

I. 預報包括：

- a. 目的：簡述。
- b. 原理：簡述實驗所應用之原理及重要之化學反應式。
- c. 藥品：需清楚瞭解所用藥品之性質，含中英文名、化學式、莫耳質量、物理與化學性質及危害毒性。
- d. 步驟：簡要條列或流程表示，繪製裝置圖，留右側 1/3 頁面書寫觀察

裝訂於
左上方

組別：
姓名：
學號：

E4. 維生素C之定量

+30 / 35

目的：簡述

一. 目的：利用氧化還原反應，測定市售維生素C藥錠及果汁中維生素C之含量 ✓

二. 原理：維生素C又稱抗壞血酸(L-ascorbic acid)為一良好還原劑，可使 $Fe^{3+} \rightarrow Fe^{2+}$ 或 $I_2 \rightarrow I^-$

2. 氧化還原滴定：(1) 滴定劑：碘酸鉀 (2) 環境：酸性 (3) 被滴定：碘化鉀 ?



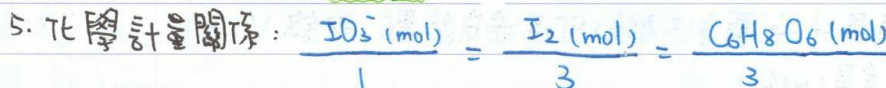
* Vit. C

3. 碘分子與抗壞血酸之氧化還原：

間接碘滴定法



4. 當抗壞血酸反應完畢，過量 I_2 和溶液中 I^- 形成 I_3^- ，且和預加在溶液中的澱粉指示劑形成藍黑色錯合物 (滴定終點)



mp./bp?. 式量可沒有單位，莫耳質量有單位 g/mol

三. 藥品	英文名稱	化學式	式量(g/mol)	外觀	d(g/cm ³)	毒性
維生素C C ₆ H ₈ O ₆	L-ascorbic acid	C ₆ H ₈ O ₆	176.12	白或黃固體	1.65	空間中濃度高會粉塵爆炸，遇Al, Zn會劇烈反應
1M 碘化鉀 NaI	sodium iodide	NaI(aq)	149.89	無色粉末	3.5(25°C)	不可吞食，刺激皮膚、眼，對水生生物毒性
0.025M 碘酸鉀 KIO ₃	potassium iodate	KIO ₃ (aq)	214	白色結晶	3.98(20°C)	遠離火源，吞食有害，刺激眼 (使用乾粉滅火)
1M 鹽酸 HCl	hydrochloric acid	HCl(aq)	36.46	無色液體	1.1(20°C)	刺激皮膚、眼、呼吸道
2% 澱粉水溶液	starch	N/A	N/A	白色固體	N/A	輕微刺激皮膚

ref: Merck SDS, TCI SDS

Merck index 沒有提? SDS有，氧化劑

編頁碼

II. 實驗紀錄：簡要、詳細、條列式記錄

- 反應物與產物之狀態與顏色。
- 進行實驗之條件。
- 操作過程。
- 反應過程顏色、沉澱、吸熱放熱等變化。
- 數據：參考實驗課本的格式或自創，但記錄整齊。

步驟：
以簡單清楚流程圖
表示，繪製裝置圖

實驗紀錄：

- 反應物與產物之狀態與顏色
- 實際進行實驗之條件
- 操作過程
- 反應過程溶液顏色、沉澱、吸熱放熱等變化

四. 步驟 +5/5

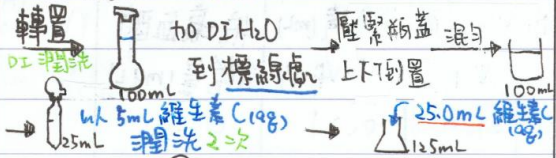
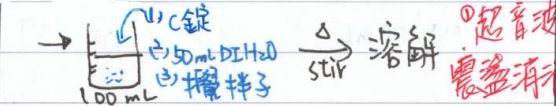
1. 維生素C錠測定：取2片C錠，記錄藥瓶標示之維生素C含量、重量。

2. 取一片C錠，稱量記錄重量

維生素C劑量 < 250mg

維生素C劑量 > 250mg

3. 以秤藥紙包裹C錠，以鐵槌擊碎



4. 125mL 加 ① 2mL, 1M NaI ② 2mL, 1M HCl ③ 1mL, 2% starch(aq)

五. 觀察

C錠顏色、外觀：淺黃、圓錠

C錠重：4.77g ✓
2張紙包裝敲碎粉都在

發泡錠不用敲碎
只需冰水溶解

用容量瓶標定
後要盡速倒出以免沾黏在瓶壁

C錠溶解後，液呈淺紫
有油狀漂浮於表面的膜

CHA SHIN

* NaI, HCl, starch
KIO₃ 均無色

4.5. 30mL, 0.025M KIO₃(aq)
乾淨 100mL 燒杯

5. 洗淨 25mL 滴定管，再用 5mL, 0.025M KIO₃ 潤洗 2 次並裝滿，記錄 V_i

* 滴定管中不可有氣泡

至 0.01 mL

6. 以 KIO₃(aq) 滴定維生素C(aq) → 藍色(不退色) 即達滴定終點

記錄 V_f

7. 取另一片C錠重複 3.~6. 完成二次重複試驗，計算維生素C含量

加 NaI, HCl, starch
液 soln 顏色不變(無色)

KIO₃: 透明澄清液
滴定時紫色出現又漸

錯過淺紫色出現不
褪的階段，滴定
完畢溶液呈深紫色

(二). 市售果汁中維生素C測定:

1. 洗淨 25 mL 果汁, 取 50.0 mL 果汁

果汁潤洗 2 次 (若含果肉 → 改用量筒)

2. 加入 2 mL, 1M NaI (2 mL, 1M HCl (1 mL, 2% starch(aq))

3. 以 $KIO_3(aq)$ 滴定果汁 → 藍色(褪去)即達滴定終點, 記錄 V_i, V_f

並計算維生素C含量 (mg/mL)

(三) 實驗完畢: 清洗移液吸管, 滴定管, 並倒夾於滴定管夾上

果汁: 灰白色液體

加入 NaI, HCl, starch 後

顏色不變

同C錠, 最終呈深紫色

助理助教章

吳美倫

要記錄, 你是 1000 mg
故 dilute to 100.0 mL
取 25.0 mL 作滴定

III. 結果報告:

a. 數據處理: 詳列計算過程, 結果須具正確有效數字及單位。

六. 實驗數據與結果

(一). C錠: 森萊富 維他命 C 1000 ✓

1. 標示劑量: 1000 mg/K (Na = 326 mg, 纖維 0.3 g)

2. 含量測定

	維生素C錠		0.025 M KIO_3 滴定體積 (mL)		抗壞血酸
	重量 (g)	V_i	V_f	$\Delta V = V_f - V_i$	含量 (mg/K)
#1	4.77 ✓	0.05	21.06	21.01	1.1×10^3
#2		0.05	21.05	21.00	1.1×10^3

