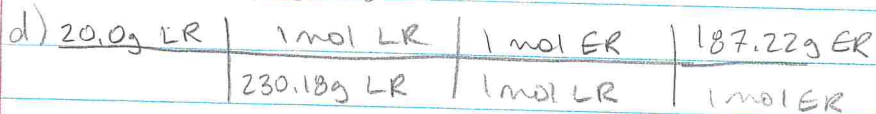
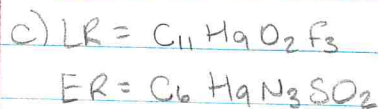
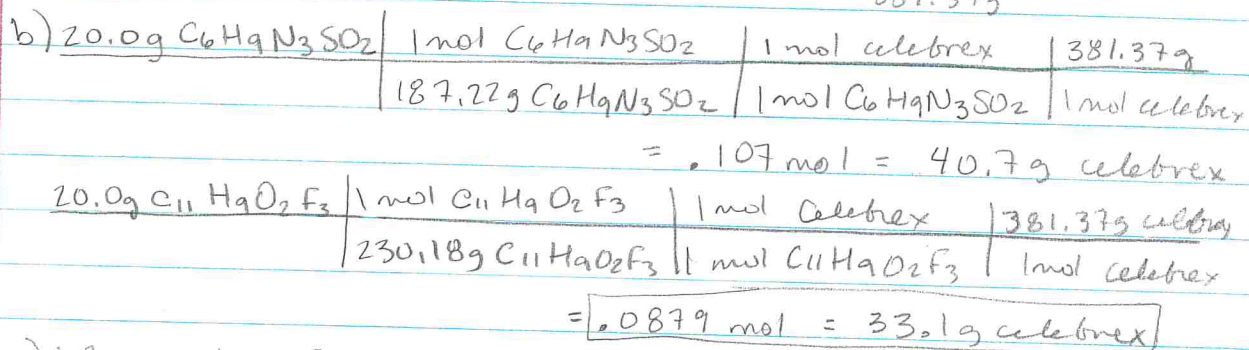
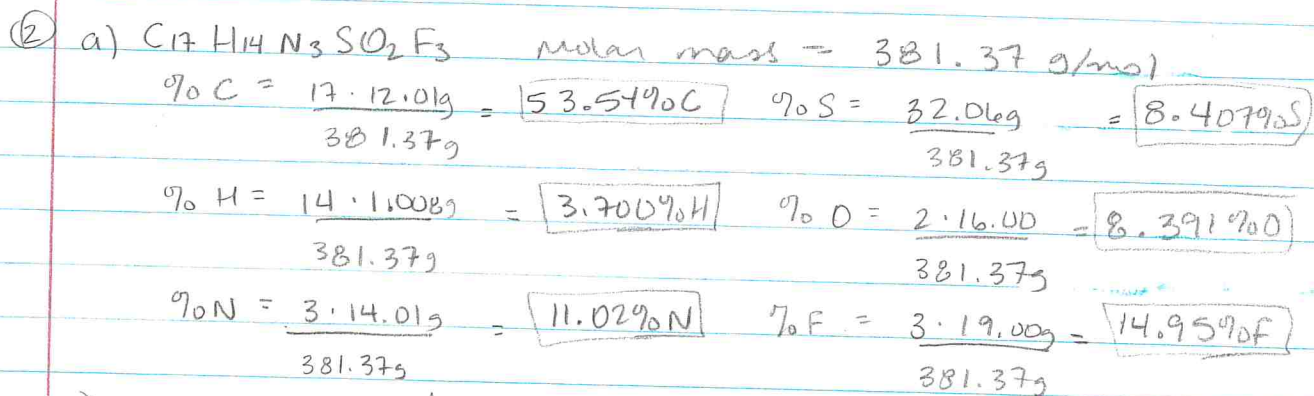
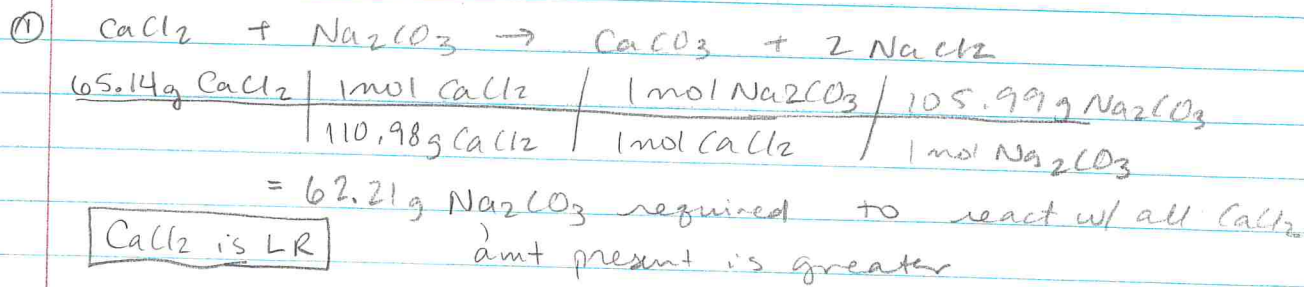


