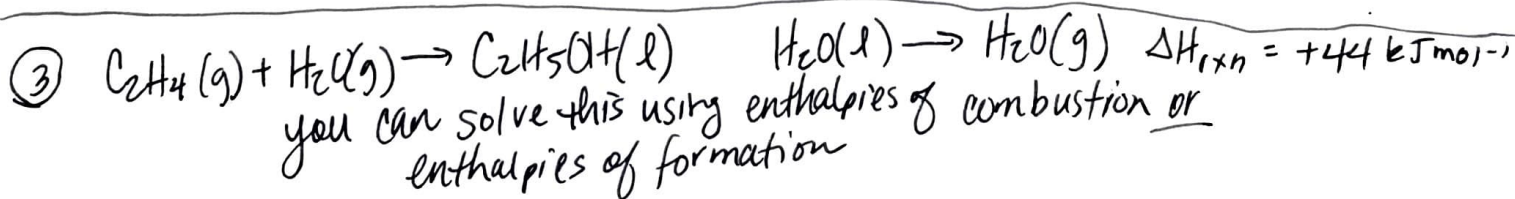
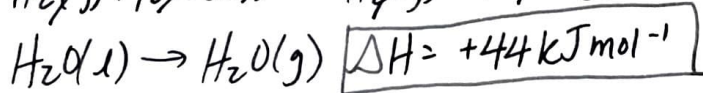
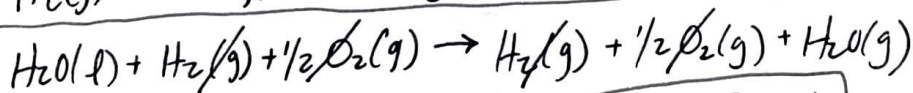
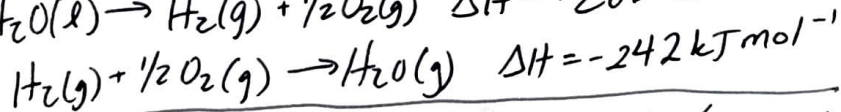
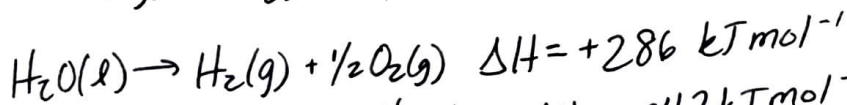
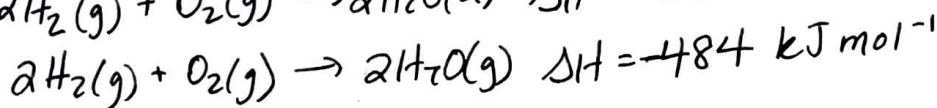
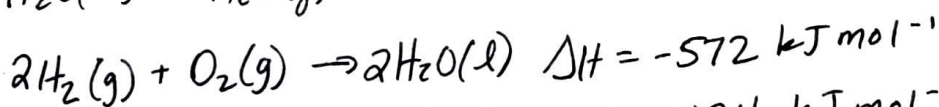
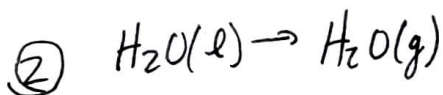
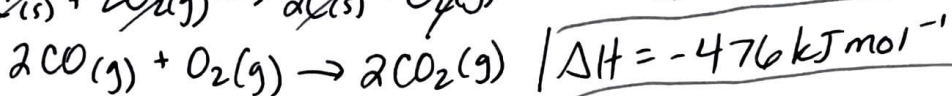
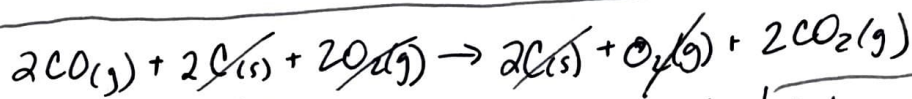
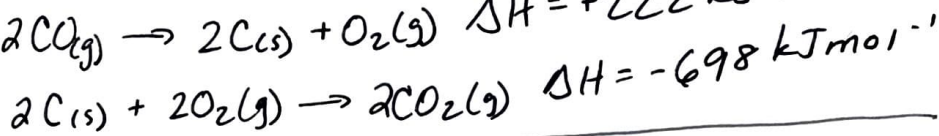
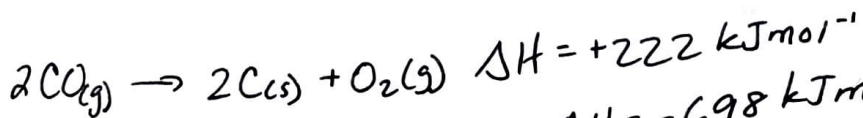
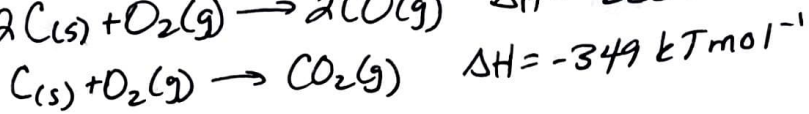
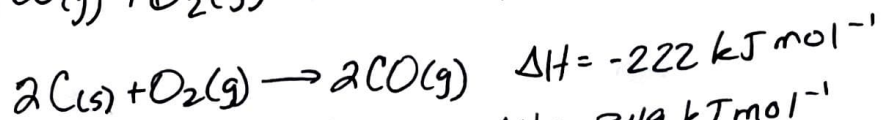
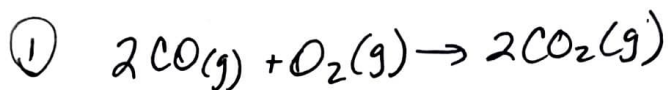
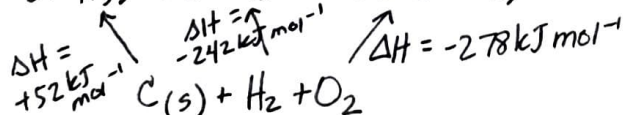
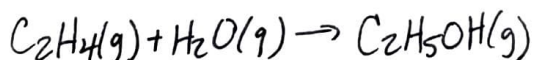


