

6 • Thermochemistry**PRACTICE TEST**

1. A system has an increase in internal energy, ΔE , of 40 kJ. If 20 kJ of work, w , is done on the system, what is the heat change, q ?
- a) +60 kJ d) -20 kJ
 b) +40 kJ e) -60 kJ
 (c) +20 kJ $\Delta E = q + w$
 $40 = q + 20$
2. A gas at 20 atm pressure with a volume of 2.0 Liters expands against a 5 atm pressure to a volume of 8.0 Liters. How much work is done by the gas? $W = -P\Delta V = (-5\text{atm})(8.0 - 2.0)\text{L}$
- (a) 30 L·atm c) 8 L·atm = -30 atm·L
 b) 18 L·atm d) 5 L·atm
3. Which equation represents the heat of formation, ΔH_f° , for MgCl_2 ?
- a) $\text{Mg}^{2+}(\text{aq}) + 2 \text{Cl}^- \rightarrow \text{MgCl}_2(\text{s})$
 b) $\text{Mg}(\text{s}) + 2 \text{Cl}(\text{g}) \rightarrow \text{MgCl}_2(\text{s})$
 c) $\text{MgCl}_2(\text{s}) \rightarrow \text{Mg}^{2+}(\text{aq}) + 2 \text{Cl}^-(\text{aq})$
 (d) $\text{Mg}(\text{s}) + \text{Cl}_2(\text{g}) \rightarrow \text{MgCl}_2(\text{s})$
4. Take a toy balloon. Quickly stretch the balloon and press it against your lower lip. What is the ΔH for the reaction:
- unstretched \rightarrow stretched**
- a) + (c) -
 b) 0 d) impossible to tell
5. Which of the following is NOT a state function?
- a) pressure c) temperature
 b) volume (d) none of these
6. The correct units for specific heat capacity:
- a) J°C (c) $\text{J/g}^\circ\text{C}$
 b) J/g d) $^\circ\text{C/g}$
7. How much heat is required to convert solid sulfur to gaseous sulfur at 298 K and 1 atm pressure? ΔH° (kJ/mol)
- $\text{S}(\text{s}) + \text{O}_2(\text{g}) \rightarrow \text{SO}_2(\text{g}) \quad -395$
 $(\text{S}(\text{g}) + \text{O}_2(\text{g}) \rightarrow \text{SO}_2(\text{g}) \quad -618) \times -1$
- a) -1013 kJ/mol (c) +223 kJ/mol
 b) -223 kJ/mol d) +618 kJ/mol
 $-395 + 618 = 223$
8. Using the ΔH_f° given below, calculate the $\Delta H_{\text{combustion}}$ for propane, C_3H_8 .
- ΔH_f° (kJ/mol)
- $\text{H}_2\text{O}(\text{l}) \quad 4(-286)$
 $\text{CO}_2(\text{g}) \quad 3(-394)$
 $\text{C}_3\text{H}_8(\text{g}) \quad -1(-104)$
- a) 576 kJ (c) -2222 kJ
 b) -576 kJ d) -2330 kJ
 $\text{C}_3\text{H}_8 + 5\text{O}_2 \rightarrow 3\text{CO}_2 + 4\text{H}_2\text{O}$
9. The heat of vaporization of methane, CH_4 , at its boiling point is 9.20 kJ/mol. How much heat energy is required to vaporize 100. g of methane at its boiling point?
- a) 1380 kJ c) 21.6 kJ
 b) 86.3 kJ (d) 57.4 kJ
 $100.\text{g CH}_4 \times \frac{1\text{mol}}{16.042\text{g}} \times \frac{9.20\text{kJ}}{\text{mol}} = 57.3\text{kJ}$
10. How much energy is required to melt 10.0 g benzene, C_6H_6 ? The heat of fusion of benzene is 2.37 kJ/mol.
- a) 3.30 kJ c) 1850 kJ
 b) 23.7 kJ (d) 0.303 kJ
 $10.0\text{g C}_6\text{H}_6 \times \frac{1\text{mol C}_6\text{H}_6}{78.108\text{g}} \times \frac{2.37\text{kJ}}{\text{mol}} = .303\text{kJ}$