# Unit 4 Study Guide



= 16.3g  $\text{C}_6\text{H}_9\text{N}_3\text{SO}_2$

$20.0\text{g} - 16.3\text{g} = 3.7\text{g } \text{C}_6\text{H}_9\text{N}_3\text{SO}_2$  | 1 mol | = 0.0199 mol  $\text{C}_6\text{H}_9\text{N}_3\text{SO}_2$

187.22g

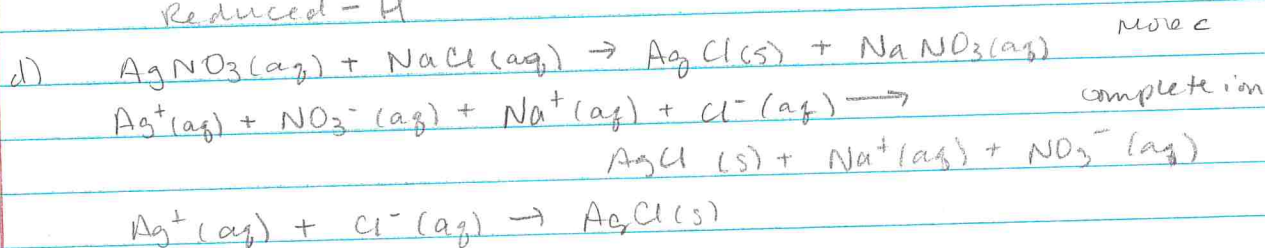
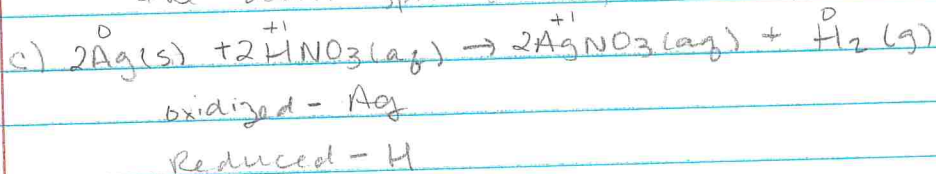


$\% \text{ yield} = \frac{A}{T} = \frac{28.0\text{g}}{33.1\text{g}} \cdot 100 = 84.6\%$

③ a)  $M_1 = 16M$      $M_2 = 6.0M$      $M_1 V_1 = M_2 V_2$   
 $V_1 = ?$      $V_2 = 50. \text{ mL}$      $V_1 = \frac{M_2 V_2}{M_1} = \frac{(6.0)(50. \text{ mL})}{16}$   
 $= \boxed{19 \text{ mL}}$

ii) Add some volume of distilled  $\text{H}_2\text{O}$  to the 50 mL volumetric flask. Use a graduated cylinder to measure 19 mL of the 16 M  $\text{HNO}_3$  stock solution + carefully add it to the  $\text{H}_2\text{O}$  in the v. flask. Add distilled  $\text{H}_2\text{O}$  until the volume of solution is at the line in the flask.

b)  $\text{NaHCO}_3$  is the best option to clean up the spill.  $\text{HCO}_3^-$  can accept a proton ( $\text{H}^+$ ) and neutralize the acid spill during clean up.