Hess's Law Worksheet (Hess Cycles)

Answer key



using enthalpies of formation:

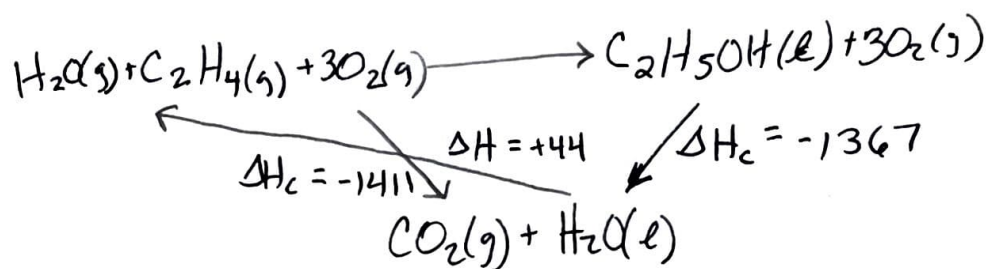
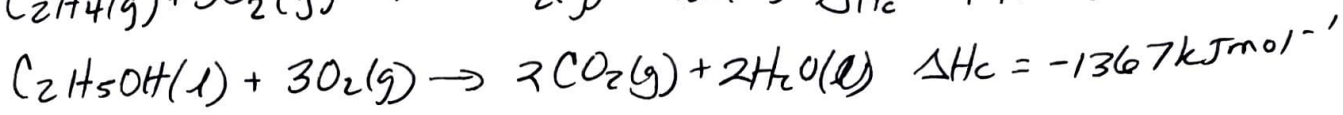
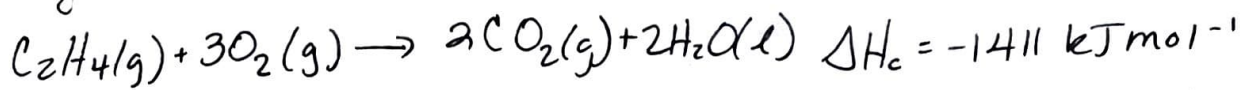


$$\Delta H_{\text{rxn}} = \sum \Delta H_f(\text{products}) - \sum \Delta H_f(\text{reactants})$$