計算在下一頁

維生素C錠中抗壞血酸平均含量: 1.1×10^3 mg/K

(二). 果汁: 蘋果 C 牌

1. 標示含量: 7.5 mg/100 mL ✓ 不太適合的試樣

2. 含量測定

	果汁取量		0.025 M KIO_3 滴定體積 (mL)		抗壞血酸
	(mL)	V_i	V_f	$\Delta V = V_f - V_i$	含量 (mg/100 mL)
	50	21.05	21.26	0.21	5.6
	50	21.27	21.46	0.19	5.0

SHE

大助教核章

計算

(一) C 錠 (高劑量) + 9/10

高劑量, 配 100mL 取 25mL

$$n_{C_6H_8O_6} = 3 n_{KIO_3} = 3 \times 0.025 \times \Delta V \times \frac{1}{1000} \times \frac{100}{25}$$

$$W_{C_6H_8O_6}^{(mg)} = n_{C_6H_8O_6} \times 176.12 \times 1000$$

$$\#1: \Delta V = 21.06 - 0.05 = 21.01 \text{ mL} \Rightarrow W_{C_6H_8O_6} = 3 \times 0.025 \times 21.01 \times \frac{1}{1000} \times \frac{100}{25} \times 176.12 \times 1000 = 1110.1 \text{ mg} \xrightarrow{s.f.=2} 1.1 \times 10^3 \text{ mg/g} \checkmark$$

$$\#2: \Delta V = 21.05 - 0.05 = 21.00 \text{ mL} \Rightarrow W_{C_6H_8O_6} = 3 \times 0.025 \times 21.00 \times \frac{1}{1000} \times \frac{100}{25} \times 176.12 \times 1000 = 1109.5 \text{ mg} \xrightarrow{s.f.=2} 1.1 \times 10^3 \text{ mg/g} \checkmark$$

$$\text{平均: } \frac{1.1 \times 10^3 + 1.1 \times 10^3}{2} = 1.1 \times 10^3 \text{ mg} \quad \text{平均 } 1.1 \times 10^3 \text{ mg/g} \checkmark$$

$$\text{誤差 } \frac{1.1 \times 10^3 - 1000}{1000} \times 100\% = 11.0\% \xrightarrow{s.f.=2} 11\% \checkmark$$

* 精簡報告, 可不談誤差討論

實驗得到之 $C_6H_8O_6$ 量較標示多, 可能原因: 1. 滴定終點判讀: 大約都多滴 1~2 滴, 溶液呈深藍紫色才停下 2. C 錠中有其他還原劑存在, 耗掉的 KIO_3 並非只和 $C_6H_8O_6$ 反應.

(二)

$$\text{果汁: } n_{C_6H_8O_6} = 3 n_{KIO_3} = 3 \times 0.025 \times \Delta V \times \frac{1}{1000} \times \frac{100}{50}$$

$$W_{C_6H_8O_6} = n_{C_6H_8O_6} \times 176.12 \times 1000$$

* $V_{\text{eg}} = 0.21 \text{ mL}$ 太少

誤差會很大, 避免用此類果汁

$$\#1: \Delta V = 21.26 - 21.05 = 0.21 \text{ mL} \Rightarrow W_{C_6H_8O_6} = 3 \times 0.025 \times 0.21 \times \frac{1}{1000} \times \frac{100}{50} \times 176.12 \times 1000 = 5.55 \text{ mg} \xrightarrow{s.f.=2} 5.6 \text{ mg}$$

$$\#2: \Delta V = 21.46 - 21.27 = 0.19 \text{ mL} \Rightarrow W_{C_6H_8O_6} = 3 \times 0.025 \times 0.19 \times \frac{1}{1000} \times \frac{100}{50} \times 176.12 \times 1000 = 5.02 \text{ mg} \xrightarrow{s.f.=2} 5.0 \text{ mg}$$

$$\text{平均: } \frac{5.6 + 5.0}{2} = 5.3 \text{ mg} \xrightarrow{s.f.=2} 5.3 \text{ mg}$$

* 列式、計算、s.f. 正確, good.

$$\text{誤差: } \frac{5.3 - 7.5}{7.5} \times 100\% = -29.5\% \xrightarrow{s.f.=2} -30\%$$

具體結論

實驗測得之 $C_6H_8O_6$ 較標示少, 可能原因: 1. 果汁中其他成份影響: 果汁中可能含有其他氧化劑, 使得實驗中加入的 KIO_3 消耗量下降, 測得 $C_6H_8O_6$ 較少

* 其他 = VitC 含量太少, V_{eg} 太少, 滴定誤差 ↑.

SHE

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- b. 原理：簡述實驗所應用之原理及重要之化學反應式。
- c. 藥品：需清楚瞭解所用藥品的性質，含中英文名、化學式、莫耳質量、物理與化學性質及危害毒性。

裝訂於
左上方



組別：
姓名：
學號：

E 8. 碘鐘實驗 - 碘鐘交響曲

37/50

目的：簡述

一. 目的：以定量 $S_2O_8^{2-}$ 與 I^- 的氧化還原反應測量 $S_2O_8^{2-}$ 與 I^- 的反應速率。學習以初始反應速率法決定反應級數、速率常數

原理：
簡述實驗所應用之原理及重要之化學反應式

二. 原理：(一) 反應速率、反應級數：[學動力學] 探討反應速率及影響因素。
反應速率：單位時間內 Δ [產物] 或 Δ [反應物]。若 $aA + bB \rightarrow cC$ ，速率 = $-\frac{d[A]}{a \cdot dt} = -\frac{d[B]}{b \cdot dt} = \frac{d[C]}{c \cdot dt}$
[反應物] 會影響速率，可用速率定律式 $rate = k[A]^m[B]^n$ 表示。k = 速率常數，m, n 分別為 A, B 的反應級數，總反應級數為 $m+n$ 。反應級數可為整、負、分數，不受平衡方程式係數影響，R 能由實驗測得。常用「初始反應速率法」，「積分作圖法」測級數
(二) $S_2O_8^{2-}$, I^- 反應速率測定：本實驗以初始反應速率法測 $S_2O_8^{2-}$, I^- 的速率定律式。利用限量 $S_2O_8^{2-}$ 作為計時劑
 $S_2O_8^{2-}(aq) + 2I^-(aq) \rightarrow 2SO_4^{2-}(aq) + I_2(aq)$ 立刻消耗掉
 $rate = k[S_2O_8^{2-}]^m [I^-]^n$
 $2S_2O_8^{2-}(aq) + I_2(aq) \rightarrow 2I^-(aq) + 2SO_4^{2-}(aq)$ 當 $[S_2O_8^{2-}] = 0 \Rightarrow I_2(aq) + I_2(aq) \rightarrow I_3(aq)$ 並和澱粉指示劑形成藍黑色錯合物 (出現時間即 Δt)
 $\Delta[S_2O_8^{2-}] = -\Delta[S_2O_8^{2-}]$
 $rate = \frac{-\Delta[S_2O_8^{2-}]}{\Delta t} = \frac{-\frac{1}{2}\Delta[S_2O_8^{2-}]}{\Delta t}$