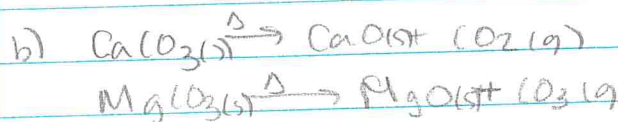


e)  $29.2598 \text{ g} - 28.7210 \text{ g} = .5388 \text{ g AgCl}$

$.5388 \text{ g AgCl}$	$1 \text{ mol AgCl}$	$1 \text{ mol AgNO}_3$	$2 \text{ mol Ag}$	$107.87 \text{ g Ag}$
	$143.32 \text{ g AgCl}$	$1 \text{ mol AgCl}$	$2 \text{ mol AgNO}_3$	$1 \text{ mol Ag}$
		from precip. eqn	From SE eqn	
$=$	$\boxed{.4055 \text{ g Ag}}$			

④ a)  $V = .0750 \text{ L}$      $PV = nRT$   
 $P = 750 \text{ mmHg}$      $n = \frac{PV}{RT} = \frac{(750)(.0750)}{(62.36)(290)} = .0031 \text{ mol CO}_2$   
 $T = 20^\circ\text{C} = 290 \text{ K}$

$.0031 \text{ mol CO}_2$	$44.01 \text{ g}$	$= \boxed{.14 \text{ g CO}_2}$
	$1 \text{ mol}$	



⑤ a) a different precipitate forms of  $PbI_2$

$$b) \frac{150 \text{ g } Pb(NO_3)_2}{331.22 \text{ g}} \times 1 \text{ mol} = 4.53 \cdot 10^{-4} \text{ mol } Pb(NO_3)_2$$

$$\frac{125 \text{ mL}}{1000 \text{ mL}} \times 1 \text{ L} \times 0.100 \text{ mol NaI} = 0.0125 \text{ mol NaI}$$

$$c) \frac{0.0125 \text{ mol NaI}}{2 \text{ mol NaI}} \times 1 \text{ mol } Pb(NO_3)_2 = 0.00625 \text{ mol } Pb(NO_3)_2$$

more than in b

$Pb(NO_3)_2$  is LR

d) all  $NO_3^-$  comes from  $Pb(NO_3)_2$  and is not used up.

$$\frac{4.53 \cdot 10^{-4} \text{ mol } Pb(NO_3)_2}{1 \text{ mol } Pb(NO_3)_2} \times 2 \text{ mol } NO_3^- = 9.06 \cdot 10^{-4} \text{ mol } NO_3^-$$
$$\frac{9.06 \cdot 10^{-4} \text{ mol } NO_3^-}{0.125 \text{ L}} = 0.00723 \text{ M } NO_3^-$$

e) upper right diagram - all  $Pb^{2+}$  is precipitated as  $PbI_2$   $\therefore$  no  $Pb^{2+}$  ions in soln.  $NO_3^-$  is a spectator ion, as is  $Na^+$ , so both are present in soln. NaI is the excess reagent, so excess  $I^-$  ions will be in soln.