$$\Delta H_{\text{rxn}} = -278 - (-242 + 52)$$

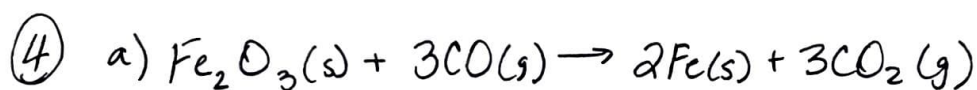
$$= -88 \text{ kJ mol}^{-1}$$

③ Continued
using enthalpies of combustion:



$$\Delta H_{\text{rxn}} = -1411 + 1367 - 44 = \boxed{-88 \text{ kJ mol}^{-1}}$$

either way you slice it,
it is -88 kJ mol^{-1} !



$$\Delta H_f \text{CO}(\text{g}) : -111 \text{ kJ mol}^{-1}$$

$$\Delta H_f \text{CO}_2(\text{g}) : -394 \text{ kJ mol}^{-1}$$

$$\Delta H_f \text{Fe}_2\text{O}_3(\text{s}) : -822 \text{ kJ mol}^{-1}$$

$$\Delta H_{\text{rxn}} = \sum \Delta H_f(\text{products}) - \sum \Delta H_f(\text{reactants})$$

$$= 3(-394) - (-822 + 3(-111))$$

$$= -1182 + 822 + 333 = \boxed{-27 \text{ kJ mol}^{-1}}$$



$$\Delta H_f \text{NH}_3 : -46.11 \text{ kJ mol}^{-1}$$

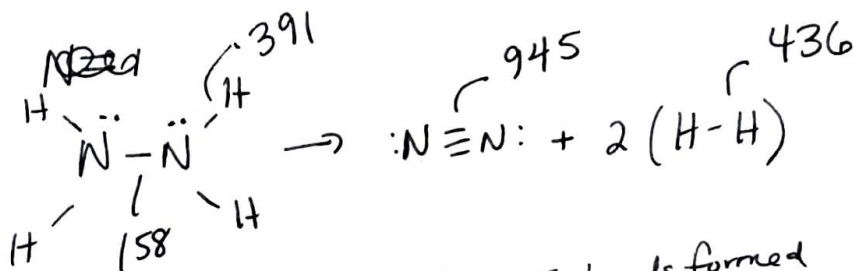
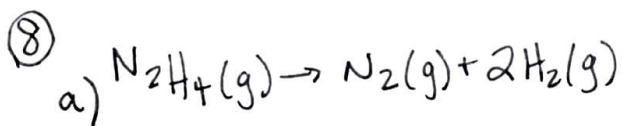
$$\Delta H_f \text{NO} : +90.7 \text{ kJ mol}^{-1}$$

$$\Delta H_f \text{H}_2\text{O}(\text{g}) : -242 \text{ kJ mol}^{-1}$$

$$\Delta H_{\text{rxn}} = (4(90.7) + -242) - (4(-46.11))$$

$$= 120.8 + 184.44$$

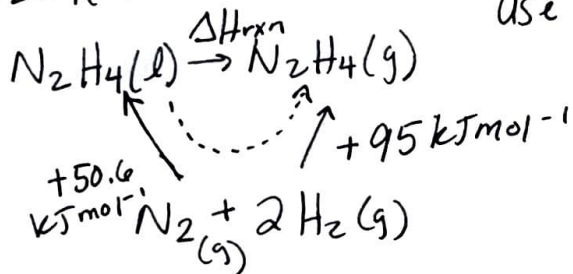
$$= \boxed{+305.24 \text{ kJ mol}^{-1}}$$



$$\begin{aligned}
 \Delta H_{rxn} &= \sum \text{bonds broken} - \sum \text{bonds formed} \\
 &= [158 + 4(391)] - [945 + 2(436)] \\
 &= 1722 - 1817 = \boxed{-95 \text{ kJ mol}^{-1}}
 \end{aligned}$$