藥品：
中英文名
化學式
莫耳質量
物理與化學性質及危害毒性

(三) 初始反應速率法：改[某反應物]。其他維持不變進行一系列試驗，求 Δ 濃度對初速率影響。本實驗每次將 $[S_2O_8^{2-}]$ 及 $[I^-]$ 增為 2 倍，其餘不變

$$\frac{r_2}{r_1} = \frac{k(2.0[S_2O_8^{2-}])^m ([I^-]_1)^n}{k([S_2O_8^{2-}]_1)^m ([I^-]_1)^n} = 2.0^m \quad ; \quad \frac{r_3}{r_1} = \frac{k([S_2O_8^{2-}]_1)^m (2.0[I^-])^n}{k([S_2O_8^{2-}]_1)^m ([I^-]_1)^n} = 2.0^n$$

三. 藥品	英文名稱	化學式	式量(M)	外觀顏色	d (g/cm ³)	毒性
0.20M 碘化鈉	sodium iodide	NaI	149.89	無色粉末	3.5(25°C)	吞食有害, 刺激皮膚、眼睛
0.10M 過硫酸鉀	potassium persulfate	K ₂ S ₂ O ₈	270.33	白色固體	2.477	氧化劑, 吞食有害, 刺激皮膚、眼、呼吸道
0.20M 氯化鈉	sodium chloride	NaCl	58.44	無色固體	2.16(25°C)	吞食有害, 刺激皮膚、眼、呼吸道
0.10M 硫酸鉀	potassium sulfate	K ₂ SO ₄	174.26	無色固體	2.662	勿吞食, 刺激皮膚、眼
2% 澱粉溶液	starch	N/A	N/A	白色固體	N/A	輕微皮膚刺激
0.0050M 硫代硫酸鈉	sodium thiosulfate	Na ₂ S ₂ O ₃	248.18	無色固體	1.74(20°C)	勿吞食, 輕微刺激皮膚、眼睛

ref: Merck 缺化性?

編頁碼

II. 實驗紀錄：簡要、詳細、條列式記錄

實驗紀錄：
 a. 反應物與產物之狀態與顏色。
 b. 進行實驗之條件。
 c. 操作過程。
 d. 反應過程顏色、沉澱、吸熱放熱等變化。

- a. 反應物與產物之狀態與顏色。
- b. 進行實驗之條件。
- c. 操作過程。
- d. 反應過程顏色、沉澱、吸熱放熱等變化。
- e. 數據：參考實驗課本的格式或自創，但記錄整齊。

步驟：
 以簡單清楚
 流程圖表示，繪製裝置圖

<p>四. 步驟. 洗淨. +5 1. 洗淨. 烘乾 圓 x 10 \xrightarrow{rt} 運行(表中) 計時試驗</p>	<p>五. 觀察.</p>
<p>① 觀察紀錄可更多 ② 再多使用一張紙 ③ 要寫出最後配方 定</p> <p>④ 要具體討論. "本實驗 rate = - - - - 指定變色 $\Delta t = 40s$. 實測... ⑤ 完整實驗報告, 要誤差討論.</p>	<p>觀察 ① 反應物 color? ② 產物 color? ③ 反應快慢... +8.</p>
<p>2. 用 10ml 移液管 準確量取 0.20M NaI, 0.20M NaCl, 0.0050M $\text{Na}_2\text{S}_2\text{O}_8$, 2% 於 圓</p>	<p>#3-1: 加成 3ml (看錯別區)</p>
<p>3. 用 5ml 移液管 準確量取 0.10M K_2SO_4, 0.10M $\text{K}_2\text{S}_2\text{O}_8$ 量取時同時計時, 加入後 圓 並搖晃 20s</p>	<p>$\Delta t = 39 - 8 = 31 (s)$</p>
<p>依 #1 混勻再靜置</p>	<p>重新取藥進行 #3</p>
<p>4. 溶液變色 \rightarrow 停止計時 \rightarrow 記錄變色時間</p>	<p>放白紙觀察變色</p>
<p>5. 用 #1 配方 重複實驗 (同一條件若 Δt 差異 $> 3s \rightarrow$ 重做)</p>	<p>反應. 玻璃出現 1(s) 後</p>
<p>6. 用 #2 & #3 分別進行不同初濃度之 重複計時實驗</p>	<p>會深到快看相</p>
<p>7. 計算求 $rate = k[S_2O_8^{2-}]^m [I^-]^n$ 中 k, m, n 值</p>	<p>Complex 越多 ④ 色愈深 拔除吸球後約 5(s) 會</p>
<p>8. 依指定 Δt 代入 7. 求得之速率定律式, 設計取量並量取, 量測 Δt 是 否符合預期</p>	<p>流完; 10(s) 會流完 輕搖 ~ 20(s) 放置. * 不用 (s) 5 s</p>
<p>9. 碘鐘交響曲: 準備一組試劑, 配合交響曲 一同反應, 觀察能否配合變色</p>	<p>5. second 10 s 為單位. 20 s</p>
<p>回收: 廢液: 指定回收桶 ✓ (10.0 ml)</p>	<p>(s) 為 solid 態.</p>

III. 結果報告：

- 數據處理：詳列計算過程，結果需有正確的有效數字及單位。
- 檢討：實驗失敗或誤差大，檢討原因及改進方法。

數據：
參考實驗課本的格式
或自創，記錄整齊。

$t = 40s$

實驗數據結果	0.20 M NaI (mL)	0.20 M NaCl (mL)	0.0050 M Na ₂ S ₂ O ₃	% starch	0.10 M K ₂ S ₂ O ₈	0.10 M K ₂ S ₂ O ₈ rxn start	反應變色時間 (Δt, s)
1. 反應時間測定 #1	2.0	2.0			2.0	2.0	11:41 → 12:47, 13:22 → 16:26 66(s)
#2	2.0	2.0	1.0	1.0	0	4.0	20:36 → 23:10, 20:11 → 20:47 34(s)
#3	4.0	0			2.0	2.0	42-17, 35(s) 14-47, 33(s)

2. 計算各反應物的起始濃度及#1~#3初速率: $rate = \frac{-\Delta[S_2O_8^{2-}]}{\Delta t} = \frac{-\Delta[S_2O_3^{2-}]}{\Delta t}$

#	混合液中反應物起始濃度 (M)			平均時間	初速率
	[S ₂ O ₈ ²⁻]	[S ₂ O ₃ ²⁻]	[I ⁻]	Δt (s)	rate (M/s)
#1	0.00050	0.020	0.040	$\frac{66+64}{2} = 65$	3.8×10^{-6}
#2	0.00050	0.040	0.040	$\frac{34+36}{2} = 35$	7.1×10^{-6}
#3	0.00050	0.020	0.080	$\frac{35+33}{2} = 34$	7.4×10^{-6}