⑥  $V = 454 \text{ mL} = 0.454 \text{ L}$

$T = 23.0^\circ\text{C} = 296.2 \text{ K}$

$P_{\text{tot}} = 712 \text{ mmHg}$

$P_{H_2O} = 19.8 \text{ mmHg}$

$M_{Zn} = ?$

$P_{H_2} = 712 \text{ mmHg} - 19.8 \text{ mmHg} = 692 \text{ mmHg}$

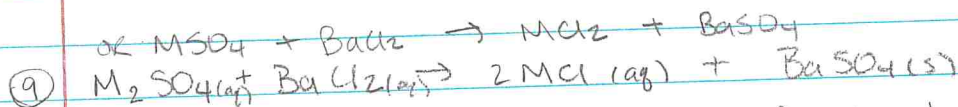
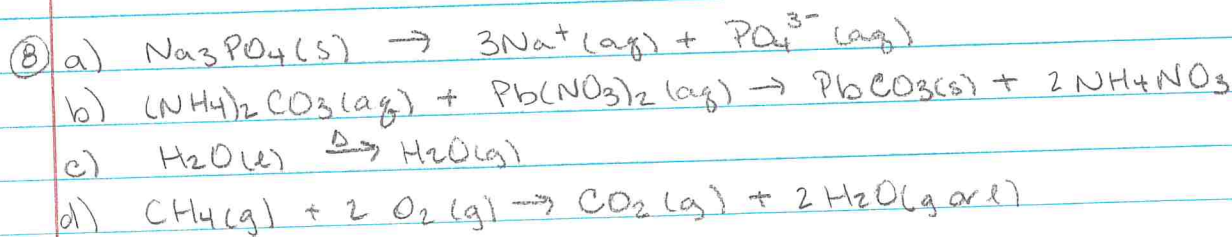
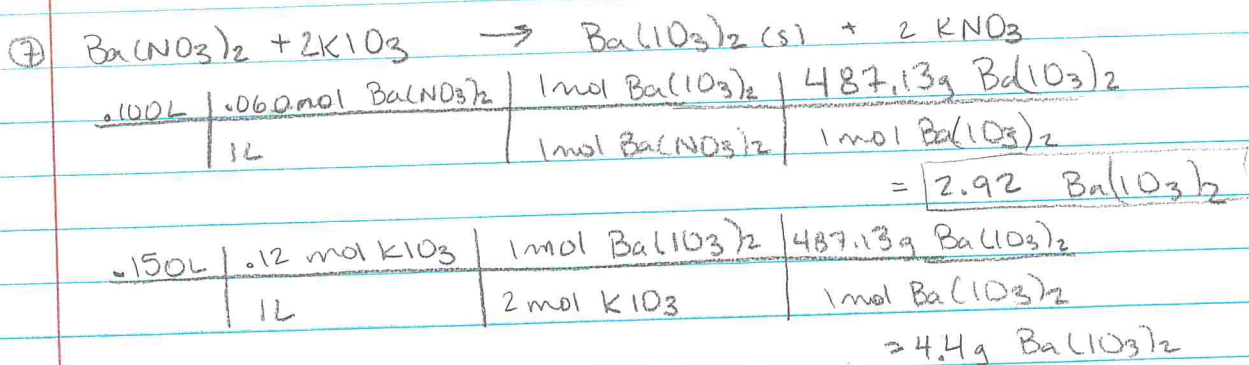
$PV = nRT$

