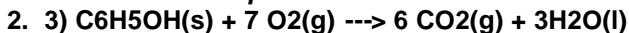


These are HIGHLY probable



When a 2.000-gram sample of pure phenol, $\text{C}_6\text{H}_5\text{OH}(s)$, is completely burned according to the equation above, 64.98 kilojoules of heat is released. Use the information in the table below to answer the questions that follow.

Substance	Standard Heat of Formation, ΔH°_f , at 25°C (kJ/mol)	Absolute Entropy, S° , at 25°C (J/mol-K)
C(graphite)	$\Delta H^\circ_f = 0.00$	$S^\circ = 5.69$
$\text{CO}_2(g)$	$\Delta H^\circ_f = -395.5$	$S^\circ = 213.6$
$\text{H}_2(g)$	$\Delta H^\circ_f = 0.00$	$S^\circ = 130.6$
$\text{H}_2\text{O}(l)$	$\Delta H^\circ_f = -285.85$	$S^\circ = 69.91$
$\text{O}_2(g)$	$\Delta H^\circ_f = 0.00$	$S^\circ = 205.0$
$\text{C}_6\text{H}_5\text{OH}(s)$	$\Delta H^\circ_f = ?$	$S^\circ = 144.0$

- (a) Calculate the molar heat of combustion of phenol in kilojoules per mole at 25°C.
- (b) Calculate the standard heat of formation, ΔH°_f , of phenol in kilojoules per mole at 25°C.
- (c) Calculate the value of the standard free-energy change, ΔG° for the combustion of phenol at 25°C.
- (d) If the volume of the combustion container is 10.0 liters, calculate the final pressure in the container when the temperature is changed to 110°C. (Assume no oxygen remains unreacted and that all products are gaseous.)

ANSWER:

(a) $2.000\text{g} \times (1 \text{ mol} / 94.113 \text{ g}) = 0.02125 \text{ mol phenol (one point)}$

Heat released per mole = $64.98\text{kJ} / 0.02125\text{mol} = 3,058 \text{ kJ/mol (one point)}$

or, $\Delta H_{\text{comb}} = -3,058 \text{ kJ/mol}$

Units not necessary

(b) $\Delta H_{\text{comb}} = -3,058 \text{ kJ/mol (one point)}$

$-3,058 \text{ kJ} = [6(-395.5) + 3(-285.85)] - [D\Delta H^\circ_f \text{ phenol}] \text{ (one point)}$

$\Delta H^\circ_f \text{ phenol} = -161 \text{ kJ (one point)}$

One point earned for correct sign of heat of combustion, one point for correct use of moles/coefficients, and one point for correct substitution

(c) $\Delta S^\circ = [3(69.91) + 6(213.6)] - [7(205.0) + 144.0] = -87.67 \text{ J/K (one point)}$

$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ = 3,058 \text{ kJ} - (298 \text{ K})(-0.08767 \text{ kJ/K}) = -3,032 \text{ kJ (one point)}$

Units not necessary; no penalty if correct except for wrong ΔH_{comb} from part (a)

(d) moles gas = $9 \times [\text{moles from part (a)}] = 9(0.02125 \text{ mol}) = 0.1913 \text{ moles gas (one point)}$

$P = (nRT) / V = [(0.193 \text{ mol})(0.0821 \text{ L atm mol}^{-1} \text{K}^{-1})(383 \text{ K})] / 10.0 \text{ L} = 0.601 \text{ atm (one point)}$

Units necessary; no penalty for using Celsius temperature if also lost point in part (c) for same error

4. Propane, C_3H_8 , is a hydrocarbon that is commonly used as fuel for cooking.

(a) Write a balanced equation for the complete combustion of propane gas, which yields $\text{CO}_2(g)$ and $\text{H}_2\text{O}(l)$

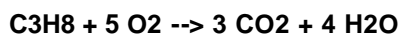
(b) Calculate the volume of air at 30°C and 1.00 atmosphere that is needed to burn completely 10.0 grams of propane. Assume that air is 21.0 percent O₂ by volume.

(c) The heat of combustion of propane is -2,220.1 kJ/mol. Calculate the heat of formation, ΔH[°]_f, of propane given that ΔH[°]_f of H₂O(l) = -285.3 kJ/mol and ΔH[°]_f of CO₂(g) = -393.5 kJ/mol.

(d) Assuming that all of the heat evolved in burning 30.0 grams of propane is transferred to 8.00 kilograms of water (specific heat = 4.18 J/g · K), calculate the increase in temperature of the water.