$[S_2O_8^{2-}] = \frac{0.0050 \times 1}{10.0} = 0.00050$ $[S_2O_3^{2-}] = \frac{0.10 \times 2.0}{10.0} = 0.020$; $\frac{0.10 \times 4.0}{10.0} = 0.040$ $[I^-] = \frac{0.20 \times 2.0}{10.0} = 0.040 M$
 $m = \frac{\log \frac{r_2}{r_1}}{\log \frac{[S_2O_8^{2-}]_2}{[S_2O_8^{2-}]_1}} = \frac{\log \frac{7.1}{3.8}}{\log \frac{0.040}{0.020}} = \log(2.0)^m \Rightarrow \log(\frac{65}{35}) = \log(2.0)^m \Rightarrow m = \log(\frac{65}{35}) / \log 2.0 \Rightarrow m = 0.893 \xrightarrow{sf=2} 0.89$
 $n = \frac{\log \frac{r_3}{r_1}}{\log \frac{[I^-]_3}{[I^-]_1}} = \frac{\log \frac{7.4}{3.8}}{\log \frac{0.080}{0.040}} = \log(2.0)^n \Rightarrow \log(\frac{34}{65}) = \log(2.0)^n \Rightarrow n = \log(\frac{65}{34}) / \log 2.0 \Rightarrow n = 0.935 \xrightarrow{sf=2} 0.94$
 $r = k [S_2O_8^{2-}]^m [I^-]^n \Rightarrow$ 將#1代入 $\Rightarrow 3.8 \times 10^{-6} = k (0.00050)^{0.89} (0.040)^{0.94} \Rightarrow k = 2.55 \times 10^3 \xrightarrow{sf=2} 2.6 \times 10^3$
 #2 $\Rightarrow 7.1 \times 10^{-6} = k (0.00050)^{0.89} (0.040)^{0.94} \Rightarrow k = 1.96 \times 10^3 \xrightarrow{sf=2} 2.0 \times 10^3$
 #3 $\Rightarrow 7.4 \times 10^{-6} = k (0.00050)^{0.89} (0.080)^{0.94} \Rightarrow k = 2.12 \times 10^3 \xrightarrow{sf=2} 2.1 \times 10^3$

3. 交響曲: If $\Delta t > \Delta t_3 \Rightarrow$ 修改#1 $\Rightarrow \frac{rate_2}{rate_1} = \frac{\Delta t_2}{\Delta t_1} = \frac{35}{65} = (\frac{2.0}{x})^{0.89} \Rightarrow \log(\frac{35}{65}) = 0.89 \log(\frac{2.0}{x})$
 指定 = 40(s). So ① 配方? ② 實測? ③ $m = 0.89$ 但代 0.9 ④ NaI 用? mL.
 If $\Delta t < \Delta t_3 \Rightarrow$ 修改#3 $\Rightarrow \frac{rate_3}{rate_2} = \frac{\Delta t_3}{\Delta t_2} = \frac{40}{34} = (\frac{2.0}{y})^{0.9} \Rightarrow \log(\frac{40}{34}) = 0.9 \log(\frac{2.0}{y})$
 $\Rightarrow \log(\frac{40}{34}) = 0.07058 \Rightarrow \frac{2.0}{y} = 1.076 \Rightarrow y = 1.70 \rightarrow 1.70 \rightarrow 1.7 (mL) \ \& \ SO_4^{2-} = 2.3 mL$
 0 誤差討論

數據處理：
詳列計算過程。
注意單位。
注意有效數字的取捨。

誤差討論及操作檢討：
與理論值比較，算出誤差百分比。
實驗失敗或誤差大，檢討原因及改進方法。
需有具體結論！

General Chemistry Experiment "Brief Version Report" Example

I. Prelab Report:

- Objective: Summarize the goal concisely.
- Principles: Indicate relevant theories and chemical reactions.
- Chemicals: Tabulate the physical and chemical properties as well as the toxicity (use The Merck Index and SDS).
- Procedures: Use a flow chart and cartoon to list the crucial operations, and draw the apparatus in this experiment (you can first watch the lab demo videos online).



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32/35

Group No. :
Name :
Dept. :
Student ID :

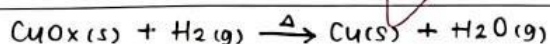
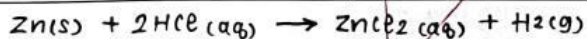
E1. DETERMINATION OF THE CHEMICAL FORMULA OF A COMPOUND

I. Objective : The purpose of this experiment is to investigate the empirical formula of copper oxide using analytical method.

Objective: Summarize the goal concisely.

II. Principles :

- 5/5
- There are two methods to determine the chemical formula of a compound, which are :
- Analytical method: the compound is broken down into its elements, then the relative amount and the ratio of its element can be determined.
 - Synthetical method: an amount of weight for element A is reacted completely with an excess of element B to find a fixed amount of the compound, then the amount of element B and the mole ratio of A and B can be found
- In this experiment, hydrogen gas will be created through the mixing of zinc granules and hydrochloric acid, then use the hydrogen gas to reduce copper oxide to elemental copper upon heating, and also determine the empirical formula of copper oxide and analyze its mass content of copper in the sample.



Principles: Indicate relevant theories and chemical reactions.

Page number

Chemicals:

- Tabulate the name, chemical formula, formula weight, physical and chemical properties as well as the toxicity.
- List your references, such as The Merck Index online, SDS *et. al.*

III. Chemicals:

NAME	FORMULA	MOLECULAR WEIGHT (g/mol)	DENSITY (g/cm ³)	BP/MP (°C)	SOLUBILITY (g/L)	PHYSICAL PROPERTIES	CHEMICAL PROPERTIES	TOXICITY
Zinc granules (鋅)	Zn	65.38	7.14	907/419	soluble in acids alkalies	shiny, silvery gray metal	insoluble in water, strong reducing agent	relatively non-toxic, over toxicity causes nausea
Copper(II)oxide (氧化銅)	CuO	79.55	6.32	2000/1326	soluble in acids	dark colored powder, granules	compound is canonicalized; insoluble in water	causes skin and eye irritation
calcium chloride (氯化鈣)	CaCl ₂	110.98	2.15	1935/782	745	white to off-white solid, odorless	absorbs moisture forms, dissolves in absorbed moisture	toxic, causes eye, skin, respiratory irritation
6M Hydrochloric acid (鹽酸)	HCl	36.46	1.05	-85.1/-114.2	823(0°C) 561(60°C)	colorless liquid, pungent odor	corrosive (pH=0)	toxic, causes eye damage, nausea, headache

95% ethanol (乙醇)	CH ₃ CH ₂ OH	46.07	0.79	78.2/-114.1	≤100	colorless liquid, pungent odor	flammable, compound is canonicalized	toxic, can cause poisoning, pain, internal bleeding
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Source: Pubchem, wikipedia, Experiments in General Chemistry (3rd edition, NTU Press)

II. Lab Notes:

- Leave ~1/3 page blank space next to the procedure section for recording results and your observations during the experiment.
- Use ball pen and avoid correction tapes (only the final reports need to be clean and legible).

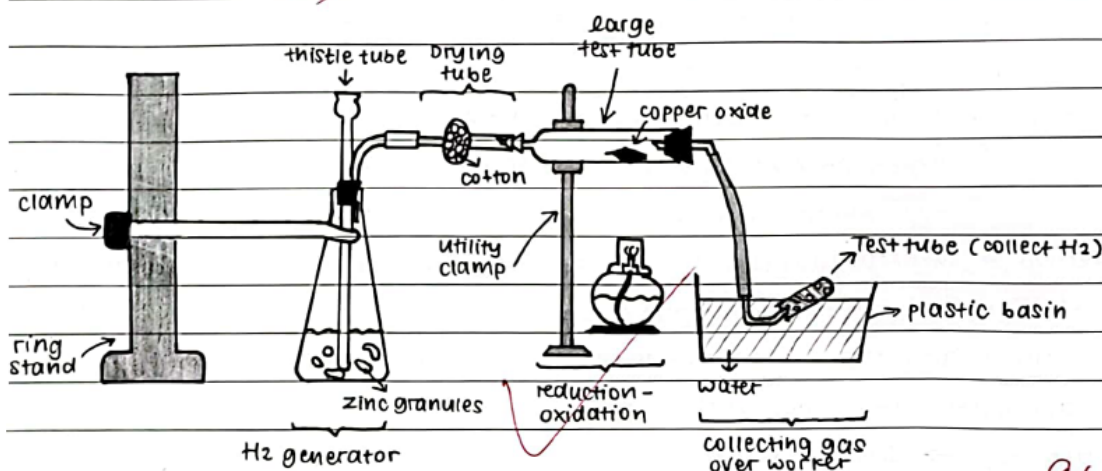
Procedures:

Use a flow chart and cartoon to list the crucial operations, and draw the apparatus in this experiment.