$$n_{H_2} = \frac{PV}{RT} = \frac{(692)(0.454)}{(62.36)(296.2)}$$

$= 0.0170 \text{ mol } H_2$

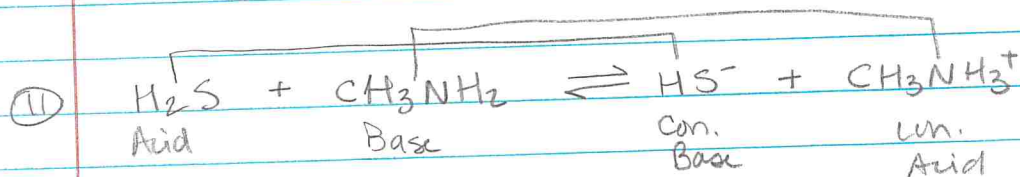
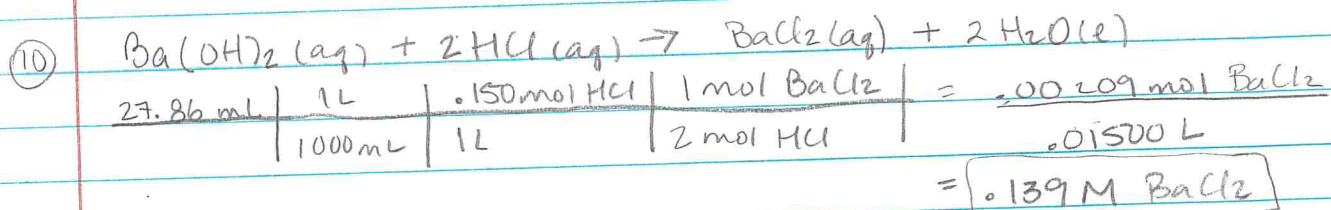
$$\frac{0.0170 \text{ mol } H_2}{1 \text{ mol } H_2} \times \frac{65.38 \text{ g}}{1 \text{ mol } Zn} = 1.06 \text{ g } Zn$$





a) Dissolve the  $\text{M}_2\text{SO}_4$  in  $\text{H}_2\text{O}$ . React w/ aqueous  $\text{BaCl}_2$  to form precipitate. Filter precipitate + dry. Use mass of precipitate ( $\text{BaSO}_4$ ) to calculate moles  $\text{M}_2\text{SO}_4$  that reacted. Use mass + moles  $\text{M}_2\text{SO}_4$  to calculate molar mass.

b) no - mixing  $\text{M}_2\text{SO}_4(aq) + \text{MgCl}_2(aq)$  would not produce a precipitate ( $\text{MgSO}_4$  is soluble)





$$\frac{7.2g H_2O}{18.02g H_2O} \left| \frac{1mol H_2O}{1mol H_2O} \right| \frac{2mol H}{1mol H_2O} = .80 mol H$$

$$\frac{7.2L CO_2}{22.4L} \left| \frac{1mol CO_2}{1mol CO_2} \right| \frac{1mol C}{1mol CO_2} = .32 mol C$$

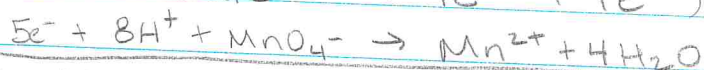
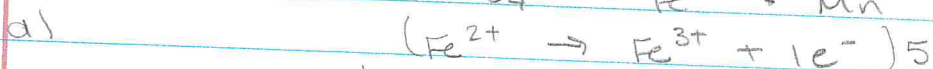
$$\frac{.32 mol C}{.32} = 1 \times 2 \quad \frac{.80 mol H}{.32} = 2.5 \times 2$$



b)  $\frac{7.2g H_2O}{18.02g H_2O} \left| \frac{1mol H_2O}{1mol H_2O} \right| \frac{13mol O_2}{10mol H_2O} \left| \frac{32.00g O_2}{1mol O_2} \right| = 17g O_2$



⑬



b) oxidized: Fe

Reduced: Mn

c)  $\frac{37.93 mL}{1000 mL} \left| \frac{1L}{1L} \right| \frac{.1350 mol MnO_4^-}{1 mol MnO_4^-} \left| \frac{5 mol Fe^{2+}}{1 mol MnO_4^-} \right|$

$= \frac{.02560 mol Fe^{2+}}{.02250 L} = 1.138 M Fe^{2+}$

Practice AP Questions

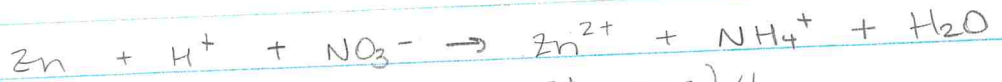
⑩  $\frac{.40 mol Al}{4 mol Al} \left| \frac{2 mol Al_2O_3}{4 mol Al} \right| = \frac{.40 \cdot 2}{4} = .2 mol Al_2O_3$