11. If ΔH for a reaction is positive, ... *- add energy*
- the reaction rate is generally very fast.
 - ~~the~~ the enthalpy change of the reverse reaction is positive. *negative*
 - the enthalpy of the products is greater than the enthalpy of the reactants. *(go uphill)*
 - the energy released during bond formation ⁽⁻⁾ is greater than the energy absorbed ⁽⁺⁾ during bonding breaking for the reaction. \therefore If $(+) > (+)$, then $(-)$

Answers:

- | | | |
|------|-------|-------|
| 1. C | 6. C | 11. C |
| 2. A | 7. C | 12. B |
| 3. D | 8. C | 13. C |
| 4. C | 9. D | 14. B |
| 5. D | 10. D | |

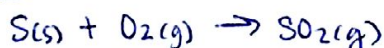
Notes:

- an increase in internal energy means an increase in P.E. of system by 40 kJ
work done ON system increases P.E., +20 kJ, so $q = +20$ kJ, too.
- $w = -P\Delta V = 5 \text{ atm} \times (8-2 \text{ L})$... the 20 atm is not used for anything.
- balloon gets warm, $\Delta H < 0$
- each of these only depends on the STATE of the substance, not on its HISTORY.
- reverse second reaction
- recall: $\text{C}_3\text{H}_8 + 5\text{O}_2 \rightarrow 3\text{CO}_2 + 4\text{H}_2\text{O}$ and use Hess's Law
- Given: 100g CH_4 , use molar mass & H_{vap} as conversion factors.
- Given: 10.0g C_6H_6 , use molar mass & H_{fus} as conversion factors.
- this is an "uphill" reaction.
- take half of first equation, reverse second equation. if you reverse and double second equation, you get TWICE the answer.
- answer = heat capacity $\times \Delta T$... you don't use 0.100 g anywhere. You would IF the question asked for MOLAR heat of combustion.
- if volume is constant, $P\Delta V$ work = 0 so $\Delta E = q + w$ becomes $\Delta E = q$.

12. Given the two equations:
- $$\frac{1}{2}(2 \text{ S(s)} + 3 \text{ O}_2(\text{g}) \rightarrow 2 \text{ SO}_3(\text{g})) \Delta H^\circ = (-790.4 \text{ kJ}) \times \frac{1}{2}$$
- $$-1 (\text{SO}_2(\text{g}) + \frac{1}{2} \text{ O}_2(\text{g}) \rightarrow \text{SO}_3(\text{g})) \Delta H^\circ = (-99.1 \text{ kJ}) \times -1$$

What is the standard enthalpy of formation for sulfur dioxide, $\text{SO}_2(\text{g})$?

- +99.1 kJ
- 296.1 kJ
- 592.2 kJ
- 839.5 kJ



13. When 0.100 g benzoic acid ($\text{HC}_6\text{H}_4\text{CO}_2$) and excess oxygen is ignited in a bomb calorimeter, the temperature of the water changes from 25.000°C to 25.225°C. The heat capacity of the calorimeter is 603 J/°C.

What is the ΔE for this reaction? $\Delta T = .225^\circ\text{C}$

- 597 J
 - 1660 J
 - 136 J
 - 149 J
- $q = \frac{603 \text{ J}}{^\circ\text{C}} \times .225^\circ\text{C} = 136 \text{ J}$*

14. Under conditions of constant volume, the heat change that occurs during a chemical reaction is equal to

- ΔH
- ΔE
- ΔT
- ΔP

$$\begin{array}{l} \Delta H = \Delta E + P\Delta V \\ \Delta H = \Delta E + P(0) \\ \Delta H = \Delta E \end{array} \quad \left| \quad \begin{array}{l} \text{OR} \\ \Delta E = q + w \\ \Delta E = q + 0 \\ \Delta E = q \end{array} \right.$$