Observation:

- The appearance and color of reactants and products.
- Conditions for actual experimentation.
- Operation procedures.
- Changes in solution, i.e. color, precipitation, endothermic, exothermic, gas evolution, etc.
- Reaction rate, fast or slow...

IV. Procedures : 5/5



- Wash large test tube, dry tube, and let cool.
- Fill drying tube with $CaCl_2$; stuff both ends with just enough cotton to prevent $CaCl_2$ from falling out.
- Measure and record weight of large test tube; add 1-1.2g copper oxide to middle of tube and measure weight (w_2).
- Measure 15 gram zinc granules to erlenmeyer flask and use 100 ml beaker to take 20 ml of 6M HCl.
- Adjust thistle tube such that end of glass pipe nearly touches bottom of flask.
 - Wet rubber tube with water before connecting it to drying tube.
- Fill 10 test tubes with water and invert and immerse into plastic basin filled with $\frac{2}{3}$ water to collect hydrogen gas.
- Pour 20 ml of 6M HCl (aq) into thistle tube; end of

V. observation : 9/10

- After pouring HCl, the Zn fizzed up and erlenmeyer flask became cloudy.
- Rubber tube in the water started bubbling
- The rubber tube has to be tight, so it produces more bubbles.

• Sound of small test tubes:

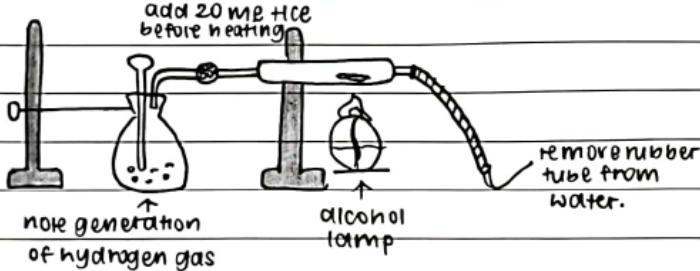
1. X	6. X	(✓ : popped) (X : didn't pop)
2. X	7. X	
3. ✓ loud	8. X	
4. X	9. ✓ subtle	
5. ✓ loud	10. X	

this tube should be immerse in solution and rubber tube not twisted, and gas flow is free from obstruction.

⑧ Collect hydrogen gas over water; bring flame to opening of inverted tube; when there's a loud "pop" sound, continue collecting gas until sound quiets.

⑨ Before heating, remove rubber tube from water; make sure it's not twisted and gas flow is free from obstruction; add another 20 ml of 6M HCl into thistle tube.

⑩ Heat large test tube containing copper oxide with an alcohol lamp until no visible changes in content and no visible moisture in tube; let system cool down



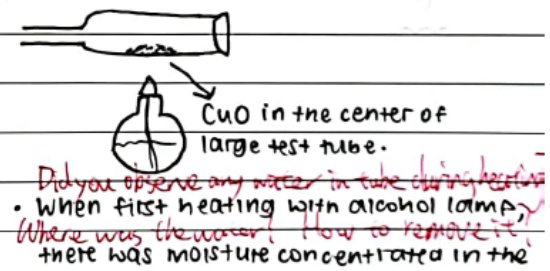
⑪ In the cooling process, maintain connection of apparatus and supply of hydrogen gas to the large test tube.

⑫ Dispose used cotton wool, CuO, unreacted Zn granules (rinsed), HCl (aq), and the produced copper.

• When collecting hydrogen gas, not all test tubes popped; only test tube 3, 5, and 9 that popped. ^{Test tube 3 and 5} "pop" sound is loud, but test tube 9 "pop" sound is subtle.

Appearance of:

- Zinc granules
 - before: solid
 - after: oxidized
- CuO
 - before: white ^{appearance?}
 - after: yellowish-white (doesn't really change)



end of large test tube.

- After 7 minutes of heating, we add 20 ml HCl again into flask.
- After 15 minutes of heating, there's slight color changes at the center part.
- After 18 minutes of heating, 2/3 part change color, and after 20 minutes, 4/5 part change
- After 25 minutes of heating, all of the copper has changed into red, but some are still brown
- After 27 minutes of heating, all are completely red.

CuO

- before: black; powder
- after: red; powder but a little rubbery.

III. Final Report:

- Observations on the reserved blank space in the prelab report.
- Record raw data with units.
- Complete the data analysis, and the required calculations in the lab manual.
- Conclusion, such as: According to the analysis of the....., it shows that.....

Experiment Data and Results:

1) Weight of empty test tube (W_1)	: <u>40.8268 g</u>
2) Weight of test tube and copper oxide (W_2)	: <u>41.9316 g</u>
3) Weight of copper oxide ($W_2 - W_1$)	: <u>$(41.9316 - 40.8268) \text{ g} = 1.1048 \text{ g}$</u>
4) Weight of test tube and copper (W_3)	: <u>41.7146 g</u>
5) Weight of copper ($W_3 - W_1$)	: <u>$(41.7146 - 40.8268) \text{ g} = 0.8878 \text{ g}$</u>
6) Weight of oxygen ($W_2 - W_3$)	: <u>$(1.1048 - 0.8878) \text{ g} = 0.2170 \text{ g}$</u>
7) Empirical formula of copper oxide	: <u>CuO</u>
↳ Calculation:	$n_{\text{Cu}} : n_{\text{O}}$
$(n = \frac{g}{M_r})$	$\frac{0.8878}{63.55} : \frac{0.2170}{16.00}$
	$0.01397 : 0.01356 = 1.030 : 1.000$
	$\frac{1}{1.000} : \frac{1.030}{1.000} \rightarrow \approx 1 : 1$

3/3

Conclusion

In this experiment, we use 16.70 gram of Zn and 20 ml of HCl to produce the hydrogen gas, which resulting in only 3 of 10 test tubes that popped. (Hydrogen and air moisture makes it popped.)

We used the hydrogen to reduce 1.1048g CuO during heating.

After heating and cooling process, we got 0.8878 g and 0.01397 mole of copper (Cu), and also 0.2170g and 0.01356 mole of oxygen. ($n_{\text{Cu}} : n_{\text{O}} \approx 1 : 1$)

So, based on the experiment, the empirical formula of copper oxide is CuO.