$\frac{.40 mol O_2}{3 mol O_2} \left| \frac{2 mol Al_2O_3}{3 mol O_2} \right| = \frac{.40 \cdot 2}{3} = \frac{.8}{3} = .27$

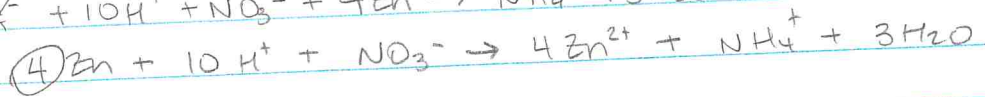
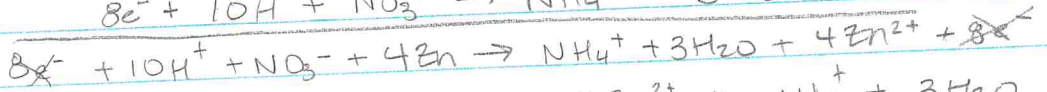
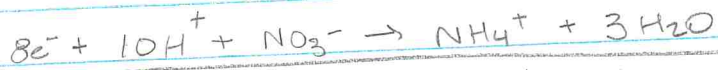
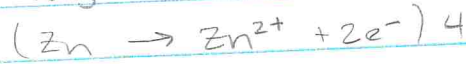


Redox

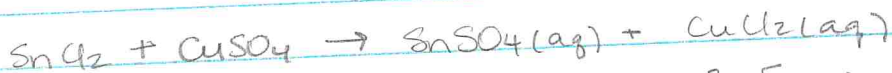
(2)



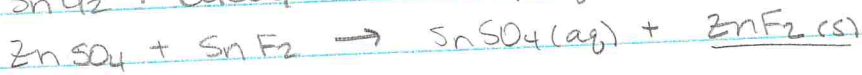
(B)



(3)



(E)



(4)



8.0g N <sub>2</sub> H <sub>4</sub>	1 mol N <sub>2</sub> H <sub>4</sub>	4 mol H <sub>2</sub> O	18.02g H <sub>2</sub> O	= $\frac{8.0 \cdot 18}{32 \cdot 2} = 9$
	32g N <sub>2</sub> H <sub>4</sub>	2 mol N <sub>2</sub> H <sub>4</sub>	1 mol H <sub>2</sub> O	

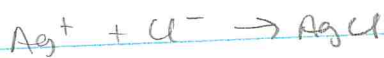
(A)

92g N <sub>2</sub> O <sub>4</sub>	1 mol N <sub>2</sub> O <sub>4</sub>	4 mol H <sub>2</sub> O	18g H <sub>2</sub> O	= $\frac{92 \cdot 4 \cdot 18}{92} = \text{more than } 9$
	92g N <sub>2</sub> O <sub>4</sub>	1 mol N <sub>2</sub> O <sub>4</sub>	1 mol H <sub>2</sub> O	

(5)

1.0 L	.10 mol NaCl	= .10 mol Cl <sup>-</sup>	} .30 mol Cl <sup>-</sup>
	.10 mol CaCl <sub>2</sub>	= .20 mol Cl <sup>-</sup>	

(C)



(6)

100. mL BaCl<sub>2</sub>

.10M BaCl<sub>2</sub>

100. mL H<sub>2</sub>SO<sub>4</sub>

.050M H<sub>2</sub>SO<sub>4</sub>



.1L	.1 mol BaCl <sub>2</sub>	= .01 mol BaCl <sub>2</sub>	= .01 mol Ba <sup>2+</sup>
	1L		

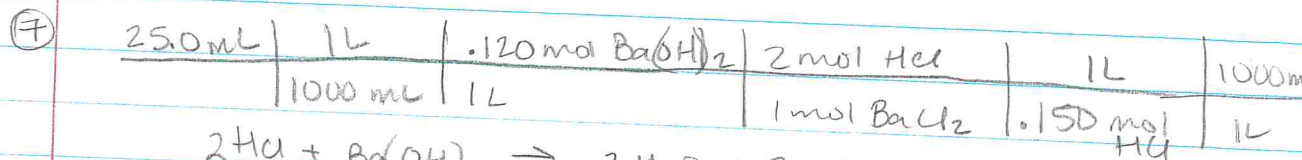
.1L	.05 mol H <sub>2</sub> SO <sub>4</sub>	= .005 mol H <sub>2</sub> SO <sub>4</sub>	= .005 mol SO <sub>4</sub> <sup>2-</sup>
	1L		