$$\Delta H_f N_2H_4(g) : +95 \text{ kJ mol}^{-1}$$

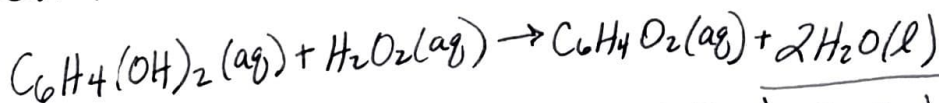
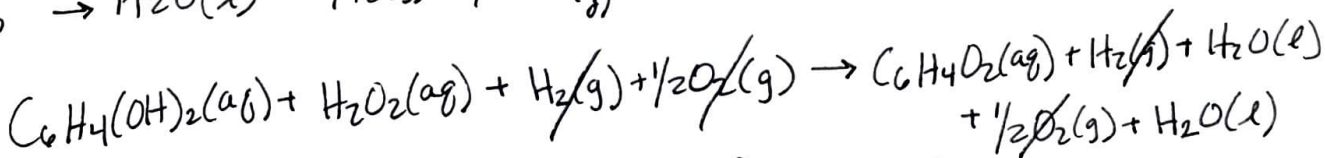
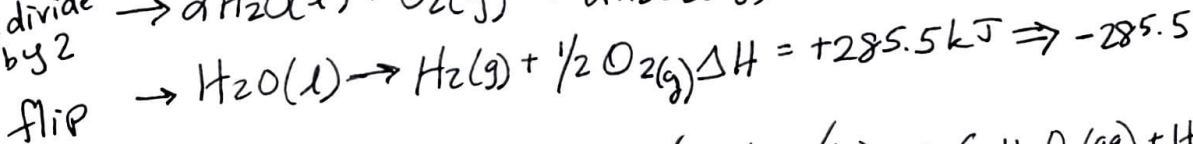
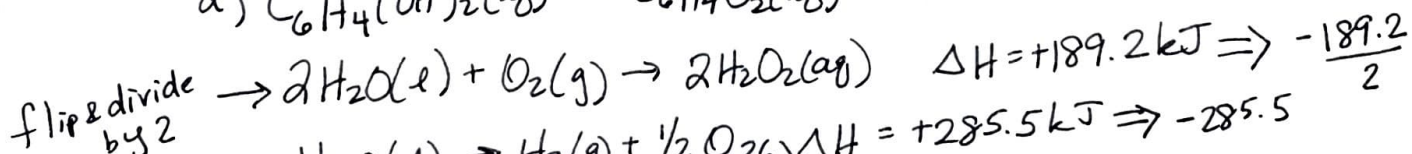
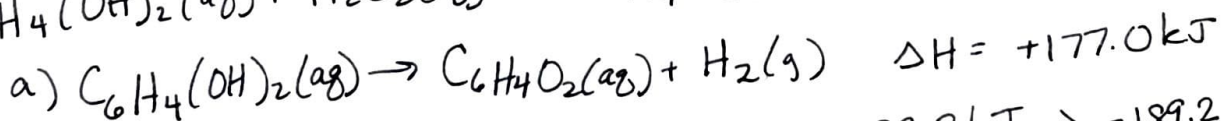
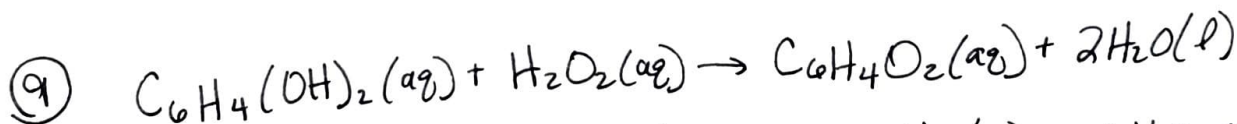
b) $\Delta H_f N_2H_4(l) : +50.6 \text{ kJ mol}^{-1}$



use ΔH_f Hess Cycle :

remember, if it goes against the arrow, flip the sign!