Signed by TA

General Chemistry Experiment "Full Version Report" Example

I. Prelab Report:

- Objective:** Summarize the goal concisely.
- Principles:** Indicate relevant theories and chemical reactions.
- Chemicals:** Tabulate the physical and chemical properties as well as the toxicity (use The Merck Index and SDS).
- Procedures:** Use a flow chart and cartoon to list the crucial operations, and draw the apparatus in this experiment (you can first watch the lab demo videos online).

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Exp 5 Heat of Reactions

Group No.:
Name:
Dept.:
Student ID

Objective: Summarize the goal concisely.

Objective: Determine the heat of neutralization and solution by home-built calorimeter and use Hess' law to calculate the molar enthalpy of formation of MgO.

Principles: Indicate relevant theories and chemical reactions.

Principles I The enthalpy of reaction and calorimeter

If the reaction occurs at constant pressure, the change of energy is called enthalpy (H) change or heat of reaction, denoted by ΔH ($\Delta H < 0$: exothermic) $\Delta H > 0$: endothermic

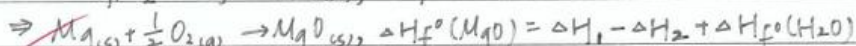
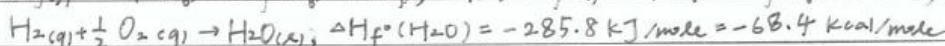
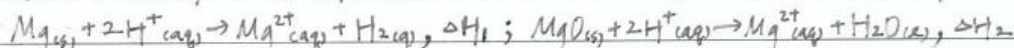
In this exp, we will set up the calorimeter by Styrofoam cups as an adiabatic system. $q_{\text{system}} = 0$

$$\Delta H = -(q_{\text{cal}} + q_{\text{aq}}) = -(C \cdot \Delta T + m_{\text{aq}} \cdot S \cdot \Delta T) \quad \text{Temperature}$$

C refers to the heat capacity, the amount of energy absorbed or released in changing its 1°C , which can be determined by: $q_{\text{c}} = q_{\text{h}} + q_{\text{cal}} = 0$. i.e. $(M_{\text{c}} \cdot S \cdot \Delta T_{\text{c}}) + (m_{\text{h}} \cdot S \cdot \Delta T_{\text{h}}) + (C \cdot \Delta T_{\text{cal}}) = 0$ $C = \frac{q}{\Delta T}$ wait!

II Application of Hess' law - Enthalpy of formation of magnesium oxide

The enthalpy of a chemical reaction is path-independent and is equal to the sum of enthalpies of the contributing steps (Hess' law). Hence, we can calculate the enthalpy of formation of magnesium oxide by the reaction and enthalpies we will measure in this exp.



Page number

Chemicals:

- Tabulate the name, chemical formula, formula weight, physical and chemical properties as well as the toxicity.
- List your references, such as The Merck Index online, SDS *et. al.*

Chemicals				
Name	Chemical formula	Density (g/cm ³)	Appearance	Toxicity and hazards
	Molecular weight	m.p./b.p (°C)	Water solubility (g/100mL H ₂ O, 25°C)	
Magnesium strips	Mg 24.3050	1.738 651/1100	silvery-white, close-packed metal; reacts (produce H ₂)	flammable solid; substance and mixture which, in contact with water, emits flammable gas
Magnesium oxide	MgO 40.31	3.6 2852/3600	white, very fine, odorless powder; practically insoluble, reacts	skin/eye damage/irritation; corrosive to metals; respiratory irritation
ammonium chloride	NH ₄ Cl 53.49	1.5274 338/520	colorless, odorless crystals or white, granular powder; 28.3	acute oral toxicity; serious eye damage/irritation
1.0 M sodium hydroxide	NaOH 40.00	1.043 318/1388	fused solid with crystalline fracture; 111	strong base; serious eye damage; skin corrosion, severe skin burn
1.0 M hydrochloric acid	HCl 36.46	1.098 -59/108	fumes in air, transparent liquid (could be yellow); 2 for 1 g	strong acid (pH < 0); serious eye damage; skin corrosion; respiratory irritation
1.0 M Acetic acid	CH ₃ COOH 60.05	1.053 16.7/118	colorless liquid (vinegar-like); miscible	flammable liquid; skin corrosion, irritation; eye damage, irritation

Ref.: Merck Index Online. Safety Data Sheet (Fisher Scientific)

II. Lab Notes:

- Leave ~1/3 page blank space next to the procedure section for recording results and your observations during the experiment.
- Use ball pen and avoid correction tapes (only the final reports need to be clean and legible).

Procedures:

Use a flow chart and cartoon to list the crucial operations, and draw the apparatus in this experiment.

Observation:

- The appearance and color of reactants and products.
- Conditions for actual experiments.
- Operation procedures.
- Changes in solution, i.e. color, precipitation, endothermic, exothermic, gas evolution, etc.
- Reaction rate, fast or slow...

100

Procedures (notes)

1. **2 styrofoam cups** wait 3 mins then record the T_1

alorimeter plastic lid
foam beaker 50.0 ml DI water
by graduated cylinder Thermometer

2. Measure exactly 50.0 ml warm water (fume hood) (10-15°C higher than T_0). Stay 3 mins. til it reaches the equilibrium. Record T_2

Always record the T_p of sample in the middle section; do not touch the cylinder

Rinse the thermoprobe with tap water after measuring warm stuffs

3. Insert the thermoprobe. Swirl the cal. Record T_p at intervals

Wash the thermoprobe and cal after each use to get rid of the residues (and wipe dry) by dispenser

4. Place (50.0 ml of 1.0 M HCl) into cal. Record T_p .

Place 50.0 ml of 1.0 M NaOH into a graduated cylinder. Record T_p .

Pour NaOH into cal. Close the lid. Insert the thermoprobe. Mix Sol.

5. Repeat step 4. But cal \rightarrow 1.0 M CH_3COOH

6. Repeat step 1. Weigh ca 3g $\text{NH}_4\text{Cl}_{(s)}$ and record its exact weight. Add NH_4Cl into the cal. Close the lid. Insert thermoprobe. Swirl the cal (must dissolve completely) continuously.

* To get equil. temp.:

Max for exothermic reaction and min for endothermic reaction

A. Cal

$t(s)$	1	2	...
$T_p(^\circ\text{C})$	30.0	30.0	...

(2nd) ... \downarrow redo

B. HCl + NaOH (Exothermic)

$t(s)$	0	5	10	15	20	25
$T_p(^\circ\text{C})$	24.6	29.6	31.1	31.0	31.0	31.1

The cups are warmer after this.