(C)

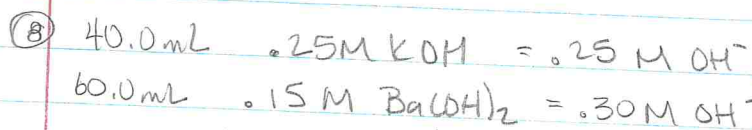
All SO<sub>4</sub><sup>2-</sup> reacts, only .005 mol Ba<sup>2+</sup> reacts, leaving .005 mol Ba<sup>2+</sup>

.005 mol Ba <sup>2+</sup>	=	.025 M
.2 L		





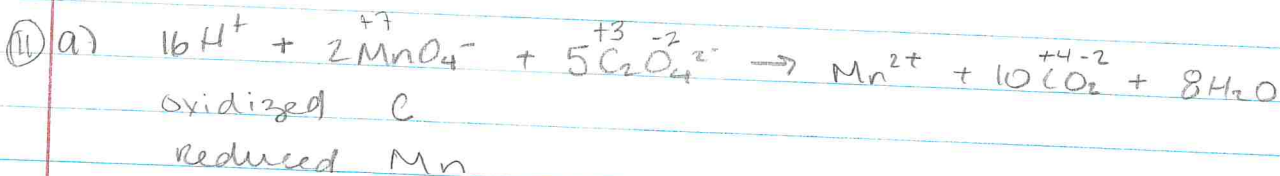
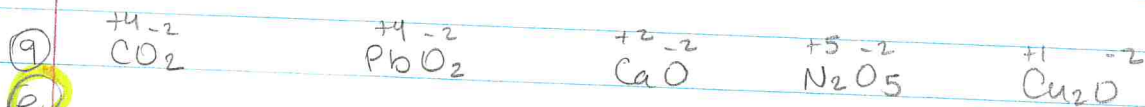
(c) 
$$= \frac{25 \cdot .120 \cdot 2 \cdot 1000}{1000 \cdot .15} = \frac{25 \cdot .120 \cdot 2}{.15} = \frac{8.5 \cdot .15 \cdot 4}{.15} = 5 \cdot 4 \cdot 2 = \textcircled{40}$$



$$[\text{OH}^-] = \frac{(.25 \frac{\text{mol}}{\text{L}} \text{OH}^-)(.04 \text{ L OH}^-) + (.30 \frac{\text{mol}}{\text{L}} \text{OH}^-)(.06 \text{ L OH}^-)}{.1 \text{ L}}$$

(c) 
$$= .25 \cdot .4 + .30 \cdot .6$$
  

$$.1 + .18 = .28 \text{ M}$$



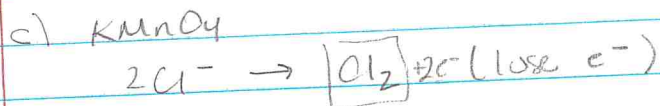
b) 
$$\frac{.01780 \text{ L MnO}_4^-}{1 \text{ L}} \cdot \frac{.0150 \text{ mol MnO}_4^-}{2 \text{ mol MnO}_4^-} \cdot \frac{5 \text{ mol C}_2\text{O}_4^{2-}}{1 \text{ mol C}_2\text{O}_4^{2-}} = .00668$$

$$.00668 \text{ mol C}_2\text{O}_4^{2-} = .00668 \text{ mol BeC}_2\text{O}_4$$

$$\frac{.00668 \text{ mol BeC}_2\text{O}_4}{1 \text{ mol BeC}_2\text{O}_4} \cdot 97.03 \text{ g BeC}_2\text{O}_4 = .648 \text{ g BeC}_2\text{O}_4 \text{ in } 20.0 \text{ mL}$$

$$\frac{.0648 \text{ g BeC}_2\text{O}_4}{20.0 \text{ mL}} = \frac{x \text{ g}}{100. \text{ mL}} \quad x = \boxed{.324 \text{ g pure BeC}_2\text{O}_4}$$