ANSWER:

a) one point



ignore phases (even when wrong)

multiples are OK

if balanced wrong, parts b and c should be consistent

b) four points

$$10.0 \text{ g C}_3\text{H}_8 \times (1 \text{ mol C}_3\text{H}_8 / 44.1 \text{ g C}_3\text{H}_8) = 0.227 \text{ mol C}_3\text{H}_8$$

$$0.227 \text{ mol C}_3\text{H}_8 \times (5 \text{ mol O}_2 / 1 \text{ mol C}_3\text{H}_8) = 1.13 \text{ mol O}_2$$

$$V = [(1.13 \text{ mol O}_2) (0.0821 \text{ L atm mol}^{-1} \text{ K}^{-1}) (303\text{K})] \div 1.00 \text{ atm} = 28.1 \text{ L O}_2$$

$$28.1 \text{ L O}_2 \times (100 \text{ L air} / 21.0 \text{ L O}_2) = 134 \text{ L air}$$

Note: answer must be consistent with part a

c) two points

$$[\Delta H^{\circ}_{\text{rxn}}] = [\Sigma] [\Delta H^{\circ}_{\text{f}} (\text{products})] - [\Sigma] [\Delta H^{\circ}_{\text{f}} (\text{reactants})]$$

$$-2,220.1 \text{ kJ} = [4 (-285.3\text{kJ}) + 3 (-393.5 \text{ kJ})] - [5 (0 \text{ kJ}) + [\Delta H^{\circ}_{\text{f}} (\text{C}_3\text{H}_8)]]$$

$$-2,220.1 \text{ kJ} = -1,141.2 \text{ kJ} - 1,180.5 \text{ kJ} - [\Delta H^{\circ}_{\text{f}} (\text{C}_3\text{H}_8)]$$

$$-2,220.1 \text{ kJ} = -2,321.7 \text{ kJ} - [\Delta H^{\circ}_{\text{f}} (\text{C}_3\text{H}_8)]$$

$$-101.6 \text{ kJ} = [\Delta H^{\circ}_{\text{f}} (\text{C}_3\text{H}_8)]$$

answer should be consistent with part a

1 point deducted if negative sign missing from answer

1 point deducted if -2,220.1 kJ substituted for [ΔH[°]_f (C₃H₈)]

no points earned if coefficients are inconsistent and not set equal to [ΔH[°]_f]

d) two points

$$30.0 \text{ g C}_3\text{H}_8 \times (1 \text{ mol C}_3\text{H}_8 / 44.1 \text{ g C}_3\text{H}_8) \times (2,220.1 \text{ kJ} / 1 \text{ mol C}_3\text{H}_8) = 1.51 \times 10^3 \text{ kJ}$$

$$1.51 \times 10^3 \text{ kJ} = 1.51 \times 10^6 \text{ J} = (8,000 \text{ g}) (4.18 \text{ J g}^{-1} \text{ K}^{-1}) ([\Delta T])$$

$$45.1 \text{ K (or } ^{\circ}\text{C)} = [\Delta T]$$

must correctly substitute into $q = mC_p[\Delta]T$ for 1 point
1 point earned if q value wrong but $[\Delta]T$ consistent

10. Compound ΔH°_f (kilocalories/mole) S° (calories/mole K)

H₂O (l) - $\Delta H^\circ_f = 68.3$ $S^\circ = 16.7$

CO₂ (g) - $\Delta H^\circ_f = 94.1$ $S^\circ = 51.1$

O₂ (g) $\Delta H^\circ_f = 0.0$ $S^\circ = 49.0$

C₃H₈ (g) $\Delta H^\circ_f = ?$ $S^\circ = 64.5$

When 1.000 gram of propane gas, C₃H₈, is burned at 25 °C and 1.00 atmosphere, H₂O(l) and CO₂(g) are formed with the evolution of 50.33 kilojoules.

(a) Write a balanced equation for the combustion reaction.

(b) Calculate the molar enthalpy of combustion, $\Delta H^\circ_{\text{comb}}$, of propane.

(c) Calculate the standard molar enthalpy of formation, ΔH°_f , of propane gas.

(d) Calculate the entropy change, $\Delta S^\circ_{\text{comb}}$, for the combustion reaction and account of the sign of $\Delta S^\circ_{\text{comb}}$.

GOOD QUESTION, NO ANSWER YET

These are probable

5. $2 \text{C}_4\text{H}_{10}(\text{g}) + 13 \text{O}_2(\text{g}) \rightarrow 8 \text{CO}_2(\text{g}) + 10 \text{H}_2\text{O}(\text{l})$

The reaction represented above is spontaneous at 25 °C. Assume that all reactants and products are in their standard states.