C. CH_3COOH + NaOH (Exothermic)

$t(s)$	0	5	10	15	20	25
$T_p(^\circ\text{C})$	24.4	29.9	31.2	31.1	31.1	31.0

Cups = warmer

D. dissolve ammonium chloride (white crystal) (endothermic)

$t(s)$	0	5	10	15	20	25	30	35
$T_p(^\circ\text{C})$	25.1	24.3	23.0	22.5	21.2	21.1	21.1	21.1

There is no ppt in the solution \rightarrow completely dissolved

B.C.E.F	0	H ₂	E. magnesium (silver colored strip) (exo.)								
7. Repeat Step 6. But ² Ca ⇒ exactly 100.0 ml of 1.0M HCl(aq)			t(s)	0	10	20	30	40	50	60	7
² Weigh c.a. 0.2 g Mg : ▲ (s-lr reactions must be mixed and reacted completely)			Tp(°)	24.6	26.5	26.6	27.2	28.8	29.9	32.9	32.
8. Repeat Step 7. But ² c.a. 2.7 g MgO : ▲			反應時有氣泡, 也有三聲, 氣泡不停冒出 (1 min 後蒸) 反應後溫度平針上有氣泡 (H ₂) soln								
9. Assume: ρ solution density 1.0 g/cm ³ (=water). Do the calculation Specific heat 1.0 cal/g.°C			F. magnesium oxide (white/pink powder) (exo.)								
			t(s)	0	5	10	15	20	25	30	35
			Tp(°)	24.5	30.1	30.8	30.9	30.9	30.8	31.0	31.2
10. Salt solutions resulting from acid-base neutralization to sink. Clean the styrofoam cups and plastic lid for reuse.			t(s)	40	50	sl. pink & yellow					
			tp(°)	31.4	31.1	29.5					

Signed by ATA

Signed by TA

III. Final Report:

- Observations on the reserved blank space in the prelab report.
- Record raw data with units.
- Complete the data analysis, and the required calculations in the lab manual.
- Conclusion, such as: According to the analysis of the....., it shows that.....

Calculation and results

HEAT OF REACTIONS

$$* | C_A | = 4.184 \text{ kJ}$$

I. Experimental Data and Results (show all calculations)

1. Heat capacity of calorimeter

second exp

50.0	Volume of cold water (V_1 , mL)	50.0	Volume of hot water (V_2 , mL)	50.0
24.8	Temp. of cold water (T_1 , °C)	24.0	Temp. of hot water (T_2 , °C)	37.2
30.5	Equilibrium temp. (T_3 , °C)	30.0	Heat capacity (C, cal/°C)	10.0

$$V_1 \times 1.0 \text{ g/mL} \times 1.0 \text{ cal/g} \cdot \text{°C} \times (T_3 - T_1) + C \times (T_3 - T_1) = V_2 \times 1.0 \text{ g/mL} \times 1.0 \text{ cal/g} \cdot \text{°C} \times (T_2 - T_3)$$

$$(50.0)(1.0)(30.0 - 24.0) + C(30.0 - 24.0) = (50.0)(1.0)(37.2 - 30.0)$$

$$\Rightarrow (6.0)C = (50.0)(7.2 - 6.0) \Rightarrow C = 10.0$$

$$(5.7)C = (50.0)(5.3 - 5.7) \Rightarrow C = 5.3$$

2. Heat of neutralization: HCl + NaOH

Volume of 1.0 M HCl (mL)	50.0	Volume of 1.0 M NaOH (mL)	50.0
Initial temp. of HCl (°C)	24.6	Initial temp. of NaOH (°C)	24.6
Eqm. temp. after reaction (°C)	31.1	Heat evolved (cal)	-6.8×10^2
Molar heat of neutralization (kJ/mol)	-57		

$$\Delta H = -[(5.3)(31.1 - 24.6) + (50.0)(1.0)(31.1 - 24.6) + (50.0)(1.0)(31.1 - 24.6)]$$

$$= -684.45 \Rightarrow -6.8 \times 10^2 \text{ (cal)}$$

$$\frac{-6.8 \times 10^2 \text{ (cal)}}{(50.0 \times 10^{-3} \text{ L}) \times (1.0 \text{ mol/L})} \times \frac{4.184 \text{ (kJ)}}{1000 \text{ (cal)}} \Rightarrow -57 \text{ kJ/mol}$$

3. Heat of neutralization: CH₃COOH + NaOH

Volume of 1.0 M CH ₃ COOH (mL)	50.0	Volume of 1.0 M NaOH (mL)	50.0
Initial temp. of CH ₃ COOH (°C)	24.4	Initial temp. of NaOH (°C)	24.6
Eqm. temp. after reaction (°C)	31.2	Heat evolved (cal)	-7.1×10^2
Molar heat of neutralization (kJ/mol)	-59		

$$\Delta H = -[(5.3)(31.2 - 24.4) + (50.0)(1.0)(31.2 - 24.4) + (50.0)(1.0)(31.2 - 24.6)]$$

$$= -706.04 \Rightarrow -7.1 \times 10^2 \text{ (cal)}$$

$$\frac{-7.1 \times 10^2 \text{ (cal)}}{(50.0 \times 10^{-3} \text{ L}) \times (1.0 \text{ mol/L})} \times \frac{4.184 \text{ (kJ)}}{1000 \text{ (cal)}} \Rightarrow -59 \text{ kJ/mol}$$

$$\frac{3.00 \text{ (g)}}{53.49} = 0.05608... \Rightarrow 5.61 \times 10^{-2}$$

4. Heat of solution: ammonium chloride $M_{H_4Cl} = 53.49$

Mass of NH ₄ Cl (g)	3.00	Volume of water (mL)	50.0
No. of moles of NH ₄ Cl	5.61×10^{-2}	Initial temp. of water (°C)	25.1
Eqm. temp. after reaction (°C)	21.1	Heat evolved (cal)	2.3×10^2
Molar heat of solution (kJ/mol)	17		

$$\Delta H = -[(5.3)(21.1 - 25.1) + (3.00 + 50.0 \frac{1.0}{\text{cm}^3}) (21.1 - 25.1)]$$

$$= 233.2 \Rightarrow 2.3 \times 10^2 \text{ (cal)}$$

5. Enthalpy of formation: magnesium oxide

$$\frac{2.3 \times 10^2}{5.61 \times 10^{-2} \text{ (mol)}} \times \frac{4.184 \text{ (kJ)}}{1000 \text{ (cal)}} \Rightarrow 17 \text{ kJ/mol}$$

$Mg = 24.305$
 $\frac{0.20 \text{ (g)}}{24.305} \Rightarrow 8.2 \times 10^{-3}$

Mass of magnesium strips (g)	0.20	Mass of magnesium oxide (g)	0.63	$MgO = 40.3$
No. of moles of Mg	8.2×10^{-3}	No. of moles of MgO	1.6×10^{-2}	$\frac{0.63}{40.3} \Rightarrow 1.6 \times 10^{-2}$
Volume of 1.0 M HCl (mL)	100.0	Volume of 1.0 M HCl (mL)	100.0	
Initial temp. of 1.0 M HCl (°C)	24.5	Initial temp. of 1.0 M HCl (°C)	24.5	
No. of moles of H ⁺	0.10	No. of moles of H ⁺	0.10	同左
Eqm. temp. after reaction (°C)	32.9	Eqm. temp. after reaction (°C)	31.4	
Heat evolved (cal)	-8.9×10^2	Heat evolved (cal)	-7.4×10^2	
Molar heat of reaction (kJ/mol)	-4.5×10^2	Molar heat of reaction (kJ/mol)	-1.9×10^2	

$\text{mole} = M \times V(L)$
 $= (1.0) \times 0.1000$
 $\Rightarrow 0.10$

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Signed by TA

$$\Delta H = -[(5.3)(32.9 - 24.5) + (0.20 + 100.0 \frac{1.0}{\text{cm}^3}) (32.9 - 24.5)]$$