- ⑫ a)  $\text{Pb}(\text{NO}_3)_2 + \text{NaCl} \rightarrow \text{PbCl}_2$   
 b)  $\text{C}_2\text{H}_5\text{OH}$  - this is a molecular compound.  
 The others are ionic + dissociate into ions  
 +  $\therefore$  conduct electricity



- ⑬ a) the Al +  $\text{CuSO}_4$  react. Al is higher on the activity series than Cu and will replace it  
 $2\text{Al} + 3\text{CuSO}_4 \rightarrow \text{Al}_2(\text{SO}_4)_3 + 3\text{Cu}$  (molecular)  
 $2\text{Al} + 3\text{Cu}^{2+} \rightarrow 2\text{Al}^{3+} + 3\text{Cu}$  (net ionic)  
 (I would accept either equation)

b) The T inside the metal container is high, as the steam (gas) is hot. As the container cools, the gas inside cools, lowering the pressure. Eventually, the pressure inside the container is lower than the pressure outside and the container implodes.

c) The rubbing alcohol absorbs heat from your skin, interrupting the intermolecular forces that hold the alcohol molecules together + causing the rubbing alcohol to evaporate

d) The baking soda neutralizes the methanoic acid.



$\text{H}_2\text{CO}_3$  decomposes to  $\text{H}_2\text{O} + \text{CO}_2$





(14)



1.625g

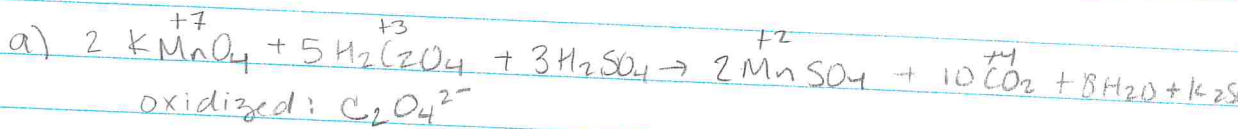
88.43 ml

.102M

$$\frac{.08843 \text{ L} \mid .102 \text{ mol NaOH} \mid 1 \text{ mol Aspirin-H}}{1 \text{ L} \mid \mid 1 \text{ mol NaOH}} = .00902 \text{ mol aspirin}$$

$$\text{Molar mass} = \frac{\text{g}}{\text{mol}} = \frac{1.625 \text{ g}}{.00902 \text{ mol}} = \boxed{180. \text{ g/mol}}$$

(15)



oxidized:  $\text{C}_2\text{O}_4^{2-}$

reduced: Mn

b) the  $\text{MnO}_4^-$  indicates the end point. When the end point has been reached, all of the purple  $\text{MnO}_4^-$  has been reduced to not purple  $\text{Mn}^{2+}$ . Adding additional  $\text{MnO}_4^-$  to the flask will result in a colored solution instead of clear.

c) need to know the mass of oxalic acid in flask and the initial volume + final volume readings on buret

d) Perform the titration, stopping when the flask shows a permanent light pink color.

Use the mass of oxalic acid in the flask to determine the moles of oxalic acid. Use stoichiometry + the balanced equation to calculate the moles of  $\text{KMnO}_4$  that reacted.

Divide the moles of  $\text{KMnO}_4$  by the volume of  $\text{KMnO}_4$  soln added in L to calculate molarity

e)  $[\text{KMnO}_4]$  would not be affected.  $\text{H}_2\text{O}$  is not a reactant +  $\therefore$  does not affect stoich. It is only used to dissolve oxalic acid