(a) Predict the sign of ΔS° for the reaction and justify your prediction.

(b) What is the sign of ΔG° for the reaction? How would the sign and magnitude of ΔG° be affected by an increase in temperature to 50 °C? Explain your answer.

(c) What must be the sign of ΔH° for the reaction at 25°C? How does the total bond energy of the reactants compare to that of the products?

(d) When the reactants are placed together in a container, no change is observed even though the reaction is known to be spontaneous. Explain this observation.

ANSWER:

a) one point

$[\Delta]S < 0$

The number of moles of gaseous products is less than the number of moles of gaseous reactant

OR

a liquid is formed from gaseous reactants.

b) one point

$[\Delta]G < 0$

ΔG becomes less negative as the temperature is increased since $\Delta S < 0$ and $\Delta G = \Delta H - T\Delta S$. The term $-T\Delta S$ adds a positive number to ΔH .

c) one point

$\Delta H < 0$

The bond energy of the reactants is less than the bond energy of the products.

d) one point

The reaction has a high activation energy

OR

is kinetically slow,

OR

a specific neutron of the needs for a catalyst or spark.

(a) State the physical significance of entropy.

(b) From each of the following pairs of substances, choose the one expected to have the greater absolute entropy. Explain your choice in each case. Assume 1 mole of each substance.

- (1) Pb(s) or C(graphite) at the same temperature and pressure
- (2) He(g) at 1 atmosphere or He(g) at 0.05 atmosphere, both at the same temperature
- (3) H₂O(l) or CH₃CH₂OH(l) at the same temperature and pressure
- (4) Mg(s) at 0 °C or Mg(s) at 150 °C, both at the same pressure.

ANSWER:

a) 2 points

Entropy (S) is a measure of randomness or disorder in a system.

b) 6 points

(4 selections at 1/2 point each and 4 accompanying explanations at 1 point each.) Selection

- (1) Pb(s) Pb has metallic bonding and is soft; atoms have large amplitude of vibration; C has atoms more localized by strong covalent bonds and is a more ordered element.
- (2) He (0.05 atm) At lower pressure (greater volume) atoms have more space in which to move.
- (3) CH₃CH₂OH(l) Ethanol is a molecule with more atoms and thus more vibrations; water has fewer atoms and is more localized by hydrogen-bonding.
- (4) Mg (150°) At higher temperature, atoms have more kinetic energy and vibrate faster and further, i.e., greater randomness occurs.

In part (a), partial credit of 1 point was allowed for confusing S with ΔS , but writing correctly in terms of "change in randomness" or "tendency toward greater randomness." In the explanations of part (b), 1/2 point was credited for correct empirical relationships involving S as a function of P or T, or mentioning relative melting-points in parts (1), (3), and (4).

9. $\text{PCl}_5(\text{g}) \rightarrow \text{PCl}_3(\text{g}) + \text{Cl}_2(\text{g})$

For the reaction above, $\Delta H^\circ = +22.1$ kilocalories per mole at 25 °C.

(a) Does the tendency of reactions to proceed to a state of minimum energy favor the formation of the products of this reaction? Explain.

(b) Does the tendency of reactions to proceed to a state of maximum entropy favor the formation of the products of this reaction? Explain.

(c) State whether an increase in temperature drives this reaction to the right, to the left, or has no effect. Explain.

(d) State whether a decrease in the volume of the system at constant temperature drives this reaction to the right, to the left, or has no effect. Explain.

ANSWER:

Four effects and 4 explanations valued at 1 point each.

a) No, the reactant is favored. Reaction is endothermic and products are at higher energy than the reactant.

b) Yes, products are favored. $\Delta S > 0$ as 2 moles of gas are produced from 1 mole of gas.

c) Right, products are favored. For the system at constant pressure: absorption of heat favors the products. An argument using LeChatelier's principle can be used with heat considered as a reactant; or $T\Delta S$ favors products as ΔG becomes more negative. For the system at constant volume: products are favored but increase in the yield of products results in an increase in pressure which would drive the reaction to the left.

No points were deducted for students who mentioned both the thermodynamic effect and the counteracting pressure effect at constant volume but were uncertain as to which effect is larger. Students who considered a system at constant volume and stated only that the increase in pressure would drive the reaction to the left received half credit.

d) Left, the reactant is favored. A decrease in the volume would increase the pressure and the strain is relieved by the reverse reaction which produces 1 mole of gas from 2 moles of gas.