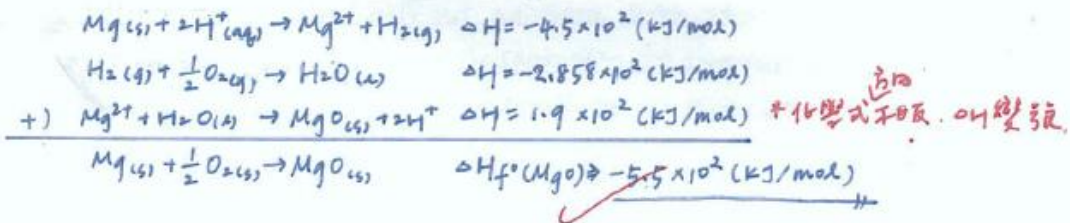
$$= -886.2 \Rightarrow -8.9 \times 10^2 \text{ (cal)}$$

$$\frac{-8.9 \times 10^2 \text{ (cal)}}{8.2 \times 10^{-3} \text{ (mol)}} \times \frac{4.184 \text{ (kJ)}}{1000 \text{ (cal)}} \Rightarrow -4.5 \times 10^2 \text{ kJ/mol}$$

$$\Delta H = -[(5.3)(31.4 - 24.5) + (0.63 + 100.0 \frac{1.0}{\text{cm}^3}) (31.4 - 24.5)]$$

$$= -730.917 \Rightarrow -7.3 \times 10^2 \text{ (cal)}$$

$$\frac{-7.3 \times 10^2 \text{ (cal)}}{1.6 \times 10^{-2} \text{ (mol)}} \times \frac{4.184 \text{ (kJ)}}{1000 \text{ (cal)}} \Rightarrow -1.9 \times 10^2 \text{ kJ/mol}$$



Molar enthalpy of formation of MgO: -5.5×10^2 (kJ/mol)

Error analysis and conclusions

Data analysis and error analysis

We have to calculate the theoretical value of each exp first.

By the Hess' law, we have: $\Delta H^\circ_{\text{rxn}} = \sum n \Delta H^\circ_f \text{ product} - \sum n \Delta H^\circ_f \text{ reactants}$ with n being the stoichiometric coefficients and ΔH°_f being the enthalpy of formation.

exp	ΔH°_f of each compound (kJ/mol)	theoretical value of $\Delta H^\circ_{\text{rxn}}$ (kJ/mol)
2. $\text{HCl}_{(aq)} + \text{NaOH}_{(aq)} \rightarrow \text{NaCl}_{(aq)} + \text{H}_2\text{O}_{(l)}$	(-167.15, -469.15, -407.27, -285.830)	$(-407.27) + (-285.830) - [(-167.15) + (-469.15)] = -56.8$
3. $\text{CH}_3\text{COOH}_{(aq)} + \text{NaOH}_{(aq)} \rightarrow \text{CH}_3\text{COONa}_{(aq)} + \text{H}_2\text{O}_{(l)}$	(-486.34, -469.15, -726.13, -285.830)	$(-726.13) + (-285.830) - [(-486.34) + (-469.15)] = -56.4$
4. $\text{NH}_4\text{Cl}_{(s)} \rightarrow \text{NH}_4^+_{(aq)} + \text{Cl}^-_{(aq)}$	(-314.5, -133.26, -167.08)	$(-133.26) + (-167.08) - (-314.5) = 14.16$
5-1. $\text{Mg}_{(s)} + 2\text{H}^+_{(aq)} \rightarrow \text{Mg}^{2+}_{(aq)} + \text{H}_2_{(g)}$	(0, 0, -467.0, 0)	-467.0 ✓
5-2. $\text{MgO}_{(s)} + 2\text{H}^+_{(aq)} \rightarrow \text{Mg}^{2+}_{(aq)} + \text{H}_2\text{O}_{(l)}$	(-601.6, 0, -467.0, -285.830)	$(-285.830) + (-467.0) - (-601.6) = -151.2$ ✓
5-3. $\text{Mg}_{(s)} + \frac{1}{2}\text{O}_2 \rightarrow \text{MgO}_{(s)}$	(0, 0, -601.6)	-601.6 ✓

Ref: Lange's Handbook of Chemistry - Fifteen Edition *Good to show detail*

Compare the theoretical value of $\Delta H^\circ_{\text{rxn}}$ with the experimental value and calculate its relative error: $\frac{\text{Absolute Error}}{\text{Actual value}} = \frac{\text{Measured value} - \text{expected value}}{\text{Actual value}}$ (expressed in %)

exp	experimental value (kJ/mol)	theoretical value (kJ/mol)	Relative Error
Heat of neutralization $\text{HCl} + \text{NaOH}$ $\text{CH}_3\text{COOH} + \text{NaOH}$	-57 -59	-56.80 -56.47	$\frac{(-57) - (-56.80)}{-56.80} \times 100(\%) = 0.35\%$ $\frac{(-59) - (-56.47)}{-56.47} \times 100(\%) = 4.5\%$
	Heat of solution NH_4Cl	17	14.16
Enthalpy of formation 5-1 5-2 5-3: MgO	-4.5×10^2 -1.9×10^2 -5.5×10^2	-467.0 -151.2 -601.6	$\frac{(-4.5 \times 10^2) - (-467.0)}{-467.0} \times 100(\%) = -3.6\%$ $\frac{(-1.9 \times 10^2) - (-151.2)}{-151.2} \times 100(\%) = 20.0\%$ $\frac{(-5.5 \times 10^2) - (-601.6)}{-601.6} \times 100(\%) = -8.6\%$

Overall, this exp exist some errors (discuss below) but it did roughly determine the heat of reaction.

- (a) First, the calorimeter is absolutely not adiabatic. With styrofoam cups in this exp, we can only roughly separate the reaction and the surrounding. So there must exist some heat loss.
- (b) The relative error of exp: $\text{HCl} + \text{NaOH}$ & $\text{CH}_3\text{COOH} + \text{NaOH}$ are quite small (better close to the true value compared with other data). Maybe that is because it only involve an ^{easy} exothermic reaction of acid-base neutralization and NaOH (strong base) enables it to react completely. Hence the error is lowered. However, the slight ^{positive} error may come from the human error (loss HCl or NaOH during exp, unattended data when recording, the temperature of our hands, et). But it ^{did} quite precisely determine the heat of neutralization.
- (c) The relative error in determining the heat of solution is up to 20%, which is high compared to other data. The main reasons of this significant positive error may derive from: loss of NH_4Cl (Although I observed no ppt. after the solution, there is still ^{some} possibility for those "white" crystals sticking on the surface, the small gaps in the white styrofoam cup), incorrect initial temperature (The weather was chill that day, I might lower the $T_i = 25.1(\%)$ I recorded)
- (d) The relative error of $\Delta H^\circ_{\text{rxn}}$ of $\text{Mg} + \text{HCl}$ is -3.6%, a negative error that is really close to the true value. The error may come from = measurement error ($\text{Mg}_{(s)} + 2\text{H}^+_{(aq)} \rightarrow \text{Mg}^{2+}_{(aq)} + \text{H}_2_{(g)}$). This reaction produces H_2 and I did observe that some little H_2 -bubb stick on the thermometer, which might imply that the device can not very precisely measure the exact temperature of the solution and therefore cause some errors.)