin irritation.	Oxidizing agent, reducing agent, acid and alkali
Benzene	Skin, eye and inhalation irritation	Oxidizing agent and acid

12. Waste List

1. Large amount of sodium hydroxide, hydrochloric acid and nitric acid are neutralized to pH7 before disposal into sink. Small amount will be disposed directly into alkaline and inorganic acid waste barrel.
2. Chlorine containing solvent, such as dichloromethane will be disposed into hydrogenated organic solvent waste barrel.
3. Other organic solvent will be disposed into non-hydrogenated organic solvent waste barrel.
4. Metal waste will be disposed into metal waste container.

13. Assessment of Significant Risks

In this experiment, highly flammable solvent are used, keep those solvents away from heat source, flame source and sparks.

Highly toxic organic solvents are used, always handle the chemicals in fume-hood to prevent breathing in toxic vapors.

Sodium hydroxide, hydrochloric acid and nitric acid are involved, which is highly corrosive can lead to severe

skin burn. Wear personal protective equipment all the times, and carry the experiment in fume-hood to avoid breathing in acid fume.

14. Safety Precaution

Safety Training Required

Safety training courses, included Chemical Safety and Hazardous Waste Management training organized by the HSEO should be attended.

Equipment Training Required

Training with technician to use the characterization instruments in lab 7101 and processing equipment in 7109 should be arranged.

Personal Protective Equipment

Laboratory coat, safety glasses, gloves, heat gloves (while handling hot samples) and goggles are worn throughout experiments

Fume-hood will be used when handling hazardous and flammable chemicals listed in the Chemical and Materials section.

In the event of emergency (e.g. chemical spillage) workers' actions will follow the standard procedures suggested by University's Chemical Safety training as well as Emergency Procedures.

Safety precautions of flammable liquids

1. Keep the flammable chemicals away from heat and ignition sources.
2. Do not leave the containers opened as vapors can flow along surfaces and travel a considerable distance to a source of ignition and flash back.

Warning sign required

1. All the chemicals will be stored based on the hazard categories with clear labeling.
2. The composition of the chemicals, date of receipt, use and expiration must be noted on label.
3. Chemical inventory will be minimized to reduce the overall risks.
4. Labels of HIGH VOLTAGE and HIGH CURRENT must be noted and be seen clearly.
5. Put up warning signs with the contact number of researcher when reactions are being carried out.

Emergency Procedures

1. Ventilated area and wash spill site after material pickup is complete. Evacuate area.
2. If inhaled the toxic fumes, remove to fresh air. If breathing is difficult, give oxygen.
3. In case of skin contact, flush with copious amounts of water for at least 15 minutes. Assure adequate flushing by separating the eyelids with fingers. Call a physician.
4. If the situation is out of control, evaluate from the scene immediately. Alert others people to evaluate from the scene immediately. Inform SEPO and security unit promptly about the incident.

15. Action in case of Abnormal or emergency situation

In the event of emergency workers' actions will follow the standard procedures suggested by University's Chemical Safety training as well as Emergency Procedures. All abnormality during experiment shall be reported to laboratory technical staff immediately.

In case of Chemical Spill

Hazardous Chemical Spill in Research Laboratory

- Alert co-workers
- Carry out emergency shut-down: shut down the analysis run via software (in low pressure analysis), or press red "Emergency Stop" button immediately for 2-3 seconds (in high pressure analysis).
- Report to laboratory technical staff immediately;
- If safe to do so, confine the spill with appropriate materials. Turn off remotely all heat/ignition sources if flammable vapor is involved.
- Ask for assistance if necessary. Press the Emergency Ventilation bottom (do not activate this bottom in case of fire).
- Inform the Security Control Centre by dialing 8999 from a safe location.
- Evacuate everyone in the affected area. Leave contaminated clothing and close the door.
- Activate local warning system to prevent others from entering the room.
- If possible, maintain a safe distance from the scene, keep the entrance or access routes in sight and help to prevent entry to the affected room.
- If conditions allow, remain to assist the emergency response team.

In case of Fire

When you hear fire alarm:

- Remain calm and check if there is any sign of fire in the vicinity.
- If you see fire or smoke, or hear the announcement asking you to evacuate, follow the evacuation procedures.
- If there is no sign of a fire, stay alert and pay attention to announcement until the fire alarm is silenced.
- Evacuate if the alarm has sounded for more than two minutes

If you discover a fire:

- Activate the fire alarm by pressing the breakglass fire alarm button located at the corridor.
- Report to Security Control Centre by dialing 8999.
- Alert other people. If SAFE to do so, try to put out the fire by firefighting equipment.
- DO NOT take any personal risk. If the fire gets beyond your control, evacuate immediately by following the procedures below.

- Close the door of the room on fire.

When someone is injured or ill:

- Call Security Control Centre (SCC) by dialing 8999.
- Call for Community Emergency Service directly by dialing (9)999 if the situation is urgent or serious and inform SCC subsequently.
- DO NOT conduct rescue operation, unless you know for sure how to perform a proper rescue and you know the situation is safe? Careless rescue operation may endanger the rescuers when, for example, the victim is inside a room filled with toxic gas, or is still in contact with live electricity.
- DO NOT move an injured person, especially when there are signs of spinal injury or bone fracture, unless it is absolutely necessary to do so for safety reason.
- Keep the injured or ill person comfortable, warm, and lying down.
- Give First aid treatment if necessary.
 1. Acid and alkali burns--flush with running water; use emergency shower if necessary. Do not attempt to neutralize.
 2. Heat or cold burns--flush with cold water.
 3. Chemical in eyes--flush eyes with emergency eyewash.
 4. Major bleeding--apply direct pressure to the wound using a clean cloth.
 5. Toxic gas inhalation--expose to fresh air.
 6. Hydrofluoric acid exposure--use antidote immediately
 7. Cyanide exposure--use antidote immediately