$$\Delta H_{rxn}^{vap} = -50.6 + 95 = \boxed{+44.4 \text{ kJ mol}^{-1}}$$



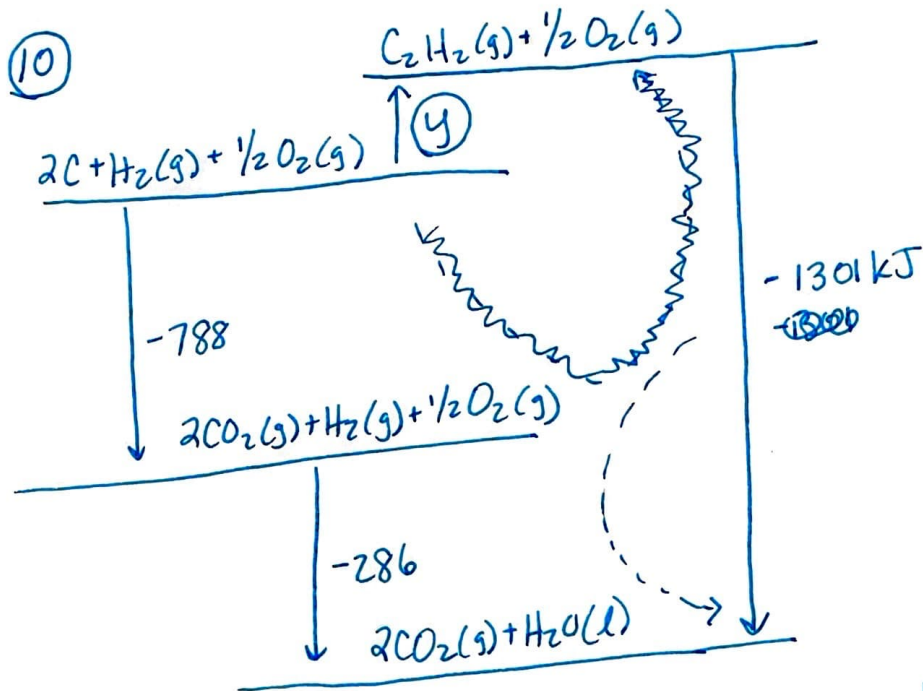
$$\Delta H_{rxn} = +177 - 94.6 - 285.5 = \boxed{-203.1 \text{ kJ}}$$

b) $q = +203.1 \text{ kJ} \quad m = 850 \text{ g} \quad T_1 = 21.8^\circ\text{C}$

$$\Delta H = -q$$

$$+203,100 \text{ J} = 850(4.184)(T_2 - 21.8)$$

$$57.11 = T_2 - 21.8 \quad \boxed{T_2 = 78.9^\circ\text{C}}$$



everything against the arrow, flip the sign!

$$y + 788 + 286 - y - 788 - 286 = -1301$$

$$-y = -1301 + 788 + 286$$

$$-y = -227$$

$$y = +227 \text{ kJ}$$

⑪ a. ΔH_{hyd} $MgSO_4$ dissolved in water (represented by ΔH_1 on the Hess cycle)

mol anhydrous $MgSO_4$: $3.0 \text{ g} \times \frac{1 \text{ mol}}{120.38 \text{ g}} = \boxed{0.0250 \text{ mol } MgSO_4}$

b. $q = mc\Delta T$
 $q = 50.0 \text{ g} (4.184) (9.7 \text{ K})$
 $= 2.029 \text{ kJ}$

$$\Delta H_1 = -2.029 \text{ kJ} / 0.0250 \text{ mol} = \boxed{-81.2 \text{ kJ mol}^{-1}}$$

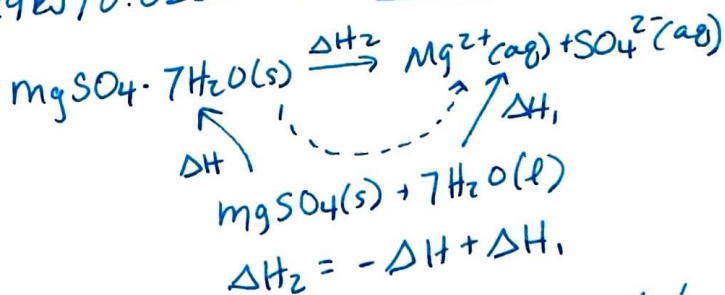
c. $\Delta H_2 = +18 \text{ kJ mol}^{-1}$

$$+18 = -\Delta H - 81.2$$

$$-\Delta H = +99.2$$

$$\boxed{\Delta H = -99.2 \text{ kJ mol}^{-1}}$$

d. % error = $\frac{-103 - (-99.2)}{-103} \times 100 = \boxed{3.7\%}$



e) loss of heat leads to a lower calculated ΔH

- incomplete rxn
- rxn not done @ STP
- absorption of heat by calorimeter